

The Alloy-Theoretic Automated Toolkit (ATAT): A User Guide

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Chapter 1

Features/Capabilities

The Alloy-Theoretic Automated Toolkit (ATAT) is a generic name that refers to a collection of alloy theory tools:

- Codes to construct cluster expansions from first-principles (maps and mmap). A cluster expansion is a very compact and efficient expression giving the energy of an substitutional alloy as a function of its configuration (i.e. which type of atom sits where on the lattice).
- Codes to perform Monte Carlo simulation (emc2 and memc2) of lattice models in order to compute thermodynamic properties of alloys, starting from a cluster expansion.
- Codes to perform lattice dynamics calculations (fitfc, fitsvsl, svsl)
- Codes to calculate electronic and magnetic free energy contributions (felec, fmag, fempmag) using simple physical or semiempirical models.
- Utilities to combine all of the above to generate free energies that include configurational, vibrational and electronic contributions (mkteci).
- Codes to generate Special Quasirandom Structures (SQS), to model disordered solid solutions (mcsqs, gensqs) and to enumerate structures (genstr).
- A large library of pre-computed SQS and structure prototypes.
- Extension of the two above tools that allow the construction of so-called reciprocal-space cluster expansion, which are useful to model the energetics of alloys exhibiting a large atomic size mismatch.
- Tensorial cluster expansions (gce).
- Elastic constant calculations (calcelas).
- Structure conversion utilities (subcells, supercells, coordinate system changes, file format, etc.) (cell-cvrt, wycked, etc.)
- Scripts to automate tasks (foreachfile, sspp, getvalue, getlines, etc.).
- Codes to generate CALPHAD databases (sqs2tdb).
- Utilities to interface the above tools with first-principles codes, such as VASP (runstruct_vasp, runstruct_abinit, runstruct_gulp, etc.).
- Job control utilities that enable the efficient use of a cluster of workstations to run the first-principles codes that provide the input to the above codes (pollmach).

Chapter 2

Credits and Licence

The Alloy-Theoretic Automated Toolkit (ATAT)¹ is a generic name that refers to a collection of alloy theory tools developed by Axel van de Walle², in collaboration with various research groups.

2.1 Collaborators and Credits

The MAPS³ (MIT Ab-initio Phase Stability) code, which automatically constructs a cluster expansion from the result of first-principles calculations, was developed by Axel van de Walle in collaboration with Prof. Gerd Ceder's group⁴ from the Department of Materials Science and Engineering at the Massachusetts Institute of Technology. MAPS consists of the following codes: `maps`, `corrump`, `genstr`, `checkcell`, `kmesh`, `cv`.

The EMC2 (Easy Monte Carlo Code), which automate the calculation of alloy thermodynamic properties via Monte Carlo simulations of lattice models, were developed by Axel van de Walle in collaboration with Prof. Mark Asta's group⁵ from the Department of Materials Science and Engineering at Northwestern University. EMC2 consists of the following codes: `emc2`, `phb`.

The CSE (Constituent Strain Extension) to both the MAPS and EMC2 codes, which implement the constituent strain formalism based on a reciprocal-space cluster expansion, was developed by Axel van de Walle in collaboration with Alex Zunger's Solid State Theory Group⁶ at the National Renewable Energy Laboratory in Golden, Colorado and in collaboration with Gus Hart⁷ from the Department of Physics and Astronomy at Northern Arizona University. CSE consists of the following files: `csfit.cc`, `predcs.cc`, `predrs.cc`, `kspacccs.cc`.

Mayeul D'Avezac at NREL has provided all the changes needed for ATAT to compile with g++ versions 4.1 and later as well as intel's c++ compiler.

Volker Blum at NREL has contributed to improve the portability of the package by providing a `perl` version of the `ch1` utility.

Dongwon Shin at Penn State has converted a large number of common lattices (found at the NRL navy web site⁸) into the `atat` format. See directory `data/str`.

Greg Pomrehn has improved the efficiency of the structure enumeration algorithm and contributed the script `mmapsrep`.

The FFT routines in the files `fftn.cc` and `fftn.h` were obtained from the NIST Guide to Available Math Software (GAMS). The origin of these routines dates back to a FORTRAN code by R. C. Singleton in 1968, later converted to C and subsequently improved by Mark Olesen and John Beale in 1995. These routines are

¹<http://www.its.caltech.edu/~avdw/atat/>

²<http://www.mit.edu/~avdw/>

³<http://www.mit.edu/~avdw/maps>

⁴<http://burgaz.mit.edu/>

⁵<http://cms.northwestern.edu/>

⁶<http://www.sst.nrel.gov/>

⁷<http://www.phy.nau.edu/~hart/>

⁸<http://cst-www.nrl.navy.mil/lattice/>

included in the `atat` package for sole purpose of providing users with the convenience of avoiding a separate download. Axel van de Walle does not claim any ownership of them or intellectual credit for them.

Some of the basic numerical routines were inspired by Numerical Recipes in C, although they were completely re-written according to the ATAT code style conventions, for instance to use 0-indexed arrays and make better use of C++ features.

Gao Zhe has contributed an interface to PWSCF (aka Quantum Espresso) that can be found in `atat/glue/qe`
 Matt Probert has contributed an interface to CASTEP that can be found in `atat/glue/castep/`

2.2 Financial Support

The development of MAPS was supported by the U.S. Department of Energy, Office of Basic Energy Sciences, under contract no. DE-F502-96ER 45571. Gerbrand Ceder acknowledges support of Union Minière through a Faculty Development Chair. Axel van de Walle acknowledges support of the National Science Foundation under program DMR-0080766 and DMR-0076097 during his stay at Northwestern University.

The development of EMC2 was supported by the NSF under program DMR-0080766.

The development of the CSE is supported by the NSF under program DMR-0080766.

The development of the tensorial cluster expansion (GCE and `gencs` code) is supported by the Center for the Science and Engineering of Materials at Caltech, an NSF-funded MRSEC and by the NSF CMMT grant DMR-0907669.

The development of the `wycked` and `getproto` codes is supported by the NSF CAREER grant DMR-1154895.

The development of the `mcsqs` code is supported by ONR grants N00014-11-1-0886 and N00014-11-1-0886.

2.3 Licence and Agreements

Any scientific article or book whose results were obtained with the codes described above must properly acknowledge their use by citing the following papers:

1. A. van de Walle and G. Ceder, “Automating First-Principles Phase Diagram Calculations⁹”, *Journal of Phase Equilibria*, **23**, 348, (2002).
2. A. van de Walle and M. Asta, “Self-driven lattice-model Monte Carlo simulations of alloy thermodynamic properties and phase diagrams¹⁰”, *Modelling Simul. Mater. Sci. Eng.* **10**, 521, (2002).
3. A. van de Walle, M. Asta and G. Ceder, “The Alloy Theoretic Automated Toolkit: A User Guide¹¹” *CALPHAD Journal*, **26**, 539, (2002).
4. A. van de Walle, “Multicomponent multisublattice alloys, nonconfigurational entropy and other additions to the Alloy Theoretic Automated Toolkit¹²”, *Calphad Journal* **33**, 266, (2009).
5. As of version 2.81, the algorithm to generate superstructures has been improved based on the ideas in G. L. W. Hart and R. W. Forcade, “Algorithm for generating derivative structures¹³,” *Phys. Rev. B* **77**, 224115, (2008).
6. If the `mcsqs` code is used: A. van de Walle and P. Tiwary and M. M. de Jong and D. L. Olmsted and M. D. Asta and A. Dick and D. Shin and Y. Wang and L.-Q. Chen and Z.-K. Liu, Efficient stochastic generation of Special Quasirandom Structures¹⁴, *Calphad Journal* **42**, 13 (2013).
7. If the constituent strain extension is used: D. B. Laks and L. G. Ferreira and S. Froyen and A. Zunger, *Phys. Rev. B* **46**, p. 12587 (1992).

⁹<http://arXiv.org/abs/cond-mat/0201511>

¹⁰<http://arXiv.org/abs/cond-mat/0201473>

¹¹<http://arxiv.org/abs/cond-mat/0212159>

¹²<http://arxiv.org/abs/0906.1608>

¹³<http://link.aps.org/abstract/PRB/v77/e224115>

¹⁴<http://dx.doi.org/10.1016/j.calphad.2013.06.006>

8. The **wycked** code uses the Wyckoff position database from the Bilbao Crystallographic Server¹⁵. Accordingly, if you use this code, you should also cite the following papers:

M. I. Aroyo, J. M. Perez-Mato, D. Orobengoa, E. Tasci, G. de la Flor, A. Kirov, “Crystallography online: Bilbao Crystallographic Server”, *Bulg. Chem. Commun.* **43** 183 (2011).

M. I. Aroyo, J. M. Perez-Mato, C. Capillas, E. Kroumova, S. Ivantchev, G. Madariaga, A. Kirov and H. Wondratschek, “Bilbao Crystallographic Server I: Databases and crystallographic computing programs”, *Z. Krist.* **221**, 15 (2006).

M. I. Aroyo, A. Kirov, C. Capillas, J. M. Perez-Mato and H. Wondratschek, “Bilbao Crystallographic Server II: Representations of crystallographic point groups and space groups”, *Acta Cryst.* **A62**, 115 (2006).

9. The **getproto** code uses processed data from the index of the Landolt-Bornstein database¹⁶. This data is available upon request by contacting avdw@alum.mit.edu¹⁷ if you can show that you have access to this database. Also, if using this data, please cite: Villars, P., Cenzual, K., Daams, J., Gladyshevskii, R., Shcherban, O., Dubensky, V., Melnichenko-Koblyuk, N., Pavlyuk, O., Stoiko, S., Sysa, L., “Landolt-Bornstein — Group III Condensed Matter: Numerical Data and Functional Relationships in Science and Technology”, Edited by Villars, P. and Cenzual, K., SpringerMaterials — The Landolt-Bornstein Database, Volume 43A1-43A10 (2013).

The files included in this distribution cannot be further distributed either in their original or in a modified form without consent of the author, Axel van de Walle (avdw@alum.mit.edu).

Users are free to modify the code solely for their personal use and are encouraged to share their improvements with the author at (avdw@alum.mit.edu¹⁸). Their contributions will be acknowledged in the present section, in future versions of this manual.

2.4 Beta testers

The following researchers have provided numerous constructive comments that have proven extremely useful to improve the quality of the program.

1. Dane Morgan (University of Wisconsin)
2. Paul Dalach (Northwestern)
3. Dinesh Balachandran (MIT, Gerd Ceder’s group)
4. Ben Burton (National Institute of Standards and Technology)
5. Gautam Ghosh (Northwestern University)
6. Nikolai Andreevich Zarkevich (University of Illinois at Urbana-Champaign, Duane Johnson’s group)
7. Volker Blum (NREL, Alex Zunger’s group)
8. Chris Woodward (Northwestern University/Air Force)
9. Zhe Liu (Northwestern University, Mark Asta’s group)
10. Yi Wang and Raymundo Arroyave (Pennsylvania State University, Long-Qing Chen and Zi-Kui Liu’s group)
11. Elif Ertekin (University of California at Berkeley, Daryl Chrzan’s group)
12. Rodrigo Barbosa Capaz (University of California at Berkeley)
13. Sundar Amancherla (General Electric)

¹⁵<http://www.cryst.ehu.es/>

¹⁶<http://www.springermaterials.com/>

¹⁷<mailto:avdw@alum.mit.edu>

¹⁸<mailto:avdw@alum.mit.edu>

Chapter 3

Getting started

3.0.1 Requirements

You need the following utilities installed:

- **g++** version 2.7.2 or later. Type `g++ --version` to verify this. This package can be downloaded from <http://www.gnu.org/>.
- GNU **make** (any version). Type `make --version` to verify this. On some systems this command may be called `gmake` or `gnumake`. This package can be downloaded from <http://www.gnu.org/>.
- A first-principle electronic structure calculation code, such as VASP¹
- You may want to use **gnuplot** to plot the output of the code. Type `gnuplot` and check the program starts (Type `q` to quit). If not, it can be downloaded from <http://www.gnuplot.info/>.
- You may need **ssh** if you have multiple machines and they are connected through an unsecure network (e.g. the internet). This package can be downloaded from <http://www.openssh.com/>.

3.0.2 Installation

If you have an earlier version of **ATAT** installed, please delete or rename the former `atat` directory before proceeding, e.g.,

```
mv atat atattold
```

Then, type

```
gunzip atatX.XX.tar.gz
```

```
tar -xvf atatX.XX.tar.gz
```

where `X.XX` is the current version number. These commands create a directory called `atat` in the current directory. It contains the whole package. For future reference, I'll call the whole access path to this directory `atat`.

Type

```
cd atat
```

and open the file `makefile` with a text editor and look for the line `BINDIR=$(HOME)/bin/`. Change `$(HOME)/bin/` to point wherever you want to put the executables. Type

```
make
```

If no error message appears, proceed with the next steps, otherwise consult Chapter 8.

```
make install
```

```
rehash (not necessary with bash shell)
```

¹<http://cms.mpi.univie.ac.at/vasp/>

3.0.3 Test MAPS with a simple example

Change to a directory of your choice (preferably empty) and type

```
cp atat/examples/cuau.in lat.in
maps -d &
```

Maps is running and waiting for a signal. Type

```
touch ready
```

to indicate that you are ready to for maps to generate a structure. Maps replies **Finding best structure...** To find the structure just created, wait for **done** to appear and type:

```
ls */wait
```

to observe that directory **0** has been created. This directory contains a file **str.out** which describes the structure whose energy needs to be calculated. The file **wait** is just a flag that allows you to find the newly created directory. Let's pretend that we have computed the energy of that structure. We need to let maps know about it. Type, for instance:

```
echo 1.1 > 0/energy (If 1.1 is the energy of the structure.)
rm 0/wait
```

Maps responds by **Finding best cluster expansion...**, followed by **done**. You can repeat the process (**touch ready**, etc.) to add more structures. Maps will update the current cluster expansion every time a new energy becomes known. (By default, maps checks every 10 sec.). For a description of the output files, type:

```
maps -h | more
```

or refer to section 7 A nice utility called **mapsrep** allows you to plot the results using gnuplot. To stop maps cleanly, type: **touch stop**

Suggestion: to clarify the output of the program, it is recommended that you run **maps** in one terminal window and type all other command in another terminal window.

3.1 Install the interface between MAPS and VASP

Type

```
ezvasp
```

and follow the instructions posted on the screen to configure this command.

To test this interface, change to a directory of your choice and type

```
maps -d &
```

(unless maps is running already in that same directory).

While MAPS automatically create files that describe the geometry of the structures (called **n/str.out**, where **n** is the structure name), we need to provide a file containing all the other parameters needed by the first-principles code. Type:

```
cp atat/glue/vasp/vasp.wrap .
```

to copy an example of such file in the current directory. For a description of these parameters, type:

```
ezvasp -h | more
```

Let's say you have a new structure in directory **0** (created by typing **touch ready**). Type:

```
cd 0
runstruct_vasp
```

When the command has terminated, the directory 0 will contain a file **energy** giving the energy of the structure. If error messages appear consult Chapter 8.

If no error messages appear, you can proceed another level up in automation. Type

```
cd .. (to go back into the main directory)
pollmach runstruct_vasp &
```

This script will automatically call the above command repeatedly. To stop it cleanly, type:

```
touch stoppoll
```

(Disregard the warning message.) If you only have access to one machine, this is as good as it gets, if you have more than one machine, read the next section. If you want to use another code than VASP, read section 3.3.

3.2 Install and test the job control utilities

This section requires some knowledge of UNIX, but it is worth it!

3.2.1 With queueing systems

Let's start with the simple and common case of a computer cluster with a queueing system. We don't give here the specific syntax of the job submission script, because it is highly system-dependent. In a nutshell, one would submit on job containing the following commands, assuming the system is using a standard implementation of MPI:

```
[some header specifying the number of nodes etc.]
maps -d & (note the "&" to run in the background.)

pollmach runstruct_vasp mpirun (you may need to specify the number of processors as mpirun -np
[number])
```

If your system does not allow a background commands in the script (or if this would be very inefficient), then you could put each command (without "&") in a separate script, but making sure they run at the same time. Either way, parallelization occurs at the level of the **vasp** command. If you wish to also parallelize by running multiple **vasp** commands simultaneously then you should submit yet another job script with

```
[some header specifying the number of nodes etc.]

pollmach runstruct_vasp mpirun
```

repeating this process as many times as you the number of simultaneous **vasp** processes you want (note that there is still only one copy of **maps** running).

3.2.2 Without queueing systems

We now cover the case where there is no queueing system and the user has full power to run on any of the nodes in the cluster.

Networking problems can be tricky and we will test various thing as we go along. You only need to perform this installation on your "master" machine that will run **maps**. All other machine (which we be called the "remote" machines) only need to have VASP (or any other ab-initio code).

Before you start, you must first make sure that it is possible to login from the master machine to the remote machines without entering a password. This is essential for the program to be able to run on its

own, without your intervention. Don't worry it is generally possible to do this without compromising the security of your system. Two commands allow you to run a command on a remote machine. If the master and remote machines are connected through a secure network (e.g. a beowolf cluster having its own local network) or if you don't care about security (for now), you can use **rsh**. Otherwise, **ssh** provides a secure way to command a machine remotely.

To set up **rsh** so that you can login without typing a password, you must have the appropriate **.rhosts** file on the remote machine. For more information, consult the **rsh** man page. (One important issue, often not mentioned in the man pages, is that you need to set the file permissions of the **.rhosts** file so that no one else but you has "write" permission: **chmod og-w ~/.rhosts**).

To set up **ssh** so that you can login without typing a password, consult the **ssh** man page, especially the section on "RSA-based host authentication". (This is the feature that makes the login secure even if no password is needed.) In general, setting this up involves creating a **.shosts** file and generating a public key files to be copied onto the remote machines.

If your username is different on the remote machine and if you use **ssh**, use the syntax **node username@host** instead. If you use **rsh**, use the syntax **node -l username host**.

Once you are able to run either **rsh node2 ls** or **ssh node2 ls** and get the content of your home directory on the remote machine (assuming that you have a remote machine called **node2**), you can proceed to the next step. Do you have the same home directory on the master and remote machines and does it have the same access path? To check this, **cd** to some arbitrary subdirectory and type:

```
node node2 ls
```

where **node** is a command provided with ATAT and **node2** is the name of the remote machine. If you want to use **ssh** instead, type

```
node -s node2 ls
```

This should print the content of the *current* directory on the master machine (not your home directory). If you get an error message or if you get the content of another directory, you will need to check if the following works. Make sure you are in a directory that does not contain too many files. If you want to use **rsh**, type:

```
node -r node2 ls
```

while if you want to use **ssh**

```
node -s -r node2 ls
```

In either case, you should get the content of the current directory before continuing on. The **node -r** command works by copying the content of the current directory on the remote machine in a temporary directory. Once the command has terminated, the new content is copied back and the remote temporary directory is deleted.

We are now ready to automate the calculations. Type

```
chl
```

and, as indicated on the screen, open the file **~/machines.rc** with a text editor. This file contains numerous comment lines explaining the format of the file and a few examples.

The commands in the first column (before the **+**) must print a single number indicating the load of the machine. It is a good check to copy and paste each of these command, one at the time, into a shell window to see if the output is a single number. In order to extract a single number out of a complicated output, the command **getvalue** is provided. It extracts the single number following the token given as an argument. The first entry, with the **none** keyword after the **+** indicates the threshold load above which a machine is considered too busy to be usable. Note that the load checking commands may quite elaborate if, for instance, you need to "rescale" the load of some machines because they have a different reporting scheme or if you want to tweak the priority given to each machine.

The second column (after the **+**) give the command prefix needed to run on each remote machine. These prefix will usually consist of the command **node** described above. It is very important that the command prefix be such that the current directory in the remote machine when the command is run is the same as on

the local machine. The best way to test that is to try the prefix in front of the `ls` command and verify that what is printed is indeed the content of the current local directory.

Once you are done with entering the information for each of your machines (you can also enter only a few and come back later to add the remaining ones), make sure to comment out the examples provided (placing a `#` at the beginning of the unwanted line). Do not comment out the first line which contains the `none` keyword.

Once you have edited the `~/.machines.rc` file, type `chl`. This should give a list of the load of all machines in the first column and a list of command prefix in the second. Next, try the command `minload`. It should give the command prefix that let you access the machine with the minimum load or `none` if no machine is available. To check, once again, that the command prefix are correct, type `'minload' ls` (make sure you use backward quotes!). This should print the content of the current directory (unless there are no machines available).

This approach could also be used with a queueing system, but this is much more advanced and requires good knowledge of scripting languages. You could read the “node list file” the queueing system assigned to the job and create a local `.machine.rc` file on-the-fly. The `-s` option of `pollmach` could prove helpful to ensure two vasp runs do run on the same processors, and avoids the need for load balancing code.

3.3 Interfacing MAPS with other first-principles codes

You need to provide a command (e.g. a shell script) called `runstruct_xxx`, where `xxx` is any name of your choice. This script should read, from the current directory, the file `str.out` describing the geometry of the structure and create the appropriate input files for the first-principles code. It should then execute the command(s) needed to run the code. If a multiple machine environment is used, the script should use the first argument passed to the script (`$1`) as command prefix to put in front of any command in order for them to be run on a remote machine. That is, if the first-principle code is called “myfp” the script should execute

```
$1 myfp
```

Once the first-principles code has terminated, the script should

- Create a file called **energy** containing the energy of the structure per unit cell of the structure (not the lattice) (this is what first-principles code usually give anyways).
- If the calculation fails, no **energy** file should not be created. Instead, an empty file called **error** should be created.

The above files must all reside in the current directory (from where the script was invoked). To follow the philosophy of the package, the additional input parameters (besides the structure geometry) needed by the first-principles code should be contained in a file called `xxx.wrap` located one (or two) levels up in the directory hierarchy, relative to the current directory.

As a starting point to write this script, have a look at the file `atat/glue/vasp/runstruct.vasp`.

Chapter 4

Version History and Bugs

1.50:

constituent strain
allow any structure names
allow new structures to be added at all times

1.66:

unit cell in 'symmetric' form, ready for ab-initio codes
constituent strain fitting code
temperature-dependent ECI in Monte Carlo code
manual in tex/html
reciprocal space Monte-Carlo (BETA)
easy-to-configure job control scripts

2.03:

vibrational and electronic entropy included.

2.50:

mixed canonical/grandcanonical multicomponent monte carlo

2.53:

reciprocal space multicomponent monte carlo (for electrostatics)

2.54:

automated patching system for c++ language qwirks

2.70:

tensorial/generalized cluster expansion (gce utility)

2.71:

fixed bug in multisublattice structure enumeration routine
(only affects cases where a pure translation maps one sublattice onto another:
omits some structures with lattice vector equal to one of the unit cell lattice vectors
cluster expansions not affected, but perhaps missed ground states or SQS, but unlikely)

Chapter 5

User guide

5.1 Introduction

First-principles calculations of alloy thermodynamic properties have been successfully employed in a variety of contexts for metallic, semi-conductor and ceramic systems, including the computation of: composition-temperature phase diagrams, thermodynamic properties of stable and metastable phases, short-range order in solid solutions, thermodynamic properties of planar defects (including surfaces or antiphase and interphase boundaries), and the morphology of precipitate microstructures [6, 5, 27, 28, 25, 3, 1].

Although the formalism that allows the calculation of thermodynamic properties from first principles has been known for decades [6, 5, 27], its practical implementation remains tedious. These practical issues limit the accuracy researchers are able to obtain without spending an unreasonable amount of their time writing input files for various computer codes, monitoring their execution and processing their output. These practical difficulties also limit the community of researchers that use these methods solely to those that possess the necessary expertise to carry out such calculations.

The Alloy Theoretic Automated Toolkit (ATAT) [22] drastically simplifies the practical use of these methods by implementing decision rules based on formal statistical analysis that free the researchers from a constant monitoring during the calculation process and automatically “glues” together the input and the output of various codes, in order to provide a high-level interface to the calculation of thermodynamic properties from first principles. In order to make this powerful toolkit available to the wide community of researchers who could benefit from it, this article presents a concise user guide to this toolkit.

5.2 Theoretical Background

While there exist numerous methodologies that enable the calculation of thermodynamic properties from first principles, we will focus on the following two-step approach (see Figure 5.1). First, a compact representation of the energetics of an alloy, known as the cluster expansion [18, 6, 5, 27], is constructed using first-principles calculations of the formation energies of various atomic arrangements. Second, the cluster expansion is used as a Hamiltonian for Monte Carlo simulations [16, 2, 8, 15] that can provide the thermodynamic properties of interest, such as the free energy of a phase or short-range-order parameters as a function of temperature and concentration. This two-step approach is essential, because the calculation of thermodynamic quantities through Monte Carlo involves averaging the property of interest over many different atomic configurations and it would be infeasible to calculate the energy of each of these configurations from first principles. The cluster expansion enables the prediction of the energy of any configuration from the knowledge of the energies of a small number of configurations (typically between 30 and 50), thus making the procedure amenable to the use of first-principles methods.

Formally, the cluster expansion is defined by first assigning occupation variables σ_i to each site i of the *parent lattice*, which is defined as the set of all the atomic sites that can be occupied by one of a few possible atomic species. In the common case of a binary alloy system, the occupation variables σ_i take the value -1 or $+1$ depending on the type of atom occupying the site. A particular arrangement of these “spins” on

the parent lattice is called a *configuration* and can be represented by a vector σ containing the value of the occupation variable for each site in the parent lattice. Although we focus here on the case of binary alloys, this framework can be extended to arbitrary multicomponent alloys (the appropriate formalism is presented in [18]).

The cluster expansion then parametrizes the energy (per atom) of the alloy as a polynomial in the occupation variables:

$$E(\sigma) = \sum_{\alpha} m_{\alpha} J_{\alpha} \left\langle \prod_{i \in \alpha'} \sigma_i \right\rangle \quad (5.1)$$

where α is a cluster (a set of sites i). The sum is taken over all clusters α that are not equivalent by a symmetry operation of the space group of the parent lattice, while the average is taken over all clusters α' that are equivalent to α by symmetry. The coefficients J_{α} in this expansion embody the information regarding the energetics of the alloy and are called the effective cluster interaction (ECI). The *multiplicities* m_{α} indicate the number of clusters that are equivalent by symmetry to α (divided by the number of lattice sites).

