

# **LMAPR2451**

## **Atomistic and nanoscopic simulations**

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# **Study of FeS<sub>2</sub> and its optical properties**

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This report aims to investigate the optical properties of FeS<sub>2</sub>. To do so, *ab initio* computations are performed on a 6-atoms orthorhombic unit cell. The convergence with respect to several structural parameters is also studied. The optical properties are then analyzed in the light of the obtained results. Finally, a comparison is made with the published research, and a discussion about the quality of the simulation is made.

**2020-2021**

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# 1 Introduction

## 1.1 Motivation

Pyrite ( $\text{FeS}_2$ ) is an interesting semiconducting material. Indeed, it can be used in numerous domains, from mechanical applications, where it is appreciated for its toughness and abrasiveness, to optical applications, where it can be used as a high-energy light absorber [1]. As pyrite is very abundant, it can become a material of choice in the industry if correct and valuable uses of the latter could be elaborated. It is depicted on Figure 1a.



(a) Pyrite



(b) Marcassite, the orthrhombic polymorph of pyrite

Figure 1:  $\text{FeS}_2$  ores

One of the promising potential application of pyrite is its integration into solar panels. Indeed, pyrite thin films could be good alternatives to the conventional silicon based solar cells, since the latter are more economically and ecologically costly [2, 3]. However, the performances of the material must be similar to or better than materials already used for solar cells, in order to make it truly competitive.

The concept of solar cell dates all the way back to 1883 [4]. However, it is interesting to recall some of the main features of such a device. A solar cell consists of photoelectric components able to convert the energy of the (solar) light into electricity through the *photovoltaic effect*. More precisely, it is composed of a *pn*-junction diode, usually made of silicon. When the depletion region of the junction is hit by solar photons, an electron is delivered in the *n*-layer and a hole is delivered in the *p*-layer (Figure 2). By doing so on a large scale, the *pn*-junction is able to generate a voltage of about 0.5 V, if the *n*-doped layer is sufficiently thin (to make the photons reach the depletion region as easily as possible). A solar panel is composed of several solar cells in series, in order to add up the voltage generated by each cell. The best materials for solar cells-related applications are the materials showing :

- a semiconducting behavior
- a high optical absorption coefficient
- a high electrical conductivity
- an abundant presence in the Earth's crust.[5]

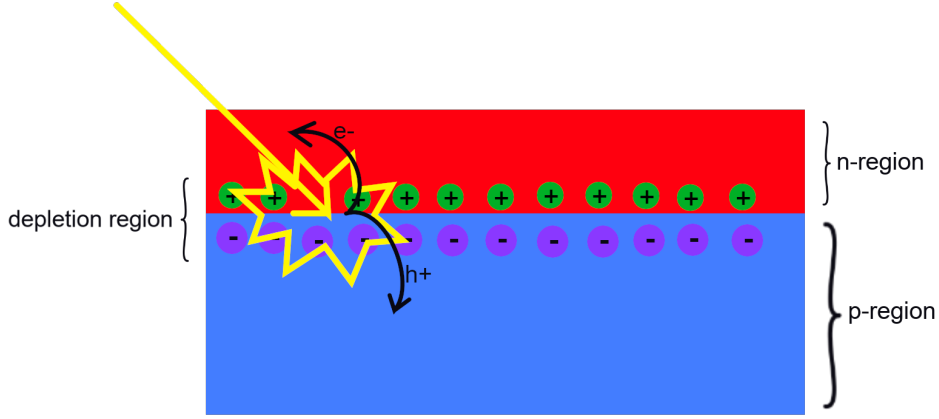


Figure 2: Simplified view of the photovoltaic effect at the  $pn$ -junction of a solar cell. The green and purple dots are the charges resulting from the recombination of charge carriers in the depletion region. When a photon hits an atom in the depletion region, an electron is released in the  $n$ -region and a hole is released in the  $p$ -region, creating a voltage when repeated multiple times.

Even if bulk pyrite shows a  $n$ -type behavior, pyrite thin-films show a  $p$ -type behavior. Only the deposition of a thin layer of  $n$ -type semiconductor on the pyrite film is needed to form a  $pn$ -heterojunction. Furthermore, the pyrite has a strong absorption coefficient. It thus allows to decrease the thickness of the pyrite film, which is helpful to improve the performances of the  $pn$ -junction, and hence, of the solar cell [6]. Furthermore, it is earth-abundant. Pyrite thus seems to be a material of choice for photovoltaic applications.

However, serious limitations prevent pyrite to be highly performant in this framework. Indeed, a very poor solar energy conversion efficiency is observed when assessing the performances of pyrite solar cells. That low efficiency is presumably due to a low photovoltage generation. The origin of that low photovoltage is highly debated among the scientific community. Several possible sources have been proposed over the years : detrimental S vacancies (although stoichiometric pyrite shows the same low photovoltage), detrimental impurities or even lattice defects [7]. No consensus has been found so far. But as pyrite could be a game-changer in the solar cell industry if its efficiency were improved, it is worth to look a little closer to the optical properties of the material.

In the following report, the properties of the  $\text{FeS}_2$  in the  $Pn\bar{m}$  space group will be studied using the **Abinit** package [8]. As the unit cell of pyrite, which contains 12 atoms, is close to the computational limits of **Abinit**, the computations will be performed on a 6-atoms orthorhombic polymorph, depicted on Figure 1b. The results will be used as a starting point for the reflection concerning the actual pyrite. First, the unit cell and its structural parameters will be described, based on the Materials Project documentation<sup>1</sup>. Then, the representation of the crystal in **Abinit** will be presented. Secondly, the pseudopotential and the approximation used in the first place will be discussed. Then, the convergence of the total energy per atom and the lattice scale parameters, with respect to the energy cut-off (**ecut**) and the number of  $k$ -points (**ngkpt**) will be studied. After that, **Abinit** computations will be performed using the converged values and the optical properties will be studied. To conclude, the obtained results will be compared with the published research.

<sup>1</sup><https://materialsproject.org/materials/mp-1522/>

## 1.2 FeS<sub>2</sub> : Overview and Abinit representation

The orthorhombic FeS<sub>2</sub> primitive cell contains 2 Fe atoms and 4 S atoms (Figure 3). The space group is Pnnm[58] in the Hermann-Mauguin notation.

Furthermore, it is a semiconductor. The energy of the indirect bandgap is about 0.978 eV[9].

Finally, the primitive cell will be represented as follow in Abinit input files :

```
acell  3.390 4.438 5.411 Angstr # the lattice vectors scaling
ntypat 2                        # there are two types of atoms in the
                                # primitive cell : Fe and S
znucl  26 16                   # Fe has 26 electrons and S has 16
natom   6                      # there are 6 atoms in the primitive cell
typat   1 1 2 2 2 2           # 2 Fe atoms and 4 S atoms
xred    0      0      0       # position of the first Fe atom in reduced coordinates
      0.5    0.5    0.5       # position of the second Fe atom
      0      0.206  0.3753    # position of the first S atom
      0      0.794  0.6247    # position of the second S atom
      0.5    0.294  0.8753    # position of the third S atom
      0.5    0.706  0.1247    # position of the fourth S atom
```

The data also comes from the Materials Project [9].

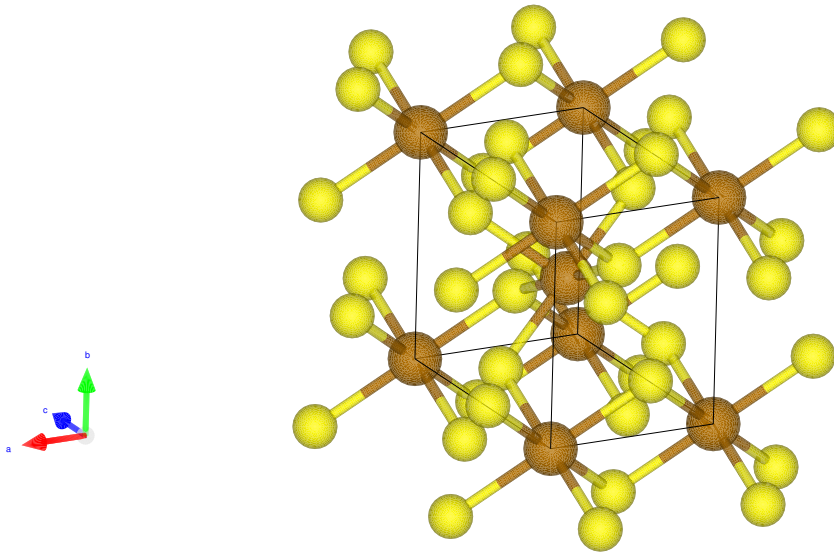


Figure 3: Primitive cell of orthorhombic FeS<sub>2</sub>.  
Materials Project (mp-1522), Vesta

## 2 Convergence studies and pseudopotentials

In DFT computations, convergence studies are very important. They are meant to ensure that the chosen input parameters, like the cut-off energy (`ecut`) or the number of  $k$ -points in the cell (determined by the sampling of the Brillouin zone `ngkpt`), allow accurate results while limiting the computational time as much. To perform such an analysis, the same computations are repeated using datasets with various values of the parameter of interest, with an increasing accuracy of the environment. The evolution of some calculated variables, like the total energy of the unit cell (`etotal`), indicates when the parameters of interest guarantee a sufficiently accurate simulation.

