

OEP-Dev 1.0.2-alpha

Novel methods for Quantum Chemistry of Extended Molecular Aggregates

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Chapter 1

GEFP-OEP

GEFP Project

The package is designed for quantum chemistry calculations. Currently, the implemented methods are:

- Density Matrix Functional Theory
- Density-Based decomposition of Mandado and Hermida-Ramon

The tutorial is under preparation.

Installation

Installation prerequisites:

- Python 3.3 or newer
- NumPy module for Python, 1.16.3 or newer
- SciPy module for Python, 1.2.1 or newer
- Psi4 1.2.0 or newer
- OEP-Dev 1.0.3 or newer

To install the GEFP package type the following commands:

```
sudo python setup.py install
```


Chapter 2

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Chapter 3

Introduction

Exploring biological phenomena at molecular scale is oftentimes indispensable to develop new drugs and intelligent materials.

Most of relevant system properties are affected by intermolecular interactions with nearby environment such as solvent or closely bound electronic chromophores. Studying such molecular aggregates requires rigorous and accurate quantum chemistry methods, the cost of which grows very fast with the number of electrons. Despite many methodologies have been devised to describe energetic and dynamical properties of **extended molecular systems** efficiently and accurately, there exist particularly difficult cases in which modelling is still challenging:

- describing electronic transitions in solution or
- when coupled with other electronic transition via resonance energy transfer,
- performing molecular dynamics at very high level of theory including dynamic electron correlation,
- vibrational frequency calculations of particular normal mode in condensed phases

and so on. The reason behind (sometimes prohibitively) high costs of fully *ab initio* calculations in the above areas is the complexity of mathematical models often based on wave functions rather than (conceptually more straightforward) electronic densities. On the other hand, it has been pointed out before that the one-electron density distributions are of particular importance in chemistry. It can be thus utilized as a means of developing a general model that re-expresses the physics of intermolecular interactions in terms of effective one-electron functions that are easier to handle in practice.

This Project will focus on finding a unified way to simplify various equations of Quantum Chemistry of extended molecular systems, i.e., molecular aggregates such as interacting chromophores and molecules solvated by water and other solvents. Indeed, one of the important difficulties encountered in Quantum Chemistry of large systems is the need of evaluation of special kind of numbers known as *electron repulsion integrals*, or in short, ERI's. In a typical calculation, the amount of ERI's can be as high as tens or even hundreds of millions (!) that unfortunately prevents from application of conventional methods when the number of particles in question is too large. In the Project, the complicated expressions involving ERI's shall be greatly simplified to reduce the computational costs as much as possible while introducing no or minor approximations to the original theories.

3.1 Research Project Methodology

In this Project the new theoretical protocol based on the one-electron effective potentials (OEP's) is developed. The main principle is to rewrite arbitrary sum of functions f of electron repulsion integrals (ERI's) by defining OEP's according to the following general prescription:

$$\sum_f \left[\left(\phi_i^A \phi_j^A || \phi_k^B \phi_l^B \right) \right] = \left(\phi_i^A | v_{kl}^B | \phi_j^A \right) \rightarrow \text{point charge or density fitting}$$

$$\sum_f \left[\left(\phi_i^A \phi_j^B || \phi_k^B \phi_l^B \right) \right] = \left(\phi_i^A | v_{kl}^B | \phi_j^B \right) \rightarrow \text{density fitting,}$$

where A and B denote different molecules and ϕ_i is the i -th molecular orbital or basis function. Here, v_{kl}^B denotes the [List of One-Electron Potentials](#) *ab initio* "OEP matrix element". The technique described above will be applied to simplify expressions for

- short-range excitation energy transfer couplings between chlorophyll subunits of reaction centres in photosynthesis
- Pauli interaction repulsion energy
- charge-transfer interaction energy
- electric field-induced charge density polarization of molecules.

The above developments might be used in fragment-based *ab initio* molecular dynamics protocols of new generation.

3.2 Expected Impact on the Development of Science, Civilization and Society

The proposed OEP's are expected to significantly develop the fragment-based methods that are widely used in physical chemistry and modelling of biologically important systems. Owing to universality of OEP's, they could find applications in many branches of chemical science: non-empirical* molecular dynamics, short-range resonance energy transfer in photosynthesis, electronic and vibrational solvatochromism, multidimensional spectroscopy and so on. In particular:

- the OEP-based models of Pauli repulsion energy and charge-transfer (CT) energy could be used to improve the computational performance of the second generation effective fragment potential method (EFP2). At present, the CT term is very time consuming and due to this reasons it is not used in most of applications of EFP2 to perform molecular dynamics simulations.
- the OEP-based model of EET couplings could significantly improve modelling of energy transfer in the light harvesting complexes. At present, short-range phenomena (Dexter mechanisms of EET) are very difficult to efficiently and quantitatively asses when performing statistical averaging and applying to large molecular aggregates. Such Dexter effects could be computed by using OEP's in much more efficient manner without losing high accuracy of parent TDFI-TI method.

- the density matrix polarization (DMS) tensors could be used in new generation fragment-based *ab initio* molecular dynamics protocols that rigorously take into consideration electron correlation effects.

Therefore, it is strongly believed that the OEP's could have an indirect impact on the design of novel drugs and materials for industry.

3.3 The OEPDev Code

To pursue the above challenges in the field of computational quantum chemistry of extended molecular aggregates, the OEPDev platform is developed. Accurate and efficient *ab initio* [models](#) based on OEP's are implemented in the OEPDev code, along with the state-of-the-art benchmark and competing methods. Written entirely in C++, OEPDev is a plugin to Psi4 quantum chemistry package. Therefore, compilation and running the OEPDev code is straightforward and follows the API interface similar to the one used in Psi4 with just a few [specific programing conventions](#). The detailed discussion about using the OEPDev code can be found in [advanced usage section](#).

Chapter 4

OEP Design.

OEP (One-Electron Potential) is associated with certain quantum one-electron operator \hat{v}^A that defines the ability of molecule A to interact in a particular way with other molecules.

Technically, OEP can be understood as a **container object** (associated with the molecule in question) that stores the information about the above mentioned quantum operator. Here, it is assumed that similar OEP object is also defined for all other molecules in a molecular aggregate.

In case of interaction between molecules A and B , OEP object of molecule A interacts directly with wavefunction object of the molecule B . Defining a Solver class that handles such interaction Wavefunction class and OEP class

the universal design of OEP-based approaches can be established and developed.

Important: OEP and Wavefunction classes should not be restricted to Hartree-Fock; in general any correlated wavefunction and derived OEP's should be allowed to work with each other.

4.1 OEP Classes

There are many types of OEP's, but the underlying principle is the same and independent of the type of intermolecular interaction. Therefore, the OEP's should be implemented by using a multi-level class design. In turn, this design depends on the way OEP's enter the mathematical expressions, i.e., on the types of matrix elements of the one-electron effective operator \hat{v}^A .

4.1.1 Structure of possible OEP-based expressions and their unification

Structure of OEP-based mathematical expressions is listed below:

Type	Matrix Element	Comment
Type 1	$(I \hat{v}^A J)$	$I \in A, J \in B$
Type 2	$(J \hat{v}^A L)$	$J, L \in B$

In the above table, I, J and K indices correspond to basis functions or molecular orbitals. Basis functions can be primary or auxiliary OEP-specialized density-fitting. Depending on the type of function and matrix element, there are many subtypes of resulting matrix elements that differ in their dimensionality. Examples are given below:

Matrix Element	DF-based form	ESP-based form
$\left(\mu \hat{v}^{A[\mu]} \sigma\right)$	$\sum_{l \in A} v_{\mu l}^A S_{l\sigma}$	$\sum_{\alpha \in A} q_{\alpha}^{A[\mu]} V_{\mu\sigma}^{(\alpha)}$
$\left(i \hat{v}^{A[i]} j\right)$	$\sum_{l \in A} v_{il}^A S_{lj}$	$\sum_{\alpha \in A} q_{\alpha}^{A[i]} V_{ij}^{(\alpha)}$
$\left(j \hat{v}^{A[i]} l\right)$	$\sum_{l \in A} S_{jl} v_{lk}^A S_{kl}$	$\sum_{\alpha \in A} q_{\alpha}^{A[i]} V_{jl}^{(\alpha)}$

In the formulae above, the OEP-part (stored by OEP instances) and the Solver-part (to be computed by the Solver) are separated. It is apparent that all OEP-parts have the form of 2nd- or 3rd-rank tensors with different class of axes (molecular orbitals, primary/auxiliary basis, atomic space). Therefore, they can be uniquely defined by a unified *tensor object* (storing double precision numbers) and unified *dimension object* storing the information of the axes classes.

In Psi4, a perfect candidate for the above is `psi4::Tensor` class declared in `psi4/libthce/thce.h`. Except from the numeric content its instances also store the information of the dimensions in a form of a vector of `psi4::Dimension` instances.

Another possibility is to use `psi::Matrix` objects, instead of `psi4::Tensor` objects, possibly putting them into a `std::vector` container in case there is more than two axes.

Chapter 5

List of One-Electron Potentials

Here I provide the list of OEP's that have been already derived within the scope of the OEPDev project.

Note

Add here a table with all the OEP types along with their symbols used in the OEPDev code (e.g., `Murrell.etal-S1` etc).

5.1 Electrostatic Energy OEP's

For electrostatic energy calculations, OEP is simply the electrostatic potential due to nuclei and electrons.

3D form:

$$v(\mathbf{r}) = \sum_x \frac{Z_x}{|\mathbf{r} - \mathbf{r}_x|} + \sum_{\mu\nu \in A} P_{\nu\mu} \int d\mathbf{r}' \frac{\phi_\mu^*(\mathbf{r}') \phi_\nu(\mathbf{r}')}{|\mathbf{r} - \mathbf{r}'|}$$

Matrix form:

$$v_{ik} = \sum_{x \in A} Z_x V_{ik}^{(x)} + \sum_{\mu\nu \in A} P_{\nu\mu} (\mu\nu|ik)$$

5.2 Pauli Repulsion OEP's

The following potentials are derived for the evaluation of the Pauli repulsion energy based on Murrel's expressions.

5.2.1 First-order contribution in overlap matrix expansion.

This contribution is simply the electrostatic potential coming from all nuclei and electron density except* from electron density from molecular orbital i that interacts with the generalized overlap density between i of molecule A and j of molecule B .

3D forms:

$$v(\mathbf{r})_{S^{-1}}^{A[i]} = - \sum_{x \in A} \frac{Z_x}{|\mathbf{r} - \mathbf{r}_x|} + \sum_{\mu \nu \in A} \left\{ D_{\nu\mu} - C_{\mu i}^* C_{\nu i} \right\} \int d\mathbf{r}' \frac{\phi_{\mu}^*(\mathbf{r}') \phi_{\nu}(\mathbf{r}')}{|\mathbf{r} - \mathbf{r}'|}$$

Matrix forms:

$$v_{\xi i}(S^{-1}) = \sum_{\kappa \in A} C_{i\kappa} \left\{ - \sum_{x \in A} V_{\kappa\xi}^{(x)} + \sum_{\mu \nu \in A} \left\{ D_{\nu\mu} - C_{\mu i}^* C_{\nu i} \right\} (\mu \nu | \xi \kappa) \right\}$$

5.2.2 Second-order contribution in overlap matrix expansion.

To be added here!

5.3 Charge-Transfer Energy OEP's

To be added here!

5.4 Excitonic Energy Transfer OEP's

The following potentials are derived for the evaluation of the short-range EET couplings based on Fujimoto's TDFI-TI method.

5.4.1 ET contributions.

3D forms:

$$\begin{aligned} v(\mathbf{r})_1^{A[\mu]} &= -C_{\mu L}^* \sum_{x \in A} \frac{Z_x}{|\mathbf{r} - \mathbf{r}_x|} + \sum_{\nu \kappa \in A} \left\{ C_{\mu L}^* D_{\nu \kappa} - \frac{1}{2} C_{\nu L}^* D_{\mu \kappa} \right\} \int d\mathbf{r}' \frac{\phi_{\nu}^*(\mathbf{r}') \phi_{\kappa}(\mathbf{r}')}{|\mathbf{r} - \mathbf{r}'|} \\ v(\mathbf{r})_2^{A[\mu]} &= C_{\kappa H} \sum_{\nu \kappa \in A} \left\{ 2C_{\nu L}^* C_{\mu H} - C_{\nu H}^* C_{\mu L} \right\} \int d\mathbf{r}' \frac{\phi_{\nu}^*(\mathbf{r}') \phi_{\kappa}(\mathbf{r}')}{|\mathbf{r} - \mathbf{r}'|} \\ v(\mathbf{r})_3^{A[\mu]} &= v(\mathbf{r})_1^{A[\mu]} + v(\mathbf{r})_1^{A[\mu]} \end{aligned}$$

Matrix forms:

$$\begin{aligned}
 v_{\mu\xi}(1) &= -C_{\mu L}^* \sum_{x \in A} V_{\mu\xi}^x + \sum_{\nu\kappa \in A} \left\{ C_{\mu L}^* D_{\nu\kappa} - \frac{1}{2} C_{\nu L}^* D_{\mu\kappa} \right\} (\nu\kappa|\mu\xi) \\
 v_{\mu\xi}(2) &= C_{\kappa H} \sum_{\nu\kappa \in A} \left\{ 2C_{\nu L}^* C_{\mu H}^* - C_{\nu H}^* C_{\mu L}^* \right\} (\nu\kappa|\mu\xi) \\
 v_{\mu\xi}(3) &= v_{\mu\xi}(1) + v_{\mu\xi}(2)
 \end{aligned}$$

5.4.2 HT contributions.

Do be derived.

5.4.3 CT contributions.

To be derived.

5.5 Full HF Interaction OEP's

The following potentials are derived for the evaluation of the full Hartree-Fock interaction energy based on the OEPDev equations.

Chapter 6

Density-fitting Specialized for OEP's

To get the ab-initio representation of a OEP, one can use a procedure similar to the typical density fitting or resolution of identity, both of which are nowadays widely used to compute electron-repulsion integrals (ERI's) more efficiently.

6.1 Fitting in Complete Space

An arbitrary one-electron potential of molecule A acting on any state vector associated with molecule A can be expanded in an *auxiliary space* centered on A as

$$v|i) = \sum_{\xi\eta} v|\xi) [\mathbf{S}^{-1}]_{\xi\eta} (\eta|i)$$

under the necessary assumption that the auxiliary basis set is *complete*. In a special case when the basis set is orthogonal (e.g., molecular orbitals) the above relation simplifies to

$$v|i) = \sum_{\xi} v|\xi) (\xi|i)$$

It can be easily shown that the above general and exact expansion can be obtained by performing a density fitting in the complete space. We expand the LHS of the first equation on this page in a series of the auxiliary basis functions scaled by the undetermined expansion coefficients:

$$v|i) = \sum_{\xi} G_{i\xi} |\xi)$$

which we shall refer here as to the matrix form of the OEP operator. By constructing the least-squares objective function

$$Z[\{G_{\xi}^{(i)}\}] = \int d\mathbf{r}_1 \left[v(\mathbf{r}_1) \phi_i(\mathbf{r}_1) - \sum_{\xi} G_{\xi}^{(i)} \phi_{\xi}(\mathbf{r}_1) \right]^2$$

and requiring that

$$\frac{\partial Z[\{G_{\xi}^{(i)}\}]}{\partial G_{\mu}^{(i)}} = 0 \text{ for all } \mu$$

we find the coefficients $G_{\xi}^{(i)}$ to be

$$\mathbf{G}^{(i)} = \mathbf{v}^{(i)} \cdot \mathbf{S}^{-1}$$

where

$$\begin{aligned} v_{\eta}^{(i)} &= (\eta | v | i) \\ S_{\eta\xi} &= (\eta | \xi) \end{aligned}$$

or explicitly

$$G_{i\xi} = \sum_{\eta} [\mathbf{S}^{-1}]_{\xi\eta} (\eta | v | i)$$

identical to what we obtained from application of the resolution of identity in space spanned by non-orthogonal complete set of basis vectors.

Since matrix elements of an OEP operator in auxiliary space can be computed in the same way as the matrix elements with any other basis function, one can formally write the following identity

$$(X | v | i) = \sum_{\xi\eta} S_{X\xi} [\mathbf{S}^{-1}]_{\xi\eta} (\eta | v | i)$$

where X is an arbitrary orbital. When the other orbital does not belong to molecule A but to the (changing) environment, it is straightforward to compute the resulting matrix element, which is simply given as

$$(j_{\in B} | v^A | i_{\in A}) = \sum_{\xi} S_{j\xi} G_{i\xi}$$

where j denotes the other (environmental) basis function.

In the above equation, the OEP-part (fragment parameters for molecule A only) and the Solver-part (subject to be computed by solver on the fly) are separated. This then forms a basis for fragment-based approach to solve Quantum Chemistry problems related to the extended molecular aggregates.

6.2 Fitting in Incomplete Space

Density fitting scheme from previous section has practical disadvantage of a nearly-complete basis set being usually very large (spanned by large amount of basis set vectors). Any non-complete basis set won't work in the previous example. Since most of basis sets used in quantum chemistry do not form a complete set, it is beneficial to design a modified scheme in which it is possible to obtain the **effective** matrix elements of the OEP operator in a **incomplete** auxiliary space. This can be achieved by minimizing the following objective function

$$Z[\{G_{\xi}^{(i)}\}] = \iint d\mathbf{r}_1 d\mathbf{r}_2 \frac{\left[v(\mathbf{r}_1) \phi_i(\mathbf{r}_1) - \sum_{\xi} G_{\xi}^{(i)} \varphi_{\xi}(\mathbf{r}_1) \right] \left[v(\mathbf{r}_2) \phi_i(\mathbf{r}_2) - \sum_{\xi} G_{\eta}^{(i)} \varphi_{\eta}(\mathbf{r}_1) \right]}{|\mathbf{r}_1 - \mathbf{r}_2|}$$

Thus requesting that

$$\frac{\partial Z[\{G_{\xi}^{(i)}\}]}{\partial G_{\mu}^{(i)}} = 0 \text{ for all } \mu$$

we find the coefficients $G_{\xi}^{(i)}$ to be

$$\mathbf{G}^{(i)} = \mathbf{b}^{(i)} \cdot \mathbf{A}^{-1}$$

where

$$b_{\eta}^{(i)} = (\eta || v_i)$$

$$A_{\eta\xi} = (\eta || \xi)$$

The symbol $||$ is to denote the operator r_{12}^{-1} and double integration over \mathbf{r}_1 and \mathbf{r}_2 . Thus, it is clear that in order to use this generalized density fitting scheme one must to compute two-centre electron repulsion integrals (implemented in [oepdev::ERI_1_1](#)) as well as four-centre asymmetric electron repulsion integrals of the type $(\alpha\beta\gamma||\eta)$ (implemented in [oepdev::ERI_3_1](#)).

Chapter 7

Implemented Models

7.1 Target Properties

Detailed list of models which is to be implemented in the OEPDev project is given below:

Table 1. Models subject to be implemented and analyzed within oep-dev.

Pauli energy	Induction energy	EET Coupling
EFP2-Pauli	EFP2-Induced Dipoles	TrCamm
Murrel et al.'s theory	Density Susceptibility	OEP-ET/HT
OEP-Murrel et al.'s		TDFI-TI
		FED
Exact (Stone's)	Exact (incl. CT)	Exact (ESD)

7.2 Target, Benchmark and Competing Models

The target models introduced in the Project shall be tested against the following benchmarks and compared with the following state-of-the-art models:

Table 2. Target models vs benchmarks and competitor models.

Target Model	Benchmarks	Competing Model
OEP-Murrel et al.'s	Murrel et al.'s	EFP2-Pauli
	Exact (Stone's)	
OEP-ET/HT + TrCamm	Exact (ESD)	TDFI-TI
	FED	FED
	TDFI-TI	
Density Susceptibility	Exact (incl. CT)	EFP2-Induced Dipoles

Chapter 8

Contributing to OEP-Dev

OepDev is a plugin to Psi4.

Therefore it should follow the programming etiquette of Psi4. Also, oep-dev has additional programming tips to make the code more versatile and easy in further development. Here, I emphasise on most important aspects regarding the **programming rules**.

8.1 Main routine and libraries

Oep-dev has only *one* source file in the plugin base directory, i.e., `main.cc`. This is the main driver routine that handles the functionality of the whole OEP testing platform: specifies options for Psi4 input file and implements test routines based on the options. Include files directly related to `main.cc` are stored in the `include` directory, where only header files are present. Options are specified in `include/oepdev_options.h` whereas macros and defines in `include/oepdev_files.h`. Other sources are stored in `MODULE/libNAME*` directories where `NAME` is the name of the library with sources and header files, whereas `MODULE` is the directory of the oep-dev module.

Things to remember:

1. **No other sources in base directory.** It is not permitted to place any new source or other files in the plugin base directory (i.e., where `main.cc` resides).
2. **Sources in library directories.** Any additional source code has to be placed in `oepdev/libNAME*` directory (either existing one or a new one; in the latter case remember to add the new `*.cc` files to `CMakeLists.txt` in the plugin base directory).
3. **Miscellanea in special directories.** If you want to add additional documentation, put it in the `doc` directory. If you want to add graphics, put it in the `images` directory.

8.2 Header files in libraries

Header files are handy in obtaining a quick glimpse of the functionality within certain library. Each library directory should contain at least one header file in oep-dev. However, header files can be problematic if not managed properly.

Things to remember:

1. **Header preprocessor variable.** Define the preprocessor variable specifying the existence of include of the particular header file. The format of such is

```
#ifndef MODULE_LIBRARY_HEADER.h
#define MODULE_LIBRARY_HEADER.h
// rest of your code goes here
#endif // MODULE_LIBRARY_HEADER.h
```

Last line is the **end** of the header file. The preprocessor variables represents the directory tree `oepdev/MODULE/LIBRARY/HEADER.h` structure (where `oepdev` is the base plugin directory). `MODULE` is the plugin module name (e.g. `oepdev`, the name of the module directory) `LIBRARY` is the name of the library (e.g. `libutil`, should be the same as library directory name) `HEADER` is the name of the header in library directory (e.g. `diis` for `diis.h` header file)

2. **Set module namespace.** To prevent naming clashes with other modules and with Psi4 it is important to operate in separate namespace (e.g. for a module).

```
namespace MODULE {
// your code goes here
} // EndNameSpace MODULE
```

For instance, all classes and functions in `oepdev` module are implemented within the namespace of the same label. Considering addition of other local namespaces within a module can also be useful in certain cases.

8.3 Environmental variables

Defining the set of intrinsic environmental variables can help in code management and conditional compilation. The oep-dev environmental variables are defined in `include/oepdev_files.h` file. Remember also about psi4 environmental variables defined in `psi4/psifiles.h` header. As a rule, the oep-dev environmental variable should have the following format:

```
OEPDEV_XXXX
```

where `XXXX` is the descriptive name of variable.

8.4 Documenting the code

Code has to be documented (at best at a time it is being created). The place for documentation is always in header files. Additional documentation can be also placed in source files. Leaving a chunk of code for a production run without documentation is unacceptable.

Use Doxygen style for documentation all the time. Remember that it supports markdown which can make the documentation even more clear and easy to understand. Additionally you can create a nice `.rst` documentation file for Sphinx program. If you are coding equations, always include formulae in the documentation!

Things to remember:

1. **Descriptions of classes, structures, global functions, etc.** Each programming object should have a description.
2. **Documentation for function arguments and return object.** Usage of functions and class methods should be explained by providing the description of all arguments (use `\param` and `\return` Doxygen keywords).
3. **One-line description of class member variables.** Any class member variable should be preceded by a one-liner documentation (starting from `///`).
4. **Do not be afraid of long names in the code.** Self-documenting code is a blessing!

8.5 Naming conventions

Naming is important because it helps to create more readable and clear self-documented code.

Some loose suggestions:

1. **Do not be afraid of long names in the code, but avoid redundancy.** Examples of good and bad names: good name: `get_density_matrix`; bad name: `get_matrix`. Unless there is only one type of matrix a particular objects can store, `matrix` is not a good name for a getter method. good name: `class Wavefunction`, bad name: `class WFN` good name: `int numberOfErrorVectors`, bad name: `int nvec`, bad name: `the_number_of_error_vectors` good name: `class EFPotential`, probably bad name: `class EffectiveFragmentPotential`. The latter might be understood by some people as a class that inherits from `EffectiveFragment` class. If it is not the case, compromise between abbreviation and long description is OK.
2. **Short names are OK in special situations.** In cases meaning of a particular variable is obvious and it is frequently used in the code locally, it can be named shortly. Examples are: `i` when iterating `no` number of occupied orbitals, `nv` number of virtual orbitals, etc.
3. **Clumped names for variables and dashed names for functions.** Try to distinguish between variable name like `sizeofOEPTypelist` and a method name `get_matrix()` (neither `size_of_OEP_type_list`, nor `getMatrix()`). This is little bit cosmetics, but helps in managing the code when it grows.
4. **Class names start from capital letter.** However, avoid only capital letters in class names, unless it is obvious. Avoid also dashes in class names (they are reserved for global functions and class methods). Examples: good name: `DIISManager`, bad name: `DIIS`. good name: `EETCouplingSolver`, bad name: `EETSolver`, very bad: `EET`.

8.6 Track timing when evaluating the code

It is useful to track time elapsed for performing a particular task by a computer. For this, use `psi::timer_on` and `psi::timer_off` functions defined in `psi4/libqt/qt.h`. Psi4 always generates the report file `timer.dat` that contains all the defined timings. For example,

```
#include "psi/libqt/qt.h"
psi::timer_on("OEP      E(Paul) Murrell-et al S1  ");
// Your code goes here
psi::timer_off("OEP      E(Paul) Murrell-et al S1  ");
```

To maintain the printout in a neat form, the timing associated with the OEPDev code should be generated via `misc/python/timing.py` utility script.

8.7 Use Object-Oriented Programming

Try to organise your creations in objects having special relationships and data structures. Encapsulation helps in producing self-maintaining code and is much easier to use. Use:

- **factory design** for creating objects
- **container design** for designing data structures
- **polymorphysm** when dealing with various flavours of one particular feature in the data structure

Note: In Psi4, factories are frequently implemented as static methods of the base classes, for example `psi::BasisSet::build` static method. It can be followed when building object factories in oep-dev too.

Chapter 9

Advanced Usage

This section is addressed for advanced users.

Make sure you have first read [the introduction](#) before proceeding.

9.1 Installation

9.1.1 Preparing Psi4

OEPDev is a Psi4 plugin. It requires

- Psi4, at least 1.2 version (git commit `9d4a61c`). Has to be modified (see below)
- Eigen3, any newer version

Note

Before compiling, make sure EFP is enabled in `CMakeLists.txt` (now it is not used in OEPDev but maybe in the future it would).

Recently, Psi4 introduced API visibility management. Only certain Psi4 classes and functions are *exposed* in the `core.so` library, that is further linked to Psi4 plugin shared library. Due to this reason, not all Psi4 functionalities can be directly used from outside Psi4. In order to access local API of Psi4 (also used in the OEPDev code) slight modification of Psi4 code and concomitant rebuild is necessary.