It can be shown that when *all* clusters α are considered in the sum, the cluster expansion is able to represent any function $E(\sigma)$ of configuration σ by an appropriate selection of the values of J_{α} . However, the real advantage of the cluster expansion is that, in practice, it is found to converge rapidly. An accuracy that is sufficient for phase diagram calculations can be achieved by keeping only clusters α that are relatively compact (*e.g.* short-range pairs or small triplets). The unknown parameters of the cluster expansion (the ECI) can then be determined by fitting them to the energy of a relatively small number of configurations obtained through first-principles computations. This approach is known as the Structure Inversion Method (SIM) or the Collony-Williams [4] method.

The cluster expansion thus presents an extremely concise and practical way to model the configurational dependence of an alloy's energy. A typical well-converged cluster expansion of the energy of an alloy consists of about 10 to 20 ECI and necessitates the calculation of the energy of around 30 to 50 ordered structures (see, for instance, [24, 9, 17]). Once the cluster expansion has been constructed, the energy of any configuration can be calculated using Equation 5.1 at a very small computational cost. This enables the use of various statistical mechanical techniques such as Monte Carlo simulations [2], the low-temperature expansion (LTE) [11, 6], the high-temperature expansion (HTE) [6], or the cluster variation method (CVM) [10, 6] to calculate thermodynamic properties and phase diagrams. The **atat** software implements Monte Carlo simulations, the LTE and the HTE.

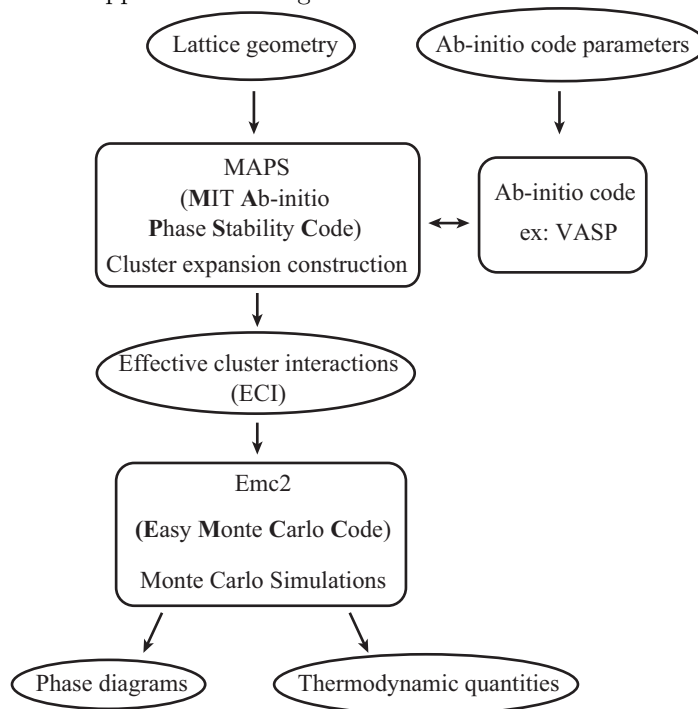
Paralleling the two-step approach described in the previous section, **atat** consists of two main computer programs (see Figure 5.1). The cluster expansion construction is performed by the MIT *Ab initio* Phase Stability (MAPS) code [23], while the Monte Carlo simulations are driven by the Easy Monte Carlo Code (EMC2), developed at Northwestern University [21]. Each of these codes will be discussed in turn.

While the present user guide describes how the **atat** software can be used to carry out all the steps necessary for the calculation of thermodynamic properties from first principles, it must be emphasized that each part of the toolkit can be used as a stand-alone code. For instance, many users may have access to an existing cluster expansion obtained through the SIM or other popular methods, such as concentration-wave-based methods (see, for instance, [7, 6, 20]). It is then straightforward to setup the appropriate input files to run the **emc2** Monte Carlo code. Alternatively, after obtaining a cluster expansion using the **maps** code, users could choose to calculate thermodynamic properties with the cluster variation method (CVM) [10, 6], as implemented in the **IMR-CVM** code [19]. The modularity of the toolkit actually extends below the level of the **maps** and **emc2** codes — many of the subroutines underlying these codes can be accessed through stand-alone utilities [22].

5.3 Cluster expansion construction using the MAPS code

The **maps** code implements the so-called Structure Inversion Method (SIM), also known as the Connolly-Williams method [4]. While the algorithms underlying the **maps** code are described in [23], the present section focuses on its practical use.

Figure 5.1: Methodology implemented in **atat** for the computation of thermodynamic properties from first principles. The automated construction of the cluster expansion is performed by the **maps** code. Whenever needed, **maps** requests the calculation of the formation energy of various atomic configurations by a first-principles code (such as **vasp**). The necessary input files are created and the resulting output files are parsed without requiring user intervention. The output of **maps** is a set of effective cluster interactions that define a computationally efficient Hamiltonian that can be used to perform Monte Carlo simulations with the **emc2** code. These simulations provide thermodynamic properties and phase diagrams that can be used to create thermodynamic databases or supplement existing ones.



5.3.1 Input files

The **maps** code needs two input files: one that specifies the geometry of the parent lattice (**lat.in**) and one that provides the parameters of the first-principles calculations (**xxxx.wrap**, where **xxxx** is the name of the first-principles code used). The clear separation between the thermodynamic and first-principles calculations is a distinguishing feature of **atat** that enables the package to be easily interfaced with any first-principles code. Table 5.1 gives two annotated examples of a lattice geometry input file. The package includes ready-made lattice files for the common lattice types (*e.g.* bcc, fcc, hcp). It also includes an utility that automatically constructs multiple lattice geometry input files for common lattices. For instance,

```
makelat Al,Ti fcc,bcc,hcp
```

creates 3 subdirectories containing the appropriate input files for each specified lattice.

The first-principles input file is usually less than 10 lines long, thanks to the dramatic improvements in the user-friendliness of most modern first-principles codes. For instance, in the case of the widely used VASP code [13, 12], a typical input file is given in Table 5.2. Examples of such input files are provided with the package. Note that **atat** contains a utility that enables the automatic construction of k -point meshes from a single parameter defining the desired target k -point density, the number of k -point per reciprocal atom (KPPRA).

5.3.2 Running the code

The **maps** code is started using the command

Table 5.1: Examples of lattice geometry input file `lat.in`. Typically, the coordinate system entry is used to define the conventional unit cell so that all other entries can be specified in the normalized coordinates that are the most natural for the symmetry of the lattice. The input lattice parameters do not need to be exact, as the first-principles code will optimize them.

Example 1: hcp Ti-Al system

3.1 3.1 5.062 90 90 120	(Coordinate system: $a b c \alpha \beta \gamma$ notation)
1 0 0	(Primitive unit cell: one vector per line
0 1 0	expressed in multiples of the above coordinate
0 0 1	system vectors)
0 0 0 Al,Ti	(Atoms in the lattice)
0.6666666 0.3333333 0.5 Al,Ti	

Example 2: rocksalt CaO-MgO pseudobinary system

4.1 4.1 4.1 90 90 90	
0 0.5 0.5	
0.5 0 0.5	
0.5 0.5 0	
0 0 0 Ca,Mg	(“Active” atoms in the lattice)
0.5 0.5 0.5 0	(“spectator” ion)

Table 5.2: Examples of first-principles code input file (example given for the `vasp` code). It is especially important to verify that the `KPPRA` parameter is set sufficiently large for the system under study.

[INCAR]	
PREC = high	
ENMAX = 200	
ISMEAR = -1	
SIGMA = 0.1	
NSW=41	
IBRION = 2	
ISIF = 3	(See <code>vasp</code> manual for a description of the above 6 parameters.)
KPPRA = 1000	(Sets the k -point density (K Point Per Reciprocal Atom))
DOSTATIC	(Performs a “static run” — see <code>vasp</code> manual)

`maps -d &`

where the option `-d` indicates that all default values of the input parameters should be used, which is what most users will ever need. (The optional parameters can be displayed by typing `maps` by itself and further help is available via the command `maps -h`.) The trailing `&` character cause the command to execute in “background” mode. In this fashion, `maps` can continuously be on the lookout, responding to various “signals”, while the user performs other tasks. (The ongoing discussion assumes that the code is run under a UNIX environment within a shell such as `sh`, `csh`, `tcsh` or `bash`.)

The process of constructing a cluster expansion from first-principles calculations can be summarized as follows.

1. Determine the parameters of the first-principles code that provide the desired accuracy.
2. Let `maps` refine the cluster expansion.
3. Decide when the cluster expansion is sufficiently accurate.

Typically, one calibrates the accuracy of the first-principles calculations using the “pure”¹ structures of the alloy system of interest. To generate the two “pure” structures, type

```
touch ready
```

This creates a file called **ready** which tells **maps** that you are ready to calculate the energy of a structure. Within 10 seconds, **maps** replies with

```
Finding best structure...
```

```
done!
```

maps has just created a directory called **0** and, within it, a file called **str.out** that contains the geometry of one of the two “pure” structures. If you type **touch ready** once more, the other “pure” structure is written to **1/str.out**. You now need to launch the first-principles code to calculate the energy of each structure. Type

```
cd 0
```

to go into the directory of the first structure. Assuming that your first-principles code is called **xxxx**, type

```
runstruct_xxxx &
```

After this command has successfully terminated, display the energy of that structure and go back to the initial directory

```
cat energy
```

```
cd ..
```

and edit the file defining the first-principles code parameters

```
emacs xxxx.wrap &
```

so that the precision of the calculation is increased (*e.g.* increase the *k*-point density or the cut-off of the plane-wave energy). Then you rerun the calculations to check by how much the calculated energy has changed:

```
cd 0
```

```
runstruct_xxxx &
```

```
cat energy (After the calculations are completed)
```

```
cd ..
```

This process is repeated until the user is satisfied with the precision of the calculation (that is, if the energy has become insensitive to changes in the input parameters within the desired accuracy).² A similar study should also be performed for the other “pure” structure (labeled structure 1) and, if one is really concerned with precision, for a few structures with intermediate concentrations.

Once the appropriate *ab initio* code parameters have been determined, the fully automated process can begin. From within the directory where **maps** was started, type

```
pollmach runstruct_xxxx
```

to start the job manager that will monitor the load on your local network of workstations and ask **maps** to generate new structures (*i.e.* atomic arrangements) whenever a processor becomes available. Note that the first time the command is run, instructions will appear on screen that explain how to configure the job dispatching system in accordance to your local computing environment. Once this configuration is complete, the above command should be invoked in the background by appending a “&” to it.

¹In the case of pseudobinary alloys with spectator ions (*e.g.* the MgO-CaO system), the “pure” structures would correspond to the structures where the sublattice of interest is entirely filled with a single type of atom.

²A key number to keep in mind is that an error of 25meV corresponds to 300K on a temperature scale.

5.3.3 Output of MAPS

While the calculations are running, you can check on the status of the best cluster expansion obtained so far. The file `log.out` contains a brief description of the status of the calculations, such as the accuracy of the cluster expansion and various warning messages. Most of the messages pertain to the accurate prediction of the so-called ground states of the alloy system. The ground states, which are the structures that have the lowest energy for each given concentration, are extremely important to predict accurately because they determine which phases will appear on the phase diagram. The four possible messages are described below.

- **Not enough known energies to fit CE.** Before displaying any results, `maps` waits until enough structural energies are known to fit a minimal cluster expansion containing only nearest-neighbor pair interactions and test its accuracy. Thus, the first cluster expansion is typically constructed after at least 4 structural energies have been computed (this number may vary as a function of the symmetry of the lattice).
- **Among structures of known energy, true ground states differ from fitted ground states.** The current cluster expansion incorrectly predicts which structures have the lowest energy for given concentrations, among structures whose first-principles energy is known. The code has built-in checks to avoid this. However, in rare instances, it may be mathematically impossible to satisfy all the constraints that the code imposes for a cluster expansion to be acceptable. This problem becomes increasingly unlikely as the number of calculated structural energies increases, so the user should just wait until the problem fixes itself.
- **Among structures of known energy, true and predicted ground states agree.** Opposite of the previous message. When this message is displayed, `maps` also displays either one of the following two messages.
- **New ground states of volume less or equal to n predicted, see `predstr.out`.** This indicates that the cluster expansion predicts that, at some concentration, there exist other structures that should have an energy even lower than the one of the structures whose energy has been calculated from first principles. In this case, `maps` will investigate the matter by generating those structures and requesting that their energy be calculated. Once again, the user should just wait for the problem to fix itself. The predicted ground states are flagged by a `g` in the `predstr.out` file, so that you can display their energy by typing

```
grep g predstr.out
```

- **No other ground states of n atoms/unit cell or less exist.** The energies of all ground states predicted by the cluster expansion have been confirmed by first-principles calculations. Because it can be computationally intensive to perform a full ground state search when interactions extend beyond the nearest-neighbor shell [6], `maps` uses a search algorithm that merely enumerates every possible structures having n atoms or less per unit cell and uses the cluster expansion to predict their energies. The upper limit n increases automatically as calculations progress.

The `log.out` file also contains two other pieces of information:

- **Concentration range used for ground state checking:** `[a,b]` This displays the user-selected range of concentration over which ground state checking is performed (which can be specified as a command-line option of the `maps` command: `-c0=a -c1=b`). It may be useful to relax the constraints that ground states be correctly reproduced over the whole concentration range when it is known that other parent lattices are stable in some concentration range. In this fashion, the code can focus on providing a higher accuracy in the concentration range where the user needs it.
- **Crossvalidation score:** `s`. This provides the predictive power of the cluster expansion. It is analogous to the root mean square error, except that it is specifically designed to estimate the error made in predicting the energy for structures not included in the least-squares fit [23]. It is defined as

$$CV = \left(\frac{1}{n} \sum_{i=1}^n \left(E_i - \hat{E}_{(i)} \right)^2 \right)^{1/2}$$

where E_i is the calculated energy of structure i , while $\hat{E}_{(i)}$ is the predicted value of the energy of structure i obtained from a least-squares fit to the $(n - 1)$ other structural energies.

The `maps` code also outputs quantitative data in various output files. The simplest way to analyze this data is by typing

```
mapsrep
```

As illustrated in Figure 5.2, this command displays, in turn

- The `log.out` file described earlier.
- The formation energy of all structures whose energy is known from first-principles calculations, as well as the predicted energy of all structures `maps` has in memory. The convex hull of the ground states among structures of known energy is overlaid while the new predicted ground states (if any) are marked by an “x”. (Note that this ground state line is only meaningful if the `log.out` file contains “Among structures of known energy, true and predicted ground states agree.”)
- The formation energy of all structures calculated from first principles and associated ground state line.
- A plot of the magnitude of the Effective Cluster Interactions (ECI) as a function of the diameter of their associated cluster (defined as the maximum distance between any two sites in the cluster). Pairs, triplets, etc. are plotted consecutively. This plot is useful to assess the convergence of the cluster expansion. When the magnitude of the ECI for the larger clusters has clearly decayed to a negligible value (relative to the nearest-neighbor pair ECI), this is indicative of a well-converged cluster expansion.
- A plot of the residuals of the fit (*i.e.* the fitting error) for each structure. This information is useful to locate potential problems in the first-principles calculations. Indeed, when first-principles calculations exhibit numerical problems, this typically results in calculated energies that are poorly reproduced by the cluster expansion.

When the user is satisfied with the results (which are constantly updated), `maps` can be stopped by creating a file called `stop` in the current directory using the command:

```
touch stop
```

while the job dispatching system can be stopped by typing:

```
touch stoppoll
```

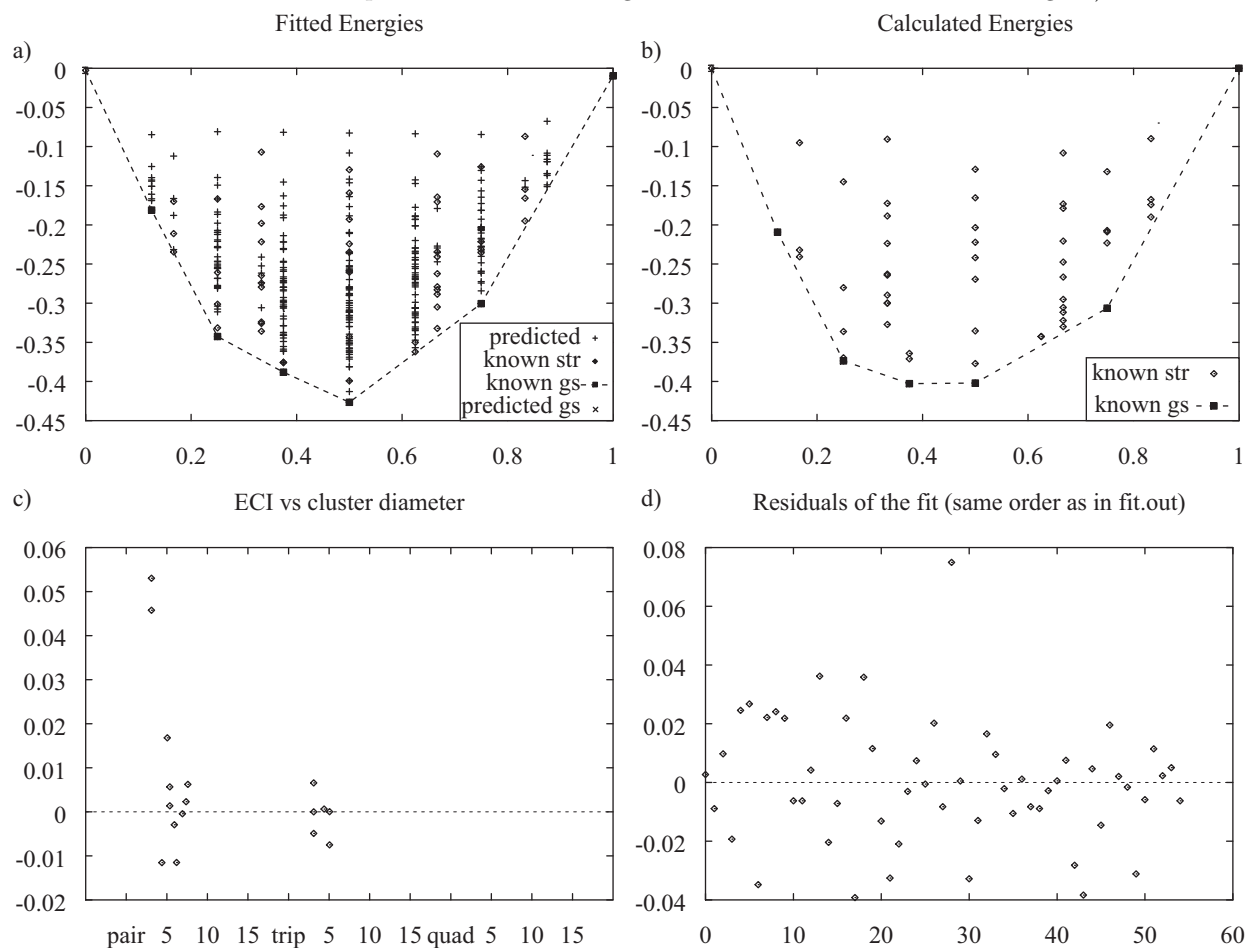
A cluster expansion can be considered satisfactory when

1. All ground states are correctly reproduced and no new ground states are predicted. (The `log.out` file would then indicate that Among structures of known energy, true and predicted ground states agree. No other ground states of n atoms/unit cell or less exist.)
2. The crossvalidation score, as given in the `log.out` file, is small (typically less than 0.025 eV).
3. Optionally, it is instructive to verify that the magnitude of the ECI decays as a function of the diameter of the corresponding cluster and as a function of the number of sites it contains.

5.4 Monte Carlo simulations

The `emc2` code implements semi-grand canonical Monte Carlo simulations [16, 2, 8, 15], where the total number of atoms is kept fixed, while the concentration is allowed to adapt to an externally imposed difference in the chemical potential of the two types of atoms. The chemical potential difference will be simply referred to as the “chemical potential” in what follows. This ensemble offers the advantage that, for any imposed

Figure 5.2: Output of the `maps` Code, as reported by the `mapsrep` command. a) Energies predicted from the cluster expansion as a function of composition for each structure generated. “`known str`” denotes structures whose energy has been calculated from first principles. “`known gs`” indicate the ground states that have so far been confirmed by first-principles calculations and the dashed line outlines the convex hull of the ground states, which serves as a threshold to detect other candidate ground states. “`predicted`” denotes structures whose energy has *not yet* been calculated from first principles. “`predicted gs`” are structures that are predicted by the cluster expansion to be ground states, although this prediction has not yet been confirmed by first-principles calculations. b) Energies calculated from first principles. “`known str`” and “`known gs`” are as in a), except that the energy calculated from first principles is reported. c) Effective Cluster Interaction (ECI) as a function of the diameter of the associated cluster and as a function of the number of sites in the cluster (*i.e.* pair, triplet, etc.). d) Residuals of the fit, that is, the difference between the first-principles energies and the energies predicted from the cluster expansion. (The abscissa refers to the line number within the output file `fit.out` listing all the structures with known energies.)



chemical potential, the equilibrium state of the system is a single phase equilibrium, free of interfaces.³ It also simplifies the process of calculating free energies through thermodynamic integration. While a detailed description of the algorithm underlying this code can be found in [21], the current section focuses on the practical usage of the code.

³If the simulation cell is commensurate with the unit cell of the phase under study, a requirement that the code automatically ensures.

5.4.1 General input parameters

The Monte Carlo code needs the following files as an input

1. A lattice geometry file (`lat.in`), which is the same as the input for `maps` (see Table 5.1).
2. Files providing the cluster expansion (the clusters used are listed in the `clusters.out` file while the corresponding ECI are listed in the `eci.out` file.). These files are automatically generated by `maps`, although users can supply their own cluster expansion, if desired. A description of the format of these files is available by typing `maps -h`.
3. A list of ground states (`gs_str.out`), which merely provide convenient starting configurations for the simulations. `maps` also automatically creates this file.

The parameters controlling the simulation are specified as command-line options. The first input parameter(s) needed by the code are the phase(s) whose thermodynamic properties are to be determined. There are two ways to invoke the Monte Carlo simulation code. When the command `emc2` is used, a single Monte Carlo simulation is run to allow the calculation of thermodynamic properties of a single phase for the whole region of chemical potential and temperature where that phase is stable. The phase of interest is specified by a command-line option of the form

`-gs= n ,`

where n is a number between -1 and $G - 1$ (inclusive), where G is the number of ground states. The value -1 indicates the disordered phase while values ranging from 0 to $G - 1$ indicate the phases associated with each ground states (0 denoting the ground state with the smallest composition). When the command `phb` is used, two Monte Carlo simulations are run simultaneously to enable the determination of the temperature-composition phase boundary associated with a given two-phase equilibrium. The two phases are specified by

`-gs1= n_1 -gs2= n_2 .`

It is possible to compute a two-phase equilibrium between phases defined on a different parent lattice. In this case, the user needs to specify the directories where the cluster expansions of each lattice resides using the options of the form

`-d1=directory 1 -d2=directory 2`

The accuracy of the thermodynamic properties obtained from Monte-Carlo simulations is determined by two parameters: The size of the simulation cell and the duration of the simulation.

The size of the simulation cell is specified by providing the radius r of a sphere through the command-line option

`-er= r .`

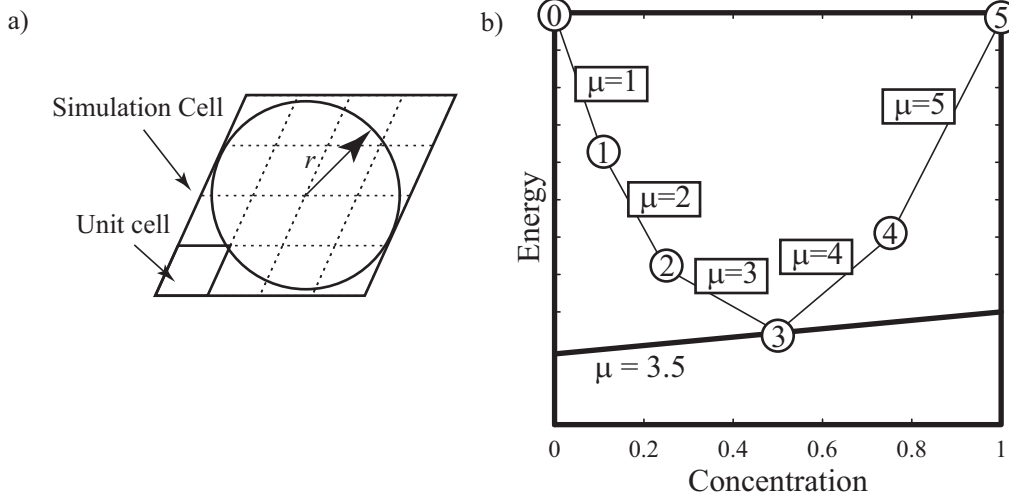
As illustrated in Figure 5.3a, the simulation cell size will be the smallest supercell that both contains that sphere and that is commensurate with the unit cell of the ground state of interest. This way of specifying the simulation cell size ensures that the system size is comparable along every direction, regardless of the crystal structure of the ground state of interest. It also frees the user from manually checking the complicated requirement of commensurability. It is important that the user check that the simulation cell size is sufficiently large for the thermodynamic properties of interest to be close to their infinite-system-size limiting value. This can be done by gradually increasing the system size until the calculated quantities become insensitive to the further increases in system size, within the desired accuracy.

