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# **SALMON document**

***Release v.2.0.2***

**SALMON developers**

**Nov 06, 2021**



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SALMON (Scalable Ab initio Light-Matter simulator for Optics and Nanoscience) is an open-source software based on first-principles time-dependent density functional theory to describe optical responses and electron dynamics in matters induced by light electromagnetic fields.



## INTRODUCTION

### 1.1 Overview

SALMON is an open-source computer program for ab-initio quantum-mechanical calculations of electron dynamics at the nanoscale that takes place in various situations of light-matter interactions. It is based on time-dependent density functional theory, solving time-dependent Kohn-Sham equation in real time and real space with norm-conserving pseudopotentials.

SALMON was born by unifying two scientific programs: ARTED, developed by Univ. Tsukuba group, that describes electron dynamics in crystalline solids, and GCEED, developed by Institute for Molecular Science group, that describes electron dynamics in molecules and nanostructures. It can thus describe electron dynamics in both isolated and periodic systems. It can also describe coupled dynamics of electrons and light-wave electromagnetic fields.

To run the program, SALMON requires MPI Fortran/C compiler with LAPACK libraries. SALMON has been tested and optimized to run in a number of platforms, including Linux PC Cluster with x86-64 CPU, supercomputer systems with Fujitsu FX100 and A64FX processors, and supercomputer system with Intel Xeon Phi (Knights Landing).

### 1.2 SALMON features

In the microscopic scale, SALMON describes electron dynamics in both isolated (molecules and nanostructures) and periodic (crystalline solids) systems, solving time-dependent Kohn-Sham equation in real time and real space with norm-conserving pseudopotential. SALMON first carries out ground-state calculations in the density functional theory to prepare initial configurations. SALMON then calculates electron dynamics induced by applied electric field. Employing a weak impulsive external field, SALMON can be used to calculate linear response properties such as a polarizability of molecules and a dielectric function of crystalline solids. Using pulsed electric fields, SALMON describes electron dynamics in matters induced by intense and ultrashort laser pulses.

SALMON is also capable of describing a propagation of electromagnetic fields of light using finite-difference time-domain method. As a unique feature of SALMON, it is possible to carry out calculations of a coupled dynamics of light electromagnetic fields and electron dynamics simultaneously.

Efficient parallelizations are implemented in the code by dividing spatial grids, orbital index, and k-points. SALMON shows a good scalability when it runs in parallel supercomputers, both for the ground state and the time evolution calculations.

- Ground state calculations
  - Kohn-Sham orbitals and energies
  - density of states
  - projected density of states
  - electron localization function

- Optical properties
  - Oscillator strength distribution (absorption spectrum)
  - dielectric function
- Light-induced electron dynamics
  - time evolution of Kohn-Sham orbitals
  - density, current
  - excitation energy
  - number density of excited carriers
- Propagation of light electromagnetic fields
  - Drude-Lorentz model
  - optical response of metasurfaces
- Simultaneous description of electron dynamics and light pulse propagation
  - light pulse propagation as well as time evolution of Kohn-Sham orbitals
  - energy transfer from pulsed light to electrons

## 1.3 License

SALMON is available under Apache License version 2.0.

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## 1.4 SALMON at Github

SALMON is developed at [GitHub.com](https://github.com)

## 1.5 List of developers

(Alphabetic order)

- Isabella Floss (TU Wien, Austria)
- Yuta Hirokawa (Prometech Software, Inc., Japan)
- Kenji Iida (Hokkaido University, Japan)
- Jun-Ichi Iwata (Advance Soft Co., Japan)



- Masashi Noda (Academeia, Japan)
- Tomohito Ootobe (National Institutes for Quantum and Radiological Science and Technology, Japan)
- Shunsuke Sato (University of Tsukuba, Japan)
- Yasushi Shinohara (University of Tokyo, Japan)
- Takashi Takeuchi (University of Tsukuba, Japan)
- Mitsuharu Uemoto (Kobe University, Japan)
- Kazuhiro Yabana (University of Tsukuba, Japan)
- Atsushi Yamada (University of Tsukuba, Japan)
- Shunsuke Yamada (University of Tsukuba, Japan)

## 1.6 Former developers

- Kazuya Ishimura
- Kyung-Min Lee
- Katsuyuki Nobusada
- Xiao-Min Tong
- Maiku Yamaguchi

## 1.7 How to cite SALMON

### 1.7.1 Suggested Citations

If you publish a paper in which SALMON makes an important contribution, please cite the SALMON code paper, Ref. [1] published in Computer Physics Communications.

We also suggest you to cite the following papers depending on your usage of SALMON.

- If you use SALMON for electron dynamics calculations of a large-size system, Ref. [2] that discusses massively parallel implementation utilizing spatial divisions will be appropriate.
- if you use SALMON to calculate electron dynamics in a unit cell of crystalline solid, Ref. [3] discussing formalism and numerical implementation will be appropriate.
- Ref. [4] is one of the first implementations of the real-time time-dependent density functional calculation, in particular, instantaneous kick for the linear response calculations.
- If you use multiscale calculation coupling Maxwell equations for the electromagnetic fields of light and electron dynamics, Ref. [5] discussing the formalism and the numerical implementation will be appropriate.
- Ref. [6] describes parallelization method for the coupled Maxwell - TDDFT calculations.
- Ref. [7] describes computational aspects of electron dynamics calculations for periodic systems in many-core processors:



## INSTALL AND RUN

### 2.1 Prerequisites

In this guide, it is assumed that readers have a basic knowledge of Linux and its command line operations. For the installation of SALMON, following packages are required.

- Fortran90/C compiler. SALMON assumes users have one of the following compilers:
  - GCC (Gnu Compiler Collection)
  - Intel Compiler
  - Fujitsu Compiler (at FX100 and A64FX)
- One of the following library packages for linear algebra:
  - Netlib BLAS/LAPACK/ScaLAPACK
  - Intel Math Kernel Library (MKL)
  - Fujitsu Scientific Subroutine Library 2 (SSL-II)
- Build tools:
  - CMake

If you use other compilers, you may need to change build scripts (CMake). See *Additional options in configure.py script*. If no numerical library is installed on your computer system, you may need to install BLAS/LAPACK by yourself. See *Troubleshooting of the Installation Process*.

For the installation of SALMON, we adopt the CMake tools as the first option. If there were any problems to use CMake tools in your environment, you may use the GNU make tools. See *Troubleshooting of the Installation Process*.

### 2.2 Download

The newest version of SALMON can be downloaded from [download page](#). You can also get the file by

```
$ wget http://salmon-tddft.jp/download/SALMON-v.2.0.1.tar.gz
```

To extract files from the downloaded file `SALMON-<VERSION>.tar.gz`, type the following command in the command-line:

```
$ tar -zxvf ./SALMON-<VERSION>.tar.gz
```

After the extraction, the following directories will be created:

```
SALMON
|- src           Source codes
|- example       Samples
|- cmakefiles    CMake related files
|- gnumakefiles  GNU Makefiles for building
```

## 2.3 Build and Install

To compile SALMON to create executable the binary files, we adopt to use CMake tools as the first option. In case you fail to build SALMON using CMake in your environment, we may use Gnu Make. See [Build using GNU Makefile](#).

### 2.3.1 Checking CMake availability

First, examine whether CMake is usable in your environment or not. Type the following in Linux command-line:

```
$ cmake --version
```

If CMake is not installed in your system, an error message such as `cmake: command not found` will appear. If CMake is installed on your system, the version number will be shown. To build SALMON, CMake of version 3.14.0 or later is required. If you confirm that CMake of version 3.14.0 or later is installed in your system, proceed to [Build using CMake](#). However, we realize that old versions of CMake are installed in many systems. If CMake is not installed or CMake of older versions is installed in your system, you need to install the new version by yourself. It is a simple procedure and explained below.

### 2.3.2 Installation of CMake (pre-compiled binary of Linux)

CMake is a cross-platform build tool. The simplest way to make CMake usable in your environment is to get the [binary distribution of CMake from the download page](#). (The file name of the binary distribution will be `cmake-<VERSION>-<PLATFORM>.tar.gz`). In standard Linux environment, a file for the platform of Linux x86\_64 will be appropriate.

To download the file, proceed as follows: We assume that you are in the directory that you extracted files from the downloaded file of SALMON, and that you will use the version 3.16.8. First get the URL of the download link from your browser, and use `wget` command in your Linux command-line:

```
$ wget https://cmake.org/files/v3.16/cmake-3.16.8-Linux-x86_64.tar.gz
```

Next, unpack the archive by:

```
$ tar -zxvf cmake-3.16.8-Linux-x86_64.tar.gz
```

and you will have the binary `make-3.16.8-Linux-x86_64/bin/cmake` in your directory.

To make the `cmake` command usable in your command-line, you need to modify the environment variable `$PATH` so that the executable of CMake are settled inside the directory specified in your `$PATH`. If you use the bash shell, you need to modify the file `~/.bashrc` that specifies the `$PATH` variable. It can be done by typing the following command in your login directory:

```
$ export PATH=<SALMON_INSTALLATION_DIRECTORY>/cmake-3.16.8-Linux-x86_64/bin:$PATH
```

and then reload the configuration by typing:

```
$ source ~/.bashrc
```

See *Installation of CMake* describes Other way of the installation.

## 2.3.3 Build using CMake

Confirming that CMake of version 3.14.0 or later can be usable in your environment, proceed the following steps. We assume that you are in the directory SALMON.

1. Create a new temporary directory `build` and move to the directory:

```
$ mkdir build
$ cd build
```

2. Execute the python script "configure.py" and then make:

```
$ python ../configure.py --arch=ARCHITECTURE --prefix=../
$ make
$ make install
```

In executing the python script, you need to specify `ARCHITECTURE` that indicates the architecture of the CPU in your computer system such as `intel-avx`. The options of the `ARCHITECTURE` are as follows:

arch	Detail	Compiler	Numerical Library
intel-knl	Intel Knights Landing	Intel Compiler	Intel MKL
intel-knc	Intel Knights Corner	Intel Compiler	Intel MKL
intel-avx	Intel Processor (Ivy-, Sandy-Bridge)	Intel Compiler	Intel MKL
intel-avx2	Intel Processor (Haswell, Broadwell ..)	Intel Compiler	Intel MKL
intel-avx512	Intel Processor (Skylake-SP)	Intel Compiler	Intel MKL
fujitsu-fx100	FX100 Supercomputer	Fujitsu Compiler	SSL-II
fujitsu-a64fx-ea	A64FX processor (Fugaku, FX1000, FX700)	Fujitsu Compiler	SSL-II

If the build is successful, you will get a file `salmon` at the top-level build directory.

## 2.4 Files necessary to run SALMON

To run SALMON, at least two kinds of files are required for any calculations. One is an input file with the filename extension `*.inp` that should be read from the standard input `stdin`. This file should be prepared in the Fortran90 namelist format. Pseudopotential files of relevant elements are also required. Depending on your purpose, some other files may also be necessary. For example, coordinates of atomic positions of the target material may be either written in the input file or prepared as a separate file.

### 2.4.1 Pseudopotentials

SALMON utilizes norm-conserving (NC) pseudopotentials. Filenames of pseudopotentials should be written in the input file.

You may find pseudopotentials of some elements in the samples prepared in *Exercises*. In SALMON, several formats of pseudopotentials may be usable (listed below). For example, pseudopotentials with an extension `.fhi` can be obtained from the ABINIT website (this is a part of previous atomic data files for the ABINIT code).

Pseudopotential	ex- ten- sion	Website
Fritz-Haber-Institute (FHI) pseudopotentials	. fhi	<a href="https://www.abinit.org/sites/default/files/PrevAtomicData/psp-links/lda_fhi.html">https://www.abinit.org/sites/default/files/PrevAtomicData/psp-links/lda_fhi.html</a> (for LDA), <a href="https://www.abinit.org/sites/default/files/PrevAtomicData/psp-links/gga_fhi.html">https://www.abinit.org/sites/default/files/PrevAtomicData/psp-links/gga_fhi.html</a> (for GGA)
Pseudopotentials for the OpenMX code	. vps	<a href="https://t-ozaki.issp.u-tokyo.ac.jp/vps_pao2019/">https://t-ozaki.issp.u-tokyo.ac.jp/vps_pao2019/</a>
Format 8 for ABINIT norm-conserving pseudopotentials	. psp8	<a href="https://www.abinit.org/psps_abinit">https://www.abinit.org/psps_abinit</a> , <a href="http://www.pseudo-dojo.org/">http://www.pseudo-dojo.org/</a>
Unified-pseudopotential-format (NC type only in SALMON)	. upf	<a href="http://www.quantum-espresso.org/pseudopotentials/unified-pseudopotential-format">http://www.quantum-espresso.org/pseudopotentials/unified-pseudopotential-format</a> , <a href="http://www.pseudo-dojo.org/">http://www.pseudo-dojo.org/</a>

## 2.4.2 input file

Input files are composed of several blocks of namelists:

```
&namelist1
  variable1 = int_value
  variable2 = 'char_value'
/
&namelist2
  variable1 = real8_value
  variable2 = int_value1, int_value2, int_value3
/
```

A block of namelists starts with `&namelist` line and ends with `/` line. The blocks may appear in any order.

Between two lines of `&namelist` and `/`, descriptions of variables and their values appear. Note that many variables have their default values so that it is not necessary to give values for all variables. Descriptions of the variables may appear at any position if they are between `&namelist` and `/`.

SALMON describes electron dynamics in systems with both isolated and periodic boundary conditions. The boundary condition is specified by the variable `iperiodic` in the namelist `&system`.

Calculations are usually achieved in two steps; first, the ground state calculation is carried out and then electron dynamics calculations in real time is carried out. A choice of the calculation mode or theory in the calculation is specified by the variable `theory` in the namelist `&calculation`. In the typical way, the ground state calculation based on DFT is first carried out specifying `theory = 'dft'`. Then the real-time electron dynamics calculation based on TDDFT is carried out specifying `theory = 'tddft_pulse'`.

In *Exercises*, we prepare six exercises that cover typical calculations feasible by SALMON. We prepare explanations of the input files of the exercises that will help to prepare input files of your own interests.

There are more than 20 groups of namelists. A complete list of namelist variables is given in the file `SALMON/manual/input_variables.md`. Namelist variables that are used in our exercises are explained at *Inputs*.

## 2.5 Run SALMON

Before running SALMON, the following preparations are required as described above: The executable file of `salmon` should be built from the source file of SALMON. An input file `inputfile.inp` and pseudopotential files should also be prepared.

The execution of the calculation can be done as follows: In single process environment, type the following command:

```
$ salmon < inputfile.inp > fileout.out
```

In multiprocess environment in which the command to execute parallel calculations using MPI is `mpiexec`, type the following command:

```
$ mpiexec -n NPROC salmon < inputfile.inp > fileout.out
```

where NPROC is the number of MPI processes that you will use.

The execution command and the job submission procedure depends much on local environment. We summarize general conditions to execute SALMON:

- SALMON runs in both single-process and multi-process environments using MPI.
- Executable file is prepared as `salmon` in the standard build procedure.
- To start calculations, `inputfile.inp` should be read through `stdin`.

### 2.5.1 MPI process distribution

SALMON provides three variables to determine the process distribution/allocation.

- `nproc_k`
- `nproc_ob`
- `nproc_rgrid(3)`

In SALMON, the process distribution is determined automatically as default. However, in many situations, an explicit assignment of the process distribution will provide a better performance than the default setting.

We recommend to distribute the processes as follows,

If you use k-points ( the number of k-points is greater than 1) and the number of the real-space grid (`num_rgrid`) is not very large (about  $16^3$ ):

- First, assign many processes to `nproc_k`.
- Then, assign the remaining processes to `nproc_ob`.
- Not dividing the spatial grid, `nproc_rgrid = 1, 1, 1`.

Else:

- First, assign the processes to `nproc_ob`.
- Then, assign the remaining processes to `nproc_rgrid`.
  - If real-space grid size (`num_rgrid(1:3) = a1(1:3) / d1(1:3)`) is equal to or larger than about  $64^3$ ,

you should find a balanced distribution between `nproc_rgrid` and `nproc_ob`.

## 2.6 Tips for large-scale calculation

We explain below some tips that will be useful to improve performance when you carry out large scale simulations using world top-level supercomputers. Therefore, the following contents will only be useful only for limited users.

### 2.6.1 Improve the performance of the eigenvalues solver

In DFT calculations of large systems, subspace diagonalization becomes the performance bottleneck in the entire calculation. Therefore, it is important to use a parallel eigenvalues solver. In SALMON, a LAPACK routine without parallelization is used for the diagonalization as default. As parallelized solvers, ScaLAPACK and EigenExa are usable. To use them, it is necessary to rebuild SALMON enabling ScaLAPACK/EigenExa. You can find the instruction in *Install and Run*.

To execute SALMON using ScaLAPACK/EigenExa, either `yn_scalapack = 'y'` or `yn_eigenexa = 'y'` should be included in the inputfile:

```
&parallel
  yn_scalapack = 'y'          ! use ScaLAPACK for diagonalization
  !yn_eigenexa  = 'y'          ! use EigenExa
  yn_scalapack_red_mem = 'y' ! to reduce the memory consumption
/
```

ScaLAPACK/EigenExa solves the eigenvalue problem with `nproc_ob` process distribution. If `nproc_ob = 1`, ScaLAPACK/EigenExa will perform in the same way as the LAPACK library.

### 2.6.2 Improve the performance of Hartree solver

For periodic systems, a Fourier transformation is used to solve the Poisson equation (to calculate the Hartree potential). In SALMON, a simple Fourier transformation without Fast Fourier Transformation (FFT) is used as default. In SALMON, a parallelized FFT routine, FFTE, is usable and works efficiently for large systems. In using FFTE, the following conditions should be satisfied:

```
num_rgrid(1) mod nproc_rgrid(2) = 0
num_rgrid(2) mod nproc_rgrid(2) = 0
num_rgrid(2) mod nproc_rgrid(3) = 0
num_rgrid(3) mod nproc_rgrid(3) = 0
```

In addition, the prime factors **for** the number of real-space grid of each direction,  $\rightarrow$  (num\_rgrid(1:3)) must be a combination of 2, 3 **or** 5.

To use FFTE, `yn_ffte = 'y'` should be included in the input file:

```
&parallel
  yn_ffte = 'y'
/
```

### 2.6.3 Improve IO performance (write/read wavefunction)

Almost all supercomputer systems provide distributed filesystems such as Lustre. Distributed filesystems are equipped with a meta-data server (MDS) and an object-storage server (OST). The OST stores real user data files, and the MDS stores the address of the user data files in the OST. When accessing to the data files in the OST, the process send a query about the OST address to MDS. Then, a network contention may occur in the query process.

In most implementations of the filesystem, the MDS that replies to the query is determined by the directory structure. For a calculation in which k-point is not used, `method_wf_distributor` and `nblock_wf_distribute` are prepared to reduce the network contention:

```
&control
  method_wf_distributor = 'slice' ! every orbital function is stored as a single file.
```

(continues on next page)



(continued from previous page)

```

nblock_wf_distribute = 32      ! files of 32 orbital functions are stored in one_
→directory.
/

```

## 2.6.4 Improve the communication performance for mesh-torus network system

Large-scale supercomputers often adopt a mesh-torus network system such as Cray dragon-fly and Fujitsu Tofu to achieve high scalability with relatively low cost. In SALMON, a special MPI process distribution (communicator creation rule) is prepared to improve the performance in large-scale mesh-torus network systems.

Currently, we provide the communicator creation rule for "Supercomputer Fugaku", which is developed by RIKEN R-CCS and Fujitsu limited. Fugaku is equipped with a 6-D mesh-torus network which is called "Tofu-D". Users may control it as a 3-D logical network. SALMON utilizes 5-D array (wavefunction(x, y, z, orbital, k-point)) as a domain for parallelization. We create a map that connects the 3-D network to the 5-D array distribution.

We introduce the following variables and conditions to assign the 3-D mesh-torus network to the 5-D array distribution:

```

PW          = nproc_ob * nproc_k
(PX, PY, PZ) = nproc_rgrid
PPN         = '# of process per node' (we recommend the value 4 in Fugaku)

Requested process shape: (PX, PY, PZ, PW)
Tofu-D network      shape: (TX, TY, TZ)
Actual process      shape: (TX * PPN, TY, TZ)

if (process_allocation == 'grid_sequential'):
    PW = PW1 * PW2 * PW3
    PW1 = (TX * PPN) / PX
    PW2 = TY / PY
    PW3 = TZ / PZ
    TX = (PX * PW1) / PPN
    TY = PY * PW2
    TZ = PZ * PW3

else if (process_allocation == 'orbital_sequential'):
    PX = PX1 * PX2 * PX3
    PX1 = (TX * PPN) / PW
    PX2 = TY / PY
    PX3 = TZ / PZ
    TX = (PW * PX1) / PPN
    TY = PY * PX2
    TZ = PZ * PX3

```

From these conditions, you can determine the suitable process distribution and the Tofu-D network shape (compute node shape). `process_allocation` input variable controls the order of the process distribution. It indicates which communications should be executed in closer processes.

- `process_allocation = 'grid_sequential'`
  - (PX, PY, PZ, PW), `nproc_rgrid` major ordering
  - improves `nproc_rgrid` related communication performance
  - communicator: `s_parallel_info::icomm_r, icomm_x, icomm_y, icomm_z, icomm_xy`
  - suitable theory: 'dft' and 'dft\_md'

- `process_allocation = 'orbital_sequential'`
  - (PW, PY, PZ, PX), `nproc_ob` major ordering
  - improves `nproc_ob` related communication performance
  - communicator: `s_parallel_info::icomm_o` and `icomm_ko`
  - suitable theory: `'tddft_response', 'tddft_pulse', 'single_scale_maxwell_tddft'` and `'multi_scale_maxwell_tddft'`

## 2.7 Troubleshooting of the Installation Process

### 2.7.1 Installation of CMake

The **CMake** is a cross-platform build tool. In order to build the SALMON from the source code, the CMake of version 3.14.0 or later is required. You may install it following one of the three instructions below.

#### Installation by package manager

If your system has a built-in package manager, you may conveniently install the CMake tools as below:

##### Debian/Ubuntu Linux

```
sudo apt-get install cmake
```

##### Fedora Linux/CentOS

```
sudo yum install cmake
```

##### openSUSE Linux

```
sudo zypper install cmake
```

#### Installation from source code

You can get the source code distribution from the [download page](#). In this time, we will use the cmake version 3.16.8 as an example. Download the archive by `wget` command and unpack it as below:

```
wget https://cmake.org/files/v3.16/cmake-3.16.8.tar.gz
tar -zxvf cmake-3.16.8.tar.gz
```

And, move to the unpacked directory and build.

```
cd cmake-3.16.8
./configure --prefix=INSTALLATION_DIRECTORY
make
make install
```

(replace `INSTALLATION_DIRECTORY` to your installation directory.)

Next, to utilize the `cmake` command, it is required that the executable are settled inside the directory specified in your `$PATH`. If you use the bash shell, edit `~/.bashrc` and append the line:

```
export PATH=INSTALLATION_DIRECTORY/bin:$PATH
```

and reload the configuration:

```
source ~/.bashrc
```

## 2.8 Appendix

### 2.8.1 Additional options in configure.py script

#### Manual specifications of compiler and environment variables

In executing `configure.py`, you may manually specify compiler and environment variables instead of specifying the architecture, for example:

```
$ python ../configure.py FC=mpiifort CC=mpiicc FFLAGS="-xAVX" CFLAGS="-restrict -xAVX"
```

The list of options of `configure.py` can be found by:

```
$ python ../configure.py --help
```

The major options are as follows:

Commandline switch	Detail
-a ARCH, -arch=ARCH	Target architecture
-enable-mpi, -disable-mpi	enable/disable MPI parallelization
-enable-scalapack, -disable-scalapack	enable/disable computations with ScaLAPACK library
-enable-eigenexa, -disable-eigenexa	enable/disable computations with RIKEN R-CCS EigenExa library
-enable-libxc, -disable-libxc	enable/disable computations with Libxc library
-with-lapack	specified LAPACK/ScaLAPACK installed directory
-with-libxc	specified Libxc installed directory
-debug	enable debug build
-release	enable release build
FC, FFLAGS	User-defined Fortran Compiler, and the compiler options
CC, CFLAGS	User-defined C Compiler, and the compiler options

In the build procedure by CMake, they search the following libraries. If the libraries don't found in the path that is specified by environment variables, they will build the required libraries automatically.

- Netlib LAPACK (includes BLAS), and ScaLAPACK
  - We will download and build the Netlib libraries as the typical implementation.
  - <http://www.netlib.org/lapack/>
  - <http://www.netlib.org/scalapack/>
- Libxc
  - <https://www.tddft.org/programs/libxc/>

EigenExa will download and build automatically even if the library is installed to your machine.

### Build for single process calculations

When using the `--arch` option, MPI parallelization is enabled as default. If you use a single processor machine, explicitly specify `--disable-mpi` in executing the python script:

```
$ python ../configure.py --arch=<ARCHITECTURE> --disable-mpi
```

### Build by user-specified compiler

If you want that specify the compiler, set the `FC` and `CC` flags in executing the python script:

```
$ python ../configure.py FC=gfortran CC=gcc
```

When `--arch` option is not used, MPI parallelization is disabled as default.

## 2.8.2 Build using GNU Makefile

If CMake build fails in your environment, we recommend you to try to use Gnu Make for the build process. First, enter the directory `gnumakefiles`:

```
$ cd SALMON/gnumakefiles
```

In the directory, Makefile files are prepared for several architectures:

- `gnu-mpi`
- `intel-mpi`
- `gnu-without-mpi`
- `intel-without-mpi`

Makefile files with `*-without-mpi` indicate that they are for single processor environment. Choose Makefile appropriate for your environment, and execute the make command:

```
$ make -f Makefile.PLATFORM
```

If the make proceeds successful, a binary file is created in the directory `SALMON/bin/`.

## EXERCISES

### 3.1 Getting started

Welcome to SALMON Exercises!