In order to expose local API used by OEPDev and hidden within Psi4 1.2, two types of small modifications are necessary:

- M1: add `PSI_API` macro after required class or function declaration in header file
- M2: add `#include "psi4/pragma.h"` line at the include section of an appropriate header file

Modification M1 is obligatory for all affected files whereas modification M2 needs to be done only in headers that do not have "psi4/pragma.h" included explicitly or implicitly. The list of Psi4 header files along with the respective changes that need to be done are listed in the table below:

Psi4 Header File	Psi4 Class	Required Changes
libfunctional/superfunctional.h	Superfunctional	M1
libscf_solver/hf.h	HF	M1
libscf_solver/rhf.h	RHF	M1
libcubeprop/csg.h	CubicScalarGrid	M1
libmints/onebody.h	OneBodyAOInt	M1
libmints/potential.h	PotentialInt	M1
libmints/multipoles.h	MultipoleInt	M1
libmints/multipolesymmetry.h	MultipoleSymmetry	M1
libmints/fjt.h	Taylor_Fjt	M1
libmints/fjt.h	Fjt	M1
libmints/oeprop.h	OEProp	M1, M2
libmints/gshell.h	GaussianShell	M1, M2

To quickly apply these modifications, use the patch files stored in `misc/patch` directory. Please make sure to use a proper patch for a chosen Psi4 version.

9.1.2 Compilation

After all the above changes have been done in Psi4 (followed by its rebuild) compile the OEPDev code by running `compile` script. Make sure Eigen3 path is set to environment variable `EIGEN3_INCLUDE_DIR` (instructions will appear on the screen). After compilation is successful, run `ctest` to check if the code works fine.

Note

It may happen, that during code development there will be symbol lookup error when importing `oepdev.so` (in such case OEPDev compiles without error but Python cannot import the module `oepdev`). In such circumstance, probably there some local Psi4 feature that is needed in OEPDev is not exposed by `PSI_API` macro. To fix this, run `c++filt [name]` where `[name]` is the mangled undefined symbol. This will show you which Psi4 class or function is not exposed and requires `PSI_API` (change M1 and perhaps M2 too). Such change requires Psi4 rebuild and recompilation of OEPDev code. In any case, please contact me and report new undefined symbol (blasiak.bartosz@gmail.com).

9.2 OEPDev Code Structure

As a plugin to Psi4, OEPDev consists of the `main.cc` file with the plugin main routine, `include/oepdev_options.h` specifying the options of the plugin, `include/oepdev_files.h` defining all global macros and environmental variables, as well as the `oepdev` directory. The latter contains the actual OEPDev code that is divided into several subdirectories called `modules`.

9.2.1 Main Routine

Before the actual OEPDev calculations are started, the wavefunction of the input molecular aggregate is computed by Psi4. See the plugin driver script `pymodule.py` for more details on how the calculation environment is initialized. Subsequently, one out of four types of target operations can be performed by the program:

1. `OEP_BUILD` - Compute the OEP effective parameters for one molecule.
2. `DMATPOL` - Compute the generalized density matrix susceptibility tensors (DMS's) for one molecule.
3. `SOLVER` - Perform calculations for a molecular aggregate. As for now, only dimers are handled.
4. `TEST` - Perform the testing routine.

The first two modes are single molecule calculations. `OEP_BUILD` uses the `oepdev::OEPotential::build` static factory to create OEP objects whereas `DMATPOL` uses the `oepdev::GenEffParFactory::build` static factory to create generalized effective fragment parameters (GEFP's) for polarization.

Note

In the future, `OEP_BUILD` will be handled also by `oepdev::GenEffParFactory::build` since OEP parameters are part of the GEFP's.

`SOLVER` requires at least molecular dimer and the `oepdev::WavefunctionUnion` object (being the Hartree product of the unperturbed monomer wavefunctions) is constructed at the beginning, which is then passed to the `oepdev::OEPDevSolver::build` static factory. `TEST` can refer to single- or multiple-molecule calculations, whereby each of the testing routines is listed in the `cmake/CCTestTestfile.cmake.in` file.

9.2.2 Modules

The source code is distributed into directories called modules:

- `liboep`
- `libgefp`
- `libsolver`
- `libints`
- `libpsi`
- `lib3d`
- `libutil`
- `libtest`

See Modules for a detailed description of each of the modules.

9.3 OEPDev Classes: Overview

9.3.1 OEP Module

The OEP module located in `oepdev/liboep` consists of the following abstract bases:

- `oepdev::OEPotential` implementing the OEP,
- `oepdev::GeneralizedDensityFit` implementing the GDF technique.

Each of the bases contains static factory method called `build` that creates instances of chosen subclasses. The module contains also a structure `oepdev::OEPTyp` which is a container storing all the data associated with a particular OEP: type name, dimensions, OEP coefficients and whether is density-fitted or not.

9.3.1.1 OEPPotential

It is a container and computer class of OEP. Among others, the most important public method is `oepdev::OEPotential::compute` which computes all the OEP's (by iterating over all possible OEP types within a chosen OEP subclass or category). OEP's can be extracted by `oepdev::OEPotential::oep` method, for instance. From protected attributes, each OE-Potential instance stores blocks of the LCAO-MO matrices associated with the occupied (`cOcc_`) and virtual (`cVir_`) MO's. It also contains the pointers to the primary, auxiliary and intermediate basis sets (`primary_`, `auxiliary_` and `intermediate_`, accordingly). Usage example:

```
#include "oepdev/liboep/oep.h"
oep = oepdev::OEPotential::build("ELECTROSTATIC ENERGY", wfn, options);
oep->compute();
oep->write_cube("V", "oep_cube_file");
```

So far, four OEPotential subclasses are implemented, from which `oepdev::ElectrostaticEnergyOEP` and `oepdev::RepulsionEnergyOEPotential` are fully operative, while the rest is under development.

9.3.1.2 GeneralizedDensityFit

9.3.2 GEFP Module

9.3.2.1 GenEffPar

9.3.2.2 GenEffParFactory

9.3.2.3 GenEffFrag

9.3.3 OEPDev Solver Module

9.3.3.1 OEPDevSolver

9.4 Developing OEP's

OEP's are implemented in a suitable subclass of the `oepdev::OEPotential` base. Due to the fact that OEP's can be density-based or ESP-based, the classes `oepdev::GeneralizedDensityFit` as well as `oepdev::ESPSolver` are usually necessary in the implementations. Handling the one-electron integrals (OEI's) and the two-electron integrals (ERI's) in AO basis is implemented in `oepdev::IntegralFactory`. In particular, potential integrals evaluated at arbitrary centres can be accessed by using the `oepdev::PotentialInt` instances. Useful iterators for looping over AO ERI's the `oepdev::ShellCombinationsIterator` and `oepdev::AOIntegralsIterator` classes. Transformations of OEI's to MO basis can be easily achieved by transforming AO integral matrices by `cOcc_` and `cVir_` members of `OEPotential` instances, e.g., by using the `psi::Matrix::doublet` or `psi::Matrix::triplet` static methods. Transformations of ERI's to MO basis can be performed by using the `psi4/libtrans/integraltransform.h` library.

It is recommended that the implementation of all the new OEP's follows the following steps:

1. **Write the class framework.** This includes choosing a proper name of a `OEPotential` subclass, sketching the constructors and a destructor, and all the necessary methods.
2. **Implement OEP types.** Each type of OEP is implemented, including the 3D vector field in case ESP-based OEP's are of use.
3. **Update base factory method.** Add appropriate entries in the `oepdev::OEPotential::build` static factory method.

Below, we shall go through each of these steps separately and discuss them in detail.

9.4.1 Drafting an OEP Subclass

This stage is the design of the overall framework of OEP subclass. The name should end with `OEPotential` to maintain the convention used so far. The template for the header file definition can be depicted as follows:

```
class SampleOEPotential : public OEPotential
{
public:
    // Purely ESP-based OEP's
    SampleOEPotential(SharedWavefunction wfn, Options& options);

    // GDF-based OEP's
    SampleOEPotential(SharedWavefunction wfn, SharedBasisSet auxiliary, SharedBasisSet intermediate,
        Options& options);

    // Necessary destructor
    virtual ~SampleOEPotential();

    // Necessary computer
    virtual void compute(const std::string& oepType) override;

    // Necessary computer
    virtual void compute_3D(const std::string& oepType,
        const double& x, const double& y, const double& z, std::shared_ptr<psi::Vector>
        & v) override;
    // Necessary printer
```

```

    virtual void print_header() const override;

private:
    // Set defaults - good practice
    void common_init();

    // Auxiliary computers - exemplary
    double compute_3D_sample_V(const double& x, const double& y, const double& z);
};

```

The constructors need to call the abstract base constructor and then specialized initializations. It is a good practice to put the specialized common initializers in a separate private method `common_init` (which is a convention in Psi4 and is adopted also in OEPDev). For instance, the exemplary constructor is show below:

```

SampleOEPotential::SampleOEPotential(SharedWavefunction wfn,
                                     SharedBasisSet auxiliary, SharedBasisSet intermediate, Options&
                                     options)
: OEPotential(wfn, auxiliary, intermediate, options)
{
    common_init();
}

void SampleOEPotential::common_init()
{
    int n1 = wfn->Ca_subset("AO", "OCC")->ncol();
    int n2 = auxiliary->nbf();
    int n3 = wfn->molecule()->natom();

    SharedMatrix mat_1 = std::make_shared<psi::Matrix>("G(S^{-1})", n2, n1);
    SharedMatrix mat_2 = std::make_shared<psi::Matrix>("G(S^{-2})", n3, n1);

    OEPTypes type_1 = {"Murrell-etal.S1", true, n1, mat_1};
    OEPTypes type_2 = {"Otto-Ladik.S2", false, n1, mat_2};

    oepTypes_[type_1.name] = type_1;
    oepTypes_[type_2.name] = type_2;
}

```

Note that the `oepdev::OEPotential::oepTypes_` attribute, which is a `std::map` of structures `oepdev::OEPTypes`, is initialized here. All the OEP types need to be stated in the constructors. Destructors usually call nothing, unless dynamically allocated memory is also of use.

It is also a good practice to already sketch the `compute` method here by adding certain private computers, like in the example below:

```

void SampleOEPotential::compute(const std::string& oepType)
{
    if (oepType == "Murrell-etal.S1") this->compute_murrell_etal_s1(); // calls private method
    else if (oepType == "Otto-Ladik.S2") this->compute_otto_ladik_s2(); // calls private method
    else throw psi::PSIEXCEPTION("OEPDEV: Error. Incorrect OEP type specified!\n"); // for safety
}

void SampleOEPotential::compute_murrell_etal_s1()
{
    psi::timer_on ("OEP      E(Paul) Murrell-etal S1  ");
    /* Your implementation goes here */
    psi::timer_off("OEP      E(Paul) Murrell-etal S1  ");
}

```

9.4.1.1 Implementing OEP Types

Implementation of the inner body of `compute` method requires populating the members of `oepTypes_` with data. This means, that for each OEP type there has to be a specific implementation of OEP parameters. GDF-based OEP's need to create the `psi::Matrix` with OEP

parameters and put them into `oepTypes_`. In the case of ESP-based OEP's `compute_3D` method has to be additionally implemented before `compute` is fully functional. To implement `compute_3D`, `oepdev::OEPotential::make_oeps3d` method is of high relevance: it creates `oepdev::OEPotential3D<T>` instances, where `T` is the OEP subclass. These instances are `oepdev::Field3D` objects that define OEP's in 3D Euclidean space. For example,

```
void SampleOEPotential::compute_otto_ladik_s2()
{
    // Switch on timer
    psi::timer.on("OEP      E(Paul) Otto-Ladik S2      ");

    // Create 3D field, automated through 'make_oeps3d'. Requires 'compute_3D' implementation.
    std::shared_ptr<OEPotential3D<OEPotential>> oeps3d = this->make_oeps3d("Otto-Ladik.S2");
    oeps3d->compute();

    // Perform ESP fit to get OEP effective charges
    ESPSolver esp(oeps3d);
    esp.set_charge_sums(0.5);
    esp.compute();

    // Put the OEP coefficients into 'oepTypes_'
    for (int i=0; i<esp.charges()->nrow(); ++i) {
        for (int o=0; o<oepTypes_["Otto-Ladik.S2"].n; ++o) {
            oepTypes_["Otto-Ladik.S2"].matrix->set(i, o, esp.charges()->get(i, o));
        }
    }

    // Switch off timer
    psi::timer.off("OEP      E(Paul) Otto-Ladik S2      ");
}

// Necessary implementation for 'make_oeps3d' to work
void SampleOEPotential::compute_3D(const std::string& oepType, const double& x, const double& y, const
double& z, std::shared_ptr<psi::Vector>& v)
{
    // Loop over all possibilities for OEP types and exclude illegal names
    if (oepType == "Otto-Ladik.S2") {

        // this computes the actual values of OEP = v(x,y,z) and stores it in 'vec_otto_ladik_s2_'
        this->compute_3D_otto_ladik_s2(x, y, z);

        // Assign final value to the buffer vector
        for (int o = 0; o < oepTypes_["Otto-Ladik.S2"].n; ++o) v->set(o, vec_otto_ladik_s2_[o]);
    }
    else if (oepType == "Murrell-et.al.S1") { /* Even if it is not ESP-based OEP, this line is necessary */}
    else {
        throw psi::PSIEXCEPTION("OEPDEV: Error. Incorrect OEP type specified!\n"); // Safety
    }
}
```

Note that `make_oeps3d` is not overridable and is fully defined in the base. Do not call `oepdev::OEPotential3D` constructors in the `OEPotential` subclass (it can be done only from the level of the abstract base where all the pointers are dynamically converted to an appropriate data type due to polymorphism)!

9.4.1.2 Abstract Base

Chapter 10

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Chapter 11

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Class Index

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Chapter 15

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15.1 File List

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Chapter 16

Module Documentation

16.1 The Generalized One-Electron Potentials Library

Implements the goal of this project: The Generalized One-Electron Potentials (OEP's). You will find here OEP's for computation of Pauli repulsion energy, charge-transfer energy and others. The routines for the generalized density fitting are also implemented here. Located at `oepdev/liboep`.

Classes

- struct `oepdev::OEType`
Container to handle the type of One-Electron Potentials.
- class `oepdev::OEPotential`
Generalized One-Electron Potential: Abstract base.
- class `oepdev::ElectrostaticEnergyOEPotential`
Generalized One-Electron Potential for Electrostatic Energy.
- class `oepdev::RepulsionEnergyOEPotential`
Generalized One-Electron Potential for Pauli Repulsion Energy.
- class `oepdev::ChargeTransferEnergyOEPotential`
Generalized One-Electron Potential for Charge-Transfer Interaction Energy.
- class `oepdev::EETCouplingOEPotential`
Generalized One-Electron Potential for EET coupling calculations.
- class `oepdev::GeneralizedDensityFit`
Generalized Density Fitting Scheme. Abstract Base.
- class `oepdev::SingleGeneralizedDensityFit`
Generalized Density Fitting Scheme - Single Fit.
- class `oepdev::DoubleGeneralizedDensityFit`
Generalized Density Fitting Scheme - Double Fit.

16.1.1 Detailed Description

16.2 The OEPDev Solver Library

Implementations of various solvers for molecular properties as a functions of unperturbed monomeric wavefunctions. This is the place all target OEP-based models are implemented and compared with benchmark and competitor models. Located at `oepdev/libsolver`.

Classes

- class `oepdev::OEPDevSolver`
Solver of properties of molecular aggregates. Abstract base.
- class `oepdev::ElectrostaticEnergySolver`
Compute the Coulombic interaction energy between unperturbed wavefunctions.
- class `oepdev::RepulsionEnergySolver`
Compute the Pauli-Repulsion interaction energy between unperturbed wavefunctions.
- class `oepdev::ChargeTransferEnergySolver`
Compute the Charge-Transfer interaction energy between unperturbed wavefunctions.
- class `oepdev::EETCouplingSolver`
Compute the EET coupling energy between unperturbed wavefunctions.
- class `oepdev::TIData`
Transfer Integral EET Data.

16.2.1 Detailed Description

16.3 The Generalized Effective Fragment Potentials Library

Implements the GEFP method, the far goal of the OEPDev project. Here you will find the containers for GEFP parameters, the density matrix susceptibility tensors and GEFP solvers. Located at `oepdev/libgefp`.

Classes

- class `oepdev::GenEffPar`
Generalized Effective Fragment Parameters. Container Class.
- class `oepdev::GenEffFrag`
Generalized Effective Fragment. Container Class.
- class `oepdev::GenEffParFactory`
Generalized Effective Fragment Factory. Abstract Base.
- class `oepdev::PolarGEFactory`
Polarization GEFP Factory. Abstract Base.
- class `oepdev::AbInitioPolarGEFactory`
Polarization GEFP Factory from First Principles. Hartree-Fock Approximation.
- class `oepdev::FFAbInitioPolarGEFactory`
Polarization GEFP Factory from First Principles: Finite-Difference Model. Arbitrary level of theory.
- class `oepdev::GeneralizedPolarGEFactory`
Polarization GEFP Factory with Least-Squares Parameterization.
- class `oepdev::UniformEFieldPolarGEFactory`
Polarization GEFP Factory with Least-Squares Parameterization.
- class `oepdev::NonUniformEFieldPolarGEFactory`
Polarization GEFP Factory with Least-Squares Parameterization.
- class `oepdev::LinearUniformEFieldPolarGEFactory`
Polarization GEFP Factory with Least-Squares Parameterization.
- class `oepdev::QuadraticUniformEFieldPolarGEFactory`
Polarization GEFP Factory with Least-Squares Parameterization.
- class `oepdev::LinearNonUniformEFieldPolarGEFactory`
Polarization GEFP Factory with Least-Squares Parameterization.
- class `oepdev::QuadraticNonUniformEFieldPolarGEFactory`
Polarization GEFP Factory with Least-Squares Parameterization.
- class `oepdev::LinearGradientNonUniformEFieldPolarGEFactory`
Polarization GEFP Factory with Least-Squares Parameterization.
- class `oepdev::QuadraticGradientNonUniformEFieldPolarGEFactory`
Polarization GEFP Factory with Least-Squares Parameterization.
- class `oepdev::UnitaryTransformedMOPolarGEFactory`
Polarization GEFP Factory with Least-Squares Scaling of MO Space.

16.3.1 Detailed Description

16.4 The Integral Package Library

Implementations of various two-, three- or four-centre two-body electron repulsion integrals via utilizing the McMurchie-Davidson recurrence scheme. Located at `oepdev/libints` and `oepdev/libpsi`.

Classes

- class `oepdev::TwoElectronInt`
General Two Electron Integral.
- class `oepdev::ERI_1_1`
2-centre ERI of the form $(a|O(2)|b)$ where $O(2) = 1/r^{12}$.
- class `oepdev::ERI_2_2`
4-centre ERI of the form $(ab|O(2)|cd)$ where $O(2) = 1/r^{12}$.
- class `oepdev::ERI_3_1`
4-centre ERI of the form $(abc|O(2)|d)$ where $O(2) = 1/r^{12}$.
- class `oepdev::TwoBodyAOInt`
- class `oepdev::IntegralFactory`
Extended `IntegralFactory` for computing integrals.
- class `oepdev::PotentialInt`
Computes potential integrals.

Macros

- `#define D1_INDEX(x, i, n) ((81*(x))+(9*(i))+(n))`
Get the index of McMurchie-Davidson-Hermite D1 coefficient stored in the `mdh_buffer_`, that is attributed to the x Cartesian coordinate from angular momentum i of function 1, and the Hermite index n .
- `#define D2_INDEX(x, i, j, n) ((1377*(x))+(153*(i))+(17*(j))+(n))`
Get the index of McMurchie-Davidson-Hermite D2 coefficient stored in the `mdh_buffer_`, that is attributed to the x Cartesian coordinate from angular momenta i, j of function 1 and 2, and the Hermite index n .
- `#define D3_INDEX(x, i, j, k, n) ((18225*(x))+(2025*(i))+(225*(j))+(25*(k))+(n))`
Get the index of McMurchie-Davidson-Hermite D3 coefficient stored in the `mdh_buffer_`, that is attributed to the x Cartesian coordinate from angular momenta i, j and k of function 1, 2 and 3, and the Hermite index n .
- `#define R_INDEX(n, l, m, j) ((14739*(n))+(867*(l))+(51*(m))+(j))`
Get the index of McMurchie-Davidson R coefficient stored in the `mdh_buffer_R_` from angular momenta n, l and m and the Boys index j .

Functions

- double `oepdev::d_N_n1_n2` (int N, int n1, int n2, double PA, double PB, double aP)
Compute McMurchie-Davidson-Hermite (MDH) coefficient for binomial expansion.
- void `oepdev::make_mdh_D1_coeff` (int n1, double aPd, double *buffer)
Compute the McMurchie-Davidson-Hermite coefficients for monomial expansion.
- void `oepdev::make_mdh_D2_coeff` (int n1, int n2, double aPd, double *PA, double *PB, double *buffer)
Compute the McMurchie-Davidson-Hermite coefficients for binomial expansion.
- void `oepdev::make_mdh_D3_coeff` (int n1, int n2, int n3, double aPd, double *PA, double *PB, double *PC, double *buffer)
Compute the McMurchie-Davidson-Hermite coefficients for trinomial expansion.
- void `oepdev::make_mdh_D2_coeff_explicit_recursion` (int n1, int n2, double aP, double *PA, double *PB, double *buffer)
Compute the McMurchie-Davidson-Hermite coefficients for binomial expansion by explicit recursion. This function makes the same changes to buffers as `oepdev::make_mdh_D2_coeff`, but implements it through explicit recursion by calls to `oepdev::d_N_n1_n2`. Therefore, it is slightly slower. Here for debugging purposes.
- void `oepdev::make_mdh_R_coeff` (int N, int L, int M, double alpha, double a, double b, double c, double *F, double *buffer)
Compute the McMurchie-Davidson R coefficients.

16.4.1 Detailed Description

Here, we define the primitive Gaussian type functions (GTO's)

$$\begin{aligned}\phi_i(\mathbf{r}) &\equiv x_A^{n_1} y_A^{l_1} z_A^{m_1} e^{-\alpha_1 r_A^2} \\ \phi_j(\mathbf{r}) &\equiv x_B^{n_2} y_B^{l_2} z_B^{m_2} e^{-\alpha_2 r_B^2} \\ \phi_k(\mathbf{r}) &\equiv x_C^{n_3} y_C^{l_3} z_C^{m_3} e^{-\alpha_3 r_C^2}\end{aligned}$$

in which $\mathbf{r}_A \equiv \mathbf{r} - \mathbf{A}$ and so on. \mathbf{A} is the centre of the GTO, α_1 its exponent, whereas n_1, l_1, m_1 the Cartesian angular momenta, with the total angular momentum $\theta_1 = n_1 + l_1 + m_1$.

In OEPDev implementations, the following definition shall be in use:

$$\begin{aligned}\mathbf{P} &\equiv \frac{\alpha_1 \mathbf{A} + \alpha_2 \mathbf{B}}{\alpha_1 + \alpha_2} \\ \mathbf{Q} &\equiv \frac{\alpha_3 \mathbf{C} + \alpha_4 \mathbf{D}}{\alpha_3 + \alpha_4} \\ \mathbf{R} &\equiv \frac{\alpha_1 \mathbf{A} + \alpha_2 \mathbf{B} + \alpha_3 \mathbf{C}}{\alpha_1 + \alpha_2 + \alpha_3} \\ \alpha_P &\equiv \alpha_1 + \alpha_2 \\ \alpha_Q &\equiv \alpha_3 + \alpha_4 \\ \alpha_R &\equiv \alpha_1 + \alpha_2 + \alpha_3\end{aligned}$$

The unnormalized products of primitive GTO's are denoted here as

$$\begin{aligned}[ij] &\equiv \phi_i(\mathbf{r})\phi_j(\mathbf{r}) \\ [ijk] &\equiv \phi_i(\mathbf{r})\phi_j(\mathbf{r})\phi_k(\mathbf{r})\end{aligned}$$

16.4.2 Hermite Operators

It is convenient to define

$$\Lambda_j(x_P; \alpha_P) \equiv \left(\frac{\partial}{\partial P_x} \right)^j = \alpha_P^{j/2} H_j(\sqrt{\alpha_P} x_P)$$

where $H_j(x)$ is the Hermite polynomial of order j evaluated at x . Introduction of the above Hermite operator can be used by invoking the recurrence relationship due to Hermite polynomial properties:

$$x_A \Lambda_j(x_P; \alpha_P) = j \Lambda_{j-1} + |\mathbf{P} - \mathbf{A}|_x \Lambda_j + \frac{1}{2\alpha_P} \Lambda_{j+1}$$

This can be directly used to derive very useful McMurchie-Davidson-Hermite coefficients as expansion coefficients of the polynomial expansions.