The duration of the simulations is automatically determined by the code from a user-specified target precision on the atomic composition of the phase, indicated by a command-line option of the form

`-dx= Δx .`

Alternatively, the user may also manually set the number n_{eq} of Monte Carlo steps the system is allowed to equilibrate before thermodynamic averages are computed over a certain number n_{avg} of Monte Carlo steps using the options

Figure 5.3: Definitions of the quantities used to specify a) the simulation cell size and b) the chemical potential.



$$-eq=n_{eq} \quad -n=n_{avg}.$$

The Monte Carlo code also needs additional parameters that specify which portion of a phase's free energy surface needs to be computed. With `emc2`, the range of temperatures to be scanned are specified in either one of the following two ways:

$$-T0=T_0 \quad -T1=T_1 \quad -dT=\Delta T \quad (\text{for steps in direct temperature})$$

$$\text{or } -T0=T_0 \quad -T1=T_1 \quad -db=\Delta(1/T) \quad (\text{for steps in reciprocal temperature}).$$

The temperature steps in reciprocal temperature ($\Delta(1/T)$) can be useful when calculations are started from infinite temperatures down to a finite temperature. The `-T1` and `-dT` (or `-db`) options can be omitted if calculations at a single temperature are desired. Since the program automatically stops when a phase transition is detected, it is not necessary to know in advance the temperature range of stability of the phase. The user only needs to ensure that the initial temperature lies within the region of stability of the phase of interest. An obvious starting point is $T_0 = 0$, since the ground state is then stable, by definition. With the `phb` code, the syntax is

$$-T=T \quad -dT=\Delta T$$

If the `-T` option is omitted, calculations start at absolute zero.⁴ The energy and temperature units used are set by specifying the Boltzmann's constant with the command-line option

$$-k=k_B.$$

A value of 8.617×10^{-5} corresponds to energies in eV and temperatures in Kelvin.

With `emc2`, the range of chemical potentials to be scanned needs to be specified. Once again, only the starting point really matters, because the code will stop when a phase transition is reached. By default, chemical potentials are given in a dimensionless form, so as to facilitate the link between the value of the chemical potential and the phase that it stabilizes. For instance, a chemical potential equal to 3.0 is such that it would stabilize a two phase equilibrium between phase number 2 and phase number 3 at absolute zero (see Figure 5.3b). A chemical potential between 3.0 and 4.0 stabilizes phase number 3 at absolute zero. While these ranges of stability are no longer exact at finite temperature, this dimensionless chemical potential still provides easy-to-interpret input parameters. The syntax is

⁴Temperature steps in reciprocal temperature are not needed, because a two-phase equilibrium never extends up to infinite temperature.

$$-\mu_0=\mu_0 \quad -\mu_1=\mu_1 \quad -d\mu=\Delta\mu$$

where $\Delta\mu$ is the chemical potential step between each new simulation. Chemical potentials can also be entered in absolute value (say in eV, if the energies are in eV) by specifying the `-abs` option. Note that the output files always give the absolute chemical potentials, so that thermodynamic quantities can be computed from them. With `phb`, the initial chemical potential is optional when starting from absolute zero because the code can determine the required value from the ground state energies. It can nevertheless be specified (in absolute value) with the `-mu= μ` option, if a finite temperature starting point is desired.

A list of the command line options of either the `emc2` or `phb` codes can be displayed by simply typing either command by itself. More detailed help is displayed using the `-h` option.

5.4.2 Examples

We now give simple examples of the usage of these commands. Consider the calculation of the free energy of the phase associated with ground state number 1 as a function of concentration and temperature. Then, the required commands could, for instance, be

```
emc2 -gs=1 -mu0=1.5 -mu1=0.5 -dmu=0.04 -T0=300 -T1=5000 -dT=50 -k=8.617e-5 -dx=1e-3 -er=50
-innerT -o=mc10.out
```

```
emc2 -gs=1 -mu0=1.5 -mu1=2.5 -dmu=0.04 -T0=300 -T1=5000 -dT=50 -k=8.617e-5 -dx=1e-3 -er=50
-innerT -o=mc12.out
```

(The only difference in the two command lines is the value of `-mu1` and the output file name, specified by the `-o` option.) These commands separately compute the two “halves” of the free energy surface, corresponding to the values of the chemical potential below and above the “middle” value of 1.5 which stabilizes ground state 1 at absolute zero. This natural separation allows you to run each half calculation on a separate processor and obtain the results in half the time. The values of `-dmu`, `-dT`, `-dx` and `-er` given here are typical values. The user should ensure that these values are such that the results are converged. Note that, thanks to the way these precision parameters are input, if satisfying values have been found for one simulation, the same values will provide a comparable accuracy for other simulations of the same system. The option `-innerT` indicates that the inner loop of the sequence of simulations scans the temperature axis while the outer loop scans the chemical potential. In this fashion, the point of highest temperature in the region of stability of the phase will be known early during the calculations. If the user is more interested in obtaining solubility limits early on, this option can be omitted and the inner loop will scan the chemical potential axis. In either cases, the code exits the inner loop (and the outer loop, if appropriate) when it encounters a phase transition.

The `emc2` code thus enables the automated calculation of the whole free energy surface of a given phase, as illustrated in Figure 5.4a. Such free energy surfaces can be used as an input to construct thermodynamic databases or supplement existing ones. To facilitate this process, a utility that converts the output of `emc2` into input files for the fitting module of ThermoCalc is provided.

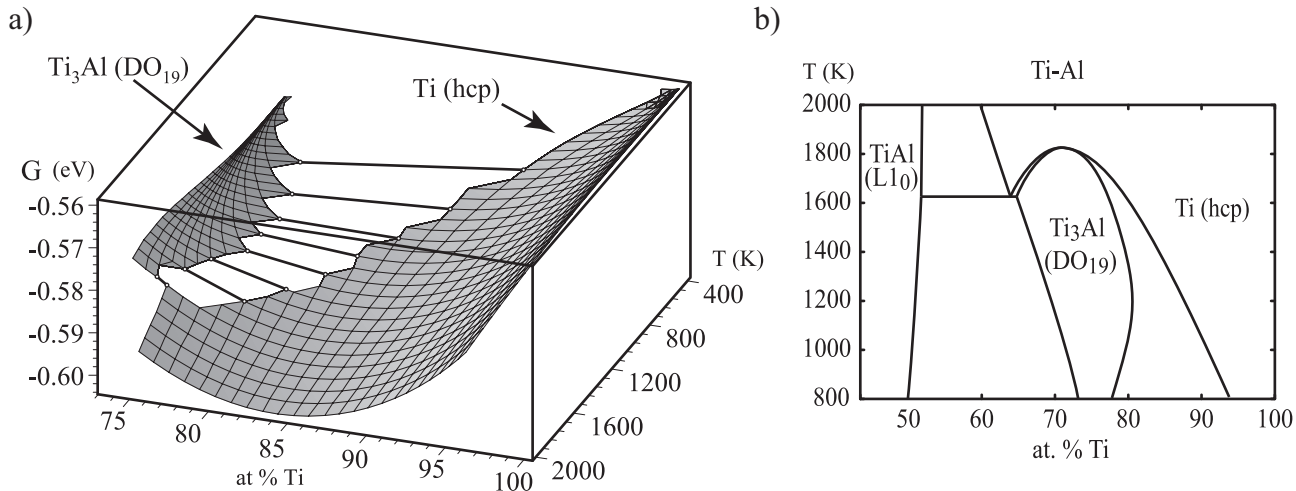
While the above examples focus on the calculation of a phase’s thermodynamic properties over its whole region of stability, one may be interested in directly computing the temperature-composition phase boundary without first constructing a full free energy surface. To accomplish this task, a typical command-line for the `phb` program would be

```
phb -gs1=0 -gs2=1 -dT=25 -dx=1e-3 -er=50 -k=8.617e-5 -ltep=5e-3 -o=ph01.out
```

This command computes the two phase equilibrium between phase 0 and phase 1, starting at absolute zero and incrementing temperature in steps of 25 K. (The `-ltep` option indicates that a Low Temperature Expansion (LTE) should be used instead of Monte Carlo simulation whenever its precision is better than 5×10^{-3} eV.) The output file `ph01.out` contains the temperature-composition phase boundary of interest, as well as the chemical potential stabilizing the two-phase equilibrium as a function of temperature. This output can be used to generate phase diagrams, as illustrated in Figure 5.4b.

The program automatically terminates when the “end” of the two-phase equilibrium has been reached. If the two-phase equilibrium disappears because of the appearance of a third phase, two new two-phase equilibria have to be separately calculated. To do so, one uses the final temperature T and chemical potential μ given in the output file as a starting point for two new `phb` runs:

Figure 5.4: Output of Monte Carlo codes. a) The `emc2` provides free energy surfaces as a function of temperature T and composition x . (For clarity, the common tangent construction (thick lines) is drawn over the calculated free energy.) b) The `phb` command generates temperature-composition phase diagrams. The calculational details underlying these results can be found in [23, 21]



```
phb -T=T -mu=μ -gs1=0 -gs2=-1 -dT=25 -dx=1e-3 -er=50 -k=8.617e-5 -o=ph0d.out
```

```
phb -T=T -mu=μ -gs1=-1 -gs2=1 -dT=25 -dx=1e-3 -er=50 -k=8.617e-5 -o=phd1.out
```

In the above example, it is assumed that the new phase appearing is the disordered phase (indicated by the number -1), which will usually be the case. Of course, it is also possible that a given two-phase equilibrium terminates because one of the two phases disappears. In this case, only one new calculation needs to be started, as in the following example:

```
phb -T=T -mu=μ -gs1=0 -gs2=2 -dT=25 -dx=1e-3 -er=50 -k=8.617e-5 -o=phd1.out
```

Note that phase 1 has been replaced by phase 2. Finally, it is also possible that the two-phase equilibrium terminates because the concentration of each phase converges to the same value, a situation which requires no further calculations. The user can easily distinguish these three cases by merely comparing the final composition of each phase.

5.4.3 Interpreting the output files

The output file of `emc2` reports the value of all calculated thermodynamic functions for each value of temperature and chemical potential scanned. The quantities reported include

- Statistical averages over Monte Carlo steps, such as energy, concentration, short-range and long-range order parameters.⁵
- Integrated statistical averages, such as the Gibbs free energy G or the semi-grand-canonical potential $\phi = G - \mu x$.
- The result of common approximations, namely, the low temperature expansion (LTE) [6, 11, 26], the mean-field (MF) approximation and the high temperature expansion (HTE) (see, for instance, [6]).

While quantities obtained from statistical averages over Monte Carlo steps are valid for all temperatures and chemical potentials, caution must be exercised when interpreting the result of the various approximations or when looking at the integrated quantities. The LTE, MF and HTE approximations are only accurate in a limited range of temperature and it is the responsibility of the user to assess this range of validity.

⁵For efficiency reasons, the long range order parameters are only calculated when starting from an ordered phase.

Also, the free energy or the semi-grand-canonical potential are obtained from thermodynamic integration and are thus only valid if the starting point of the integration is chosen appropriately. By default, the low temperature expansion value is used as a starting point whenever the phase of interest is a ground state, while the high temperature expansion is used when the phase of interest is the disordered state. Hence, to obtain absolute values of the semi-grand-canonical potential, one must ensure that the calculations are started at a sufficiently low temperature (or sufficiently high temperature, in the case of the disordered phase). This can be checked by comparing the Monte Carlo estimates with the LTE (or HTE) estimates and verifying that they agree for the first few steps of the thermodynamic integration. A user-specified starting point for ϕ (*e.g.* obtained from an earlier Monte Carlo simulation) can be indicated using the option

```
-phi0= $\phi$ 
```

Note that, unlike `emc2`, the `phb` code automatically makes use of the low temperature expansion whenever it is sufficiently accurate in order to save a considerable amount of computational time.

By default, the code reports the thermodynamic quantities associated with the semi-grand-canonical ensemble, such as the semi-grand-canonical potential ϕ . The command-line option `-can`, instructs the code to add μx to all appropriate thermodynamic quantities, so that the code outputs the more commonly used canonical quantities, such as the Gibbs free energy G and the internal energy E .

5.5 Recent developments

The multicomponent version of `maps` is `mmaps` and the multicomponent version of `emc2` is `memc2`. The `phb` code does not have a multicomponent version. All other utilities can handle multicomponent systems.

Although the present tutorial does not discuss the topic, `atat` also implements reciprocal space cluster expansions and, in particular, the constituent strain formalism [14], see the command reference for `csfit`. `atat` can also calculate nonconfigurational contributions to the free energy, such as lattice vibrations and electronic excitations, see the command reference for `fitsvsl`, `svsl`, `fitfc` and `felec`.

5.6 Conclusion

The Alloy Theoretic Automated Toolkit (ATAT) drastically simplifies the practical implementation of the Connolly-Williams method, in combination with semi-grand-canonical Monte Carlo simulations, thus providing a high-level interface to the calculation of thermodynamic properties from first principles. This toolkit enables researcher to focus on higher-level aspects of first-principles thermodynamic calculations by encapsulating the intricate details of the calculations under an easy-to-use interface. It also makes these powerful methodologies readily available to the wide community of researchers who could benefit from it.

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Chapter 6

Interfacing with ThermoCalc

A description of the steps necessary to convert the output of the Monte Carlo code `emc2` to input files suitable for the PARROT module of ThermoCalc are described in an online tutorial by Gautam Ghosh available in: `atat/doc/emc2tc.pdf`. These steps are partially automated in the script `atat/glue/tc/emc2tc` included in the ATAT distribution.

Chapter 7

Command Reference

7.1 Man pages

7.1.1 apb

This code creates an APB from a given structure file and computes its APB energy using ECIs.

Reference:

R. Sun and A. van de Walle.

Automating impurity-enhanced antiphase boundary energy calculations from ab initio Monte Carlo.

Calphad 53, 20 (2016).

- * The structure file is assumed to have the following form.
Orient the cell such that the APB plane is defined by T1 x T2:

```
a    0    0
0    b    0
0    0    c
T1x T1y T1z
T2x T2y T2z
T3x T3y T3z
```

- * The APB structure is generated by duplicating atoms along T3 and shifting them by the slip vector.
- * To obtain the APB energy, the user should first perform a cluster expansion on the system and provide clusters.out and eci.out as input.
- * In Monte Carlo mode, the concentration of the initial structure is fixed in conccons.in. The temperature and number of averaging steps are used to write control.in. The file name of the perfect and defective structures MC snapshot is str?????.out and str????_apb.out, respectively. After MC, the APBs are generated and their energies are computed.
- * The output format is:
[id] e0 e1 gamma
where
[id] = structure ID (only shown in Monte Carlo mode)

```

e0 = total energy (eV/atom) of perfect cell
e1 = total energy (eV/atom) of defective cell
gamma = APB energy (mJ/m^2)

```

7.1.2 corrdump

->What does this program do?

- 1) It reads the lattice file (specified by the -l option).
- 2) It determines the space group of this lattice and writes it to the sym.out file.
- 3) It finds all symmetrically distinct clusters that satisfy the conditions specified by the options -2 through -6.
For instance, if -2=2.1 -3=1.1 is specified, only pairs shorter than 2.1 units and triplets containing no pairs longer than 1.1 will be selected.
- 4) It writes all clusters found to clusters.out.
If the -c option is specified, clusters are read from clusters.out instead.
- 5) It reads the structure file (specified by the -s option).
- 6) It determines, for that structure, the correlations associated with all the clusters chosen earlier.
This information is then output on one line, in the same order as in the clusters.out file. See below for conventions used to calculate correlations.
- 7) It writes the files corrdump.log containing the list of all adjustments needed to map the (possibly relaxed) structure onto the ideal lattice.

->File formats

Lattice and structure files

Both the lattice and the structure files have a similar structure.

First, the coordinate system a,b,c is specified, either as

```
[a] [b] [c] [alpha] [beta] [gamma]
```

or as:

```
[ax] [ay] [az]
```

```
[bx] [by] [bz]
```

```
[cx] [cy] [cz]
```

Then the lattice vectors u,v,w are listed, expressed in the coordinate system just defined:

```
[ua] [ub] [uc]
```

```
[va] [vb] [vc]
```

```
[wa] [wb] [wc]
```

Finally, atom positions and types are given, expressed in the same coordinate system as the lattice vectors:

```
[atom1a] [atom1b] [atom1c] [atom1type]
```

```
[atom2a] [atom2b] [atom2c] [atom1type]
```

etc.

In the lattice file:

- The atom type is a comma-separated list of the atomic symbols of the atoms that can sit the lattice site.
- In a binary, the first symbol listed is assigned a spin of -1.
- In general, ordering of the atom symbol corresponds to value of $s=0,1,\dots$ in the table 'Convention used to calculate the correlations' below.
- When only one symbol is listed, this site is ignored for the purpose of calculating correlations, but not for determining symmetry.
- The atomic symbol 'Vac' is used to indicate a vacancy.

In the structure file:

- The atom type is just a single atomic symbol (which, of course, has to be among the atomic symbols given in the lattice file).
- Vacancies do not need to be specified.

Examples

The fcc lattice of the Cu-Au system:

```
1 1 1 90 90 90
0 0.5 0.5
0.5 0 0.5
0.5 0.5 0
0 0 0 Cu,Au
```

The Cu₃Au L1₂ structure:

```
1 1 1 90 90 90
1 0 0
0 1 0
0 0 1
0 0 0 Au
0.5 0.5 0 Cu
0.5 0 0.5 Cu
0 0.5 0.5 Cu
```

A lattice for the Li_x Co_y Al_(1-y) O₂ system:

```
0.707 0.707 6.928 90 90 120
0.3333 0.6667 0.3333
-0.6667 -0.3333 0.3333
0.3333 -0.3333 0.3333
0 0 0 Li,Vac
0.3333 0.6667 0.0833 0
0.6667 0.3333 0.1667 Co,Al
0 0 0.25 0
```

Symmetry file format (sym.out)

[number of symmetry operations]

3x3 matrix: point operation

1x3 matrix: translation

repeat, etc.

Note that if you enter more than one unit cell of the lattice, sym.out will contain some pure translations as symmetry elements.

Cluster file format (clusters.out)

for each cluster:

[multiplicity]

[length of the longest pair within the cluster]

[number of points in cluster]

[coordinates of 1st point] [number of possible species-2] [cluster function]

[coordinates of 2nd point] [number of possible species-2] [cluster function]

etc.

repeat, etc.

(Multiplicity and length are ignored when reading in the clusters.out file.)

For each 'point' the following convention apply

-The coordinates are expressed in the coordinate system given in the first line (or the first 3 lines) of the lat.in file.

-The 'number of possible species' distinguishes between binaries, ternaries, etc...

Since each site can accomodate any number of atom types, this is specified for each point of the cluster.

-In multicomponent system, the cluster function are numbered from 0 to number of possible species-2.

In the simple of a binary system [number of possible species-2] [cluster function] are just 0 0.

For a ternary, the possible values are 1 0 and 1 1.

All the utilities that are not yet multicomponent-ready just ignore the entries [number of possible species-2] [cluster function]

Convention used to calculate the correlations:

The cluster functions in a m-component system are defined as

function '0' : $-\cos(2\pi \cdot 1 \cdot s/m)$

function '1' : $-\sin(2\pi \cdot 1 \cdot s/m)$

.

.

.

$-\cos(2\pi [m/2] s/m)$

$-\sin(2\pi [m/2] s/m)$ <--- the last sin() is omitted if m is even

where the occupation variable s can take any values in {0,...,m-1}

and [...] denotes the 'round down' operation.

Note that, these functions reduce to the single function $(-1)^s$ in the binary case.

Special options:

-sym: Just find the space group and then abort.

-clus: Just find space group and clusters and then abort.

-z: To find symmetry operations, atoms are considered to lie on top of one another when they are less than this much apart.

-sig: Number of significant digits printed.

->Cautions

When vacancies are specified, the program may not be able to warn you that the structure and the lattice just don't fit.

Carefully inspect the corrdump.log file!

If the structure has significant cell shape relaxations, the program will be unable to find how it relates to the ideal lattice. The problem gets worse as the supercell size of the structure gets bigger.

There is no limit on how far an atom in a structure can be from the ideal lattice site. The program first finds the atom that can be the most unambiguously assigned to a lattice site. It then finds the next best assignement and so on. This is actually a pretty robust way to do this. But keep in mind that the -z option does NOT control this process.

7.1.3 cv

Help for cv

Command-line options:

- g: Favor cluster choices where the ground state line is correct.
(It is suggested to turn this feature on if you have trouble getting the right ground states).
- w: Weight structures by $w/(\text{struct_energy} - \text{gs_energy} + w)$
(recommended value: start with 1. and decrease down to about 0.010 (eV) if you have trouble getting the correct ground states.
Default: all structure have equal weight)
- p: Penalty for structures that lie below the ground state line.
(recommended value: start with 1. and increase up to about 20. if you have trouble getting the correct ground states.
Default: 0)

-> corr.in

table of correlations, one structure per line
(exactly the output of corrdump)

-> energy.in

column of energy values, one per line, in the same order as in the corr.in file.
If energy is unknown, put 'x'.

-> include.in

This file can be omitted.

It gives a list of clusters than must be part of the cluster expansion.
(Each number in this file indicates a column in the corr.in file.)

-> weight.in

This file can be omitted.

It indicates the weight that must be given to each structure.
Usually this file is copied from the weight_sug.out file obtained from a previous run.
Each weight in this file is multiplied by
 $\text{gs_prec}/(\text{structure_energy} - \text{ground_state_energy} + \text{gs_prec})$

(see above).

-> clusters.out

A list of the clusters (in the same format as generated by either maps or corrdump)

OUTPUT FILES

-> fit.out

Contains the results of the fit, one structure per line and each line has the following information:

concentration energy fitted_energy weight index
'concentration' lies between 0 and 1.
'energy' is the true formation energy.
'fitted_energy' is the predicted formation energy.
'weight' is the weight of this structure in the fit.
'index' is the structure number (not counting structure of unknown energies)

-> predstr.out

Contains the predicted energy of all structures whose true energy is unknown. Format: one structure per line, and each line has the following information:

concentration predicted_energy -1 status
the '-1' is for compatibility with maps.
status is 'u' (for compatibility with maps) and a 'g' is appended to status if that structure is predicted to be a ground state.
To list all predicted ground states, type
grep 'g' predstr.out

-> gs.out

The ground states, as determined by energies given in energy.in. Each line is a concentration, energy pair.

-> gs_fitted.out

The ground states, as determined by energies predicted from the cluster expansion. (Structures for which the true energy is unknown are omitted.)
Each line is a concentration, energy pair.

-> weight_sug.out

If the ground states are incorrectly predicted by the cluster expansion, this file contains suggested weight that should correct the problem.

just type
cp weight_sug.out weight.in
and rerun the program.

-> weight_used.out

Weights actually used to do the fit.

-> fit.gnu

A gnuplot script that display all the relevant information.

Just type:

gnuplot fit.gnu

-> eci.out

The values of the fitted eci, in the same order as in the corr.in file.

7.1.4 emc2

Easy Monte Carlo Code (emc2)

by Axel van de Walle

Command line parameters:

-T0 : initial temperature

-T1 : final temperature

-dT : temperature step

-db : inverse temperature step

Temperatures are given in units of energy unless the -k option is set.

-dT and -db are mutually exclusive. Which option you use determines whether T or 1/T are uniformly spaced. The output file always shows T.

Avoid setting -T0 or -T1 to 0: The output will be nonsensical.

-k : Sets boltzman's constant (default k=1). This only affects how temperatures are converted in energies. -k=8.617e-5 lets you enter temperatures in kelvins when energies are in eV.
(You can also select this value by using the -keV option.)

-mu0 : initial chemical potential

-mu1 : final chemical potential

By default, these are input as dimensionless values defined as follows: If x is integer, x is the chemical potential that stabilizes a two phase equilibrium between ground state x-1 and x. (Ground states are numbered starting at 0.) Fractional values interpolate linearly between these integer values. Negative values or values larger than the number of ground states - 1 are allowed. The values of mu are found by linear extrapolation. If the -abs option is specified, the value of mu are in units of energy per spin value (as in the output file). (The ground states are read in from the file gs_str.out .)

If there are only two ground states, the only correction performed is to shift mu so that mu=0 stabilizes a two-phase equilibrium between the two ground states.

If there are less than 2 ground states (!), no correction is made.

-dmu : chemical potential step (expressed in the same units as mu0 and mu1).

NOTE: If mu1 is omitted, only a scan through T is performed keeping mu=mu0.

If T1 is omitted, only scan through μ is performed keeping $T=T_0$.

- gs : Gives the index of the ground state to use as an initial spin configuration (the leftmost pure element is numbered 0). If the index is -1, the disordered state with an average spin of zero is used as the starting configuration.
- phi0 : value of the grand canonical potential at the initial point of the run. If left unspecified, it is set to the grand canonical potential of given by either the 1-spin Low Temperature Expansion or the High Temperature Expansion, depending whether the initial state is an ordered or a disordered phase.
- er : enclosed radius. The Monte Carlo cell will be the smallest possible supercell of unit cell of the initial configuration such that a sphere of that radius fits inside it. This allows you to specify the system size in a structure-independent fashion.
- innerT : by default, the inner loop is over values μ while the outer loop is over values of T . This flag permutes that order.
- eq : number of equilibration passes. (Equilibration is performed at the beginning of each step of the outer loop.)
- n : number of Monte Carlo passes performed to evaluate thermodynamic quantities at each step of the inner loop.
- dx: instead of specifying -eq and -n, you can ask the code to equilibrate and run for a time such that the average concentration is accurate within the target precision specified by -dx=.
- tstat : Critical value of the test for discontinuity. The code is able to catch phase transformations (most of the time) when going from an ordered phase to another or to the disordered state. This involves a statistical test which a user-specified confidence level. The default, 3, corresponds to a 0.4% chance of finding a transition where there is none. (Refer to a standard normal table.) -tstat=0 turns off this option. Suggestion: if a phase transition is undetected, first try to reduce dT or $d\mu$ or increase n or decrease dx before toying around with this parameter. Also: beware of hysteresis.
- sigdig : Number of significant digits printed. Default is 6.
- o: Name of the output file (default: mc.out).

Tricks:

To read parameters from a file, use:

```
emc2 'cat inputfile'
```

where inputfile contains the commands line options.

To selectively display a few of the output quantities, use:

```
emc2 [options] | cut -f n1,n2,n3,...
where n1,n2,n3,... are the column number desired (see below).
```

Input files:

```
lat.in      : description of the lattice.
clusters.out : describes the clusters.
eci.out     : provides the ECI.
gs_str.out  : a list of ground states, in increasing order of concentration.
```

These 4 files can be created by maps.