In the following subsections, the convergence of the total energy per atom and of the lattice scale parameters will be studied with respect to `ecut` and `ngkpt`.

### 2.1 Additional parameters

To begin, we will use the *Local Density Approximation* (LDA) functional. The pseudopotentials that will be used are retrieved from the PseudoDojo [10]. Two pseudopotentials are used : the NC SR (ONCVSP v0.4.1) LDA standard pseudopotential relative to Fe, and the same one but relative to S (both in `psp8` format). The pseudopotentials respectively consider 16 electrons ( $3s^23p^64s^23d^6$  for Fe) and 6 electrons ( $3s^23p^4$  for S) so that the unit cell contains 56 electrons.

It is also important to properly define the parameters ruling the SCF procedure. The most important one is `nstep`, defining the number of allowed SCF cycles to reach convergence. It is set to 100 in the first place. `toldfe`, the difference of total energy between two SCF cycles defining the moment when convergence is reached, is set to `1.0d-10` Ha. `toldfe` is chosen as the structural relaxation is not performed yet (in that case, `toldff` or `tolrff` is preferably used). Although it is not mandatory, the SCF procedure can be preconditioned by specifying the macroscopic dielectric constant (`diemac`). It is used to speed up the SCF procedure. A value of 24 is chosen, accordingly to the Materials Project [9].

Finally, the parameters of the  $k$ -points grid must be specified. `kptopt` is set to 1 in order to take advantage of the symmetry of the unit cell. By setting `prtkpt` to 1, Abinit will generate a set of  $k$ -point grids, that will be helpful to optimize further computations and convergence studies.

```
pseudos "pdj_nc_sr_041_lda_standard_psp8/Fe.psp8, pdj_nc_sr_041_lda_standard_psp8/S.psp8"

# parameters of the SCF procedure :
nstep      100      # maximal number of SCF cycles
toldfe     1.0d-10  # SCF procedure will stop when the difference of total energy
                   # between two iterations will be lower than toldfe Hartree
diemac     24.0     # preconditioning of the SCF procedure.

# parameters of the k-points grid :
kptopt     1
prtkpt     1
```

### 2.2 Convergence of the total energy per atom as a function of the cut-off energy (`ecut`)

In the next computations, the default Monkhorst-Pack grids will be used. 4  $k$ -points will be sampled along the longest lattice direction (`[0 0 1]`). To keep a constant density of  $k$ -points

along each axis, 3  $k$ -points will be sampled along directions  $[1\ 0\ 0]$  and  $[0\ 1\ 0]$ . The input lattice parameters are thus :

```
ngkpt    3 3 4
nshiftk  1
shiftk    0.5 0.5 0.5
```

The study of the convergence of the simulation was performed by running **Abinit** with the input file `1522_3_ecutConv.abi` (Appendix A). The total energy per atom of the unit cell is then plotted versus `ecut` (which spans from 10 [Ha] to 59 [Ha]) (Figure 4). Upper and lower bounds of  $\pm 0.5$  [mHa] are set around the last obtained value. It allows to determine the first `ecut` for which the convergence is reasonable. In the present case, `ecut` = 40 [Ha].

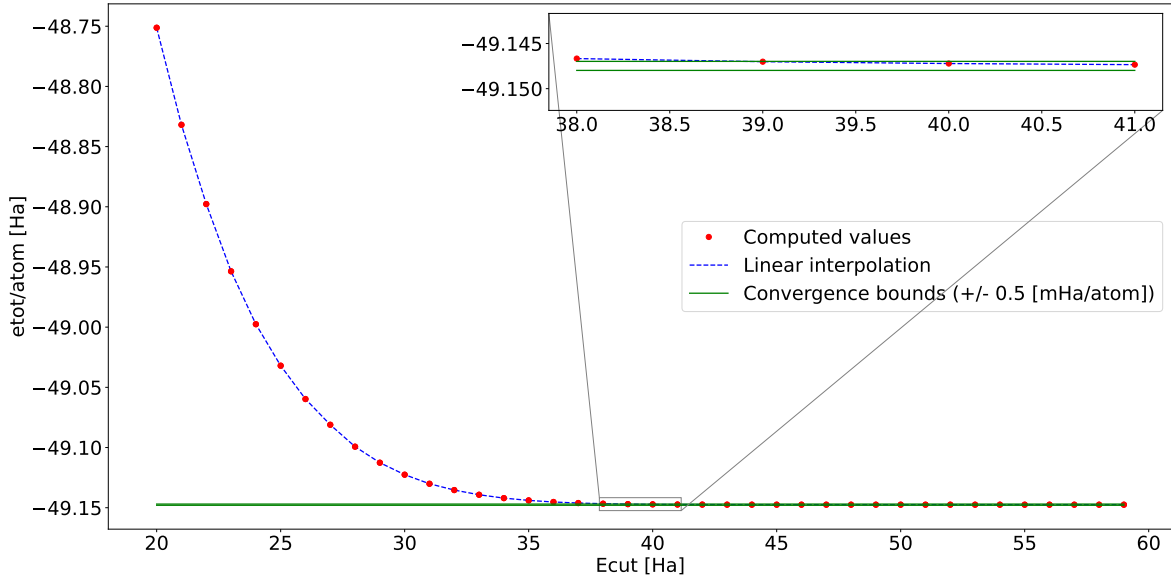


Figure 4: Total energy per atom as a function of the cut-off energy.

The `ecut` range is truncated for visual reasons.

Linear interpolations are shown as a guide for the eyes.

### 2.3 Convergence of the total energy per atom as a function of the number of $k$ -points (ngkpt)

Alternatively, the same kind of study is performed with respect to the number of  $k$ -points in the Brillouin zone. **Abinit** is run with the input file `1522_4_nkpConv.abi` (Appendix B). The energy per atom can be plotted versus the `ngkpt` parameter (Figure 5). However, as the biggest and the smallest lattice scale parameters differs from 60% in size, additional values of `ngkpt` keeping a similar  $k$ -sampling density along each axis are also tested.

The first converged `ngkpt` value is `[3 3 3]`.

Concerning the convergence of the total energy per atom, the couple of values (`ecut`, `ngkpt`) that will be used in the further computations is thus `(40 [Ha], [3 3 3])`.

### 2.4 Convergence of `acell` as a function of the cut-off energy

The convergence of the length scales of the unit cell was performed using a dataset of `ecut` values between 20 [Ha] and 50 [Ha], `ngkpt` = `[3 3 3]` and `ecutsm` = 0.5 [Ha]. The input file

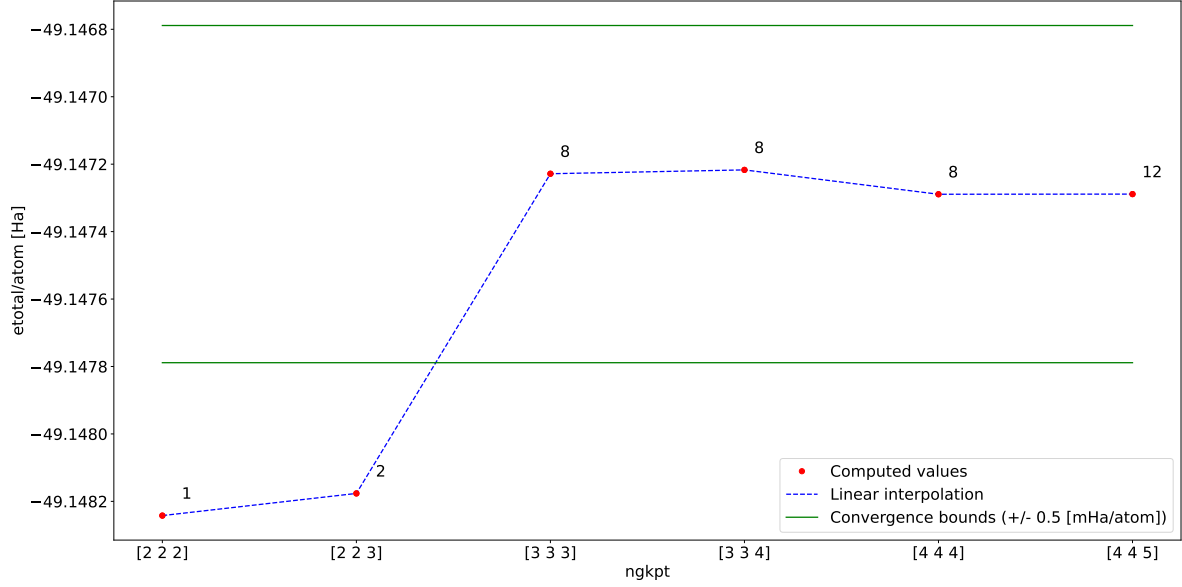


Figure 5: Total energy per atom as a function of the number of  $k$ -points in the  $k$ -points grid. The number near each dot represents the corresponding number of  $k$ -points.

used is `1522_6_acellEcutConv.abi` (Appendix C). The results are displayed on Figure 6. The first converged value is 30 [Ha].