16.4.2.1 Polynomial Expansions as Hermite Series

By using the previous relation, it is possible to express the following expansions in Hermite series:

$$\begin{aligned}x_A^{n_1} &= \sum_{N=0}^{n_1} d_N^{n_1} \Lambda_N(x_A; \alpha_A) \\ x_A^{n_1} x_B^{n_2} &= \sum_{N=0}^{n_1+n_2} d_N^{n_1 n_2} \Lambda_N(x_P; \alpha_P) \\ x_A^{n_1} x_B^{n_2} x_C^{n_3} &= \sum_{N=0}^{n_1+n_2+n_3} d_N^{n_1 n_2 n_3} \Lambda_N(x_R; \alpha_R)\end{aligned}$$

The recurrence relationships can be easily found and they read

$$d_N^{n_1+1} = \frac{1}{2\alpha_A} d_{N-1}^{n_1} + (N+1) d_{N+1}^{n_1}$$

as well as

$$\begin{aligned}d_N^{n_1+1, n_2} &= \frac{1}{2\alpha_P} d_{N-1}^{n_1 n_2} + |\mathbf{P} - \mathbf{A}|_x d_N^{n_1 n_2} + (N+1) d_{N+1}^{n_1 n_2} \\ d_N^{n_1, n_2+1} &= \frac{1}{2\alpha_P} d_{N-1}^{n_1 n_2} + |\mathbf{P} - \mathbf{B}|_x d_N^{n_1 n_2} + (N+1) d_{N+1}^{n_1 n_2}\end{aligned}$$

and

$$\begin{aligned} d_N^{n_1+1,n_2,n_3} &= \frac{1}{2\alpha_R} d_{N-1}^{n_1 n_2 n_3} + |\mathbf{R} - \mathbf{A}|_x d_N^{n_1 n_2 n_3} + (N+1) d_{N+1}^{n_1 n_2 n_3} \\ d_N^{n_1,n_2+1,n_3} &= \frac{1}{2\alpha_R} d_{N-1}^{n_1 n_2 n_3} + |\mathbf{R} - \mathbf{B}|_x d_N^{n_1 n_2 n_3} + (N+1) d_{N+1}^{n_1 n_2 n_3} \\ d_N^{n_1,n_2,n_3+1} &= \frac{1}{2\alpha_R} d_{N-1}^{n_1 n_2 n_3} + |\mathbf{R} - \mathbf{C}|_x d_N^{n_1 n_2 n_3} + (N+1) d_{N+1}^{n_1 n_2 n_3} \end{aligned}$$

respectively. The first elements are given by

$$\begin{aligned} d_0^0 &= 1 \\ d_0^{00} &= 1 \\ d_0^{000} &= 1 \end{aligned}$$

By using the above formalisms, it is straightforward to express the doublet of primitive GTO's as

$$[ij] = E_{ij} \sum_{N=0}^{n_1+n_2} \sum_{L=0}^{l_1+l_2} \sum_{M=0}^{m_1+m_2} d_N^{n_1 n_2} d_L^{l_1 l_2} d_M^{m_1 m_2} \Lambda_N(x_P) \Lambda_L(y_P) \Lambda_M(z_P) e^{-\alpha_P r_P^2}$$

Analogously, the triplet of primitive GTO's is given by

$$[ijk] = E_{ijk} \sum_{N=0}^{n_1+n_2+n_3} \sum_{L=0}^{l_1+l_2+l_3} \sum_{M=0}^{m_1+m_2+m_3} d_N^{n_1 n_2 n_3} d_L^{l_1 l_2 l_3} d_M^{m_1 m_2 m_3} \Lambda_N(x_R) \Lambda_L(y_R) \Lambda_M(z_R) e^{-\alpha_R r_R^2}$$

The multiplicative constants are given by

$$\begin{aligned} E_{ij}(\alpha_1, \alpha_2) &= \exp \left[-\frac{\alpha_1 \alpha_2}{\alpha_1 + \alpha_2} |\mathbf{A} - \mathbf{B}|^2 \right] \\ E_{ijk}(\alpha_1, \alpha_2, \alpha_3) &= \exp \left[-\frac{\alpha_1 \alpha_2}{\alpha_1 + \alpha_2} |\mathbf{A} - \mathbf{B}|^2 \right] \exp \left[-\frac{(\alpha_1 + \alpha_2) \alpha_3}{\alpha_1 + \alpha_2 + \alpha_3} |\mathbf{P} - \mathbf{C}|^2 \right] \end{aligned}$$

16.4.3 One-Body Integrals over Hermite Functions

The fundamental Hermite integrals that appear during computations of any kind of one-body integrals over GTO's are as follows

$$[NLM|\Theta(1)] \equiv \int d\mathbf{r}_1 \Theta(\mathbf{r}_1) \Lambda_N(x_{1P}; \alpha_P) \Lambda_L(y_{1P}; \alpha_P) \Lambda_M(z_{1P}; \alpha_P) e^{-\alpha_P r_{1P}^2}$$

It immediately follows that the overlap, dipole, quadrupole and potential integrals are given as

$$\begin{aligned} [NLM|1] &= \delta_{N0} \delta_{L0} \delta_{M0} \left(\frac{\pi}{\alpha_P} \right)^{3/2} \\ [NLM|x_C] &= [\delta_{N1} + |\mathbf{P}\mathbf{C}|_x \delta_{N0}] \delta_{L0} \delta_{M0} \left(\frac{\pi}{\alpha_P} \right)^{3/2} \\ [NLM|x_C^2] &= \left[2\delta_{N2} + 2|\mathbf{P}\mathbf{C}|_x \delta_{N1} + \left(|\mathbf{P}\mathbf{C}|_x^2 + \frac{1}{2\alpha_P} \right) \delta_{N0} \right] \delta_{L0} \delta_{M0} \left(\frac{\pi}{\alpha_P} \right)^{3/2} \\ [NLM|x_C y_C] &= (\delta_{N1} + |\mathbf{P}\mathbf{C}|_x \delta_{N0}) (\delta_{L1} + |\mathbf{P}\mathbf{C}|_y \delta_{L0}) \delta_{M0} \left(\frac{\pi}{\alpha_P} \right)^{3/2} \\ [NLM|r_C^{-1}] &= \frac{2\pi}{\alpha_P} R_{NLM} \end{aligned}$$

The coefficients R_{NLM} are discussed in separate section below.

16.4.4 Two-Body Integrals over Hermite Functions

The fundamental Hermite integrals that appear during computations of any kind of two-electron integrals over GTO's are as follows

$$[N_1 L_2 M_2 | N_2 L_2 M_2] \equiv \iiint d\mathbf{r}_1 d\mathbf{r}_2 \Lambda_{N_1}(x_{1P}; \alpha_P) \Lambda_{L_1}(y_{1P}; \alpha_P) \Lambda_{M_1}(z_{1P}; \alpha_P) \Lambda_{N_2}(x_{2Q}; \alpha_Q) \Lambda_{L_2}(y_{2Q}; \alpha_Q) \Lambda_{M_2}(z_{2Q}; \alpha_Q)$$

The above formula dramatically reduces to the following

$$[N_1 L_2 M_2 | N_2 L_2 M_2] = \lambda (-)^{N_2+L_2+M_2} R_{N_1+N_2, L_1+L_2, M_1+M_2}$$

with

$$\lambda \equiv \frac{2\pi^{5/2}}{\alpha_P \alpha_Q \sqrt{\alpha_P + \alpha_Q}}$$

To compute the $R_{N_1+N_2, L_1+L_2, M_1+M_2}$ coefficients, the parameter T is given by

$$T = \frac{\alpha_P \alpha_Q}{\alpha_P + \alpha_Q} |\mathbf{P} - \mathbf{Q}|^2$$

16.4.5 The R(N,L,M) Coefficients

The R coefficients are defined as

$$R_{NLM} \equiv \left(\frac{\partial}{\partial a} \right)^N \left(\frac{\partial}{\partial b} \right)^L \left(\frac{\partial}{\partial c} \right)^M \int_0^1 e^{-Tu^2} du$$

with

$$T \equiv \alpha (a^2 + b^2 + c^2)$$

By extending the above definition to more general

$$R_{NLMj} \equiv (-\sqrt{\alpha})^{N+L+M} (-2\alpha)^j \int_0^1 u^{N+L+M+2j} H_N(au\sqrt{\alpha}) H_L(bu\sqrt{\alpha}) H_M(cu\sqrt{\alpha}) e^{-Tu^2} du$$

one can see that

$$R_{000j} = (-2\alpha)^j F_j(T)$$

The Boys function is here given by

$$F_j(T) \equiv \int_0^1 u^{2j} e^{-Tu^2} du$$

and its efficient implementation can be discussed elsewhere. In Psi4, `psi::Taylor.Fjt` class is used for this purpose.

Now, it is possible to show that the following recursion relationships are true:

$$R_{0,0,M+1,j} = cR_{0,0,M,j+1} + MR_{0,0,M-1,j+1}$$

$$R_{0,L+1,M,j} = bR_{0,L,M,j+1} + LR_{0,L-1,M,j+1}$$

$$R_{N+1,L,M,j} = aR_{N,L,M,j+1} + NR_{N-1,L,M,j+1}$$

This scheme is implemented in OEPDev.

16.4.6 Function Documentation

16.4.6.1 `d.N.n1.n2()`

```
double oepdev::d.N.n1.n2 (
    int N,
    int n1,
    int n2,
    double PA,
    double PB,
    double aP )
```

Parameters

<i>N</i>	- increment in the summation of MDH series
<i>n1</i>	- angular momentum of first function
<i>n2</i>	- angular momentum of second function
<i>PA</i>	- cartesian component of P-A distance
<i>PB</i>	- cartesian component of P-B distance
<i>aP</i>	- free parameter of MDH expansion

Returns

the McMurchie-Davidson-Hermite coefficient

16.4.6.2 `make_mdh.D1.coeff()`

```
void oepdev::make_mdh.D1.coeff (
    int n1,
    double aPd,
    double * buffer )
```

Parameters

<i>n1</i>	- angular momentum of first function
<i>aPd</i>	- parameter equal to 0.500/Pa where Pa is exponent
<i>buffer</i>	- the McMurchie-Davidson-Hermite 3-dimensional array (raveled to vector): <ul style="list-style-type: none"> • axis 0: dimension 3 (x, y or z Cartesian component) • axis 1: dimension n1+1 (0 to n1) • axis 2: dimension n1+1 (0 to n1)

See also

[D1.INDEX](#)

16.4.6.3 make_mdh_D2_coeff()

```
void oepdev::make_mdh_D2_coeff (
    int n1,
    int n2,
    double aPd,
    double * PA,
    double * PB,
    double * buffer )
```

Parameters

<i>n1</i>	- angular momentum of first function
<i>n2</i>	- angular momentum of second function
<i>aPd</i>	- parameter equal to 0.500/Pa where Pa is exponent
<i>PA</i>	- cartesian components of P-A distance
<i>PB</i>	- cartesian components of P-B distance
<i>buffer</i>	- the McMurchie-Davidson-Hermite 4-dimensional array (raveled to vector): <ul style="list-style-type: none"> • axis 0: dimension 3 (x, y or z Cartesian component) • axis 1: dimension n1+1 (0 to n1) • axis 2: dimension n2+1 (0 to n2) • axis 3: dimension n1+n2+1 (0 to n1+n2)

See also

[D2.INDEX](#)

16.4.6.4 make_mdh_D2_coeff_explicit_recursion()

```
void oepdev::make_mdh_D2_coeff_explicit_recursion (
    int n1,
    int n2,
    double aP,
    double * PA,
    double * PB,
    double * buffer )
```

Parameters

<i>n1</i>	- angular momentum of first function
<i>n2</i>	- angular momentum of second function
<i>aPd</i>	- parameter equal to 0.500/Pa where Pa is exponent
<i>PA</i>	- cartesian components of P-A distance
<i>PB</i>	- cartesian components of P-B distance
<i>buffer</i>	- the McMurchie-Davidson-Hermite 4-dimensional array (raveled to vector): <ul style="list-style-type: none"> • axis 0: dimension 3 (x, y or z Cartesian component) • axis 1: dimension $n1+1$ (0 to $n1$) • axis 2: dimension $n2+1$ (0 to $n2$) • axis 3: dimension $n1+n2+1$ (0 to $n1+n2$)

See also

[D2.INDEX](#)

16.4.6.5 make_mdh_D3_coeff()

```
void oepdev::make_mdh_D3_coeff (
    int n1,
    int n2,
    int n3,
    double aPd,
    double * PA,
    double * PB,
    double * PC,
    double * buffer )
```

Parameters

<i>n1</i>	- angular momentum of first function
<i>n2</i>	- angular momentum of second function
<i>n3</i>	- angular momentum of third function
<i>aPd</i>	- parameter equal to 0.500/Pa where Pa is exponent
<i>PA</i>	- cartesian components of P-A distance
<i>PB</i>	- cartesian components of P-B distance
<i>PC</i>	- cartesian components of P-C distance

Parameters

<i>buffer</i>	- the McMurchie-Davidson-Hermite 5-dimensional array (raveled to vector): <ul style="list-style-type: none"> • axis 0: dimension 3 (x, y or z Cartesian component) • axis 1: dimension n_1+1 (0 to n_1) • axis 2: dimension n_2+1 (0 to n_2) • axis 3: dimension n_3+1 (0 to n_3) • axis 4: dimension $n_1+n_2+n_3+1$ (0 to $n_1+n_2+n_3$)
---------------	---

See also

[D3.INDEX](#)

16.4.6.6 `make_mdh_R_coeff()`

```
void oepdev::make_mdh_R_coeff (
    int N,
    int L,
    int M,
    double alpha,
    double a,
    double b,
    double c,
    double * F,
    double * buffer )
```

Parameters

<i>N</i>	- increment in the summation of MDH series along <i>x</i> direction
<i>L</i>	- increment in the summation of MDH series along <i>y</i> direction
<i>M</i>	- increment in the summation of MDH series along <i>z</i> direction
<i>alpha</i>	- alpha parameter of R coefficient
<i>a</i>	- x component of PQ vector of R coefficient
<i>b</i>	- y component of PQ vector of R coefficient
<i>c</i>	- z component of PQ vector of R coefficient
<i>F</i>	- array of Boys function values for given alpha and PQ

Parameters

<i>buffer</i>	<div>- the McMurchie-Davidson 4-dimensional array (raveled to vector):<ul style="list-style-type: none">• axis 0: dimension $N+1$• axis 1: dimension $L+1$• axis 2: dimension $M+1$• axis 3: dimension $N+L+M+1$ (j-th element)</div>
---------------	---

16.5 The Three-Dimensional Vector Fields Library

Handles all sorts of scalar distributions in 3D Euclidean space, such as general vector potentials defined at particular collection of points. In this Module, you will also find handling both random and ordered points collections in a form of a G09 cube, as well as handling G09 Cube files. You will also find solvers used to fit the generalized multipole moments of a generalized density distribution, such as the electrostatic potential (ESP) fitting method. Located at `oepdev/lib3d`.

Classes

- class `oepdev::MultipoleConvergence`
Multipole Convergence.
- class `oepdev::DMTPole`
Distributed Multipole Analysis Container and Computer. Abstract Base.
- class `oepdev::Camm`
Cumulative Atomic Multipole Moments.
- class `oepdev::ESPSolver`
Charges from Electrostatic Potential (ESP). A solver-type class.
- class `oepdev::Points3DIterator`
Iterator over a collection of points in 3D space. Abstract base.
- class `oepdev::CubePoints3DIterator`
Iterator over a collection of points in 3D space. g09 Cube-like order.
- class `oepdev::RandomPoints3DIterator`
Iterator over a collection of points in 3D space. Random collection.
- class `oepdev::PointsCollection3D`
Collection of points in 3D space. Abstract base.
- class `oepdev::RandomPointsCollection3D`
Collection of random points in 3D space.
- class `oepdev::CubePointsCollection3D`
G09 cube-like ordered collection of points in 3D space.
- class `oepdev::Field3D`
General Vector Field in 3D Space. Abstract base.
- class `oepdev::ElectrostaticPotential3D`
Electrostatic potential of a molecule.
- class `oepdev::OEPotential3D< T >`
Class template for OEP 3D fields.

Typedefs

- using `oepdev::SharedField3D` = `std::shared_ptr< oepdev::Field3D >`

Functions

- `oepdev::OEPotential3D< T >::OEPotential3D` (const int &ndim, const int &np, const double &padding, std::shared_ptr< T > oep, const std::string &oepType)
Construct random spherical collection of 3D field of type T.
- `oepdev::OEPotential3D< T >::OEPotential3D` (const int &ndim, const int &nx, const int &ny, const int &nz, const double &px, const double &py, const double &pz, std::shared_ptr< T > oep, const std::string &oepType, psi::Options &options)
Construct ordered 3D collection of 3D field of type T.
- virtual `oepdev::OEPotential3D< T >::~~OEPotential3D` ()
Destructor.
- virtual void `oepdev::OEPotential3D< T >::print` () const
Print information of the object to Psi4 output.
- virtual std::shared_ptr< psi::Vector > `oepdev::OEPotential3D< T >::compute_xyz` (const double &x, const double &y, const double &z)
Compute a value of 3D field at point (x, y, z)

16.5.1 Detailed Description

16.5.2 Function Documentation

16.5.2.1 OEPotential3D() [1/2]

```
template<class T >
oepdev::OEPotential3D< T >::OEPotential3D (
    const int & ndim,
    const int & np,
    const double & padding,
    std::shared_ptr< T > oep,
    const std::string & oepType )
```

The points are drawn according to uniform distrinution in 3D space.

Parameters

<i>ndim</i>	- dimensionality of 3D field (1: scalar field, >2: vector field)
<i>np</i>	- number of points to draw
<i>padding</i>	- spherical padding distance (au)
<i>oep</i>	- OEP object of type T
<i>oepType</i>	- type of OEP

16.5.2.2 OEPotential3D() [2/2]

```
template<class T >
oepdev::OEPotential3D< T >::OEPotential3D (
    const int & ndim,
    const int & nx,
    const int & ny,
    const int & nz,
    const double & px,
    const double & py,
    const double & pz,
    std::shared_ptr< T > oep,
    const std::string & oepType,
    psi::Options & options )
```

The points are generated according to Gaussian cube file format.

Parameters

<i>ndim</i>	- dimensionality of 3D field (1: scalar field, >2: vector field)
<i>nx</i>	- number of points along x direction
<i>ny</i>	- number of points along y direction
<i>nz</i>	- number of points along z direction
<i>px</i>	- padding distance along x direction
<i>py</i>	- padding distance along y direction
<i>pz</i>	- padding distance along z direction
<i>oep</i>	- OEP object of type T
<i>oepType</i>	- type of OEP
<i>options</i>	- Psi4 options object

16.6 The Density Functional Theory Library

Implements the OEPDev ab initio DFT methods. Located at `oepdev/libdft`. Currently, this library is empty.

16.7 The OEPDev Utilities

Contains utility functions such as printing OEPDev preamble to the output file, class for wavefunction union, DIIS converger, CPHF Solver, SCF solver for external electrostatic perturbations, and others. You will also find here various iterators to go through orbital shells while computing ERI, or iterators over ERI itself. Located at `oepdev/libutil`.

Classes

- struct `oepdev::CISData`
Container to handle the CIS wavefunction parameters.
- class `oepdev::CISComputer`
CISComputer.
- class `oepdev::R_CISComputer`
- class `oepdev::U_CISComputer`
- class `oepdev::CPHF`
CPHF solver class.
- class `oepdev::DIISManager`
DIIS manager.
- class `oepdev::ShellCombinationsIterator`
Iterator for Shell Combinations. Abstract Base.
- class `oepdev::AOIntegralsIterator`
Iterator for AO Integrals. Abstract Base.
- class `oepdev::AllAOShellCombinationsIterator_4`
Loop over all possible ERI shells in a shell quartet.
- class `oepdev::AllAOShellCombinationsIterator_2`
Loop over all possible ERI shells in a shell doublet.
- class `oepdev::AllAOIntegralsIterator_4`
Loop over all possible ERI within a particular shell quartet.
- class `oepdev::AllAOIntegralsIterator_2`
Loop over all possible ERI within a particular shell doublet.
- struct `oepdev::PerturbCharges`
Structure to hold perturbing charges.
- class `oepdev::RHFPerturbed`
RHF theory under electrostatic perturbation.
- struct `oepdev::ABCD`
Simple structure to hold the Fourier series expansion coefficients.
- struct `oepdev::Fourier9`
Simple structure to hold the Fourier series expansion coefficients for N=4.
- class `oepdev::UnitaryOptimizer`
Find the optimim unitary matrix of quadratic matrix equation.

- class [oepdev::UnitaryOptimizer_4_2](#)
Find the optimum unitary matrix for quartic-quadratic matrix equation with trace.
- class [oepdev::WavefunctionUnion](#)
Union of two Wavefunction objects.

Macros

- `#define OEPDEV_USE_PSI4_DIIS_MANAGER 0`
Use DIIS from Psi4 (1) or OEPDev (0)?
- `#define OEPDEV_MAX_AM 8`
L_max.
- `#define OEPDEV_N_MAX_AM 17`
2L_max+1
- `#define OEPDEV_CRIT_ERI 1e-9`
*ERI criterion for E12, E34, E123 and lambda*EXY coefficients.*
- `#define OEPDEV_SIZE_BUFFER_R 250563`
*Size of R buffer (OEPDEV_N_MAX_AM*OEPDEV_N_MAX_AM*OEPDEV_N_MAX_AM*OEPDEV_N_MAX_AM*3)*
- `#define OEPDEV_SIZE_BUFFER_D2 3264`
Size of D2 buffer (3(OEPDEV_MAX_AM+1)*(OEPDEV_MAX_AM+1)*OEPDEV_N_MAX_AM)*
- `#define OEPDEV_AU_KcalPerMole 627.509`
Energy converters.
- `#define OEPDEV_AU_CMRec 219474.63`
- `#define OEPDEV_AU_EV 27.21138`

Typedefs

- using [oepdev::SharedShellsIterator](#) = std::shared_ptr< [ShellCombinationsIterator](#) >
Iterator over shells as shared pointer.
- using [oepdev::SharedAOIntsIterator](#) = std::shared_ptr< [AOIntegralsIterator](#) >
Iterator over AO integrals as shared pointer.

Functions

- PSI_API void [oepdev::preamble](#) (void)
Print preamble for module OEPDEV.
- template<typename... Args>
std::string [oepdev::string_sprintf](#) (const char *format, Args... args)
Format string output. Example: std::string text = oepdev::string_sprintf("Test %3d, %13.5f", 5, -10.5425);.
- PSI_API std::shared_ptr< SuperFunctional > [oepdev::create_superfunctional](#) (std::string name, Options &options)

Set up DFT functional.

- PSI.API std::shared_ptr< Molecule > [oepdev::extract_monomer](#) (std::shared_ptr< const Molecule > molecule_dimer, int id)

Extract molecule from dimer.

- PSI.API double [oepdev::compute_distance](#) (psi::SharedVector v1, psi::SharedVector v2)

Compute distance between two points in nD space.

- PSI.API std::shared_ptr< Wavefunction > [oepdev::solve_scf](#) (std::shared_ptr< Molecule > molecule, std::shared_ptr< BasisSet > primary, std::shared_ptr< BasisSet > auxiliary, std::shared_ptr< SuperFunctional > functional, Options &options, std::shared_ptr< PSIO > psio, bool compute_mints=false)

Solve RHF-SCF equations for a given molecule in a given basis set.

- PSI.API std::shared_ptr< Wavefunction > [oepdev::solve_scf_sad](#) (std::shared_ptr< Molecule > molecule, std::shared_ptr< BasisSet > primary, std::shared_ptr< BasisSet > auxiliary, std::vector< std::shared_ptr< BasisSet >> sad, std::vector< std::shared_ptr< BasisSet >> sad_fit, std::shared_ptr< SuperFunctional > functional, Options &options, std::shared_ptr< PSIO > psio, bool compute_mints=false)

Solve RHF-SCF equations for a given molecule in a given basis set.

- PSI.API double [oepdev::average_moment](#) (std::shared_ptr< psi::Vector > moment)

Compute the scalar magnitude of multipole moment.

- PSI.API std::vector< std::shared_ptr< psi::Matrix > > [oepdev::calculate_JK](#) (std::shared_ptr< psi::Wavefunction > wfn, std::shared_ptr< psi::Matrix > C)

Compute the Coulomb and exchange integral matrices in MO basis.

- PSI.API std::vector< std::shared_ptr< psi::Matrix > > [oepdev::calculate_JK_r](#) (std::shared_ptr< psi::Wavefunction > wfn, std::shared_ptr< psi::IntegralTransform > tr, std::shared_ptr< psi::Matrix > Dij)

Compute the Coulomb and exchange integral matrices in MO basis.

- PSI.API std::shared_ptr< psi::Matrix > [oepdev::calculate_der_D](#) (std::shared_ptr< psi::Wavefunction > wfn, std::shared_ptr< psi::IntegralTransform > tr, std::shared_ptr< psi::Matrix > C, std::vector< std::shared_ptr< psi::Matrix >> A)

Compute the derivative of exchange-correlation energy wrt the density matrix in MO-A basis.

- PSI.API double [oepdev::calculate_e_xc](#) (std::shared_ptr< psi::Wavefunction > wfn, std::shared_ptr< psi::IntegralTransform > tr, std::shared_ptr< psi::Matrix > f, std::shared_ptr< psi::Matrix > C)

Compute the exchange-correlation energy from ERI in MO-SCF basis.

- PSI.API std::shared_ptr< psi::Matrix > [oepdev::matrix_power_derivative](#) (std::shared_ptr< psi::Matrix > A, double g, double step)

Compute the contracted derivative of power of a square and symmetric matrix.

- std::shared_ptr< psi::Matrix > [oepdev::calculate_DFI_Vel](#) (std::shared_ptr< psi::IntegralFactory > f_aabb, std::shared_ptr< psi::IntegralFactory > f_abab, std::shared_ptr< psi::Matrix > d_b)

Compute the Effective DFI Potential Matrix Due To Electrons.

- PSI.API std::shared_ptr< psi::Matrix > [oepdev::calculate_DFI_Vel_JK](#) (std::shared_ptr< psi::IntegralFactory > f_aabb, std::shared_ptr< psi::IntegralFactory > f_abab, std::shared_ptr< psi::Matrix > d_b)

Compute the Effective DFI Coulomb+Exchange Potential Matrix Due To Electrons.

- PSI_API std::shared_ptr< psi::Matrix > oepdev::calculate_DFI_Vel_J (std::shared_ptr< psi::IntegralFactory > f_aabb, std::shared_ptr< psi::Matrix > d_b)

Compute the Effective DFI Coulomb Potential Matrix Due To Electrons.

16.7.1 Detailed Description

16.7.2 Function Documentation

16.7.2.1 _calculate_DFI_Vel()

```
std::shared_ptr< psi::Matrix > oepdev::_calculate_DFI_Vel (
    std::shared_ptr< psi::IntegralFactory > f_aabb,
    std::shared_ptr< psi::IntegralFactory > f_abab,
    std::shared_ptr< psi::Matrix > d_b )
```

Potential is felt by molecule A and induced by electrons in molecule B.

Parameters

<i>f_aabb</i>	- IntegralFactory of type (AA BB)
<i>f_abab</i>	- IntegralFactory of type (AB AB)
<i>d_b</i>	- one-particle density matrix in AO basis of B

Returns

- V_el(B) matrix in AO basis set of A

If f_abab is nullptr, then only Coulomb matrix is computed. Otherwise, also exchange contribution is computed.

16.7.2.2 average_moment()

```
PSI_API double oepdev::average_moment (
    std::shared_ptr< psi::Vector > moment )
```

Parameters

<i>moment</i>	- multipole moment vector with unique matrix elements. Now supported only for dipole and quadrupole.
---------------	--

Returns

- the average multipole moment value.

The magnitudes of multipole moments are defined here as follows:

- The dipole moment magnitude is just a norm

$$|\mu| \equiv \sqrt{\mu_x^2 + \mu_y^2 + \mu_z^2}$$

- The quadrupole moment magnitude refers to the traceless moment in Buckingham convention

$$|\Theta| \equiv \sqrt{\Theta_{zz}^2 + \frac{1}{3}(\Theta_{xx} - \Theta_{yy})^2 + \frac{4}{3}(\Theta_{xy}^2 + \Theta_{xz}^2 + \Theta_{yz}^2)}$$

In the above equation, the quadrupole moment elements refer to its traceless form.

16.7.2.3 calculate_der_D()

```
PSI_API std::shared_ptr< psi::Matrix > oepdev::calculate_der_D (
    std::shared_ptr< psi::Wavefunction > wfn,
    std::shared_ptr< psi::IntegralTransform > tr,
    std::shared_ptr< psi::Matrix > C,
    std::vector< std::shared_ptr< psi::Matrix >> A )
```

Reads the existing MO ERI's.

Parameters

<i>wfn</i>	- Wavefunction object
<i>tr</i>	- IntegralTransform object
<i>C</i>	- Transformation matrix MO-B::MO-A (columns are MO-A basis)
<i>A</i>	- Vector of matrices $A^{\wedge}(n)_{-}\{\text{bd}\}$

Returns

- derivative matrix in MO-A basis

16.7.2.4 calculate_DFI_Vel_J()

```
PSI_API std::shared_ptr< psi::Matrix > oepdev::calculate_DFI_Vel_J (
    std::shared_ptr< psi::IntegralFactory > f_aabb,
    std::shared_ptr< psi::Matrix > d_b )
```

Potential is felt by molecule A and induced by electrons in molecule B.

Parameters

<i>f_aabb</i>	- IntegralFactory of type (AA BB)
<i>d.b</i>	- one-particle density matrix in AO basis of B

Returns

- $V_{el}(B)$ matrix in AO basis set of A

16.7.2.5 calculate_DFI_Vel_JK()

```
PSI_API std::shared_ptr< psi::Matrix > oepdev::calculate_DFI_Vel_JK (
    std::shared_ptr< psi::IntegralFactory > f_aabb,
    std::shared_ptr< psi::IntegralFactory > f_abab,
    std::shared_ptr< psi::Matrix > d.b )
```

Potential is felt by molecule A and induced by electrons in molecule B.