See maps documentation (maps -h) for a description of the formats.

Optional input file:

```
teci.out    : if present, provides temperature-dependent eci (overrides eci.out)
              (Note that, even when the teci.out file is used, column 6 of the output file
              reflects only the configurational contribution to the heat capacity.)
```

The format this file is:

```
[maximum temperature: Tm]
[number of temperatures where eci are provided: nT]
[ECIs for temperature 0]
[ECIs for temperature Tm/(nT-1)]
[ECIs for temperature 2*Tm/(nT-1)]
...
[ECIs for temperature Tm]
```

Note that these numbers can be separated by blanks or newlines, as desired.

Format of the output file:

Each line contains, in order:

- 1) T: Temperature
- 2) mu: chemical potential (derivative of the free energy wrt to average spin)
- 3) E-mu*x: Average energy (per spin)
- 4) x: Average Concentration [ranges from -1 to 1]
- 5) phi: grand canonical potential
- 6) E2: Variance of the energy (proportional to heat capacity)
- 7) x2: Variance of the concentration (proportional to susceptibility)
- 8) E_lte-mu*x_lte: calculated with a one spin Low Temperature Expansion
- 9) x_lte
- 10) phi_lte
- 11) E_mf-mu*x_mf: calculated with the Mean Field approximation
- 12) x_mf
- 13) phi_mf
- 14) E_hte-mu*x_hte: calculated with a High Temperature Expansion (ideal solution + polynomial in x)
- 15) x_hte
- 16) phi_hte
- 17) lro: Long Range Order parameter of the initial ordered phase (=0 if initial phase is disordered).
- 18-) corr: the average correlations associated with each cluster.

NOTE: to obtain 'canonical' rather than grand canonical quantities,

use the `-g2c` option. This has the effect of adding μ^*x to columns 3,5,8,10,11,13,14,16.

7.1.5 felec

This code calculates the electronic free energy within the one-electron and temperature-independent bands approximations.

It needs an `dos.out` input file (whose name can be changed with `-dos` option) that has the following format:

```
[number of electron in unitcell]
[energy width of each bins used to calculate the dos]
[a multiplicative scale factor to adjust units]
[the density in each bin, in states/unit cell/energy] <- repeated
```

The code calculates the electronic free energy from temperature T_0 to T_1 in steps of dT .

- The defaults are $T_0=0$ $T_1=2000$ $dT=100$.
- If a file `Trange.in` exists in the upper directory, it is used to set T_0, T_1, dT :
`Trange.in` must contain two numbers on one line separated by a space: T_1 ($T_1/dT+1$).
 Note that $T_0=0$ always.
 For phase diagram calculations, you must use this method to specify the temperature range.
- These defaults can be overridden by the `-T0`, `-T1` and `-dT` options.

The output files contain the free energy per unit cell.

`felec.log` contain temperature and corresponding free energy on each line.

`felec` contains the free energies only.

`plotdos.out` contains the dos (col 1: energy normalized so that $E_f=0$, col 2: DOS)

-> For including electronic entropy in phase diagram calculations

You are likely to use this code as follow:

```
#first create the Trange.in file for up to 2000K in intervals of 100K:
```

```
echo 2000 21 > Trange.in
```

```
#This executes the svsl code in each subdirectory containing dos.out but no error file.
```

```
foreachfile -e dos.out pwd \; felec [options if desired]
```

```
#constructs a cluster expansion of the electronic free energy (eci are in felec.eci)
```

```
clusterexpand felec
```

```
#add the energetic eci from eci.out to the electronic eci from felec.eci and create the teci.out
```

```
#file that will be read by the Monte Carlo code.
```

```
mkteci felec.eci
```

```
#you can even combine vibrational and electronic eci:
```

```
mkteci fvib.eci felec.eci
```

7.1.6 fempmag

I. Ohnuma et al., Phase equilibria in the Fe-Co binary system, *Acta Materialia* 50 (2002) 379.

7.1.7 fitfc

Calculates vibrational properties by fitting a spring model to reaction forces resulting from imposed atomic displacements.

The examples below are given assuming that one uses the vasp code, although other ab initio codes would work as well.

The calculations proceed as follows:

- 1) You first need to fully relax the structure of interest. The code expects the relaxed geometry in the file `str_relax.out`. It also expects a `str.out` file containing the unrelaxed geometry (which may be the same as the relaxed geometry, if you wish). The unrelaxed geometry will be used to determine the neighbor shells and measure distances between atoms. Typically the user would specify the `str.out` file, then obtain the `str_relax.out` file by running an ab initio code with a command of the form


```
runstruct_vasp
```

 (making sure the `vasp.wrap` file indicates that all degrees of freedom must be relaxed).
- 2) Generation of the perturbations.
 - 2a) A typical command line is as follows:


```
fitfc -er=11.5 -ns=3 -ms=0.02 -dr=0.1
```

`-er` specifies how far apart the periodic images of the displaced atom must be. The code then finds the size of the supercell satisfying this constraint. Distances are measured according to the atomic positions given in `str.out` and in the same units.
 `-ns` specifies the number of different strain at which phonon calculations will be performed.
 (`-ns=1` implies a purely harmonic model, the default while values greater than 1 will invoke a quasiharmonic model)
 `-ms` specifies the maximum strain (0.02 signifies a 2% elongation along every direction).
 `-dr` the magnitude of the displacement of the perturbed atom.
 The above command writes out a series of subdirectories `vol_*`, one for each level of strain.
 If the structure has cubic symmetry or if you are willing to assume that thermal expansion is isotropic or if you only wish to use the harmonic approximation, the `fitfc` command should be invoked with the `-nrr` option (do Not ReRelax) and you can now skip to step 3).
 - 2b) Each volume subdirectory now contains a `str.out` file which is stretched version of the main `str_relax.out` file provided. You then need to run the ab initio code to rerelease the geometry at the various levels of imposed strain and obtain the energy as a function of strain. Typically, this is achieved by typing:


```
pollmach runstruct_vasp &
```

 (make sure that the `vasp.wrap` file is modified so that all degrees of freedom except volume are allowed to relax.)
 After this command each subdirectory will contain an energy and a `str_relax.out` file.
 Type

```
touch stoppoll
after all energies have been calculated.
The runstruct_vasp command can also be executed manually in each
subdirectory or as follows:
    foreachfile wait runstruct_vasp \; rm wait
```

- 2c) Now you need to reinvoke fitfc to generate perturbations of the atomic position for each level of strain.

```
fitfc -er=11.5 -ns=3 -ms=0.02 -dr=0.1
```

This is exactly the same command as before but the code notices the presence on the new files and can proceed further.

- 3) At this point the files generated are arranged as follows.

At the top level, there is one subdirectory per level of strain (vol_*, where * is the strain in percent), and in each subdirectory, there are a number of subsubdirectories, each containing a different perturbation. The perturbation names have the form p<+/-><dr>_<er>_<index>, where <pertmag> is the number given by the -dr option, <er> by the -er option and <index> is a number used to distinguish between different perturbations. Two perturbations that differ only by their signs are sometimes generated and are distinguished by a + or - prefix.

If you want to ensure that the third-order force constants cancel out exactly in the fit, you need to consider both perturbations.

Otherwise, only the 'positive' perturbation will be sufficient.

Note that whenever the third-order terms cancel out by symmetry, only the 'positive' perturbation will be generated.

You then need to use the ab initio code to calculate reaction forces for each perturbation.

This will typically be accomplished by typing

```
pollmach runstruct_vasp &
```

(make sure that the vasp.wrap file indicates that no degrees of freedom are allowed to relax and the smearing is used for Brillouin zone integration. Do not use the DOSTATIC option.)

- 4) Fitting the force constants and phonon calculations.

This mode is invoked with the -f option.

In addition, you need to specify the range of the springs included in the fit using the -fr=... option.

Usually, the range specified with -fr should be not more than half the distance specified with the -er option earlier.

Distances are measured according to the atomic positions given in str.out and in the same units.

It is a good idea to try different values of -fr (starting with the nearest neighbor bond length) and check that the vibrational properties converge as -fr is increased.

- 5) Sometimes you will get the message 'Unstable modes found. Aborting.'

This indicates that the structure considered may be mechanically unstable. If, in addition, you see the warning 'Warning: p... is an unstable mode.', then the structure is certainly unstable. Otherwise, it may be a artifact

of the fitting procedure.

To find out, you can tell the code to generate perturbations along the unstable directions and let your ab initio code calculate the reaction forces which can then be included in the fit to settle this issue.

First, to view the unstable modes, use the -fu option: the output is in vol_*/unstable.out and has the form:

```
u [index] [nb_atom] [kpoint] [branch] [frequency]
  [displacements...]
```

where [index] is a reference number, nb_atom is the size of the supercell needed to represent this mode (the other entries are self-explanatory).

(If this file contains only entries with nb_atom=too_large, you need to increase the -mau option beyond its default of 64.)

You can pick one of these modes to written out to disk with the option -gu=[index] where [index] is the one reported in the unstable.out file.

Using a negative index gives the sine instead of cosine phase of the mode.

You can run your ab initio code in the subdirectory generated (named vol_*/p_<uns_<dr>_<kmesh>_<number>) and rerun fitfc -f -fr=...

If you see 'Warning: p... is an unstable mode.' then you have found a true instability. If you only see 'Unstable modes found. Aborting.' you may repeat the process until the message disappear or a truly unstable mode is found.

Note: If you want to generate a phonon DOS even if there are unstable modes, use the -fn option. The unstable modes will be shown as negative frequencies.

-> Phonon Dispersion curves

The -df=inputfile option invokes the phonon dispersion curve module.

The syntax of the input file is:

```
[nb of points] [kx1] [ky1] [kz1] [kx2] [ky2] [kz2]
repeat...
```

Each line of input defines one segment (kx1,ky1,kz1)-(kx2,ky2,kz2) along which the dispersion curve is to be calculated.

[nb of points] specifies the number of points sampled along the segment.

The coordinates are in multiple of the reciprocal cell defined by the axes in the file specified by the -si option (or, by default, in the str.out file).

(The k-point coordinates are appropriately strained by the amount needed for the str.out file to be identical to the str_relax.out file.)

The phonon frequencies are output in the eigenfreq.out file, in the vol_* subdirectories.

Output files:

fitfc.log : A general log file.

vol_*/vdos.out : the phonon density of states for each volume considered

vol_*/fc.out : The force constants.

For each force constant a summary line gives:

(i) the atomic species involved

(ii) the 'bond length'

(iii) the stretching and bending terms

Then, each separate component of the force constant is given and, finally, their sum.

vol_*/svib_ht : gives the high-temperature limit of the vibrational entropy (in units of kB/atom) in the harmonic approximation, excluding the configuration-independent contribution at each unit cell volume considered (so, this just -3 times the

average log phonon frequency).
 vol_*/fvib : gives the harmonic vibrational free energy (in eV) at that volume
 as a function of temperature.

The following 3 files give the output of the quasiharmonic approximation if ns>1 and the harmonic approximation if ns=1:

fitfc.out : gives along each row, the temperature, the free energy,
 and the linear thermal expansion
 (e.g. 0.01 means that the lattice has expanded by 1%
 at that temperature).
 fvib : gives only the free energy
 svib : gives only the entropy (in energy/temperature, by default, eV/K)
 eos0.out : equation of state at 0K (one strain,energy pair per line)
 eos0.gnu : gnuplot script to plot polynomial & data

Caution:

When using fitfc as part of a cluster expansion, make sure you use the -me0 option to ensure that the static energy (included in eci.out) is not double-counted.

7.1.8 fitsvsl

This code determines bond stiffness vs bond length relationship for the purpose of calculating vibrational properties (with the svsl code).

It requires the following files as an input.

- 1) A lattice file (by default, lat.in, but this can be overridden with the -l option) which allows the code to determine what chemical bonds are present in the system.
 The format is as described in the maps documentation (see maps -h).
- 2) A list of directories containing structures that will be used to calculate force constants (by default strname.in, but this can be overridden with the -dn option).
 Each of the listed directory must contain
 - a) a str.out file containing an ideal unrelaxed structure that will be used to automatically determine the nearest neighbor shell,
 - b) a str_relax.out file containing the relaxed structure that will be used to calculate bond lengths and that the code will perturb in various ways to determine the force constants.

The code can operate in two modes: a structure generation mode and a fitting mode (indicated by the -f option).

In structure generation mode:

All the above input files are needed and the option
 -er must be specified in order to indicate the size of the supercells generated.
 The -er option indicates how far from each other a displaced atom must be from its periodic images, the code will infer the smallest supercell satisfying this constraint. Typically, -er should be 3 or 4 times the nearest neighbor distance. All of these distances are measured using the ideal structures (*str.out).

The following parameters have reasonable default values which can be overridden:

- dr specifies the displacement of the perturbed atom, which is 0.2 Angstrom by default.
- ms specifies the maximum (linear) expansion of the structures for the purpose of lengthing the bond length. For instance -ms=0.01 (the default) indicates that the

supercells will be stretched by up to 1% isotropically.
 -ns indicates the number of intermediate lattice parameters sampled (by default 2, which is the minimum in order to be able to determine the length dependence of bond stiffness).

After the structure generation step:

Each of the directory specified in strname.in will contain multiple subdirectories, each of which contains

- a) the ideal unrelaxed supercell in a str_ideal.out file.
- b) the relaxed but unperturbed supercell in a str_unpert.out file.
- c) the actual geometry of perturbed supercell calculation in a str.out file.

The appropriate first-principles calculations can be performed using the other utilities in the atat package, such as runstruct_vasp. Of course a corresponding vasp.wrap file must given in order to provide the input parameters for the first-principles calculations.

Make sure that these parameters indicate a static run (no relaxations!).

After all (or some) of these calculations are done, each subdirectory will contain

- a force.out file containing the forces acting on each atoms
- a str_relax.out file containing the atomic positions (in the same order as in force.out)

The fitting mode (-f option) of ftsvsl then needs to be used.

The lattice file (e.g. lat.in) must be present and the code will look for all files of the form */force.out and */*/force.out and the corresponding files */str*.out and */*/str*.out .

The code will then use that information to create the length-dependent force constants (this may take a few minutes) and outputs them in

slspring.out

Here is an example of the format of this file:

```
Al Al      (gives the type of bond)
  2        (2 parameters: linear fit is used)
50.28971   \ polynomial coefficients of the stiffness vs length relationship for stretching term
7.88973    / (typically, stiffness is in eV/Angstrom^2 and length is in Angstrom)
  2        \
6.12722    | idem for bending term
-1.01641   /
Ti Al      (repeat for each type of chemical bond)
  3
etc.
```

The only option controlling this process is -op, which specifies the order of the polynomial used to fit the length dependence (by default -op=1 and a linear fit is used).

(Contact the author for information about the -sf option.)

Diagnostic files are also output:

```
fitsvsl.log          (a log file)
fitsvsl.gnu and f_*.dat (to plot the s vs l relationship)
```

7.1.9 gce

->What does this program do?

- 1) It reads the lattice file (specified by the -l option). (See format below.)
- 2) It determines the space group of this lattice and writes it to the sym.out file.

- 3) It finds all symmetrically distinct clusters-'object' pairs that satisfy the conditions specified by the options -2 through -6.
 For instance, if -2=2.1 -3=1.1 is specified, only pair clusters shorter than 2.1 units and triplet clusters containing no pairs longer than 1.1 will be selected.
 This produces a generalized cluster expansion (GCE) representing a relationship between a property of type 'object' and a crystal structure.
 The 'object' can be anything obeying well-defined symmetry rules.
 By default the 'object' is a tensor of rank defined in the file gcetensor.in (see below for format).
- 4) It writes all clusters found to clusters.out.
 If the -c option is specified, clusters are read from clusters.out instead.
- 5) It reads the structure file (specified by the -s option).
- 6) It determines, for that structure, the correlations associated with all the clusters chosen earlier.
 This information is then output on one line, in the same order as in the clusters.out file. See below for conventions used to calculate correlations.
- 7) It writes the files corrdump.log containing the list of all adjustments needed to map the (possibly relaxed) structure onto the ideal lattice.

->File formats

Lattice and structure files

Both the lattice and the structure files have a similar structure.

First, the coordinate system a,b,c is specified, either as

[a] [b] [c] [alpha] [beta] [gamma]

or as:

[ax] [ay] [az]

[bx] [by] [bz]

[cx] [cy] [cz]

Then the lattice vectors u,v,w are listed, expressed in the coordinate system just defined:

[ua] [ub] [uc]

[va] [vb] [vc]

[wa] [wb] [wc]

Finally, atom positions and types are given, expressed in the same coordinate system as the lattice vectors:

[atom1a] [atom1b] [atom1c] [atom1type]

[atom2a] [atom2b] [atom2c] [atom1type]

etc.

In the lattice file:

-The atom type is a comma-separated list of the atomic symbols of the atoms that can sit the lattice site.

-The first symbol listed is assigned a spin of -1.

-When only one symbol is listed, this site is ignored for the purpose of calculating correlations, but not for determining symmetry.

-The atomic symbol 'Vac' is used to indicate a vacancy.

In the structure file:

- The atom type is just a single atomic symbol
(which, of course, has to be among the atomic symbols given in the
lattice file).
- Vacancies do not need to be specified.

Examples

The fcc lattice of the Cu-Au system:

```
1 1 1 90 90 90
0 0.5 0.5
0.5 0 0.5
0.5 0.5 0
0 0 0 Cu,Au
```

The Cu₃Au L1₂ structure:

```
1 1 1 90 90 90
1 0 0
0 1 0
0 0 1
0 0 0 Au
0.5 0.5 0 Cu
0.5 0 0.5 Cu
0 0.5 0.5 Cu
```

A lattice for the Li_x Co_y Al_(1-y) O₂ system:

```
0.707 0.707 6.928 90 90 120
0.3333 0.6667 0.3333
-0.6667 -0.3333 0.3333
0.3333 -0.3333 0.3333
0 0 0 Li,Vac
0.3333 0.6667 0.0833 0
0.6667 0.3333 0.1667 Co,Al
0 0 0.25 0
```

File format of gcetensor.in

[rank]

[list of pairs of indices indicating which simultaneous index permutations leave the tensor invariant]

[next list, etc...]

Examples

for strain or stress:

```
2
0 1
```

for elastic constants:

```
4
0 1
2 3
0 2 1 3
```

Symmetry file format (sym.out)

[number of symmetry operations]

3x3 matrix: point operation

1x3 matrix: translation

repeat, etc.

Note that if you enter more than one unit cell of the lattice, sym.out will contain some pure translations as symmetry elements.

Cluster file format (clusters.out)

for each cluster:

[multiplicity]

[length of the longest pair within the cluster]

[number of points in cluster]

[coordinates of 1st point] [number of possible species-2] [cluster function]

[coordinates of 2nd point] [number of possible species-2] [cluster function]

etc.

[keyword describing type 'object' in the GCE, for instance: tensor]

['object' data]

repeat, etc.

When the 'object' is a tensor the 'object' data is:

[rank]

[3's repeated 'rank' times]

[3^rank]

[elements of the tensor]

(Multiplicity and length are ignored when reading in the clusters.out file.)

For each 'point' the following convention apply

-The coordinates are expressed in the coordinate system given in the first line (or the first 3 lines) of the lat.in file.

-The 'number of possible species' distinguishes between binaries, ternaries, etc...

Since each site can accomodate any number of atom types, this is specified for each point of the cluster.

-In multicomponent system, the cluster function are numbered from 0 to number of possible species-2.

In the simple of a binary system [number of possible species-2] [cluster function] are just 0 0.

For a ternary, the possible values are 1 0 and 1 1.

All the utilities that are not yet multicomponent-ready just ignore the entries [number of possible species-2] [cluster function]

Convention used to calculate the correlations:

The cluster functions in a m-component system are defined as

function '0' : $-\cos(2\pi \cdot 1 \cdot s/m)$

function '1' : $-\sin(2\pi \cdot 1 \cdot s/m)$

.

.

.

$-\cos(2\pi [m/2] \cdot s/m)$

$-\sin(2\pi [m/2] \cdot s/m)$ <--- the last sin() is omitted if m is even

where the occupation variable s can take any values in {0,...,m-1}

and [...] denotes the 'round down' operation.

Note that, these functions reduce to the single function $(-1)^s$ in the binary case.

Special options:

```
-sym: Just find the space group and then abort.
-clus: Just find space group and clusters and then abort.
-z:   To find symmetry operations, atoms are considered to lie on
      top of one another when they are less than this much apart.
-sig: Number of significant digits printed.
```

->Cautions

When vacancies are specified, the program may not be able to warn you that the structure and the lattice just don't fit. Carefully inspect the corrdump.log file!

If the structure has significant cell shape relaxations, the program will be unable to find how it relates to the ideal lattice. The problem gets worse as the supercell size of the structure gets bigger.

There is no limit on how far an atom in a structure can be from the ideal lattice site. The program first finds the atom that can be the most unambiguously assigned to a lattice site. It then finds the next best assignment and so on. This is actually a pretty robust way to do this. But keep in mind that the -z option does NOT control this process.

7.1.10 gensqs

IMPORTANT NOTE: Please see mcsqs command for a better, easier-to-use SQS generator.

This code requires 3 input files:

- 1) A lattice file (by default lat.in) in the same format as for maps or corrdump.
- 2) A cluster file (by default clusters.out), as generated with the corrdump utility.
- 3) A target correlation file (by default tcorr.out) which contains the value of desired correlations for each of the clusters listed in the cluster file.

A typical caling sequence would be:

the following command can be used to generate the target correlation file tcorr.out

```
corrdump -noe -2=maxradius -rnd -s=conc.in > tcorr.out
```

```
# where maxradius is the length of the longest pair desired
# and where conc.in contains an (ordered) structure having the
# desired concentration.
# The geometry of the structure in the conc.in file is not crucial -
# only the average concentration on each sublattice will be used.
```

```
# CAUTION: Here, a 'sublattice' is a set of symmetrically equivalent point clusters.
# If your system contains multiple sublattices (as evidenced by multiple
# point clusters in clusters.out, make sure that your conc.in file sets
# the composition of each sublattice correctly! This can be verified
# by looking at the point correlations output.
```

```
#this looks for possible sqs of 8 atoms/ cell
gensqs -n=8 > sqs8.out
```

```
corrump -2=anotherradius -3=anotherradius -noe -s=sqs8.out
# this helps you decide which sqs is best based on other correlations
# associated with clusters (pairs and triplets) of diameter less than
# anotherradius.
```

Caution:

gensqs only generates structures containing exactly the number of atoms per unit cell specified by the
(If an SQS with a smaller unit cell exists, it will not be listed.)

If you give too many correlations to match, the code may not
output anything.

Finding an 8-atom sqs takes a few minutes, an 16-atom sqs, a few hours
and a 32-atom sqs, a few days!

The exact speed depends on the symmetry of the lattice and on your
computer.

7.1.11 maps

-> What does this program do?

It gradually constructs a increasingly more accurate cluster expansion.
A user-provided script running concurrently is responsible for notifying
maps when computer time is available. maps creates files describing
structures whose energy should be calculated. The user-provided script
sets up the runs needed to calculate the energy of these structures.
As maps becomes aware of more and more structural energies, it gradually
improves the precision of the cluster expansion, which is continuously
written to an output file.

The code terminates when a stop file is created by typing, for instance,
touch stop

NOTE: Fully functional scripts are included with the package:

pollmach and runstruct_vasp.

For for more information type

```
pollmach
runstruct_vasp -h
```

-> Format of the input file defining the lattice (specified by the -l option)

First, the coordinate system a,b,c is specified, either as

```
[a] [b] [c] [alpha] [beta] [gamma]
```

or as:

```
[ax] [ay] [az]
```

```
[bx] [by] [bz]
```

```
[cx] [cy] [cz]
```

Then the lattice vectors u,v,w are listed, expressed in the coordinate system just defined:

```
[ua] [ub] [uc]
[va] [vb] [vc]
[wa] [wb] [wc]
```

Finally, atom positions and types are given, expressed in the same coordinate system as the lattice vectors:

```
[atom1a] [atom1b] [atom1c] [atom1types]
[atom2a] [atom2b] [atom2c] [atom2types]
etc.
```

- The atom type is a comma-separated list of the atomic symbols of the atoms that can sit the lattice site.
- The first symbol listed is assigned a spin of -1 and the second, a spin of 1.
- When only one symbol is listed, this site is ignored for the purpose of calculating correlations, but not for determining symmetry.

Examples:

The fcc lattice of the Cu-Au system:

```
3.8 3.8 3.8 90 90 90
0 0.5 0.5
0.5 0 0.5
0.5 0.5 0
0 0 0 Cu,Au
```

A lattice for the $\text{Li}_x\text{Co}_y\text{Al}_{(1-y)}\text{O}_2$ system:

```
0.707 0.707 6.928 90 90 120
0.3333 0.6667 0.3333
-0.6667 -0.3333 0.3333
0.3333 -0.3333 0.3333
0 0 0 Li,Vac
0.3333 0.6667 0.0833 0
0.6667 0.3333 0.1667 Co,Al
0 0 0.25 0
```

Running the above example requires the multicomponent version of maps, called mmaps.

Optional input file: ref_energy.in

Contains the reference energy (per site) to be subtracted to get formation energies. The first line is the $c=0$ energy, the second line is the $c=1$ energy. If this file is omitted, the energies of leftmost and rightmost structures (on the concentration axis) are taken.

Optional input file: nbclusters.in

Allows the user to manually select which clusters to include in the fit.

This file should contains:

```
number of pairs to include
number of triplets to include
etc.
```

This file can be changed while maps is running. However, you must type touch refresh to tell maps to reread it.

-> Output files

maps.log

Contains possible warnings:

```
'Not enough known energies to fit CE'
'True ground states not = fitted ground states'
'New ground states predicted, see predstr.out'
```

These warning should disappear as more structural energies become available and the following messages should be displayed:

```
'Among structures of known energy, true and predicted ground states agree.'
'No other ground states of xx atoms/unit cell or less exist.'
```

This file also gives the crossvalidation score of the current fit (before the weighting is turned on in order to get the correct ground states).

fit.out

Contains the results of the fit, one structure per line and each line has the following information:

```
concentration energy fitted_energy (energy-fitted_energy) weight index
'concentration' lies between 0 and 1.
'energy' is per site (a site is a place where more than one atom type can lie)
'weight' is the weight of this structure in the fit.
'index' is the name of the directory associated with this structure.
```

predstr.out

Contains the predicted energy (per site) of all structures maps has in memory but whose true energy is unknown or has been flagged with error.