## 2.5 Convergence of `acell` as a function of the number of $k$ -points

The convergence of the length scales of the unit cell was analyzed with `ecut` = 40 [Ha] and `ecutsm` = 0.5 [Ha]. Additional values of `ngkpt` accounting for the conservation of a similar  $k$ -sampling density along the axis of the unit cell are also tested. The input file used is `1522_3_acellNgkptConv.abi` (Appendix D). The results are displayed on Figure 7. It can be seen that [2 2 2] is already in the limits of 0.2% of the length. Therefore, the value [2 2 2] is considered as the first converged value.

## 2.6 Summary

	ecut [Ha]	ngkpt
Convergence of <code>etotal/atom</code>	40	[3 3 3]
Convergence of <code>acell</code>	30	[2 2 2]

Furthermore, the most accurate value for `acell` and `xred`, obtained with `ngkpt` = [4 4 5] (`nkpt` = 12) and `ecut` = 40 [Ha], is

<code>acell(1)</code>	6.2645 Bohr
<code>acell(2)</code>	8.2011 Bohr
<code>acell(3)</code>	9.9992 Bohr

or

<code>acell(1)</code>	3.315 Angstroms
<code>acell(2)</code>	4.3398 Angstroms
<code>acell(3)</code>	5.2913 Angstroms



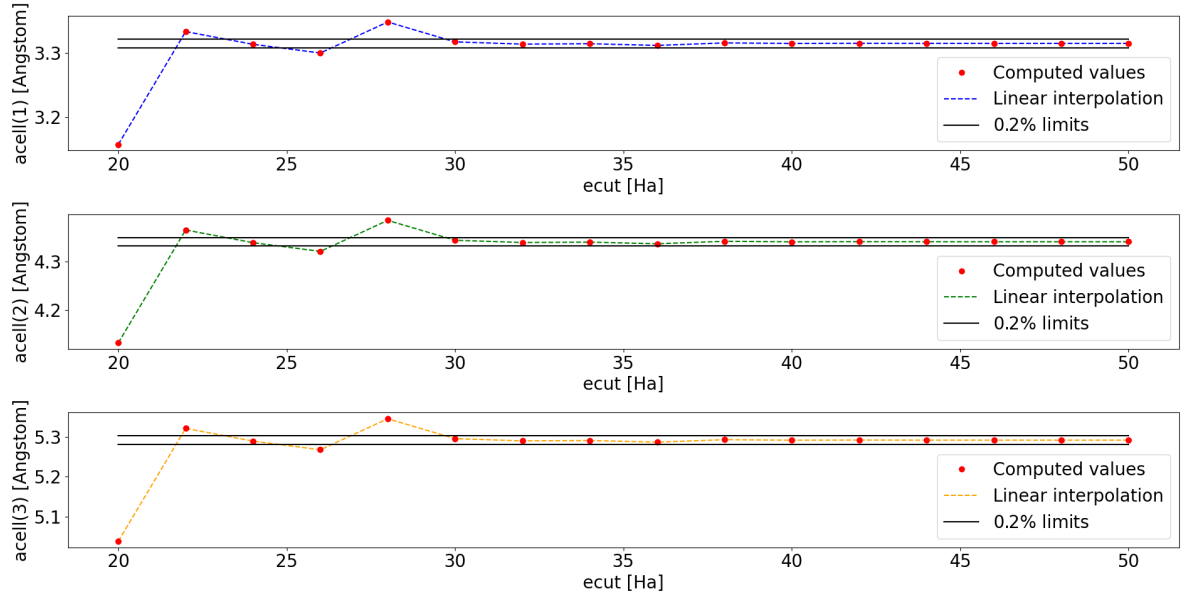


Figure 6: Convergence of the length scales  $\text{acell}$  as a function of the cut-off energy  $\text{ecut}$ .

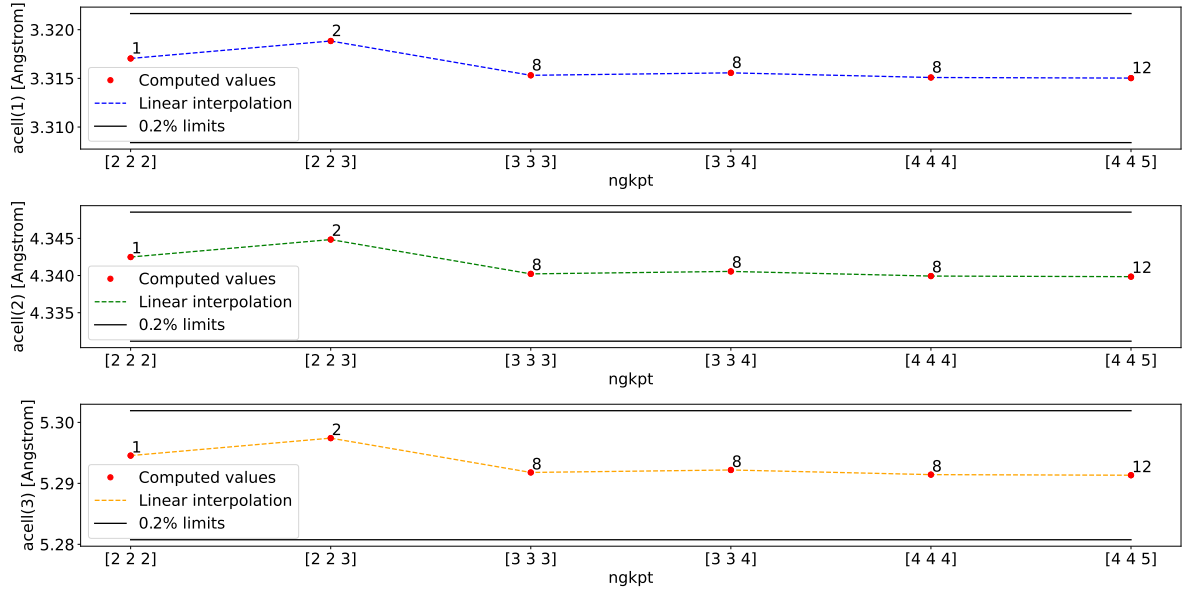


Figure 7: Convergence of the length scales  $\text{acell}$  as a function of the number of  $k$ -points in the  $k$ -points grid. The number near each dot represents the corresponding number of  $k$ -points.

and

	xred(1)	xred(2)	xred(3)
Fe #1	0	0	0
Fe #2	0.5	0.5	0.5
S #1	0	2.0809E-01	3.7374E-01
S #2	0	7.9191E-01	6.2626E-01
S #3	0.5	2.9191E-01	8.7374E-01
S #4	0.5	7.0809E-01	1.2626E-01

### 3 Optical properties

#### 3.1 Electronic band structure

The band structure of the material was computed as follows. First, an optimal  $k$ -path is generated in the Brillouin zone of a simple orthorhombic lattice Figure 8 using the **Abistruct** module of **Abipy** (Appendix E).

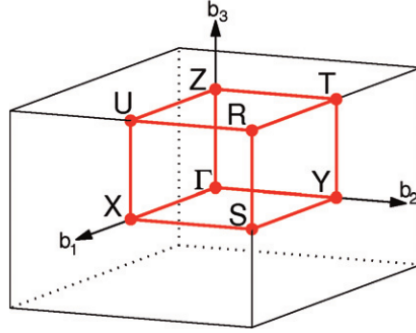


Figure 8: Brillouin zone for a simple orthorhombic lattice [11]

To obtain the band structure of the material, a SCF computation of the electronic density is run in the first place. Then, the band structure is computed using a non-SCF computation based on the previously generated electronic density file. The structural parameters of the unit cell are the ones provided by **Absitstruct**. **ecut** and **ngkpt** are chosen accordingly to the previous convergence studies.

For the SCF computation, **prtden** is set to 1, in order to generate the DEN files.

For the band structure computation, **iscf** is set to -2 and **getden** is set to -1 to read the electronic densities from the previously generated DEN file. As the pseudopotentials used only consider 56 electrons in the whole unit cell, 27 electronic bands are completely filled. The 28th band is thus the valence band. To obtain a clear representation of the band gap of the material, 35 electronic bands are defined. **kptopt** is set to -15 accordingly to the  $k$ -path provided by **Abistruct**. **ndivsm** is set to 30, and **nbdbuf** is set to -2 to place the last 2% bands of highest energy in a buffer, preventing **Abinit** to spent too much time trying to converge them.

The input file **1522\_14\_bs.abi** is available in Appendix F.

The band structure is represented on Figure 9. The Fermi energy and the energy of the bandgap were also retrieved from the computed data. It is compared below with the data from the Materials Project [9]. The band structure provided by the Materials Project, which can be found in Appendix G, is compared with the computed one on Figure 10.