In these exercises, we explain the use of SALMON from the very beginning, taking a few samples that cover applications of SALMON in several directions. We assume that you are in the computational environment of UNIX/Linux OS. First you need to download and install SALMON in your computational environment. If you have not yet done it, do it following the instruction, [download](#) and *Install and Run*.

As described in *Install and Run*, you are required to prepare at least an input file and pseudopotential files to run SALMON. In the following, we present input files for several sample calculations and provide a brief explanation of the input keywords that appear in the input files. You may modify the input files to execute for your own calculations. Pseudopotential files of elements that appear in the samples are also attached. We also present explanations of main output files.

We present 10 exercises.

First 3 exercises (Exercise-1 ~ 3) are for an isolated molecule, acetylene C<sub>2</sub>H<sub>2</sub>. If you are interested in learning electron dynamics calculations in isolated systems, please look into these exercises. In SALMON, we usually calculate the ground state solution first using a static density functional theory (DFT). This is illustrated in *Exercise-1*. After finishing the ground state calculation, two exercises of electron dynamics calculations based on time-dependent density functional theory (TDDFT) are prepared. *Exercise-2* illustrates the calculation of linear optical responses in real time, obtaining polarizability and photoabsorption of the molecule. *Exercise-3* illustrates the calculation of electron dynamics in the molecule under a pulsed electric field.

Next 3 exercises (Exercise-4 ~ 6) are for a crystalline solid, silicon. If you are interested in learning electron dynamics calculations in extended periodic systems, please look into these exercises. *Exercise-4* illustrates the ground state calculation of the crystalline silicon based on DFT. *Exercise-5* illustrates the calculation of linear response properties of the crystalline silicon based on TDDFT to obtain the dielectric function. *Exercise-6* illustrates the calculation of electron dynamics in the crystalline silicon induced by a pulsed electric field.

Exercise-7 is for a simultaneous calculation of the propagation of a pulsed light and electronic motion in a bulk silicon, coupling Maxwell equations for the electromagnetic fields of the pulsed light and the electron dynamics in the unit cells based on TDDFT. This calculation is quite time-consuming and is recommended to execute using massively parallel supercomputers. *Exercise-7* illustrates the calculation of a linearly polarized pulsed light irradiating normally on a surface of a bulk silicon.

Next 2 exercises (Exercise-8 ~ 9) are for geometry optimization based on DFT and Ehrenfest molecular dynamics based on TDDFT for an isolated molecule, acetylene C<sub>2</sub>H<sub>2</sub>. *Exercise-8* illustrates the geometry optimization in the ground state. *Exercise-9* illustrates the Ehrenfest molecular dynamics induced by a pulsed electric field.

*Exercise-10* is for a macroscopic light propagation through a metallic nanosphere solving Maxwell equations. The optical response of the nanosphere is described by a dielectric function. Finite-Difference Time-Domain (FDTD) method is used to calculate the three-dimensional light propagation.

Input files of exercises are included in SALMON, in the directory `SALMON/samples/exercise_##_<description>/`.

## 3.2 C2H2 (isolated molecules)

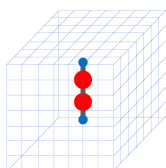
### 3.2.1 Exercise-1: Ground state of C2H2 molecule

In this exercise, we learn the calculation of the ground state of acetylene (C2H2) molecule, solving the static Kohn-Sham equation. This exercise will be useful to learn how to set up calculations in SALMON for any isolated systems such as molecules and nanoparticles.

Acetylene molecule is a linear chain molecule composed of two Carbon atoms and two Hydrogen atoms.



In SALMON, we use a three-dimensional (3D) uniform grid system to express physical quantities such as electron orbitals.

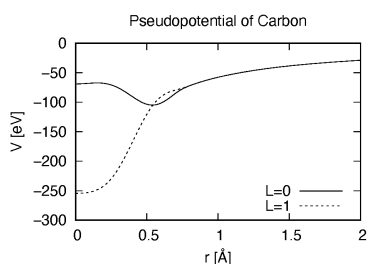


#### Input files

To run the code, following files in the directory `SALMON/samples/exercise_01_C2H2_gs/` are used:

file name	description
<i>C2H2_gs.inp</i>	input file that contains input keywords and their values
<i>C_rps.dat</i>	pseudopotential file for carbon atom
<i>H_rps.dat</i>	pseudopotential file for hydrogen atom

Pseudopotential files are needed for two elements, Carbon (C) and Hydrogen (H). The pseudopotential depends on the angular momentum, and looks as follows (for Carbon).



In the input file *C2H2\_gs.inp*, input keywords are specified. Most of them are mandatory to execute the ground state calculation. This will help you to prepare an input file for other systems that you want to calculate. A complete list of the input keywords that can be used in the input file can be found in [List of input keywords](#).

```
#####
! #####
! Exercise 01: Ground state of C2H2 molecule
! #####
```

(continues on next page)

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```

!-----!
! * The detail of this exercise is explained in our manual(see chapter: 'Exercises').
!
! The manual can be obtained from: https://salmon-tddft.jp/documents.html
!
! * Input format consists of group of keywords like:
!
!     &group
!
!     input keyword = xxx
!
!     /
!
! (see chapter: 'List of input keywords' in the manual)
!
!-----!
! * Conversion from unit_system = 'a.u.' to 'A_eV_fs':
!
! Length: 1 [a.u.] = 0.52917721067      [Angstrom]
!
! Energy: 1 [a.u.] = 27.21138505        [eV]
!
! Time   : 1 [a.u.] = 0.02418884326505 [fs]
!
! #####
!###!

&calculation
!type of theory
theory = 'dft'
/

```

*theory* specifies which theoretical method is used in the calculation.

```

&control
!common name of output files
sysname = 'C2H2'
/

```

*sysname* is a prefix for filenames of output files.

```

&units
!units used in input and output files
unit_system = 'A_eV_fs'
/

```

*unit\_system* specifies which unit system is used in the input and output files.

```
&system
!periodic boundary condition
yn_periodic = 'n'

!number of elements, atoms, electrons and states(orbitals)
nelem = 2
natom = 4
nelec = 10
nstate = 6
/
```

*yn\_periodic* specifies whether or not periodic boundary condition is applied.

*nelem* is the number of elements in the system.

*natom* is the number of atoms in the system.

*nelec* is the number of electrons in the system.

*nstate* is the number of orbitals that are used in the calculation.

```
&pseudo
!name of input pseudo potential file
file_pseudo(1) = './C_rps.dat'
file_pseudo(2) = './H_rps.dat'

!atomic number of element
izatom(1) = 6
izatom(2) = 1

!angular momentum of pseudopotential that will be treated as local
lloc_ps(1) = 1
lloc_ps(2) = 0
!--- Caution -----!
! Indices must correspond to those in &atomic_coor. !
!-----!
/
```

*file\_pseudo(n)* specifies the filename of the pseudopotential file of the n-th element.

*izatom(n)* is the atomic number of the n-th element.

*lloc\_ps(n)* specifies which angular momentum component is chosen as the local potential for the n-th element.

```
&functional
!functional('PZ' is Perdew-Zunger LDA: Phys. Rev. B 23, 5048 (1981).)
xc = 'PZ'
/
```

*xc* specifies the exchange-correlation potential to be used in the calculation.



```

&rgrid
!spatial grid spacing(x,y,z)
dl(1:3) = 0.25d0, 0.25d0, 0.25d0

!number of spatial grids(x,y,z)
num_rgrid(1:3) = 64, 64, 64
/

```

*dl(i)* specifies the spatial grid spacing in i-th direction.

*num\_rgrid(i)* specifies the number of grid points in i-th direction.

```

&scf
!maximum number of scf iteration and threshold of convergence
nscf      = 300
threshold = 1.0d-9
/

```

*nscf* specifies the maximum number of SCF iterations.

*threshold* specifies the threshold to judge the convergence.

```

&analysis
!output of all orbitals, density,
!density of states, projected density of states,
!and electron localization function
yn_out_psi  = 'y'
yn_out_dns  = 'y'
yn_out_dos  = 'y'
yn_out_pdos = 'y'
yn_out_elf  = 'y'
/

```

*yn\_out\_psi*, *yn\_out\_dns*, *yn\_out\_dos*, *yn\_out\_pdos*, *yn\_out\_elf* specify output files that are generated after the calculation.

```

&atomic_coor
!cartesian atomic coordinates
'C'    0.000000    0.000000    0.599672    1
'H'    0.000000    0.000000    1.662257    2
'C'    0.000000    0.000000   -0.599672    1
'H'    0.000000    0.000000   -1.662257    2
!--- Format -----!
! 'symbol' x y z index(correspond to that of pseudo potential) !
!-----!
/

```

*&atomic\_coor* specifies spatial coordinates of atoms.

## Execusion

In a multiprocess environment, calculation will be executed as

```
$ mpiexec -n NPROC salmon < C2H2_gs.inp > C2H2_gs.out
```

where NPROC is the number of MPI processes. A standard output will be stored in the file `C2H2_gs.out`.

## Output files

After the calculation, following output files and a directory are created in the directory that you run the code in addition to the standard output file,

name	description
<i>C2H2_info.data</i>	information on ground state solution
<i>C2H2_eigen.data</i>	orbital energies
<i>C2H2_k.data</i>	k-point distribution (for isolated systems, only gamma point is described)
<i>data_for_restart</i>	directory where files used in the real-time calculation are contained
<i>psi_ob1.cube, psi_ob2.cube, ...</i>	electron orbitals
<i>dns.cube</i>	a cube file for electron density
<i>dos.data</i>	density of states
<i>pdos1.data, pdos2.data, ...</i>	projected density of states
<i>elf.cube</i>	electron localization function (ELF)
<i>PS_C_KY_n.dat</i>	information on pseudopotential file for carbon atom
<i>PS_H_KY_n.dat</i>	information on pseudopotential file for hydrogen atom

You may download the above files (zipped file, except for the directory *data\_for\_restart*) from:

[https://salmon-tddft.jp/webmanual/v\\_2\\_0\\_1/exercise\\_zip\\_files/01\\_C2H2\\_gs.zip](https://salmon-tddft.jp/webmanual/v_2_0_1/exercise_zip_files/01_C2H2_gs.zip)

We first explain the standard output file. In the beginning of the file, input variables used in the calculation are shown.

```
#####
# SALMON: Scalable Ab-initio Light-Matter simulator for Optics and Nanoscience
#
#                               Version 2.0.1
#
#####
Libxc: [disabled]
theory= dft
use of real value orbitals = T
=====
MPI distribution:
nproc_k      :          1
nproc_ob     :          1
nproc_rgrid  :          1          1          2
OpenMP parallelization:
number of threads :        256
.....
```

After that, the SCF loop starts. At each iteration step, the total energy as well as orbital energies and some other quantities are displayed.

```

-----
iter=      1      Total Energy=    -197.59254070      Gap=    -20.17834599      Vh iter=
↪234
   1      -29.9707      2      -28.3380      3      -13.0123      4      5.8457
   5      -9.9213      6      -14.3326
iter and int_x|rho_i(x)-rho_i-1(x)|dx/nelec      =      1 0.31853198E+00
Ne=    10.00000000000000
-----
iter=      2      Total Energy=    -280.97950515      Gap=     -9.59770609      Vh iter=
↪247
   1      -17.4334      2      -24.4941      3      -20.1872      4      0.8020
   5      -3.4058      6      -8.7957
iter and int_x|rho_i(x)-rho_i-1(x)|dx/nelec      =      2 0.54493263E+00
Ne=    10.00000000000000
-----
iter=      3      Total Energy=    -295.67034640      Gap=     -6.90359156      Vh iter=
↪229
   1      -16.0251      2      -19.7759      3      -17.6765      4      -0.9015
   5      -2.9323      6      -7.8050
iter and int_x|rho_i(x)-rho_i-1(x)|dx/nelec      =      3 0.13010987E+00
Ne=    10.00000000000000

```

When the convergence criterion is satisfied, the SCF calculation ends.

```

-----
iter=    162      Total Energy=    -339.69525272      Gap=      6.78870999      Vh iter=
↪  1
   1      -18.4106      2      -13.9966      3      -12.4163      4      -7.3386
   5      -7.3386      6      -0.5498
iter and int_x|rho_i(x)-rho_i-1(x)|dx/nelec      =    162 0.50237787E-08
Ne=     9.99999999999999
-----
iter=    163      Total Energy=    -339.69525269      Gap=      6.78870999      Vh iter=
↪  1
   1      -18.4106      2      -13.9966      3      -12.4163      4      -7.3386
   5      -7.3386      6      -0.5498
iter and int_x|rho_i(x)-rho_i-1(x)|dx/nelec      =    163 0.69880308E-09
Ne=     9.99999999999999
#GS converged at 164 : 0.69880308E-09

```

Next, the force acting on ions and some other information related to orbital energies are shown.

```

===== force =====
  1 -0.33652081E-05  0.16854696E-04 -0.59496450E+00
  2 -0.59222259E-06  0.24915590E-05  0.57651725E+00
  3 -0.37839836E-05  0.20304090E-04  0.59493028E+00
  4 -0.86779607E-06  0.39560274E-05 -0.57651738E+00
orbital energy information-----
Lowest occupied orbital -0.676576619015730
Highest occupied orbital (HOMO) -0.269686750876529
Lowest unoccupied orbital (LUMO) -2.020624936948345E-002
Highest unoccupied orbital -2.020624936948345E-002
HOMO-LUMO gap 0.249480501507045
Physically upper bound of eps(omega) 0.656370369646246
-----
Lowest occupied orbital[eV] -18.4105868958642
Highest occupied orbital (HOMO)[eV] -7.33855002098465

```

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```

Lowest unoccupied orbital (LUMO) [eV] -0.549840032009334
Highest unoccupied orbital[eV] -0.549840032009334
HOMO-LUMO gap[eV] 6.78870998897532
Physically upper bound of eps(omega) [eV] 17.8607468638548
-----
writing restart data...
writing completed.

```

In the directory `data_for_restart`, files that will be used in the next-step time evolution calculations are stored.

Other output files include following information.

### **C2H2\_info.data**

Calculated orbital and total energies as well as parameters specified in the input file are shown.

### **C2H2\_eigen.data**

Orbital energies.

```

#esp: single-particle energies (eigen energies)
#occ: occupation numbers, io: orbital index
# 1:io, 2:esp[eV], 3:occ

```

### **C2H2\_k.data**

k-point distribution(for isolated systems, only gamma point is described).

```

# ik: k-point index
# kx,ky,kz: Reduced coordinate of k-points
# wk: Weight of k-point
# 1:ik[none] 2:kx[none] 3:ky[none] 4:kz[none] 5:wk[none]
# coefficients (2*pi/a [a.u.]) in kx, ky, kz

```

### **psi\_ob1.cube, psi\_ob2.cube, ...**

Cube files for electron orbitals. The number in the filename indicates the index of the orbital. Atomic unit is adopted in all cube files.

### **dns.cube**

A cube file for electron density.

### **dos.data**

A file for density of states. The units used in this file are affected by the input parameter, `unit_system` in `&unit`.

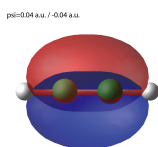
### **elf.cube**

A cube file for electron localization function (ELF).

We show several image that are created from the output files.

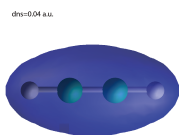
- **Highest occupied molecular orbital (HOMO)**

The output files `psi_ob1.cube`, `psi_ob2.cube`, ... are used to create the image.



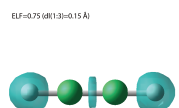
- **Electron density**

The output files *dns.cube*, ... are used to create the image.



- **Electron localization function**

The output files *elf.cube*, ... are used to create the image.



### 3.2.2 Exercise-2: Polarizability and photoabsorption of C2H2 molecule

In this exercise, we learn the linear response calculation in the acetylene (C2H2) molecule, solving the time-dependent Kohn-Sham equation. The linear response calculation provides the polarizability and the oscillator strength distribution of the molecule. This exercise should be carried out after finishing the ground state calculation that was explained in *Exercise-1*.

Polarizability  $\alpha_{\mu\nu}(t)$  is the basic quantity that characterizes optical responses of molecules and nano-particles, where  $\mu, \nu$  indicate Cartesian components,  $\mu, \nu = x, y, z$ . The polarizability  $\alpha_{\mu\nu}(t)$  relates the  $\mu$  component of the electric dipole moment at time  $t$ ,  $p_\mu(t)$ , with the  $\nu$  component of the electric field at time  $t'$ ,

$$p_\mu(t) = \sum_{\nu=x,y,z} \alpha_{\mu\nu}(t-t') E_\nu(t').$$

We introduce a frequency-dependent polarizability by the time-frequency Fourier transformation of the polarizability,

$$\tilde{\alpha}_{\mu\nu}(\omega) = \int dt e^{i\omega t} \alpha_{\mu\nu}(t).$$

The imaginary part of the frequency-dependent polarizability is related to the photoabsorption cross section  $\sigma(\omega)$  by

$$\sigma(\omega) = \frac{4\pi\omega}{c} \frac{1}{3} \sum_{\mu=x,y,z} \text{Im} \tilde{\alpha}_{\mu\mu}(\omega).$$

The photoabsorption cross section is also related to the oscillator strength distribution by

$$\sigma(\omega) = \frac{2\pi^2 e^2}{mc} \frac{df(\omega)}{d\omega}.$$

In SALMON, the polarizability is calculated in time domain. First the ground state orbital  $\phi_i(\mathbf{r})$  that satisfies the Kohn-Sham equation,

$$H_{\text{KS}} \phi_i(\mathbf{r}) = \epsilon_i \phi_i(\mathbf{r}),$$

is prepared. Then an impulsive force given by the potential

$$V_{\text{ext}}(\mathbf{r}, t) = I \delta(t) z,$$

is applied to all electrons in the C2H2 molecule along the molecular axis which we take  $z$  axis.  $I$  is the magnitude of the impulse, and  $\delta(t)$  is the Dirac's delta function. The orbital is distorted by the impulsive force at  $t = 0$ . Immediately after the impulse is applied, the orbital becomes

$$\psi_i(\mathbf{r}, t = 0_+) = e^{iIz/\hbar} \phi_i(\mathbf{r}).$$

After the impulsive force is applied at  $t = 0$ , a time evolution calculation is carried out without any external fields,

$$i\hbar \frac{\partial}{\partial t} \psi_i(\mathbf{r}, t) = H_{\text{KS}}(t) \psi_i(\mathbf{r}, t).$$

During the time evolution, the electric dipole moment given by

$$p_z(t) = \int d\mathbf{r} (-ez) \sum_i |\psi_i(\mathbf{r}, t)|^2,$$

is monitored. After the time evolution calculation, a time-frequency Fourier transformation is carried out for the electric dipole moment to obtain the frequency-dependent polarizability by

$$\tilde{\alpha}_{zz}(\omega) = -\frac{e}{i} \int dt e^{i\omega t} p_z(t).$$

## Input files

To run the code, following files are necessary:

file name	description
<i>C2H2_rt_response.inp</i>	input file that contain input keywords and their values.
<i>C_rps.dat</i>	pseudopotential file for carbon
<i>H_rps.dat</i>	pseudopotential file for hydrogen
<i>restart</i>	directory created in the ground state calculation (rename the directory from <i>data_for_restart</i> to <i>restart</i> )

First three files are prepared in the directory `SALMON/samples/exercise_02_C2H2_lr/`. The file *C2H2\_rt\_response.inp* that contains input keywords and their values. The pseudopotential files should be the same as those used in the ground state calculation. In the directory *restart*, those files created in the ground state calculation and stored in the directory *data\_for\_restart* are included. Therefore, copy the directory as `cp -R data_for_restart restart` if you calculate at the same directory as you did the ground state calculation.

In the input file *C2H2\_rt\_response.inp*, input keywords are specified. Most of them are mandatory to execute the linear response calculation. This will help you to prepare the input file for other systems that you want to calculate. A complete list of the input keywords that can be used in the input file can be found in [List of input keywords](#).

```
#####
#####!
! Exercise 02: Polarizability and photoabsorption of C2H2 molecule
!
!-----!
! * The detail of this exercise is explained in our manual(see chapter: 'Exercises').
!
! The manual can be obtained from: https://salmon-tddft.jp/documents.html
!
! * Input format consists of group of keywords like:
!
!      &group
!
!      input keyword = xxx
!
!      /
!
! (see chapter: 'List of input keywords' in the manual)
!
!-----!
#####
```

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```

! * Conversion from unit_system = 'a.u.' to 'A_eV_fs':
↳ !
!   Length: 1 [a.u.] = 0.52917721067      [Angstrom]
↳ !
!   Energy: 1 [a.u.] = 27.21138505      [eV]
↳ !
!   Time   : 1 [a.u.] = 0.02418884326505 [fs]
↳ !
!-----!
↳ ---!
! * Copy the ground state data directory('data_for_restart') (or make symbolic link)
↳ !
!   calculated in 'samples/exercise_01_C2H2_gs/' and rename the directory to 'restart/'
↳ ' !
!   in the current directory.
↳ !
!#####
↳###!

&calculation
!type of theory
theory = 'tddft_response'
/

```

*theory* specifies which theoretical method is used in the calculation.

```

&control
!common name of output files
sysname = 'C2H2'
/

```

*sysname* is a prefix for filenames of output files.

```

&units
!units used in input and output files
unit_system = 'A_eV_fs'
/

```

*unit\_system* specifies which unit system is used in the input and output files.

```

&system
!periodic boundary condition
yn_periodic = 'n'

!number of elements, atoms, electrons and states(orbitals)
nelem = 2
natom = 4

```

(continues on next page)

(continued from previous page)

```
nelec = 10
nstate = 6
/
```

*yn\_periodic* specifies whether or not periodic boundary condition is applied.

*nelem* is the number of elements in the system.

*natom* is the number of atoms in the system.

*nelec* is the number of electrons in the system.

*nstate* is the number of orbitals that are used in the calculation.

```
&pseudo
!name of input pseudo potential file
file_pseudo(1) = './C_rps.dat'
file_pseudo(2) = './H_rps.dat'

!atomic number of element
izatom(1) = 6
izatom(2) = 1

!angular momentum of pseudopotential that will be treated as local
lloc_ps(1) = 1
lloc_ps(2) = 0
!--- Caution -----!
! Indices must correspond to those in &atomic_coor. !
!-----!
/
```

*file\_pseudo(n)* specifies the filename of the pseudopotential file of the n-th element.

*izatom(n)* is the atomic number of the n-th element.

*lloc\_ps(n)* specifies which angular momentum component is chosen as the local potential for the n-th element.

```
&functional
!functional('PZ' is Perdew-Zunger LDA: Phys. Rev. B 23, 5048 (1981).)
xc = 'PZ'
/
```

*xc* specifies the exchange-correlation potential to be used in the calculation.

```
&rgrid
!spatial grid spacing(x,y,z)
dl(1:3) = 0.25d0, 0.25d0, 0.25d0

!number of spatial grids(x,y,z)
num_rgrid(1:3) = 64, 64, 64
/
```



*dl(i)* specifies the spatial grid spacing in i-th direction.

*num\_rgrid(i)* specifies the number of grid points in i-th direction.

```
&tgrid
!time step size and number of time grids(steps)
dt = 1.25d-3
nt = 5000
/
```

*dt* specifies the time step.

*nt* is the number of time steps for the time propagation.

```
&emfield
!envelope shape of the incident pulse('impulse': impulsive field)
ae_shape1 = 'impulse'

!polarization unit vector(real part) for the incident pulse(x,y,z)
epdir_rel(1:3) = 0.0d0, 0.0d0, 1.0d0
!---- Caution -----!
! Defenition of the incident pulse is wrritten in:                !
! https://www.sciencedirect.com/science/article/pii/S0010465518303412 !
!-----!
/
```

*ae\_shape1* specifies the envelope of the field. For a linear response calculation, *ae\_shape1*='impulse' is used. It indicates that a weak impulsive perturbation is applied at  $t = 0$ .

*epdir\_rel(i)* specifies the i-th component of the real part of the polarization unit vector.

```
&analysis
!energy grid size and number of energy grids for output files
de      = 1.0d-2
nenergy = 3000
/
```

*de* specifies the energy grid size for frequency-domain analysis.

*nenergy* specifies the number of energy grid points for frequency-domain analysis.

```
&atomic_coor
!cartesian atomic coodinates
'C'    0.000000    0.000000    0.599672    1
'H'    0.000000    0.000000    1.662257    2
'C'    0.000000    0.000000   -0.599672    1
'H'    0.000000    0.000000   -1.662257    2
!---- Format -----!
! 'symbol' x y z index(correspond to that of pseudo potential) !
```

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```
!-----!
/
```

*&atomic\_coor* specifies spatial coordinates of atoms.

## Excursion

Before execution, remember to copy the directory *restart* that is created in the ground state calculation as *data\_for\_restart* in the present directory. In a multiprocess environment, calculation will be executed as

```
$ mpiexec -n NPROC salmon < C2H2_rt_response.inp > C2H2_rt_response.out
```

where NPROC is the number of MPI processes. A standard output will be stored in the file *C2H2\_rt\_response.out*.