Format: one structure per line, and each line has the following information:

```
concentration energy predicted_energy index status
index is the structure number (or -1 if not written to disk yet).
energy is the calculated energy (or 0 if unknown).
status is either
    b for busy (being calculated),
    e for error,
    u for unknown (not yet calculated) or
    g if that structure is predicted to be a ground state (g can be combined with the above).
To list all predicted ground states, type
grep 'g' predstr.out
```

gs.out

Lists the ground state energies, one structure per line and each line has the following information:

```
concentration energy fitted_energy index
```

gs_str.out

Lists the ground state structures, in the same format as the n/str.out files (see below). Each structure is terminated by the word 'end' on a line by itself, followed by a blank line.

eci.out

Lists the eci. (They have already been divided by multiplicity.)
The corresponding clusters are in clusters.out

clusters.out

For each cluster, the first line is the multiplicity, the second line is the cluster diameter, and the third line is the number of points in the cluster. The remaining lines are the coordinates of the points in the cluster (in the coordinate system specified in the input file defining the lattice). A blank line separates each cluster.

n/str.out

Same format as the lattice file, except that

- The coordinate system is always written as 3x3 matrix
- Only one atom is listed for each site.

ref_energy.out

Reference energies used to calculate formation energies.

(Usually: energy of the pure end members OR values given in ref_energy.in if provided.)

The standard output reports the current progress of the calculations.

During the fit of the cluster expansion, each line of numbers displayed has the following meaning:

- 1) The current number of point, pair, triplet, etc. clusters
- 2) An indicator of whether the predicted ground states agree with the true ones (1) or not (0)
- 3) The CV score.

-> Communication protocol between maps and the script driving the energy method code (e.g. ab initio code)
(Only those who want to customize the code need to read this section.
The scripts described in this section are provided with the atat distribution in the glue/ subdirectory.)

Unless otherwise specified all files mentioned reside in the directory where maps was started. All paths are relative to the startup directory.

+The script should first wait for computer time to be available before creating a file called 'ready'.
-Upon detecting that the 'ready' file has been created, maps responds by creating a subdirectory 'n' (where 'n' is a number) and a file 'n/str.out' containing a description of a structure whose energy needs to be calculated.
-maps creates a file called 'n/wait' to distinguish this directory from other ones created earlier.
-maps deletes the 'ready' file.
+Upon detecting that the 'ready' file has disappeared, the script should now look for the 'n/wait' file, start the calculations in the directory 'n' and delete file 'n/wait'.
+If anything goes wrong in the calculations, the script should create a file 'n/error'.
+When the calculations terminate successfully, the energy per unit cell of the structure should be copied to the file 'n/energy'.
(NOTE: use energy per unit cell of the structure NOT per unit cell of the lattice).
-maps continuously scans all the subdirectories 'n' for 'n/energy' or 'n/error' files and updates the cluster expansion accordingly.
-maps updates the cluster expansion whenever a file called 'refresh' is created (maps then deletes it).
-maps terminates when a 'stop' file is created.

Note that the script can ask maps to create new structure directories even before the energy of the current structure has been found.

Note that human intervention is allowed: an 'n/error' file can be manually created if an error is later found in a run.

Users can also manually step up all runs if they wish so, as long as they follow the protocol.

Example of script

(portions in `/* */` have to be filled in with the appropriate code):

```
#!/bin/csh

while (! -e stop)
  /* check machine load here */
  if ( /* load low enough */ ) then
    touch ready
    while (-e ready)
      sleep 30
    end
    cd 'ls */wait | sed 's+/.+++g' | head -1'
    rm -f wait
    /* convert str.out to the native format of ab initio code */
    /* in background: run code and create either energy file or error file */
    cd ..
  endif
  sleep 180
end
```

-> Using maps with vasp

The script `runstruct_vasp`, when run from within directory 'n',

- 1) converts 'vasp.wrap' and 'n/str.out' into all the necessary files to run vasp,
- 2) runs vasp
- 3) extract all the information from the output files and writes in a format readable by maps.

An example of `vasp.wrap` is:

```
[INCAR]
PREC = high
ISMear = -1
SIGMA = 0.1
NSW=41
IBRION = 2
ISIF = 3
KPPRA = 1000
DOSTATIC
```

See `ezvasp` documentation for more information.

-> Importing structures into maps

MAPS continuously scans all the first-level subdirectories containing a file called `str.out` and tries to map them onto superstructures of the lattice provided. This lets you 'import' structures from another source

into MAPS. A word of caution: the imported structures must be unrelaxed and no effort is made to rotate or scale them in order to match the lattice (aside from space group symmetry operations).

7.1.12 mcsqs

This code uses a Monte Carlo algorithm to find a Special Quasirandom Structure (SQS). In writing this code, Axel van de Walle benefited from the input of Alexey Dick, Dongwon Shin and Yi Wang.

If you use this code in a publication, please cite:

A. van de Walle, P. Tiwary, M. de Jong, D.L. Olmsted, M. Asta, A. Dick, D. Shin, Y. Wang, L.-Q. Chen, Z. Efficient stochastic generation of special quasirandom structures, *Calphad Journal* 42, pp. 13-18 (2013)

-> This code requires 2 input files:

- 1) A file defining the random state (by default `rndstr.in`) in a format similar to the `lat.in` that is needed for `maps` or `corrump` but with partial occupation of the sites (see below).
- 2) A cluster file (by default `clusters.out`), generated, for instance, with the command line:
`mcsqs -2=... -3=... etc.`
 where `-2=... -3=...` indicate the range of pairs, triplets etc.
 The code display the clusters found (format: nb of points, diameter, multiplicity).

Internally, this actually calls the command

```
corrump -l=rndstr.in -ro -noe -nop -clus -2=... -3=... ; getclus
```

Note that the (new) `-ro` option allows `corrump` to read the same input file as `mcsqs` (here `rndstr.in`).

The `-noe` and `-nop` skip the empty and point clusters that are not used by `mcsqs`

`-clus` indicates to generate clusters only

`-2=... -3=...` indicate the range of pairs, triplets etc.

`getclus` just indicates to write out the clusters (nb of points, length, multiplicity)

See `corrump -h` for more info.

-> Output files:

`bestsqs.out`

The best SQS found so far (in standard ATAT structure file format - see `mmaps -h` for a description)

`bestcorr.out`

For each of the clusters selected,

its number of pointd (1st column)

its diameter (2nd column)

the correlations of the best SQS found so far (3rd column)

along with the target disordered state correlation (4th column)

and the difference between the two (5th column).

`rndstrgrp.out`

A file containing the same info as the input file defining the random state (`rndstr.in`),

but with symmetrically equivalent sites grouped together and separated by blank lines.

(This helps determine which sites can have the same occupations.)

`mcsqs.log`

A log file.

The code stops if a perfect match is found (all correlations requested match the disordered state), but this may never happen if there are too many clusters in `clusters.out` or `-n` (the number of atoms) is too small. In any case, the `bestsqs.out` and `bestcorr.out` always contain the best solution found so far. Stopping the code prematurely if the solution is satisfactory is fine.

-> Optional files:

Creating a file called stopsqs stops the code cleanly.

Creating a file called sqsparam.in (or as specified by the -pf option) with 4 numbers in it allows you to set the -wr, -wn, -wd and -T options during runtime. (for backward compatibility, if 2 numbers are specified, they set the -wr and -T options while the remaining options are set to their defaults.)

The -rc option lets you specify supercells to use (via a sqscell.out file). Useful if brute force enumeration takes too long.

The sqscell.out usually comes from a previous mcsqs run but you can create it yourself. Note that the sqscell.out file must start with the number supercells you will provide and that supercells must be expressed in multiple of the axes defined in the rndstr.in file (in particular, all numbers entered would typically be integers).

[If your are using a version prior to 3.07, these should be entered in cartesian coordinates.]

-> Format of the input file defining the lattice (specified by the -l option)

First, the coordinate system a,b,c is specified, either as

[a] [b] [c] [alpha] [beta] [gamma]

or as:

[ax] [ay] [az]

[bx] [by] [bz]

[cx] [cy] [cz]

Then the lattice vectors u,v,w are listed, expressed in the coordinate system just defined:

[ua] [ub] [uc]

[va] [vb] [vc]

[wa] [wb] [wc]

Finally, atom positions and types are given, expressed in the same coordinate system as the lattice vectors:

[atom1a] [atom1b] [atom1c] [atomtype11]=[occupation11],[atomtype12]=[occupation12],...

[atom2a] [atom2b] [atom2c] [atomtype21]=[occupation21],[atomtype22]=[occupation22],...

etc.

Examples:

The fcc lattice of the Cu-Au system; request for an SQS at composition 0.5:

3.8 3.8 3.8 90 90 90

0 0.5 0.5

0.5 0 0.5

0.5 0.5 0

0 0 0 Cu=0.5,Au=0.5

A lattice for the Li_x Co_y Al_(1-y) O₂ system; request for an SQS with different composition on each sublattice:

0.707 0.707 6.928 90 90 120

0.3333 0.6667 0.3333

-0.6667 -0.3333 0.3333

0.3333 -0.3333 0.3333

0 0 0 Li=0.75,Vac=0.25

0.3333 0.6667 0.0833 0

0.6667 0.3333 0.1667 Co=0.25,Ni=0.25,Al=0.5

0 0 0.25 0

-> The objective function used is:

$$\frac{(\text{sum (absolute difference from the disordered state correlations)} * \exp(-\text{wd} * (\text{diameter of cluster})) * (\text{wn}^{(n)}))}{(\text{normalization for weights to sum up to 1 over clusters bigger than the smallest nonmatching cluster})}$$

-sum over p wr* wn^(p-2)*(diameter of the smallest cluster of p points of less whose correlation does not match)

The wr, wn and wd are set through the -wr, -wn and -wd options respectively (or as the 3 first numbers of the input file). The diameter of a cluster is the length of the longest pair contained in a cluster (normalized by nearest neighbor distance).

See routine calc_objective_func in mcsqs.c++ for a more precise description.

This objective function reflect researchers' desire to perfectly match all correlation up to some range. Adding the absolute difference serves to guide the algorithm in the right direction to extend the range of perfect match.

This objective function is a generalization of the one found in the Calphad paper cited above (reduces to the original one for small clusters).

-> Post-processing

If you have obtained an SQS and would like to see how good/bad the correlations you have NOT included in the objective function are, you can use a command of the form:

```
corrump -l=rndstr.in -ro -noe -2=... -3=... -s=bestsqs.out
and compare to the output (for a perfectly random solid solution) of
corrump -l=rndstr.in -ro -noe -2=... -3=... -s=bestsqs.out -rnd
```

Beware that the above commands overwrite the clusters.out file.

Make sure to NOT include the -nop option (which is incompatible with the -rnd option).

The getclus command is useful to get a summary of what the clusters are (number of points, diameter, multiplicity).

7.1.13 memc2

MultiComponent Easy Monte Carlo Code (emc2)
by Axel van de Walle

Command line parameters:

- er : enclosed radius. The Monte Carlo cell will be the smallest possible supercell of unit cell of the initial configuration such that a sphere of that radius fits inside it. This allows you to specify the system size in a structure-independent fashion.
- cf : Control file specifying the ranges of temperature and chemical potentials scanned (default: control.in). See file format below.
- eq : number of equilibration passes. (Equilibration is performed at the beginning of each step of the outer loop.)
- n : number of Monte Carlo passes performed to evaluate thermodynamic quantities at each step of the inner loop.

- tp : instead of specifying -eq and -n, you can ask the code to equilibrate and run for a time such that the average energy is accurate within the target precision specified by -tp=.
- aq : Alternative quantity that must meet the tolerance specified by -tp.
0: energy (default), 1: long-range order, 2-: concentrations, correlations
- gs : Gives the index of the ground state to use as an initial spin configuration (starting at 0). The ground states are listed in gs_str.out.
If the index is -1, the disordered state with equiatomic composition is used.
(If -gs=-1, the LTE and MF approximations are not calculated.
If you want the LTE and MF outputs, use one of the 'pure' end members as a starting point.)
- tstat : Critical value of the test for discontinuity. The code is able to catch phase transformations (most of the time) when going from an ordered phase to another or to the disordered state. This involves a statistical test which a user-specified confidence level. The default, 3, corresponds to a 0.4% chance of finding a transition where there is none. (Refer to a standard normal table.) -tstat=0 turns off this option. Suggestion: if a phase transition is undetected, first try to reduce temperature or chem. pot. steps or increase n or decrease tp before toying around with this parameter. Also: beware of hysteresis.
- sigdig : Number of significant digits printed. Default is 6.
- o: Name of the output file (default: mc.out).
- k : Sets boltzman's constant (default k=1). This only affects how temperatures are converted in energies. -k=8.617e-5 lets you enter temperatures in kelvins when energies are in eV.
(You can also select this value by using the -keV option.)
- mft : Mean field threshold. If the grand canonical potential obtained via the mean field approximation and the low temperature expansion are less than this threshold, Monte Carlo simulation are skipped and mean field values are used instead. (They replace the MC values in the output file.
The column labelled use_mf is equal to 1 when this substitution is made.
Note: When use_mf=1, the correlations in the output file are incorrectly set to their values in an ideal fully ordered structure.
All other columns are reliable.
- mftq: Quantity that must meet the tolerance specified by -mft.
0: phi (default) , 1: energy , 2: long-range order, 3-: concentrations

Tricks:

To read parameters from a file, use:
memc2 'cat inputfile'
where inputfile contains the commands line options.

To selectively display a few of the output quantities, use:
`memc2 [options] | cut -f n1,n2,n3,...`
 where `n1,n2,n3,...` are the column number desired (see below).

-> Input files:

`control.in` : range of temperature and chemical potentials to scan.

The first line of this file specifies the initial conditions and has the format:

`temperature chemical_potentials_1 ... chemical_potentials_n`

where the chemical potentials are for each specie.

Each subsequent line of this file indicates one of the axes along which to scan and has the format:

`temperature chemical_potentials_1 ... chemical_potentials_n number_of_steps`

where `number_of_steps` is the the number of steps made between the initial conditions and the final conditions given on the line.

Example: to scan the region in (T,mu1,mu2,mu3)-space defined by

`100 <= T < 200 and 0 <= mu1 < 1.0 and 0 <= mu2 < 0.5 and mu3 == 0.0`

with a 10x5x5 grid

the `control.in` file should be

`100 0.0 0.0 0.0 0.0`

`200 0.0 0.0 0.0 0.0 10`

`100 0.0 0.5 0.0 5`

`100 1.0 0.0 0.0 5`

Note: Temperatures are given in units of energy unless the `-k` or `-eV` options are set.

By default, the finals conditions are excluded from the scan (in the example above the chemical potentials scanned in the last line are 0.0 0.2 0.4 0.6 0.8)

This behavior can be changed with the `-il` option, to give: 0.0 0.25 0.5 0.75 1.0)

Alternatively, the `-hf` option gives: 0.1 0.3 0.5 0.7 0.9 .

NOTE: The `mmaps` code generates a file called `chempot.out` which contains special values of the chemical potential that stabilize various types of equilibria. These values are useful guidelines to select relevant regions in `mu`-space to scan.

-> Other input files:

`lat.in` : description of the lattice.

`clusters.out` : describes the clusters.

`eci.out` : provides the ECI.

`gs_str.out` : a list of ground states, in no particular order.

These 4 files can be created by `maps`.

See `maps` documentation (`maps -h`) for a description of the formats.

-> Optional input files:

`teci.out` : if present, provides temperature-dependent `eci` (overrides `eci.out`)
 (Note that, even when the `teci.out` file is used, column 6 of the output file reflects only the configurational contribution to the heat capacity.)

The format this file is:

[maximum temperature: `Tm`]

[number of temperatures where `eci` are provided: `nT`]

[ECIs for temperature 0]

[ECIs for temperature `Tm/(nT-1)`]

```
[ECIs for temperature 2*Tm/(nT-1)]
...
[ECIs for temperature Tm]
```

Note that these numbers can be separated by blanks or newlines, as desired.

conconcs.in : Specifies linear constraints on the composition that spin flips must obey.
 (The code will generate multi-spin flips if necessary.)
 Here is an example of such file:
 $1.0*Al - 1.0*Li + 0.1*Co = 0.5$
 Please use rational numbers only - infinite loop will result otherwise.
 Composition is determined by the user-supplied initial configuration
 (via the -g or -is options).

-> Format of the output file

The file mheader.out gives the content of column of the output file.
 The following abbreviations are used:

```
T:      temperature
mu(A):  chemical potentials of specie A
x(A):   concentration of specie A
E:      energy (per active site)
Egc:    grand canonical energy (per active site) , i.e.,  $E - \sum_i \mu(i)*x(i)$ 
lro:    long Range Order parameter of the initial ordered phase
        (=0 if initial phase is disordered)
F:      Helmholtz free energy (per active site)
phi:    grand canonical potential:  $F - \sum_i \mu(i)*x(i)$ 
use_mf: flag that is 1 if mf values have been used instead of mc values
corr_n_d: the average correlations associated with each cluster
          n is the number of points in the cluster
          d is the diameter of the cluster
```

Suffixes:

```
_lte: obtained with low temperature expansion
_mf:  obtained with mean field approximation
_mc:  obtained with Monte Carlo simulations
```

NOTE: to obtain 'canonical' rather than grand canonical quantities (adding $\mu*x$),
 to all (free) energies use the -g2c option.

NOTE: an 'active site' is one that can host more than one possible atom type.
 Energy and free energy are reported per 'active site' not per total sites.

7.1.14 mmap

-> What does this program do?

This is the multicomponent version of the maps code.

It gradually constructs a increasingly more accurate cluster expansion.
 A user-provided script running concurrently is responsible for notifying
 maps when computer time is available. maps creates files describing
 structures whose energy should be calculated. The user-provided script
 sets up the runs needed to calculate the energy of these structures.

As maps becomes aware of more and more structural energies, it gradually improves the precision of the cluster expansion, which is continuously written to an output file.

The code terminates when a stop file is created by typing, for instance, touch stop

NOTE: Fully functional scripts are included with the package:

pollmach and runstruct_vasp.

For more information type

```
pollmach
runstruct_vasp -h
```

-> Format of the input file defining the lattice (specified by the -l option)

First, the coordinate system a,b,c is specified, either as

```
[a] [b] [c] [alpha] [beta] [gamma]
```

or as:

```
[ax] [ay] [az]
```

```
[bx] [by] [bz]
```

```
[cx] [cy] [cz]
```

Then the lattice vectors u,v,w are listed, expressed in the coordinate system just defined:

```
[ua] [ub] [uc]
```

```
[va] [vb] [vc]
```

```
[wa] [wb] [wc]
```

Finally, atom positions and types are given, expressed in the same coordinate system as the lattice vectors:

```
[atom1a] [atom1b] [atom1c] [atom1types]
```

```
[atom2a] [atom2b] [atom2c] [atom2types]
```

etc.

-The atom type is a comma-separated list of the atomic symbols of the atoms that can sit the lattice site.

-When only one symbol is listed, this site is ignored for the purpose of calculating correlations, but not for determining symmetry.

Examples:

The fcc lattice of the Cu-Au system:

```
3.8 3.8 3.8 90 90 90
```

```
0 0.5 0.5
```

```
0.5 0 0.5
```

```
0.5 0.5 0
```

```
0 0 0 Cu,Au
```

A lattice for the Li_x Co_y Al_(1-y) O₂ system:

```
0.707 0.707 6.928 90 90 120
```

```
0.3333 0.6667 0.3333
```

```
-0.6667 -0.3333 0.3333
```

```
0.3333 -0.3333 0.3333
```

```
0 0 0 Li,Vac
```

```
0.3333 0.6667 0.0833 0
```

```
0.6667 0.3333 0.1667 Co,Al
```

```
0 0 0.25 0
```

Optional input file: `ref_energy.in`

Contains the reference energy (per active site, i.e. those that can host more than one specie) to be subtracted to get formation energies.

The atomic reference energies must be in the same order as in the `atoms.out` file.

If this file is omitted, the energies of the structures with the most extreme compositions are used to determine the reference energies, which are then output to the `ref_energy.out` file.

Optional input file: `nbclusters.in`

Allows the user to manually select which clusters to include in the fit.

This file should contains:

number of pairs to include
number of triplets to include
etc.

This file can be changed while `maps` is running. However, you must type `touch refresh` to tell `maps` to reread it.

Optional input file: `crange.in` (or as specified by the `-cr` option)

If present, this file

selects the range of concentration over which the cluster expansion is to be fitted. This controls both where the correct ground states are enforced in the fitting process and the range of concentration of the generated structures. Occasionally, a structure outside that range is generated, to verify the ground state hull.

Here is an example of such file:

```
1.0*Al -1.0*Li +0.1*Co >= 0.5
```

Multiple constraints can be listed (on separate lines).

Make sure to include a numerical prefactor for each species even if it is 1.0. Do not put a space between '-' and a number.

Note that the `crange.in` file is read (if present) even if the `-cr` option is not specified.

Optional input file: `weights.in`

Forces specific weights to assign to each structure when fitting its energy.

These weights may be updated by the code internally to better reproduce the ground state line.

Format: one weight per line, in the same order as in `fit.out`

This should only be used to fine-tune the fit at the end when no new structures will be added (since `weights.in` would have to change every time a new structure is added).

-> Output files

`maps.log`

Contains possible warnings:

```
'Not enough known energies to fit CE'
'True ground states not = fitted ground states'
'New ground states predicted, see predstr.out'
```

These warning should disappear as more structural energies become available and the following messages should be displayed:

```
'Among structures of known energy, true and predicted ground states agree.'
'No other ground states of xx atoms/unit cell or less exist.'
```

This file also gives the crossvalidation score of the current fit (before the weighting is turned on in order to get the correct ground states).

atoms.out

Lists all atomic species given in the input files.

fit.out

Contains the results of the fit, one structure per line and each line has the following information:

concentration energy fitted_energy (energy-fitted_energy) weight index
'concentration' a vector of the atom fraction of all species (in the same order as in atoms.out).

'energy' is per site (a site is a place where more than one atom type can sit)

'weight' is the weight of this structure in the fit.

'index' is the name of the directory associated with this structure.

predstr.out

Contains the predicted energy (per site) of all structures maps has in memory but whose true energy is unknown.

Format: one structure per line, and each line has the following information:

concentration predicted_energy index status
index is the structure number (or -1 if not written to disk yet).
status is either

b for busy (being calculated),

e for error,

u for unknown (not yet calculated) or

g if that structure is predicted to be a ground state (g can be combined with the above).

To list all predicted ground states, type

grep 'g' predstr.out

gs.out

Lists the ground state energies, one structure per line and each line has the following information:

concentration energy fitted_energy (energy-fitted_energy) index

gs_connect.out

Indicates which ground states touch each face of the ground state convex hull.

gs_str.out

Lists the ground state structures, in the same format as the n/str.out files (see below). Each structure is terminated by the word 'end' on a line by itself, followed by a blank line.

chempot.out

This file contains

(i) the values of the chemical potentials that stabilize all fixed-composition multiphase equilibria of the system at OK.
(e.g. in a n-nary system, all the n-phase equilibria)

(ii) values of the chemical potentials that stabilize each of the ground states at OK.

This data is useful to set up Monte Carlo simulations with the memc2 code.

eci.out

Lists the eci. (They have already been divided by multiplicity.)

The corresponding clusters are in clusters.out

clusters.out

For each cluster, the first line is the multiplicity, the second line is the cluster diameter, and the third line is the number of points in the cluster. The remaining lines are the coordinates of the points in the cluster (in the coordinate system specified in the input file defining the lattice). A blank line separates each cluster.

n/str.out

Same format as the lattice file, except that
 -The coordinate system is always written as 3x3 matrix
 -Only one atom is listed for each site.

ref_energy.out

Reference energies used to calculate formation energies.
 (Usually: energy of the pure end members OR values given in ref_energy.in if provided.)

The standard output reports the current progress of the calculations.

During the fit of the cluster expansion, each line of numbers displayed has the following meaning:

- 1) The current number of point, pair, triplet, etc. clusters
- 2) An indicator of whether the predicted ground states agree with the true ones (1) or not (0)
- 3) The CV score (per atom, as of version 2.88; per cell earlier).

-> Communication protocol between maps and the script driving the energy method code (e.g. ab initio code)
 (Only those who want to customize the code need to read this section.
 The scripts described in this section are provided with the atat distribution in the glue/ subdirectory.)

Unless otherwise specified all files mentioned reside in the directory where maps was started. All paths are relative to the startup directory.

+The script should first wait for computer time to be available before creating a file called 'ready'.
 -Upon detecting that the 'ready' file has been created, maps responds by creating a subdirectory 'n' (where 'n' is a number) and a file 'n/str.out' containing a description of a structure whose energy needs to be calculated.
 -maps creates a file called 'n/wait' to distinguish this directory from other ones created earlier.
 -maps deletes the 'ready' file.
 +Upon detecting that the 'ready' file has disappeared, the script should now look for the 'n/wait' file, start the calculations in the directory 'n' and delete file 'n/wait'.
 +If anything goes wrong in the calculations, the script should create a file 'n/error'.
 +When the calculations terminate successfully, the energy per unit cell of the structure should be copied to the file 'n/energy'.
 (NOTE: use energy per unit cell of the structure NOT per unit cell of the lattice).
 -maps continuously scans all the subdirectories 'n' for 'n/energy' or 'n/error' files and updates the cluster expansion accordingly.
 -maps updates the cluster expansion whenever a file called 'refresh' is created

(maps then deletes it).
 -maps terminates when a 'stop' file is created.

Note that the script can ask maps to create new structure directories even before the energy of the current structure has been found.

Note that human intervention is allowed: an 'n/error' file can be manually created if an error is later found in a run.

Users can also manually step up all runs if they wish so, as long as they follow the protocol.