Besides a difference of 2.45 eV concerning the Fermi energies, the results of the computation seems consistent regarding the data of the Materials Project. **why ???**

	Computed	Materials Project
Fermi energy [eV]	10.13183	7.6763
Band gap energy [eV]	0.8674	0.8807

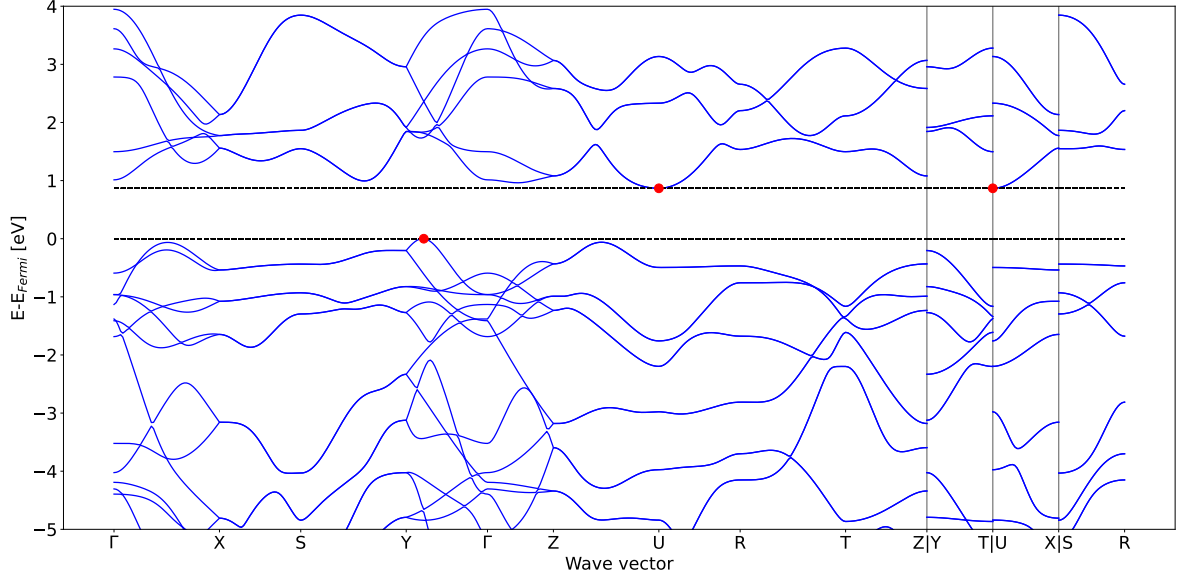


Figure 9: Band structure of  $\text{FeS}_2$ . The red dots represent the valence band minimums and the conduction band maximum.

### 3.2 Electronic density of states (DOS)

The electronic density of states was obtained by first running a SCF computation to generate a density file as for the computation of the bandstructure. In that case, `shiftk` was set to `[0 0 0]` as recommended in the Abinit documentation [12].

Then, the DOS was computed using the tetrahedron methode with `prtdos = 2` and `dosdeltae = 5.0d-5` Ha. `ngkpt` was set to `[8 8 8]` to ensure a good resolution in the DOS plot. The input file used is available in Appendix H. Several plots were obtained when trying to display the density of states of the material. Each of the plot presents a different level of smearing. The smearing is controlled by the  $k$ -sampling of the Brillouin zone and the step of energy that was chosen during the computation of the density of states. Indeed, the  $k$ -sampling and the energy step have both an influence on the resolution of the plot, as the DOS plot is obtained with the sum of Gaussian spreads localized on the eigenvalues composing each band, and for each  $k$ -point. Therefore, the smaller the energy step, the more detailed (and as it will be seen, the rougher) the pattern of density of states, as long as the  $k$ -sampling is sufficiently high (if the number of  $k$ -points or the energy step is too small, there will be undesired gaps in the DOS plot).

By default, `Abipy` smears the the plot of the DOS, no matter what step of energy was chosen fort the computation. It will thus be compared to the DOS pattern provided by the Materials Project, as the smearing is of the same order of magnitude there. The comparison is displayed on Figure 11. It can be seen that the two plots present the same features : a drop of the DOS at  $E = 0$  eV representing the bandgap, and similar peaks around the bandgap. However, substantial differences subsist :

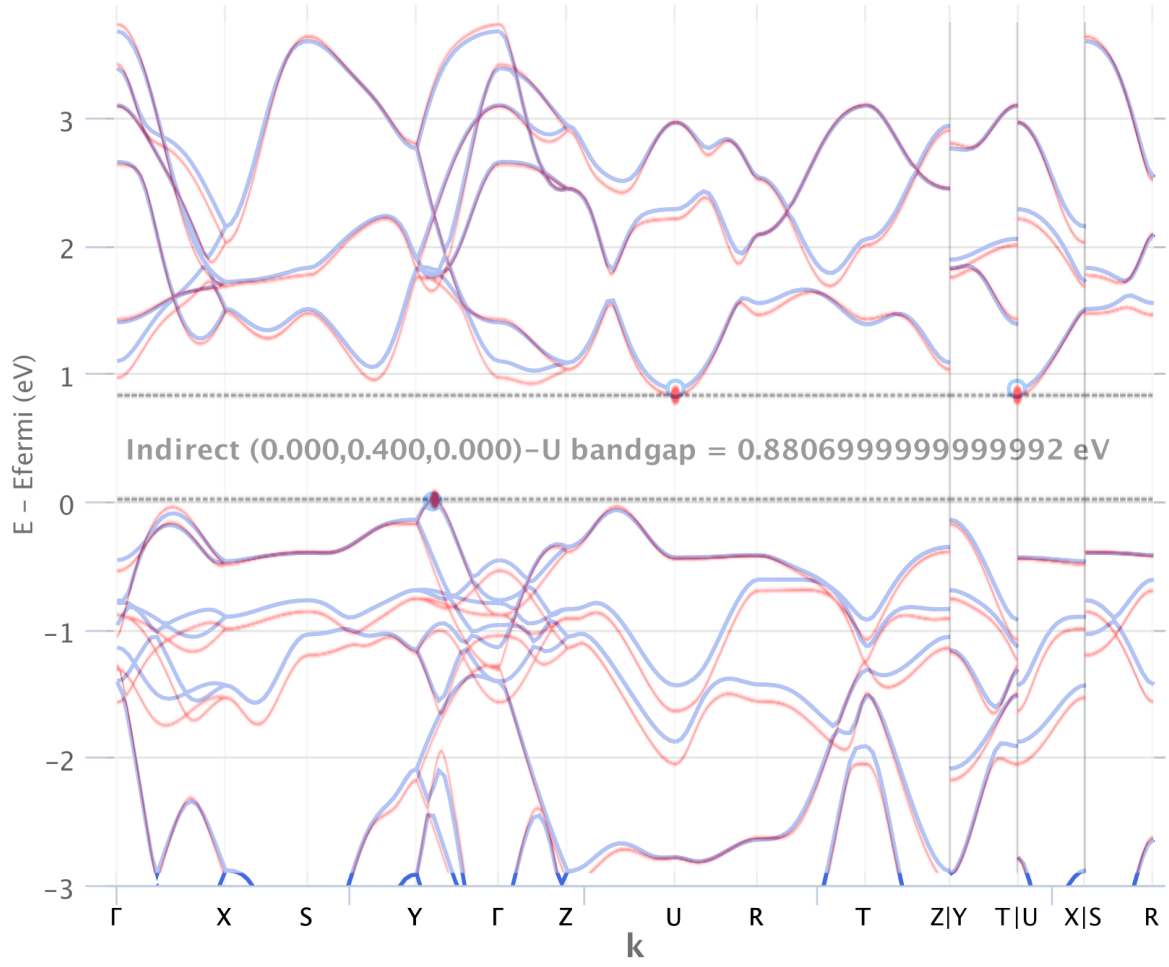
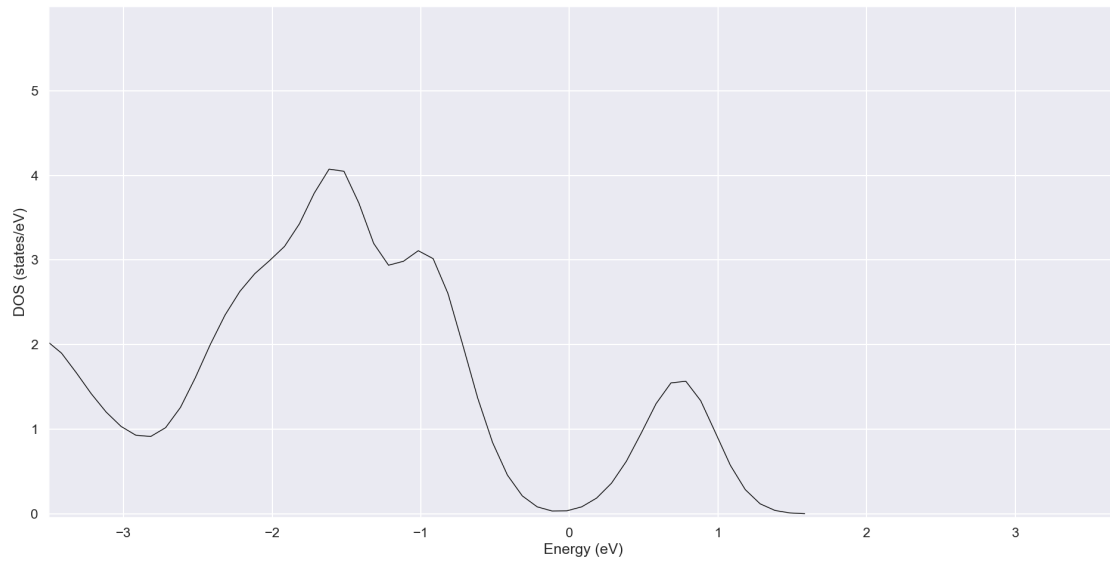
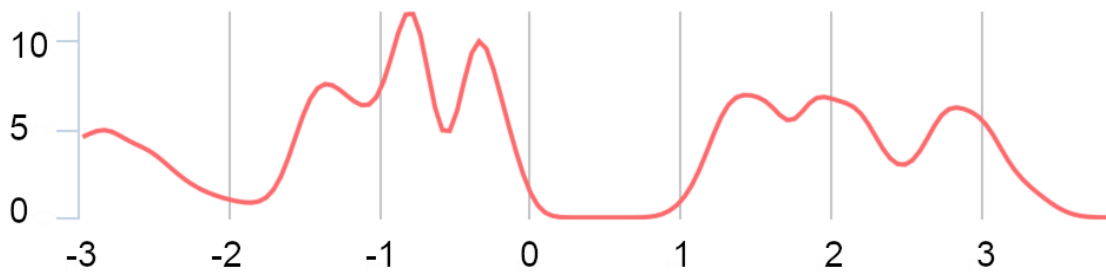


Figure 10: Blue : Band structure provided by the Materials Project. Red : Computed band structure.