## Output files

After the calculation, following output files are created in the directory that you run the code,

file name	description
<i>C2H2_response.data</i>	polarizability and oscillator strength distribution as functions of energy
<i>C2H2_rt.data</i>	components of change of dipole moment (electrons/plus definition) and total dipole moment (electrons/minus + ions/plus) as functions of time
<i>C2H2_rt_energy.data</i>	components of total energy and difference of total energy as functions of time
<i>PS_C_KY_n.dat</i>	information on pseudopotential file for carbon atom
<i>PS_H_KY_n.dat</i>	information on pseudopotential file for hydrogen atom

You may download the above files (zipped file) from:

[https://salmon-tddft.jp/webmanual/v\\_2\\_0\\_1/exercise\\_zip\\_files/02\\_C2H2\\_lr.zip](https://salmon-tddft.jp/webmanual/v_2_0_1/exercise_zip_files/02_C2H2_lr.zip)

We first explain the standard output file. In the beginning of the file, input variables used in the calculation are shown.

```
#####
# SALMON: Scalable Ab-initio Light-Matter simulator for Optics and Nanoscience
#
#                                     Version 2.0.1
#####
Libxc: [disabled]
theory= tddft_response

Total time step      =      5000
```

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```

Time step[fs]          = 1.2500000000000000E-003
Energy range           = 3000
Energy resolution[eV] = 1.0000000000000000E-002
Field strength[a.u.] = 1.0000000000000000E-002
  use of real value orbitals = F
=====
.....

```

After that, the time evolution loop starts. At every 10 iteration steps, the time, dipole moments in three Cartesian directions, the total number of electrons, the total energy, and the number of iterations solving the Poisson equation are displayed.

time-step	time[fs]	Dipole moment (xyz) [A]			electrons
↪ Total energy[eV]	iterVh				
#-----					
10	0.01250000	-0.56521137E-07	-0.28812833E-07	-0.25558983E-01	10.00000000
↪ -339.68150366	34				
20	0.02500000	-0.19835467E-06	-0.10147641E-06	-0.45169126E-01	9.99999999
↪ -339.68147442	49				
30	0.03750000	-0.37937911E-06	-0.19537418E-06	-0.57843871E-01	9.99999999
↪ -339.68146891	45				
40	0.05000000	-0.56465010E-06	-0.29324906E-06	-0.64072126E-01	9.99999999
↪ -339.68146804	38				
50	0.06250000	-0.73343753E-06	-0.38431758E-06	-0.65208422E-01	9.99999999
↪ -339.68146679	25				
60	0.07500000	-0.87559727E-06	-0.46276791E-06	-0.62464066E-01	9.99999999
↪ -339.68146321	35				
70	0.08750000	-0.98769124E-06	-0.52594670E-06	-0.56740338E-01	9.99999998
↪ -339.68145535	20				
80	0.10000000	-0.10701350E-05	-0.57309375E-06	-0.48483747E-01	9.99999998
↪ -339.68144840	40				
90	0.11250000	-0.11253992E-05	-0.60455485E-06	-0.38296037E-01	9.99999998
↪ -339.68144186	21				

Explanations of other output files are given below:

### C2H2\_rt.data

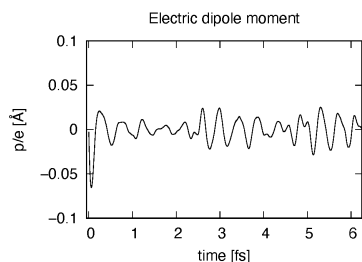
Results of time evolution calculation for vector potential, electric field, and dipole moment. In the first several lines, explanations of included data are given.

```

# Real time calculation:
# Ac_ext: External vector potential field
# E_ext: External electric field
# Ac_tot: Total vector potential field
# E_tot: Total electric field
# ddm_e: Change of dipole moment (electrons/plus definition)
# dm: Total dipole moment (electrons/minus + ions/plus)
# 1:Time[fs] 2:Ac_ext_x[fs*V/Angstrom] 3:Ac_ext_y[fs*V/Angstrom] 4:Ac_ext_z[fs*V/
↪Angstrom] 5:E_ext_x[V/Angstrom] 6:E_ext_y[V/Angstrom] 7:E_ext_z[V/Angstrom] 8:Ac_
↪tot_x[fs*V/Angstrom] 9:Ac_tot_y[fs*V/Angstrom] 10:Ac_tot_z[fs*V/Angstrom] 11:E_tot_
↪x[V/Angstrom] 12:E_tot_y[V/Angstrom] 13:E_tot_z[V/Angstrom] 14:ddm_e_x[Angstrom]
↪15:ddm_e_y[Angstrom] 16:ddm_e_z[Angstrom] 17:dm_x[Angstrom] 18:dm_y[Angstrom] 19:dm_
↪z[Angstrom]

```

Using first column (time in femtosecond) and 19th column (dipole moment in z direction), the following graph can be drawn.



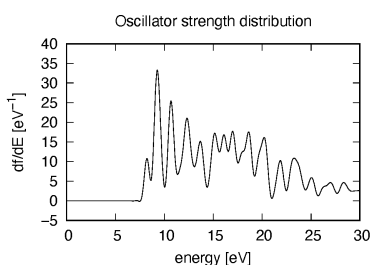
The dipole moment shows oscillations in femtosecond time scale that reflect electronic excitations.

### C2H2\_response.data

Time-frequency Fourier transformation of the dipole moment gives the polarizability and the strength function.

```
# Fourier-transform spectra:
# alpha: Polarizability
# df/dE: Strength function
# 1:Energy[eV] 2:Re(alpha_x)[Augstrom^2/V] 3:Re(alpha_y)[Augstrom^2/V] 4:Re(alpha_
↪ z)[Augstrom^2/V] 5:Im(alpha_x)[Augstrom^2/V] 6:Im(alpha_y)[Augstrom^2/V] 7:Im(alpha_
↪ z)[Augstrom^2/V] 8:df_x/dE[none] 9:df_y/dE[none] 10:df_z/dE[none]
```

Using first column (energy in electron-volt) and 10th column (oscillator strength distribution in  $z$  direction), the following graph can be drawn.



There appears many peaks above the HOMO-LUMO gap energy. The strong excitation appears at around 9.3 eV.

### C2H2\_rt\_energy.data

Energies are stored as functions of time.

```
# Real time calculation:
# Eall: Total energy
# Eall0: Initial energy
# 1:Time[fs] 2:Eall[eV] 3:Eall-Eall0[eV]
```

$E_{all}$  and  $E_{all}-E_{all0}$  are total energy and electronic excitation energy, respectively.

## 3.2.3 Exercise-3: Electron dynamics in C2H2 molecule under a pulsed electric field

In this exercise, we learn the calculation of the electron dynamics in the acetylene (C2H2) molecule under a pulsed electric field, solving the time-dependent Kohn-Sham equation. As outputs of the calculation, such quantities as the total energy and the electric dipole moment of the system as functions of time are calculated. This tutorial should be carried out after finishing the ground state calculation that was explained in [Exercise-1](#).

In the calculation, a pulsed electric field specified by the following vector potential will be used,

$$A(t) = -\frac{E_0}{\omega} \hat{z} \cos^2 \frac{\pi}{T} \left( t - \frac{T}{2} \right) \sin \omega \left( t - \frac{T}{2} \right), \quad (0 < t < T).$$

The electric field is given by  $E(t) = -(1/c)(dA(t)/dt)$ . The parameters that characterize the pulsed field such as the amplitude  $E_0$ , frequency  $\omega$ , pulse duration  $T$ , polarization direction  $\hat{z}$ , are specified in the input file. In the time dependent Kohn-Sham equation, the external field is included as the scalar potential,  $V_{\text{ext}}(\mathbf{r}, t) = eE(t)z$ .

## Input files

To run the code, following files are necessary:

file name	description
<i>C2H2_rt_pulse.inp</i>	input file that contain input keywords and their values.
<i>C_rps.dat</i>	pseudopotential file for carbon
<i>H_rps.dat</i>	pseudopotential file for hydrogen
<i>restart</i>	directory created in the ground state calculation (rename the directory from <i>data_for_restart</i> to <i>restart</i> )

First three files are prepared in the directory `SALMON/samples/exercise_03_C2H2_rt/`. The file *C2H2\_rt\_pulse.inp* that contains input keywords and their values. The pseudopotential files should be the same as those used in the ground state calculation. In the directory *restart*, those files created in the ground state calculation and stored in the directory *data\_for\_restart* are included. Therefore, copy the directory as `cp -R data_for_restart restart` if you calculate at the same directory as you did the ground state calculation.

In the input file *C2H2\_rt\_pulse.inp*, input keywords are specified. Most of them are mandatory to execute the calculation of electron dynamics induced by a pulsed electric field. This will help you to prepare the input file for other systems and other pulsed electric fields that you want to calculate. A complete list of the input keywords that can be used in the input file can be found in [List of input keywords](#).

```
#####
<###!
! Exercise 03:  Electron dynamics in C2H2 molecule under a pulsed electric field
<  !
!-----!
<---!
! * The detail of this exercise is explained in our manual(see chapter: 'Exercises').
<  !
!   The manual can be obtained from: https://salmon-tddft.jp/documents.html
<  !
! * Input format consists of group of keywords like:
<  !
!      &group
<  !
!      input keyword = xxx
<  !
!      /
<  !
!   (see chapter: 'List of input keywords' in the manual)
<  !
!-----!
<---!
! * Conversion from unit_system = 'a.u.' to 'A_eV_fs':
<  !
!   Length: 1 [a.u.] = 0.52917721067      [Angstrom]
<  !
```

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```

!   Energy: 1 [a.u.] = 27.21138505      [eV]
↪   !
!   Time   : 1 [a.u.] = 0.02418884326505 [fs]
↪   !
!-----!
↪---!
! * Copy the ground state data directory('data_for_restart') (or make symbolic link)
↪   !
!   calculated in 'samples/exercise_01_C2H2_gs/' and rename the directory to 'restart/'
↪   !
!   in the current directory.
↪   !
!#####
↪###!

&calculation
!type of theory
theory = 'tddft_pulse'
/

```

*theory* specifies which theoretical method is used in the calculation.

```

&control
!common name of output files
sysname = 'C2H2'
/

```

*sysname* is a prefix for filenames of output files.

```

&units
!units used in input and output files
unit_system = 'A_eV_fs'
/

```

*unit\_system* specifies which unit system is used in the input and output files.

```

&system
!periodic boundary condition
yn_periodic = 'n'

!number of elements, atoms, electrons and states(orbitals)
nelem = 2
natom = 4
nelec = 10
nstate = 6
/

```

*yn\_periodic* specifies whether or not periodic boundary condition is applied.

*nelem* is the number of elements in the system.

*natom* is the number of atoms in the system.

*nelec* is the number of electrons in the system.

*nstate* is the number of orbitals that are used in the calculation.

```
&pseudo
!name of input pseudo potential file
file_pseudo(1) = './C_rps.dat'
file_pseudo(2) = './H_rps.dat'

!atomic number of element
izatom(1) = 6
izatom(2) = 1

!angular momentum of pseudopotential that will be treated as local
lloc_ps(1) = 1
lloc_ps(2) = 0
!--- Caution -----!
! Indices must correspond to those in &atomic_coor. !
!-----!
/
```

*file\_pseudo(n)* specifies the filename of the pseudopotential file of the n-th element.

*izatom(n)* is the atomic number of the n-th element.

*lloc\_ps(n)* specifies which angular momentum component is chosen as the local potential for the n-th element.

```
&functional
!functional('PZ' is Perdew-Zunger LDA: Phys. Rev. B 23, 5048 (1981).)
xc = 'PZ'
/
```

*xc* specifies the exchange-correlation potential to be used in the calculation.

```
&rgrid
!spatial grid spacing(x,y,z)
dl(1:3) = 0.25d0, 0.25d0, 0.25d0

!number of spatial grids(x,y,z)
num_rgrid(1:3) = 64, 64, 64
/
```

*dl(i)* specifies the spatial grid spacing in i-th direction.

*num\_rgrid(i)* specifies the number of grid points in i-th direction.

```
&tgrid
!time step size and number of time grids(steps)
dt = 1.25d-3
nt = 5000
/
```

*dt* specifies the time step.

*nt* is the number of time steps for the time propagation.

```
&emfield
!envelope shape of the incident pulse('Ecos2': cos^2 type envelope for scalar_
↪potential)
ae_shape1 = 'Acos2'

!peak intensity(W/cm^2) of the incident pulse
I_wcm2_1 = 5.00d13

!duration of the incident pulse
tw1 = 6.00d0

!mean photon energy(average frequency multiplied by the Planck constant) of the_
↪incident pulse
omega1 = 3.10d0

!polarization unit vector(real part) for the incident pulse(x,y,z)
epdir_rel(1:3) = 0.00d0, 0.00d0, 1.00d0
!--- Caution -----!
! Defenition of the incident pulse is wrritten in: !
! https://www.sciencedirect.com/science/article/pii/S0010465518303412 !
!-----!
/
```

*ae\_shape1* specifies the envelope of the field.

*I\_wcm2\_1* specify the intensity of the pulse in unit of W/cm<sup>2</sup>.

*tw1* specifies the duration of the pulse.

*omega1* specifies the mean photon energy of the pulse.

*epdir\_rel(i)* specifies the i-th component of the real part of the polarization unit vector.

```
&analysis
!energy grid size and number of energy grids for output files
de = 1.0d-2
nenergy = 10000
/
```

*de* specifies the energy grid size for frequency-domain analysis.

*nenergy* specifies the number of energy grid points for frequency-domain analysis.



```

&atomic_coor
!cartesian atomic coordinates
'C'    0.000000    0.000000    0.599672    1
'H'    0.000000    0.000000    1.662257    2
'C'    0.000000    0.000000   -0.599672    1
'H'    0.000000    0.000000   -1.662257    2
!--- Format -----!
! 'symbol' x y z index(correspond to that of pseudo potential) !
!-----!
/

```

*&atomic\_coor* specifies spatial coordinates of atoms.

## Excursion

Before execution, remember to copy the directory *restart* that is created in the ground state calculation as *data\_for\_restart* in the present directory. In a multiprocess environment, calculation will be executed as

```
$ mpiexec -n NPROC salmon < C2H2_rt_pulse.inp > C2H2_rt_pulse.out
```

where NPROC is the number of MPI processes. A standard output will be stored in the file *C2H2\_rt\_pulse.out*.

## Output files

After the calculation, following output files are created in the directory that you run the code,

file name	description
<i>C2H2_pulse.data</i>	dipole moment as functions of energy
<i>C2H2_rt.data</i>	components of change of dipole moment (electrons/plus definition) and total dipole moment (electrons/minus + ions/plus) as functions of time
<i>C2H2_rt_energy.data</i>	components of total energy and difference of total energy as functions of time
<i>PS_C_KY_n.dat</i>	information on pseudopotential file for carbon atom
<i>PS_H_KY_n.dat</i>	information on pseudopotential file for hydrogen atom

You may download the above files (zipped file) from:

[https://salmon-tddft.jp/webmanual/v\\_2\\_0\\_1/exercise\\_zip\\_files/03\\_C2H2\\_rt.zip](https://salmon-tddft.jp/webmanual/v_2_0_1/exercise_zip_files/03_C2H2_rt.zip)

We first explain the standard output file. In the beginning of the file, input variables used in the calculation are shown.

```

#####
# SALMON: Scalable Ab-initio Light-Matter simulator for Optics and Nanoscience
#
#                                     Version 2.0.1

```

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```
#####
Libxc: [disabled]
theory= tddft_pulse

Total time step      =      5000
Time step[fs]        =  1.250000000000000E-003
Energy range         =      10000
Energy resolution[eV]=  1.000000000000000E-002
Laser frequency      =  3.10[eV]
Pulse width of laser=  6.00000000[fs]
Laser intensity       =  0.50000000E+14[W/cm^2]
  use of real value orbitals = F
=====
.....
```

After that, the time evolution loop starts. At every 10 iteration steps, the time, dipole moments in three Cartesian directions, the total number of electrons, the total energy, and the number of iterations solving the Poisson equation are displayed.

time-step	time[fs]	Dipole moment (xyz) [A]			electrons
→ Total energy[eV]	iterVh				
#-----					
10	0.01250000	-0.57275542E-07	-0.29197105E-07	-0.74600728E-06	10.00000000
→ -339.69524047	1				
20	0.02500000	-0.20616352E-06	-0.10537273E-06	-0.10256205E-04	10.00000000
→ -339.69524348	1				
30	0.03750000	-0.40063325E-06	-0.20597522E-06	-0.47397133E-04	10.00000000
→ -339.69524090	3				
40	0.05000000	-0.59093535E-06	-0.30630513E-06	-0.13774845E-03	10.00000000
→ -339.69524287	1				
50	0.06250000	-0.75588343E-06	-0.39552925E-06	-0.31097825E-03	10.00000000
→ -339.69523949	5				
60	0.07500000	-0.89221538E-06	-0.47142217E-06	-0.59735355E-03	10.00000000
→ -339.69523784	11				
70	0.08750000	-0.99769538E-06	-0.53192187E-06	-0.10253308E-02	10.00000000
→ -339.69523285	5				
80	0.10000000	-0.10738281E-05	-0.57676878E-06	-0.16195168E-02	9.99999999
→ -339.69522482	19				
90	0.11250000	-0.11250289E-05	-0.60722757E-06	-0.23985719E-02	9.99999999
→ -339.69521092	2				

Explanations of the files are given below:

### C2H2\_rt.data

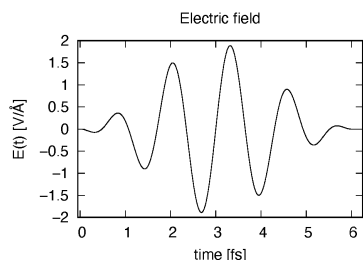
Results of time evolution calculation for vector potential, electric field, and dipole moment. In the first several lines, explanations of data included data are given.

```
# Real time calculation:
# Ac_ext: External vector potential field
# E_ext: External electric field
# Ac_tot: Total vector potential field
# E_tot: Total electric field
# ddm_e: Change of dipole moment (electrons/plus definition)
# dm: Total dipole moment (electrons/minus + ions/plus)
# 1:Time[fs] 2:Ac_ext_x[fs*V/Angstrom] 3:Ac_ext_y[fs*V/Angstrom] 4:Ac_ext_z[fs*V/
→Angstrom] 5:E_ext_x[V/Angstrom] 6:E_ext_y[V/Angstrom] 7:E_ext_z[V/Angstrom] 8:Ac_
→tot_x[fs*V/Angstrom] 9:Ac_tot_y[fs*V/Angstrom] 10:Ac_tot_z[fs*V/Angstrom] 11:E_tot
→x[V/Angstrom] 12:E_tot_y[V/Angstrom] 13:E_tot_z[V/Angstrom] 14:ddm_e_x[Angstrom]
→15:ddm_e_y[Angstrom] 16:ddm_e_z[Angstrom] 17:dm_x[Angstrom] 18:dm_y[Angstrom] 19:dm
38z[Angstrom]
```

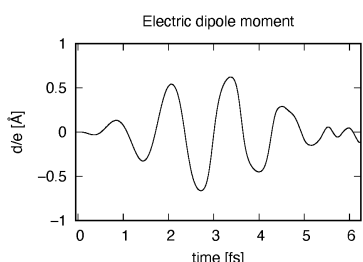
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The applied electric field is drawn using the first column (time in femtosecond) and the 7th column (electric field in  $z$  direction in Volt per Angstrom).



The induced dipole moment is drawn using the first column (time in femtosecond) and 19th column (dipole moment in  $z$  direction). It shows an oscillation similar to the applied electric field. However, the response is not linear since the applied electric field is rather strong.

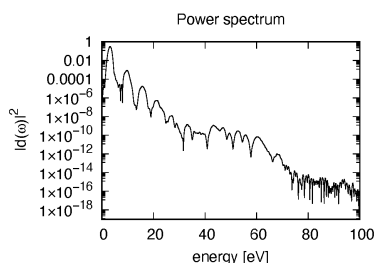


### C2H2\_pulse.data

Time-frequency Fourier transformation of the dipole moment. In the first several lines, explanations of data included data are given.

```
# Fourier-transform spectra:
# energy: Frequency
# dm: Dipole moment
# 1:energy[eV] 2:Re(dm_x)[fs*Angstrom] 3:Re(dm_y)[fs*Angstrom] 4:Re(dm_
↪ z)[fs*Angstrom] 5:Im(dm_x)[fs*Angstrom] 6:Im(dm_y)[fs*Angstrom] 7:Im(dm_
↪ z)[fs*Angstrom] 8:|dm_x|^2[fs^2*Angstrom^2] 9:|dm_y|^2[fs^2*Angstrom^2] 10:|dm_z|^
↪ 2[fs^2*Angstrom^2]
```

The spectrum of the induced dipole moment,  $|d(\omega)|^2$  is shown in logarithmic scale as a function of the energy,  $\hbar\omega$ . High harmonic generations are visible in the spectrum.

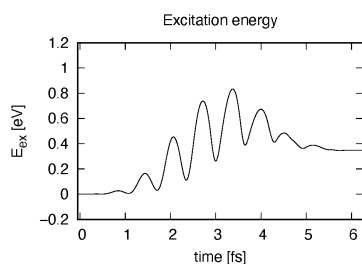


### C2H2\_rt\_energy.data

Energies are stored as functions of time. In the first several lines, explanations of data included data are given.

```
# Real time calculation:
# Eall: Total energy
# Eall0: Initial energy
# 1:Time[fs] 2:Eall[eV] 3:Eall-Eall0[eV]
```

$E_{all}$  and  $E_{all}-E_{all0}$  are total energy and electronic excitation energy, respectively. The figure below shows the electronic excitation energy as a function of time, using the first column (time in femtosecond) and the 3rd column ( $E_{all}-E_{all0}$ ). Although the frequency is below the HOMO-LUMO gap energy, electronic excitations take place because of nonlinear absorption process.



### Additional exercise

If we change parameters of the applied electric field, we find a drastic change in the electronic excitations. In the example below, we increase the intensity from  $I_{wcm2\_1} = 5.00d13$  to  $I_{wcm2\_1} = 1.00d12$  and changes the frequency from  $\omega_{gal} = 3.10d0$  to  $\omega_{gal} = 9.28d0$ . The new frequency corresponds to the resonant excitation energy seen in the linear response analysis shown in [Exercise-2](#).

The change in the input file is shown below.

```
&emfield
!envelope shape of the incident pulse('Ecos2': cos^2 type envelope for scalar_
↪potential)
ae_shape1 = 'Acos2'

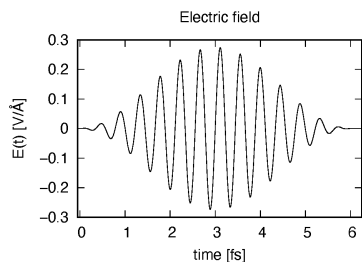
!peak intensity(W/cm^2) of the incident pulse
I_wcm2_1 = 1.00d12

!duration of the incident pulse
tw1 = 6.00d0

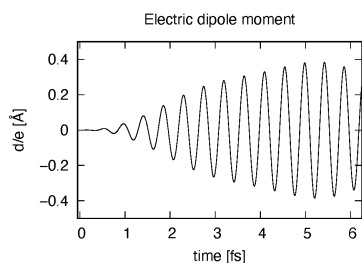
!mean photon energy(average frequency multiplied by the Planck constant) of the_
↪incident pulse
omega1 = 9.28d0

!polarization unit vector(real part) for the incident pulse(x,y,z)
epdir_rel(1:3) = 0.00d0, 0.00d0, 1.00d0
```

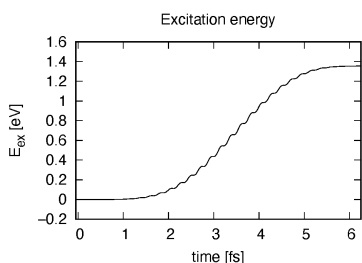
The applied electric field shows a rapid oscillation.



The induced dipole moment also shows a rapid oscillation and does not decrease even though the electric field decreases. This is because the frequency of the applied electric field coincides with the excitation energy of the molecule.



The electronic excitation energy also shows a monotonic increase. Although the strength of the applied electric field is much smaller than the previous case, the amount of the excitation energy is larger, again due to the resonant excitation.



### 3.3 Crystalline silicon (periodic solids)

#### 3.3.1 Exercise-4: Ground state of crystalline silicon

In this exercise, we learn the the ground state calculation of the crystalline silicon of a diamond structure. Calculation is done in a cubic unit cell that contains eight silicon atoms. This exercise will be useful to learn how to set up calculations in SALMON for any periodic systems such as crystalline solid.

#### Input files

To run the code, following files in samples are used:

file name	description
<i>Si_gs.inp</i>	input file that contains input keywords and their values
<i>Si_rps.dat</i>	pseudopotential file for silicon atom

In the input file *Si\_gs.inp*, input keywords are specified. Most of them are mandatory to execute the ground state

calculation. This will help you to prepare an input file for other systems that you want to calculate. A complete list of the input keywords that can be used in the input file can be found in *List of input keywords*.

```
#####
↳###!
! Exercice 04: Ground state of crystalline silicon(periodic solids)
↳ !
!-----!
↳---!
! * The detail of this exercice is expained in our manual(see chapter: 'Exercises').
↳ !
! The manual can be obtained from: https://salmon-tddft.jp/documents.html
↳ !
! * Input format consists of group of keywords like:
↳ !
! &group
↳ !
! input keyword = xxx
↳ !
! /
↳ !
! (see chapter: 'List of input keywords' in the manual)
↳ !
!-----!
↳---!
! * Conversion from unit_system = 'a.u.' to 'A_eV_fs':
↳ !
! Length: 1 [a.u.] = 0.52917721067 [Angstrom]
↳ !
! Energy: 1 [a.u.] = 27.21138505 [eV]
↳ !
! Time : 1 [a.u.] = 0.02418884326505 [fs]
↳ !
!#####
↳###!

&calculation
!type of theory
theory = 'dft'
/
```

*theory* specifies which theoretical method is used in the calculation.

```
&control
!common name of output files
sysname = 'Si'
/
```

*sysname* is a prefix for filenames of output files.

```
&units
!units used in input and output files
unit_system = 'a.u.'
/
```

*unit\_system* specifies which unit system is used in the input and output files.

```
&system
!periodic boundary condition
yn_periodic = 'y'

!grid box size(x,y,z)
al(1:3) = 10.26d0, 10.26d0, 10.26d0

!number of elements, atoms, electrons and states(bands)
nelem = 1
natom = 8
nelec = 32
nstate = 32
/
```

*yn\_periodic* specifies whether or not periodic boundary condition is applied.

*al(i)* specifies the side length of the unit cell.

*nelem* is the number of elements in the system.

*natom* is the number of atoms in the system.

*nelec* is the number of electrons in the system.

*nstate* is the number of orbitals that are used in the calculation.