Parameters

<i>f_aabb</i>	- IntegralFactory of type (AA BB)
<i>f_abab</i>	- IntegralFactory of type (AB AB)
<i>d.b</i>	- one-particle density matrix in AO basis of B

Returns

- $V_{el}(B)$ matrix in AO basis set of A

16.7.2.6 calculate_e_xc()

```
PSI_API double oepdev::calculate_e_xc (
    std::shared_ptr< psi::Wavefunction > wfn,
    std::shared_ptr< psi::IntegralTransform > tr,
    std::shared_ptr< psi::Matrix > f,
    std::shared_ptr< psi::Matrix > C )
```

Reads the existing MO ERI's.

Parameters

<i>wfn</i>	- Wavefunction object
<i>tr</i>	- IntegralTransform object
<i>f</i>	- f_{ij} matrix in MO-NEW basis
<i>C</i>	- Transformation matrix MO-SCF::MO-NEW (columns are MO-A basis)

Returns

- Exchange-correlation energy

16.7.2.7 calculate_JK()

```
PSI_API std::vector< std::shared_ptr< psi::Matrix > > oepdev::calculate_JK
(
    std::shared_ptr< psi::Wavefunction > wfn,
    std::shared_ptr< psi::Matrix > C )
```

Transforms the AO ERI's based on provided C matrix.

Parameters

<i>wfn</i>	- Wavefunction object
<i>C</i>	- molecular orbital coefficients (AO x MO)

Returns

- vector with J_{ij} and K_{ij} matrix

16.7.2.8 calculate_JK_r()

```
PSI_API std::vector< std::shared_ptr< psi::Matrix > > oepdev::calculate_JK_r
(
    std::shared_ptr< psi::Wavefunction > wfn,
    std::shared_ptr< psi::IntegralTransform > tr,
    std::shared_ptr< psi::Matrix > Dij )
```

Reads the existing MO ERI's.

Parameters

<i>wfn</i>	- Wavefunction object
<i>tr</i>	- IntegralTransform object
<i>D</i>	- density matrix in MO basis

Returns

- vector with J_{ij} and K_{ij} matrix

16.7.2.9 compute_distance()

```
PSI_API double oepdev::compute_distance (
    psi::SharedVector v1,
    psi::SharedVector v2 )
```

Parameters

<i>v1</i>	- vector 1
<i>v2</i>	- vector 2

Returns

distance The vectors have to have the same length.

16.7.2.10 create_superfunctional()

```
PSI_API std::shared_ptr< SuperFunctional > oepdev::create_superfunctional (
    std::string name,
    Options & options )
```

Now it accepts only pure HF functional.

Parameters

<i>name</i>	name of the functional ("HF" is now only available)
<i>options</i>	psi::Options object

Returns

psi::SharedSuperFunctional object with functional.

Examples:

[example_scf_perturb.cc](#).

16.7.2.11 extract_monomer()

```
PSI_API std::shared_ptr< Molecule > oepdev::extract_monomer (
    std::shared_ptr< const Molecule > molecule_dimer,
    int id )
```


Parameters

<i>molecule_dimer</i>	psi::SharedMolecule object with dimer
<i>id</i>	index of a molecule (starts from 1)

Returns

psi::SharedMolecule object with indicated monomer

16.7.2.12 matrix_power_derivative()

```
PSI_API std::shared_ptr< psi::Matrix > oepdev::matrix_power_derivative (
    std::shared_ptr< psi::Matrix > A,
    double g,
    double step )
```

The contracted matrix derivative is defined here as

$$\mathbf{D} = \frac{d\mathbf{A}^\gamma}{d\mathbf{A}} : \mathbb{I}$$

where \mathbb{I} is the identity matrix. The derivative, which is the fourth-rank tensor, is computed by the forward 2-centre finite difference formula,

$$f' = (f(h) - f(0)) / h$$

- if γ is non-integer, input matrix has to be positive-definite.

Parameters

<i>A</i>	- Matrix
<i>g</i>	- Power
<i>step</i>	- Differentiation step h

Returns

- Contracted derivative (matrix)

16.7.2.13 solve_scf()

```
PSI_API std::shared_ptr< Wavefunction > oepdev::solve_scf (
    std::shared_ptr< Molecule > molecule,
    std::shared_ptr< BasisSet > primary,
```

```
std::shared_ptr< BasisSet > auxiliary,
std::shared_ptr< SuperFunctional > functional,
Options & options,
std::shared_ptr< PSIO > psio,
bool compute_mints = false )
```

Parameters

<i>molecule</i>	psi::SharedMolecule object with molecule
<i>primary</i>	shared primary basis set
<i>auxiliary</i>	shared auxiliary basis set
<i>functional</i>	DFT functional
<i>options</i>	psi::Options object
<i>psio</i>	psi::PSIO object
<i>compute_mints</i>	Compute integrals (write IWL TOC entry - necessary when transforming integrals)

Returns

psi::SharedWavefunction SCF wavefunction of the molecule

16.7.2.14 solve_scf_sad()

```
PSIAPI std::shared_ptr< Wavefunction > oepdev::solve_scf_sad (
    std::shared_ptr< Molecule > molecule,
    std::shared_ptr< BasisSet > primary,
    std::shared_ptr< BasisSet > auxiliary,
    std::vector< std::shared_ptr< BasisSet >> sad,
    std::vector< std::shared_ptr< BasisSet >> sad_fit,
    std::shared_ptr< SuperFunctional > functional,
    Options & options,
    std::shared_ptr< PSIO > psio,
    bool compute_mints = false )
```

Parameters

<i>molecule</i>	psi::SharedMolecule object with molecule
<i>primary</i>	shared primary basis set
<i>auxiliary</i>	shared auxiliary basis set
<i>sad</i>	SAD basis set list
<i>sad_fit</i>	SAD DF fitting basis set list
<i>functional</i>	DFT functional
<i>options</i>	psi::Options object
<i>psio</i>	psi::PSIO object
<i>compute_mints</i>	Compute integrals (write IWL TOC entry - necessary when transforming integrals)

Returns

psi::SharedWavefunction SCF wavefunction of the molecule

16.8 The OEPDev Testing Platform Library

Testing platform at C++ level of code. You should add more tests here when developing new functionalities, theories or models. Located at `oepdev/libtest`.

Classes

- class `oepdev::test::Test`
Manages test routines.

16.8.1 Detailed Description

Chapter 17

Namespace Documentation

17.1 `gefp.basis.optimize` Namespace Reference

Classes

- class [DFBasis](#)
- class [DFBasisOptimizer](#)
- class [OEP](#)
- class [OEP_CT](#)
- class [OEP_FockLike](#)
- class [OEP_Pauli](#)

Functions

- def **make_bastempl** (templ, param)
- def **oeffitbasis** (mol, role='ORBITAL')
- def **removeComments** (string)

17.1.1 Detailed Description

Auxiliary Basis Set Optimization Library.

The auxiliary basis sets for generalized density fitting (GDF) are here optimized.

17.2 `gefp.density.dfi` Namespace Reference

Classes

- class [DFI](#)
- class [DFI_J](#)

- class [DFIJK](#)
- class [SCF](#)

Variables

- int **MAX.NBF** = 128

17.2.1 Detailed Description

Demonstrates the use of Psi4 from Python level.

Useful notes:

- Use psi4.core module for most of the work
- Useful modules within psi4.core:
 - MintsHelper
 - Molecule
 - BasisSet
 - ExternalPotential
 - others
- Psi4 defines its own matrix type (psi4.core.Matrix).
Extracting numpy.array is easy:


```
numpy_array = numpy.asarray(psi4_matrix)
```

 Creating Psi4 matrix from array is also easy:


```
psi4_matrix = psi4.core.Matrix.from_array(numpy_array)
```
- To compute 1-el potential matrix for a set of charges
use ExternalPotential (charge positions are to be provided in Angstroms)
unless charges are just nuclei within the basis set (in this case use of ao_potential of MintsHelper is easier).
- ao_potential method of MintsHelper is limited only for nuclei within the same basis (the nuclei are taken from the first basis set axis, for example:


```
mints = MintsHelper(basis_X)
mints.ao_potential()          -> nuclei taken from basis of mints object (b
mints.ao_potential(basis_1, basis_2) -> nuclei taken from basis_1
```
- Psi4 has efficient and easy to use method of defining fragments within a molecule (u
Defining ghost atoms and extracting fragment i in the multimer-centred basis set is
(method extract_subsets(...) of psi4.core.Molecule)

17.3 oepdev Namespace Reference

OEPEDev module namespace.

Classes

- struct [ABCD](#)
Simple structure to hold the Fourier series expansion coefficients.
- class [AbInitioPolarGEFactory](#)
Polarization GEFP Factory from First Principles. Hartree-Fock Approximation.
- class [AllAOIntegralsIterator_2](#)
Loop over all possible ERI within a particular shell doublet.

- class [AllAOIntegralsIterator_4](#)
Loop over all possible ERI within a particular shell quartet.
- class [AllAOShellCombinationsIterator_2](#)
Loop over all possible ERI shells in a shell doublet.
- class [AllAOShellCombinationsIterator_4](#)
Loop over all possible ERI shells in a shell quartet.
- class [AOIntegralsIterator](#)
Iterator for AO Integrals. Abstract Base.
- class [CAMM](#)
Cumulative Atomic Multipole Moments.
- class [ChargeTransferEnergyOEPotential](#)
Generalized One-Electron Potential for Charge-Transfer Interaction Energy.
- class [ChargeTransferEnergySolver](#)
Compute the Charge-Transfer interaction energy between unperturbed wavefunctions.
- class [CISComputer](#)
CISComputer.
- struct [CISData](#)
Container to handle the CIS wavefunction parameters.
- class [CPHF](#)
CPHF solver class.
- class [CubePoints3DIterator](#)
Iterator over a collection of points in 3D space. g09 Cube-like order.
- class [CubePointsCollection3D](#)
G09 cube-like ordered collection of points in 3D space.
- class [DIISManager](#)
DIIS manager.
- class [DMTPole](#)
Distributed Multipole Analysis Container and Computer. Abstract Base.
- class [DoubleGeneralizedDensityFit](#)
Generalized Density Fitting Scheme - Double Fit.
- class [EETCouplingOEPotential](#)
Generalized One-Electron Potential for EET coupling calculations.
- class [EETCouplingSolver](#)
Compute the EET coupling energy between unperturbed wavefunctions.
- class [ElectrostaticEnergyOEPotential](#)
Generalized One-Electron Potential for Electrostatic Energy.
- class [ElectrostaticEnergySolver](#)
Compute the Coulombic interaction energy between unperturbed wavefunctions.
- class [ElectrostaticPotential3D](#)
Electrostatic potential of a molecule.

- class [ERI_1_1](#)
2-centre ERI of the form $(a|O(2)|b)$ where $O(2) = 1/r12$.
- class [ERI_2_2](#)
4-centre ERI of the form $(ab|O(2)|cd)$ where $O(2) = 1/r12$.
- class [ERI_3_1](#)
4-centre ERI of the form $(abc|O(2)|d)$ where $O(2) = 1/r12$.
- class [ESPSolver](#)
Charges from Electrostatic Potential (ESP). A solver-type class.
- class [FFAbInitioPolarGEFactory](#)
Polarization GEFP Factory from First Principles: Finite-Difference Model. Arbitrary level of theory.
- class [Field3D](#)
General Vector Field in 3D Space. Abstract base.
- struct [Fourier9](#)
Simple structure to hold the Fourier series expansion coefficients for $N=4$.
- class [GenEffFrag](#)
Generalized Effective Fragment. Container Class.
- class [GenEffPar](#)
Generalized Effective Fragment Parameters. Container Class.
- class [GenEffParFactory](#)
Generalized Effective Fragment Factory. Abstract Base.
- class [GeneralizedDensityFit](#)
Generalized Density Fitting Scheme. Abstract Base.
- class [GeneralizedPolarGEFactory](#)
Polarization GEFP Factory with Least-Squares Parameterization.
- class [IntegralFactory](#)
Extended [IntegralFactory](#) for computing integrals.
- class [LinearGradientNonUniformEFieldPolarGEFactory](#)
Polarization GEFP Factory with Least-Squares Parameterization.
- class [LinearNonUniformEFieldPolarGEFactory](#)
Polarization GEFP Factory with Least-Squares Parameterization.
- class [LinearUniformEFieldPolarGEFactory](#)
Polarization GEFP Factory with Least-Squares Parameterization.
- class [MultipoleConvergence](#)
Multipole Convergence.
- class [NonUniformEFieldPolarGEFactory](#)
Polarization GEFP Factory with Least-Squares Parameterization.
- class [OEPDevSolver](#)
Solver of properties of molecular aggregates. Abstract base.
- class [OEPotential](#)

- Generalized One-Electron Potential: Abstract base.*
- class [OEPotential3D](#)
 - Class template for OEP 3D fields.*
- struct [OEType](#)
 - Container to handle the type of One-Electron Potentials.*
- struct [PerturbCharges](#)
 - Structure to hold perturbing charges.*
- class [Points3DIterator](#)
 - Iterator over a collection of points in 3D space. Abstract base.*
- class [PointsCollection3D](#)
 - Collection of points in 3D space. Abstract base.*
- class [PolarGEFactory](#)
 - Polarization GEFP Factory. Abstract Base.*
- class [PotentialInt](#)
 - Computes potential integrals.*
- class [QuadraticGradientNonUniformEFieldPolarGEFactory](#)
 - Polarization GEFP Factory with Least-Squares Parameterization.*
- class [QuadraticNonUniformEFieldPolarGEFactory](#)
 - Polarization GEFP Factory with Least-Squares Parameterization.*
- class [QuadraticUniformEFieldPolarGEFactory](#)
 - Polarization GEFP Factory with Least-Squares Parameterization.*
- class [R_CISComputer](#)
- class [RandomPoints3DIterator](#)
 - Iterator over a collection of points in 3D space. Random collection.*
- class [RandomPointsCollection3D](#)
 - Collection of random points in 3D space.*
- class [RepulsionEnergyOEPotential](#)
 - Generalized One-Electron Potential for Pauli Repulsion Energy.*
- class [RepulsionEnergySolver](#)
 - Compute the Pauli-Repulsion interaction energy between unperturbed wavefunctions.*
- class [RHFPerturbed](#)
 - RHF theory under electrostatic perturbation.*
- class [ShellCombinationsIterator](#)
 - Iterator for Shell Combinations. Abstract Base.*
- class [SingleGeneralizedDensityFit](#)
 - Generalized Density Fitting Scheme - Single Fit.*
- class [TIData](#)
 - Transfer Integral EET Data.*
- class [TwoBodyAOInt](#)
- class [TwoElectronInt](#)

General Two Electron Integral.

- class [U.CISComputer](#)
- class [UniformEFieldPolarGEFactory](#)

Polarization GEFP Factory with Least-Squares Parameterization.

- class [UnitaryOptimizer](#)

Find the optimum unitary matrix of quadratic matrix equation.

- class [UnitaryOptimizer_4_2](#)

Find the optimum unitary matrix for quartic-quadratic matrix equation with trace.

- class [UnitaryTransformedMOPolarGEFactory](#)

Polarization GEFP Factory with Least-Squares Scaling of MO Space.

- class [WavefunctionUnion](#)

Union of two Wavefunction objects.

Typedefs

- using **SharedField3D** = std::shared_ptr< [oepdev::Field3D](#) >
- using **SharedWavefunction** = std::shared_ptr< [Wavefunction](#) >
- using **SharedBasisSet** = std::shared_ptr< [BasisSet](#) >
- using **SharedMatrix** = std::shared_ptr< [Matrix](#) >
- using **SharedVector** = std::shared_ptr< [Vector](#) >
- using **SharedDMTPole** = std::shared_ptr< [DMTPole](#) >
- using **SharedLocalizer** = std::shared_ptr< [Localizer](#) >
- using **SharedCISData** = std::shared_ptr< [CISData](#) >
- using **SharedWavefunctionUnion** = std::shared_ptr< [WavefunctionUnion](#) >
- using **SharedOEPotential** = std::shared_ptr< [OEPotential](#) >
- using **SharedDMTPConvergence** = std::shared_ptr< [oepdev::MultipoleConvergence](#) >
- using **SharedMolecule** = std::shared_ptr< [psi::Molecule](#) >
- using **SharedMOSpace** = std::shared_ptr< [psi::MOSpace](#) >
- using **SharedMOSpaceVector** = std::vector< std::shared_ptr< [psi::MOSpace](#) > >
- using **SharedIntegralTransform** = std::shared_ptr< [psi::IntegralTransform](#) >
- using **SharedIntegralFactory** = std::shared_ptr< [IntegralFactory](#) >
- using **SharedTwoBodyAOInt** = std::shared_ptr< [TwoBodyAOInt](#) >
- using [SharedShellsIterator](#) = std::shared_ptr< [ShellCombinationsIterator](#) >
- *Iterator over shells as shared pointer.*
- using [SharedAOIntsIterator](#) = std::shared_ptr< [AOIntegralsIterator](#) >
- *Iterator over AO integrals as shared pointer.*
- using **SharedSuperFunctional** = std::shared_ptr< [SuperFunctional](#) >

Functions

- double [d.N.n1.n2](#) (int N, int n1, int n2, double PA, double PB, double aP)
Compute McMurchie-Davidson-Hermite (MDH) coefficient for binomial expansion.
- void [make_mdh_D2_coeff_explicit_recursion](#) (int n1, int n2, double aP, double *PA, double *PB, double *buffer)
Compute the McMurchie-Davidson-Hermite coefficients for binomial expansion by explicit recursion. This function makes the same changes to buffers as [oepdev::make_mdh_D2_coeff](#), but implements it through explicit recursion by calls to [oepdev::d.N.n1.n2](#). Therefore, it is slightly slower. Here for debugging purposes.
- void [make_mdh_D1_coeff](#) (int n1, double aPd, double *buffer)
Compute the McMurchie-Davidson-Hermite coefficients for monomial expansion.
- void [make_mdh_D2_coeff](#) (int n1, int n2, double aPd, double *PA, double *PB, double *buffer)
Compute the McMurchie-Davidson-Hermite coefficients for binomial expansion.
- void [make_mdh_D3_coeff](#) (int n1, int n2, int n3, double aPd, double *PA, double *PB, double *PC, double *buffer)
Compute the McMurchie-Davidson-Hermite coefficients for trinomial expansion.
- void [make_mdh_R_coeff](#) (int N, int L, int M, double alpha, double a, double b, double c, double *F, double *buffer)
Compute the McMurchie-Davidson R coefficients.
- constexpr std::complex< double > **operator**""_i (unsigned long long d)
- constexpr std::complex< double > **operator**""_i (long double d)
- PSI.API void [preamble](#) (void)
Print preamble for module OEPDEV.
- PSI.API std::shared_ptr< SuperFunctional > [create_superfunctional](#) (std::string name, Options &options)
Set up DFT functional.
- PSI.API std::shared_ptr< Molecule > [extract_monomer](#) (std::shared_ptr< const Molecule > molecule_dimer, int id)
Extract molecule from dimer.
- PSI.API double [compute_distance](#) (psi::SharedVector v1, psi::SharedVector v2)
Compute distance between two points in nD space.
- PSI.API std::shared_ptr< Wavefunction > [solve_scf](#) (std::shared_ptr< Molecule > molecule, std::shared_ptr< BasisSet > primary, std::shared_ptr< BasisSet > auxiliary, std::shared_ptr< SuperFunctional > functional, Options &options, std::shared_ptr< PSIO > psio, bool compute_mints=false)
Solve RHF-SCF equations for a given molecule in a given basis set.
- PSI.API std::shared_ptr< Wavefunction > [solve_scf_sad](#) (std::shared_ptr< Molecule > molecule, std::shared_ptr< BasisSet > primary, std::shared_ptr< BasisSet > auxiliary, std::vector< std::shared_ptr< BasisSet > > sad, std::vector< std::shared_ptr< BasisSet > > sad_fit, std::shared_ptr< SuperFunctional > functional, Options &options, std::shared_ptr< PSIO > psio, bool compute_mints=false)
Solve RHF-SCF equations for a given molecule in a given basis set.

- PSI.API double [average_moment](#) (std::shared_ptr< psi::Vector > moment)
Compute the scalar magnitude of multipole moment.
- PSI.API std::vector< std::shared_ptr< psi::Matrix > > [calculate_JK](#) (std::shared_ptr< psi::Wavefunction > wfn, std::shared_ptr< psi::Matrix > C)
Compute the Coulomb and exchange integral matrices in MO basis.
- PSI.API std::vector< std::shared_ptr< psi::Matrix > > [calculate_JK_r](#) (std::shared_ptr< psi::Wavefunction > wfn, std::shared_ptr< psi::IntegralTransform > tr, std::shared_ptr< psi::Matrix > Dij)
Compute the Coulomb and exchange integral matrices in MO basis.
- PSI.API std::shared_ptr< psi::Matrix > [calculate_der_D](#) (std::shared_ptr< psi::Wavefunction > wfn, std::shared_ptr< psi::IntegralTransform > tr, std::shared_ptr< psi::Matrix > C, std::vector< std::shared_ptr< psi::Matrix > > A)
Compute the derivative of exchange-correlation energy wrt the density matrix in MO-A basis.
- PSI.API double [calculate_e_xc](#) (std::shared_ptr< psi::Wavefunction > wfn, std::shared_ptr< psi::IntegralTransform > tr, std::shared_ptr< psi::Matrix > f, std::shared_ptr< psi::Matrix > C)
Compute the exchange-correlation energy from ERI in MO-SCF basis.
- PSI.API std::shared_ptr< psi::Matrix > [matrix_power_derivative](#) (std::shared_ptr< psi::Matrix > A, double g, double step)
Compute the contracted derivative of power of a square and symmetric matrix.
- std::shared_ptr< psi::Matrix > [_calculate_DFI_Vel](#) (std::shared_ptr< psi::IntegralFactory > f_aabb, std::shared_ptr< psi::IntegralFactory > f_abab, std::shared_ptr< psi::Matrix > d_b)
Compute the Effective DFI Potential Matrix Due To Electrons.
- PSI.API std::shared_ptr< psi::Matrix > [calculate_DFI_Vel_JK](#) (std::shared_ptr< psi::IntegralFactory > f_aabb, std::shared_ptr< psi::IntegralFactory > f_abab, std::shared_ptr< psi::Matrix > d_b)
Compute the Effective DFI Coulomb+Exchange Potential Matrix Due To Electrons.
- PSI.API std::shared_ptr< psi::Matrix > [calculate_DFI_Vel_J](#) (std::shared_ptr< psi::IntegralFactory > f_aabb, std::shared_ptr< psi::Matrix > d_b)
Compute the Effective DFI Coulomb Potential Matrix Due To Electrons.
- template<typename... Args>
std::string [string_sprintf](#) (const char *format, Args... args)
Format string output. Example: std::string text = oepdev::string_sprintf("Test %3d, %13.5f", 5, -10.5425);

17.3.1 Detailed Description

Contains all the functionalities for the development of the Generalized One-Electroc Potentials (OEP's).

17.4 psi Namespace Reference

Psi4 package namespace.

Typedefs

- using **SharedBasisSet** = std::shared_ptr< BasisSet >
- using **SharedMolecule** = std::shared_ptr< Molecule >
- using **SharedMatrix** = std::shared_ptr< Matrix >
- using **SharedWavefunction** = std::shared_ptr< Wavefunction >

Functions

- PSI_API int [read_options](#) (std::string name, Options &options)
Options for the OEPDev plugin.
- void **export_dmt** (py::module &)
- void **export_cphf** (py::module &)
- void **export_solver** (py::module &)
- void **export_util** (py::module &)
- void **export_oep** (py::module &)
- PSI_API SharedWavefunction [oepdev](#) (SharedWavefunction ref_wfn, Options &options)
Main routine of the OEPDev plugin.
- **PYBIND11_MODULE** ([oepdev](#), m)

17.4.1 Detailed Description

Contains all Psi4 functionalities.

17.4.2 Function Documentation

17.4.2.1 oepdev()

```
PSI_API SharedWavefunction psi::oepdev (  
    SharedWavefunction ref_wfn,  
    Options & options )
```

Created with intention to test various models of the interaction energy between two molecules, described by the Hartree-Fock-Roothaan-Hall theory or the configuration interaction with singles theory.

In particular, the plugin tests the models of:

1. the Pauli repulsion and CT interaction energy (Project II)
2. the Induction interaction energy (Project III)
3. the excitation energy transfer couplings (Project I)

against benchmarks (exact or reference solutions). The list of implemented models can be found in [Implemented Models](#) .

Parameters

<i>ref_wfn</i>	shared wavefunction of a dimer
<i>options</i>	psi::Options object

Returns

psi::SharedWavefunction (either ref_wfn or wavefunction union)

17.4.2.2 read_options()

```
PSI_API int psi::readOptions (
    std::string name,
    Options & options )
```

Parameters

<i>name</i>	name of driver function
<i>options</i>	psi::Options object

Returns

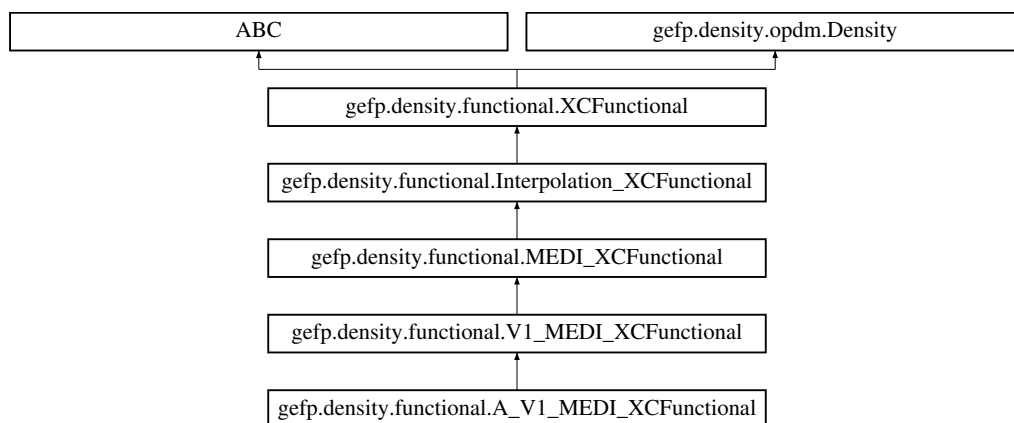
true

Chapter 18

Class Documentation

18.1 `gefp.density.functional.A_V1_MEDI_XCFunctional` Class Reference

Inheritance diagram for `gefp.density.functional.A_V1_MEDI_XCFunctional`:



Public Member Functions

- `def __init__ (self, coeff, kmax)`
- `def abbr (self)`
- `def compute_a0 (self, n)`

Static Public Member Functions

- `def name ()`

Additional Inherited Members

18.1.1 Detailed Description

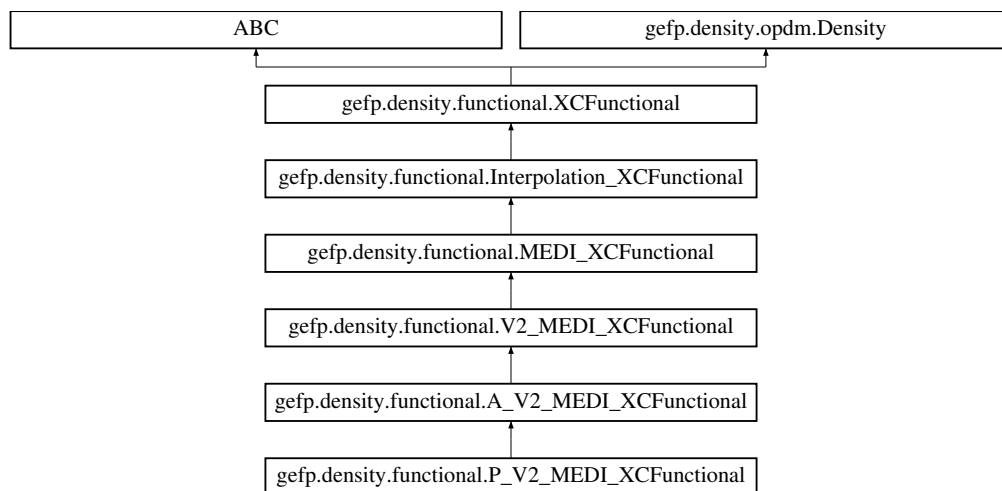
The New Class of Exchange-Correlation Functionals:
Interpolation Functionals with Monotonous Exponential Decay.