Example of script

(portions in /* */ have to be filled in with the appropriate code):

```
#!/bin/csh

while (! -e stop)
  /* check machine load here */
  if ( /* load low enough */ ) then
    touch ready
    while (-e ready)
      sleep 30
    end
    cd 'ls */wait | sed 's+/.+++g' | head -1'
    rm -f wait
    /* convert str.out to the native format of ab initio code */
    /* in background: run code and create either energy file or error file */
    cd ..
  endif
  sleep 180
end
```

-> Using maps with vasp

The script runstruct_vasp, when run from within directory 'n',

- 1) converts 'vasp.wrap' and 'n/str.out' into all the necessary files to run vasp,
- 2) runs vasp
- 3) extract all the information from the output files and writes in a format readable by maps.

An example of vasp.wrap is:

```
[INCAR]
PREC = high
ISMear = -1
SIGMA = 0.1
NSW=41
IBRION = 2
ISIF = 3
KPPRA = 1000
DOSTATIC
```

See ezvasp documentation for more information.

-> Importing structures into maps

MAPS continuously scans all the first-level subdirectories containing a file called `str.out` and tries to map them onto superstructures of the lattice provided. This lets you 'import' structures from another source into MAPS. A word of caution: the imported structures must be unrelaxed and no effort is made to rotate or scale them in order to match the lattice (aside from space group symmetry operations).

7.1.15 phb

This is Monte Carlo code which automatically follows a given phase boundary.

Input files: (see maps documentation)

<code>lat.in</code>	(describes the lattice)
<code>gs_str.out</code>	(lists the ground states)
<code>eci.out</code>	(ECI)
<code>clusters.out</code>	(clusters)

You have to provide, on the command line, the following parameters:

-The two phases which are in equilibrium.

For instance,

`-gs1=0 -gs2=1`

If there are n ground states, phases are numbered from 0 to $n-1$.

These ground states are read in from `gs_str.out`

The disordered state is labelled by the number -1.

If the two phases are on different lattices, you need to specify the path that give access to the files

<code>lat.in</code>
<code>gs_str.out</code>
<code>eci.out</code>
<code>clusters.out</code>

for each lattice. For instance

`-d1=../fcc/ -d2=../hcp/`

If either or both options are omitted, the files are read from the current directory.

-The starting temperature and chemical potential

for instance,

`-T=100 -mu=0.125`

(Make sure to set blatzmann's constant appropriately.)

(If you tracing the phase boundary between two ordered phases, starting from OK, you do not need to specify a starting T and μ .)

-The temperature step

for instance `-dT=50`

-The 'enclosed radius', which sets the system size

for instance, `-er=35`

(see `emc2` documentation for more information)

-The precision of the calculation

This is expressed as the desired precision of the average concentration.

For instance, `-dx=1e-3`

The code automatically finds the equilibration time and the number Monte Carlo steps needed to obtain the target standard deviation of the average concentration.

There are a number of optional parameters as well.

`-ltep`: The low temperature expansion is used to find the free energy at low temperature. `-ltep` gives the maximum error allowed before Monte Carlo is used instead of LTE.

`-dmu`: The step in chemical potential used when scanning in search of the phase boundary.

for instance, `-dmu=0.005`

Sometimes, the algorithm loses track of the phase boundary (because of statistical errors). When that happens, it scans a range of values of the chemical potential in search of the boundary of the hysteresis loop associated with the first order transition of interest. It then positions itself in the middle of it. `dmu` is the step size used for that search. Note that the code will automatically shrink `dmu` if needed. By default, `dmu` is automatically set to the formation energy of a disordered alloy times 0.01.

`-mug` lets you specify a small difference in chemical potential between the phases, to make the code less sensitive to accidental phase transition.

`-k` sets Boltzmann's constant (see `emc2` documentation)

`-keV`

`-dn` indicates that the boundary must be followed downward (decreasing `T`)

Output file:

Column	Value
1	temperature
2	chemical potential
3	concentration of phase 1 in $[-1,1]$
4	concentration of phase 2 in $[-1,1]$
5	energy of phase 1
6	energy of phase 2

7.1.16 `predcs`

This code fits a reciprocal-space cluster expansion of the constituent strain energy for binary systems with cubic symmetry.

USAGE:

`csfit` needs, as an input, 4 files.

- 1) A `lat.in` file defining the geometry of the lattice (same format as `maps`).
- 2) Two `str_relax.out` residing in given directories (`-pa` and `-pb` options, or, by

- default 0/ and 1/) and providing the relaxed geometry of the two pure end members for the system.
- 3) A `dir.in` file listing the directions along which to compute epitaxial deformation energy in order to fit the expansion. Directions are specified as miller indices in the coordinate system defined in `lat.in` . Each index must be separated by a space and each direction must be on a separate line.

The code then generates subdirectories (in 0/ and 1/ or any other directory you specify) with deformed structures whose energy will be used to fit the expansion. The `-np` and `-nl` options let you control the number of structures generated: `-np` provides the number of intermediate epitaxial strain (perpendicular to specified directions) considered between the lattice parameters of the 2 pure elements while `-nl` provides the number of different values of strain along the directions specified. The range of strains spanned is given by the `-ml` option (+/- that value, 0 strain being the strain that keeps volume constant). The `-ns` option controls the numerical energy minimization (it is the number of mesh points used in the search for a minimum) and has no effect of the number of generated structures.

All default values are very reasonable. You may want to increase `-ml` and `-nl` for systems with a large size mismatch.

The files generated are compatible with the 'pollmach' automatic job control utility. If you use vasp, you would typically type

```
csfit -d &
pollmach runstruct_vasp -w csvasp.wrap &
```

where the `csvasp.wrap` contains the parameters needed for a vasp run that does not relax the structure geometry (see `atat/examples` directory).

WARNING: you are free to rerun `csfit` many times, adding new lines in `dir.in` but you CANNOT rerun it with different `-ml` `-np` `-nl` settings without first deleting the subdirectories generated.

The `csfit` programs waits until all calculations are done and then fits the expansion and writes the results to the `cs.in` file. The `cs.log` contains the constituent strain as a function of concentration (rows) for long period superlattices along the specified directions (columns).

You can use the resulting expansion in maps by using the `-p=cs` option and in `emc2` or `phb` by using the `-ks=cs` option.

CONVENTIONS:

We expand the concentration-dependent reciprocal space ECI associated with constituent strain as:

$$J_{\{CS\}}(x,k) = \sum_{l=0,2,4,\dots} a_{\{l\}}(x) K_{\{l\}}(k)$$

where $|K_{\{l\}}(k)|^2$ is normalized to integrate to one over the unit spherical shell $|k|=1$.

By combining Equations (12) and (19) in V. Ozolins, C. Wolverton, and A. Zunger, Phys. Rev. B, 57, 6427 (1998), we see that the coefficients $a_{\{l\}}(x)$ are related to the $c_{\{l\}}(x)$ coefficients defined in this paper through:

$$a_{\{l\}}(x) = c_{\{l\}}(x) / (4x(1-x))$$

The configuration-dependent constituent strain energy is given by

$$\Delta E_{\{CS\}}(\sigma) = \sum_{\{k\}} \Delta J_{\{CS\}}(x, k) |S(\sigma, k)|^2 \exp(-|k|^2/k_c^2)$$

(see Equation (22) in C. Wolverton, V. Ozolins, A. Zunger, J. Phys: Condens. Matter, 12, 2749 (2000)).

The $S(\sigma, k)$ is computed according to the following convention:

$$S(\sigma, k) = \frac{\sum_{\{j \text{ in unit cell of structure} \}} S_j \exp(i 2 \pi k \cdot R_j) * (\text{number of atom in unit cell of parent lattice})}{(\text{number of atom in unit cell of structure})}$$

$\Delta E_{\{CS\}}(\sigma)$ is thus given per unit cell of the parent lattice.

$1/k_c$ is given in the same unit of length as in the lat.in file (for instance, Angstroms).

The output file cs.in contains:

```
[1/k_c, set to 0 to turn off 'attenuation']
[Number of kubic harmonic to use, e.g. 2 to use K_0 and K_4]
[Number n of mesh point in concentration grid, including x=0 and x=1]

a_{0}(0)
a_{0}(1/(n-1))
a_{0}(2/(n-1))
.
.
.
a_{0}(1)

a_{4}(0)
a_{4}(1/(n-1))
a_{4}(2/(n-1))
.
.
.
a_{4}(1)

a_{6}(0)
a_{6}(1/(n-1))
a_{6}(2/(n-1))
.
.
.
a_{6}(1)

etc.
```

Additional output files

cs.log

Contains the constituent strain energy (cse) of superstructures along the specified directions as a function of concentration.

Format:

[concentration] [cse along direction 1 of dir.in] [cse along direction 2 of dir.in] etc.
repeat for each concentration

csdebug.out

Contains the raw energies for each calculation.

The outer loop is on the directions of the dir.in file.

The middle loop is on the element (which pure end member).

The inner loop is on the stretching perpendicular to the k-vector.

Each line contains the energy for various amount of stretching along the direction parallel to the k-vector.

7.1.17 svsl

This code calculates the vibrational free energy of a structure using the Stiffness VS Length method.

Before using this code, you will probably first need to use the fitsvsl code, which generates the length-dependent force constants that the present needs as an input.

It requires, as an input, 3 files:

- 1) A 'relaxed' structure (the default file name is str_relax.out, but it can be overridden with the -rs option).
This provides the actual atomic positions used in the calculations.
The format of this file is as described in the maps documentation (see maps -h).
- 2) An 'unrelaxed' structure (the default file name is str.out, but it can be overridden with the -us option).
This provides the ideal atomic positions that are used to automatically determine which atoms lie in the nearest neighbor shell. This file can be the same as the relaxed structure but then the determination of the nn shell may be less reliable.
- 3) A spring constant file (the default is to lookup in slspring.out and ../slspring.out but this can be overridden with the -sp option).
This file provides the bond stiffness vs bond length relationships that are used to determine the force constants of the springs joining neighboring atoms.
The format of this file is described in the documentation of the fitsvsl code, which is a utility that fits such spring constants.

A number of optional files can be given as well.

- 4) An input file defining the atomic masses. By default, the code looks up in the ~/.atat.rc file to determine the directory where atat is installed and then loads the file data/masses.in. This behavior can be overridden with the -m option.
- 5) By default, the bulk modulus is calculated from the force constants but it can also be read from a file (whose name is specified with the -bf option) or specified on the command line with the -b option.

The parameters that govern the accuracy of the calculations are as follows.

The default values are all reasonable.

The code uses the quasiharmonic approximation account for thermal expansion. For this purpose, it calculates the vibrational free for a range of lattice parameters from the 0K value to the (1+ms) larger, where ms is the number specified in the -ms option. The -ns option gives the number of volumes considered. Setting -ns=1 selects the harmonic (instead of the quasiharmonic) approximation.

The code calculates the vibrational free energy from temperature T0 to T1 in steps of dT.

a) The defaults are T0=0 T1=2000 dT=100.

b) If a file Trange.in exists in the upper directory, it is used to set T0,T1,dT:

Trange.in must contain two numbers on one line separated by a space: T1 (T1/dT+1).

Note that T0=0 always.

For phase diagram calculations, you must use this method to specify the temperature range.

c) These defaults can be overridden by the -T0, -T1 and -dT options.

The kpoint sampling is specified with the -kp option. The actual number of kpoints used is the number give divided by the number of atoms in the cell.

-> Phonon Dispersion curves

The -df=inputfile option invokes the phonon dispersion curve module.

The syntax of the input file is:

[nb of points] [kx1] [ky1] [kz1] [kx2] [ky2] [kz2]

repeat...

Each line of input defines one segment (kx1,ky1,kz1)-(kx2,ky2,kz2)

along which the dispersion curve is to be calculated.

[nb of points] specifies the number of points sampled along the segment.

The coordinates are in multiple of the reciprocal cell defined by the axes in the file specified by the -us option (or, by default, in the str.out file).

(The k-point coordinates are appropriately strained

by the amount needed for the str.out file to be identical to the str_relax.out file.)

The phonon frequencies are output in the eigenfreq.out file.

Other parameters can be altered, if needed.

The physical constants are set by default for

force constants input in eV/Angstrom²

temperature in K

free energy in eV

frequencies in Hz

masses in a.u.

They can be altered by the -hb, -kb, -cfk and -mu options.

By default the code give free energies per unit cell, but the -pa option gives them per atom.

The -sc option can provide a multiplicative factor for other conventions (e.g. per formula unit).

By default, the code aborts whenever unstable modes are found, unless the -fn option is specified.

Contact the author for information about the -df and -sf options.

The output files are as follows:

svsl.log : a log file giving some of the intermediate steps of the calculations

vdos.out: the phonon density of states for each lattice parameter considered (unstable modes appear as negative frequencies).

svsl.out: gives along each row, the temperature, the free energy, and the linear thermal expansion (e.g. 0.01 means that the lattice has expansion by 1% at that temperature).
 fvib: gives only the free energy

-> For including vibrations in phase diagram calculations

You are likely to use this code as follow:

```
#first create the Trange.in file for up to 2000K in intervals of 100K:
echo 2000 21 > Trange.in

#This executes the svsl code in each subdirectory containing str_relax.out but no error file.
foreachfile -e str_relax.out pwd \; svsl [options if desired]

#constructs a cluster expansion of the vibrational free energy (eci are in fvib.eci)
clusterexpand fvib

#add the energetic eci from eci.out to the vibrational eci from fvib.eci and create the teci.out
#file that will be read by the Monte Carlo code.
mkteci fvib.eci
```

7.2 Command line options

7.2.1 apb

AntiPhase Boundary 3.20, by Ruoshi Sun and Axel van de Walle
 Automating impurity-enhanced antiphase boundary energy calculations from ab initio Monte Carlo,
 Calphad 53, 20 (2016)

File options:

```
-l=[string] Input file: lattice (Default: lat.in)
-s=[string] : structure (Default: str.out)
-o=[string] Output file: APB structure (Default: str_apb.out)
-og=[string] : energies (Default: gamma_apb.out)
```

APB options:

```
-f Generate APB structure file and exit; do not compute APB energy (Default: Off)
-sx=[real] APB slip vector: x-component (Default: 0)
-sy=[real] : y-component (Default: 0)
-sz=[real] : z-component (Default: 0)
```

Monte Carlo options:

```
-mc Run Monte Carlo (Default: Off)
-eq=[int] Number of: equilibration passes (Default: -1)
-n=[int] : averaging passes (Default: -1)
-T=[real] Temperature (Default: 0)
```

Other options:

```
-d Use all default values (Default: Off)
-h Display more help (Default: Off)
```

7.2.2 cellcvrt

cellcvrt 3.20, by Axel van de Walle

Performs various conversions/changes on structure (or lattices).

Reads from stdin, writes to stdout.

```

-c          Use cartesian coordinates (Default: Off)
-f          Use fractional coordinates (Default: Off)
-cc         Find conventional cell and use it as coordinate system (Default: Off)
-abc        Use a b c alpha beta gamma format (Default: Off)
-ro         Read fractional Occupation (Default: Off)
-u=[string] User-specified coordinate system input file (optional) (Default: )
-uss=[string] User-specified supercell (Default: )
-ss=[int]   Simple supercell (multiple of axes in all directions) (Default: 0)
-fc         Fix cell (to make it as symmetric as possible) (Default: Off)
-s          Look for smaller unit cell (Default: Off)
-sh=[string] Shift all atoms (default 0,0,0).
-ja=[real]  Jitter all atoms positions by a random amount less than specified (Default: 0)
-jc=[real]  Jitter all cell parameters by a random amount less than the fraction specified (Default: 0)
-sd=[int]   Seed for random number generation (default: use clock)
-sg=[string] Space group file (Default: )
-gsg        Generate Space Group (Default: Off)
-wi         Wrap all atoms inside unit cell (Default: Off)
-wiu        Wrap all atoms inside unit cell and undo shift (Default: Off)
-rr         Remove redundant atoms (Default: Off)
-rra        Remove redundant atoms and average (Default: Off)
-ar         Add redundant atoms (Default: Off)
-osf=[string] Original setting file (optional) (Default: )
-fsf=[string] Final setting file (optional) (Default: )
-r          Print reciprocal unit cell (Default: Off)
-fs=[int]   Index of first structure to process (default: 1)
-ns=[int]   Number of structures to process (Default: 2147483647)
-slf=[string] Structure list file (Default: )
-pn         Print the number of atoms in the structure only (Default: Off)
-pv         Print volume of cell only (Default: Off)
-sym        Print spacegroup (Default: Off)
-ppg        Print point group name (Default: Off)
-pbv        Print Bravais Lattice name (Default: Off)
-sc=[real]  Scale factor (default: 1)
-sig=[int]  Number of significant digits to print in output files (Default: 6)
-z=[real]   Tolerance for checking overlap (Default 1e-3)

```

7.2.3 checkcell

check cell distortion 3.20, by Axel van de Walle

```

-d          default (Default: Off)
-q          be quiet (Default: Off)
-p          print strain (Default: Off)

```

7.2.4 corrdump

CORrelation DUMPer 3.20, by Axel van de Walle

```

-h          Display more help (Default: Off)
-2=[real]   Maximum distance between two points within a pair (Default: 0)
-3=[real]   Maximum distance between two points within a triplet (Default: 0)
-4=[real]   Maximum distance between two points within a quadruplet (Default: 0)
-5=[real]   Maximum distance between two points within a quintuplet (Default: 0)
-6=[real]   Maximum distance between two points within a sextuplet (Default: 0)

```

```

-l=[string] Input file defining the lattice (default: lat.in)
-s=[string] Input file defining the structure (default: str.out)
-pc        Print composition only (Default: Off)
-pcm       Print composition matrix only (Default: Off)
-fast      Use fast algo to calculate correlations of 'simple' supercells (Default: Off)
-skipr     Skip algorithm robust to relaxations (Default: Off)
-sym       Just find space group (Default: Off)
-clus      Just find clusters (Default: Off)
-c         Read cluster file instead of writing it (Default: Off)
-cf=[string] File name of the cluster file (default: clusters.out)
-z=[real]  Tolerance for finding symmetry operations (default: 1e-3)
-sig=[int] Number of significant digits printed (default: 5)
-noe       Do not include empty cluster (Default: Off)
-nop       Do not include point cluster(s) (Default: Off)
-noc       Do not include any cluster(s) (Default: Off)
-eca       Calculate Energy of Clusters of Atoms (Default: Off)
-wu=[string] Write Unrelaxed structure into specified file (Default: )
-rnd       Print correlation of the random state of the same composition as the input structure (D
-eci=[string] Predict quantity using ECI in specified file (Default: )
-mi        Multiplicities are already included in ECI file (Default: Off)
-crff=[string] Select correlation functions (default: trigo)
-nb        Print structure number (Default: Off)
-ro        Read lattice file containing occupation variables (this code does not make use of them)

```

7.2.5 csfit

Constituent Strain FITter 3.20, by Axel van de Walle

```

-h          Display more help (Default: Off)
-nc=[int]   Number of points in concentration mesh (default 50)
-ns=[int]   Number of points in mesh used to look for energy minimum (default 100)
-np=[int]   Number of points in stretching mesh in the direction perpendicular to the k-vector(default
-nl=[int]   Number of points in stretching mesh in the direction parallel to the k-vector (default 3)
-ml=[real]  Maximum parallel stretching (default 0.05)
-l=[string] Input file defining the lattice (default: lat.in)
-pa=[string] Directory containing the pure A calculations (default 0/)
-pb=[string] Directory containing the pure B calculations (default 1/)
-ds=[string] File containing a list of stretching directions (default: dir.in)
-t=[int]    Time between disk reads in sec (default: 10 sec)
-sig=[int]  Number of significant digits to print in output files (Default: 6)
-d          Use all default values (Default: Off)

```

7.2.6 cv

Cross-Validation code 3.20, by Axel van de Walle

```

-g          Favor cluster choices where the ground state line is correct (Default: Off)
-w=[real]   Weight structures by w/(struct_energy-gs_energy+w) (Default: 1000)
-p=[real]   Penalty for structures that lie below the ground state line (Default: 0)
-d          Use all default values (Default: Off)
-h          Display Help (Default: Off)

```

7.2.7 emc2

Eazy Monte Carlo Code 3.20, by Axel van de Walle

```

-h          Help (Default: Off)

```

```

-mu0=[real]    initial chemical potential (Default: Off)
-T0=[real]    initial temperature (Default: Off)
-mu1=[real]    final chemical potential (Default: Off)
-T1=[real]    final temperature (Default: Off)
-dmu=[real]    chemical potential step (Default: Off)
-dT=[real]    temperature step (Default: Off)
-db=[real]    inverse temperature step (Default: Off)
-cm           Set Canonical mode (Default: Off)
-x=[real]     Set concentration (Default: Off)
-abs         take chemical potentials as absolute quantities (as in mc.out) (Default: Off)
-phi0=[real]  initial (grand) canonical potential (Default: Off)
-er=[real]    set the system size so that a sphere of that radius must fit inside the simulation cell
-eq=[int]     number of equilibration passes (Default: -1)
-n=[int]      number of averaging passes (Default: -1)
-dx=[real]    Target precision for the average concentration (optional, replaces -n and -eq) (Default: 1e-4)
-aq=[int]     Alternative quantity that must meet the tolerance specified by -dx. 0: energy, 1: concentration
-gs=[int]     which ground state to use as initial config (-gs=-1 to use random state, c=1/2) (Default: -1)
-innerT      inner loop over T (Default: Off)
-tstat=[real] Critical value of the test for discontinuity (Default: 3)
-sigdig=[int] Number of significant digits printed (Default: 6)
-q           Quiet (do not write to stdout) (Default: Off)
-o=[string]   Output file (default: mc.out)
-oss=[string] Output snapshot file (default: mcsnapshot.out)
-opss=[string] Output periodic snapshot files (default: do not write)
-k=[real]     Boltzman's constant (conversion factor from T to energy) (Default: 1)
-keV         Set Boltzman's constant to 8.617e-5 so that temperature is in K when energy is in eV (Default: 1)
-sd=[int]     Seed for random number generation (default: use clock)
-dl          Drop the last data point of each inner loop (after the phase transition occurred) (Default: Off)
-g2c         Convert output to canonical rather than grand-canonical quantities (Default: Off)
-is=[string]  File name containing a user-specified initial configuration (replaces -gs) (Default: )
-ks=[string]  Specify how k space ECI are calculated (e.g. -ks=cs). (Default: )

```

7.2.8 felec

Electronic free energy calculator 3.20, by Axel van de Walle

```

-dos=[string] DOS input file name (default: dos.out)
-T0=[real]    Minimum temperature (default: 0)
-T1=[real]    Maximum temperature (default: 2000)
-dT=[real]    Temperature step (default: 100)
-kb=[real]    Boltzman's constant (default: in eV/K)
-pa          Output free energy per atom instead of per unit cell (Default: Off)
-sc=[real]    Correction factor if spectator ion are present (Default: 1)
-sd=[real]    Smooth DOS with a Gaussian this width (Default: 0)
-nf          Do not calculate free energy (Default: Off)
-sig=[int]    Number of significant digits to print in output files (Default: 5)
-h           Display more help (Default: Off)
-d           Use all default values (Default: Off)

```

7.2.9 fitfc

FIT Force Constants 3.20, by Axel van de Walle

```

-f          Fit force constants (otherwise, generate perturbations) (Default: Off)
-si=[string] Input file defining the ideal structure (default: str.out)
-sr=[string] Input file defining the relaxed structure (default: str_relax.out)

```

```

-er=[real]      Minimum distance between displaced atoms (Default: 0)
-ernn=[real]    Minimum distance between displaced atoms in multiple of nearest neighbor distance (Default: 0)
-fr=[real]      Force constant range (Default: 0)
-frnn=[real]    Force constant range (in multiple of nearest neighbor distance) (Default: 0)
-dr=[real]      Displacement of the perturbed atom (default: 0.2)
-ms=[real]      Strain of the maximum volume sampled (default: 0.01)
-dv             Specified strain above is volumetric (Default: Off)
-ns=[int]       Number of volume sample (default: 2)
-nrr           Do not rerelease structures at each new volume (Default: Off)
-cs           Check for Singular matrix in fit (Default: Off)
-ncs          No check for singular matrix in fit (this is the default, option included for backward compatibility)
-wm           Write fitting Matrices (Default: Off)
-sf=[string]    Extra strain file (Default: )
-m=[string]     Input file defining the atomic masses (default: ${atadir}/data/masses.in)
-T0=[real]     Minimum temperature (default: 0)
-T1=[real]     Maximum temperature (default: 2000)
-dT=[real]     Temperature step (default: 100)
-P=[real]      Pressure (in GPa, default: 0)
-kp=[real]     Number of k-points per reciprocal atom (default: 1000)
-kx=[int]      k-point mesh (along 1st recip lat. vect.) if <>0 then overrides -kp=... (Default: 0)
-ky=[int]      k-point mesh (along 2nd recip lat. vect.) (Default: 0)
-kz=[int]      k-point mesh (along 3rd recip lat. vect.) (Default: 0)
-fp=[int]      Power of polynomial to fit free energy (Default: -1)
-s0           Use a gamma-centered k-point mesh (default shift: 1/2 1/2 1/2) (Default: Off)
-sx=[real]     k-point shift (along 1st recip lat. vect.) (Default: 0.5)
-sy=[real]     k-point shift (along 2nd recip lat. vect.) (Default: 0.5)
-sz=[real]     k-point shift (along 3rd recip lat. vect.) (Default: 0.5)
-df=[string]   Phonon dispersion curve calculation input file. (Default: )
-hp=[real]     Planck's constant (default in (eV s))
-kb=[real]     Boltzman's constant (default in eV/K)
-cfk=[real]    Conversion factor for force constants into energy/dist^2 (default: converts eV/A^2 into eV/Å^2)
-mu=[real]     Mass units (default: converts a.u. mass into kg)
-rl=[real]     Robust Length algorithm parameter for soft modes (beta) (Default: 0)
-rls=[real]    Quadratic strain-dependence of robust length for positive strain (beta) (Default: 0)
-cP=[real]     Conversion factor from pressure*volume into eV (Default: 0.00624151)
-pa           Output free energy per atom instead of per unit cell (Default: Off)
-pe           Print energy in 3rd column of fitfc.out (Default: Off)
-sc=[real]     Correction factor if spectator ion are present (default: 1)
-me0          Subtract energy at OK (Default: Off)
-no           Print number of atom in supercell only (Default: Off)
-w2           Flag negative perturbation directions with wait2 instead of wait file (Default: Off)
-fn           Force continuation of calculations even if unstable (Default: Off)
-fu           Find unstable modes (Default: Off)
-gu=[int]     Generate unstable modes number n (Default: 0)
-mau=[int]     Maximum number of atom per supercell for unstable mode generation (Default: 64)
-sfc=[int]     Simplify force constants: 1=stretching+bending, 2=symmetric (Default: 0)
-apd          Atom-Projected DOS (Default: Off)
-sig=[int]     Number of significant digits printed (default: 5)
-z=[real]     Tolerance for finding symmetry operations (default: 1e-3)
-h           Display more help (Default: Off)