- First, the plot displayed by **Abipy** (Figure 11a) isn't as wide as the one provided by the Materials Project (Figure 11b). This is due to the fact that only the first conduction band was computed with **Abinit** in the first case, while additional conduction bands were considered on the Materials Project.
- Secondly, the amplitude of the peaks isn't the same for the two graphs. Once again, it is presumably due to smearing reasons. The main drawback of these two sources (**Abipy** and the Materials Project) is that the smearing (or the energy step and the sampling of the Brillouin zone) is a black box. However, it can also be due to the method that was used to generate the data (but as it is not specified on the Materials Project, it's difficult to be sure).
- Lastly, it can be seen that the peaks on the plot provided by the Materials Project are higher in terms of DOS than on the plot calculated by **Abinit** and displayed by **Abipy**.  
**why ???**



(a) Electronic DOS (displayed using **Abipy**)

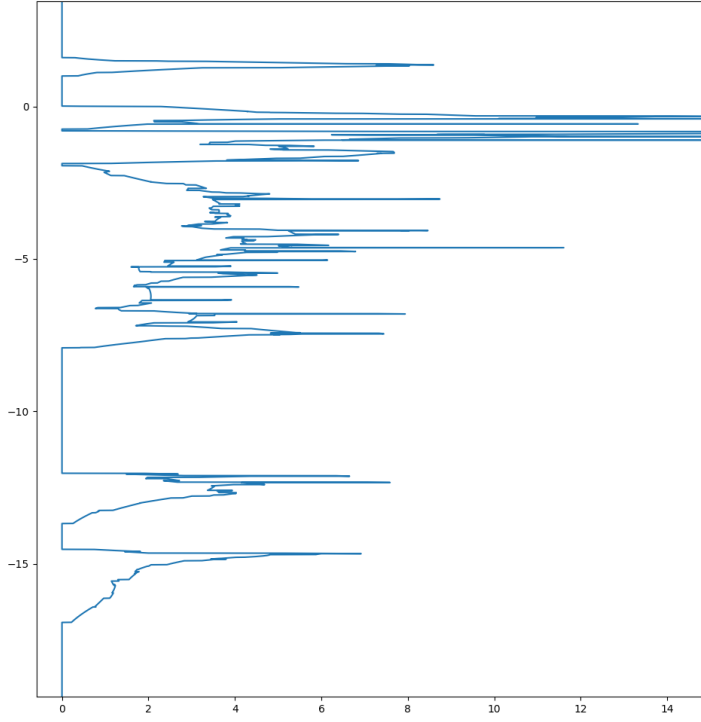


(b) Electronic DOS provided by the Materials Project

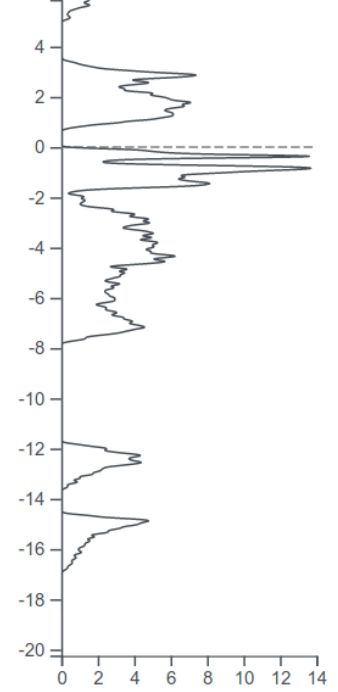
Figure 11: Comparison of the most smeared DOS patterns

Then, the plot of the electronic DOS was displayed using a handmade **Python** script, using the `_DS2_GSR.nc` output file generated by **Abinit** (here **DS2** is explained by the fact that the DOS was computed in the second dataset of the run, the first one being the SCF density computation).

It was then compared with a plot of the DOS whose smearing seems to be significantly lower than for the two previous examples. The latter plot was obtained on the Topological Materials Database [13, 14, 15, 16, 17]. The two plots are compared on Figure 12. The goal here is to make the comparison between two less smeared plots.



(a) Electronic DOS plotted using **Python** and the `_DS2_GSR.nc` **Abinit** output file



(b) Electronic DOS provided by the Topological Materials Database

Figure 12: Comparison of less smeared DOS patterns

One can see here that the plots also show the same features. Some differences can however be mentioned :

- On the plot obtained using **Abinit** and **Python** (Figure 12a), the peaks above the bandgap are not as developed as on the plot provided by the Topological Materials Database (Figure 12b). This is due to the same reason as previously : only the first conduction band has been considered during the **Abinit** computation, while it is clearly not the case for the other plot.
- On the plot obtained with **Python**, peaks of high amplitude can be seen at several energies (for example just below the bandgap). After further investigation, the peaks span until a DOS of  $\sim 65$  states/eV, implying that the amplitude of those peaks is probably a computational artefact. Such peaks are not visible on the other plot, because it has been smeared a little, demonstrating the utility of the smearing of the plot of the electronic DOS.

### 3.3 Phonon dispersion

The phonon dispersion of the material was computed as follows. First, a list of convenient  $q$ -points has to be generated. This is done by generating a list of  $k$ -points. To do so, we run

a standard SCF computation. However in this case the SCF procedure is dummy. Indeed, `nstep` and `nline` are set to 1, preventing **Abinit** to run more than one iteration during the computation, as explained in [18]. The input file is available in Appendix I. Note that `shiftk` was set to `[0 0 0]` and not `[0.5 0.5 0.5]` as in the previous computations. This is done to force **Abinit** to include the  $\Gamma$  point in the list to be generated.

Once the list of  $k$ -points is generated, we use it as a list of  $q$ -points for the further computations. The next step is to compute the phonon spectrum over the obtained  $q$ -points. A few datasets are used :

- The first one is a ground-state self-consistency computation that will be used for all the other datasets.
- The second dataset is used to compute the response function of the  $\frac{d}{dk}$  wave functions.
- The third one is used to compute the response function at  $\Gamma$  of an electric field perturbation.
- The other datasets are used to compute the phonon response at the other  $q$ -points. As only one  $q$ -point can be considered at a time, we need as many additional datasets as  $q$ -points.

After the run, several databases are generated, corresponding to the specified datasets. The input file is available in Appendix J. Those databases are merged and analyzed using the **MRGDDB** and **ANADDB** utilities. During the latter analysis, **ANADDB** computes different physical properties based on the databases containing the derivatives of the total energy (merged using **MRGDDB**). The properties to be computed are specified by a series of *flags*. For example :

- `ifcflag` is used to compute the interatomic force constants, which will be used to interpolate the phonon spectrum used in the band structure.
- `dipdip` is used to handle the dipole-dipole interactions separately from the interatomic forces, leading to more accurate results.
- `eivec` is used to generate an output file containing the frequencies for the different  $q$ -points, allowing to plot the phonon dispersion (referred to as the `_B2EPS.freq` file).
- `prtdos` allows to output a file containing the phonon density of states (see Figure 15).

The input file must also contain two specific lists of  $q$ -points. The first one is the  $q$ -path along which the phonon dispersion will be computed. A previous list of  $k$ -points used to plot the electronic band structure was used. The second list of  $q$ -points contains actually one single  $q$ -point :  $\Gamma$ . Indeed, a discontinuity can be seen in the phonon band structure at this specific point. The frequencies computed at this point will be hardcoded in the `_B2EPS.freq` file to get a proper plot<sup>2</sup>.

Then, using the generated `_B2EPS.freq` file containing the frequencies for all the  $q$ -points of the specified  $q$ -path, it is possible to produce a final file containing the band structure. It was plotted using **Abipy** (Figure 13) and was compared with the phonon dispersion plot provided by the Materials Project (Figure 14).

---

<sup>2</sup>Note that the discontinuity at  $\Gamma$  is still there on Figure 13 and Figure 14. Indeed, no package was able to open the hacked `.eps` file properly.