The documentation for this class was generated from the following file:

- `gefp/gefp/density/functional.py`

18.2 `gefp.density.functional.A_V2_MEDI_XCFunctional` Class Reference

Inheritance diagram for `gefp.density.functional.A_V2_MEDI_XCFunctional`:



Public Member Functions

- `def __init__ (self, coeff, kmax)`
- `def abbr (self)`
- `def compute_a0 (self, n)`
- `def energy_P_costly (self, x)`
- `def gradient_P (self, x)`
- `def gradient_P_approximate (self, x)`

Static Public Member Functions

- `def name ()`

Additional Inherited Members

18.2.1 Detailed Description

The New Class of Exchange-Correlation Functionals:
Interpolation Functionals with Monotonous Exponential Decay.

The documentation for this class was generated from the following file:

- [gefp/gefp/density/functional.py](#)

18.3 oepdev::ABCD Struct Reference

Simple structure to hold the Fourier series expansion coefficients.

```
#include <unitary_optimizer.h>
```

Public Attributes

- double **A**
- double **B**
- double **C**
- double **D**

18.3.1 Detailed Description

The documentation for this struct was generated from the following file:

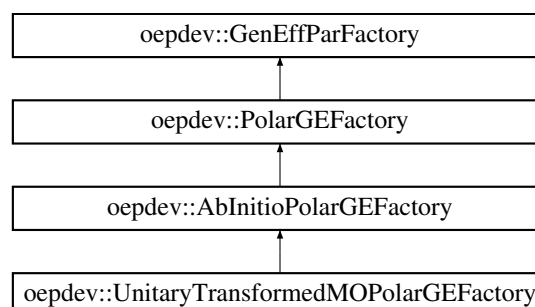
- [oepdev/libutil/unitary_optimizer.h](#)

18.4 oepdev::AbInitioPolarGEFactory Class Reference

Polarization GEFP Factory from First Principles. Hartree-Fock Approximation.

```
#include <gefp.h>
```

Inheritance diagram for oepdev::AbInitioPolarGEFactory:



Public Member Functions

- **AbInitioPolarGEFactory** (std::shared_ptr< psi::Wavefunction > [wfn](#), psi::Options &opt)
- virtual std::shared_ptr< [GenEffPar](#) > [compute](#) (void)

Compute the density matrix susceptibility tensors.

Additional Inherited Members

18.4.1 Detailed Description

Implements creation of the density matrix susceptibility tensors for which $\mathbf{X} = \mathbf{1}$. Guarantees the idempotency of the density matrix up to first-order in LCAO-MO variation. The density matrix susceptibility tensor is represented by:

$$\delta D_{\alpha\beta} = \sum_i \mathbf{B}_{\alpha\beta}^{(i;1)} \cdot \mathbf{F}(\mathbf{r}_i)$$

where $\mathbf{B}_{\alpha\beta}^{(i;1)}$ is the density matrix dipole polarizability defined for the distributed LMO site at \mathbf{r}_i . Its explicit form is given by

$$\mathbf{B}_{\alpha\beta}^{(i;1)} = C_{\alpha i}^{(0)} \mathbf{b}_{\beta}^{(i;1)} C_{\beta i}^{(0)} \mathbf{b}_{\alpha}^{(i;1)} - \sum_{\gamma} \left(D_{\alpha\gamma}^{(0)} C_{\beta i}^{(0)} + D_{\beta\gamma}^{(0)} C_{\alpha i}^{(0)} \right) \mathbf{b}_{\gamma}^{(i;1)}$$

where the susceptibility of the LCAO-MO coefficient is given by

$$b_{\alpha;w}^{(i;1)} = \frac{1}{4} \sum_u^{x,y,z} [\alpha_i]_{uw} \left[[\mathbf{L}_i]_{\text{Left}}^{-1} \right]_{u;\alpha}$$

for $w = x, y, z$. The auxiliary tensor \mathbb{L} is defined as

$$\mathbb{L} = \mathbf{C}^{(0)\text{T}} \cdot \mathbb{M} \cdot \left(\mathbf{1} - \mathbf{D}^{(0)} \right)$$

where \mathbb{M} is the dipole integral vector of matrices in AO representation. The left inverse of the i -th element is defined as

$$[\mathbf{L}_i]_{\text{Left}}^{-1} \equiv [\mathbf{L}_i^{\text{T}} \cdot \mathbf{L}_i]^{-1} \cdot \mathbf{L}_i^{\text{T}}$$

Note that $\mathbf{L}_i \equiv [\mathbb{L}]_i$ is a $n \times 3$ matrix, whereas its left inverse is a $3 \times n$ matrix with n being the size of the AO basis set.

The documentation for this class was generated from the following files:

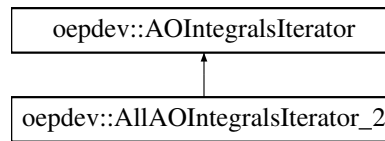
- oepdev/libgefp/[gefp.h](#)
- oepdev/libgefp/gefp_polar_abinitio.cc

18.5 oepdev::AllAOIntegralsIterator_2 Class Reference

Loop over all possible ERI within a particular shell doublet.

```
#include <integrals_iter.h>
```

Inheritance diagram for oepdev::AllAOIntegralsIterator_2:



Public Member Functions

- [AllAOIntegralsIterator_2](#) (const [ShellCombinationsIterator](#) *shellIter)
Construct by shell iterator (const object)
- [AllAOIntegralsIterator_2](#) (std::shared_ptr< [ShellCombinationsIterator](#) > shellIter)
Construct by shell iterator (pointed by shared pointer)
- void [first](#) ()
First iteration.
- void [next](#) ()
Next iteration.
- int [i](#) () const
Grab the current integral i index.
- int [j](#) () const
Grab the current integral j index.
- int [index](#) () const

Additional Inherited Members

18.5.1 Detailed Description

Constructed by providing a const reference or shared pointer to an AllAOShellCombinationsIterator object.

See also

[AllAOShellCombinationsIterator_2](#)

18.5.2 Constructor & Destructor Documentation

18.5.2.1 AllAOIntegralsIterator_2() [1/2]

```

AllAOIntegralsIterator_2::AllAOIntegralsIterator_2 (
    const ShellCombinationsIterator * shellIter )
  
```

Parameters

<i>shellIter</i>	- shell iterator object
------------------	-------------------------

18.5.2.2 AllAOIntegralsIterator_2() [2/2]

```
AllAOIntegralsIterator_2::AllAOIntegralsIterator_2 (
    std::shared_ptr< ShellCombinationsIterator > shellIter )
```

Parameters

<i>shellIter</i>	- shell iterator object
------------------	-------------------------

18.5.3 Member Function Documentation

18.5.3.1 index()

```
int oepdev::AllAOIntegralsIterator_2::index (
    void ) const [inline], [virtual]
```

Grab the current index of integral value stored in the buffer

Implements [oepdev::AOIntegralsIterator](#).

The documentation for this class was generated from the following files:

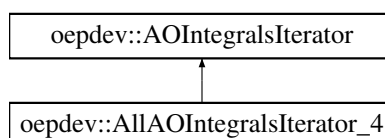
- oepdev/libutil/[integrals_iter.h](#)
- oepdev/libutil/integrals_iter.cc

18.6 oepdev::AllAOIntegralsIterator_4 Class Reference

Loop over all possible ERI within a particular shell quartet.

```
#include <integrals_iter.h>
```

Inheritance diagram for oepdev::AllAOIntegralsIterator_4:



Public Member Functions

- [AllAOIntegralsIterator_4](#) (const [ShellCombinationsIterator](#) *shellIter)
Construct by shell iterator (const object)
- [AllAOIntegralsIterator_4](#) (std::shared_ptr< [ShellCombinationsIterator](#) > shellIter)
Construct by shell iterator (pointed by shared pointer)
- void [first](#) ()
First iteration.
- void [next](#) ()
Next iteration.
- int [i](#) () const
Grab the current integral i index.
- int [j](#) () const
Grab the current integral j index.
- int [k](#) () const
Grab the current integral k index.
- int [l](#) () const
Grab the current integral l index.
- int [index](#) () const

Additional Inherited Members

18.6.1 Detailed Description

Constructed by providing a const reference or shared pointer to an AllAOShellCombinationsIterator object.

See also

[AllAOShellCombinationsIterator_4](#)

18.6.2 Constructor & Destructor Documentation

18.6.2.1 AllAOIntegralsIterator_4() [1/2]

```
AllAOIntegralsIterator_4::AllAOIntegralsIterator_4 (
    const ShellCombinationsIterator * shellIter )
```

Parameters

<i>shellIter</i>	- shell iterator object
------------------	-------------------------

18.6.2.2 AllAOIntegralsIterator_4() [2/2]

```
AllAOIntegralsIterator_4::AllAOIntegralsIterator_4 (
    std::shared_ptr< ShellCombinationsIterator > shellIter )
```

Parameters

<i>shellIter</i>	- shell iterator object
------------------	-------------------------

18.6.3 Member Function Documentation

18.6.3.1 index()

```
int oepdev::AllAOIntegralsIterator_4::index (
    void ) const [inline], [virtual]
```

Grab the current index of integral value stored in the buffer

Implements [oepdev::AOIntegralsIterator](#).

The documentation for this class was generated from the following files:

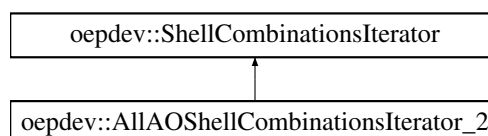
- oepdev/libutil/[integrals_iter.h](#)
- oepdev/libutil/integrals_iter.cc

18.7 oepdev::AllAOShellCombinationsIterator_2 Class Reference

Loop over all possible ERI shells in a shell doublet.

```
#include <integrals_iter.h>
```

Inheritance diagram for oepdev::AllAOShellCombinationsIterator_2:



Public Member Functions

- [AllAOShellCombinationsIterator_2](#) (SharedBasisSet [bs_1](#), SharedBasisSet [bs_2](#))

Iterate over shell doublets. Construct by providing basis sets for each axis. The basis sets must be defined for the same molecule.

- [AllAOShellCombinationsIterator_2](#) (std::shared_ptr< [IntegralFactory](#) > integrals)

Construct by providing integral factory.

- [AllAOShellCombinationsIterator_2](#) (const [IntegralFactory](#) &integrals)
- [AllAOShellCombinationsIterator_2](#) (std::shared_ptr< psi::IntegralFactory > integrals)

Construct by providing integral factory.

- [AllAOShellCombinationsIterator_2](#) (const psi::IntegralFactory &integrals)
- void [first](#) ()

First iteration.

- void [next](#) ()

Next iteration.

- void [compute_shell](#) (std::shared_ptr< [oepdev::TwoBodyAOInt](#) > tei) const

Compute ERI's for the current shell. The eris are stored in the buffer of the argument object.

- void **compute_shell** (std::shared_ptr< psi::TwoBodyAOInt > tei) const
- int [P](#) () const

Grab the current shell P index.

- int [Q](#) () const

Grab the current shell Q index.

Additional Inherited Members

18.7.1 Detailed Description

Constructed by providing [IntegralFactory](#) object or shared pointers to two basis set spaces.

18.7.2 Constructor & Destructor Documentation

18.7.2.1 AllAOShellCombinationsIterator_2() [1/5]

```
AllAOShellCombinationsIterator_2::AllAOShellCombinationsIterator_2 (
    SharedBasisSet bs_1,
    SharedBasisSet bs_2 )
```

Parameters

<i>bs_1</i>	- basis set of axis 1
<i>bs_2</i>	- basis set of axis 2

18.7.2.2 AllAOShellCombinationsIterator_2() [2/5]

```
oepdev::AllAOShellCombinationsIterator_2::AllAOShellCombinationsIterator_2 (
    std::shared_ptr< IntegralFactory > integrals )
```

Parameters

<i>integrals</i>	- OepDev integral factory object
------------------	----------------------------------

18.7.2.3 AllAOShellCombinationsIterator_2() [3/5]

```
AllAOShellCombinationsIterator_2::AllAOShellCombinationsIterator_2 (
    const IntegralFactory & integrals )
```

This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

18.7.2.4 AllAOShellCombinationsIterator_2() [4/5]

```
AllAOShellCombinationsIterator_2::AllAOShellCombinationsIterator_2 (
    std::shared_ptr< psi::IntegralFactory > integrals )
```

Parameters

<i>integrals</i>	- Psi4 integral factory object
------------------	--------------------------------

18.7.2.5 AllAOShellCombinationsIterator_2() [5/5]

```
AllAOShellCombinationsIterator_2::AllAOShellCombinationsIterator_2 (
    const psi::IntegralFactory & integrals )
```

This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

18.7.3 Member Function Documentation**18.7.3.1 compute_shell()**

```
void AllAOShellCombinationsIterator_2::compute_shell (
    std::shared_ptr< oepdev::TwoBodyAOInt > tei ) const [virtual]
```


Parameters

<i>tei</i>	- two electron AO integral
------------	----------------------------

Implements [oepdev::ShellCombinationsIterator](#).

The documentation for this class was generated from the following files:

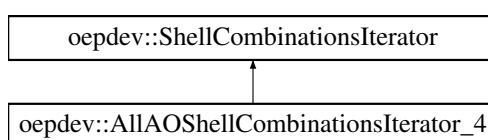
- oepdev/libutil/integrals_iter.h
- oepdev/libutil/integrals_iter.cc

18.8 oepdev::AllAOShellCombinationsIterator_4 Class Reference

Loop over all possible ERI shells in a shell quartet.

```
#include <integrals_iter.h>
```

Inheritance diagram for oepdev::AllAOShellCombinationsIterator_4:



Public Member Functions

- [AllAOShellCombinationsIterator_4](#) (SharedBasisSet [bs_1](#), SharedBasisSet [bs_2](#), SharedBasisSet [bs_3](#), SharedBasisSet [bs_4](#))
Iterate over shell quartets. Construct by providing basis sets for each axis. The basis sets must be defined for the same molecule.
- [AllAOShellCombinationsIterator_4](#) (std::shared_ptr< [IntegralFactory](#) > integrals)
Construct by providing integral factory.
- [AllAOShellCombinationsIterator_4](#) (const [IntegralFactory](#) &integrals)
- [AllAOShellCombinationsIterator_4](#) (std::shared_ptr< psi::IntegralFactory > integrals)
Construct by providing integral factory.
- [AllAOShellCombinationsIterator_4](#) (const psi::IntegralFactory &integrals)
- void [first](#) ()
Do the first iteration.
- void [next](#) ()
Do the next iteration.
- void [compute_shell](#) (std::shared_ptr< [oepdev::TwoBodyAOInt](#) > [tei](#)) const
- void [compute_shell](#) (std::shared_ptr< psi::TwoBodyAOInt > [tei](#)) const
- int [P](#) () const
Grab the current shell P index.

- `int Q () const`
Grab the current shell Q index.
- `int R () const`
Grab the current shell R index.
- `int S () const`
Grab the current shell S index.

Additional Inherited Members

18.8.1 Detailed Description

Constructed by providing [IntegralFactory](#) object or shared pointers to four basis set spaces.

18.8.2 Constructor & Destructor Documentation

18.8.2.1 AllAOShellCombinationsIterator_4() [1/5]

```
AllAOShellCombinationsIterator_4::AllAOShellCombinationsIterator_4 (
    SharedBasisSet bs_1,
    SharedBasisSet bs_2,
    SharedBasisSet bs_3,
    SharedBasisSet bs_4 )
```

Parameters

<i>bs_1</i>	- basis set of axis 1
<i>bs_2</i>	- basis set of axis 2
<i>bs_3</i>	- basis set of axis 3
<i>bs_4</i>	- basis set of axis 4

18.8.2.2 AllAOShellCombinationsIterator_4() [2/5]

```
oepdev::AllAOShellCombinationsIterator_4::AllAOShellCombinationsIterator_4 (
    std::shared_ptr< IntegralFactory > integrals )
```

Parameters

<i>integrals</i>	- OepDev integral factory object
------------------	----------------------------------

18.8.2.3 AllAOShellCombinationsIterator_4() [3/5]

```
AllAOShellCombinationsIterator_4::AllAOShellCombinationsIterator_4 (
    const IntegralFactory & integrals )
```

This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

18.8.2.4 AllAOShellCombinationsIterator_4() [4/5]

```
AllAOShellCombinationsIterator_4::AllAOShellCombinationsIterator_4 (
    std::shared_ptr< psi::IntegralFactory > integrals )
```

Parameters

<i>integrals</i>	- OepDev integral factory object
------------------	----------------------------------

18.8.2.5 AllAOShellCombinationsIterator_4() [5/5]

```
AllAOShellCombinationsIterator_4::AllAOShellCombinationsIterator_4 (
    const psi::IntegralFactory & integrals )
```

This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

18.8.3 Member Function Documentation**18.8.3.1 compute_shell()** [1/2]

```
void AllAOShellCombinationsIterator_4::compute_shell (
    std::shared_ptr< oepdev::TwoBodyAOInt > tei ) const [virtual]
```

Compute integrals in a current shell. Works both for [oepdev::TwoBodyAOInt](#) and [psi::TwoBodyAOInt](#)

Parameters

<i>tei</i>	- two body integral object
------------	----------------------------

Implements [oepdev::ShellCombinationsIterator](#).

18.8.3.2 compute_shell() [2/2]

```
void oepdev::AllAOShellCombinationsIterator_4::compute_shell (
    std::shared_ptr< psi ::TwoBodyAoint > tei ) const [virtual]
```

Compute integrals in a current shell. Works both for [oepdev::TwoBodyAoint](#) and [psi::TwoBodyAoint](#)

Parameters

<i>tei</i>	- two body integral object
------------	----------------------------

Implements [oepdev::ShellCombinationsIterator](#).

The documentation for this class was generated from the following files:

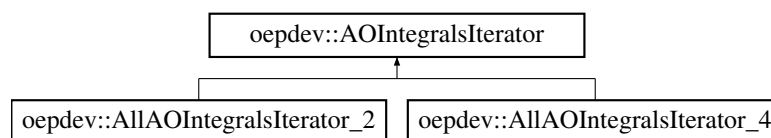
- [oepdev/libutil/integrals_iter.h](#)
- [oepdev/libutil/integrals_iter.cc](#)

18.9 oepdev::AOIntegralsIterator Class Reference

Iterator for AO Integrals. Abstract Base.

```
#include <integrals_iter.h>
```

Inheritance diagram for [oepdev::AOIntegralsIterator](#):



Public Member Functions

- [AOIntegralsIterator](#) ()
Base Constructor.
- virtual [~AOIntegralsIterator](#) ()
Base Destructor.
- virtual void [first](#) (void)=0
Do the first iteration.
- virtual void [next](#) (void)=0
Do the next iteration.
- virtual int [i](#) (void) const
Grab i-th index.
- virtual int [j](#) (void) const
Grab j-th index.

- virtual int `k` (void) const
Grab k-th index.
- virtual int `l` (void) const
Grab l-th index.
- virtual int `index` (void) const =0
Grab index in the integral buffer.
- virtual bool `is_done` (void)
Returns the status of an iterator.

Static Public Member Functions

- static std::shared_ptr< [AOIntegralsIterator](#) > `build` (const [ShellCombinationsIterator](#) *shellIter, std::string mode="ALL")
- static std::shared_ptr< [AOIntegralsIterator](#) > `build` (std::shared_ptr< [ShellCombinationsIterator](#) > shellIter, std::string mode="ALL")

Protected Attributes

- bool `done`
The status of an iterator.

18.9.1 Detailed Description

18.9.2 Member Function Documentation

18.9.2.1 `build()` [1/2]

```
std::shared_ptr< AOIntegralsIterator > AOIntegralsIterator::build (
    const ShellCombinationsIterator * shellIter,
    std::string mode = "ALL" ) [static]
```

Build AO integrals iterator from current state of iterator over shells

Parameters

<code>shellIter</code>	- iterator over shells - either "ALL" or "UNIQUE" (iterate over all or unique integrals)
------------------------	--

Returns

iterator over AO integrals

18.9.2.2 build() [2/2]

```
std::shared_ptr< AOIntegralsIterator > AOIntegralsIterator::build (
    std::shared_ptr< ShellCombinationsIterator > shellIter,
    std::string mode = "ALL" ) [static]
```

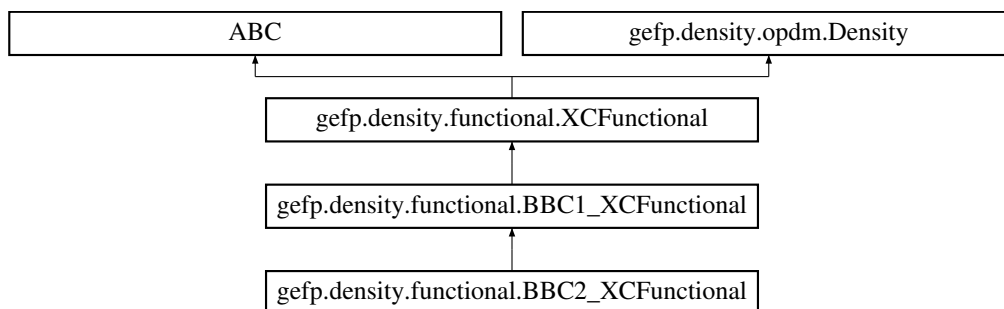
This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

The documentation for this class was generated from the following files:

- oepdev/libutil/[integrals_iter.h](#)
- oepdev/libutil/integrals_iter.cc

18.10 gefp.density.functional.BBC1_XCFunctional Class Reference

Inheritance diagram for gefp.density.functional.BBC1_XCFunctional:



Public Member Functions

- def **__init__** (self)
- def **abbr** (self)

Static Public Member Functions

- def **name** ()
- def **fij** (n)

Additional Inherited Members

18.10.1 Detailed Description

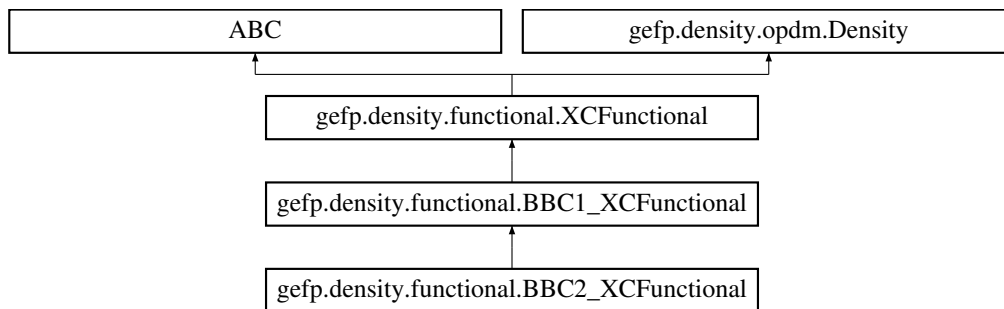
The BBC1 Exchange-Correlation Functional.

The documentation for this class was generated from the following file:

- gefp/gefp/density/functional.py

18.11 gefp.density.functional.BBC2_XCFunctional Class Reference

Inheritance diagram for gefp.density.functional.BBC2_XCFunctional:



Public Member Functions

- `def __init__ (self)`
- `def abbr (self)`

Static Public Member Functions

- `def name ()`
- `def fij (n)`

Additional Inherited Members

18.11.1 Detailed Description

The BBC2 Exchange-Correlation Functional.

The documentation for this class was generated from the following file:

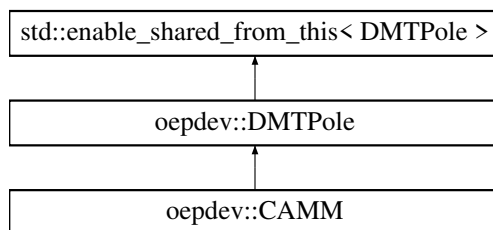
- `gefp/gefp/density/functional.py`

18.12 oepdev::CAMM Class Reference

Cumulative Atomic Multipole Moments.

```
#include <dmt.h>
```

Inheritance diagram for oepdev::CAMM:



Public Member Functions

- [CAMM](#) (psi::SharedWavefunction wfn, int n)
Construct [CAMM DMTPole](#) object.
- virtual void [compute](#) (psi::SharedMatrix D, bool transition, int n)
Compute DMTP's from the one-particle density matrix.
- virtual void [print_header](#) (void) const
Print the header.

Additional Inherited Members

18.12.1 Detailed Description

Cumulative atomic multipole representation of the molecular charge distribution. Method of Sokalski and Poirier. Ref.: W. A. Sokalski and R. A. Poirier, *Chem. Phys. Lett.*, 98(1) **1983**

Methodology.

The distributed multipole moments are computed in the following way:

- first the atomic additive multipole moments (AAMM's) with origins set to the global coordinate system origin are computed. AO basis set partitioning is used to distribute the AAMM's onto the atomic centres.
- subsequently, the AAMM's origins are moved to the corresponding atomic site.

The computation of the AAMM's is performed according to the following prescription:

$$M_{uw\dots z}^{(A)}(\mathbf{0}) = \sum_{\alpha \in A} \sum_{\beta \in \text{all AO's}} D_{\alpha\beta}^{\text{OED}} \langle \alpha | \mathcal{M}_{uw\dots z}(\mathbf{0}) | \beta \rangle$$

where $M_{uw\dots z}^{(A)}$ denotes the $(uw\dots z)$ -th component of the multipole centered at atomic site A , the symbol $\mathcal{M}(\mathbf{0})$ is the associated quantum mechanical operator and $D_{\alpha\beta}^{\text{OED}}$ is the (generalized) one-particle density matrix element in AO basis (Greek indices).

Recentering of the multipole moments is described in the documentation of [oepdev::DMTPole::recenter](#).