```

7.2.10 fitsvsl

Fit Stiffness VS Length transferable force constants 3.20, by Axel van de Walle


```

-f          Fit force constants (otherwise, generate perturbations) (Default: Off)
-l=[string] Input file defining the lattice (default: lat.in)
-dn=[string] Input file listing the directories containing the structures used to calculate force constants
-er=[real]  Minimum distance between displaced atoms (Default: 0)
-dr=[real]  Displacement of the perturbed atom (default: 0.2)
-ms=[real]  Strain of the maximum volume sampled (default: 0.01)
-ns=[int]   Number of volume sample (default: 2)
-op=[int]   Order of the polynomial used to fit stiffness vs length (Default: 1)
-dd         Use direction-dependent force constants (Default: Off)
-pc=[int]   Maximum power of composition used in fit (default: no composition dependence)
-msl=[real] Maximum spring length (Default: Off)
-sf=[string] Extra strain file (Default: )
-eqt=[real] Tolerance for excluding force constants in plots. (Default: 2)
-sig=[int]  Number of significant digits printed (default: 5)
-z=[real]   Tolerance for finding symmetry operations (default: 1e-3)
-ncs       No check for singular matrix in fit (Default: Off)
-od        Use older 1-level only directory tree (Default: Off)
-h         Display more help (Default: Off)

```

7.2.11 fixcell

fixcell 3.20, by Axel van de Walle

```

-d          Use all default values (Default: Off)
-c          read and print cell in cartesian only (no axes specified) (Default: Off)
-z=[real]   Tolerance for finding symmetry operations (default: 1e-3)
-b          print bravais lattice type only and conventional (potentially not primitive) cell (Default: Off)
-sig=[int]  Number of significant digits to print in output files (Default: 6)

```

7.2.12 gce

Generalized Cluster Expansion 3.20, by Axel van de Walle

```

-h          Display more help (Default: Off)
-2=[real]   Maximum distance between two points within a pair (Default: 0)
-3=[real]   Maximum distance between two points within a triplet (Default: 0)
-4=[real]   Maximum distance between two points within a quadruplet (Default: 0)
-5=[real]   Maximum distance between two points within a quintuplet (Default: 0)
-6=[real]   Maximum distance between two points within a sextuplet (Default: 0)
-l=[string] Input file defining the lattice (default: lat.in)
-s=[string] Input file defining the structure (default: str.out)
-pc         Print composition only (Default: Off)
-sym        Just find space group (Default: Off)
-clus       Just find clusters (Default: Off)
-c          Read clusters.out file instead of writing it (Default: Off)
-z=[real]   Tolerance for finding symmetry operations (default: 1e-3)
-sig=[int]  Number of significant digits printed (default: 5)
-noe        Do not include empty cluster (Default: Off)
-nop        Do not include point cluster(s) (Default: Off)
-rnd        Print correlation of the random state of the same composition as the input structure (Default: Off)
-eci=[string] Predict quantity using ECI in specified file (Default: )
-mi         Multiplicities are already included in ECI file (Default: Off)
-crf=[string] Select correlation functions (default: trigo)
-gce=[string] Select type of Generalized Cluster Expansion (default: tensor)
-nb         Print structure number (Default: Off)

```

7.2.13 gens

Generalized Constituent Strain 3.20, by Axel van de Walle

```
-h          Display more help (Default: Off)
-l=[string] Input file defining the lattice   (default: lat.in)
-s          Specify point group directly via generator (in sym.in) (Default: Off)
-z=[real]   Tolerance for finding symmetry operations (default: 2e-4)
-r=[int]    Maximum Rank of the harmonic (Default: 1)
-mr=[int]   Minimum Rank of the harmonic (default 1)
-sig=[int]  Number of significant digits printed (default: 5)
```

7.2.14 gensqs

GENerate Special Quasirandom Structures 3.20, by Axel van de Walle

```
-n=[int]    nb of atom/unit cell (Default: 0)
-sig=[int]  Number of significant digits to print in output files (Default: 6)
-cf=[string] Input file defining the clusters (default: clusters.out)
-tc=[string] Input file defining the target correlations (default: tcorr.out)
-tol=[real] Tolerance for matching correlations (default: 1e-3)
-l=[string] Input file defining the lattice (default: lat.in)
-rc         Read unit cells from file (Default: Off)
-pc         Print number of supercells only, and quit (Default: Off)
-tp=[int]   Total number of processes (for parallel operation) (Default: 1)
-ip=[int]   Index of current process (0,...,tp-1) (for parallel operation) (Default: 0)
-crf=[string] Select correlation functions (default: trigo)
-h          Display more help (Default: Off)
```

7.2.15 genstr

GENerate STRuctures 3.20, by Axel van de Walle

```
-n=[int]    maximum nb of atom/unit cell (Default: 0)
-sig=[int]  Number of significant digits to print in output files (Default: 6)
-2d         Find supercells along a and b axes only (Default: Off)
-l=[string] Input file defining the lattice (default: lat.in)
```

7.2.16 infdet

INFlection DETection method 3.20, by Axel van de Walle

```
-egt=[real] Epicycle Gradient Tolerance (Default: 1)
-ets=[real] Epicycle Trial Step (Default: 0.1)
-ems=[real] Epicycle Maximum Step (Default: 0.785398)
-eml=[real] Epicycle Multiplier in Line minimization (Default: 2)
-eil=[int]  Epicycle maximum number of Iterations in Line minimizations (Default: 40)
-eic=[int]  Epicycle maximum number of Iterations in Conjugate gradient (Default: 40)
-ebf=[int]  Epicycle motion Bad step Forgiveness (Default: 1)
-el=[real]  Epicycle Length (Default: 0.01)
-ism=[real] Ion motion Steepest descent Multiplier (Default: 0.01)
-isi=[int]  Ion motion Steepest maximum number of Iterations (Default: 0)
-igt=[real] Ion motion Gradient Tolerance (Default: 1)
-its=[real] Ion motion Trial Step (Default: 0.003)
-ims=[real] Ion motion Maximum Step (Default: 0.01)
-iml=[real] Ion motion Multiplier in Line minimization (Default: 2)
-iil=[int]  Ion motion maximum number of Iterations in Line minimizations (Default: 40)
-iic=[int]  Ion motion maximum number of Iterations in Conjugate gradient (Default: 40)
-ibf=[int]  Ion motion Bad step Forgiveness (Default: 1)
```

```

-ics=[real]    Ion motion Curvature constraint Stiffness (0=automatic) (Default: 0)
-ff=[real]     Force scale Factor (Default: 4)
-2d            Considers only 2d strain and stress (Default: Off)
-st=[int]      Sleep Time between read access (Default: 5)
  -r           Relax to minimum (for debugging only) (Default: Off)
  -db          debug (Default: Off)
-aex           Analytical Example (for debugging) (Default: Off)
-t1=[real]     Temp param 1 (Default: 1.9)
-t2=[real]     Temp param 2 (Default: -2)
  -d          Use all default values (Default: Off)

```

7.2.17 kmesh

```

-d            Use all default options (Default: Off)
-q            Quiet! (Default: Off)
-r            Round output to next largest integer (Default: Off)
-e            Force nb of kpoints to be an even number (Default: Off)

```

7.2.18 lsfit

Least-Square FIT 3.20, by Axel van de Walle

```

-x=[string]   x file (Default: )
-y=[string]   y file (Default: )
-w=[string]   weight file (Default: )
-r=[string]   regularization parameters file (Default: )
-pf           Perfect Fit algorithm (Default: Off)
-pw=[string]  number of powers of each column to regress on (Default: )
-s=[string]   select columns to regress on (one number by column, 0: ignore, 1: use) (Default: )
-1           add a column of ones as regressor (Default: Off)
-colin        Ignore colinear columns in x file (Default: Off)
-cv           Print crossvalidation score (Default: Off)
-se           Print standard errors (Default: Off)
-ty           Print trues values of y (Default: Off)
-p           Print predicted values of y (Default: Off)
-e           Print prediction error (Default: Off)

```

7.2.19 maps

MIT Ab initio Phase Stability (MAPS) code 3.20, by Axel van de Walle

```

-h            Display more help (Default: Off)
-l=[string]   Input file defining the lattice (default: lat.in)
-z=[real]     Tolerance for finding symmetry operations (default: 1e-3)
-c=[real]     Exponent of the order of complexity (default: 3)
-t=[int]      Time between disk reads in sec (default: 10 sec)
-m=[int]      Maximum number of points in cluster (default 4)
-g=[int]      Extend ground state search up to structures having at least that many atoms. (Default: 0)
-c0=[real]    [c0,c1] is the concentration range where ground states must be correct. (Default: 0)
-c1=[real]    (Default: 1)
-mv=[int]     Maximum volume (number of atom per unit cell) allowed when generating structures (default
-2d          Find supercells along a and b axes only (Default: Off)
  -p=[string] Predictor plugins to use (examples: -p=cs or -p=cs_es) (Default: )
-ks=[string]  same as -p (Default: )
-fa=[string]  Select fitting algorithm (default: built-in)
-q           Quiet mode (do not print status to stderr) (Default: Off)

```

-sig=[int] Number of significant digits to print in output files (Default: 6)
 -d Use all default values (Default: Off)

7.2.20 mcsqs

Monte Carlo generator of Special Quasirandom Structures 3.20, by Axel van de Walle

-n=[int] nb of atom/unit cell (Default: 0)
 If -n is not specified, generate clusters of the following sizes:
 -2=[real] Maximum distance between two points within a pair (Default: 0)
 -3=[real] Maximum distance between two points within a triplet (Default: 0)
 -4=[real] Maximum distance between two points within a quadruplet (Default: 0)
 -5=[real] Maximum distance between two points within a quintuplet (Default: 0)
 -6=[real] Maximum distance between two points within a sextuplet (Default: 0)
 -l=[string] Input file defining the random structure (default: rndstr.in)
 -cf=[string] Input file defining the clusters (default: clusters.out)
 -tcf=[string] Input file defining the target multibody correlations (default: internally calculated via
 -tol=[real] Tolerance for matching correlations (default: 1e-3)
 -wr=[real] Weight assigned to range of perfect correlation match in objective function (default 1)
 -wn=[real] Multiplicative decrease in weight per additional point in cluster (default 1)
 -wd=[real] Exponent of decay in weight as function of cluster diameter (default 0)
 -T=[real] Temperature (default 1)
 -pf=[string] Input file defining the optimization parameters (default: sqsparam.in)
 -rc Read supercells from file sqscell.out (default: generate internally and write to sqscell
 -ip=[int] Index of current process (for parallel operation) (Default: -1)
 -best Collect best SQS among the outputs of prior parallel runs (Default: Off)
 -crf=[string] Select correlation functions (default: trigo)
 -sd=[int] Seed for random number generation (default: use clock)
 -rt=[int] Read parameter file every (rt) step (default:10000)
 -2d Generate only supercells in the plane of a,b axes (Default: Off)
 -sig=[int] Number of significant digits to print in output files (default: 6)
 -h Display more help (Default: Off)

7.2.21 memc2

Multicomponent Eazy Monte Carlo Code 3.20, by Axel van de Walle

-h Help (Default: Off)
 -er=[real] Set the system size so that a sphere of that radius must fit inside the simulation cell
 -cf=[string] Control file specifying the ranges of temperature and chem. pot. scanned (default: con
 -eq=[int] Number of equilibration passes (Default: -1)
 -n=[int] Number of averaging passes (Default: -1)
 -tp=[real] Target precision (optional, replaces -n and -eq) (Default: 0)
 -aq=[int] Quantity that must meet the tolerance specified by -tp. 1: energy (default), 2: long-r
 -gs=[int] which ground state to use as initial config (-gs=-1 to use random state) (Default: -2)
 -k=[real] Boltzman's constant (conversion factor from T to energy) (Default: 1)
 -keV Set Boltzman's constant to 8.617e-5 so that temperature is in K when energy is in eV (1
 -g2c Convert output to canonical rather than grand-canonical quantities (Default: Off)
 -phi0=[real] initial (grand) canonical potential (default: from mean field approx.)
 -tstat=[real] Critical value of the test for discontinuity (Default: 3)
 -mft=[real] Mean field threshold (if |lte-mf|<mft, use mf values instead of mc) (Default: -1)
 -mftq=[int] Quantity that must meet the tolerance specified by -mft.0: phi (default) , 1: energy ,
 -sigdig=[int] Number of significant digits printed (Default: 6)
 -q Quiet (do not write to stdout) (Default: Off)
 -o=[string] Output file (default: mc.out)
 -oss=[string] Output snapshot file (default: mcsnapshot.out)

```

-opss=[string] Output periodic snapshot files (default: do not write)
-sd=[int]      Seed for random number generation (default: use clock)
-dl           Drop the last data point of each inner loop (after the phase transition occurred) (Default: )
-is=[string]   File name containing a user-specified initial configuration (replaces -gs) (Default: )
-hf           Shift all coordinates by half a grid point in scan (e.g. 2 steps in [0,1] give 0.25,0.75)
-il           Include last coordinate in scan (e.g. 3 steps in [0,1] gives 0,0.5,1 instead of 0,0.33,1)
-ts=[string]   Triangular scanning. Specify list of composition axes (e.g. -ts=1,2). (Default: )
-crf=[string]  Select correlation functions (default: trigo)
-df=[int]      Max distance between flips (in unit cells) in grand-canonical mode (Default: -1)
-mf=[int]      Minimum number of flips in grand-canonical mode (experimental) (Default: 0)
-rw           Use random walk algorithm (experimental) (Default: Off)
-fdT=[real]    Temperature step for finite differences (Default: 0.01)
-fdmu=[real]   Chemical potential step for finite differences (Default: 0.01)
-ks=[string]   Specify how k space ECI are calculated (e.g. -ks=cs). (Default: )

```

7.2.22 mmaps

MIT Ab initio Phase Stability (MAPS) code 3.20, by Axel van de Walle

```

-h           Display more help (Default: Off)
-l=[string]  Input file defining the lattice (default: lat.in)
-z=[real]    Tolerance for finding symmetry operations (default: 1e-3)
-c=[real]    Exponent of the order of complexity (default: 3)
-t=[int]     Time between disk reads in sec (default: 10 sec)
-m=[int]     Maximum number of points in cluster (default 4)
-g=[int]     Extend ground state search up to structures having at least that many atoms. (Default: 0)
-cr=[string] Concentration region input file (Default: crange.in)
-he=[real]   Highest predicted energy, above ground state hull, allowed when generating structures (default: 0)
-mv=[int]    Maximum volume (number of atom per unit cell) allowed when generating structures (default: 1000)
-2d         Find supercells along a and b axes only (Default: Off)
-p=[string]  Predictor plugins to use (examples: -p=es or -p=es_cs) (Default: )
-ks=[string] same as -p (Default: )
-fa=[string] Select fitting algorithm (default: built-in)
-crf=[string] Select correlation functions (default: trigo)
-pn=[string] Property to cluster expand (default: energy)
-ig         Ignore whether cluster expansion predicts correct ground states (Default: Off)
-pa         Quantity to expand is already per atom (Default: Off)
-q          Quiet mode (do not print status to stderr) (Default: Off)
-sig=[int]  Number of significant digits to print in output files (Default: 6)
-d          Use all default values (Default: Off)

```

7.2.23 nnshell

Nearest Neighbor Shell 3.20, by Axel van de Walle and Paul Dalach

```

-l=[string] Input file defining the lattice (default: lat.in)
-lb         List bond types in given structures (Default: Off)
-sig=[int]  Number of significant digits to print in output files (Default: 5)
-r=[real]   Define a max radius for neighbor search (Default: 0)
-d          Use all default values (Default: On)

```

7.2.24 nntouch

nntouch 3.20, by Axel van de Walle

Scales a lattice so that nearest neighbor atom just touch.

```

-l=[string] Input file defining the lattice (default: lat.in)

```

```

-r=[string] Input file defining the atomic radii (default: rad.in)
-s=[real]   Multiplicative scaling factor (Default: 1)
-sig=[int]  Number of significant digits to print in output files (Default: 6)
-d          Use all default values (Default: Off)

```

7.2.25 pdef

Point DEFect generator 3.20, by Axel van de Walle

```

-l=[string] Input file defining the lattice (default: lat.in)
-s=[string] Input file defining the lattice (default: str.out)
-p=[string] Prefix of the directories that will contain the defected structures (Default pdef)
-er=[real]  Minimum distance between point defects (Default: +0.00000)
-sig=[int]  Number of significant digits printed (default: 5)
-h          Display more help (Default: Off)

```

7.2.26 phb

PHase Boundary 3.20, by Axel van de Walle

```

-h          Help (Default: Off)
-mu=[real]  initial chemical potential (Default: Off)
-T=[real]   initial temperature (Default: Off)
-dmu=[real] chemical potential adjustment step (Default: Off)
-dT=[real]  temperature step (Default: Off)
-mug=[real] Gap between the mu in phase 1 and mu in phase 2 (default: 0)
-ltep=[real] threshold free energy precision to use MC instead of LTE (in units of T) (default: always)
-er=[real]  enclosed radius (Default: 0)
-gs1=[int]  ground state for phase #1 (Default: -2)
-gs2=[int]  ground state for phase #2 (Default: -2)
-d1=[string] directory for phase #1 (default: current dir)
-d2=[string] directory for phase #2 (default: current dir)
-lro        print long range order parameter (Default: Off)
-tstat=[real] Critical value of the test for discontinuity (Default: 3)
-smax=[real] Maximum step (experimental feature) (Default: 1e+50)
-sigdig=[int] Number of significant digits printed (Default: 6)
-q          Quiet (do not write to stdout) (Default: Off)
-o=[string] Output file (default: mc.out)
-k=[real]   Boltzman's constant (conversion factor from T to energy) (Default: 1)
-keV        Set Boltzman's constant to 8.617e-5 so that temperature is in K when energy is in eV (Default: 1)
-sd=[int]   Seed for random number generation (default: use clock)
-dn         Go down in temperature (Default: Off)
-dx=[real]  Concentration Precision (Default: 0)
-ks=[string] Specify how k space ECI are calculated (e.g. -ks=cs). (Default: )

```

7.2.27 strpath

STRucture PATH generator 3.20, by Axel van de Walle

```

-s1=[string] Input file defining structure 1 (default: str1.out)
-s2=[string] Input file defining structure 2 (default: str2.out)
-ni=[int]    Number of images (Default: 0)
-f=[real]    Interpolation fraction in [0,1] (Default: 0.5)
-df=[real]   Dimer distance [0,1] (Default: 0)
-dp=[string] Directory name pattern (default 00)
-ci          Calculate inflection point (Default: Off)
-cil         Calculate inflection point location (Default: Off)

```

```
-sig=[int]    Number of significant digits printed (default: 5)
-h           Display more help (Default: Off)
-d           Use all default values (Default: Off)
```

7.2.28 svsl

Vibrational free energy calculator using the Stiffness VS Length method 3.20, by Axel van de Walle

```
-l=[string]  Input file defining the lattice (defaults: lat.in, ../lat.in, str.out)
-us=[string] Input file defining the unrelaxed structure (default: str.out)
-rs=[string] Input file defining the relaxed structure (default: str_relax.out)
-sp=[string] Input file defining the springs (default: slspring.out)
-m=[string]  Input file defining the atomic masses (default: ${atadir}/data/masses.in)
-b=[real]    Bulk modulus (Default: 0)
-bf=[string] Bulk modulus file (Default: )
-ms=[real]   Maximum stretching of the lattice parameter (default: 0.05)
-ns=[int]    Number of lattice parameter stretching step (default: 1 => harmonic approximation)
-T0=[real]   Minimum temperature (default: 0)
-T1=[real]   Maximum temperature (default: 2000)
-dT=[real]   Temperature step (default: 100)
-kp=[real]   Number of k-points per reciprocal atom (default: 1000)
-s0          Use a gamma-centered k-point mesh (default shift: 1/2 1/2 1/2) (Default: On)
-sx=[real]   k-point shift (along 1st recip lat. vect.) (Default: 0.5)
-sy=[real]   k-point shift (along 2nd recip lat. vect.) (Default: 0.5)
-sz=[real]   k-point shift (along 3rd recip lat. vect.) (Default: 0.5)
-hp=[real]   Planck's constant (default in (eV s))
-kb=[real]   Boltzman's constant (default in eV/K)
-cfk=[real]  Conversion factor for force constants into energy/dist^2 (default: converts eV/A^2 into J)
-mu=[real]   Mass units (default: converts a.u. mass into kg)
-rl=[real]   Robust Length algorithm parameter for soft modes (beta) (Default: 0)
-pa          Output free energy per atom instead of per unit cell (Default: Off)
-sc=[real]   Correction factor if spectator ion are present (default: 1)
-fn          Force continuation of calculations even if unstable (Default: Off)
-df=[string] Phonon dispersion curve calculation input file. (Default: )
-sf=[string] Extra strain file (Default: )
-msl=[real]  Maximum spring length (Default: Off)
-sig=[int]   Number of significant digits to print in output files (Default: 5)
-up=[string] User-specified Potential Plug-in (optional) (Default: )
-d           Use all default values (Default: Off)
-h           Display more help (Default: Off)
```

7.2.29 triph

TRIangulate PHase 3.20, by Axel van de Walle

```
-t=[string]  Input table file (Default: tab.in)
-sc=[real]   Scale for the unique axis (1st) (Default: 1000)
-col=[int]   Color (Default: 0)
-nfn         do Not fix Normal vector (Default: Off)
-bf          generate back face (Default: Off)
-sig=[int]   Number of significant digits printed (default: 5)
-ef=[string] Export format (default: vtk)
-h           Display more help (Default: Off)
-d           Use all default values (Default: Off)
```

7.2.30 getclus

Syntax: `getclus [-e] [ecifile]`

extracts, from `clusters.out`, the cluster sizes, lengths and multiplicities
 if `ecifile` is given, it also prints the `eci` therein
 if `-e` is specified, it prints `eci` from `eci.out`

7.2.31 mapsrep

Display the output of maps in graphical form.

Syntax: `mapsrep [-e]`

the optional switch `-e` tells the code to plot structures flagged with error too.
 Thanks to Ruoshi Sun and Mike Widom for updating the script for recent versions of `gnuplot`.

7.2.32 mmapsrep

Display the output of `mmaps` in graphical form.

Syntax: `mapsrep [-e] c1 c2 c3`

where `c1,c2,c3` are the column numbers of the compositions to plot
 the optional switch `-e` tells the code to plot structures flagged with error too.

7.2.33 str2xyz

Utility to convert structure files into xyz files suitable for viewing with `rasmol`

Syntax: `str2xyz [-v] na nb nc [scale] infile`

`-v` (optional) calls `rasmol` (otherwise, xyz file is written to `stdout`)
`na nb nc` define the number of periodic repetition to plot
`scale` (optional) scales the structure to adjust which bonds are plotted

7.2.34 makelat

MAKE LATtice, by Axel van de Walle

Utility setting up directories with maps input files for a given alloy system
 and for one or multiple lattices.

Syntax: `makelat [-o] [-s=scale] atom1,atom2,... lattice1,lattice2,...`

Make sure that there are commas but no spaces between atoms and similarly between the lattices.

Examples:

```
makelat Al,Ni bcc,fcc
makelat Ca,Mg:O rocksalt
```

(note the `:` to separate sublattices)

The default scale is 1. It can be changed to allow for units other than angstroms.

The `-o` option outputs to `strout`.

Lattices available:

B2 bcc bcc-oct bcc-tet diamond fcc fcc-oct fcc-octtet fcc-tet hcp hcp-oct hcp-tet L11 rocksalt sc wurtz

Other lattices can be added in the `/home/avdw/avdw/atat/data/` directory or
 new atomic radius can be added in the `/home/avdw/avdw/atat/data/radii.in` file.

7.2.35 clusterexpand

Syntax: clusterexpand [-e] [-pa] [-s "1,0,1, ..."] filename
 Cluster expands the quantity stored in */filename
 Uses the clusters in clusters.out
 Outputs to filename.eci filename.ecimult
 ecimult contains the eci multiplied by multiplicity
 the option -e causes the structures flagged with an error file to be ignored
 the option -pa indicates that the quantity is per atom already
 so that the code must NOT divide it by the number of atoms
 the option -s lets you select which cluster to include (1) or exclude (0)
 the option -g selects a generalized (tensor) cluster expansion
 the -cv option prints the cv score.

7.2.36 mkteci

Syntax: mkteci file1.eci ...
 where file1.eci etc. is a list of T-dependent eci files obtained with
 the clusterexpand command, e.g.
 clusterexpand fvib in the case of vibrational free energy or
 clusterexpand felec in the case of electronic free energy.
 Usually the file list is: fvib.eci felec.eci
 ^vibrational ^electronic free energy

7.2.37 clusterpredict

Syntax: clusterpredict [-pa] file
 see clusterexpand for more info.
 This command is to be run from within a specific structure directory containing a str.out file.

7.2.38 checkrelax

Prints the amount of relaxation each structure in */str_relax.out has
 undergone, relative to */str.out (or */str_sup.out if it exists)
 Typically, values above 0.1 are considered too much for the
 cluster expansion to be applicable and the offending structures
 should be disabled using the command:
 touch structure_number/error
 where structure_number is the appropriate directory name.
 The option -1 checks relaxations for one structure only in the current directory.