```
&pseudo
!name of input pseudo potential file
file_pseudo(1) = './Si_rps.dat'

!atomic number of element
izatom(1) = 14

!angular momentum of pseudopotential that will be treated as local
lloc_ps(1) = 2
!--- Caution -----!
! Index must correspond to those in &atomic_red_coor.    !
!-----!
/
```

*file\_pseudo(n)* specifies the filename of the pseudopotential file of the n-th element.

*izatom(n)* is the atomic number of the n-th element.

*lloc\_ps(n)* specifies which angular momentum component is chosen as the local potential for the n-th element.

```
&functional
!functional('PZ' is Perdew-Zunger LDA: Phys. Rev. B 23, 5048 (1981).)
xc = 'PZ'
/
```

*xc* specifies the exchange-correlation potential to be used in the calculation.

```
&rgrid
!number of spatial grids(x,y,z)
num_rgrid(1:3) = 12, 12, 12
/
```

*num\_rgrid(i)* specifies the number of real-space grid point in i-th direction.

```
&kgrid
!number of k-points(x,y,z)
num_kgrid(1:3) = 4, 4, 4
/
```

*num\_kgrid(i)* specifies the number of k-points for i-th direction discretizing the Brillouin zone.

```
&scf
!maximum number of scf iteration and threshold of convergence
nscf      = 300
threshold = 1.0d-9
/
```

*nscf* specifies the maximum number of SCF iterations.

*threshold* specifies the threshold to judge the convergence.

```
&atomic_red_coor
!cartesian atomic reduced coordinates
'Si'      .0      .0      .0      1
'Si'      .25     .25     .25     1
'Si'      .5      .0      .5      1
'Si'      .0      .5      .5      1
'Si'      .5      .5      .0      1
'Si'      .75     .25     .75     1
'Si'      .25     .75     .75     1
'Si'      .75     .75     .25     1
!--- Format -----!
! 'symbol' x y z index(correspond to that of pseudo potential) !
!-----!
/
```



*&atomic\_red\_coor* specifies spatial coordinates of atoms in reduced coordinate system.

## Output files

After the calculation, following output files and a directory are created in the directory that you run the code,

name	description
<i>Si_info.data</i>	information on ground state solution
<i>Si_eigen.data</i>	energy eigenvalues of orbitals
<i>Si_k.data</i>	k-point distribution
<i>PS_Si_KY_n.dat</i>	information on pseudopotential file for silicon atom
<i>data_for_restart</i>	directory where files used in the real-time calculation are contained

You may download the above files (zipped file, except for the directory *data\_for\_restart*) from:

[https://salmon-tddft.jp/webmanual/v\\_2\\_0\\_1/exercise\\_zip\\_files/04\\_bulkSi\\_gs.zip](https://salmon-tddft.jp/webmanual/v_2_0_1/exercise_zip_files/04_bulkSi_gs.zip)

Main results of the calculation such as orbital energies are included in *Si\_info.data*. Explanations of the *Si\_info.data* and other output files are below:

### Si\_info.data

Calculated orbital and total energies as well as parameters specified in the input file are shown in this file.

### Si\_eigen.data

1 particle energies.

```
#esp: single-particle energies (eigen energies)
#occ: occupation numbers, io: orbital index
# 1:io, 2:esp[a.u.], 3:occ
```

### Si\_k.data

k-point distribution.

```
# ik: k-point index
# kx,ky,kz: Reduced coordinate of k-points
# wk: Weight of k-point
# 1:ik[none] 2:kx[none] 3:ky[none] 4:kz[none] 5:wk[none]
# coefficients (2*pi/a [a.u.]) in kx, ky, kz
```

## 3.3.2 Exercise-5: Dielectric function of crystalline silicon

In this exercise, we learn the linear response calculation of the crystalline silicon of a diamond structure. Calculation is done in a cubic unit cell that contains eight silicon atoms. This exercise should be carried out after finishing the ground state calculation that was explained in *Exercise-4*. An impulsive perturbation is applied to all electrons in the unit cell along  $z$  direction. Since the dielectric function is isotropic in the diamond structure, calculated dielectric function should not depend on the direction of the perturbation. During the time evolution, electric current averaged over the unit cell volume is calculated. A time-frequency Fourier transformation of the electric current gives us a frequency-dependent conductivity. The dielectric function may be obtained from the conductivity using a standard relation.

## Input files

To run the code, following files in samples are used:

file name	description
<i>Si_rt_response.inp</i>	input file that contain input keywords and their values
<i>Si_rps.dat</i>	pseudopotential file of silicon
<i>restart</i>	directory created in the ground state calculation (rename the directory from <i>data_for_restart</i> to <i>restart</i> )

In the input file *Si\_rt\_response.inp*, input keywords are specified. Most of them are mandatory to execute the calculation. This will help you to prepare the input file for other systems that you want to calculate. A complete list of the input keywords can be found in [List of input keywords](#).

```
#####
!#####
! Exercise 05: Dielectric function of crystalline silicon
!
!-----!
!---!
! * The detail of this exercise is explained in our manual(see chapter: 'Exercises').
!
! The manual can be obtained from: https://salmon-tddft.jp/documents.html
!
! * Input format consists of group of keywords like:
!
!      &group
!      input keyword = xxx
!
!      /
!
! (see chapter: 'List of input keywords' in the manual)
!
!-----!
!---!
! * Conversion from unit_system = 'a.u.' to 'A_eV_fs':
!
! Length: 1 [a.u.] = 0.52917721067      [Angstrom]
!
! Energy: 1 [a.u.] = 27.21138505        [eV]
!
! Time   : 1 [a.u.] = 0.02418884326505 [fs]
!
!-----!
!---!
! * Copy the ground state data directory('data_for_restart') (or make symbolic link)
!
! calculated in 'samples/exercise_04_bulkSi_gs/' and rename the directory to
! 'restart/'!
! in the current directory.
!
!#####
!#####
```

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```
&calculation
!type of theory
theory = 'tddft_response'
/
```

*theory* specifies which theoretical method is used in the calculation.

```
&control
!common name of output files
sysname = 'Si'
/
```

*sysname* is a prefix for filenames of output files.

```
&units
!units used in input and output files
unit_system = 'a.u.'
/
```

*unit\_system* specifies which unit system is used in the input and output files.

```
&system
!periodic boundary condition
yn_periodic = 'y'

!grid box size(x,y,z)
al(1:3) = 10.26d0, 10.26d0, 10.26d0

!number of elements, atoms, electrons and states(bands)
nelem = 1
natom = 8
nelec = 32
nstate = 32
/
```

*yn\_periodic* specifies whether or not periodic boundary condition is applied.

*al(i)* specifies the side length of the unit cell.

*nelem* is the number of elements in the system.

*natom* is the number of atoms in the system.

*nelec* is the number of electrons in the system.

*nstate* is the number of orbitals that are used in the calculation.

```
&pseudo
!name of input pseudo potential file
file_pseudo(1) = './Si_rps.dat'

!atomic number of element
izatom(1) = 14

!angular momentum of pseudopotential that will be treated as local
lloc_ps(1) = 2
!--- Caution -----!
! Index must correspond to those in &atomic_red_coor.  !
!-----!
/
```

*file\_pseudo(n)* specifies the filename of the pseudopotential file of the n-th element.

*izatom(n)* is the atomic number of the n-th element.

*lloc\_ps(n)* specifies which angular momentum component is chosen as the local potential for the n-th element.

```
&functional
!functional('PZ' is Perdew-Zunger LDA: Phys. Rev. B 23, 5048 (1981).)
xc = 'PZ'
/
```

*xc* specifies the exchange-correlation potential to be used in the calculation.

```
&rgrid
!number of spatial grids(x,y,z)
num_rgrid(1:3) = 12, 12, 12
/
```

*num\_rgrid(i)* specifies the number of real-space grid point in i-th direction.

```
&kgrid
!number of k-points(x,y,z)
num_kgrid(1:3) = 4, 4, 4
/
```

*num\_kgrid(i)* specifies the number of k-points for i-th direction discretizing the Brillouin zone.

```
&tgrid
!time step size and number of time grids(steps)
dt = 0.08d0
nt = 6000
/
```

*dt* specifies the time step.

*nt* is the number of time steps for the time propagation.

```
&emfield
!envelope shape of the incident pulse('impulse': impulsive field)
ae_shape1 = 'impulse'

!polarization unit vector(real part) for the incident pulse(x,y,z)
epdir_re1(1:3) = 0.00d0, 0.00d0, 1.00d0
!--- Caution -----!
! Defenition of the incident pulse is wrritten in:           !
! https://www.sciencedirect.com/science/article/pii/S0010465518303412 !
!-----!
/
```

*ae\_shape1* specifies the envelope of the field. For a linear response calculation, as\_shape1='impulse' is used. It indicates that a weak impulsive perturbation is applied at  $t = 0$ .

*epdir\_re1(i)* specifies the i-th component of the real part of the polarization unit vector.

```
&analysis
!energy grid size and number of energy grids for output files
de      = 4.0d-4
nenergy = 5000
/
```

*de* specifies the energy grid size for frequency-domain analysis.

*nenergy* specifies the number of energy grid points for frequency-domain analysis.

```
&atomic_red_coor
!cartesian atomic reduced coodinates
'Si'      .0      .0      .0      1
'Si'      .25     .25     .25     1
'Si'      .5      .0      .5      1
'Si'      .0      .5      .5      1
'Si'      .5      .5      .0      1
'Si'      .75     .25     .75     1
'Si'      .25     .75     .75     1
'Si'      .75     .75     .25     1
!--- Format -----!
! 'symbol' x y z index(correspond to that of pseudo potential) !
!-----!
/
```

*&atomic\_red\_coor* specifies spatial coordinates of atoms in reduced coordinate system.

## Output files

After the calculation, following output files are created in the directory that you run the code,

file name	description
<i>Si_response.data</i>	Fourier spectra of the conductivity and dielectric functions
<i>Si_rt.data</i>	vector potential, electric field, and matter current as functions of time
<i>Si_rt_energy</i>	components of total energy and difference of total energy as functions of time
<i>PS_Si_KY_n.dat</i>	information on pseudopotential file for silicon atom

You may download the above files (zipped file) from:

[https://salmon-tddft.jp/webmanual/v\\_2\\_0\\_1/exercise\\_zip\\_files/05\\_bulkSi\\_lr.zip](https://salmon-tddft.jp/webmanual/v_2_0_1/exercise_zip_files/05_bulkSi_lr.zip)

Explanations of the output files are described below:

### Si\_response.data

Time-frequency Fourier transformation of the macroscopic current gives the conductivity of the system. Then the dielectric function is calculated.

```
# Fourier-transform spectra:
# sigma: Conductivity
# eps: Dielectric constant
# 1:Energy[a.u.] 2:Re(sigma_x)[a.u.] 3:Re(sigma_y)[a.u.] 4:Re(sigma_z)[a.u.]
↪5:Im(sigma_x)[a.u.] 6:Im(sigma_y)[a.u.] 7:Im(sigma_z)[a.u.] 8:Re(eps_x)[none]
↪9:Re(eps_y)[none] 10:Re(eps_z)[none] 11:Im(eps_x)[none] 12:Im(eps_y)[none]
↪13:Im(eps_z)[none]
```

### Si\_rt.data

Results of time evolution calculation for vector potential, electric field, and matter current density.

```
# Real time calculation:
# Ac_ext: External vector potential field
# E_ext: External electric field
# Ac_tot: Total vector potential field
# E_tot: Total electric field
# Jm: Matter current density (electrons)
# 1:Time[a.u.] 2:Ac_ext_x[a.u.] 3:Ac_ext_y[a.u.] 4:Ac_ext_z[a.u.] 5:E_ext_x[a.u.] 6:E_
↪ext_y[a.u.] 7:E_ext_z[a.u.] 8:Ac_tot_x[a.u.] 9:Ac_tot_y[a.u.] 10:Ac_tot_z[a.u.]
↪11:E_tot_x[a.u.] 12:E_tot_y[a.u.] 13:E_tot_z[a.u.] 14:Jm_x[a.u.] 15:Jm_y[a.u.]
↪16:Jm_z[a.u.]
```

### Si\_rt\_energy

$E_{all}$  and  $E_{all}-E_{all0}$  are total energy and electronic excitation energy, respectively.

```
# Real time calculation:
# Eall: Total energy
# Eall0: Initial energy
# 1:Time[a.u.] 2:Eall[a.u.] 3:Eall-Eall0[a.u.]
```

## Input files

file name	description
<i>Si_rt_pulse.inp</i>	input file that contain input keywords and their values.
<i>Si_rps.dat</i>	pseudopotential file for Carbon
<i>restart</i>	directory created in the ground state calculation (rename the directory from <i>data_for_restart</i> to <i>restart</i> )

```

#####
↳###!
! Exercice 06: Electron dynamics in crystalline silicon under a pulsed electric_
↳field !
!-----!
↳---!
! * The detail of this exercice is explained in our manual(see chapter: 'Exercises')._
↳!
! The manual can be obtained from: https://salmon-tddft.jp/documents.html
↳!
! * Input format consists of group of keywords like:
↳!
! &group
↳!
! input keyword = xxx
↳!
! /
↳!
! (see chapter: 'List of input keywords' in the manual)
↳!
!-----!
↳---!
! * Conversion from unit_system = 'a.u.' to 'A_eV_fs':
↳!
! Length: 1 [a.u.] = 0.52917721067 [Angstrom]
↳!
! Energy: 1 [a.u.] = 27.21138505 [eV]
↳!
! Time : 1 [a.u.] = 0.02418884326505 [fs]
↳!

```

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```

!-----!
! * Copy the ground state data directory('data_for_restart') (or make symbolic link)
!
!   calculated in 'samples/exercise_04_bulkSi_gs/' and rename the directory to
!   'restart/'!
!   in the current directory.
!
!#####
!###!

&calculation
!type of theory
theory = 'tddft_pulse'
/

```

*theory* specifies which theoretical method is used in the calculation.

```

&control
!common name of output files
sysname = 'Si'
/

```

*sysname* is a prefix for filenames of output files.

```

&units
!units used in input and output files
unit_system = 'a.u.'
/

```

*unit\_system* specifies which unit system is used in the input and output files.

```

&system
!periodic boundary condition
yn_periodic = 'y'

!grid box size(x,y,z)
al(1:3) = 10.26d0, 10.26d0, 10.26d0

!number of elements, atoms, electrons and states(bands)
nelem = 1
natom = 8
nelec = 32
nstate = 32
/

```



*yn\_periodic* specifies whether or not periodic boundary condition is applied.

*al(i)* specifies the side length of the unit cell.

*nelem* is the number of elements in the system.

*natom* is the number of atoms in the system.

*nelec* is the number of electrons in the system.

*nstate* is the number of orbitals that are used in the calculation.

```
&pseudo
!name of input pseudo potential file
file_pseudo(1) = './Si_rps.dat'

!atomic number of element
izatom(1) = 14

!angular momentum of pseudopotential that will be treated as local
lloc_ps(1) = 2
!--- Caution -----!
! Index must correspond to those in &atomic_red_coor.    !
!-----!
/
```

*file\_pseudo(n)* specifies the filename of the pseudopotential file of the n-th element.

*izatom(n)* is the atomic number of the n-th element.

*lloc\_ps(n)* specifies which angular momentum component is chosen as the local potential for the n-th element.

```
&functional
!functional('PZ' is Perdew-Zunger LDA: Phys. Rev. B 23, 5048 (1981).)
xc = 'PZ'
/
```

*xc* specifies the exchange-correlation potential to be used in the calculation.

```
&rgrid
!number of spatial grids(x,y,z)
num_rgrid(1:3) = 12, 12, 12
/
```

*num\_rgrid(i)* specifies the number of real-space grid point in i-th direction.

```
&kgrid
!number of k-points(x,y,z)
num_kgrid(1:3) = 4, 4, 4
/
```

*num\_kgrid(i)* specifies the number of k-points for i-th direction discretizing the Brillouin zone.

```
&tgrid
!time step size and number of time grids(steps)
dt = 0.08d0
nt = 6000
/
```

*dt* specifies the time step.

*nt* is the number of time steps for the time propagation.

```
&emfield
!envelope shape of the incident pulse('Acos2': cos^2 type envelope for vector_
↪potential)
ae_shape1 = 'Acos2'

!peak intensity(W/cm^2) of the incident pulse
I_wcm2_1 = 5.0d11

!duration of the incident pulse
tw1 = 441.195136248d0

!mean photon energy(average frequency multiplied by the Planck constant) of the_
↪incident pulse
omega1 = 0.05696145187d0

!polarization unit vector(real part) for the incident pulse(x,y,z)
epdir_re1(1:3) = 0.0d0, 0.0d0, 1.0d0
!--- Caution -----!
! Defenition of the incident pulse is wrritten in:                !
! https://www.sciencedirect.com/science/article/pii/S0010465518303412 !
!-----!
/
```

*ae\_shape1* specifies the envelope of the field.

*I\_wcm2\_1* specify the intensity of the pulse in unit of  $\text{W}/\text{cm}^2$ .

*tw1* specifies the duration of the pulse.

*omega1* specifies the mean photon energy of the pulse.

*epdir\_re1(i)* specifies the i-th component of the real part of the polarization unit vector.

```
&atomic_red_coor
!cartesian atomic reduced coodinates
'Si'      .0      .0      .0      1
'Si'      .25     .25     .25     1
'Si'      .5      .0      .5      1
'Si'      .0      .5      .5      1
'Si'      .5      .5      .0      1
'Si'      .75     .25     .75     1
```

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```
'Si'      .25      .75      .75      1
'Si'      .75      .75      .25      1
!--- Format -----!
! 'symbol' x y z index(correspond to that of pseudo potential) !
!-----!
/
```

*&atomic\_red\_coor* specifies spatial coordinates of atoms in reduced coordinate system.

## Output files

After the calculation, following output files are created in the directory that you run the code,

file name	description
<i>Si_pulse.data</i>	matter current and electric field as functions of energy
<i>Si_rt.data</i>	vector potential, electric field, and matter current as functions of time
<i>Si_rt_energy</i>	components of total energy and difference of total energy as functions of time
<i>PS_Si_KY_n.dat</i>	information on pseudopotential file for silicon atom

You may download the above files (zipped file) from:

[https://salmon-tddft.jp/webmanual/v\\_2\\_0\\_1/exercise\\_zip\\_files/06\\_bulkSi\\_rt.zip](https://salmon-tddft.jp/webmanual/v_2_0_1/exercise_zip_files/06_bulkSi_rt.zip)

Explanations of the output files are described below:

### Si\_pulse.data

Time-frequency Fourier transformation of the matter current and electric field.

```
# Fourier-transform spectra:
# energy: Frequency
# Jm: Matter current
# E_ext: External electric field
# E_tot: Total electric field
# 1:energy[a.u.] 2:Re(Jm_x)[a.u.] 3:Re(Jm_y)[a.u.] 4:Re(Jm_z)[a.u.] 5:Im(Jm_x)[a.u.]
↪ 6:Im(Jm_y)[a.u.] 7:Im(Jm_z)[a.u.] 8:|Jm_x|^2[a.u.] 9:|Jm_y|^2[a.u.] 10:|Jm_z|^2[a.u.]
↪ 11:Re(E_ext_x)[a.u.] 12:Re(E_ext_y)[a.u.] 13:Re(E_ext_z)[a.u.] 14:Im(E_ext_x)[a.u.]
↪ 15:Im(E_ext_y)[a.u.] 16:Im(E_ext_z)[a.u.] 17:|E_ext_x|^2[a.u.] 18:|E_ext_y|^2[a.u.]
↪ 19:|E_ext_z|^2[a.u.] 20:Re(E_ext_x)[a.u.] 21:Re(E_ext_y)[a.u.] 22:Re(E_ext_z)[a.u.]
↪ 23:Im(E_ext_x)[a.u.] 24:Im(E_ext_y)[a.u.] 25:Im(E_ext_z)[a.u.] 26:|E_ext_x|^2[a.u.]
↪ 27:|E_ext_y|^2[a.u.] 28:|E_ext_z|^2[a.u.]
```

### Si\_rt.data

Results of time evolution calculation for vector potential, electric field, and matter current density.

```
# Real time calculation:
# Ac_ext: External vector potential field
# E_ext: External electric field
# Ac_tot: Total vector potential field
# E_tot: Total electric field
```

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```
# Jm: Matter current density (electrons)
# 1:Time[a.u.] 2:Ac_ext_x[a.u.] 3:Ac_ext_y[a.u.] 4:Ac_ext_z[a.u.] 5:E_ext_x[a.u.] 6:E_
↪ext_y[a.u.] 7:E_ext_z[a.u.] 8:Ac_tot_x[a.u.] 9:Ac_tot_y[a.u.] 10:Ac_tot_z[a.u.]_
↪11:E_tot_x[a.u.] 12:E_tot_y[a.u.] 13:E_tot_z[a.u.] 14:Jm_x[a.u.] 15:Jm_y[a.u.]_
↪16:Jm_z[a.u.]
```

**Si\_rt\_energy**

*Eall* and *Eall-Eall0* are total energy and electronic excitation energy, respectively.

```
# Real time calculation:
# Eall: Total energy
# Eall0: Initial energy
# 1:Time[a.u.] 2:Eall[a.u.] 3:Eall-Eall0[a.u.]
```

## 3.4 Maxwell + TDDFT multiscale simulation

### 3.4.1 Exercise-7: Pulsed-light propagation through a silicon thin film

In this exercise, we learn the calculation of the propagation of a pulsed light through a thin film of crystalline silicon. We consider a silicon thin film of 42 nm thickness, and an irradiation of a few-cycle, linearly polarized pulsed light normally on the thin film. This exercise should be carried out after finishing the ground state calculation that was explained in [Exercise-4](#). The pulsed light locates in the vacuum region in front of the thin film. The parameters that characterize the pulsed light such as magnitude and frequency are specified in the input file.

**Input files**

To run the code, following files in samples are used:

file name	description
<i>Si_rt_multiscale.inp</i>	input file that contain input keywords and their values.
<i>Si_rps.dat</i>	pseudopotential file for silicon
<i>restart</i>	directory created in the ground state calculation (rename the directory from <i>data_for_restart</i> to <i>restart</i> )

In the input file *Si\_rt\_multiscale.inp*, input keywords are specified. Most of them are mandatory to execute the calculation. This will help you to prepare the input file for other systems that you want to calculate. A complete list of the input keywords can be found in [List of input keywords](#).

```
!#####
↪###!
! Exercise 07: Maxwell+TDDFT multiscale simulation
↪ !
! (Pulsed-light propagation through a silicon thin film)
↪ !
!-----!
↪---!
! * The detail of this exercise is explained in our manual(see chapter: 'Exercises')._
↪ !
```

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```

!   The manual can be obtained from: https://salmon-tddft.jp/documents.html
↳ !
! * Input format consists of group of keywords like:
↳ !
!   &group
↳ !
!       input keyword = xxx
↳ !
!   /
↳ !
!   (see chapter: 'List of input keywords' in the manual)
↳ !
!-----!
↳ ---!
! * Conversion from unit_system = 'a.u.' to 'A_eV_fs':
↳ !
!   Length: 1 [a.u.] = 0.52917721067      [Angstrom]
↳ !
!   Energy: 1 [a.u.] = 27.21138505        [eV]
↳ !
!   Time   : 1 [a.u.] = 0.02418884326505 [fs]
↳ !
!-----!
↳ ---!
! * Copy the ground state data directory('data_for_restart') (or make symbolic link)
↳ !
!   calculated in 'samples/exercise_04_bulkSi_gs/' and rename the directory to
↳ 'restart/!'
!   in the current directory.
↳ !
!#####
↳###!

&calculation
!type of theory
theory = 'multi_scale_maxwell_tddft'
/

```

*theory* specifies which theoretical method is used in the calculation.

```

&control
!common name of output files
sysname = 'Si'
/

```

*sysname* is a prefix for filenames of output files.

```

&units
!units used in input and output files

```

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```
unit_system = 'a.u.'  
/
```

*unit\_system* specifies which unit system is used in the input and output files.

```
&system  
!periodic boundary condition  
yn_periodic = 'y'  
  
!grid box size(x,y,z)  
al(1:3) = 10.26d0, 10.26d0, 10.26d0  
  
!number of elements, atoms, electrons and states(bands)  
nelem = 1  
natom = 8  
nelec = 32  
nstate = 32  
/
```

*yn\_periodic* specifies whether or not periodic boundary condition is applied.

*al(i)* specifies the side length of the unit cell.

*nelem* is the number of elements in the system.

*natom* is the number of atoms in the system.

*nelec* is the number of electrons in the system.

*nstate* is the number of orbitals that are used in the calculation.

```
&pseudo  
!name of input pseudo potential file  
file_pseudo(1) = './Si_rps.dat'  
  
!atomic number of element  
izatom(1) = 14  
  
!angular momentum of pseudopotential that will be treated as local  
lloc_ps(1) = 2  
!--- Caution -----!  
! Index must correspond to those in &atomic_red_coor.  !  
!-----!  
/
```

*file\_pseudo(n)* specifies the filename of the pseudopotential file of the n-th element.

*izatom(n)* is the atomic number of the n-th element.

*lloc\_ps(n)* specifies which angular momentum component is chosen as the local potential for the n-th element.

```
&functional
!functional('PZ' is Perdew-Zunger LDA: Phys. Rev. B 23, 5048 (1981).)
xc = 'PZ'
/
```

*xc* specifies the exchange-correlation potential to be used in the calculation.

```
&rgrid
!number of spatial grids(x,y,z)
num_rgrid(1:3) = 12, 12, 12
/
```

*num\_rgrid(i)* specifies the number of real-space grid point in i-th direction.

```
&kgrid
!number of k-points(x,y,z)
num_kgrid(1:3) = 4, 4, 4
/
```

*num\_kgrid(i)* specifies the number of k-points for i-th direction discretizing the Brillouin zone.

```
&tgrid
!time step size and number of time grids(steps)
dt = 0.08d0
nt = 6000
/
```

*dt* specifies the time step.

*nt* is the number of time steps for the time propagation.