The documentation for this class was generated from the following files:

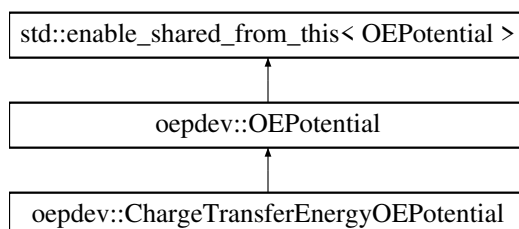
- [oepdev/lib3d/dmtp.h](#)
- [oepdev/lib3d/dmtp_camm.cc](#)

18.13 oepdev::ChargeTransferEnergyOEPotential Class Reference

Generalized One-Electron Potential for Charge-Transfer Interaction Energy.

```
#include <oep.h>
```

Inheritance diagram for oepdev::ChargeTransferEnergyOEPotential:



Public Member Functions

- **ChargeTransferEnergyOEPotential** (SharedWavefunction [wfn](#), SharedBasisSet auxiliary, SharedBasisSet intermediate, Options &options)
- **ChargeTransferEnergyOEPotential** (SharedWavefunction [wfn](#), Options &options)
- virtual void [compute](#) (const std::string &oepType) override
Compute matrix forms of all OEP's within a specified OEP type.
- virtual void [compute_3D](#) (const std::string &oepType, const double &x, const double &y, const double &z, std::shared_ptr< psi::Vector > &v) override
Compute value of potential in point x, y, z and save at v.
- virtual void [print_header](#) () const override
Header information.

Additional Inherited Members

18.13.1 Detailed Description

Contains the following OEP types:

- `Otto-Ladik.V1.GDF` - DF-based term (group I)
- `Otto-Ladik.V3.CAMM-nj` - CAMM-based term (group III; truncated on distributed charges)

Group II terms do not require any particular OEP's due to great simplification of this term. Atomic numbers and LMO centroids are sufficient.

The documentation for this class was generated from the following files:

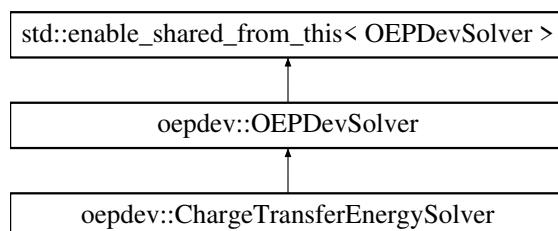
- oepdev/liboep/[oep.h](#)
- oepdev/liboep/oep_energy_ct.cc

18.14 oepdev::ChargeTransferEnergySolver Class Reference

Compute the Charge-Transfer interaction energy between unperturbed wavefunctions.

```
#include <solver.h>
```

Inheritance diagram for oepdev::ChargeTransferEnergySolver:



Public Member Functions

- **ChargeTransferEnergySolver** (SharedWavefunctionUnion wfn_union)
- virtual double [compute_oep_based](#) (const std::string &method="DEFAULT")
Compute property by using OEP's.
- virtual double [compute_benchmark](#) (const std::string &method="DEFAULT")
Compute property by using benchmark method.

Additional Inherited Members

18.14.1 Detailed Description

The implemented methods are shown below

Table 18.15: Methods available in the Solver

Keyword	Method Description
Benchmark Methods	
OTTO_LADIK	<i>Default.</i> CT energy at HF level from Otto and Ladik (1975).
EFP2	CT energy at HF level from EFP2 model.
OEP-Based Methods	
OTTO_LADIK	<i>Default.</i> OEP-based Otto-Ladik expressions.

In order to construct this solver, **always** use the [OEPDevSolver::build](#) static factory method.

Below the detailed description of the implemented equations is given for each of the above provided methods. In the formulae across, it is assumed that the orbitals are real. The Coulomb notation for electron repulsion integrals (ERI's) is adopted; i.e.,

$$(ac|bd) = \iint d\mathbf{r}_1 d\mathbf{r}_2 \phi_a(\mathbf{r}_1) \phi_c(\mathbf{r}_1) \frac{1}{r_{12}} \phi_b(\mathbf{r}_2) \phi_d(\mathbf{r}_2)$$

Greek subscripts denote basis set orbitals whereas *Italic* subscripts denote the occupied molecular orbitals.

The CT energy between molecules *A* and *B* is given by

$$E^{\text{CT}} = E^{A^+B^-} + E^{A^-B^+}$$

Benchmark Methods

CT energy at HF level by Otto and Ladik (1975).

For a closed-shell system, CT energy equation of Otto and Ladik becomes

$$E^{A^+B^-} \approx 2 \sum_{i \in A}^{\text{Occ}_A} \sum_{n \in B}^{\text{Vir}_B} \frac{V_{in}^2}{\epsilon_i - \epsilon_n}$$

where

$$\begin{aligned} V_{in} = & V_{in}^B + 2 \sum_{j \in B}^{\text{Occ}_B} (in|jj) - \sum_{k \in A}^{\text{Occ}_A} S_{kn} \left\{ V_{ik}^B + 2 \sum_{j \in B}^{\text{Occ}_B} (ik|jj) \right\} \\ & - \sum_{j \in B}^{\text{Occ}_B} \left[S_{ij} \left\{ V_{nj}^A + 2 \sum_{k \in A}^{\text{Occ}_A} (1 - \delta_{ik})(nj|kk) \right\} + (nj|ij) \right] + \sum_{k \in A}^{\text{Occ}_A} \sum_{j \in B}^{\text{Occ}_B} S_{kj} (1 - \delta_{ik})(ik|nj) \end{aligned}$$

and analogously the twin term.

CT energy at HF level by EFP2.

In EFP2 method, CT energy is given as

$$E^{A^+B^-} \approx 2 \sum_{i \in A}^{\text{Occ}_A} \sum_{n \in B}^{\text{Vir}_B} \frac{V_{in}^2}{F_{ii} - T_{nn}}$$

where

$$V_{in}^2 = \frac{V_{in}^{EF,B} - \sum_{m \in A}^{\text{All}_A} V_{im}^{EF,B} S_{mn}^B}{1 - \sum_{m \in A}^{\text{All}_A} S_{mn}^2} \left\{ V_{in}^{EF,B} - \sum_{m \in A}^{\text{All}_A} V_{im}^{EF,B} S_{mn} + \sum_{j \in B}^{\text{Occ}_B} S_{ij} \left(T_{nj} - \sum_{m \in A}^{\text{All}_A} S_{nm} T_{mj} \right) \right\}$$

and analogously the twin term.

OEP-Based Methods

OEP-Based Otto-Ladik's theory

After introducing OEP's, the original Otto-Ladik's theory is reformulated *without* approximation as

$$E^{A+B-} \approx 2 \sum_{i \in A}^{\text{Occ}_A} \sum_{n \in B}^{\text{Vir}_B} \frac{\left(V_{in}^{\text{DF}} + V_{in}^{\text{ESP,A}} + V_{in}^{\text{ESP,B}} \right)^2}{\epsilon_i - \epsilon_n}$$

where

$$\begin{aligned} V_{in}^{\text{DF}} &= \sum_{\eta \in B}^{\text{Aux}_B} S_{i\eta} G_{\eta n}^B \\ V_{in}^{\text{ESP,A}} &= \sum_{k \in A}^{\text{Occ}_A} \sum_{j \in B}^{\text{Occ}_B} S_{kj} \sum_{x \in A} V_{nj}^{(x)} q_{ik}^{(x)} \\ V_{in}^{\text{ESP,B}} &= - \sum_{k \in A}^{\text{Occ}_A} S_{kn} V_{ik}^B \end{aligned}$$

The OEP matrix for density fitted part is given by

$$G_{\eta n}^B = \sum_{\eta' \in B}^{\text{Aux}_B} [\mathbf{S}^{-1}]_{\eta\eta'} \left\{ V_{\eta'n}^B + \sum_{j \in B}^{\text{Occ}_B} [2(\eta'n|jj) - (\eta'j|nj)] \right\}$$

The OEP ESP-A charges are fit to reproduce the OEP potential

$$v_{ik}^A(\mathbf{r}) \equiv (1 - \delta_{ik}) \int \frac{\phi_i(\mathbf{r}') \phi_k(\mathbf{r}')}{|\mathbf{r} - \mathbf{r}'|} d\mathbf{r}' - \delta_{ik} \left(\sum_{x \in A} \frac{-Z_x}{|\mathbf{r} - \mathbf{r}_x|} + 2 \sum_{k \in A}^{\text{Occ}_A} \int \frac{\phi_k(\mathbf{r}') \phi_k(\mathbf{r}')}{|\mathbf{r} - \mathbf{r}'|} d\mathbf{r}' - 2 \int \frac{\phi_i(\mathbf{r}') \phi_i(\mathbf{r}')}{|\mathbf{r} - \mathbf{r}'|} d\mathbf{r}' \right)$$

so that

$$v_{ik}^A(\mathbf{r}) \cong \sum_{x \in A} \frac{q_{ik}^{(x)}}{|\mathbf{r} - \mathbf{r}_x|}$$

The OEP ESP-B charges are fit to reproduce the electrostatic potential of molecule *B* (they are standard ESP charges).

18.14.2 Member Function Documentation

18.14.2.1 compute_benchmark()

```
double ChargeTransferEnergySolver::compute_benchmark (
    const std::string & method = "DEFAULT" ) [virtual]
```

Each solver object has one `DEFAULT` benchmark method

Parameters

<i>method</i>	- benchmark method
---------------	--------------------

Implements [oepdev::OEPDevSolver](#).

18.14.2.2 compute_oep_based()

```
double ChargeTransferEnergySolver::compute_oep_based (
    const std::string & method = "DEFAULT" ) [virtual]
```

Each solver object has one DEFAULT OEP-based method.

Parameters

<i>method</i>	- flavour of OEP model
---------------	------------------------

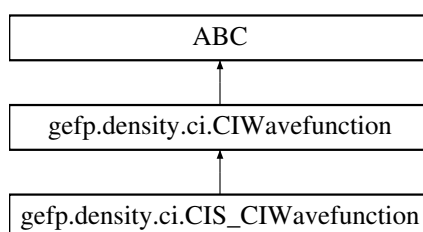
Implements [oepdev::OEPDevSolver](#).

The documentation for this class was generated from the following files:

- oepdev/libsolver/[solver.h](#)
- oepdev/libsolver/solver_energy_ct.cc

18.15 gefp.density.ci.CIS_CIWavefunction Class Reference

Inheritance diagram for gefp.density.ci.CIS_CIWavefunction:



Public Member Functions

- def **__init__** (self, ref_wfn, E, W)
- def **make_ci_l** (self)

Additional Inherited Members

The documentation for this class was generated from the following file:

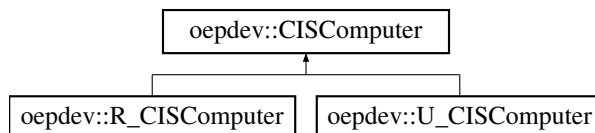
- `gefp/gefp/density/ci.py`

18.16 oepdev::CISComputer Class Reference

[CISComputer](#).

```
#include <cis.h>
```

Inheritance diagram for oepdev::CISComputer:



Public Member Functions

- virtual `~CISComputer` ()
Destructor.
- virtual void `compute` (void)
Solve the CIS problem.
- virtual void `clear_dpd` (void)
Clear DPD instance.
- int `nstates` (void) const
Get the total number of excited states.
- SharedVector `eigenvalues` () const
Get the CIS eigenvalues.
- SharedVector `E` () const
- SharedMatrix `eigenvectors` () const
Get the CIS eigenvectors.
- SharedMatrix `U` () const
- std::pair< double, double > `U_homo_lumo` (int l, int h=0, int l=0) const
*Get the HOMO+*h*->LUMO+*l* CIS coefficient for a given excited state l for spin alpha and beta.*
- SharedMatrix `Da_mo` (int i) const
Compute MO one-particle alpha density matrix for state i
- SharedMatrix `Db_mo` (int i) const
Compute MO one-particle beta density matrix for state i
- SharedMatrix `Da_ao` (int i) const
Compute AO one-particle alpha density matrix for state i
- SharedMatrix `Db_ao` (int i) const
Compute AO one-particle beta density matrix for state i

- SharedDMTPole [cammm](#) (int j, bool symmetrize=false) const
Compute CAMM for j excited state.
- SharedMatrix [Ta_ao](#) (int j) const
*Compute MO one-particle alpha 0->*j* transition density matrix.*
- SharedMatrix [Tb_ao](#) (int j) const
*Compute MO one-particle beta 0->*j* transition density matrix.*
- SharedMatrix [Ta_ao](#) (int i, int j) const
*Compute MO one-particle alpha i->*j* transition density matrix.*
- SharedMatrix [Tb_ao](#) (int i, int j) const
*Compute MO one-particle beta i->*j* transition density matrix.*
- SharedDMTPole [trcammm](#) (int j, bool symmetrize=true) const
*Compute TrCAMM for 0->*j* transition.*
- SharedDMTPole [trcammm](#) (int i, int j, bool symmetrize=true) const
*Compute TrCAMM for i->*j* transition.*
- SharedVector [transition_dipole](#) (int j) const
*Compute transition dipole moment for 0->*j* transition.*
- SharedVector [transition_dipole](#) (int i, int j) const
*Compute transition dipole moment for i->*j* transition.*
- double [oscillator_strength](#) (int j) const
*Compute oscillator strength for 0->*j* transition.*
- double [oscillator_strength](#) (int i, int j) const
*Compute oscillator strength for i->*j* transition.*
- void [determine_electronic_state](#) (int &l)
Determine electronic state.
- std::shared_ptr< [CISData](#) > [data](#) (int l, bool symmetrize_trcammm=false)
Return CIS data structure for a given excited state l

Static Public Member Functions

- static std::shared_ptr< [CISComputer](#) > [build](#) (const std::string &type, std::shared_ptr< psi::Wavefunction > wfn, psi::Options &opt, const std::string &reference="")
Build CIS Computer.

Static Public Attributes

- static const std::vector< std::string > [reference_types](#) = {"RHF", "UHF"}
Slater determinant possible references, that are implemented.

Protected Member Functions

- **CISComputer** (std::shared_ptr< psi::Wavefunction > wfn, psi::Options &opt, psi::IntegralTransform::TransformationType trans_type)
- virtual void **prepare_for_cis_** (void)
- virtual void **build_hamiltonian_** (void)=0
- virtual void **diagonalize_hamiltonian_** (void)
- virtual void **set_beta_** (void)=0
- virtual void **transform_integrals_** (void)

Protected Attributes

- std::shared_ptr< psi::Wavefunction > **ref_wfn_**
Reference wavefunction.
- psi::Options **options_**
Psi4 Options.
- const int **nmo_**
Number of MO's.
- const int **naocc_**
Number of alpha occupied MO's.
- const int **nbocc_**
Number of beta occupied MO's.
- const int **navir_**
Number of alpha virtual MO's.
- const int **nbvir_**
Number of beta virtual MO's.
- int **ndets_**
Number of excited determinants.
- SharedMatrix **H_**
CIS Excited State Hamiltonian in Slater determinantal basis.
- SharedMatrix **U_**
CIS Coefficients U_{uI} for each excited state I and basis Slater determinant u
- SharedVector **E_**
Electronic excitation energies E_I wrt ground state.
- SharedVector **eps.a.o_**
- SharedVector **eps.a.v_**
- SharedVector **eps.b.o_**
- SharedVector **eps.b.v_**
- const psi::IntegralTransform::TransformationType **transformation_type_**
MO Integral Transformation Type.
- std::shared_ptr< psi::IntegralTransform > **inttrans_**

18.16.1 Detailed Description

18.16.2 Member Function Documentation

18.16.2.1 build()

```
std::shared_ptr< CISComputer > oepdev::CISComputer::build (
    const std::string & type,
    std::shared_ptr< psi::Wavefunction > wfn,
    psi::Options & opt,
    const std::string & reference = "" ) [static]
```

Parameters

<i>type</i>	- Type of computer
<i>wfn</i>	- Psi4 wavefunction
<i>opt</i>	- Psi4 options
<i>reference</i>	- Reference Slater determinant (RHF, UHF available).

Available computer types:

- RESTRICTED or RCIS - RHF wavefunction is used as reference state
- UNRESTRICTED or UCIS - UHF wavefunction is used as reference state

Implementation

The CIS Hamiltonian in the basis space of singly-excited Slater determinants is constructed from canonical molecular orbitals (CMO's)

$$\begin{aligned}\langle \Phi_0 | \mathcal{H} | \Phi_i^a \rangle &= 0 \\ \langle \Phi_j^b | \mathcal{H} | \Phi_i^a \rangle &= \delta_{ij} \delta_{ab} (\epsilon_a - \epsilon_i) + \langle a j | i b \rangle - \langle a j | b i \rangle\end{aligned}$$

where i labels the occupied CMO's whereas a labels the virtual CMO's. In the above equation, $\langle a j | i b \rangle$ is the 2-electron 4-centre integral in physicist's notation. After integrating out the spin coordinate, four blocks of Hamiltonian are explicitly given as

$$\begin{aligned}\langle \Phi_j^b | \mathcal{H} | \Phi_i^a \rangle &= \delta_{ij} \delta_{ab} (\epsilon_a - \epsilon_i) + [i a | j b] - [a b | i j] \\ \langle \Phi_{\bar{j}}^{\bar{b}} | \mathcal{H} | \Phi_{\bar{i}}^{\bar{a}} \rangle &= \delta_{\bar{i}\bar{j}} \delta_{\bar{a}\bar{b}} (\epsilon_{\bar{a}} - \epsilon_{\bar{i}}) + [\bar{i} \bar{a} | \bar{j} \bar{b}] - [\bar{a} \bar{b} | \bar{i} \bar{j}] \\ \langle \Phi_{\bar{j}}^{\bar{b}} | \mathcal{H} | \Phi_i^a \rangle &= [i a | \bar{j} \bar{b}] \\ \langle \Phi_j^b | \mathcal{H} | \Phi_{\bar{i}}^{\bar{a}} \rangle &= [\bar{i} \bar{a} | j b]\end{aligned}$$

where the $[ia|jb]$ is the 2-electron 4-centre integral in the chemist's (Coulomb) notation.

Such matrix is diagonalized yielding the excitation energies (wrt HF ground state) as well as the CIS coefficients

$$\sum_{ij} \sum_{ab} t_{i,I}^a H_{ij}^{ab} t_{j,J}^b = E_I \delta_{IJ}$$

where the summations above extend over alpha and beta electron spin labels and $t_{i,I}^a$ is the CIS amplitude for the i th excited state, associated with the $i \rightarrow a$ excitation with respect to the HF reference determinant. Note that E_I is *not* the excited state energy, but the energy relative the the HF reference energy.

Transition density matrix

AO basis transition density from ground (HF) to excited (CIS) state is given by

$$P_{\mu\nu}^{(g \rightarrow e)} = \sum_i \sum_a^{\text{Occ Vir}} t_{i,e}^a C_{\nu i} C_{\mu a} + \sum_i \sum_{\bar{a}}^{\text{Occ Vir}} t_{i,e}^{\bar{a}} C_{\nu i} C_{\mu \bar{a}}$$

Excited state density matrix

CMO basis excited state density matrix for alpha spin is given by

Analogous expression is given for the beta spin.

AO representation of the CMO excited state density matrix is

$$P_{\mu\nu}^{(e)} = \sum_{pq} C_{\mu p} P_{pq}^{(e)} C_{\nu q} + \sum_{\bar{p}\bar{q}} C_{\mu \bar{p}} P_{\bar{p}\bar{q}}^{(e)} C_{\nu \bar{q}}$$

which is the sum of alpha and beta density matrices in CMO basis transformed to AO basis.

The CMO excited state density matrix for spin alpha is given by

$$P_{pq}^{(e)} = \begin{cases} \delta_{pq} - \sum_a^{\text{Vir}} t_{p,e}^a t_{q,e}^a & \text{for } p, q \in \text{Occ} \\ \sum_i^{\text{Occ}} t_{i,e}^p t_{i,e}^q & \text{for } p, q \in \text{Vir} \\ 0 & \text{otherwise} \end{cases}$$

The beta spin density matrix is generated analogously as above.

The cumulative atomic multipole moments ([Camm](#)) are computed from the excited state density matrices in AO basis. The nuclear contribution is included.

Transition multipole moments

The transition dipole moment is computed from the AO transition density matrix and the dipole integrals in AO basis, i.e.,

$$\langle \Phi_0 | \hat{\mu}_u | \Psi_e \rangle = \text{Tr} [\mathbf{d}^{(u)} \cdot \mathbf{P}^{g \rightarrow e}]$$

Oscillator strength is computed from the transition dipole moment via

$$f^{g \rightarrow e} = \frac{2}{3} E_e \left| \langle \Phi_0 | \hat{\mu} | \Psi_e \rangle \right|^2$$

Transition cumulative atomic multipole moments (TrCamm) are computed from the transition density matrices in AO basis. The nuclear contribution is not included.

Note

Useful options:

- `CIS_NSTATES` - Number of lowest-energy excited states to include. Default: `-1` (means all states are saved)
- For UHF references, SAD guess might lead to triplet instabilities. It is then better to set `CORE` as the UHF guess

The documentation for this class was generated from the following files:

- oepdev/libutil/cis.h
- oepdev/libutil/cis_base.cc

18.17 oepdev::CISData Struct Reference

Container to handle the CIS wavefunction parameters.

```
#include <cis.h>
```

Public Attributes

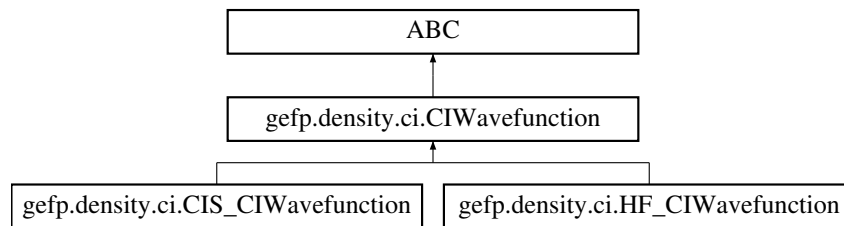
- double `E_ex`
Excitation energy.
- double `t_homo_lumo`
CIS HOMO-LUMO amplitude.
- SharedMatrix `Pe`
Excited state density matrix (sum of alpha and beta)
- SharedMatrix `Peg`
Transition ground-to-excited state density matrix (sum of alpha and beta)
- SharedDMTPole `trcamm`
TrCamm.
- SharedDMTPole `camm_homo`
Camm for HOMO orbital.
- SharedDMTPole `camm_lumo`
Camm for LUMO orbital.

The documentation for this struct was generated from the following file:

- oepdev/libutil/cis.h

18.18 gefp.density.ci.CIWavefunction Class Reference

Inheritance diagram for gefp.density.ci.CIWavefunction:



Public Member Functions

- def **__init__** (self, ref_wfn, E, W)
- def **make_ci_l** (self)
- def **overlap** (self, other)

Public Attributes

- **ref_wfn**
- **ci_e**
- **ci_c**
- **ca_o**
- **cb_o**
- **ca_v**
- **cb_v**
- **bfs**
- **naocc**
- **nbocc**
- **nmo**
- **navir**
- **nbvir**
- **ci_l**
- **ndet**

The documentation for this class was generated from the following file:

- gefp/gefp/density/ci.py

18.19 oepdev::CPHF Class Reference

[CPHF](#) solver class.

```
#include <cphf.h>
```

Public Member Functions

- [CPHF](#) (SharedWavefunction ref_wfn, Options &[options](#))
Constructor.
- [~CPHF](#) ()
Desctructor.
- void [compute](#) (void)
run the calculations
- void [print](#) (void) const
print to output file
- int [nocc](#) (void) const
get the number of occupied orbitals
- std::shared_ptr< Wavefunction > [wfn](#) (void) const
grab the wavefunction
- Options & [options](#) (void) const
grab the Psi4 options
- std::shared_ptr< Matrix > [polarizability](#) (void) const
retrieve the molecular (total) polarizability
- std::shared_ptr< Matrix > [polarizability](#) (int i) const
retrieve the i-th orbital-associated polarizability
- std::shared_ptr< Matrix > [polarizability](#) (int i, int j) const
retrieve the charge-transfer polarizability associated with orbitals i and j
- std::shared_ptr< Matrix > [X](#) (int x) const
retrieve the X operator O-V perturbation matrix in AO basis for x-th component
- std::vector< std::shared_ptr< Matrix > > [X](#) (void) const
retrieve the X operator O-V perturbation matrix in AO basis for all three Cartesian components
- std::shared_ptr< Matrix > [X_mo](#) (int x) const
retrieve the X operator O-V perturbation matrix in MO basis for x-th component
- std::vector< std::shared_ptr< Matrix > > [X_mo](#) (void) const
retrieve the X operator O-V perturbation matrix in MO basis for all three Cartesian components
- std::shared_ptr< Matrix > [F_mo](#) (int x) const
retrieve the F operator O-V perturbation matrix in MO basis for x-th component
- std::vector< std::shared_ptr< Matrix > > [F_mo](#) (void) const
retrieve the F operator O-V perturbation matrix in MO basis for all three Cartesian components
- std::shared_ptr< Matrix > [T](#) (void) const
retrieve the transformation from old to new MO's
- std::shared_ptr< Matrix > [Cocc](#) (void) const
retrieve the Cocc
- std::shared_ptr< Matrix > [Cvir](#) (void) const
retrieve the Cvir
- std::shared_ptr< Vector > [lmo_centroid](#) (int i) const

retrieve the i -th orbital (LMO) centroid

- `std::shared_ptr< Localizer > localizer` (void) const
retrieve the orbital localizer

Protected Attributes

- `std::shared_ptr< psi::Wavefunction > _wfn`
Wavefunction object.
- `std::shared_ptr< Localizer > _localizer`
Orbital localizer.
- `const int _no`
Number of occupied orbitals.
- `const int _nv`
Number of virtual orbitals.
- `const int _nn`
Number of basis functions.
- `long int _memory`
Memory.
- `int _maxiter`
Maximum number of iterations.
- `double _conv`
CPHF convergence threshold.
- `bool _with_diis`
whether use DIIS or not
- `const int _diis_dim`
Size of subspace.
- `std::shared_ptr< BasisSet > _primary`
Primary Basis Set.
- `std::shared_ptr< Matrix > _cocc`
Occupied orbitals.
- `std::shared_ptr< Matrix > _cvir`
Virtual orbitals.
- `std::shared_ptr< Vector > _eps_occ`
Occupied orbital energies.
- `std::shared_ptr< Vector > _eps_vir`
Virtual orbital energies.
- `std::vector< std::shared_ptr< oepdev::DIISManager > > _diis`
the DIIS managers for each perturbation operator x , y and z
- `Options & _options`
Options.

- `std::shared_ptr< Matrix > _molecularPolarizability`
Total (molecular) polarizability tensor.
- `std::vector< std::shared_ptr< Vector > > _orbitalCentroids`
LMO centroids.
- `std::vector< std::shared_ptr< Matrix > > _orbitalPolarizabilities`
orbital-associated polarizability tensors
- `std::vector< std::vector< std::shared_ptr< Matrix > > > _orbitalChargeTransferPolarizabilities`
orbital-orbital charge-transfer polarizability tensors
- `std::vector< std::shared_ptr< Matrix > > _X_OV_ao_matrices`
Perturbation X Operator O->V matrices in AO basis.
- `std::vector< std::shared_ptr< Matrix > > _X_OV_mo_matrices`
Perturbation X Operator O->V matrices in MO basis.
- `std::vector< std::shared_ptr< Matrix > > _F_OV_mo_matrices`
Electric Field Operator O->V matrices in MO basis.
- `std::shared_ptr< psi::Matrix > _T`
Transformation from old to new MO's.