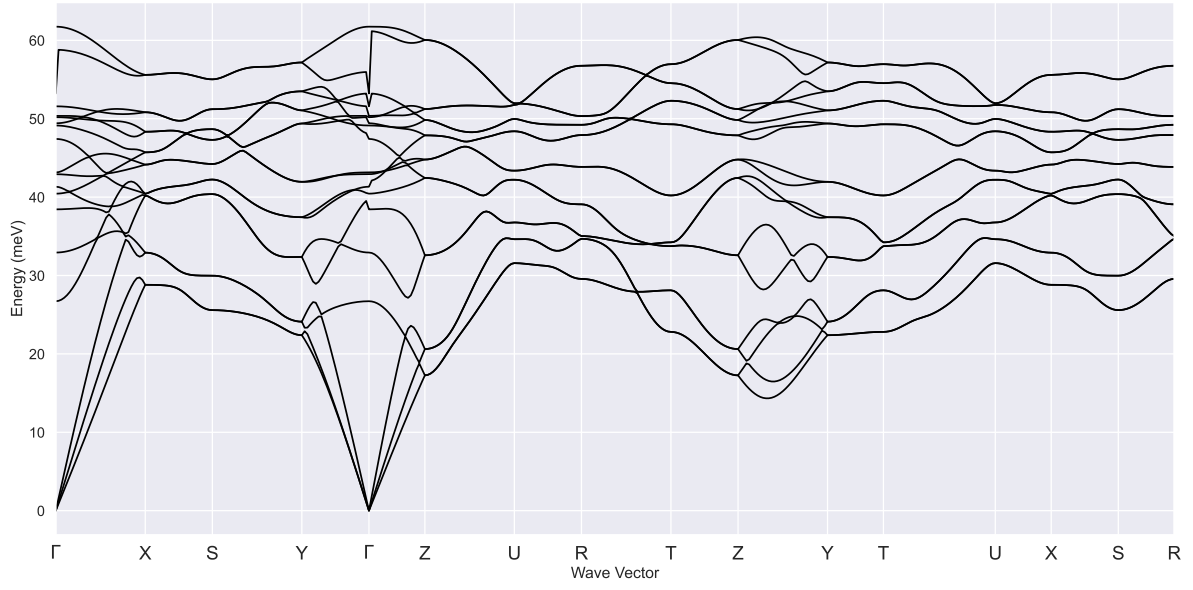


Figure 13: Computed phonon dispersion plotted with **Abipy**

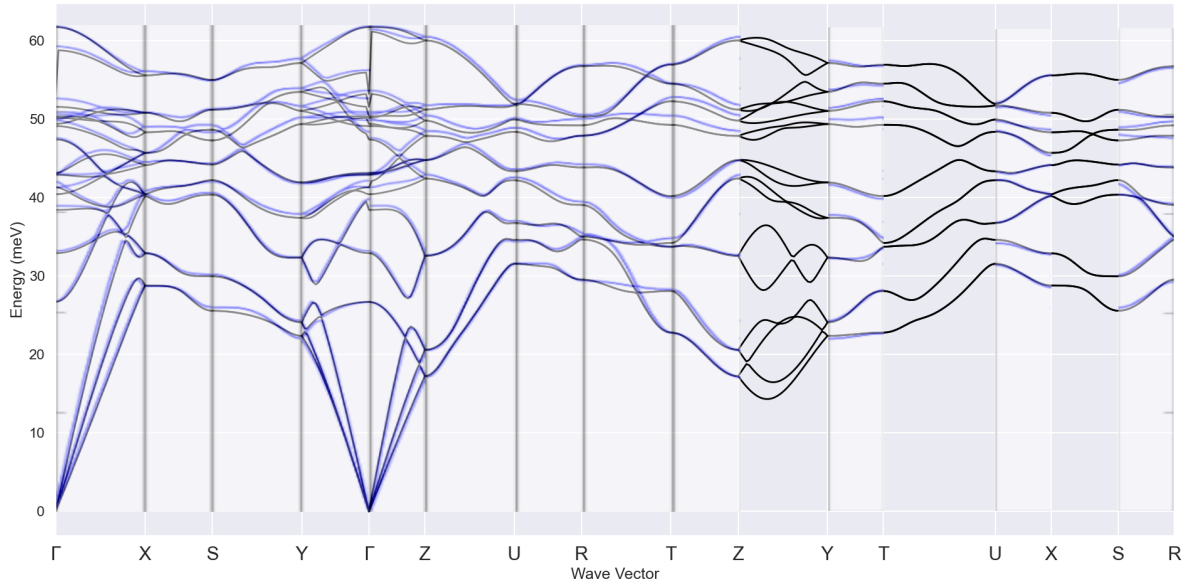


Figure 14: Comparison between the computed phonon dispersion (in black) and the phonon dispersion obtained on the Materials Project (in blue). The superposition of the plots is not complete as the plot provided by the Materials Project is truncated at some  $q$ -points.



### 3.4 Phonon density of states

By setting `prtdos = 2`, it is possible to compute the phonon DOS using the tetrahedron method. The plot is obtained with `Abipy`, and is compared in Figure 15 with the phonon DOS plot provided on the Materials Project. It can be seen that the results are in good agreement, as the plots show the same features, and only differ by the smearing level of the plot.

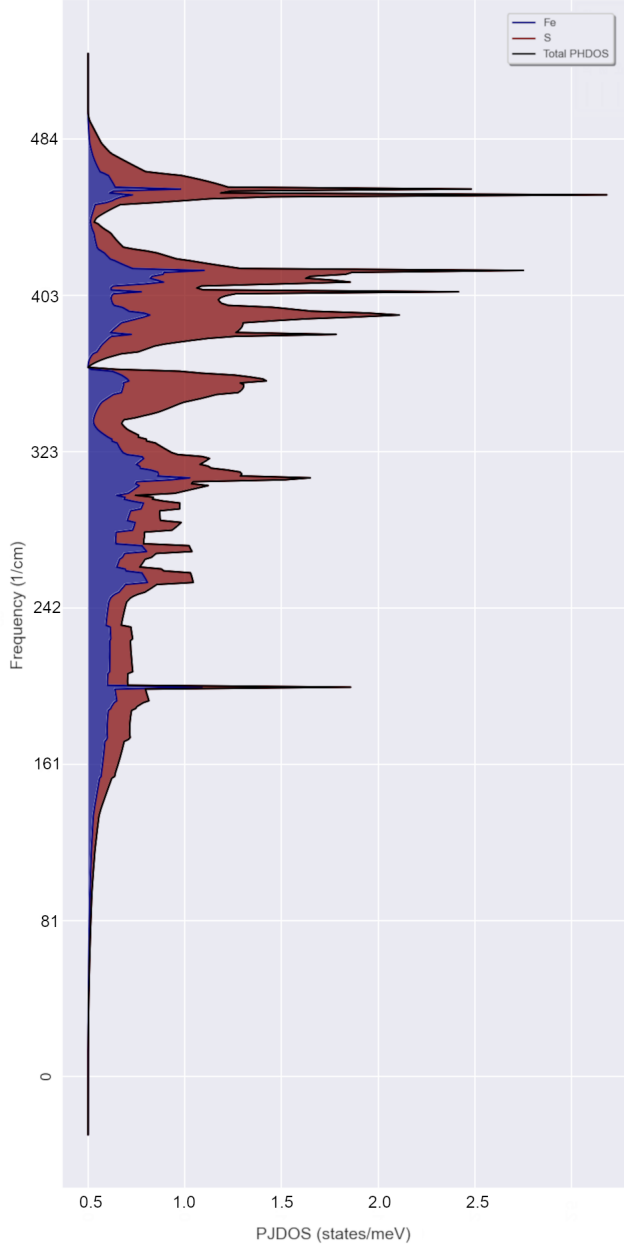
### 3.5 Absorption spectrum

### 3.6 Macroscopic consequences

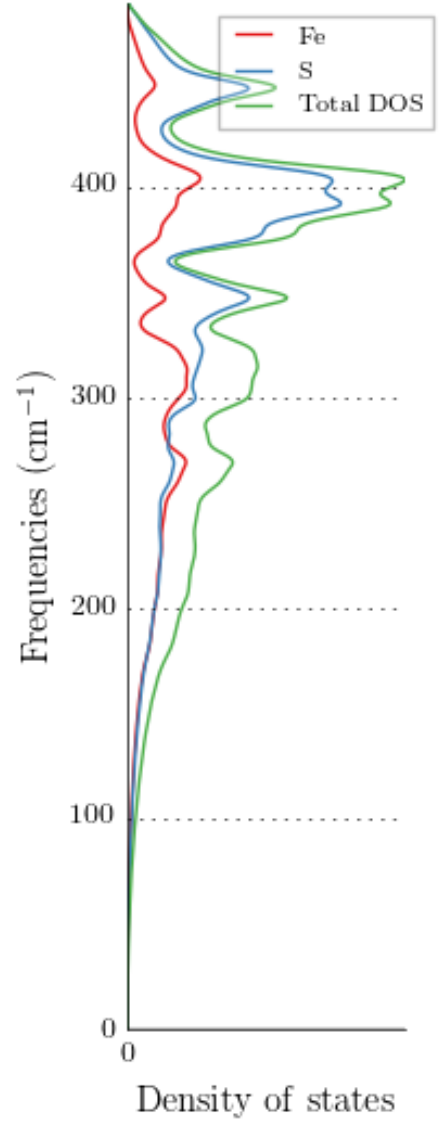
## 4 Conclusion

\\TODO

- Summary of the study - What has been learned ?
- What implications in real life applications ?
- What further studies could be done ?