This command measures the magnitude of the strain that need to be applied
 to the unit cell of str.cell to get the unit cell of str_relax.out,
 neglecting (i) isotropic scaling and (ii) rigid rotation.
 The numbers reported are the sqrt of the sum of each element of the strain tensor squared.

7.2.39 calcbulk

Calculates bulk modulus.
 Syntax: calcbulk mins maxs ds runcommand
 where mins and maxs are the minimum and maximum strains (e.g. 0.00 0.01)
 ds is the step size (e.g. 0.005)
 runcommand is the command to run the ab initio code (e.g. runstruct_vasp)

7.2.40 mcdroplast

This script reads the output file of a mc run and deletes the last data point of each inner loop.
(The point after a phase transition occurred.)
Same effect as the -dl option of emc2.

7.2.41 str2cif

Converts an ATAT structure file (from stdin) to the CIF format (stdout).

7.2.42 wycked

WYCKoff position EDitor, by Axel van de Walle

Syntax:

```
wycked [spcgrp] [list_of_wyckoff_position_letter_without_seperators]
```

The output is meant to first be edited to set the free parameters a,b,c,alpha,beta,gamma,x,y,z,etc in each of the lines starting with #const and processed by sspp to create a valid ATAT structure file.

ACKNOWLEDGEMENTS

This code makes use of a Wyckoff position database downloaded from the Bilbao Crystallographic Server at <http://www.cryst.ehu.es/>

Accordingly, if you use this code, you should cite, in addition to the usual ATAT citations, the following papers:

- M. I. Aroyo, J. M. Perez-Mato, D. Orobengoa, E. Tasci, G. de la Flor, A. Kirov
"Crystallography online: Bilbao Crystallographic Server"
Bulg. Chem. Commun. 43(2) 183-197 (2011).
- M. I. Aroyo, J. M. Perez-Mato, C. Capillas, E. Kroumova, S. Ivantchev, G. Madariaga, A. Kirov and H. Wondratschek,
"Bilbao Crystallographic Server I: Databases and crystallographic computing programs"
Z. Krist. 221, 1, 15-27 (2006). doi:10.1524/zkri.2006.221.1.15
- M. I. Aroyo, A. Kirov, C. Capillas, J. M. Perez-Mato and H. Wondratschek
"Bilbao Crystallographic Server II: Representations of crystallographic point groups and space groups", Acta Cryst. A62, 115-128 (2006). doi:10.1107/S0108767305040286

7.2.43 getproto

GET PROTOtypes from database, by Axel van de Walle

Syntax:

```
getproto [string]
```

Returns all structure prototypes matching [string] in ATAT's prototype database.

[string] can be a regular expression, e.g. ^A3

The last 2 columns of the output can be used as arguments to the wycked code to generate the structure (up to some adjustable parameters not constrained by symmetry).

ACKNOWLEDGEMENTS

This file contains processed data from the index of the Landolt-Bornstein database available at <http://www.springermaterials.com/>

If using this data, please cite:

Villars, P., Cenzual, K., Daams, J., Gladyshevskii, R., Shcherban, O., Dubenskyy, V., Melnichenko-Koblyuk, N., Pavlyuk, O., Stoiko, S., Sysa, L.

"Landolt-Bornstein - Group III Condensed Matter: Numerical Data and Functional Relationships in Science and Technology",

Villars, P., Cenzual, K. (ed.).

SpringerMaterials - The Landolt-Bornstein Database (<http://www.springermaterials.com>)
 Volume 43A1-43A10
 2013

7.2.44 sqs2tdb

sqs2tdb version 3.20 by Axel van de Walle and with contributions to the SQS database from Ruoshi Sun, Qijun Hong and Sara Kadkhodaei.

Usage: sqs2tdb [options] where the available options are:

-h : More help.

-cp : Copy SQS from the database to the current directory. Must specify:

-lv=[level]

level indicates how fine of composition mesh to use (0: only end members, 1: mid points, etc)

-l=[lattice]

Available lattices

AUCU_L10
 BCC_A2
 BETASN_A5
 CDI2_C6
 CHI_A12
 CR3SI_A15
 CSCL_B2
 CUPRITE_C3
 CUPT_B22
 DIAMOND_A4
 FCC_A1
 FE3AL_D03
 FLUORITE_C1
 GAMMA_L12
 HCP_A3
 HEUSLER_D022
 HEUSLER_L21
 hidden
 LIQUID
 MGCU2_C15
 NIAS_D81
 OMEGA_C32
 PEROVSKITE_E21
 PYRITE_C2
 REO3_D09
 ROCKSALT_B1
 RUTILE_C4
 SIGMA_D8B
 WURTZITE_B4
 ZINCBLLENDE_B3
 ZINTL_B32
 ZIRCONIA_TET

-fit : Fit solution model from SQS energies (to be run within the desired lattice directory)

-tdb : Combine solution models from one or multiple lattices into a single .tdb file.

-mk : Generate input file for SQS generation with mcsqs. (Usually not needed - only for experts.)

To be run in the SQS database directory:

/home/avdw/avdw/atat/data/sqsdb/[lattice name]

Note that `-h,-mk,-cp,-fit,-tdb` are mutually exclusive.

EXAMPLES

-> A simple session of `sqs2tdb` could be:

```
echo Al,Ni > species.in
```

```
sqs2tdb -cp -l=FCC_A1 -lv=1
```

```
sqs2tdb -cp -l=FCC_A1 -lv=1
```

```
cd FCC_A1
```

```
foreachfile wait pwd \; runstruct_vasp
```

```
sqs2tdb -fit
```

Edit the file `terms.in` to read

```
1,0
```

```
2,0
```

```
sqs2tdb -fit
```

```
cd ..
```

(repeat the above for other lattices if necessary)

```
sqs2tdb -tdb
```

The file `AL_NI.tdb` would contain a valid `.tdb` file.

-> In a more advanced example, the `runstruct_vasp` command could be replaced by a job script submission containing the `runstruct_vasp` command.

-> When there are lattice instabilities, the `runstruct_vasp` could be replaced by `robustrelax_vasp -mk` #this is to be run only once (see `robustrelax_vasp -h` for instructions)
`robustrelax_vasp -neb`

-> If phonons are to be included (for end members) one could do:

```
foreachfile endmem pwd \; fitfc -si=str_relax.out -ernn=4 -ns=1 -nrr
```

```
foreachfile -d 3 wait \; runstruct_vasp -w vaspf.wrap
```

where `vaspf.wrap` contains parameters for a static run to calculate forces.

```
foreachfile endmem pwd \; fitfc -si=str_relax.out -f -frnn=2
```

```
foreachfile endmem pwd \; robustrelax_vasp -vib
```

```
sqs2tdb -fit
```

7.2.45 symbrklib

Usage: `symbrklib [-s fraction] [element] [initial_structure] [final_structure]`

Currently, the initial and final structures can be one of `bcc,fcc,hcp`.

The `-s` option (optional) indicates to just generate a small distortion (a given fraction along the inst

This command is intended to be run before `robustrelax_xxxx`

7.2.46 `getvalue`

Syntax: `getvalue [label]`

Script that scans standard input for a given label and write, to standard output, the numerical value following every occurrence of that label.

Any number of `:` = spaces or tabs can separate the label and the number.

Example: `getvalue "Final energy" < gulp.output`

7.2.47 `getlines`

Syntax: `getlines [-h] [-af | -bf | -bt | -jaf | -jbf | -bt] string1 string2`

Copies selected lines of standard input to standard output.

`-h` print this help.

`-af` print all lines located *after* a line containing `string1`

`-bf` print all lines located *before* a line containing `string1`

`-bt` print all lines located between a line containing `string1` and a line containing `string2`
(If there are multiple blocks of lines bracketed by `[begin string]` and `[end string]` they are all copied.)

The prefix `j` means "just", indicating that the line(s) containing the matching string(s) is (are) omitted.
Don't forget to quote your strings if they contain spaces.

7.2.48 `nltoblank`

Converts each newline character to a space.

7.2.49 `blanktonl`

Converts each contiguous sequence of blanks to a newline character.

7.2.50 `foreachfile`

Syntax: `foreachfile [-e] [-d depth] filename command`

Execute the specified command in every first-level subdirectory containing the file `filename`.

The `-e` option cause `foreachfile` to skip directories containing the file "error".

The `-d` option specifies to go down to lower level subdirectories as well (the default is 1).

7.2.51 `calc`

Syntax: `calc "mathematical expression"`

7.2.52 `sspp`

Simple symbolic pre-processor

reads from stdin, writes to stdout (also writes temporary file `tmp.awk`)

Interprets simple definitions and evaluates numerical expressions with constants definitions.

invoking `sspp -nb` produces a output file where all blank lines have been removed

input file syntax:

understands any C pre-processor commands (see `man cpp`)

also understands `#const [var1]=[value2]; [var2]=[value2]`

the variables `var1 ...` can be used later in the file.

Expression to be evaluated must be enclosed in braces `{}`

except when they are in a `#const` declaration.
 The brace characters are entered as `\{` and `\}`
 Understands `c++` comment char `//`

Examples:

```
#const x=2*3.14
The value is {1+cos(x)}
#define PLUS(x,y) ((x)+(y))
The answer is {PLUS(1,2)}
Braces are \{ \}
```

Implementation details:

Any line beginning with `#const` or `#awk`
 is interpreted as an `awk` command (see `man awk`).
 Expression in braces are `awk` expressions.
 The file `tmp.awk` is the file sent to `awk` for interpretation.

BUGS

Never start a line with `%`
`#` is not a comment character

7.2.53 chl

`chl [-m alternative_machine_config_file]`
 Checks the load on the remote machines specified in the `~/.machines.rc` file
 or as overridden by the `-m` option
 The first column of output is the load and the commands following the `+` sign
 give the command prefix needed to run on that machine
 A default `~/.machines.rc` is created the first time the command is run.
 See comments therein for the format of the file.

7.2.54 minload

Syntax: `minload [-m alternate_machine_config_file] [-w]`
 Returns the command prefix needed to run a command on the remote
 machine having the minimum load, among the
 machines listed in the `~/.machines.rc` file
 (or as specified by the `-m` option).
 A default `~/.machines.rc` file is created the first time
 the `chl` command is run and the file format is described
 in the comments therein.

7.2.55 node

This command is basically like `rsh`, except that the command
 makes sure that the current directory on the remote machine
 is the same as on the local machine.
 (If the two machines do not share the same disk space, use the `-r` option.)

Syntax: `node [-s] [-r] user@host command`
`-s`: use `ssh` instead of `rsh`
`-r`: copy files in current directory to remote host before running command

if user is the same on remote host, use may use
 node [-s] [-r] host command

7.2.56 pollmach

Syntax: pollmach [-f] [-e] [-m alternate_machine_configuration_file] [-s] [-w] command_to_run

This command periodically checks the load on various machines (specified in the file `~/machines.rc`) and looks for subdirectories containing a file called "wait". If the load on a machine is sufficiently low, it executes `command_to_run` on that machine, in the directory where the oldest wait file lies. A default `~/machines.rc` file is created the first time the `chl` command is run. The format of the file is explained in the comments therein. If no `~/machines.rc` file exists, `command_to_run` is run sequentially on the local computer in each directory containing a "wait" file. This is the single-machine mode.

When using this command with maps, `command_to_run` will typically be `runstruct_vasp`

The `-m` option overrides the default `~/machines.rc` (it is optional).

The `-e` option specifies that the command should terminate when no more "wait" files are to be found (only works in single-machine (sequential) mode, when no `~/machines.rc` file is given).

The `-f` option forces the command to run even if of `pollmach_is_running` file exists.

The `-s` option cause pollmach to self-manage the load (it does not poll the nodes for their loads but instead keeps track of where it sends a job and does not send any other job to that node until it is finished). The first column in the `machine_configuration_file` is ignored and can be omitted.

The `-w` option tells pollmach to wait until all sub processes have finished before quitting.

To cleanly stop this program, type
 touch stoppoll

7.2.57 fixeol

converts text files to unix (-u) or to pc (-p) format.

Syntax: fixeol -u|-p [file1, file2 ...]

7.2.58 runstruct-vasp

runstruct_vasp [-w file] [-d depth] [-p] [-lu] [-ng] [-nr] [-ex] cmdprefix

where file is an optional alternate wrap file (default: `vasp.wrap`)

If the wrap file is not found in the current directory, it searches in the parent directories, up to number specified by depth (default is 5). For a description of the syntax of the `vasp.wrap` file, type
 ezvasp -h
 and look for options under the '[INCAR]' section.

-p means preserve vasp output file

-lu means look up one directory for WAVECAR to use as starting point.

-ng means do not generate a vasp.in file, use the existing one.
 -nr means do not run vasp, just generate input files
 -ex means do not generate vasp.in, do not run vasp, but extract info from vasp output file
 cmdprefix is the prefix needed for vasp to run on a remote machine,
 such as "node -s node2"

7.2.59 ezvasp

Syntax: ezvasp [-c] [-s] [-n] [-p "..."] input_file

- c: Cell patch. This option will convert the cell you entered into a cell that has the symmetry needed for vasp to work properly. The atom positions are altered correspondingly.
- s: Static run. After the relaxations are completed, this option makes vasp start again, turning off relaxation and turning on tetrahedron integration with bloech correction. Can achieve the same effect by including DOSTATIC in the INCAR portion of the input file.
- n: do Not run vasp. If you want to create the input files now and run them later, perhaps on another machine.
- p: name of the vasp command ("vasp" by default). You can include additional prefix such as:
 ezvasp -p "rsh remote.machine nice +5 /usr/local/bin/vasp" vasp.in

Here is an example of input file:

```
[INCAR]
SYSTEM = FeNi
ISPIN = 2
PREC = HIGH
ISMEAR = 1
NSW=41
SIGMA = 0.1
IBRION = 2
ISIF = 3
ENMAX = 400
EDIFF = 1e-6
EDIFFG = 1e-4
KPPRA = 1000
USEPOT = PAWGGA
DOSTATIC
SUBATOM = s/Fe$/Fe_h/g
KSCHEME = GAMMA
MAGATOM =

[POSCAR]
SOME TITLE
3.55
-.5000000000    -.5000000000    -1.0000000000
 .5000000000    -1.0000000000    -.5000000000
-.5000000000    .5000000000    1.0000000000
1 2
Direct
.33333 .33333 .00000 Fe+5
.00000 .00000 .00000 Vac
.66667 .66667 .00000 Fe-5
```


End of example.

Note that there 5 new tokens in the INCAR section.

- 1) KPPRA stands for "K-Point Per Reciprocal Atom". This is a way to automatically set the k-point mesh for a number of similar systems. Here is how it works. If you know that you need 500 k-points for a 2-atom structure, then you type in 1000. If you now try a similar structure with 4 atoms, the code automatically use 250 k-points. The mesh along the three axes is automatically chosen to make the mesh as uniform as possible. (For the algorithm, see `atat/src/kmesh.cc`).
You can also specify KPPRA = UPDIR which indicates that the KPOINTS file found one directory above should be used.
(This is useful for cell distortions used in elastic constant calculations of constituent strain runs).
If this token is omitted, you need to specify a [KPOINT] section in the input file that contains what you want the KPOINT file to be.
- 2) DOSTATIC indicates that a static run must be done after the relaxation run (NSW tag is removed and the NSW token are adjusted accordingly). The output files named *.relax contain the result of the (first) relaxation run while the files named *.static contain the results of the (second) static run.
When DOSTATIC is not specified, all output files are named *.static, whether the VASP input parameters allow relaxations or not.
- 3) KSCHEME specifies the type mesh to use: Gamma shift or Monkhorst-Pack. The default is Gamma if KSCHEME is not specified.
- 4) MAGATOM is similar to the usual token MAGMOM used to specify the moment on each atom. The novelty is that it sets the magnetic moments in the POSCAR section. (In version 3.15 and above, this token can be omitted. Having spins specified in the POSCAR section turns this on automatically. It also set ISPIN=2 automatically.)
The usual MAGMOM token is still available (but you cannot use both MAGMOM and MAGATOM).
- 5) USEPOT
selects the potentials used: LDA GGA PAWLDA PAWGGA PAWPBE (or any other potential defined in `.ezvasp`)
By default, LDA (without PAW) is used.
For backward compatibility, you can also use the tag DOGGA instead of USEPOT = GGA
- 6) The SUBATOM token lets you specify a sed-like substitution command to be applied to the name of the atomic species.
For instance, SUBATOM= s/Fe\$/Fe_h/g will cause ezvasp to use the Fe_h pseudopotential instead of the Fe one. Multiple SUBATOM token can be given, one for each substitution. Don't forget the s/ and /g and the \$ after the first element name (to avoid F matching Fe, or Y matching Yb).

There are new features in the POSCAR section as well.

- 1) The line that specify the number of atoms (here 1 2) can be omitted (it is ignored).
- 2) The atomic species are specified to the right of the atom positions.
The corresponding POTCAR are automatically copied. You can specify pseudopotentials like Li_h or O_s
- 3) The number following the species is the magnetic moment on the atom (if MAGMOM is in the INCAR section).
Identical species with different moments will be considered symmetrically distinct to allow for symmetry-breaking relaxations.
- 4) Do not specify "selective dynamics". It will be written to the INCAR file automatically if you include T's and F's after the atom species, e.g.:
0.5 0.5 0.5 Al T T F
- 5) The element "Vac" stands for vacancy. It is ignored for everything but to compute the k-point mesh with the KPPRA option.

7.2.60 robustrelax-vasp

This command implements the "inflection-detection" method introduced in

"The free energy of mechanically unstable phases" by

A. van de Walle, Q. Hong, S. Kadkhodaei and R. Sun

Nat. Com. [forthcoming] (2015).

It calculates the energy of a structure that is potentially mechanically unstable.

In a nutshell: the method finds either a local minimum (for a mechanically stable phase)

or the inflection point on a path joining the unrelaxed and fully relaxed structures

(for a mechanically unstable phase).

Syntax: `robustrelax_vasp [options] command_prefix`

where options are

<code>-id</code>	Inflection Detection method
<code>-idop "options"</code>	Options passed to the <code>infdet</code> command.
<code>-neb</code>	Use the Nudged Elastic Band method to find minimum energy path Otherwise: simply interpolate linearly.
<code>-ex</code>	Only extract data from VASP files. Do not run anything.
<code>-c [real]</code>	Relaxation magnitude cutoff needed to activate robust relax algorithm (default: 0).
<code>-ni [integer]</code>	Number of images in the minimum energy path (default: 6).
<code>-rc [string]</code>	Command for full relaxation (Default: <code>runstruct_vasp -w vasp.wrap</code>).
<code>-vc [string]</code>	Command for volume only relaxation (Default: <code>runstruct_vasp -w vaspvol.wrap</code>).
<code>-cc [string]</code>	Command for chain (neb) calculations (Default: <code>runstruct_vasp -w vaspneb.wrap</code>).
<code>-sc [string]</code>	Command for static runs (Default: <code>runstruct_vasp -w vaspstatic.wrap</code>).
<code>-ic [string]</code>	Command for inflection-detection runs (<code>-id</code> option) (Default: <code>runstruct_vasp -p -w vasp</code>).
<code>-idf [real]</code>	Starting point fraction for inflection detection (default: 0.5)
<code>-ja</code>	
<code>-jc</code>	By how much to jitter atoms (<code>-ja</code>) and cell parameters (<code>-jt</code>) for the initial conditions for inflection-detection method.
<code>-vib</code>	Update vibrational data in <code>svib_ht</code> file from <code>vol_0/svib_ht</code> file.
<code>-mk</code>	Make auxiliary vasp input files (<code>vaspvol.wrap</code> , <code>vaspneb.wrap</code> , <code>vaspstatic.wrap</code> , <code>vaspf.w</code>) Do not run anything.
<code>-cld</code>	clean files for a full restart.
<code>-d</code>	Use all defaults values.
<code>command_prefix</code>	Prefix needed for vasp to run on a remote machine or in parallel (e.g. <code>mpirun</code>).

Note: the commands specified by `-rc`, `-vc`, `-cc`, `-sc` can be set to empty `"` to skip the corresponding calculation step (useful to restart an aborted run).

This command does the following:

- 1) Fully relax the structure (in `str_hint.out` or `str.out`) to create `str_relax.out`, just like `runstruct_vasp`.
- 2) If the relaxation cutoff (`-c=...`) not exceeded, stop. Typically, should be set to `-c=0.05`.
- 3) Relax the volume only of structure in `str_sup.out` or `str.out`. Results are in directory `00/`
- 4) Do a neb calculation between the two structures obtained in step 1 and 3 (if `-neb` is specified) or just a linear interpolation between them. Results are in directories `??/`
Note that you need VTST Tools from <http://theory.cm.utexas.edu/vtsttools/> for the neb method with variable cell shape.
- 5) Do static calculations in each directory `??/`
- 6) Find the inflection point on the path optimized in step 4. If it does not exist, find the minimum energy along that path.
- 7) Report the energy found in 6 in the file `energy` and the corresponding geometry in `str_relax.out`

Notes:

- 1) `str_beg.out` and `str_end.out` contain the extremities of the path.
If you want to restart from scratch, delete these files.
- 2) For high symmetry phases (e.g. bcc,fcc,hcp) you may need to break the symmetry of the phase for it to work.
For that purpose, you can use, e.g.,
`symbrklib fcc bcc W`
or
`cellcvrt -ja=0.01 -jc=0.01 < str.out > str_hint.out .`
Sometimes a supercell is needed (a phonon analysis will tell you that - or `symbrklib` for bcc,fcc,hcp).

Chapter 8

Troubleshooting

8.1 Forum

Please see the new ATAT forum at <http://alum.mit.edu/www/avdw/forum> for answers to common (and uncommon!) questions.

8.2 FAQ

Q. I can do a least-squares by hand with a much lower mean square error (MSE) than MAPS can. Yet, you claim that MAPS is “optimal”. How is this possible?

A. The MSE is not a very good measure of the predictive power of a cluster expansion. It can be made arbitrarily close to zero but merely including enough ECI in the fit! But having a small MSE is no guarantee that the error on the predicted energy of structures not included in the fit is small. The cross-validation (CV) score, which is used in MAPS, is a better measure of the predictive power of a least-squares fit. The CV score does not systematically decrease as the number of ECI increases and it has been shown that choosing the number of ECI such that the CV score is minimized is an asymptotically optimal strategy. Researchers often have a tendency to use too many ECI because this appears to reduce the error, but this is an illusion. MAPS gives you an unbiased estimate of the prediction error, and the magnitude of that number is larger than many would like to believe.

Q. MAPS is frozen. What is going on?

A. MAPS sometimes needs a few minutes to generate new clusters or structures. This is a complex operation, be patient. This generation can occur in the middle of the calculations (finding the best cluster expansion or the best structures) because MAPS only generates these clusters or structures when they are needed.

Q. Where are the results?

A. In a variety of files whose descriptions you can view by typing `maps -h | more`. A utility called `mapsrep` lets you display the most useful output data using `gnuplot`.

Q. What files do I need to get started with MAPS?

A. A `lat.in` file to specify the geometry of the lattice and a “wrapping” file that specifies the parameters of the first-principles code. If you are using `vasp`, this file is usually called `vasp.wrap`. Examples of those files can be found in the `example` directory.

Q. I already have structures and energies. What if I just want to fit them?

A. You need a `lat.in` file to specify the geometry of the lattice and you need to create one subdirectory for each structure (name them as you wish). In each subdirectory, you need a `str.out` file to specify the geometry of the structure and an `energy` file to indicate the energy of that structure. For the format of those files, type `maps -h | more`. Note that the energy must be per unit cell of the structure (not the lattice).

Q. When I type `maps` I only get the command line option help. How can I run it?

A. Type `maps -d` to use all the default options. You can also specify any option you want. Either way, the help message will not be displayed and the code will run. If the help is still displayed, it is because there is

an error in the options you specified.

Q. When I superimpose the energies of `fit.out` with the energies obtained from the Monte Carlo code, I can see that they don't match. What is happening?

A. The "energies" given by the Monte Carlo code are in fact $E - \mu x$ so you need to add μx to get the true energy. Use the `-g2c` option to tell the Monte Carlo code to output the true energy.

Q. How can I verify that the structures do not relax to a superstructure of a different type of lattice (e.g. fcc becoming bcc)?

A. Run the utility called `checkrelax`. It will give you a list all structures along with a measure of the strain they each experienced during relaxation.

Q. When I look at the calculated and fitted energies in the `fit.out` file there are structures where the difference is very large.

A. Make sure that these structures have properly relaxed. A common problem is to start the first-principles calculations with an unrelaxed geometry where the atoms are too close to one another. Restart the energy calculations of the offending structures by using larger starting volumes.

Q. My input files seem correct but ATAT utilities crash while reading them.

A. Make sure your files have unix-style end-of-lines. The `fixeol -u yourinputfile` command should fix that issue.

8.3 Common mistakes

- Write the energy per atom (or per unit cell of the lattice) in the `*/energy` files. One should write the energy per unit cell of the structure (that is, in the way a first-principles code usually gives it).
- Errors in the `lat.in` file. Use the `corrddump -sym ; more sym.out` command to analyse the symmetry of the lattice and see if it correspond to your expectations. You can also use `corrddump -2=10 -clus ; more clusters.out` to verify the symmetrically distinct pairs are what you expect them to be.
- When the structures are not generated by MAPS (i.e. you typed them it), beware of typos. MAPS will detect if a structure is incompatible with the lattice but it will not detect another very common mistake: putting the energy and str.out files that correspond to different structures in the same directory. To check for this error, look at the residual plot generated by `mapsrep`.
- When copying the `lat.in` files provided as examples, make sure you make all the required changes. (i) Change the atom names. (ii) Set the lattice parameters (first 3 numbers in the file) to values that are at least as large as the true lattice parameter (otherwise, first-principles code can be unable to properly relax the geometry of the structure is the atoms are too close to one another initially).

Chapter 9

Organization of the Toolkit

- src (source code)
- glue (utilities to interface MAPS with other codes)
- glue/jobctrl (utilities allowing maps to automatically launch jobs)
- glue/vasp (interface between vasp and maps)
- doc (user guide and documentation regarding the inner workings of the code)