```
&emfield
!envelope shape of the incident pulse('Acos2': cos^2 type envelope for vector_
↪potential)
ae_shape1 = 'Acos2'

!peak intensity(W/cm^2) of the incident pulse
I_wcm2_1 = 1.0d12

!duration of the incident pulse
tw1 = 441.195136248d0

!mean photon energy(average frequency multiplied by the Planck constant) of the_
↪incident pulse
omega1 = 0.05696145187d0
```

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```

!polarization unit vector(real part) for the incident pulse(x,y,z)
epdir_re1(1:3) = 0.0d0, 0.0d0, 1.0d0
!--- Caution -----!
! Defenition of the incident pulse is wrriten in:                !
! https://www.sciencedirect.com/science/article/pii/S0010465518303412 !
!-----!
/

```

*ae\_shape1* specifies the envelope of the field.

*I\_wcm2\_1* specify the intensity of the pulse in unit of W/cm<sup>2</sup>.

*tw1* specifies the duration of the pulse.

*omega1* specifies the mean photon energy of the pulse.

*epdir\_re1(i)* specifies the i-th component of the real part of the polarization unit vector.

```

&multiscale
!number of macro grids in electromagnetic analysis for x, y, and z directions
nx_m = 8
ny_m = 1
nz_m = 1

!macro grid spacing for x, y, and z directions
hx_m = 100.0d0
hy_m = 100.0d0
hz_m = 100.0d0

!number of macroscopic grids for vacumm region
!(nxvac1_m is for negative x-direction in front of material)
!(nxvacr_m is for positive x-direction behind material)
nxvac1_m = 1000
nxvacr_m = 1000
/

```

*nx\_m, ny\_m, nz\_m* specify the number of macroscopic grid points inside the material.

*hx\_m, hy\_m, hz\_m* specify the grid spacing of macroscopic coordinates.

*nxvac1\_m / nxvacr\_m* specifies the number of grid points in the vacuum region in the left / right side of the material.

```

&maxwell
!boundary condition of electromagnetic analysis
!first index(1-3 rows) corresponds to x, y, and z directions
!second index(1-2 columns) corresponds to bottom and top of the directions
!('abc' is absorbing boundary condition)
boundary_em(1,1) = 'abc'
boundary_em(1,2) = 'abc'
/

```

*boundary\_em(i,n)* specifies the boundary condition for the electromagnetic analysis. The first index i corresponds to the x,y, and z direction. The second index n specifies left or right side of the material.



```

&atomic_red_coor
!cartesian atomic reduced coordinates
'Si'      .0      .0      .0      1
'Si'      .25     .25     .25     1
'Si'      .5      .0      .5      1
'Si'      .0      .5      .5      1
'Si'      .5      .5      .0      1
'Si'      .75     .25     .75     1
'Si'      .25     .75     .75     1
'Si'      .75     .75     .25     1
!---- Format -----!
! 'symbol' x y z index(correspond to that of pseudo potential) !
!-----!
/

```

*&atomic\_red\_coor* specifies spatial coordinates of atoms in reduced coordinate system.

## Output files

After the calculation, new directory *multiscale/* is created, then, following output files are created in the directory,

file name	description
<i>Si_m/mxxxxxx/Si_rt.data</i>	vector potential, electric field, and matter current at macroscopic position xxxxxx as functions of time
<i>Si_m/mxxxxxx/Si_rt_energy.data</i>	components of total energy and difference of total energy at macroscopic position xxxxxx as functions of time
<i>Si_m/mxxxxxx/PS_Si_KY_n.dat</i>	information on pseudopotential file for silicon atom at macroscopic position xxxxxx
<i>Si_RT_Ac/Si_Ac_yyyyyy.data</i>	vector potential, electric field, magnetic field, electromagnetic current density at time step yyyyyy as function of space
<i>Si_wave.data</i>	amplitudes of incident, reflected, and transmitted wave

You may download the above files (zipped file) from:

[https://salmon-tddft.jp/webmanual/v\\_2\\_0\\_1/exercise\\_zip\\_files/07\\_bulkSi\\_ms.zip](https://salmon-tddft.jp/webmanual/v_2_0_1/exercise_zip_files/07_bulkSi_ms.zip)

Explanations of the output files are described below:

**Si\_m/mxxxxxx/Si\_rt.data**

The number in the file name specifies the macroscopic position. Results of time evolution calculation for vector potential, electric field, and matter current density.

```
# Real time calculation:
# Ac_ext: External vector potential field
# E_ext: External electric field
# Ac_tot: Total vector potential field
# E_tot: Total electric field
# Jm: Matter current density (electrons)
# 1:Time[a.u.] 2:Ac_ext_x[a.u.] 3:Ac_ext_y[a.u.] 4:Ac_ext_z[a.u.] 5:E_ext_x[a.u.] 6:E_
→ext_y[a.u.] 7:E_ext_z[a.u.] 8:Ac_tot_x[a.u.] 9:Ac_tot_y[a.u.] 10:Ac_tot_z[a.u.]
→11:E_tot_x[a.u.] 12:E_tot_y[a.u.] 13:E_tot_z[a.u.] 14:Jm_x[a.u.] 15:Jm_y[a.u.]
→16:Jm_z[a.u.]
```

**Si\_m/mxxxxxx/Si\_rt\_energy.data**

The number in the file name specifies the macroscopic position. *Eall* and *Eall-Eall0* are total energy and electronic excitation energy, respectively.

```
# Real time calculation:
# Eall: Total energy
# Eall0: Initial energy
# 1:Time[a.u.] 2:Eall[a.u.] 3:Eall-Eall0[a.u.]
```

**Si\_RT\_Ac/Si\_Ac\_yyyyyy.data**

The number in the file name specifies the iteration number. Various quantities at a time are shown as function of macroscopic position.

```
# Multiscale TDDFT calculation
# IX, IY, IZ: FDTD Grid index
# x, y, z: Coordinates
# Ac: Vector potential field
# E: Electric field
# Jem: Electromagnetic current density
# 1:IX[none] 2:IY[none] 3:IZ[none] 4:Ac_x[a.u.] 5:Ac_y[a.u.] 6:Ac_z[a.u.] 7:E_x[a.u.]
→8:E_y[a.u.] 9:E_z[a.u.] 10:B_x[a.u.] 11:B_y[a.u.] 12:B_z[a.u.] 13:Jem_x[a.u.]
→14:Jem_y[a.u.] 15:Jem_z[a.u.] 16:E_em[a.u./vol] 17:E_abs[a.u./vol]
```

**Si\_wave.data**

Amplitudes of incident, reflected, and transmitted wave.

```
# 1D multiscale calculation:
# E_inc: E-field amplitude of incident wave
# E_ref: E-field amplitude of reflected wave
# E_tra: E-field amplitude of transmitted wave
# 1:Time[a.u.] 2:E_inc_x[a.u.] 3:E_inc_y[a.u.] 4:E_inc_z[a.u.] 5:E_ref_x[a.u.] 6:E_
→ref_y[a.u.] 7:E_ref_z[a.u.] 8:E_tra_x[a.u.] 9:E_tra_y[a.u.] 10:E_tra_z[a.u.]
```

## 3.5 Geometry optimization and Ehrenfest molecular dynamics

### 3.5.1 Exercise-8: Geometry optimization of C2H2 molecule

In this exercise, we learn the calculation of geometry optimization of acetylene (C2H2) molecule, solving the static Kohn-Sham equation. This exercise will be useful to learn how to set up calculations in SALMON for any isolated systems such as molecules and nanoparticles.

#### Input files

To run the code, following files in samples are used:

file name	description
<i>C2H2_opt.inp</i>	input file that contains input keywords and their values
<i>C_rps.dat</i>	pseudopotential file for carbon atom
<i>H_rps.dat</i>	pseudopotential file for hydrogen atom

In the input file *C2H2\_opt.inp*, input keywords are specified. Most of them are mandatory to execute the geometry optimization. This will help you to prepare an input file for other systems that you want to calculate. A complete list of the input keywords that can be used in the input file can be found in [List of input keywords](#).

```
#####
↳###!
! Exercise 08: Geometry optimization of C2H2 molecule
↳ !
!-----!
↳---!
! * The detail of this exercise is explained in our manual(see chapter: 'Exercises')..
↳ !
!   The manual can be obtained from: https://salmon-tddft.jp/documents.html
↳ !
! * Input format consists of group of keywords like:
↳ !
!   &group
↳ !
!   input keyword = xxx
↳ !
!   /
↳ !
!   (see chapter: 'List of input keywords' in the manual)
↳ !
!-----!
↳---!
! * Conversion from unit_system = 'a.u.' to 'A_eV_fs':
↳ !
!   Length: 1 [a.u.] = 0.52917721067 [Angstrom]
↳ !
!   Energy: 1 [a.u.] = 27.21138505 [eV]
↳ !
!   Time : 1 [a.u.] = 0.02418884326505 [fs]
↳ !
!#####
↳###!
```

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```
&calculation
!type of theory
theory = 'dft'

!geometry optimization option
yn_opt = 'y'
/
```

*theory* specifies which theoretical method is used in the calculation.

*yn\_opt* is a switch to carry out the structure optimization.

```
&control
!common name of output files
sysname = 'C2H2'
/
```

*sysname* is a prefix for filenames of output files.

```
&units
!units used in input and output files
unit_system = 'A_eV_fs'
/
```

*unit\_system* specifies which unit system is used in the input and output files.

```
&system
!periodic boundary condition
yn_periodic = 'n'

!grid box size(x,y,z)
al(1:3) = 12.0d0, 12.0d0, 16.0d0

!number of elements, atoms, electrons and states(orbitals)
nelem = 2
natom = 4
nelec = 10
nstate = 6
/
```

*yn\_periodic* specifies whether or not periodic boundary condition is applied.

*al(i)* specifies the spatial box size of the cubiod cell.

*nelem* is the number of elements in the system.

*natom* is the number of atoms in the system.

*nelec* is the number of electrons in the system.

*nstate* is the number of orbitals that are used in the calculation.

```
&pseudo
!name of input pseudo potential file
file_pseudo(1) = './C_rps.dat'
file_pseudo(2) = './H_rps.dat'

!atomic number of element
izatom(1) = 6
izatom(2) = 1

!angular momentum of pseudopotential that will be treated as local
lloc_ps(1) = 1
lloc_ps(2) = 0
!--- Caution -----!
! Indices must correspond to those in &atomic_coor. !
!-----!
/
```

*file\_pseudo(n)* specifies the filename of the pseudopotential file of the n-th element.

*izatom(n)* is the atomic number of the n-th element.

*lloc\_ps(n)* specifies which angular momentum component is chosen as the local potential for the n-th element.

```
&functional
!functional('PZ' is Perdew-Zunger LDA: Phys. Rev. B 23, 5048 (1981).)
xc = 'PZ'
/
```

*xc* specifies the exchange-correlation potential to be used in the calculation.

```
&rgrid
!spatial grid spacing(x,y,z)
dl(1:3) = 0.20d0, 0.20d, 0.20d0
/
```

*dl(i)* specifies the spatial grid spacing in i-th direction.

```
&scf
!maximum number of scf iteration and threshold of convergence for ground state_
↪ calculation
nscf      = 300
threshold = 1.0d-9
/
```

*nscf* specifies the maximum number of SCF iterations.

*threshold* specifies the threshold to judge the convergence.

```
&opt
!threshold(maximum force on atom) of convergence for geometry optimization
convrg_opt_fmax = 1.0d-3
/

&atomic_coor
!cartesian atomic coordinates
'C'    0.0    0.0    0.6  1  y
'H'    0.0    0.0    1.7  2  y
'C'    0.0    0.0   -0.6  1  y
'H'    0.0    0.0   -1.7  2  y
!--- Format -----!
! 'symbol' x y z index(correspond to that of pseudo potential) y/n !
!--- Caution -----!
! final index(y/n) determines free/fix for the atom coordinate.    !
!-----!
/
```

*&atomic\_coor* specifies spatial coordinates of atoms.

## Output files

After the calculation, following output files and a directory are created in the directory that you run the code,

name	description
<i>C2H2_info.data</i>	information on ground state solution
<i>C2H2_eigen.data</i>	1 particle energies
<i>C2H2_trj.xyz</i>	atomic coordinates during the geometry optimization
<i>C2H2_k.data</i>	k-point distribution (for isolated systems, only gamma point is described)
<i>data_for_restart</i>	directory where files used in the real-time calculation are contained
<i>PS_C_KY_n.dat</i>	information on pseudopotential file for carbon atom
<i>PS_H_KY_n.dat</i>	information on pseudopotential file for hydrogen atom

You may download the above files (zipped file, except for the directory *data\_for\_restart*) from:

[https://salmon-tddft.jp/webmanual/v\\_2\\_0\\_1/exercise\\_zip\\_files/08\\_C2H2\\_opt.zip](https://salmon-tddft.jp/webmanual/v_2_0_1/exercise_zip_files/08_C2H2_opt.zip)

Main results of the calculation such as orbital energies are included in *C2H2\_info.data*. Explanations of the *C2H2\_info.data* and other output files are below:

### **C2H2\_info.data**

Calculated orbital and total energies as well as parameters specified in the input file are shown in this file.

### **C2H2\_eigen.data**

1 particle energies.

```
#esp: single-particle energies (eigen energies)
#occ: occupation numbers, io: orbital index
# 1:io, 2:esp[eV], 3:occ
```

### C2H2\_trj.xyz

The atomic coordinates during the geometry optimization in xyz format.

### C2H2\_k.data

k-point distribution(for isolated systems, only gamma point is described).

```
# ik: k-point index
# kx,ky,kz: Reduced coordinate of k-points
# wk: Weight of k-point
# 1:ik[none] 2:kx[none] 3:ky[none] 4:kz[none] 5:wk[none]
# coefficients (2*pi/a [a.u.]) in kx, ky, kz
```

## 3.5.2 Exercise-9: Ehrenfest molecular dynamics of C2H2 molecule

In this exercise, we learn the calculation of the molecular dynamics in the acetylene (C2H2) molecule under a pulsed electric field, solving the time-dependent Kohn-Sham equation and the Newtonian equation. As outputs of the calculation, time-evolution of the electron density as well as molecular structures and associated quantities such as the electron and ion kinetic energies, the electric dipole moment of the system and temperature as functions of time are calculated.. This tutorial should be carried out after finishing the geometry optimization that was explained in [Exercise-8](#). In the calculation, a pulsed electric field that has  $\cos^2$  envelope shape is applied. The parameters that characterize the pulsed field such as magnitude, frequency, polarization direction, and carrier envelope phase are specified in the input file.

### Input files

To run the code, following files in samples are used. The directory *restart* is created in the ground state calculation as *data\_for\_restart*. Pseudopotential files are already used in the geometry optimization. Therefore, *C2H2\_md.inp* that specifies input keywords and their values for the pulsed electric field and molecular dynamics calculations is the only file that the users need to prepare.

file name	description
<i>C2H2_md.inp</i>	input file that contain input keywords and their values.
<i>C_rps.dat</i>	pseudopotential file for carbon
<i>H_rps.dat</i>	pseudopotential file for hydrogen
<i>restart</i>	directory created in the geometry optimization (rename the directory from <i>data_for_restart</i> to <i>restart</i> )

In the input file *C2H2\_md.inp*, input keywords are specified. Most of them are mandatory to execute the calculation of electron dynamics induced by a pulsed electric field. This will help you to prepare the input file for other systems and other pulsed electric fields with molecular dynamics calculation that you want to calculate. A complete list of the input keywords that can be used in the input file can be found in [List of input keywords](#).

```

#####
!###!
! Exercise 09: Ehrenfest molecular dynamics of C2H2 molecule
!
!-----!
!---!
! * The detail of this exercise is explained in our manual(see chapter: 'Exercises').
!
! The manual can be obtained from: https://salmon-tddft.jp/documents.html
!
! * Input format consists of group of keywords like:
!
!     &group
!
!         input keyword = xxx
!
!     /
!
! (see chapter: 'List of input keywords' in the manual)
!
!-----!
!---!
! * Conversion from unit_system = 'a.u.' to 'A_eV_fs':
!
! Length: 1 [a.u.] = 0.52917721067      [Angstrom]
!
! Energy: 1 [a.u.] = 27.21138505      [eV]
!
! Time   : 1 [a.u.] = 0.02418884326505 [fs]
!
!-----!
!---!
! * Ehrenfest-MD option is still trial.
!
! * Copy the ground state data directory ('data_for_restart') (or make symbolic link)
!
! calculated in 'samples/exercise_08_C2H2_opt/' and rename the directory to
! 'restart/' !
! in the current directory.
!
!#####
!###!

&calculation
!type of theory
theory = 'tddft_pulse'

!molecular dynamics option
yn_md = 'y'
/

```

*theory* specifies which theoretical method is used in the calculation.

*yn\_md* is a switch for Ehrenfest molecular dynamics.



```
&control
!common name of output files
sysname = 'C2H2'
/
```

*sysname* is a prefix for filenames of output files.

```
&units
!units used in input and output files
unit_system = 'A_eV_fs'
/
```

*unit\_system* specifies which unit system is used in the input and output files.

```
&system
!periodic boundary condition
yn_periodic = 'n'

!grid box size(x,y,z)
al(1:3) = 12.0d0, 12.0d0, 16.0d0

!number of elements, atoms, electrons and states(orbitals)
nelem = 2
natom = 4
nelec = 10
nstate = 6
/
```

*yn\_periodic* specifies whether or not periodic boundary condition is applied.

*al(i)* specifies the spatial box size of the cubiod cell.

*nelem* is the number of elements in the system.

*natom* is the number of atoms in the system.

*nelec* is the number of electrons in the system.

*nstate* is the number of orbitals that are used in the calculation.

```
&pseudo
!name of input pseudo potential file
file_pseudo(1) = './C_rps.dat'
file_pseudo(2) = './H_rps.dat'

!atomic number of element
izatom(1) = 6
izatom(2) = 1

!angular momentum of pseudopotential that will be treated as local
lloc_ps(1) = 1
```

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```

lloc_ps(2) = 0
!--- Caution -----!
! Indices must correspond to those in &atomic_coor. !
!-----!
/

```

*file\_pseudo(n)* specifies the filename of the pseudopotential file of the n-th element.

*izatom(n)* is the atomic number of the n-th element.

*lloc\_ps(n)* specifies which angular momentum component is chosen as the local potential for the n-th element.

```

&functional
!functional('PZ' is Perdew-Zunger LDA: Phys. Rev. B 23, 5048 (1981).)
xc = 'PZ'
/

```

*xc* specifies the exchange-correlation potential to be used in the calculation.

```

&rgrid
!spatial grid spacing(x,y,z)
dl(1:3) = 0.20d0, 0.20d0, 0.20d0
/

```

*dl(i)* specifies the spatial grid spacing in i-th direction.

```

&tgrid
!time step size and number of time grids(steps)
dt = 1.00d-3
nt = 5000
/

```

*dt* specifies the time step.

*nt* is the number of time steps for the time propagation.

```

&emfield
!envelope shape of the incident pulse('Ecos2': cos^2 type envelope for scalar_
↪potential)
ae_shapel = 'Ecos2'

!peak intensity(W/cm^2) of the incident pulse
I_wcm2_1 = 1.00d8

!duration of the incident pulse
twl = 6.00d0

```

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```

!mean photon energy(average frequency multiplied by the Planck constant) of the_
↪incident pulse
omega1 = 9.28d0

!polarization unit vector(real part) for the incident pulse(x,y,z)
epdir_rel(1:3) = 0.00d0, 0.00d0, 1.00d0

!carrier envelope phase of the incident pulse
!(phi_cep1 must be 0.25 + 0.5 * n(integer) when ae_shape1 = 'Ecos2')
phi_cep1 = 0.75d0
!--- Caution -----!
! Defenition of the incident pulse is wrritten in:                !
! https://www.sciencedirect.com/science/article/pii/S0010465518303412 !
!-----!
/

```

*ae\_shape1* specifies the envelope of the field.

*I\_wcm2\_1* specify the intensity of the pulse in unit of W/cm<sup>2</sup>.

*tw1* specifies the duration of the pulse.

*omega1* specifies the mean photon energy of the pulse.

*epdir\_rel(i)* specifies the i-th component of the real part of the polarization unit vector.

*phi\_cep1* specifies the carrier-envelope phase of the pulse.

```

&md
!ensemble
ensemble = 'NVE'

!set of initial velocities
yn_set_ini_velocity = 'y'

!setting temperature [K] for NVT ensemble, velocity scaling,
!and generating initial velocities
temperature0_ion_k = 300.0d0

!time step interval for updating pseudopotential
step_update_ps = 20
/

```

*ensemble* specifies the choice of the ensemble.

*yn\_set\_ini\_velocity* is a switch to prepare initial velocity for atoms.

*temperature0\_ion\_k* specifies the temperature that is used to generate initial velocity of ions.

*step\_update\_ps* specifies the time step interval to update projector for the nonlocal pseudopotential.

## Output files

After the calculation, following output files are created in the directory that you run the code,

file name	description
<i>C2H2_pulse.data</i>	dipole moment as functions of energy
<i>C2H2_rt.data</i>	components of change of dipole moment (electrons/plus definition) and total dipole moment (electrons/minus + ions/plus) as functions of time
<i>C2H2_rt_energy.data</i>	components of total energy and difference of total energy as functions of time
<i>C2H2_trj.xyz</i>	Trajectory of atoms(ions): Atomic coordinates, velocities, and forces are printed
<i>PS_C_KY_n.dat</i>	information on pseudopotential file for carbon atom
<i>PS_H_KY_n.dat</i>	information on pseudopotential file for hydrogen atom

You may download the above files (zipped file) from:

[https://salmon-tddft.jp/webmanual/v\\_2\\_0\\_1/exercise\\_zip\\_files/09\\_C2H2\\_md.zip](https://salmon-tddft.jp/webmanual/v_2_0_1/exercise_zip_files/09_C2H2_md.zip)

Explanations of the files are described below:

### C2H2\_pulse.data

Time-frequency Fourier transformation of the dipole moment.

```
# Fourier-transform spectra:
# energy: Frequency
# dm: Dipole moment
# 1:energy[eV] 2:Re(dm_x)[fs*Angstrom] 3:Re(dm_y)[fs*Angstrom] 4:Re(dm_
↪z)[fs*Angstrom] 5:Im(dm_x)[fs*Angstrom] 6:Im(dm_y)[fs*Angstrom] 7:Im(dm_
↪z)[fs*Angstrom] 8:|dm_x|^2[fs^2*Angstrom^2] 9:|dm_y|^2[fs^2*Angstrom^2] 10:|dm_z|^
↪2[fs^2*Angstrom^2]
```

### C2H2\_rt.data

Results of time evolution calculation for vector potential, electric field, and dipole moment.

```
# Real time calculation:
# Ac_ext: External vector potential field
# E_ext: External electric field
# Ac_tot: Total vector potential field
# E_tot: Total electric field
# ddm_e: Change of dipole moment (electrons/plus definition)
# dm: Total dipole moment (electrons/minus + ions/plus)
# 1:Time[fs] 2:Ac_ext_x[fs*V/Angstrom] 3:Ac_ext_y[fs*V/Angstrom] 4:Ac_ext_z[fs*V/
↪Angstrom] 5:E_ext_x[V/Angstrom] 6:E_ext_y[V/Angstrom] 7:E_ext_z[V/Angstrom] 8:Ac_
↪tot_x[fs*V/Angstrom] 9:Ac_tot_y[fs*V/Angstrom] 10:Ac_tot_z[fs*V/Angstrom] 11:E_tot_
↪x[V/Angstrom] 12:E_tot_y[V/Angstrom] 13:E_tot_z[V/Angstrom] 14:ddm_e_x[Angstrom]
↪15:ddm_e_y[Angstrom] 16:ddm_e_z[Angstrom] 17:dm_x[Angstrom] 18:dm_y[Angstrom] 19:dm_
↪z[Angstrom]
```

### C2H2\_rt\_energy.data

*Eall* and *Eall-Eall0* are total energy and electronic excitation energy, respectively.

```
# Real time calculation:
# Eall: Total energy
# Eall0: Initial energy
# Tion: Kinetic energy of ions
# Temperature_ion: Temperature of ions
# E_work: Work energy of ions(sum f*dr)
# 1:Time[fs] 2:Eall[eV] 3:Eall-Eall0[eV] # 4:Tion[eV] 5:Temperature_ion[K] 6:E_
↪work[eV]
```

### C2H2\_trj.xyz

Atomic coordinates [Angstrom], velocities [a.u.] and forces [a.u.] are printed along the time evolution in xyz format.

## 3.6 FDTD simulation(electromagnetic analysis)

### 3.6.1 Exercise-10: Pulsed electric field response of a metallic nanosphere in classical electromagnetism(FDTD simulation)

In this exercise, we learn the pulsed electric field response in the metallic nanosphere, solving the time-dependent Maxwell equations. As outputs of the calculation, the time response of the electromagnetic field is calculated. A pulsed electric field that has  $\cos^2$  envelope shape is applied. The parameters that characterize the pulsed field such as magnitude, frequency, polarization direction, and carrier envelope phase are specified in the input file.

#### Input files

To run the code, the input file *classicEM\_rt\_pulse.inp* that contains input keywords and their values for the pulsed electric field calculation is required. The shape file of the metallic nanosphere *shape.cube* is also required.

The shape file can be generated by program *FDTD\_make\_shape* in SALMON utilities: <https://salmon-tddft.jp/utilities.html>

'shape.inp' is an input file for 'FDTD\_make\_shape' to generate 'shape.cube'.

The input files are in samples

file name	description
<i>classicEM_rt_pulse.inp</i>	input file that contain input keywords and their values.
<i>shape.cube</i>	shape file for fdtd
<i>shape.inp</i>	input file for FDTD_make_shape

In the input file *classicEM\_rt\_pulse.inp*, input keywords are specified. Most of them are mandatory to execute the linear response calculation. This will help you to prepare the input file for other systems that you want to calculate. A complete list of the input keywords that can be used in the input file can be found in [List of input keywords](#).