(a) Computed phonon DOS plotted using Abipy



(b) Phonon DOS provided by the Materials Project

Figure 15: Comparison of the phonon DOS

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## A Convergence of the total energy per atom as a function of ecut

Name of the input file : 1522\_3\_ecutConv.abi

```
acell      3.390 4.438 5.411 Angstr
ntypat     2
znuc1      26 16
natom       6
typat      1 1 2 2 2 2
xred        0      0      0
            0.5    0.5    0.5
            0      0.206  0.3753
            0      0.794  0.6247
            0.5    0.294  0.8753
            0.5    0.706  0.1247
```

```
pseudos    "Fe.psp8,S.psp8"
```

```
# parameters of the SCF procedure :
```

```
nstep      100                      # maximal number of SCF cycles
toldfe     1.0d-10                  # SCF procedure will stop when the difference of total
                                     # energy between two iterations will be lower than
                                     # toldfe Hartree
diemac     24.0                     # preconditioning of the SCF procedure.
```

```
# parameters for generating the k-points grids :
```

```
kptopt     1
ngkpt      3 3 4
nshiftk     1
shiftk     0.5 0.5 0.5
```

```
ndtset     50
ecut:      10
ecut+      1
```

## B Convergence of the total energy per atom as a function of ngkpt

Name of the input file : 1522\_4\_nkpConv.abi

```
acell      3.390 4.438 5.411 Angstr
ntypat     2
znucl      26 16
natom       6
typat      1 1 2 2 2 2
xred        0      0      0
            0.5    0.5    0.5
            0      0.206  0.3753
            0      0.794  0.6247
            0.5    0.294  0.8753
            0.5    0.706  0.1247
ecut       40                                # the converged value for ecut

pseudos    "/Fe.psp8,S.psp8"

# parameters of the SCF procedure :
nstep      100                                # maximal number of SCF cycles
toldfe     1.0d-10                            # SCF procedure will stop when the difference of total
                                                # energy between two iterations will be lower than
                                                # toldfe Hartree
diemac     24.0                               # preconditioning of the SCF procedure.

# parameters for generating the k-points grids :
getwfk     -1
kptopt      1
ndtset      6
ngkpt1     2 2 2
ngkpt2     2 2 3
ngkpt3     3 3 3
ngkpt4     3 3 4
ngkpt5     4 4 4
ngkpt6     4 4 5
nshiftk     1
shiftk     0.5 0.5 0.5
```

## C Convergence of the lattice scale parameters as a function of ecut

Name of the input file : 1522\_6\_acellEcutConv.abi

```
acell      3.390 4.438 5.411 Angstr
ntypat     2
znuc1      26 16
natom      6
typat      1 1 2 2 2 2
xred       0      0      0
           0.5    0.5    0.5
           0      0.206  0.3753
           0      0.794  0.6247
           0.5    0.294  0.8753
           0.5    0.706  0.1247
```

```
ndtset     16
ecut:      20
ecut+      2
```

```
pseudos    "/Fe.psp8,S.psp8"
```

```
# parameters of the SCF procedure :
```

```
nstep      100                      # maximal number of SCF cycles
toldff      1.0d-6                  # SCF procedure will stop when the difference of total
                                     # energy between two iterations will be lower than
                                     # toldfe Hartree
diemac      24.0                    # preconditioning of the SCF procedure.
```

```
ionmov      2
optcell     1
dilatmx     1.05
ecutsm      0.5
kptopt      1
ngkpt       3 3 3
nshiftk     1
shiftk      0.5 0.5 0.5
getwfk      -1
```

## D Convergence of the lattice scale parameters as a function of ngkpt

Name of the input file : 1522\_3\_acellNgkptConv.abi

```
acell      3.390 4.438 5.411 Angstr
ntypat     2
znucl      26 16
natom       6
typat      1 1 2 2 2 2
xred        0      0      0
            0.5    0.5    0.5
            0      0.206  0.3753
            0      0.794  0.6247
            0.5    0.294  0.8753
            0.5    0.706  0.1247
```

```
ndtset      6
ngkpt1      2 2 2
ngkpt2      2 2 3
ngkpt3      3 3 3
ngkpt4      3 3 4
ngkpt5      4 4 4
ngkpt6      4 4 5
```

```
ecut        40
pseudos     "/Fe.psp8,S.psp8"
```

# parameters of the SCF procedure :

```
nstep       100                      # maximal number of SCF cycles
toldff      1.0d-6                   # SCF procedure will stop when the difference of total
                                     # energy between two iterations will be lower than
                                     # toldfe Hartree
diemac      24.0                     # preconditioning of the SCF procedure.
```

```
ionmov      2
optcell     1
dilatmx     1.05
ecutsm      0.5
nshiftk     1
shiftk      0.5 0.5 0.5
getwfk      -1
```



## E Band structure : $k$ -path generation using Absitrustruct (Abipy)

Command used : abistruct.py kpath input.abi

Input file : input.abi :

```
acell      3.315 4.3398 5.2913 Angstr # the converged values
ntypat     2
znucl      26 16
natom       6
typat      1 1 2 2 2 2
xred                                     # the converged values
          0  0      0
          0.5 0.5    0.5
          0  0.20809 0.37374
          0  0.79191 0.62626
          0.5 0.29191 0.87374
          0.5 0.70809 0.12626
pseudos    "/Fe.psp8,S.psp8"
nstep      100
toldfe     1.0d-10
diemac     24.0
kptopt      1
ngkpt      3 3 3
nshiftk     1
shiftk     0.5 0.5 0.5
getwfk     -1
```

```

Output :
# Abinit structure
natom      6
ntypat     2
typat      1 1 2 2 2 2
znuc1      26 16
xred                                              # the converged values
          0   0   0
          0.5 0.5   0.5
          0   0.20809 0.37374
          0   0.79191 0.62626
          0.5 0.29191 0.87374
          0.5 0.70809 0.12626
acell      1.0 1.0 1.0
rprim
          6.2644421031 0.0000000000 0.0000000000
          0.0000000000 8.2010334357 0.0000000000
          0.0000000000 0.0000000000 9.9991078432
# tolwfr 1e-20 iscf -2 # NSCF run
# To read previous DEN file, use:  getden -1 or specify filename via getden_path "out_DEN"
# K-PATH IN REDUCED COORDINATES:
ndivsm     10
kptopt     -15
kptbounds
          +0.00000 +0.00000 +0.00000          # $\Gamma$
          +0.50000 +0.00000 +0.00000          # X
          +0.50000 +0.50000 +0.00000          # S
          +0.00000 +0.50000 +0.00000          # Y
          +0.00000 +0.00000 +0.00000          # $\Gamma$
          +0.00000 +0.00000 +0.50000          # Z
          +0.50000 +0.00000 +0.50000          # U
          +0.50000 +0.50000 +0.50000          # R
          +0.00000 +0.50000 +0.50000          # T
          +0.00000 +0.00000 +0.50000          # Z
          +0.00000 +0.50000 +0.00000          # Y
          +0.00000 +0.50000 +0.50000          # T
          +0.50000 +0.00000 +0.50000          # U
          +0.50000 +0.00000 +0.00000          # X
          +0.50000 +0.50000 +0.00000          # S
          +0.50000 +0.50000 +0.50000          # R

```

## F Band structure computation

```

ndtset      2
natom       6
typas       1 1 2 2 2 2
xred
            0.0000000000 0.0000000000 0.0000000000
            0.5000000000 0.5000000000 0.5000000000
            0.0000000000 0.2080900000 0.3737400000
            0.0000000000 0.7919100000 0.6262600000
            0.5000000000 0.2919100000 0.8737400000
            0.5000000000 0.7080900000 0.1262600000
ecut         40.0                                     # converged value
# definition of the SCF procedure
nstep        100
diemac       24.0
# first dataset : SCF density computation
kptopt1      1
nshiftk1     1
shiftk1      0.5 0.5 0.5
ngkpt1       3 3 3                                     # converged value
prtden1      1
tolwfr1      1.0d-10
# second dataset : band structure computation
iscf2        -2
getden2      -1
kptopt2      -15                                     # number of k-segments
nband2       35                                     # number of bands
nbdbuf       -2
ndivsm2      30
kptbounds
            +0.00000 +0.00000 +0.00000             # $\Gamma$
            +0.50000 +0.00000 +0.00000             # X
            +0.50000 +0.50000 +0.00000             # S
            +0.00000 +0.50000 +0.00000             # Y
            +0.00000 +0.00000 +0.00000             # $\Gamma$
            +0.00000 +0.00000 +0.50000             # Z
            +0.50000 +0.00000 +0.50000             # U
            +0.50000 +0.50000 +0.50000             # R
            +0.00000 +0.50000 +0.50000             # T
            +0.00000 +0.00000 +0.50000             # Z
            +0.00000 +0.50000 +0.00000             # Y
            +0.00000 +0.50000 +0.50000             # T
            +0.50000 +0.00000 +0.50000             # U
            +0.50000 +0.00000 +0.00000             # X
            +0.50000 +0.50000 +0.00000             # S
            +0.50000 +0.50000 +0.50000             # R
tolwfr2      1.0d-12
enunit2      1                                     # to get the energies in eV