18.19.1 Detailed Description

Solves [CPHF](#) equations (now only for RHF wavefunction). Computes molecular and polarizabilities associated with the localized molecular orbitals (LMO).

Note

Useful options:

- `CPHF_CONVER` - convergence of [CPHF](#). Default: `1e-8 (au)`
- `CPHF_CONVER` - maximum number of iterations. Default: `50`
- `CPHF_DIIS` - wheather use DIIS or not. Default: `true`
- `CPHF_DIIS_DIM` - dimension of iterative subspace. Default: `3`
- `CPHF_LOCALIZE` - localize the molecular orbitals? Default: `true`
- `CPHF_LOCALIZER` - set orbital localization method. Available: `BOYS` and `PIPEK_MEZEY`. Default: `BOYS`

18.19.2 Constructor & Destructor Documentation

18.19.2.1 CPHF()

```
oepdev::CPHF::CPHF (
    SharedWavefunction ref_wfn,
    Options & options )
```

Parameters

<i>ref_wfn</i>	reference HF wavefunction
<i>options</i>	set of Psi4 options

The documentation for this class was generated from the following files:

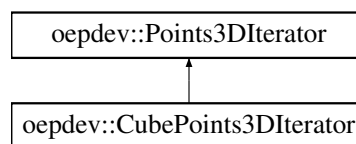
- oepdev/libutil/cphf.h
- oepdev/libutil/cphf.cc

18.20 oepdev::CubePoints3DIterator Class Reference

Iterator over a collection of points in 3D space. g09 Cube-like order.

```
#include <space3d.h>
```

Inheritance diagram for oepdev::CubePoints3DIterator:



Public Member Functions

- **CubePoints3DIterator** (const int &nx, const int &ny, const int &nz, const double &dx, const double &dy, const double &dz, const double &ox, const double &oy, const double &oz)
- virtual void [first](#) ()
Initialize first iteration.
- virtual void [next](#) ()
Step to next iteration.

Protected Attributes

- const int **nx_**
- const int **ny_**
- const int **nz_**
- const double **dx_**
- const double **dy_**
- const double **dz_**
- const double **ox_**
- const double **oy_**

- const double **oz_**
- int **ii_**
- int **jj_**
- int **kk_**

Additional Inherited Members

18.20.1 Detailed Description

Note: Always create instances by using static factory method from [Points3DIterator](#). Do not use constructor of this class.

The documentation for this class was generated from the following files:

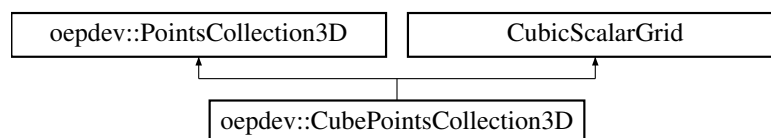
- oepdev/lib3d/space3d.h
- oepdev/lib3d/space3d.cc

18.21 oepdev::CubePointsCollection3D Class Reference

G09 cube-like ordered collection of points in 3D space.

```
#include <space3d.h>
```

Inheritance diagram for oepdev::CubePointsCollection3D:



Public Member Functions

- **CubePointsCollection3D** ([Collection](#) collectionType, const int &nx, const int &ny, const int &nz, const double &px, const double &py, const double &pz, psi::SharedBasisSet bs, psi::Options &options)
- virtual void [print](#) () const
Print the information to Psi4 output file.
- virtual void **write_cube_file** (psi::SharedMatrix v, const std::string &name, const int &col=0)

Additional Inherited Members

18.21.1 Detailed Description

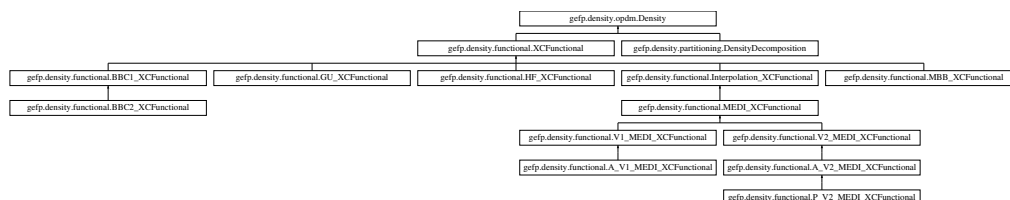
Note: Do not use constructors of this class explicitly. Instead, use static factory methods of the superclass to create instances.

The documentation for this class was generated from the following files:

- oepdev/lib3d/space3d.h
- oepdev/lib3d/space3d.cc

18.22 gefp.density.opdm.Density Class Reference

Inheritance diagram for gefp.density.opdm.Density:



Public Member Functions

- def **__init__** (self, D=None, jk=None)
- def **matrix** (self)
- def **set_D** (self, D)
- def **set_jk** (self, jk)
- def **compute_1el_energy** (self, D, Hcore)
- def **compute_2el_energy** (self, D_left, D_right, type='j')
- def **generalized_JK** (self, D, type='j')
- def **natural_orbitals** (cls, D, S=None, C=None, orthogonalize_mo=True, order='descending', return_ao_orthogonal=False, renormalize=False, no_cutoff=False, ignore_large_n=False, n_eps=5.0E-5)
- def **generalized_density** (cls, n, c, g=1.0)
- def **orthogonalize_OPDM** (cls, D, S)
- def **deorthogonalizer** (cls, S)
- def **orthogonalizer** (cls, S)

18.22.1 Detailed Description

Electron Density

Handles the Electron Density Distribution.

Usage as container class:

- 1) Initialize container object:

```
density = Density(D = None, jk = None)
```

where:

- o D - the density matrix in AO or MO basis
- o jk - the psi4::JK object for AO basis JK calculations

- 2) Grab the density matrix

```
D = density.matrix()
```

- 3) Computations in AO basis:

- o compute 1-electron energy (does not require JK object to be set)

```
e_1 = density.compute_1el_energy(D, V1)
```

The below require jk to be set:

- o compute 2-electron energy (J-type expression)

```
e_2j = density.compute_2el_energy(D_left, D_right, type='j')
```

- o compute 2-electron energy (K-type expression)

```
e_2k = density.compute_2el_energy(D_left, D_right, type='k')
```

- o compute J matrix (or K matrix if type=='k'):

```
J = density.generalized_JK(D, type='j')
```

Usage as method class.

Using 'Density' as a class of methods do not require object initialization.
The list of class methods is given below:

- o Density.natural_orbitals - compute natural orbitals
- o Density.generalized_density - compute generalized OPDM
- o Density.orthogonalize_OPDM - compute orthogonalized OPDM
- o Density.orthogonalizer - compute orthogonalizer matrix
- o Density.deorthogonalizer - compute deorthogonalizer rmatrix

Usage:

```
result = Density.'class method name'
```

See respective documentation for each of them for further details.

Last Revis

18.22.2 Member Function Documentation

18.22.2.1 `natural_orbitals()`

```
def gefp.density.opdm.Density.natural_orbitals (
    cls,
    D,
    S = None,
    C = None,
    orthogonalize_mo = True,
    order = 'descending',
    return_ao_orthogonal = False,
    renormalize = False,
    no_cutoff = False,
    ignore_large_n = False,
    n_eps = 5.0E-5 )
```

 Compute the Natural Orbitals from a given OPDM

Usage:

```
n, c = Density.natural_orbitals(D, S = None, C = None,
                                orthogonalize_mo = True,
                                order = 'descending',
                                return_ao_orthogonal = False,
                                renormalize = False,
                                no_cutoff = False,
                                ignore_large_n = False,
                                n_eps = 5.0E-5)
```

where:

- o D - OPDM in AO or MO basis
- o S - overlap integrals in AO or MO basis
- o C - LCAO-MO transformation matrix
- o orthogonalize_mo - whether to transform D from AO to certain MO basis and diagonalize
- o order - order in which eigenvalues (occupancies) are sorted. Eigenvalues are sorted in descending order
- o return_ao_orthogonal - whether to return NO's in oAO basis set or not
- o renormalize - renormalize to integer number of electrons
- o no_cutoff - cut-off threshold for occupancies
- o ignore_large_n - raise ValueError if $(1.0 + n_eps) < n < (0.0 - n_eps)$
- o n_eps - tolerance for occupancy deviation

 Examples:

- 1) NO's in AO (non-orthogonal, original) basis from D in AO basis

```
n, c = Density.natural_orbitals(D, S, C, orthogonalize_mo = True, n_eps = 0.001)
```

```
D: ndarray of shape (AO x AO)
S: ndarray of shape (AO x AO)
C: ndarray of shape (AO x MO)
```

```
--> transformation D (MO x MO) = C.T S D S C and its diagonalization
--> transformation of transformation matrix from MO to AO basis
```

```
n: ndarray of shape (NO)
c: ndarray of shape (AO x NO)
```

2) NO's in certain orthogonal MO basis from D in the same MO basis

```
n, c = Density.natural_orbitals(D, None, None, orthogonalize_mo = False, n_eps = 0.
```

```
D: ndarray of shape (MO x MO)
```

```
--> diagonalization of D
```

```
n: ndarray of shape (NO)
c: ndarray of shape (MO x NO)
```

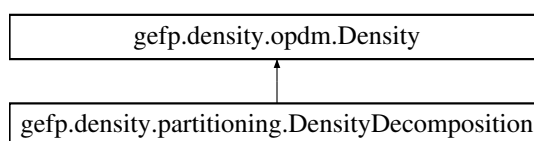
Last Revision: Gundelf

The documentation for this class was generated from the following file:

- gefp/gefp/density/opdm.py

18.23 gefp.density.partitioning.DensityDecomposition Class Reference

Inheritance diagram for gefp.density.partitioning.DensityDecomposition:



Public Member Functions

- `def __init__ (self, aggregate, method='hf', acbs=True, jk_type='direct', no_cutoff=0.000, xc_scale=1.0, l_dds=True, cc_relax=True, verbose=False, n_eps=5.0E-5, kwargs)`
- `def compute (self, polar_approx=True)`
- `def deformation_density (self, name)`
- `def compute_monomers (self)`
- `def compute_full_QM (self)`
- `def compute_densities (self)`
- `def compute_coulomb (self)`

- def **compute_pauli** (self)
- def **compute_polar** (self)
- def **compute_polar_approx** (self)
- def **__repr__** (self)
- def **print_out** (self)
- def **doublet** (self, A, B)
- def **triplet** (self, A, B, C)
- def **matrix_power** (self, M, x, eps=1.0e-6)
- def **rms** (self, m1, m2)

Public Attributes

- **aggregate**
- **method**
- **data**
- **matrix**
- **vars**
- **xc_recommended**
- **kwargs**
- **acbs**
- **no_cutoff**
- **n_eps**
- **cc_relax**
- **l_dds**
- **verbose**
- **xc_scale**
- **monomers_computed**
- **densities_computed**
- **energy_coulomb_computed**
- **energy_pauli_computed**
- **energy_polar_computed**
- **energy_polar_approx_computed**
- **energy_ind_computed**
- **energy_disp_computed**
- **energy_ct_compute**
- **energy_full_QM_computed**
- **dms_ind_computed**
- **dms_disp_computed**
- **dms_ct_computed**
- **bfs**
- **global_jk**
- **nmo_t**
- **nbf_t**

18.23.1 Detailed Description

Density-Based Decomposition Scheme of Mandado and Hermida-Ramon with partitioning of polarization deformation density into induction, dispersion and charge-transfer contributions.

```
--> DDS <--
--> Density Decomposition Scheme <--
```

References:

- * Mandado and Hermida-Ramon, J. Chem. Theory Comput. 2011, 7, 633-641. (JCTC 2011)
 - * B_lasiak, J. Chem. Phys. 2018 149 (16), 164115. (JCP 2018)
-

Constructor arguments and options:

- o aggregate - Psi4 molecular aggregate with at least two fragments
 - o method - QM method (hf, mp2, cc2, ccSD)
 - o acbs - use aggregate-centred basis set for calculations of wavefunctions. Otherwise use monomer-centred basis sets and composite Hadamard addition of AO spaces. ACBS=False result in no correction for BSSE.
 - o jk_type - type of Psi4 JK object.
 - o no_cutoff - cutoff for natural occupancies threshold. All natural orbitals with occupancies less or equal to the threshold will be neglected.
 - o xc_scale - scaling parameter for exchange-correlation density
 - o l_dds - compute also linear DDS total interaction energy
 - o kwargs - additional Psi4-relevant options.
-

Usage example:

```
solver = DensityDecomposition(aggr, method='hf',
                              acbs=True,
                              jk_type='direct',
                              no_cutoff=0.000,
                              xc_scale=1.0,
                              l_dds=True,
                              n_eps=5.0E-5,
                              cc_relax=True,
                              verbose=True,
                              **kwargs)

solver.compute(polar_approx=False)

dD_pauli = solver.deformation_density('pau')
dD_pol   = solver.deformation_density('pol')
dD       = solver.deformation_density('fqm')

# dictionaries:
# 1. accessing variables
solver.vars
# 2. accessing aggregate data
solver.matrix
# 3. accessing unperturbed fragment data (expert)
solver.data
```

```
print(solver)
solver.print_out()
```

Created : Gundelf

Last Revision: Gundelf

18.23.2 Member Function Documentation

18.23.2.1 compute()

```
def gefp.density.partitioning.DensityDecomposition.compute (
    self,
    polar_approx = True )
```

\

Perform the full density and interaction energy decompositions.

Options:

- o polar_approx - in addition to exact polarization energy, compute also the approximated polarization energy using NO-expansion of exchange-correlation 2-electron density and exact Pauli, polarization and unperturbed 1-electron densities.
- Default: True

Notes:

- o Exact polarization energy is always computed as a difference between the full QM interaction energy and all the remaining energies (Coulombic, exchange and repulsion energies).

18.23.2.2 deformation_density()

```
def gefp.density.partitioning.DensityDecomposition.deformation_density (
    self,
    name )
```

\

Compute the deformation 1-particle density matrix.
Returns density matrix in AO basis of entire molecular aggregate.

Possible <name> entries:

- o fqm - full QM deformation density
- o pau - Pauli-repulsion denformation density
- o pol - polarization deformation density
- o ind - induction part of polarization deformation density

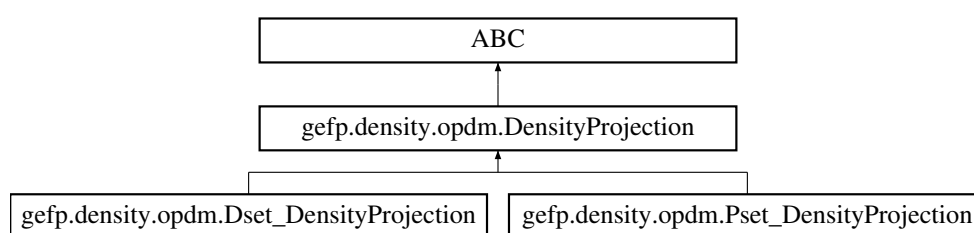
- o `dis` - dispersion part of polarization deformation density
- o `ct` - charge-transfer part of polarization deformation density

The documentation for this class was generated from the following file:

- `gefp/gefp/density/partitioning.py`

18.24 gefp.density.opdm.DensityProjection Class Reference

Inheritance diagram for `gefp.density.opdm.DensityProjection`:



Public Member Functions

- `def __init__ (self, np, S)`
- `def compute (self, n, c)`

Static Public Member Functions

- `def create (np, dtype='p', S=None)`

18.24.1 Detailed Description

\
 Gradient Projection Algorithms.
 Ref.: Pernal, Cancas, J. Chem. Phys. 2005

Usage:

```
proj = DensityProjection.create(np, dtype='p', S=None)
n, c = proj.compute(n, c, S)
```

The documentation for this class was generated from the following file:

- `gefp/gefp/density/opdm.py`

18.25 gefp.basis.optimize.DFBasis Class Reference

Public Member Functions

- def **__init__** (self, mol, templ_file='templ.dat', param_file='param.dat', bounds_file=None, constraints=())
- def **basisset** (self, param=None)
- def **print** (self, param=None)
- def **save** (self, out='oepfit.gbs', param=None)

Public Attributes

- **mol**
- **templ**
- **param**
- **n_param**
- **bounds**
- **constraints**
- **basis**

Static Public Attributes

- float **exp_lower_bound** = 0.01
- float **exp_upper_bound** = 10000.0
- float **ctr_lower_bound** = -2.0
- float **ctr_upper_bound** = 2.0

18.25.1 Detailed Description

Basis set object to be optimized.

Notes:

- o Default bounds can be modified by resetting static variables
DFBasis.exp_lower_bound
DFBasis.exp_upper_bound
DFBasis.ctr_lower_bound
DFBasis.ctr_upper_bound
prior to calling DFBasis if not using the driver.gdf_basis_optimizer.

The documentation for this class was generated from the following file:

- gefp/gefp/basis/optimize.py

18.26 gefp.basis.optimize.DFBasisOptimizer Class Reference

Public Member Functions

- `def __init__ (self, oep)`
- `def fit (self, maxiter=1000)`
- `def compute_error (self, basis, rms=False)`

Public Attributes

- `oep`
- `basis_fit`
- `param`

18.26.1 Detailed Description

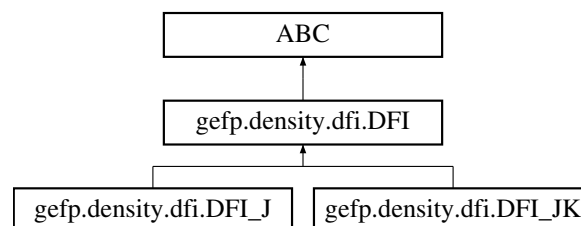
Method that optimizes DF basis set.

The documentation for this class was generated from the following file:

- `gefp/gefp/basis/optimize.py`

18.27 gefp.density.dfi.DFI Class Reference

Inheritance diagram for gefp.density.dfi.DFI:



Public Member Functions

- `def __init__ (self, frags)`
- `def run (self, maxit=100, conv=1.0e-5, verbose_scf=False, conv_scf=1.0e-5, maxit_scf=100, damp_scf=0.14, ndamp_scf=0)`
- `def aggregate (self)`
- `def wfn (self, i)`
- `def epsilon (self, i)`
- `def Cocc (self, i)`

- def **C** (self, i)
- def **D** (self, i)
- def **F** (self, i)
- def **V** (self, i)
- def **E** (self, i)

Static Public Member Functions

- def **create** (frags, j_only=False)

Public Attributes

- **enuc**
- **en_0**

18.27.1 Detailed Description

Density Fragment Interaction (DFI) Method

Demo for SCF-DFI method (closed shells).

Usage:

```
dfi = DFI(fragment_1, fragment_2, [...]) # OR: dfi = DFI(fragments)
dfi.run(maxit=30, conv=1.0e-5, verbose_scf=False, conv_scf=1.0e-5, maxit_scf=100, damp_scf=0.5)
```

Notes:

- o fragment_i is a psi4.core.Molecule with one Psi4 Fragment
 - o fragments is a psi4.core.Molecule with multiple Psi4 Fragments ('--' separator in input)
 - o SCF of unperturbed molecule is solved by Psi4, while the subsequent SCF's in DFI iteration are solved by SCF class instances of this Demo.
-

Last Revision: Gund

18.27.2 Member Function Documentation

18.27.2.1 run()

```
def gefp.density.dfi.DFI.run (
    self,
    maxit = 100,
    conv = 1.0e-5,
    verbose_scf = False,
    conv_scf = 1.0e-5,
```

```
maxit_scf = 100,  
damp_scf = 0.14,  
ndamp_scf = 0 )
```

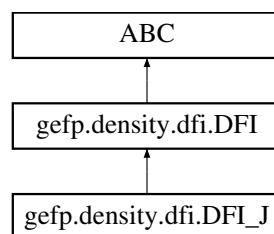
Runs DFI iterations

The documentation for this class was generated from the following file:

- gefp/gefp/density/dfi.py

18.28 gefp.density.dfi.DFI_J Class Reference

Inheritance diagram for gefp.density.dfi.DFI_J:



Public Member Functions

- `def __init__ (self, frags)`

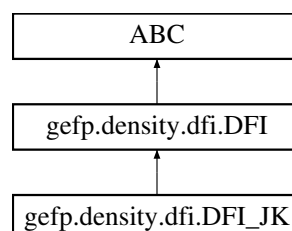
Additional Inherited Members

The documentation for this class was generated from the following file:

- gefp/gefp/density/dfi.py

18.29 gefp.density.dfi.DFI_JK Class Reference

Inheritance diagram for gefp.density.dfi.DFI_JK:



Public Member Functions

- `def __init__ (self, frags)`

Additional Inherited Members

The documentation for this class was generated from the following file:

- `gefp/gefp/density/dfi.py`

18.30 oepdev::DIISManager Class Reference

DIIS manager.

```
#include <diis.h>
```

Public Member Functions

- [DIISManager](#) (int dim, int na, int nb)
- [~DIISManager](#) ()
Destructor.
- void [put](#) (const std::shared_ptr< const Matrix > &error, const std::shared_ptr< const Matrix > &vector)
- void [compute](#) (void)
- void [update](#) (std::shared_ptr< Matrix > &other)

18.30.1 Detailed Description

Instance can interact directly with the process of solving vector quantities in iterative manner. One needs to pass the dimensions of solution vector as well as the DIIS subspace size. The iterative procedure requires providing the current vector and also an estimate of the error vector. The updated DIIS vector can be copied to an old vector through the Instance.

18.30.2 Constructor & Destructor Documentation

18.30.2.1 DIISManager()

```
oepdev::DIISManager::DIISManager (
    int dim,
    int na,
    int nb )
```

Constructor.

Parameters

<i>dim</i>	Size of DIIS subspace
<i>na</i>	Number of solution rows
<i>nb</i>	Number of solution columns

18.30.3 Member Function Documentation

18.30.3.1 compute()

```
void oepdev::DIISManager::compute (
    void )
```

Perform DIIS interpolation.

18.30.3.2 put()

```
void oepdev::DIISManager::put (
    const std::shared_ptr< const Matrix > & error,
    const std::shared_ptr< const Matrix > & vector )
```

Put the current solution to the DIIS manager.

Parameters

<i>error</i>	Shared matrix with current solution error
<i>vector</i>	Shared matrix with current solution vector

18.30.3.3 update()

```
void oepdev::DIISManager::update (
    std::shared_ptr< Matrix > & other )
```

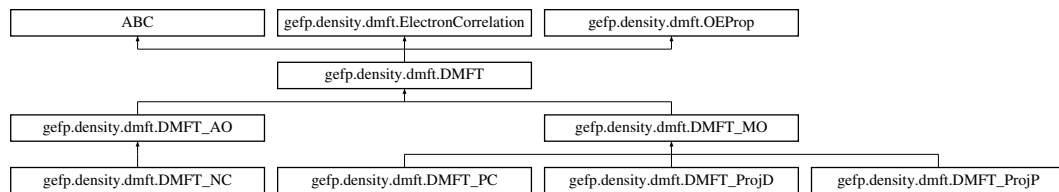
Update solution vector. Pass the Shared pointer to current solution. Then it will be overridden by the updated DIIS solution.

The documentation for this class was generated from the following files:

- oepdev/libutil/[diis.h](#)
- oepdev/libutil/diis.cc

18.31 gefp.density.dmft.DMFT Class Reference

Inheritance diagram for gefp.density.dmft.DMFT:



Public Member Functions

- `def __init__ (self, wfn, xc_functional, v_ext=default_v_ext, guess=default_guess, step_mode=default_step_mode)`
- `def create (cls, wfn, xc_functional=default_xc_functional, v_ext=default_v_ext, guess=default_guess, algorithm=default_algorithm, step_mode=default_step_mode, kwargs)`
- `def run (self, conv=default_convergence, maxiter=default_maxiter, verbose=default_verbose_run, g_0=default_g0, g=default_g, restart=False, kwargs)`
- `def set_gradient_mode (self, exact=False, approx=False, num=False)`
- `def E (self)`
- `def D (self)`
- `def Dmo (self)`
- `def Dao (self)`
- `def C (self)`
- `def N (self)`
- `def scalar_correlation (self)`
- `def matrix_correlation (self)`
- `def dipole (self)`
- `def quadrupole (self)`
- `def abbr (self)`

Static Public Member Functions

- `def name ()`

Static Public Attributes

- `default_xc_functional = XCFunctional.create('hf')`
- `string default_algorithm = 'proj-p'`
- `float default_convergence = 0.00001`
- `int default_maxiter = 100`
- `bool default_verbose_run = True`

- float **default_g0** = 0.0001
- **default_v_ext** = None
- string **default_guess** = 'hcore'
- string **default_step_mode** = 'search'
- float **default_g** = 0.01

18.31.1 Detailed Description

\

The Density Matrix Functional Theory. Abstract Base.

Usage:

```
# init
dmft = DMFT.create(wfn,
                   xc_functional = DMFT.default_xc_functional,
                   v_ext         = DMFT.default_v_ext         ,
                   guess         = DMFT.default_guess         ,
                   algorithm      = DMFT.default_algorithm     ,
                   step_mode      = DMFT.default_step_mode     ,
                   **kwargs)

# options
dmft.set_gradient_mode(exact=False, approx=False, num=False)

# run
dmft.run(conv       = DMFT.default_convergence,
         maxit      = DMFT.default_maxiter    ,
         verbose    = DMFT.default_verbose_run,
         g_0        = DMFT.default_g0         ,
         g          = DMFT.default_g          ,
         restart    = False                   ,
         **kwargs):
```

Options (init):

- o wfn - psi4.core.Wavefunction object. Must contain SCF LCAO-MO coefficients
- o xc_functional - XCFunctional object. Default: HF functional object.
- o v_ext - External potential in AO basis. Default is no potential.
- o guess - Guess for the density:
 - o 'hcore' - diagonalize Hcore Hamiltonian (default)
 - o 'current' - use the current density matrix stored in 'wfn'
- o algorithm - DMFT algorithm to converge the density matrix.
 - o 'proj-d' - Projected gradient algorithm on D-sets. Suitable only for
 - o 'proj-p' - Projected gradient algorithm on P-sets. Suitable for any
 - o 'nc' - Direct optimization within n and C parameter space. Suit
 - o 'pc' - Direct optimization within p and C parameter space. Not
- o step_mode - How to search for next guess: estimate step length in steepest desce
 - o 'search' - Compute from two last density guesses (default)
 - o 'constant' - Apply constant step.

Options (optional setup): relevant only for 'proj-p' algorithm

- o exact - Compute exact derivatives of XC energy wrt P

- o approx - Compute approximate derivatives of XC energy wrt P
 - o num - Compute numerically derivatives of XC energy wrt P
- If using this, set only one of the above three to True.