```
#####
↪###!
! Exercise 10: Pulsed electric field response of a metallic nanosphere
↪!
!           in classical electromagnetism(FDTD simulation)
↪!
!-----!
↪---!
! * The detail of this excercise is expained in our manual(see chapter: 'Exercises').
↪!
```

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```

!   The manual can be obtained from: https://salmon-tddft.jp/documents.html
!
! * Input format consists of group of keywords like:
!
!   &group
!
!       input keyword = xxx
!
!   /
!
!   (see chapter: 'List of input keywords' in the manual)
!
!-----!
!---!
! * Conversion from unit_system = 'a.u.' to 'A_eV_fs':
!
!   Length: 1 [a.u.] = 0.52917721067      [Angstrom]
!
!   Energy: 1 [a.u.] = 27.21138505       [eV]
!
!   Time   : 1 [a.u.] = 0.02418884326505 [fs]
!
!-----!
!---!
! * The read-in file 'shape_file' in &maxwell category can be generated by program
!
!   'FDTD_make_shape' in SALMON utilities(https://salmon-tddft.jp/utilities.html).
!
!   'shape.inp' is an input file for 'FDTD_make_shape' to generate 'shape.cube'.
!
! * Results can be visualized by program 'FDTD_make_figani' in SALMON utilities.
!
!#####
!###!

&calculation
!type of theory
theory = 'maxwell'
/

```

*theory* specifies which theoretical method is used in the calculation.

```

&control
!name of directory where output files are contained
base_directory = 'result'
/

```

*base\_directory* specifies the directory name where output files are generated.

```
&units
!units used in input and output files
unit_system = 'A_eV_fs'
/
```

*unit\_system* specifies which unit system is used in the input and output files.

```
&system
!periodic boundary condition
yn_periodic = 'n'
/
```

*yn\_periodic* specifies whether or not periodic boundary condition is applied.

```
&emfield
!envelope shape of the incident pulse('Ecos2': cos^2 type envelope for scalar_
↪potential)
ae_shape1 = 'Ecos2'

!peak intensity(W/cm^2) of the incident pulse
I_wcm2_1 = 1.00d8

!duration of the incident pulse
tw1 = 4.60d0

!mean photon energy(average frequency multiplied by the Planck constant) of the_
↪incident pulse
omega1 = 5.49d0

!polarization unit vector(real part) for the incident pulse(x,y,z)
epdir_rel(1:3) = 0.00d0, 0.00d0, 1.00d0

!carrier envelope phase of the incident pulse
!(phi_cep1 must be 0.25 + 0.5 * n(integer) when ae_shape1 = 'Ecos2')
phi_cep1 = 0.75d0
!--- Caution -----!
! Defenition of the incident pulse is wrritten in:      !
! https://www.sciencedirect.com/science/article/pii/S0010465518303412 !
!-----!
/
```

*ae\_shape1* specifies the envelope of the field.

*I\_wcm2\_1* specify the intensity of the pulse in unit of W/cm<sup>2</sup>.

*tw1* specifies the duration of the pulse.

*omega1* specifies the mean photon energy of the pulse.

*epdir\_rel(i)* specifies the i-th component of the real part of the polarization unit vector.

*phi\_cep1* specifies the carrier-envelope phase of the pulse.

```

&maxwell
!box size and spacing of spatial grid(x,y,z)
al_em(1:3) = 120d0, 120d0, 120d0
dl_em(1:3) = 1.2d0, 1.2d0, 1.2d0

!time step size and number of time grids(steps)
dt_em = 2.30d-4
nt_em = 20000

!name of input shape file and number of media in the file
shape_file = './shape.cube'
media_num = 1

!*** MEDIA INFORMATION(START) *****!
!type of media(media ID)
media_type(1) = 'lorentz-drude'
!--- Au described by Lorentz-Drude model -----!
! The parameters are determined from:           !
! (https://www.osapublishing.org/ao/abstract.cfm?uri=ao-37-22-5271) !
!-----!

!number of poles and plasma frequency of media(media ID)
pole_num_ld(1) = 6
omega_p_ld(1) = 9.030d0

!oscillator strength, collision frequency,
!and oscillator frequency of media(media ID,pole ID)
f_ld(1,1:6) = 0.760d0, 0.024d0, 0.010d0, 0.071d0, 0.601d0, 4.384d0
gamma_ld(1,1:6) = 0.053d0, 0.241d0, 0.345d0, 0.870d0, 2.494d0, 2.214d0
omega_ld(1,1:6) = 0.000d0, 0.415d0, 0.830d0, 2.969d0, 4.304d0, 13.32d0
!*** MEDIA INFORMATION(END) *****!

!*** SOURCE INFORMATION(START) *****!
!type of method to generate the incident pulse
!('source': incident current source)
wave_input = 'source'

!location of source(x,y,z)
source_loc1(1:3) = -37.8d0, 0.0d0, 0.0d0

!propagation direction of the incident pulse(x,y,z)
ek_dir1(1:3) = 1.0d0, 0.0d0, 0.0d0
!*** SOURCE INFORMATION(END) *****!

!*** OBSERVATION INFORMATION(START) *****!
!number of observation points
obs_num_em = 1

!time step interval for sampling
obs_samp_em = 20

!location of observation point(observation ID,x,y,z)
obs_loc_em(1,1:3) = 0.0d0, 0.0d0, 0.0d0

!output flag for electromagnetic field distribution(observation ID)
yn_obs_plane_em(1) = 'n'
!--- Make of animation file -----!

```

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```

! When yn_obs_plane_em(1) = 'y', animation file can be made      !
! by program 'FDTD_make_figani' in SALMON utilities.              !
! The animation file visualizes electromagnetic field distributions !
! on the cross-section including the observation point            !
! whose location is determined by obs_loc_em.                    !
!-----!
!*** OBSERVATION INFORMATION (END) *****!
/

```

*al\_em(i)* specifies the lengths of three sides of the cuboid where the grid points are prepared.

*dl\_em(i)* specifies the grid spacings in three Cartesian directions.

*dt\_em* specifies the time step of the time evolution calculation.

*nt\_em* specifies the number of time steps in the calculation.

*shape\_file* specifies the filename of the shape file.

*media\_num* specifies the number of the types of media that is provided in the shape file.

*media\_type(n)* specifies the type of the n-th media.

*pole\_num\_ld(n)* and *omega\_p\_ld(n)* specify the number of poles and the plasmas frequency of the n-th media, respectively.

*f\_ld(n,m)*, *omega\_ld(n,m)*, *gamma\_ld(n,m)* specify the oscillator strength, oscillator frequency, and collision frequency of the m-th pole of the n-th media, respectively.

*wave\_input* specifies an electric current source that is used for the generation of the pulse.

*source\_loc1(i)* specifies the coordinate of the current source.

*ek\_dir1(i)* specifies the propagation direction of the pulse.

*obs\_num\_em* specifies the number of the observing point.

*obs\_samp\_em* specifies the sampling interval.

*obs\_loc\_em(n,i)* specifies the coordinate of n-th observing point.

*yn\_obs\_plane\_em(n)* is a switch to output the electromagnetic fields on the xy, yz, and xz planes that include the n-th observation point.

## Output files

After the calculation, following output files are created in the directory 'result',

file name	description
<i>obs0_info.data</i>	information to generate animation
<i>obs1_at_point_rt.data</i>	components of electric and magnetic fields as functions of time

You may download the above files (zipped file) from:

[https://salmon-tddft.jp/webmanual/v\\_2\\_0\\_1/exercise\\_zip\\_files/10\\_classicEM\\_rt.zip](https://salmon-tddft.jp/webmanual/v_2_0_1/exercise_zip_files/10_classicEM_rt.zip)

Explanations of the files are described below:

### **obs0\_info.data**

This file is used to generate animation files by using SALMON utilities: <https://salmon-tddft.jp/utilities.html>

### **obs1\_at\_point\_rt.data**

Results of time evolution calculation for electric and magnetic fields at observation point 1.

```
# Real time calculation:
# E: Electric field
# H: Magnetic field
# 1:Time[fs] 2:E_x[V/Angstrom] 3:E_y[V/Angstrom] 4:E_z[V/Angstrom] 5:H_x[A/Angstrom]
↪ 6:H_y[A/Angstrom] 7:H_z[A/Angstrom]
```

## LIST OF INPUT KEYWORDS

'[Trial]' : These options are not tested well

### 4.1 &calculation

#### 4.1.1 theory

character, default=""

Choice of a theory to be used in the calculation.

Options:

- dft / ground state calculation based on DFT
- dft\_md / ab initio MD simulations based on DFT (electronic ground state)
- tddft\_response / linear response TDDFT calculation in real time
- tddft\_pulse / simulations under pulsed electric field based on TDDFT
- single\_scale\_maxwell\_tddft / single-scale simulation coupling Maxwell and TDDFT
- multi\_scale\_maxwell\_tddft / multiscale simulation coupling Maxwell and TDDFT
- maxwell / electromagnetic analysis using finite difference time domain (FDTD) method
- dft\_k\_expand / convert checkpoint data of dft with k-points calculation to that of larger supercell system with gamma-point

#### 4.1.2 yn\_md

[Trial] character, default='n'

Available for theory='dft' (ground-state MD) and theory='tddft\_pulse' (Ehrenfest MD).

Switch for molecular dynamics calculation.

Options:

- 'y' / enable
- 'n' / disable

#### 4.1.3 yn\_opt

[Trial] character, default='n'

Available for theory='dft'.

Switch for geometry optimization.

Options:

'y' / enable  
'n' / disable

## 4.2 &control

### 4.2.1 sysname

character, default='default'

Available for all options of `theory`.  
A prefix of output files.

### 4.2.2 base\_directory

character, default='./'

Available for all options of `theory`.  
Name of a directory where major output files are stored.

### 4.2.3 yn\_restart

character, default='n'

Available for the DFT/TDDFT based options of `theory`.  
Whether to continue previous calculation (restart) or start a new calculation.  
Options:  
'y' / enable (restart)  
'n' / disable (new calculation)

### 4.2.4 directory\_read\_data

character, default='restart'

Available for `yn_restart='y'`.  
Directory name to read data that are required in the present calculation (restart) and were generated in previous calculations. For TDDFT based options, it specifies the name of the directory containing ground state results that were stored in 'data\_for\_restart'. When restarting from a checkpoint, it specifies the name of the directory that contains the checkpoint data.

### 4.2.5 yn\_self\_checkpoint

character, default='n'

Available for the DFT/TDDFT based options of `theory`.  
With this option, each process writes/reads the restart (and checkpoint) data independently (self data format) so that the restart cost is reduced for large systems. Note that the number of processes and their assignments must be unchanged in restarting. The data is written out into 'checkpoint\_gs\_XXXXXX/' (DFT) or 'checkpoint\_rt\_XXXXXX/' (TDDFT).  
Options:

'y' / enable  
'n' / disable

### 4.2.6 checkpoint\_interval

integer, default=-1

Available for the DFT/TDDFT based options of `theory`.

Interval of time steps (iteration steps) to write down the checkpoint data during the time-propagation (SCF iteration). Checkpoint data will not be written if a negative value is set.

### 4.2.7 yn\_reset\_step\_restart

character, default='n'

Available for `yn_restart='y'` in the DFT/TDDFT based options of `theory`.

With this option, the counter of the SCF iteration step (for DFT) or the counter of the time propagation step (for TDDFT) is reset to 0 at the restart. In the SCF iteration, the density data in the previous SCF iteration step are abandoned.

### 4.2.8 read\_gs\_restart\_data

character, default='all'

Available for `yn_restart='y'` with `theory='dft'`.

Specify which data are read in the restart. Specified data that are generated in the previous calculation and are contained in the restart (or checkpoint) directory are used in restarting the SCF iteration of DFT. The default option 'all' indicates the complete restart. In other options, a part of restart data are used (other data are prepared in the same way as in the initial SCF step).

Options:

`all` / all of restart data are read

`all:single` / same as `all` option but the data is read in the single file format even though the self data format is specified with `yn_self_checkpoint='y'` (i.e., the restart data is read in the single file format while written out in the self format)

`rho_inout` / only electron densities including those of previous iteration steps are read (from `rho_inout.bin` file)

`rho_inout:single` / same as `rho_inout` option but the data is read in the single file format even though the self data format is specified with `yn_self_checkpoint='y'`

`rho` / only the latest electron density is read (from user-made data)

`wfn` / only orbital wavefunctions are read

### 4.2.9 write\_gs\_restart\_data

character, default='all'

Available for `theory='dft'`.

Options

`all` / all of restart data are written out

`rho_inout` / only electron densities including those of previous iteration steps are written out

`wfn` / only orbital wavefunctions are written out

`checkpoint_only` / the restart data are outputted only in the self data format (separated data for each process) at the last step into '`checkpoint_gs_XXXXXX`' directory (`yn_self_checkpoint='y'` is required) without generating the restart data into '`data_for_restart`' directory in the single file format.

Output data files are written out in the restart (or checkpoint) directory.

The default option '`all`' gives the complete set of restart data.

#### 4.2.10 `time_shutdown`

[Trial] `real(8)`, default=-1d0

Available for the DFT/TDDFT based options of `theory`.

Timer for automatic shutdown. The unit is second.

If a negative time is set, the automatic shutdown will not be performed.

#### 4.2.11 `method_wf_distributor`

character, default='single'

Available for the DFT/TDDFT based options of `theory`.

A method to save/load orbital wavefunctions.

Options

`single`: all orbital wavefunctions are saved(loaded) to(from) a single file.

`slice`: each orbital wavefunction is saved(loaded) to(from) a file. This choice reduces the I/O costs, and increase the flexibility in handling files for large systems.

#### 4.2.12 `nblock_wf_distribute`

integer, default='16'

Available for `method_wf_distributor='slice'`.

In the 'slice' mode, `nblock_wf_distribute` files are saved in one directory.

### 4.3 &units

#### 4.3.1 `unit_system`

character, default='au'

Unit system to be used in input variables and some of output files.

If `unit_system = 'A_eV_fs'` is chosen, Angstrom for length, eV for energy, and fs for time are adopted.

For isolated systems specified by `yn_periodic = 'n'` in `&system`, a unit of 1/eV is used for the output files of DOS and PDOS if `unit_system = 'A_eV_fs'` is specified, while atomic unit is used if not. For other output files, the Angstrom/eV/fs units are used irrespective of the input keyword. For periodic systems specified by `yn_periodic = 'n'` in `&system`, the unit system specified by this input keyword is used for most output files. To confirm the unit, see the first few lines of output files.

Options:

'au' or 'a.u.' / atomic unit system

'A\_eV\_fs' / Angstrom-eV-fs unit system

## 4.4 &parallel

### 4.4.1 nproc\_k

### 4.4.2 nproc\_ob

### 4.4.3 nproc\_rgrid(3)

integer, default=0

Options:

nproc\_k/ Number of MPI parallelization for k-points of electron orbitals.

nproc\_ob/ Number of MPI parallelization for orbital index of electron orbitals.

nproc\_rgrid(3) '/ Number of MPI parallelization for each direction of real-space grid that are used for electron orbitals and density.

Defaults are 0 for nproc\_k/nproc\_ob and (0, 0, 0) for nproc\_rgrid. In the default choice, MPI assignment is achieved automatically. Users can specify nproc\_k, nproc\_ob, and nproc\_rgrid manually. In that case, there are several constraints that should be fulfilled:

nproc\_k must be set to 1 for &system/yn\_periodic='n'.

nproc\_k and nproc\_ob must be set to 1 for theory='maxwell'.

$\text{nproc\_k} * \text{nproc\_ob} * \text{nproc\_rgrid}(1) * \text{nproc\_rgrid}(2) * \text{nproc\_rgrid}(3) =$   
total number of processes.

### 4.4.4 yn\_ffte

character, default='n'

Available for &system/yn\_periodic='y'

For periodic systems, SALMON uses Fourier transformation to solve a poisson equation.

This switch selects if FFTE library is used or not. If FFTE is not used, the Fourier transformation in a simple algorithm is carried out.

Options

'y' / enable

'n' / disable

To enable it, following relations must be satisfied.

$\text{mod}(\text{num\_rgrid}(1), \text{nproc\_rgrid}(2)) == 0$

$\text{mod}(\text{num\_rgrid}(2), \text{nproc\_rgrid}(2)) == 0$

$\text{mod}(\text{num\_rgrid}(2), \text{nproc\_rgrid}(3)) == 0$

$\text{mod}(\text{num\_rgrid}(3), \text{nproc\_rgrid}(3)) == 0$

### 4.4.5 yn\_scalapack

character, default='n'

Available for `&calculation/theory='dft'` or `'dft_md'`

To calculate large systems, an eigenvalue problem in the subspace diagonalization becomes a bottle-neck in the ground state calculation. In SALMON, ScaLAPACK library can be used to solve the eigenvalue problem.

To enable it, it is necessary to link ScaLAPACK library when you build SALMON.

Options:

`'y'` / enable

`'n'` / disable

#### 4.4.6 `yn_eigenexa`

character, default=`'n'`

Available for `&calculation/theory='dft'` or `'dft_md'`

SALMON can use RIKEN R-CCS EigenExa library to solve eigenvalue problem in subspace diagonalization. It is more efficient than ScaLAPACK to diagonalize matrices of large dimension. To enable it, it is necessary to link both ScaLAPACK and EigenExa libraries when you build SALMON.

Options:

`'y'` / enable

`'n'` / disable

#### 4.4.7 `yn_diagonalization_red_mem`

character, Default=`'n'`

Available for `&parallel/yn_scalapack='y'` or `&parallel/yn_eigenexa='y'`

This option reduces memory consumption in using ScaLAPACK/EigenExa libraries.

Options:

`'y'` / enable

`'n'` / disable

#### 4.4.8 `process_allocation`

character, default=`'grid_sequential'`

This controls the order of process allocation.

Options:

`'grid_sequential'` / real-space grid major ordering.

`'orbital_sequential'` / orbital-space major ordering.

Suggestion:

`&calculation/theory='dft'` or `'dft_md' / 'orbital_sequential'`

`&calculation/theory='tddft*' or '*maxwell_tddft' /`

`'grid_sequential'`



## 4.5 &system

### 4.5.1 yn\_periodic

character, default='n'

Available for all options of `theory`.

Specify boundary condition for electron orbitals.

Options:

'y' / periodic systems (crystalline solids)

'n' / isolated systems (molecules and nano-particles)

### 4.5.2 spin

character, default='unpolarized'

Available for the DFT/TDDFT based options of `theory`.

It specifies the spin state of the system, spin-unpolarized (closed shell) and spin-polarized (open shell).

Options

'unpolarized' / spin-unpolarized systems (default)

'polarized' / spin-polarized systems

### 4.5.3 al(3)

real(8), default=0d0

Available for the DFT/TDDFT based options of `theory`.

Spatial box size or lattice constants for cuboid cell (x, y, z).

For nonorthogonal cell, see `al_vec1(3)`, `al_vec2(3)`, `al_vec3(3)`.

### 4.5.4 al\_vec1(3)

### 4.5.5 al\_vec2(3)

### 4.5.6 al\_vec3(3)

real(8), default=0d0

Available for `yn_periodic = 'y'` in the DFT/TDDFT based options of `theory`.

Primitive lattice vectors for nonorthogonal cell. For cuboid cell, see `al(3)`.

### 4.5.7 nstate

integer, default=0

Available for the DFT/TDDFT based options of `theory`.

of orbitals/bands to be calculated. In the time evolution calculation of dielectrics, only occupied orbitals are evolved even when more `nstate` is specified.

### 4.5.8 nelec

integer, default=0

Available for the DFT/TDDFT based options of `theory`.

Number of valence electrons in the system.

### 4.5.9 nelec\_spin(2)

integer, Default=0

Available for the DFT/TDDFT based options of `theory`.

Number of up/down-spin electrons are specified by `nelec_spin(1)/nelec_spin(2)`.

This option is incompatible with `nelec`. (If `nelec_spin` is specified, `nelec` is ignored.)

### 4.5.10 temperature

real(8), default=-1d0

Available for DFT-based options of `theory`.

It specifies the temperature for electrons. The value must be given using the unit of energy as specified in `&units/unit_system`.

The kelvin unit can also be used by the keyword `temperature_k` instead of `temperature` (see next).

Occupation numbers are updated in every SCF step in the following way.

`temperature < 0` / the occupation numbers are fixed by `nelec` (appropriate for systems with energy gap).

`temperature = 0` / redistribution of the occupation numbers by the step function (metallic system at zero temperature).

`temperature > 0` / redistribution of the occupation numbers by the Fermi-Dirac distribution function.

### 4.5.11 temperature\_k

[Trial] real(8), default=-1d0

Available for DFT-based options of `theory`.

The same as `temperature` but kelvin is used as the unit.

### 4.5.12 nelem

integer, default=0

Available for the DFT/TDDFT based options of `theory`.

Number of atomic elements in the system.

### 4.5.13 natom

integer, default=0

Available for the DFT/TDDFT based options of `theory`.

Number of atoms in the system.

#### 4.5.14 file\_atom\_red\_coor

[Trial] character, default='none'

Available for the DFT/TDDFT based options of `theory`.

Name of the file that contains atomic positions given in reduced coordinates. This option is incompatible with `&system/file_atom_coor`, `&atomic_coor`, and `&atomic_red_coor`.

#### 4.5.15 file\_atom\_coor

[Trial] character, default='none'

Available for the DFT/TDDFT based options of `theory`.

Name of the file that contains atomic Cartesian coordinates (The unit is specified by `&units/unit_system`). This option is incompatible with `&system/file_atom_coor`, `&atomic_coor`, and `&atomic_red_coor`.

#### 4.5.16 yn\_spinorbit

[Trial] character, default='n'

Options

'y' / enable

'n' / disable

#### 4.5.17 yn\_symmetry

[Trial] character, default='n'

Available for orthogonal cell system with the DFT/TDDFT based options of `theory`.

Symmetry option. Pre-generated input file, "sym.dat", is necessary. (details are not explained in the current manual)

Options

(e.g.) 'yy' / symmetry option is applied for the x and y direction (under applied electric field in the z-direction)

'n' / disable

#### 4.5.18 absorbing\_boundary

[Trial] character, default='none'

Available for the TDDFT based option of `theory` with orthogonal unit cell.

Absorbing boundary condition for electrons. (T. Nakatsukasa et al., J. Chem. Phys., 114, 2550 (2001))

Options:

'none' / disable (default)

'z' / absorbing boundary region is set in z direction for `'yn_periodic' = 'y'`

### 4.5.19 imaginary\_potential\_w0

real(8), default='0d0'

Available when absorbing\_boundary options is not 'none'.  
Strength of the absorbing (imaginary) potential.

### 4.5.20 imaginary\_potential\_dr

real(8), default='0d0'

Available when absorbing\_boundary options is not 'none'.  
Thickness of the absorbing (imaginary) potential. For absorbing\_boundary='z', the absorbing region is  $0 < z < \text{imaginary\_potential\_dr}$  and  $\text{al}(3) - \text{imaginary\_potential\_dr} < z < \text{al}(3)$

## 4.6 &atomic\_red\_coor

Atomic coordinates in periodic systems ('yn\_periodoc = 'y') are specified in reduced coordinates using the following format:

```
'Si' 0.00 0.00 0.00 1
'Si' 0.25 0.25 0.25 1
...
```

Here, the information of atoms is ordered in row, the first row for the first atom, etc. The number of rows must be equal to &system/natom. Atomic species are written in the first column although they are not used in the calculation. The second, third and fourth columns are reduced coordinates for the first, second and third directions, respectively. The fifth column is a serial number of the atom species, which is defined in &pseudo.

This option is incompatible with &system/file\_atom\_red\_coor, &system/file\_atom\_coor, and &atomic\_coor.

## 4.7 &atomic\_coor

Atomic coordinates are specified in the same way as atomic\_red\_coor but with length dimension. The unit chosen by &units/unit\_length is applied.

This option is incompatible with &system/file\_atom\_red\_coor, &system/file\_atom\_coor, and &atomic\_red\_coor.

## 4.8 &pseudo

### 4.8.1 izatom(:)

integer, default=-1

Available for the DFT/TDDFT based options of theory.  
Atomic number of the element. The size of array is equal to &system/nelem.

### 4.8.2 file\_pseudo(:)

character, default='none'

Available for the DFT/TDDFT based options of `theory`.

File name of the pseudopotential file. The size of array is equal to `&system/nelem`.

### 4.8.3 lmax\_ps(:)

integer, default=-1

Available for the DFT/TDDFT based options of `theory`.

Maximum angular momentum of pseudopotential projectors.

If not given, values specified in the pseudopotential file will be used. The size of array is equal to `&system/nelem`.

### 4.8.4 lloc\_ps(:)

integer, default=-1

Available for the DFT/TDDFT based options of `theory`.

Angular momentum of the pseudopotential that will be treated as local. The size of array is equal to `&system/nelem`.

### 4.8.5 yn\_psmask(:)

[Trial] character, default='n'

Available for the DFT/TDDFT based options of `theory`.

Fourier filtering for pseudopotentials. The size of array is equal to `&system/nelem`.

Options:

'y' / enable

'n' / disable

### 4.8.6 alpha\_mask(:)

[Trial] real(8), default=0.8d0

Available for the DFT/TDDFT based options of `theory`.

Parameter for the Fourier filtering of the pseudopotential. The size of array is equal to `&system/nelem`.

### 4.8.7 gamma\_mask(:)

[Trial] real(8), default=1.8d0

Available for the DFT/TDDFT based options of `theory`.

Parameter for the Fourier filtering of the pseudopotential. The size of array is equal to `&system/nelem`.

### 4.8.8 eta\_mask(:)

[Trial] real(8), default=15.0d0

Available for the DFT/TDDFT based options of `theory`.

Parameter for the Fourier filtering of the pseudopotential. The size of array is equal to `&system/nelem`.

## 4.9 &functional

### 4.9.1 xc

character, default='none'

Available for the DFT/TDDFT based options of `theory`.

Exchange-correlation functional to be used.

In the present version, functionals 'PZ', 'PZM' and 'TBmBJ' are available for both `yn_periodic = 'y'` and `'n'` calculations in the adiabatic approximation.

Options:

'PZ': Perdew-Zunger LDA :Phys. Rev. B 23, 5048 (1981).

'PZM': Perdew-Zunger LDA with modification to improve smooth connection between high density form and low density one. :J. P. Perdew and Alex Zunger, Phys. Rev. B 23, 5048 (1981).

'TBmBJ': Tran-Blaha meta-GGA exchange with Perdew-Wang correlation. :Fabien Tran and Peter Blaha, Phys. Rev. Lett. 102, 226401 (2008). John P. Perdew and Yue Wang, Phys. Rev. B 45, 13244 (1992). This potential is known to provide a reasonable description for the bandgap of various insulators. For this choice, the additional mixing parameter 'cval' may be specified. See below.

### 4.9.2 cval

real(8), default=-1d0

Available for `xc='TBmBJ'`.

Mixing parameter in Tran-Blaha meta-GGA exchange potential. If `cval` is set to a minus value, the mixing-parameter is evaluated by the formula in the original paper [Phys. Rev. Lett. 102, 226401 (2008)],  $\langle |\nabla \rho(\mathbf{r}; t)| / \rho(\mathbf{r}; t) \rangle$ . However, note that the value may be different from that in all electron calculations.

### 4.9.3 cname

### 4.9.4 xname

character, default='none'

Available for `theory='XXX'`.

XXX

### 4.9.5 alibxc

### 4.9.6 alibx

### 4.9.7 alibc

character, default='none'

Available for the DFT/TDDFT based options of `theory`.

Since version 1.1.0, exchange-correlation functionals in Libxc library

(<http://www.tddft.org/programs/libxc/>) have been usable in SALMON. At present, usable functionals are limited to LDA and GGA. For periodic systems, meta-GGA functionals are usable as well. To specify the exchange-correlation potentials of Libxc library, there are two ways. If the exchange and correlation potentials are given separately, you need to specify both `alibx` and `alibc` separately. If the exchange and correlation potentials are given as a combined set, you need to specify `alibxc`. We show below an example:

```
&functional
    alibx = 'LDA_X'
    alibc = 'LDA_C_PZ'
```

Note that, the hybrid functionals (hybrid gga/mgga) are not supported.

To use libxc libraries, `--enable-libxc` option must be added in executing configure. The available option of the exchange-correlation functionals are listed in the LibXC website. [See

<http://www.tddft.org/programs/libxc/functionals/>]

## 4.10 &rgrid

### 4.10.1 dl(3)

real(8), default=0d0

Available for the DFT/TDDFT based options of `theory`.

Spacing of real-space grids.

This cannot be used together with `&rgrid/num_rgrid`.

### 4.10.2 num\_rgrid(3)

integer, default=0

Available for the DFT/TDDFT based options of `theory`.

Number of real-space grids for each direction.

This cannot be used together with `&rgrid/dl`.

## 4.11 &kgrid

### 4.11.1 num\_kgrid(3)

integer, default=1

Available for `yn_periodic='y'` in the DFT/TDDFT based options of `theory`.  
Number of k-points (grid points of k-vector) for each direction discretizing the Brillouin zone.

### 4.11.2 `file_kw`

character, default='none'

Available for `yn_periodic='y'` in the DFT/TDDFT based options of `theory`.  
File name for a file that includes user specified k-points. This file will be read if `num_kgrid` is equal to 0 or negative values. The file should be described in the following format :