```

## G Band structure : Materials Project

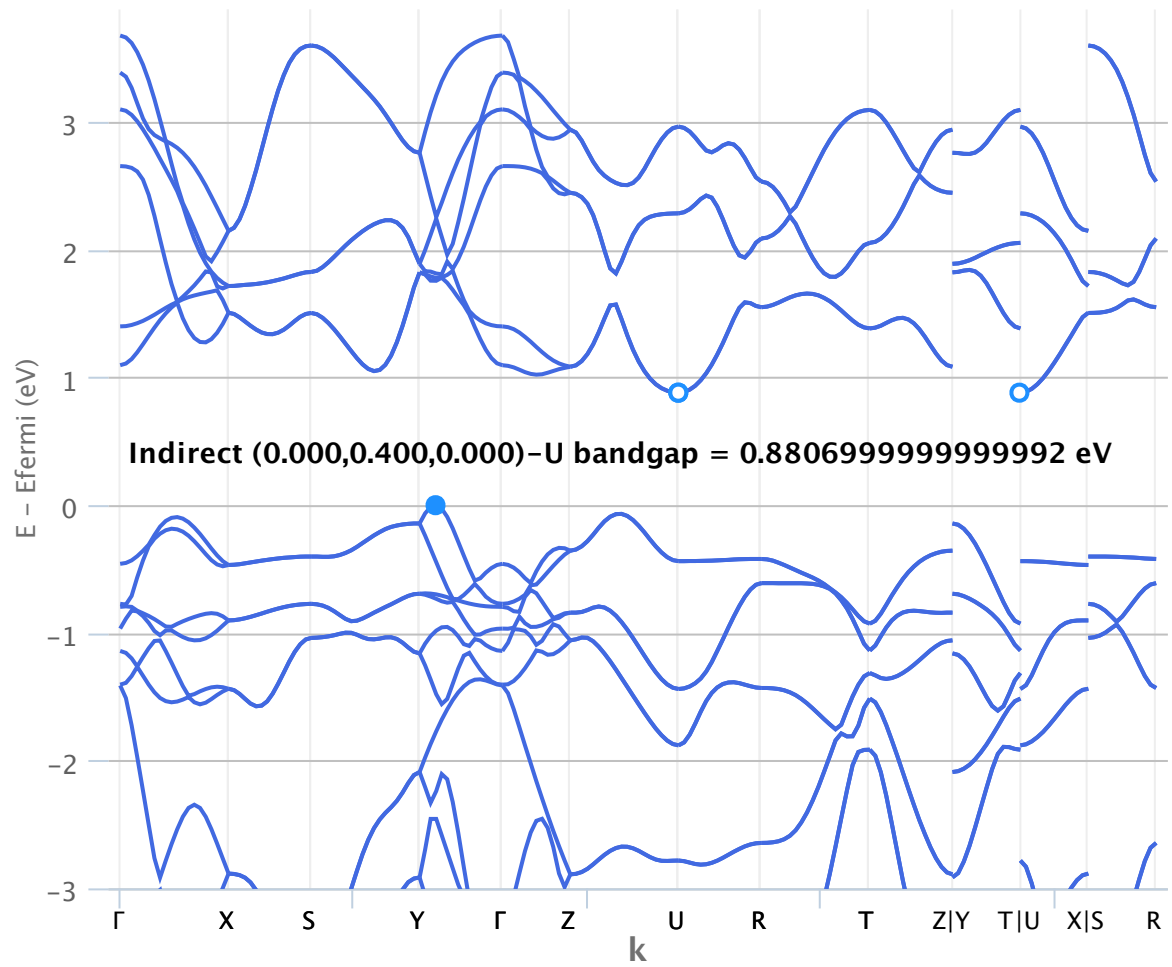


Figure 16: Band structure of orthorhombic FeS<sub>2</sub> provided by the Materials Project

## H Electronic density of states

```

# computation of the electronic DOS
ndtset      2
# definition of the unit cell
acell       1.0 1.0 1.0
rprim
            6.2644421031 0.0000000000 0.0000000000
            0.0000000000 8.2010334357 0.0000000000
            0.0000000000 0.0000000000 9.9991078432
# definition of the atoms types
ntypat      2
znucl       26 16
pseudos     "Fe.psp8,S.psp8"
# definition of the atoms
natom       6
typat       1 1 2 2 2 2
xred
            0.0000000000 0.0000000000 0.0000000000
            0.5000000000 0.5000000000 0.5000000000
            0.0000000000 0.2080900000 0.3737400000
            0.0000000000 0.7919100000 0.6262600000
            0.5000000000 0.2919100000 0.8737400000
            0.5000000000 0.7080900000 0.1262600000
ecut        40.0                                     # converged value
# first dataset : SCF density computation
# definition of the SCF procedure
tolddf1     1.0d-10
nstep       100
diemac      24.0
# Brillouin zone sampling
kptopt1     1
nshiftk1    1
shiftk1     0 0 0                                     # use of a non-shifted k-grid
ngkpt1      8 8 8                                     # higher than the converged value to
                                                         # obtain a good resolution in the
                                                         # DOS plot

prtden1     1
# second dataset : DOS computation
prtdos2     2                                     # computing the DOS using the
                                                         # wtetrahedron method

getden2     -1
iscf2       -3
tolwfr2     1.0d-12
dosdeltae2  5.0d-5

```

## I Phonons : Generation of a list of $q$ -points

```
# Pnm [58] FeS2 : computation of the q points
ngkpt    3 3 3
nshiftk  1
shiftk    0 0 0
# dummy SCF procedure
nstep    1
nline    1
diemac   24.0
# common input variables
acell    1.0 1.0 1.0
rprim
          6.2644421031 0.0000000000 0.0000000000
          0.0000000000 8.2010334357 0.0000000000
          0.0000000000 0.0000000000 9.9991078432
ntypat   2
znucl    26 16
pseudos  "Fe.psp8,S.psp8"
natom     6
typat    1 1 2 2 2 2
xred
          0.0000000000 0.0000000000 0.0000000000
          0.5000000000 0.5000000000 0.5000000000
          0.0000000000 0.2080900000 0.3737400000
          0.0000000000 0.7919100000 0.6262600000
          0.5000000000 0.2919100000 0.8737400000
          0.5000000000 0.7080900000 0.1262600000
ecut     40.0                                     # converged value
nband    35
tolvrs   1.0d-18                                # dummy variable as there is only one
                                                # step in the SCF procedure
```

## J Phonons : Generation of the derivative databases

```
ndtset    10
# dataset 1 :  GS self-consistent computation
getwfk1   0
kptopt1   1
nqpt1     0
tolvrs1   1.0d-18
rfphon1    0
# Q vectors for all datasets
nqpt      1
qpt2      0.00000000E+00 0.00000000E+00 0.00000000E+00
qpt3      0.00000000E+00 0.00000000E+00 0.00000000E+00
qpt4      3.33333333E-01 0.00000000E+00 0.00000000E+00
qpt5      0.00000000E+00 3.33333333E-01 0.00000000E+00
qpt6      3.33333333E-01 3.33333333E-01 0.00000000E+00
qpt7      0.00000000E+00 0.00000000E+00 3.33333333E-01
qpt8      3.33333333E-01 0.00000000E+00 3.33333333E-01
qpt9      0.00000000E+00 3.33333333E-01 3.33333333E-01
qpt10     3.33333333E-01 3.33333333E-01 3.33333333E-01
# dataset 2 :  response function calculation of d/dk wave function
iscf2     -3
kptopt2    2
rfphon2    0
rfelld2    2
tolwfr2    1.0d-22
# dataset 3 :  response function calculation at Q=0 phonons and electric field perturbation
getddk3    2
kptopt3    2
rfelld3    3
```

```

# datasets 4-10 :  finite-wave-vector phonon calculations (default for all datasets)
getwfk    1
kptopt    3
rfphon    1
rfatpol   1 6
rfdir     1 1 1
tolvrs    1.0d-18
ntypat    2
znucl     26 16
pseudo    "Fe.psp8,S.psp8"
natom     6
typat     1 1 2 2 2 2
xred       0.0000000000 0.0000000000 0.0000000000
           0.5000000000 0.5000000000 0.5000000000
           0.0000000000 0.2080900000 0.3737400000
           0.0000000000 0.7919100000 0.6262600000
           0.5000000000 0.2919100000 0.8737400000
           0.5000000000 0.7080900000 0.1262600000
ecut      40.0
nband     35
diemac    24.0
ngkpt     3 3 3
nshiftk   1
shiftk    0.5 0.5 0.5

```



## K Phonons : Computation of the band structure

```
ifcflag  1
ifcout   0
brav      1
ngqpt    3 3 3
nqshift   1
q1shift  0.0 0.0 0.0
dipdip    1
eivec     4
prtdos    2
nph1l     401
# the q-path is not written for visual reasons
# \Gamma-X-S-Y-\Gamma-Z-U-R-T-Z-Y-T-U-X-S-R
# retrieved from the electronic bandstructure computation
nph2l     1
qph2l     1.0 1.0 1.0
ng2qpt    3 3 3
```