```
8 #(number of k-points)
1 -0.50 -0.50 -0.50 0.1250 #(id, kx, ky, kz, weight)
2 -0.50 -0.50 0.00 0.1250
3 -0.50 0.00 -0.50 0.1250
4 -0.50 0.00 0.00 0.1250
5 0.00 -0.50 -0.50 0.1250
6 0.00 -0.50 0.00 0.1250
7 0.00 0.00 -0.50 0.1250
8 0.00 0.00 0.00 0.1250
```

## 4.12 `&tgrid`

### 4.12.1 `nt`

integer, Default=0

Available for 'dft\_md' and TDDFT-based options of `theory`.  
Number of total time steps for real-time propagation.

### 4.12.2 `dt`

real(8), Default=0d0

Available for 'dft\_md' and TDDFT-based options of `theory`.  
Time step size.

### 4.12.3 `gram_schmidt_interval`

integer, default=-1

Available for TDDFT-based options of `theory`.  
Interval of a time step for the Gram-Schmidt orthonormalization of the orbitals during time evolution calculations. If this is set to a negative value, no Gram-Schmidt orthogonalization will be achieved. If this is set to zero, the Gram-Schmidt orthogonalization is carried out once at the initial step only. Usually this Gram-Schmidt orthogonalization is not necessary and should not be used.



## 4.13 &propagation

### 4.13.1 n\_hamil

**integer, default=4**

Available for TDDFT-based options of `theory`.

Order of the Taylor expansion adopted for the propagation operator.

### 4.13.2 propagator

**character, default=middlepoint**

Available for TDDFT-based options of `theory`.

Choice of the propagator in the time evolution calculation.

Options:

`middlepoint` / Hamiltonian at midpoint of two-times is used in the propagation if

`yn_predictor_corrector = 'y'`. Hamiltonian at the time  $t$  is used if

`yn_predictor_corrector = 'n'`.

`aetrs` / time-reversal symmetry propagator. [M.A.L. Marques, A. Castro, G.F. Bertsch, and A.

Rubio, Comput. Phys. Commun., 151 60 (2003)].

### 4.13.3 yn\_predictor\_corrector

**character, default='n'**

Available for TDDFT-based options of `theory`.

Switch of the predictor-corrector method of TDDFT.

For meta-GGA functionals (`xc='tbmbj'`), the predictor corrector is automatically used even with

`yn_predictor_corrector='n'`.

Options:

`'y'` / enable

`'n'` / disable

### 4.13.4 yn\_fix\_func

**[Currently not available] character, default='n'**

Available for `'dft_md'` and TDDFT-based options of `theory`.

Switch not to update the Hamiltonian during the time evolution, i.e., ground state Hamiltonian is used during the propagation.

Options:

`'y'` / enable

`'n'` / disable

## 4.14 &scf

### 4.14.1 method\_init\_wf

character, default='gauss'

Available for 'dft' and 'dft\_md' options of `theory`.

The generation method of the initial orbitals at the beginning of the SCF iteration in DFT calculations. For a stable calculation of very large systems, multiple gaussian functions are preferred for a stable calculation.

Options:

- `gauss` / single gauss function per orbital centered at a position determined by random numbers
- `gauss2` / two gauss functions per orbital centered at positions determined by random numbers
- `gauss3` / three gauss functions per orbital centered at positions determined by random numbers
- `gauss4` / four gauss functions per orbital centered at positions determined by random numbers
- `gauss5` / five gauss functions per orbital centered at positions determined by random numbers
- `gauss10` / ten gauss functions per orbital centered at positions determined by random numbers
- `random` / a random number is assigned at each real-space grid point of orbitals

### 4.14.2 method\_init\_density

[Trial] character, default='wf'

Available for 'dft' and 'dft\_md' options of `theory`.

Specifying how to generate the initial density to start the SCF iteration in the DFT calculation.

Supported for limited formats of pseudopotentials ('KY' and 'UPF').

Options:

- `wf` / generate from the initial wavefunctions (cf. `method_init_wf`).
- `pp` / generate from a superposition of the pseudo-atom densities.

### 4.14.3 iseed\_number\_change

integer, default=0

Available for 'dft' and 'dft\_md' options of `theory`.

Change a seed of random numbers that are used to generate initial orbitals. The value specified by this parameter is added to the seed.

### 4.14.4 nscf

integer, Default=300

Available for 'dft' and 'dft\_md' options of `theory`.

Number of maximum SCF iterations in the DFT calculation.

### 4.14.5 method\_min

character, Default='cg'

Available for 'dft' and 'dft\_md' options of `theory`.

Method for updating orbitals in the SCF iteration. At present only conjugate gradient method is implemented.

Options:

`cg` / Conjugate-Gradient(CG) method

#### 4.14.6 `ncg`

integer, default=4

Available for 'dft' and 'dft\_md' options of `theory`.

Number of iterations of conjugate-gradient method in the SCF iteration.

#### 4.14.7 `ncg_init`

integer, default=4

Available for 'dft' and 'dft\_md' options of `theory`.

Number of iterations of conjugate-gradient method for the first SCF step.

#### 4.14.8 `method_mixing`

character, default='broyden'

Available for 'dft' and 'dft\_md' options of `theory`.

Method to update density/potential in the scf iteration.

Options:

`simple` / Simple mixing method

`broyden` / modified Broyden method

`pulay` / Pulay method

#### 4.14.9 `mixrate`

real(8), default=0.5d0

Available for `method_mixing='simple'` in 'dft' and 'dft\_md' options of `theory`.

Mixing ratio for simple mixing.

#### 4.14.10 `nmemory_mb`

integer, default=8

Available for `method_mixing='broyden'` in 'dft' and 'dft\_md' options of `theory`.

Number of previous densities to be stored in the SCF iteration using the modified Broyden method. This must be less than 21.

#### 4.14.11 `alpha_mb`

real(8), default=0.75d0

Available for `method_mixing='broyden'` in 'dft' and 'dft\_md' options of `theory`.

A parameter of the modified Broyden method.

#### **4.14.12 nmemory\_p**

integer, default=4

Available for `method_mixing='pulay'` in `'dft'` and `'dft_md'` options of `theory`.  
Number of previous densities to be stored in the SCF iteration using the Pulay method.

#### **4.14.13 beta\_p**

real(8), default=0.75d0

Available for `method_mixing='pulay'` in `'dft'` and `'dft_md'` options of `theory`.  
A parameter of the mixing rate of the Pulay method.

#### **4.14.14 yn\_auto\_mixing**

character, default='n'

Available for `'dft'` and `'dft_md'` options of `theory`.  
Switch to change the mixing rate automatically (i.e. automatic adjustments of `mixrate/alpha_mb/beta_p`)  
Options:  
    'y' / enable  
    'n' / disable

#### **4.14.15 update\_mixing\_ratio**

real(8), default=3.0d0

Available for `yn_auto_mixing='y'` in `'dft'` and `'dft_md'` options of `theory`.  
Threshold for the change of the mixing rate in `yn_auto_mixing='y'` option. The mixing-rate is reduced to half when the ratio of the density differences between the current and previous iteration steps is larger than `update_mixing_ratio`.

#### **4.14.16 yn\_subspace\_diagonalization**

character, default='y'

Available for `'dft'` and `'dft_md'` options of `theory`.  
Switch for the subspace diagonalization during SCF iterations.  
Options:  
    'y' / enable  
    'n' / disable

#### 4.14.17 convergence

character, default='rho\_dne'

Available for 'dft' and 'dft\_md' options of `theory`.

Specify a quantity that is used for convergence check of the SCF iteration.

Options:

- 'rho\_dne' / Convergence is checked by  $\sum_{\text{ix}} |\rho_{\text{iter}}(\text{ix}) - \rho_{\text{iter-1}}(\text{ix})| dx / N$ . N is &system/nelec.
- 'norm\_rho' / Convergence is checked by the square of the norm of the density difference,  $\|\rho_{\text{iter}}(\text{ix}) - \rho_{\text{iter-1}}(\text{ix})\|^2 = \sum_{\text{ix}} |\rho_{\text{iter}}(\text{ix}) - \rho_{\text{iter-1}}(\text{ix})|^2$ .
- 'norm\_rho\_dng' / Convergence is checked by  $\|\rho_{\text{iter}}(\text{ix}) - \rho_{\text{iter-1}}(\text{ix})\|^2 / (\text{number of grids})$ . "dng" means "divided by number of grids".
- 'norm\_pot' / Convergence is checked by  $\|V_{\text{local\_iter}}(\text{ix}) - V_{\text{local\_iter-1}}(\text{ix})\|^2$ , where  $V_{\text{local}}$  is  $V_{\text{h}} + V_{\text{xc}} + V_{\text{ps\_local}}$ .
- 'pot\_dng' / Convergence is checked by  $\|V_{\text{local\_iter}}(\text{ix}) - V_{\text{local\_iter-1}}(\text{ix})\|^2 / (\text{number of grids})$ .

#### 4.14.18 threshold

real(8), default=1d-17 [a.u.] (for convergence='rho\_dne') and -1 (for other options of convergence))

Available for 'dft' and 'dft\_md' options of `theory`.

Threshold of convergence that is specified by `convergence` keyword.

#### 4.14.19 nscf\_init\_redistribution

integer, default=10

Available for 'dft' and 'dft\_md' options of `theory`.

Number of initial iterations during which a redistribution of the occupation number is suppressed in the finite temperature calculation.

#### 4.14.20 nscf\_init\_no\_diagonal

integer, default=10

Available for &scf/yn\_subspace\_diagonalization='y' in 'dft' option of `theory`.

Number of initial iterations during which the subspace diagonalization will not be carried out.

#### 4.14.21 nscf\_init\_mix\_zero

integer, default=-1

Available for 'dft' option of `theory`.

The density will not be mixed (i.e. fixed) during the given number of the SCF iteration, that is, orbitals are optimized without updating the density.

### 4.14.22 conv\_gap\_mix\_zero

real(8), default=99999d0

Available if `nscf_init_mix_zero` is positive value in the 'dft' option of `theory`.

Specify a condition to quit the fixed density iteration forced by `step_initial_mix_zero` option.

Mixing of the density will start after the band-gap energy exceeds this parameter for consecutive five SCF iteration steps.

## 4.15 &emfield

### 4.15.1 trans\_longi

character, default='tr'

Available for `yn_periodic='y'` in 'maxwell' and TDDFT based options of `theory`.

Specify the treatment of the polarization in the time evolution calculation.

Options:

'tr' / Transverse

'lo' / longitudinal

'2d' / 2D maxwell-TDDFT method (for more details, see `film_thickness` of &maxwell)

### 4.15.2 ae\_shape1

### 4.15.3 ae\_shape2

character, Default='none'

Available for 'maxwell' and TDDFT based options of `theory`.

Envelope shape of the first/second pulse. 'Acos2' indicates a cosine square envelope for vector potential, and 'Ecos2' a cosine square envelope for electric field.

Options:

'impulse' / A weak impulsive field is applied at  $t = 0$ . This will be used to explore linear response properties. The magnitude of the impulse can be specified by `e_impulse`.

'Acos2' / Envelope of  $\cos^2$  for a vector potential.

'Acos3' / Envelope of  $\cos^3$  for a vector potential.

'Acos4' / Envelope of  $\cos^4$  for a vector potential.

'Acos6' / Envelope of  $\cos^6$  for a vector potential.

'Acos8' / Envelope of  $\cos^8$  for a vector potential.

'Ecos2' / Envelope of  $\cos^2$  for an electric field.

'Asin2cos' [Trial] / Envelope of  $\sin^2$  with cosine type oscillation for a vector potential.

'Asin2\_cw' [Trial] / Envelope of  $\sin^2$  at the beginning and continuous wave after that for a vector potential (for 'ae\_shape1' only).

'input' [Trial] / read the vector potential as a numerical table with `file_input1` option (for 'ae\_shape1' only).

'none' / no incident field is applied.

If 'Ecos2' is adopted, 'phi\_cep1' must be chosen either 0.75 or 0.25, since otherwise the time integral of the electric field (vector potential at the end of the pulse) does not vanish. There is no such restriction for 'Acos2' pulses.

For `yn_periodic='n'`, available choices are limited to 'impulse', 'Acos2', and 'Ecos2'.

#### 4.15.4 file\_input1

character, default=""

Available if `ae_shape1='input'` is specified and `theory='tddft_pulse'`.

Name of an input file that contains user-defined vector potential. The file must be a numerical table separated by blank, having four columns; the first column is time and second to fourth columns are  $A_x/c$ ,  $A_y/c$ ,  $A_z/c$ , respectively. All the quantities are written using the units specified by `unit_system`. '#' and '!' may be used for a comment line.

Note that a linear interpolation will be applied when the time step differs from that used in the calculation.

#### 4.15.5 e\_impulse

real(8), Default=1d-2 a.u.

Available for 'maxwell' and TDDFT based options of `theory`.

Magnitude of the impulse in the impulsive perturbation. This variable has the dimension of momentum,  $\text{energy} \cdot \text{time} / \text{length}$ .

#### 4.15.6 E\_amplitude1

#### 4.15.7 E\_amplitude2

real(8), default=0d0

Available for 'maxwell' and TDDFT based options of `theory`.

Maximum amplitude of electric field for the first/second pulse. This variable has the dimension of electric field,  $\text{energy} / (\text{length} \cdot \text{charge})$ . This cannot be set with `&emfield/I_wcm2_1` (`I_wcm2_2`) simultaneously.

#### 4.15.8 I\_wcm2\_1

#### 4.15.9 I\_wcm2\_2

real(8), default=-1d0

Available for 'maxwell' and TDDFT based options of `theory`.

Maximum intensity ( $\text{W}/\text{cm}^2$ ) of the first/second pulse. This variable cannot be set with `&emfield/E_amplitude1` (`E_amplitude2`) simultaneously. For this quantity, a unit of  $\text{W}/\text{cm}^2$  is adopted irrespective of `&units\unit_system`.

#### 4.15.10 tw1

#### 4.15.11 tw2

real(8), default=0d0

Available for 'maxwell' and TDDFT based options of `theory`.

Duration of the first/second pulse (edge-to-edge time length).

Note that this is not the FWHM duration.

#### 4.15.12 omega1

#### 4.15.13 omega2

real(8), default=0d0

Available for 'maxwell' and TDDFT based options of `theory`.

Mean photon energy (average frequency multiplied by the Planck constant) of the first/second pulse.

#### 4.15.14 ekdir\_re1(3)

#### 4.15.15 ekdir\_re2(3)

real(8), default=1d0, 0d0, 0d0

Available for 'maxwell' and TDDFT based options of `theory`.

Real part of the polarization unit vector for the first/second pulse.

#### 4.15.16 ekdir\_im1(3)

#### 4.15.17 ekdir\_im2(3)

real(8), default=0d0

Available for 'maxwell' and TDDFT based options of `theory`.

Imaginary part of the polarization unit vector for the first/second pulse. Using both real 'ekdir\_re1' and imaginary 'ekdir\_im1' parts of the polarization vector, circularly and general ellipsoidary polarized pulses may be described.

#### 4.15.18 phi\_cep1

#### 4.15.19 phi\_cep2

real(8), default=0d0

Available for 'maxwell' and TDDFT based options of `theory`.

Carrier envelope phase of the first/second pulse. It specifies the CEP in unit of  $2\pi$ .



### 4.15.20 `t1_t2`

`real(8), default=0d0`

Available for 'maxwell' and TDDFT based options of `theory`.  
Time-delay between the first and the second pulses.

### 4.15.21 `t1_start`

`real(8), default=0d0`

Available for 'maxwell' and TDDFT based options of `theory`.  
Shift the starting time of the first pulse. (this is not available for multiscale option).

### 4.15.22 `num_dipole_source`

`integer, default=0`

Available for TDDFT based options of `theory`.  
Number of radiation sources to mimic optical near fields. Maximum number is 2.

### 4.15.23 `vec_dipole_source(3,num_dipole_source)`

`real(8), default=0d0`

Available for TDDFT based options of `theory`.  
Dipole vectors of the radiation sources mimicing optical near fields.

### 4.15.24 `cood_dipole_source(3,num_dipole_source)`

`real(8), default=0d0`

Available for TDDFT based options of `theory`.  
Coordinates of the radiation sources mimicing optical near fields.

### 4.15.25 `rad_dipole_diele`

`real(8), default=2d0 [a.u.]`

Available for TDDFT based options of `theory`.  
Radii of dielectric spheres of the radiation sources mimicing optical near fields.

## 4.16 `&singlescale[Trial]`

### 4.16.1 `method_singlescale`

`character, default='3d'`

Available for `theory='single_scale_maxwell_tddft'`.  
Type of single-scale Maxwell-TDDFT method.

Options:

'3d' / 3-dimensional FDTD + TDDFT

'1d' / 1-dimensional FDTD (along the z axis) + TDDFT

'1d\_fourier' / '1d' with 3D Fourier component of the vector potential

#### 4.16.2 cutoff\_G2\_emfield

real(8), default=-1d0

Available for `theory='single_scale_maxwell_tddft'`.

Cutoff energy of Fourier component of the vector potential when `method_singlescale='1d_fourier'`.

#### 4.16.3 yn\_symmetrized\_stencil

[Trial] character, default='n'

Available for `theory='single_scale_maxwell_tddft'`.

Switch to symmetrize the finite differences for the product of vector potential and orbitals,  $(\nabla A(r) \cdot \psi(r))$ . This option improves hermiticity of the Hamiltonian although computational cost increases.

#### 4.16.4 yn\_put\_wall\_z\_boundary

[Trial] character, default='n'

Available for DFT/TDDFT based options of `theory`.

Option to put potential wall near the boundary planes at  $z=0$  and  $z=\text{'&system/al(3)'}$ . This potential prevents electrons from crossing the  $z$ -boundary plane. In the single-scale Maxwell-TDDFT method, the electron density on the  $z$ -boundary plane harms the norm conservation of electrons due to the discontinuity of the vector potential. The wall is described using the square of cosine function.

Options:

'y' / put the potential wall

'n' / no potential wall

#### 4.16.5 wall\_height

real(8), default=100.0 [eV]

Available for `yn_put_wall_z_boundary='y'`.

The height of the potential wall.

#### 4.16.6 wall\_width

real(8), default=5.0 [Angstrom]

Available for `yn_put_wall_z_boundary='y'`.

The width of the potential wall defined by the length from the potential peak ( $z=0$  and  $z=\text{'&system/al(3)'}$ ) to the edge.

## 4.17 &multiscale

### 4.17.1 fdtddim

[Trial] character, default='1d'

Available for `theory='multi_scale_maxwell_tddft'` with `yn_periodic='y'`  
Dimension of macroscopic scale system (Maxwell(FDTD) calculation) in multiscale Maxwell-TDDFT method.

Options:

'3d' / 3-dimensional FDTD for macroscopic electromagnetism [currently not available]

'1d' / 1-dimensional FDTD (along the  $x$ -axis) for macroscopic electromagnetism

### 4.17.2 nx\_m

integer, default=1

Available for `theory='multi_scale_maxwell_tddft'` with `yn_periodic='y'`

Number of macroscopic grid points inside materials for  $x$ -direction.

### 4.17.3 ny\_m

### 4.17.4 nz\_m

[Trial] integer, default=1)

Available for `theory='multi_scale_maxwell_tddft'` with `yn_periodic='y'`

Number of macroscopic grid points inside materials for ( $y/z$ )-direction.

### 4.17.5 hx\_m

real(8), default=0d0

Available for `theory='multi_scale_maxwell_tddft'` with `yn_periodic='y'`

Grid spacing of macroscopic coordinate for  $x$ -direction.

Variable `hx_m` is deprecated, and will be moved to `&units/dl_em(1)`

### 4.17.6 hy\_m

### 4.17.7 hz\_m

[Trial] real(8), default=0d0

Available for `theory='multi_scale_maxwell_tddft'` with `yn_periodic='y'`

Grid spacing of macroscopic coordinate for ( $y/z$ )-direction.

Variable `hy_m` and `hz_m` are deprecated, and will be moved to `&units/dl_em(2:3)`

### 4.17.8 nxvac1\_m

### 4.17.9 nxvacr\_m

integer, default=1/0

Available for `theory='multi_scale_maxwell_tddft'` with `yn_periodic='y'`

Number of macroscopic grid points for vacuum region.

`nxvac1_m` / `nxvacr_m` specifies the number for negative / positive  $x$ -direction in front of the material.

## 4.18 &maxwell

### 4.18.1 al\_em(3)

real(8), default=0d0

Available for `theory='maxwell'`.

Size of simulation box in electromagnetic analysis.

### 4.18.2 dl\_em(3)

real(8), default=0d0

Available for `theory='maxwell'` and `theory='multi_scale_maxwell_tddft'`.

Spacing of real-space grids in electromagnetic analysis.

### 4.18.3 dt\_em

real(8), default=0

Available for `theory='maxwell'`.

Time step for electromagnetic analysis.

### 4.18.4 nt\_em

integer, default=0

Available for `theory='maxwell'`.

Number of total time steps of time propagation in electromagnetic analysis.

### 4.18.5 boundary\_em(3,2)

character, default='default'

Available for `theory='maxwell'` and `theory='multi_scale_maxwell_tddft'`.

Boundary condition in electromagnetic analysis. The first index(1-3 rows) corresponds to  $x$ ,  $y$ , and  $z$  axes. The second index(1-2 columns) corresponds to bottom and top of the axes.

Options:

'abc' / absorbing boundary

'pec' / perfect electric conductor

'periodic' / periodic boundary

If `&system/yn_periodic='n'`, 'default', 'abc', and 'pec' can be chosen, where 'default' automatically chooses 'abc'. If `&system/yn_periodic='y'`, 'default', 'abc', and 'periodic' can be chosen, where 'default' automatically chooses 'periodic'. If `theory='maxwell'`, perfectly matched layer(PML) is used for 'abc'.

#### 4.18.6 shape\_file

character, default='none'

Available for `theory='maxwell'`.

Name of input shape file in electromagnetic analysis. The shape file can be generated by using `FDTD_make_shape` in SALMON utilities (<https://salmon-tddft.jp/utilities.html>).

#### 4.18.7 media\_num

integer, default=0

Available for `theory='maxwell'`.

Number of media in electromagnetic analysis.

#### 4.18.8 media\_type(:)

character, default='vacuum'

Available for `theory='maxwell'`.

`media_type(n)` specifies type of n-th media in electromagnetic analysis.

Options:

- 'vacuum'
- 'constant media'
- 'pec'
- 'lorentz-drude'

If 'lorentz-drude' is chosen, linear response calculation is feasible by setting `&emfield/ae_shape1` or `ae_shape2='impulse'`.

#### 4.18.9 epsilon\_em(:)

real(8), Default=1d0

Available for `theory='maxwell'` and for TDDFT based options of theory with `trans_longi='2d'`.

For `theory='maxwell'`, `epsilon_em(n)` specifies relative permittivity of n-th media in electromagnetic analysis.

For TDDFT based options of theory with `trans_longi='2d'`, the relative permittivity of the transparent media on both sides of the film is specified by `epsilon_em(1)` and `epsilon_em(2)`, respectively.

#### 4.18.10 mu\_em(:)

real(8), default=1d0

Available for `theory='maxwell'`.

`mu_em(n)` specifies relative permeability of n-th media in electromagnetic analysis.

#### 4.18.11 sigma\_em(:)

real(8), default=0d0

Available for `theory='maxwell'`.

`sigma_em(n)` specifies conductivity of n-th media in electromagnetic analysis.

#### 4.18.12 pole\_num\_ld(:)

integer, default=1

Available for `theory='maxwell'`.

`pole_num_ld(n)` specifies number of poles of n-th media, available for

`type_media(n)='lorentz-drude'` in electromagnetic analysis.

#### 4.18.13 omega\_p\_ld(:)

real(8), default=0d0

Available for `theory='maxwell'`.

`omega_p_ld(n)` specifies plasma frequency of n-th media, available for

`type_media(n)='lorentz-drude'` in electromagnetic analysis.

#### 4.18.14 f\_ld(:,:)

real(8), default=0d0

Available for `theory='maxwell'`.

`f_ld(n,m)` specifies m-th oscillator strength of n-th media, available for

`type_media='lorentz-drude'` in electromagnetic analysis. The first index is the media ID

whose maximum value is given by `media_num`. The second index is the pole ID whose maximum

value is given by `pole_num_ld(n)`.

#### 4.18.15 gamma\_ld(:,:)

real(8), default=0d0

Available for `theory='maxwell'`.

`gamma_ld(n,m)` specifies m-th collision frequency of n-th media, available for

`type_media(n)='lorentz-drude'` in electromagnetic analysis. The first index is the media ID

whose maximum value is given by `media_num`. The second index is the pole ID whose maximum

value is given by `pole_num_ld(n)`.

#### 4.18.16 omega\_ld(:,:)

real(8), default=0d0

Available for `theory='maxwell'`.

`omega_ld(n,m)` specifies m-th oscillator frequency of n-th media, available for `type_media(n)='lorentz-drude'` in electromagnetic analysis. The first index is the media ID whose maximum value is given by `media_num`. The second index is the pole ID whose maximum value is given by `pole_num_ld(n)`.

#### 4.18.17 wave\_input

character, default='none'

Available for `theory='maxwell'`.

If 'source', the incident pulse in electromagnetic analysis is generated by the incident current source.

#### 4.18.18 ek\_dir1(3)

#### 4.18.19 ek\_dir2(3)

real(8), default=0d0

Available for `theory='maxwell'`.

Propagation direction of the first/second pulse.

#### 4.18.20 source\_loc1(3)

#### 4.18.21 source\_loc2(3)

real(8), default=0d0

Available for `theory='maxwell'`.

Location of the incident current source of the first/second pulse. Note that the coordinate system ranges from  $-a_{em}/2$  to  $a_{em}/2$  for `&system/yn_periodic='n'` while ranges from 0 to  $a_{em}$  for `&system/yn_periodic='y'`.

#### 4.18.22 obs\_num\_em

integer, default=0

Available for `theory='maxwell'`.

Number of observation points in electromagnetic analysis. From the obtained results, figure and animation files can be generated by using SALMON utilities (<https://salmon-tddft.jp/utilities.html>).

#### 4.18.23 obs\_samp\_em

integer, default=1

Available for `theory='maxwell'`.

Sampling time-step of the observation in electromagnetic analysis.

#### 4.18.24 obs\_loc\_em(:,3)

real(8), default=0d0

Available for `theory='maxwell'`.

`obs_loc_em(n,1:3)=x,y,z` specifies location of the n-th observation point in electromagnetic analysis. Note that the coordinate system ranges from  $-a_{em}/2$  to  $a_{em}/2$  for `&system/yn_periodic='n'` while ranges from 0 to  $a_{em}$  for `&system/yn_periodic='y'`.

#### 4.18.25 yn\_obs\_plane\_em(:)

character, default='n'

Available for `theory='maxwell'`.

Specify whether or not to generate output of the electromagnetic fields on the planes (xy, yz, and xz planes) for n-th observation point. This option must be 'y' for generating animation files by using `FDTD_make_figani` in SALMON utilities (<https://salmon-tddft.jp/utilities.html>).

Options:

'y'  
'n'

#### 4.18.26 yn\_obs\_plane\_integral\_em(:)

character, default='n'

Available for `theory='maxwell'`.

Specify whether or not to generate output of the spatial integration of electromagnetic fields on the planes (xy, yz, and xz planes) for n-th observation point.

Options:

'y'  
'n'

#### 4.18.27 yn\_wf\_em

character, default='y'

Available for `theory='maxwell'`.

Switch of a window function for linear response calculation. Available for `&calculation/theory=maxwell`.

Options:

'y'  
'n'

#### 4.18.28 film\_thickness

real(8), default=0d0

Available for TDDFT based options of `theory` with `trans_longi='2d'`.  
Thickness of the film for the 2D maxwell-TDDFT method.



The relative permittivity of the transparent media on both sides of the film can be specified by `epsilon_em(1)` and `epsilon_em(2)`, respectively.

#### 4.18.29 `media_id_pml(3:2)`

integer, default=0

Available for `theory='maxwell'`.

Media ID used in PML. The first index(1-3 rows) corresponds to *x*, *y*, and *z* axes. The second index(1-2 columns) corresponds to bottom and top of the axes.

#### 4.18.30 `media_id_source1`

#### 4.18.31 `media_id_source2`

integer, default=0

Available for `theory='maxwell'`.

Media ID used in incident current source1/source2 to generate the first/second pulse.

### 4.19 `&analysis`

#### 4.19.1 `projection_option`

character, default='no'

Available for TDDFT based options of `theory`.

Methods of projection to analyze the excited states (e.g. the number of excited electrons).

Options:

'no' / no projection.

'gs' / projection to eigenstates of ground-state Hamiltonian.

'rt' / projection to eigenstates of instantaneous Hamiltonian. [currently not available]

#### 4.19.2 `out_projection_step`

integer, default=100

Available for TDDFT based options of `theory`.

Results of the projection analysis will be outputted every `out_projection_step` step during the time-propagation.

#### 4.19.3 `nenergy`

integer, default=1000

Number of energy grid points for frequency-domain analysis. This parameter is used, for examples, in `theory='tddft_response'` and `theory='maxwell'`.

#### 4.19.4 de

real(8), Default=0.01d0 (eV)

Energy grid size for frequency-domain analysis.

This parameter is used, for examples, in `theory='tddft_response'` and `theory='maxwell'`.

#### 4.19.5 out\_rt\_energy\_step

integer, default=10

Available for the TDDFT based option of `theory`.

Total energy is calculated and printed every `out_rt_energy_step` time steps.

#### 4.19.6 yn\_out\_psi

character, default='n'

Available for `theory='dft'`.

Switch for output of orbitals.

Options:

'y' / enable

'n' / disable

The format of the output is specified by `&analysis/format_voxel_data`.

#### 4.19.7 yn\_out\_dos

character, default='n'

Available for `theory='dft'`.

Switch for output of density of states.

Options:

'y' / enable

'n' / disable

#### 4.19.8 yn\_out\_pdos

character, default='n'

Available for `theory='dft'`.

Switch for output of projected density of states.

Options:

'y' / enable

'n' / disable

#### 4.19.9 yn\_out\_dos\_set\_fe\_origin

character, default='n'

Available when `yn_out_dos='y'` or `yn_out_pdos='y'`.

Switch to set the Fermi energy to zero.

Options:

'y' / enable

'n' / disable

This option is not available if the temperature is not set in the calculation.

#### 4.19.10 out\_dos\_start

real(8), default=-1d10 (eV)

#### 4.19.11 out\_dos\_end

real(8), default=1d10 (eV)

Available when `yn_out_dos='y'` or `yn_out_pdos='y'`.

Lower/Upper bound of the energy range for the density of states spectra.

If this value is lower/higher than a specific value near the lowest/highest energy level, this parameter is re-set to the value.

#### 4.19.12 out\_dos\_nenergy

integer, default=601

Available when `yn_out_dos='y'` or `yn_out_pdos='y'`.

Number of energy points sampled in the density of states spectra.

#### 4.19.13 out\_dos\_function

character, default='gaussian'

Available when `yn_out_dos='y'` or `yn_out_pdos='y'`.

Choice of the smearing function for the density of states spectra.

Options:

gaussian / Gaussian function

lorentzian / Lorentzian function

#### 4.19.14 out\_dos\_width

real(8), default=0.1d0 [eV]

Available when `yn_out_dos='y'` or `yn_out_pdos='y'`.

Smearing width used in the density of states spectra.

#### 4.19.15 yn\_out\_dns

character, default='n'

Available for `theory='dft'`.

Switch to output electron density distribution of the ground state.

Options:

'y' / enable

'n' / disable

#### 4.19.16 yn\_out\_dns\_rt

character, default='n'

Available when `theory='dft_md'` or `'theory=tddft_pulse'`.

Switch to output electron density distribution during the time-propagation.

Options:

'y' / enable

'n' / disable

#### 4.19.17 out\_dns\_rt\_step

integer, default=50

Available when `theory='dft_md'` or `'theory=tddft_pulse'`.

Density is outputted every `out_dns_rt_step` steps.

#### 4.19.18 yn\_out\_dns\_ac\_je

character, default='n'

Available for `theory='single_scale_maxwell_tddft'`.

Switch to print the electron density, vector potential, electronic current, and ionic coordinates every `outdns_dns_ac_je_step` time steps.

Options:

'y' / enable

'n' / disable

The data written in binary format are divided into files corresponding to the space-grid parallelization number.

#### 4.19.19 out\_dns\_ac\_je\_step

integer, default=50

Available for `theory='single_scale_maxwell_tddft'`.

Electron density, vector potential, electronic current, and ionic coordinates are outputted every `outdns_dns_ac_je_step` time steps.

yn\_out\_dns\_trans

[currently not available] character default='n'

Available for `theory='tddft_pulse'`.

Switch to calculate transition density at specified frequency  $\omega$  (specified by `out_dns_trans_energy`),  $\text{drho}(\mathbf{r}, \omega) = \text{FT}(\rho(\mathbf{r}, t) - \rho_{\text{gs}}(\mathbf{r})) / T$ .

Options:

'y' / enable

'n' / disable

### 4.19.20 out\_dns\_trans\_energy

[currently not available] real(8), default=1.55d0 [eV]

Available for `theory='tddft_pulse'`.

A frequency to output  $\text{drho}(\mathbf{r}, \omega) = \text{FT}(\rho(\mathbf{r}, t) - \rho_{\text{gs}}(\mathbf{r})) / T$ .

### 4.19.21 yn\_out\_elf

character, default='n'

Available for `theory='dft'`.

Switch to output the electron localization function.

Options:

'y' / enable

'n' / disable

### 4.19.22 yn\_out\_elf\_rt

character, default='n'

Available for `theory='dft_md', 'tddft_pulse'`.

Switch to output the electron localization function during the time propagation every

`out_elf_rt_step` time steps.

Options:

'y' / enable

'n' / disable

### 4.19.23 out\_elf\_rt\_step

integer, default=50

Available for `theory='dft_md', 'tddft_pulse'`.

Electron localization function during the time propagation is outputted every `out_elf_rt_step` time steps.

### 4.19.24 yn\_out\_estatic\_rt

character, default='n'

Available for `theory='tddft_pulse'`.

Switch to print the static electric field during the time propagation every `out_estatic_rt_step` time steps.

Options:

'y' / enable

'n' / disable

#### 4.19.25 out\_estatic\_rt\_step

integer, default=50

Available for `theory='tddft_pulse'`.

The static electric field during the time propagation is outputed every `out_estatic_rt_step` time steps.

#### 4.19.26 yn\_out\_rvf\_rt

character, default='n'

Available for TDDFT based options and 'dft\_md' option of `theory`.

Switch to print the coordinates[A], velocities[au], forces[au] of atoms during time-propagation in `SYSname_trj.xyz` every `out_rvf_rt_step` time steps.

Options:

'y' / enable

'n' / disable

If `yn_md='y'`, this option is automatically turned on.

#### 4.19.27 out\_rvf\_rt\_step

integer, default=10

Available for TDDFT based options and 'dft\_md' option of `theory`.

The coordinates[A], velocities[au], forces[au] of atoms during time-propagation are outputed in `SYSname_trj.xyz` every `out_rvf_rt_step` time steps.

#### 4.19.28 yn\_out\_tm

[Trial] character, default='n'

Available for `yn_periodic='y'` with `theory='dft'`.

Switch to calculate and print the transition matrix elements between occupied and virtual orbitals to `SYSname_tm.data` after the ground state calculation.

Options:

'y' / enable

'n' / disable

#### 4.19.29 out\_ms\_step

integer, default=100

Available for `theory='multi_scale_maxwell_tddft'`.

Some quantities are printed every `out_ms_step` time step in the Maxwell-TDDFT multiscale calculations.

### 4.19.30 format\_voxel\_data

character, default='cube'

Available for `yn_out_psi='y'`, `yn_out_dns(_rt)='y'`, `yn_out_dns_ac_je='y'`,  
`yn_out_elf(_rt)='y'`, `yn_out_estatic_rt='y'`.

Option of the file format for three-dimensional volumetric data.

'avs' / AVS format

'cube' / cube format

'vtk' / vtk format

### 4.19.31 nsplit\_voxel\_data

integer, default=1

Available for `format_voxel_data='avs'`.

Number of separated files for three dimensional data.

### 4.19.32 yn\_lr\_w0\_correction

[Trial] character, default='n'

Available for `yn_periodic='y'` and `trans_longi='tr'` with  
`theory='tddft_response'`.

Apply correction around zero frequency of dielectric function to suppress numerical error.

Options:

'y' / enable

'n' / disable

### 4.19.33 yn\_out\_perflog

character, default='y'

Available for all `theory`

Switch to print the performance log of routines and modules.

Options:

'y' / enable

'n' / disable

### 4.19.34 format\_perflog

character, default='stdout'

Available for `yn_out_perflog = 'y'`

The output format of performance log.

Options:

'stdout' / standard output unit

'text' / save as a text file

'csv' / save as a csv format file

## 4.20 &poisson

### 4.20.1 layout\_multipole

character, Default=3

Available for `yn_periodic='n'` in DFT and TDDFT based options of `theory`.

This parameter specifies how to achieve multipole expansion in the Hartree potential calculation.

Options:

- 1/ A single pole at the center.
- 2/ Multipoles are set at each center of atoms.
- 3/ Multipoles are set at the center of mass of electrons in prepared cuboids in each process.

### 4.20.2 num\_multipole\_xyz(3)

integer, default=0

Available for `yn_periodic='n'` in DFT and TDDFT based options of `theory`.

Number of multipoles. When default is set, the number of multipoles is calculated automatically.

### 4.20.3 lmax\_multipole

[Trial] integer, default=4

Available for `yn_periodic='n'` in DFT and TDDFT based options of `theory`.

A maximum order of the multipole expansion to prepare boundary condition of Poisson equation.

### 4.20.4 threshold\_cg

real(8), default=1d-15 [a.u.]

Available for `yn_periodic='n'` in DFT and TDDFT based options of `theory`.

A threshold for the convergence of the Hartree-cg calculation. A quantity examined is given by  $\|tV_h(i) - tV_h(i-1)\|^2 / (\text{number of grids})$ .

## 4.21 &ewald

### 4.21.1 newald

integer, default=4

Available for `yn_periodic='y'` in DFT/TDDFT based options of `theory`.

Parameter of the Ewald method for the ion-ion Coulombic interaction. Short-range part of the Ewald sum is calculated within `newald`-th nearest neighbor cells.



### 4.21.2 aewald

real(8), default=0.5d0 [a.u.]

Available for `yn_periodic='y'` in DFT/TDDFT based options of `theory`.

Square of range separation parameter for Ewald method (This parameter is given only in atomic unit).

### 4.21.3 cutoff\_r

real(8), default=-1d0

Available for `yn_periodic='y'` in DFT/TDDFT based options of `theory`.

Cut-off length in real-space. The length is automatically determined if `cutoff_r < 0`.

### 4.21.4 cutoff\_r\_buff

real(8), default=2d0 [a.u.]

Available for `yn_periodic='y'` in `yn_md='y'` or in `theory='dft_md'`.

Buffer length in radius for book-keeping for real-space interaction.

### 4.21.5 cutoff\_g

real(8), Default=-1d0

Available for `yn_periodic='y'` in DFT/TDDFT based options of `thddeory`.

Cut-off in G-space in the Ewald method. No cut-off in default.

## 4.22 &opt[Trial]

### 4.22.1 nopt

integer, default=100

Available for `yn_opt='y'` in `theory='dft'`.

The maximum step number of geometry optimization.

### 4.22.2 convrg\_opt\_fmax

real(8), default=1d-3 (a.u.)

Available for `yn_opt='y'` in `theory='dft'`.

Convergence threshold of geometry optimization is specified for the maximum force acting on atoms.

### 4.22.3 max\_step\_len\_adjust

real(8), default=-1d0

Available for `yn_opt='y'` in `theory='dft'`.

Set maximum optimization step length (if positive number is given)

## 4.23 &md[Trial]

### 4.23.1 ensemble

character, default='NVE'

Available for `yn_md='y'` or `theory='dft_md'`.

Ensemble in MD option:

Options:

NVE/ NVE ensemble (constant energy and volume system)

NVT/ NVT ensemble (constant temperature and volume system)

### 4.23.2 thermostat

character, default='nose-hoover'

Available for `yn_md='y'` or `theory='dft_md'`.

Thermostat in "NVT" option:

Options:

nose-hoover/ Nose-Hoover thermostat

### 4.23.3 step\_velocity\_scaling

integer, default=-1

Available for `yn_md='y'` or `theory='dft_md'`.

Time step interval for velocity-scaling. Velocity-scaling is applied if this is set to positive.

### 4.23.4 step\_update\_ps

integer, default=10

Available for `yn_md='y'` or `theory='dft_md'`.

Time step interval for updating pseudopotential (Larger number reduces computational time but increases inaccuracy).

### 4.23.5 temperature0\_ion\_k

real(8), Default=298.15d0 [K]

Available for `yn_md='y'` or `theory='dft_md'`.

Setting ionic temperature in unit of [K] for NVT ensemble, velocity scaling and generating initial velocities.

### 4.23.6 yn\_set\_ini\_velocity

character, Default='n'

Available for `yn_md='y'` or `theory='dft_md'`.

Switch to generate initial velocities.

Options:

y/ Generate initial velocity with Maxwell-Boltzman distribution

n/ disable

### 4.23.7 file\_ini\_velocity

[Trial] character, default='none'

Available for `yn_md='y'` or `theory='dft_md'`.

File name for reading initial velocities. This is read if the file name is given, then, the priority is higher than use of `set_ini_velocity` and restart data of velocities. The format is simply `vx(iatom)` `vy(iatom)` `vz(iatom)` in each line. The order of atoms must be the same as the given coordinates in the main input file. In case of using nose-hoover thermostat, a thermostat variable should be put at the last line (all atomic unit).

### 4.23.8 thermostat\_tau

real(8), default=1d0 [fs]

Available for `yn_md='y'` or `theory='dft_md'`.

Parameter in Nose-Hoover method: controlling time constant for temperature.

### 4.23.9 yn\_stop\_system\_mom

character, default='n'

Available for `yn_md='y'` or `theory='dft_md'`.

Center of mass is fixed every time step.

Options:

y/ enable

n/ disable

## 4.24 &jellium

### 4.24.1 yn\_jm

character, default='n'

Available for the DFT/TDDFT based options of `theory`.

Switch to use jellium model.

Options:

y/ enable

n/ disable

When `yn_jm='y'`, `&functional/xc` must be 'pz'.

### 4.24.2 yn\_charge\_neutral\_jm

character, default='y'

Available for `yn_jm='y'` in the DFT/TDDFT based options of `theory`.

Option to enforce exact charge neutrality :

Options:

y/ enable. `rs_bohr_jm` is modified to fulfill exact charge neutrality.

n/ disable. `rs_bohr_jm` is not modified, and there may appears small charge-neutrality error.

### 4.24.3 `yn_output_dns_jm`

character, default='y'

Available for `yn_jm='y'` in the DFT/TDDFT based options of `theory`.

Switch to output positive background charge density.

Options:

y/ enable

n/ disable

### 4.24.4 `shape_file_jm`

character, default='none'

Available for `yn_jm='y'` in the DFT/TDDFT based options of `theory`.

Name of input shape file that contains positive background charge density to be used in the jellium model calculations. The shape file can be generated by using `FDTD_make_shape` in SALMON utilities (<https://salmon-tddft.jp/utilities.html>). When `shape_file_jm='none'`, the shape of the positive background charge density is specified by `sphere_nion_jm` and `sphere_loc_jm` which generate spherical shapes.

### 4.24.5 `num_jm`

integer, Default=0

Available for `yn_jm='y'` in the DFT/TDDFT based options of `theory`.

When `shape_file_jm` is not 'none', `num_jm` specifies number of media used in the jellium model.

When `shape_file_jm='none'`, `num_jm` specifies number of spherical shapes.

### 4.24.6 `rs_bohr_jm(:)`

real(8), default=0d0

Available for `yn_jm='y'` in the DFT/TDDFT based options of `theory`.

When `shape_file_jm` is not 'none', `rs_bohr_jm(n)` specifies the Wigner-Seitz radius of n-th media. When `shape_file_jm='none'`, `rs_bohr_jm(n)` specifies the Wigner-Seitz radius of n-th sphere.

### 4.24.7 `sphere_nion_jm(:)`

integer, default=0

Available for `yn_jm='y'` and `shape_file_jm='none'` in the DFT/TDDFT based options of `theory`. `sphere_nion_jm(n)` specifies ion number for n-th sphere. At present, only neutral systems can be treated.

### 4.24.8 sphere\_loc\_jm(:,3)

real(8), default=0d0

Available for `yn_jm='y'` and `shape_file_jm='none'` in the DFT/TDDFT based options of theory. `sphere_loc_jm(n,1:3)=x,y,z` specifies location of center of mass for n-th sphere. Note that the coordinate system ranges from  $-a/2$  to  $a/2$  for `&system/yn_periodic='n'` while ranges from 0 to  $a$  for `&system/yn_periodic='y'`.

## 4.25 &code

### 4.25.1 yn\_want\_stencil\_hand\_vectorization

character, default='y'

Switch to use hand-vectorized optimization code of stencil in the hamiltonian calculation.

SALMON checks if the calculation can use the hand-vectorized code. If it fails, SALMON will use a typical implementation.

### 4.25.2 yn\_want\_communication\_overlapping

character, default='n'

Available for `theory='tddft*' or '*maxwell_tddft'`

Switch to use computation/communication overlap algorithm to improve the performance of stencil in the hamiltonian calculation.

SALMON checks if the calculation can use the overlap algorithm. If it fails, SALMON will use a non-overlap algorithm.

### 4.25.3 stencil\_openmp\_mode

character, default='auto'

This option selects an OpenMP parallelization mode of stencil in the hamiltonian calculation.

Options:

- `auto` / SALMON decides the parallelization target automatically.
- `orbital` / OpenMP parallelization is applied to orbital (and k-point) loop.
- `rgrid` / OpenMP parallelization is applied to real-space grid loop.

### 4.25.4 current\_openmp\_mode

character, default='auto'

This selects an OpenMP parallelization mode of the current calculation.

Options:

- `auto` / SALMON decides the parallelization target automatically.
- `orbital` / OpenMP parallelization is applied to orbital (and k-point) loop.
- `rgrid` / OpenMP parallelization is applied to real-space grid loop.

#### 4.25.5 `force_openmp_mode`

character, default='auto'

This selects an OpenMP parallelization mode of the force calculation.

Options:

`auto` / SALMON decides the parallelization target automatically.

`orbital` / OpenMP parallelization is applied to orbital (and k-point) loop.

`rgrid` / OpenMP parallelization is applied to real-space grid loop.

## RELEASE HISTORY

### 5.1 Release Notes

- Release note of SALMON ver. 2:  
<https://github.com/SALMON-TDDFT/SALMON2/releases>
- Release note of SALMON ver. 0 and 1:  
<https://github.com/SALMON-TDDFT/SALMON/releases>

### 5.2 Details of Minor Changes

Followings are history of fixed bugs and changes of models/inputs/outputs after releasing v.2.0.0. (not complete list currently)

#### 5.2.1 Fixed bugs

(Fixed in v.2.0.2)

- Small noise on the total energy in TDDFT calculation (that is seen with weak pulse around e.g.  $I=1\text{d9 W/cm}^2$ ) has been removed.
- The printed absolute values of electron density in cube format has been fixed..
- Printing of the external field in TDDFT calculation of the isolated system has a bug in v.2.0.0 and v.2.0.1. It has been fixed in v.2.0.2.
- The file reading option of the external electric field in TDDFT calculation ("file\_input1" in &emfield) has been fixed.
- Invalid occupation number printed in SYSNAME\_ovlp.data file for projection option with non-uniform k-points has been fixed.

(Fixed in v.2.0.0)

- The imaginary part of wavefunction was not printed in cube format until v.1.2.1

(Fixed in v.?.?.?)

- Abnormal calculation that sometimes happens if zero value is included in the atomic coordinate in the input with "A\_eV\_fs" has been fixed.

## 5.2.2 Changes of models/inputs/outputs

(v.2.0.2)

- The definition of the total energy of the periodic system printed in TDDFT calculation has changed: The electric field energy is included until v.2.0.1. It has not been included from v.2.0.2.
- The directory names generated by "method\_wf\_distributor=slice" option have changed from v.2.0.2.



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