Options (run):

- o conv - Energy convergence (default 0.00001)
 - o maxit - Maximum number of iterations (default 100)
 - o verbose - Print detailed information or not (default True)
 - o g0 - Initial SD step size (default 0.0001)
 - o g - Constant SD step size (default 0.01)
 - o restart - Wheather to restart the calculations or not (default False)
-

18.31.2 Member Function Documentation

18.31.2.1 create()

```
def gefp.density.dmft.DMFT.create (
    cls,
    wfn,
    xc_functional = default_xc_functional,
    v_ext = default_v_ext,
    guess = default_guess,
    algorithm = default_algorithm,
    step_mode = default_step_mode,
    kwargs )

\
Create DMFT solver.
```

18.31.2.2 run()

```
def gefp.density.dmft.DMFT.run (
    self,
    conv = default_convergence,
    maxit = default_maxiter,
    verbose = default_verbose_run,
    g_0 = default_g0,
    g = default_g,
    restart = False,
    kwargs )

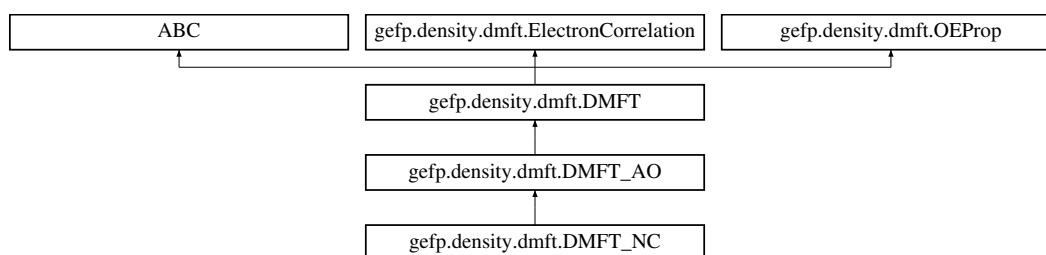
\
Run the DMFT calculations.
```

The documentation for this class was generated from the following file:

- gefp/gefp/density/dmft.py

18.32 gefp.density.dmft.DMFT_AO Class Reference

Inheritance diagram for gefp.density.dmft.DMFT_AO:



Public Member Functions

- def **__init__** (self, wfn, xc_functional, v_ext, guess, step)
- def **dipole** (self)
- def **Dmo** (self)

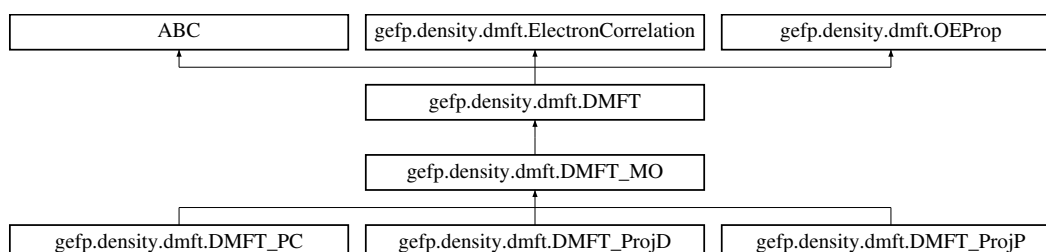
Additional Inherited Members

The documentation for this class was generated from the following file:

- gefp/gefp/density/dmft.py

18.33 gefp.density.dmft.DMFT_MO Class Reference

Inheritance diagram for gefp.density.dmft.DMFT_MO:



Public Member Functions

- def **__init__** (self, wfn, xc_functional, v_ext, guess, step)

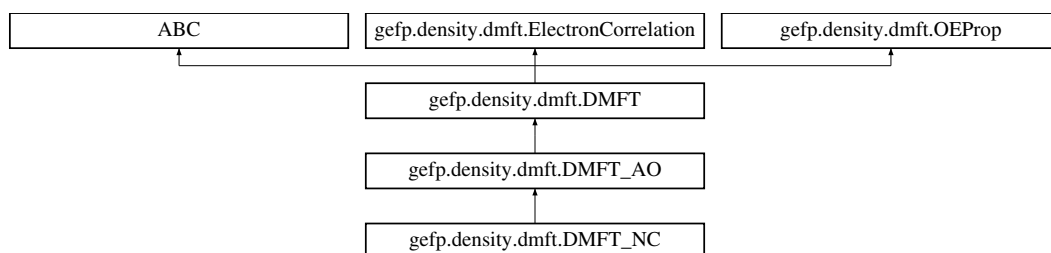
Additional Inherited Members

The documentation for this class was generated from the following file:

- `gefp/gefp/density/dmft.py`

18.34 `gefp.density.dmft.DMFT_NC` Class Reference

Inheritance diagram for `gefp.density.dmft.DMFT_NC`:



Public Member Functions

- `def __init__ (self, wfn, xc_functional, v_ext, guess, step)`
- `def abbr (self)`

Static Public Member Functions

- `def name ()`

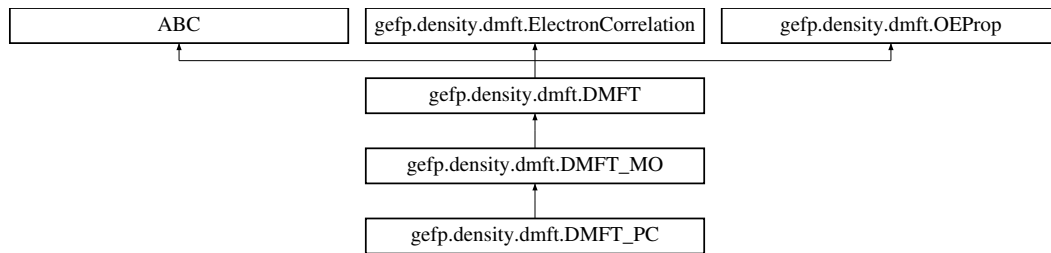
Additional Inherited Members

The documentation for this class was generated from the following file:

- `gefp/gefp/density/dmft.py`

18.35 `gefp.density.dmft.DMFT_PC` Class Reference

Inheritance diagram for `gefp.density.dmft.DMFT_PC`:



Public Member Functions

- `def __init__ (self, wfn, xc_functional, v_ext, guess, step)`
- `def abbr (self)`

Static Public Member Functions

- `def name ()`

Public Attributes

- **g**

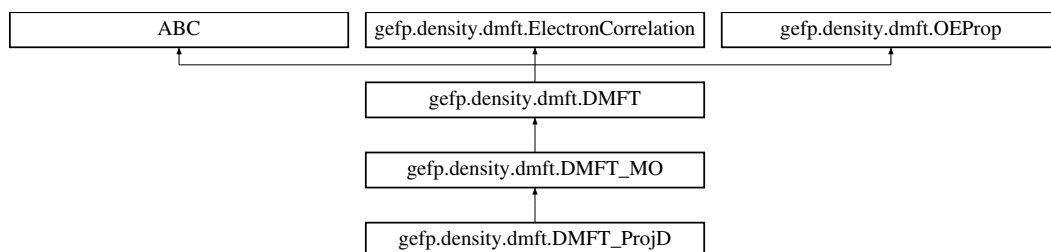
Additional Inherited Members

The documentation for this class was generated from the following file:

- `gefp/gefp/density/dmft.py`

18.36 gefp.density.dmft.DMFT_ProjD Class Reference

Inheritance diagram for gefp.density.dmft.DMFT_ProjD:



Public Member Functions

- `def __init__ (self, wfn, xc_functional, v_ext, guess, step)`
- `def abbr (self)`

Static Public Member Functions

- def **name** ()

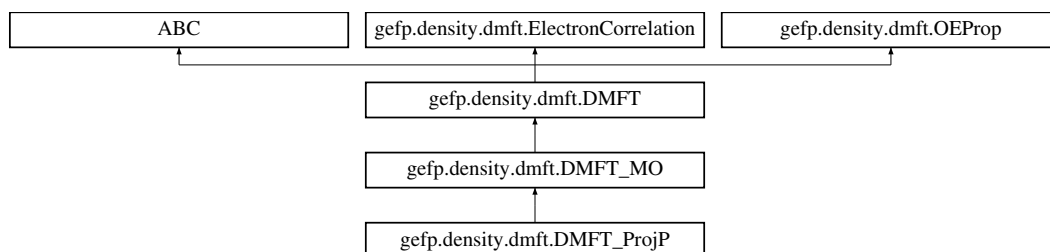
Additional Inherited Members

The documentation for this class was generated from the following file:

- gefp/gefp/density/dmft.py

18.37 gefp.density.dmft.DMFT_ProjP Class Reference

Inheritance diagram for gefp.density.dmft.DMFT_ProjP:



Public Member Functions

- def **__init__** (self, wfn, xc_functional, v_ext, guess, step)
- def **abbr** (self)

Static Public Member Functions

- def **name** ()

Additional Inherited Members

The documentation for this class was generated from the following file:

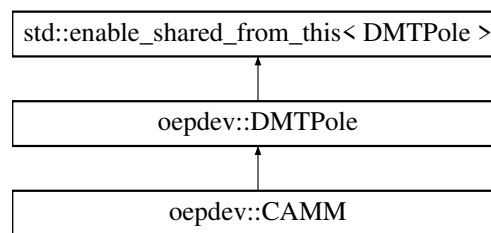
- gefp/gefp/density/dmft.py

18.38 oepdev::DMTPole Class Reference

Distributed Multipole Analysis Container and Computer. Abstract Base.

```
#include <dmt.h>
```

Inheritance diagram for oepdev::DMTPole:



Public Member Functions

- virtual `~DMTPole ()`
Destructor.
- virtual bool `has_charges () const`
Has distributed charges?
- virtual bool `has_dipoles () const`
Has distributed dipoles?
- virtual bool `has_quadrupoles () const`
Has distributed quadrupoles?
- virtual bool `has_octupoles () const`
Has distributed octupoles?
- virtual bool `has_hexadecapoles () const`
Has distributed hexadecapoles?
- virtual `psi::SharedMatrix centres () const`
Get the positions of distribution centres.
- virtual `psi::SharedMatrix origins () const`
Get the positions of distribution origins.
- virtual `psi::SharedVector centre (int x) const`
*Get the position of the *x*th distribution centre.*
- virtual `psi::SharedVector origin (int x) const`
*Get the position of the *x*th distribution origin.*
- virtual `std::vector< psi::SharedMatrix > charges () const`
Get the distributed charges.
- virtual `std::vector< psi::SharedMatrix > dipoles () const`
Get the distributed dipoles.
- virtual `std::vector< psi::SharedMatrix > quadrupoles () const`
Get the distributed quadrupoles.
- virtual `std::vector< psi::SharedMatrix > octupoles () const`
Get the distributed octupoles.
- virtual `std::vector< psi::SharedMatrix > hexadecapoles () const`
Get the distributed hexadecapoles.

- virtual `psi::SharedMatrix` `charges` (int i) const
Get the distributed charges for the ith distribution.
- virtual `psi::SharedMatrix` `dipoles` (int i) const
Get the distributed dipoles for the ith distribution.
- virtual `psi::SharedMatrix` `quadrupoles` (int i) const
Get the distributed quadrupoles for the ith distribution.
- virtual `psi::SharedMatrix` `octupoles` (int i) const
Get the distributed octupoles for the ith distribution.
- virtual `psi::SharedMatrix` `hexadecapoles` (int i) const
Get the distributed hexadecapoles for the ith distribution.
- virtual int `n_sites` () const
Get the number of distributed sites.
- virtual int `n_dmtp` () const
Get the number of distributions.
- void `set_charges` (std::vector< `psi::SharedMatrix` > M)
Set the distributed charges.
- void `set_dipoles` (std::vector< `psi::SharedMatrix` > M)
Set the distributed dipoles.
- void `set_quadrupoles` (std::vector< `psi::SharedMatrix` > M)
Set the distributed quadrupoles.
- void `set_octupoles` (std::vector< `psi::SharedMatrix` > M)
Set the distributed octupoles.
- void `set_hexadecapoles` (std::vector< `psi::SharedMatrix` > M)
Set the distributed hexadecapoles.
- void `set_charges` (`psi::SharedMatrix` M, int i)
Set the distributed charges for the ith distribution.
- void `set_dipoles` (`psi::SharedMatrix` M, int i)
Set the distributed dipoles for the ith distribution.
- void `set_quadrupoles` (`psi::SharedMatrix` M, int i)
Set the distributed quadrupoles for the ith distribution.
- void `set_octupoles` (`psi::SharedMatrix` M, int i)
Set the distributed octupoles for the ith distribution.
- void `set_hexadecapoles` (`psi::SharedMatrix` M, int i)
Set the distributed hexadecapoles for the ith distribution.
- virtual void `recenter` (`psi::SharedMatrix` new_origins)
- void `translate` (`psi::SharedVector` transl)
Translate the DMTP sets.
- void `rotate` (`psi::SharedMatrix` rotmat)
Rotate the DMTP sets.
- void `superimpose` (`psi::SharedMatrix` ref_xyz, std::vector< int > suplist)

Superimpose the DMTP sets.

- void [compute](#) (std::vector< psi::SharedMatrix > D, std::vector< bool > t)
- void [compute](#) (void)
- std::shared_ptr< [MultipoleConvergence](#) > [energy](#) (std::shared_ptr< [DMTPole](#) > other, [MultipoleConvergence::ConvergenceLevel](#) max_clevel=[MultipoleConvergence::R5](#))

Evaluate the generalized interaction energy.

- std::shared_ptr< [MultipoleConvergence](#) > [potential](#) (std::shared_ptr< [DMTPole](#) > other, [MultipoleConvergence::ConvergenceLevel](#) max_clevel=[MultipoleConvergence::R5](#))

Evaluate the generalized potential.

- virtual void [print_header](#) () const =0

Print the header.

- void [print](#) () const

Print the contents.

Static Public Member Functions

- static std::shared_ptr< [DMTPole](#) > [build](#) (const std::string &type, std::shared_ptr< psi::Wavefunction > wfn, int n=1)

Build an empty DMTP object from the wavefunction.

- static [MultipoleConvergence::ConvergenceLevel](#) [determine_dmtp_convergence_level](#) (const std::string &option)

Protected Member Functions

- [DMTPole](#) (std::shared_ptr< psi::Wavefunction > wfn, int n)

Construct an empty DMTP object from the wavefunction.

- virtual void [compute](#) (psi::SharedMatrix D, bool transition, int i)=0

Compute DMTP's from the one-particle density matrix.

- void [compute_integrals](#) ()

Compute multipole integrals.

- void [compute_order](#) ()

Compute maximum order of the integrals.

- virtual void [recenter](#) (psi::SharedMatrix new_origins, int i)

Change origins of the distributed multipole moments of ith set.

- virtual void [allocate](#) ()

Initialize and allocate memory.

Protected Attributes

- `std::string` [name_](#)
Name of the distribution method.
- `psi::SharedMolecule` [mol_](#)
Molecule associated with this DMTP.
- `psi::SharedWavefunction` [wfn_](#)
Wavefunction associated with this DMTP.
- `psi::SharedBasisSet` [primary_](#)
Basis set (primary)
- `int` [nDMTPs_](#)
Number of DMTP's.
- `int` [nSites_](#)
Number of DMTP sites.
- `int` [order_](#)
Maximum order of the multipole.
- `std::vector< psi::SharedMatrix >` [mplnts_](#)
Multipole integrals.
- `bool` [hasCharges_](#)
Has distributed charges?
- `bool` [hasDipoles_](#)
Has distributed dipoles?
- `bool` [hasQuadrupoles_](#)
Has distributed quadrupoles?
- `bool` [hasOctupoles_](#)
Has distributed octupoles?
- `bool` [hasHexadecapoles_](#)
Has distributed hexadecapoles?
- `psi::SharedMatrix` [centres_](#)
DMTP centres.
- `psi::SharedMatrix` [origins_](#)
DMTP origins.
- `std::vector< psi::SharedMatrix >` [charges_](#)
DMTP charges.
- `std::vector< psi::SharedMatrix >` [dipoles_](#)
DMTP dipoles.
- `std::vector< psi::SharedMatrix >` [quadrupoles_](#)
DMTP quadrupoles.
- `std::vector< psi::SharedMatrix >` [octupoles_](#)
DMTP octupoles.
- `std::vector< psi::SharedMatrix >` [hexadecapoles_](#)
DMTP hexadecapoles.

Friends

- class [MultipoleConvergence](#)

18.38.1 Detailed Description

Handles the distributed multipole expansions up to hexadecapoles. Distributed centres as well as DMTP origins are allowed to be located in arbitrary points in space. The object describes a set of N DMTP's, that can be generated by providing one-particle density matrices in AO basis. Nuclear contributions can be switched on or off separately for each DMTP within a set. The following operations on the DMTP sets are available through the API:

- translation
- rotation
- superimposition
- recentering the origins
- computing the generalized property from another DMTP set

See also

[MultipoleConvergence](#)

18.38.2 Constructor & Destructor Documentation

18.38.2.1 DMTPole()

```
oepdev::DMTPole::DMTPole (
    std::shared_ptr< psi::Wavefunction > wfn,
    int n ) [protected]
```

Parameters

<i>wfn</i>	- wavefunction
<i>n</i>	- number of DMTP sets

Do not use this constructor. Use the [DMTPole::build](#) method.

18.38.3 Member Function Documentation

18.38.3.1 build()

```
std::shared_ptr< DMTPole > oepdev::DMTPole::build (
    const std::string & type,
    std::shared_ptr< psi::Wavefunction > wfn,
    int n = 1 ) [static]
```

Parameters

<i>type</i>	- DMTP method. Available: CAMM .
<i>wfn</i>	- wavefunction
<i>n</i>	- number of DMTP sets

Returns

DMTP distribution

18.38.3.2 compute() [1/2]

```
void oepdev::DMTPole::compute (
    std::vector< psi::SharedMatrix > D,
    std::vector< bool > t )
```

Compute DMTP's from the set of the one-particle density matrices.

Parameters

<i>D</i>	- list of one-particle density matrices
<i>t</i>	- list of flags determining if density is of transition type or not

18.38.3.3 compute() [2/2]

```
void oepdev::DMTPole::compute (
    void )
```

Compute DMTP's from the *sum* of the ground-state alpha and beta one-particle density matrices (t=false, i=0).

Results in a usual DMTP analysis of a molecule's charge density distribution.

18.38.3.4 determine_dmt_p_convergence_level()

```
MultipoleConvergence::ConvergenceLevel oepdev::DMTPole::determine_dmt_p_convergence_level
```

```
(
    const std::string & option ) [static]
```

Determine the CAMM convergence for a given global option

Parameters

<i>option</i>	- string for option
---------------	---------------------

18.38.3.5 energy()

```
std::shared_ptr< MultipoleConvergence > oepdev::DMTPole::energy (
    std::shared_ptr< DMTPole > other,
    MultipoleConvergence::ConvergenceLevel max_clevel = MultipoleConvergence::R5
)
```

Parameters

<i>other</i>	- interacting DMTP distribution.
<i>max_clevel</i>	- maximum convergence level (see below).

Returns

The generalized interaction energy convergence (A.U. units)

The following convergence levels are available:

- `MultipoleConvergence::R1`: includes qq terms.
- `MultipoleConvergence::R2`: includes dq terms and above.
- `MultipoleConvergence::R3`: includes qQ, dd terms and above.
- `MultipoleConvergence::R4`: includes qO, dQ terms and above.
- `MultipoleConvergence::R5`: includes qH, dO, QQ terms and above.

18.38.3.6 potential()

```
std::shared_ptr< MultipoleConvergence > oepdev::DMTPole::potential (
    std::shared_ptr< DMTPole > other,
    MultipoleConvergence::ConvergenceLevel max_clevel = MultipoleConvergence::R5
)
```

Parameters

<i>other</i>	- interacting DMTP distribution.
<i>max_clevel</i>	- maximum convergence level (see below).

Returns

The generalized potential convergence (A.U. units)

The following convergence levels are available:

- `MultipoleConvergence::R1`: includes qq terms.
- `MultipoleConvergence::R2`: includes dq terms and above.
- `MultipoleConvergence::R3`: includes qQ, dd terms and above.
- `MultipoleConvergence::R4`: includes qO, dQ terms and above.
- `MultipoleConvergence::R5`: includes qH, dO, QQ terms and above.

18.38.3.7 recenter()

```
void oepdev::DMTPole::recenter (
    psi::SharedMatrix new_origins ) [virtual]
```

Change origins of the distributed multipole moments of all sets

Parameters

<i>new_origins</i>	- matrix with coordinates of the new origins $\{\mathbf{r}_{\text{new}}\}$.
--------------------	--

Note

The number of origins has to be equal to the number of distributed centres.

Recentering of the multipoles affects the distributed dipoles and higher moments. The moments are given as

$$\begin{aligned}
 q_{\text{new}} &= q_{\text{old}} \\
 \mu_{\text{new}} &= \mu_{\text{old}} - q_{\text{old}} \Delta^{(1)} \\
 \Theta_{\text{new}} &= \Theta_{\text{old}} + q_{\text{old}} \Delta^{(2)} - \sum_{\mathcal{P}_2} \mathcal{P}_2 \left[(q_{\text{old}} \mathbf{r}_{\text{old}} + \mu_{\text{old}}) \otimes \Delta^{(1)} \right] \\
 \Omega_{\text{new}} &= \Omega_{\text{old}} - q_{\text{old}} \Delta^{(3)} + \sum_{\mathcal{P}_3} \mathcal{P}_3 \left[(q_{\text{old}} \mathbf{r}_{\text{old}} + \mu_{\text{old}}) \otimes \Delta^{(2)} \right] - \sum_{\mathcal{P}_6} \mathcal{P}_6 \left[(q_{\text{old}} \mathbf{r}_{\text{old}}^2 + \mu_{\text{old}} \otimes \mathbf{r}_{\text{old}} + \Theta_{\text{old}}) \otimes \Delta^{(1)} \right] \\
 \Xi_{\text{new}} &= \Xi_{\text{old}} + q_{\text{old}} \Delta^{(4)} - \sum_{\mathcal{P}_3} \mathcal{P}_3 \left[(q_{\text{old}} \mathbf{r}_{\text{old}} + \mu_{\text{old}}) \otimes \Delta^{(3)} \right] + \sum_{\mathcal{P}_3} \mathcal{P}_3 \left[(q_{\text{old}} \mathbf{r}_{\text{old}}^2 + \mu_{\text{old}} \otimes \mathbf{r}_{\text{old}} + \Theta_{\text{old}}) \otimes \Delta^{(2)} \right] - \sum_{\mathcal{P}_6} \mathcal{P}_6 \left[(q_{\text{old}} \mathbf{r}_{\text{old}}^2 + \mu_{\text{old}} \otimes \mathbf{r}_{\text{old}} + \Theta_{\text{old}}) \otimes \Delta^{(1)} \right]
 \end{aligned}$$

where

$$\begin{aligned}\Delta^{(1)} &\equiv \mathbf{r}_{\text{new}} - \mathbf{r}_{\text{old}} \\ \Delta^{(2)} &\equiv \mathbf{r}_{\text{new}}^2 - \mathbf{r}_{\text{old}}^2 \\ \Delta^{(3)} &\equiv \mathbf{r}_{\text{new}}^3 - \mathbf{r}_{\text{old}}^3 \\ \Delta^{(4)} &\equiv \mathbf{r}_{\text{new}}^4 - \mathbf{r}_{\text{old}}^4\end{aligned}$$

In the above equations, the distributed centre label was omitted (redundant) as each distributed site of multipoles is independent of the others. TODO - Finish for octupoles and hexadecapoles!
-> define the permutation operators!

18.38.4 Friends And Related Function Documentation

18.38.4.1 MultipoleConvergence

```
friend class MultipoleConvergence [friend]
```

Convergence of multipole moment series.

The documentation for this class was generated from the following files:

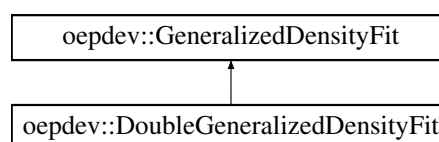
- oepdev/lib3d/dmtp.h
- oepdev/lib3d/dmtp_base.cc

18.39 oepdev::DoubleGeneralizedDensityFit Class Reference

Generalized Density Fitting Scheme - Double Fit.

```
#include <oep_gdf.h>
```

Inheritance diagram for oepdev::DoubleGeneralizedDensityFit:



Public Member Functions

- **DoubleGeneralizedDensityFit** (std::shared_ptr< psi::BasisSet > bs_auxiliary, std::shared_ptr< psi::BasisSet > bs_intermediate, std::shared_ptr< psi::Matrix > v_vector)
- std::shared_ptr< psi::Matrix > **compute** (void)

Perform the generalized density fit.

Additional Inherited Members

18.39.1 Detailed Description

The density fitting map projects the OEP onto an arbitrary (not necessarily complete) auxiliary basis set space through application of the self energy minimization technique. The resulting three-electron repulsion integrals are computed by utilizing the resolution of identity in an intermediate, nearly-complete basis set space, hence performing an internal density fitting in nearly complete basis. Refer to [density fitting specialized for OEP's](#) for more details.

18.39.2 Determination of the OEP matrix

Coefficients \mathbf{G} are computed by using the following relation

$$\mathbf{G} = \mathbf{A}^{-1} \cdot \mathbf{R} \cdot \mathbf{H}$$

where the intermediate projection matrix is given by

$$\mathbf{H} = \mathbf{S}^{-1} \cdot \mathbf{V}$$

In the above equations,

$$A_{\xi\xi'} = (\xi||\xi')$$

$$R_{\xi\varepsilon} = (\xi||\varepsilon)$$

$$S_{\varepsilon\varepsilon'} = (\varepsilon|\varepsilon')$$

$$V^{\varepsilon i} = (\varepsilon|\hat{v}i)$$

The following labeling convention is used here:

- i denotes the arbitrary state vector
- ξ denotes the auxiliary basis set element
- ε denotes the intermediate (nearly complete) basis set element

In the above, $|$ denotes the single integration over electron coordinate, i.e.,

$$(a|b) \equiv \int d\mathbf{r} \phi_a^*(\mathbf{r}) \phi_b(\mathbf{r})$$

whereas $||$ acts as shown below:

$$(a||b) \equiv \iint d\mathbf{r}' d\mathbf{r}'' \frac{\phi_a^*(\mathbf{r}') \phi_b(\mathbf{r}'')}{|\mathbf{r}' - \mathbf{r}''|}$$

The spatial form of the potential operator \hat{v} can be expressed by

$$v(\mathbf{r}) \equiv \int d\mathbf{r}' \frac{\rho(\mathbf{r}')}{|\mathbf{r}' - \mathbf{r}|}$$

with $\rho(\mathbf{r})$ being the effective one-electron density associated with \hat{v} .

18.39.2.1 Theory behind the double GDF scheme

In order to perform the generalized density fitting in an incomplete auxiliary basis set, one must apply the following formula:

$$\mathbf{G} = \mathbf{A}^{-1} \cdot \mathbf{B}$$

where one encounters the need of evaluation of the following *three-electron integrals*

$$B_{\xi i} = (\xi || \hat{v} i) \equiv \iiint d\mathbf{r}' d\mathbf{r}'' d\mathbf{r}''' \phi_{\xi}^*(\mathbf{r}') \frac{1}{|\mathbf{r}' - \mathbf{r}''|} \rho(\mathbf{r}''') \frac{1}{|\mathbf{r}''' - \mathbf{r}''|} \phi_i(\mathbf{r}'')$$

Computation of all the necessary integrals of this kind is very costly and impractical for larger molecules. However, one can use the same trick that is a kernel of the OEP technique introduced in the OEPDev project, i.e., introduce the effective potential in order to get rid of one integration. This can be done by performing the generalized density fitting in the nearly complete intermediate basis

$$\hat{v} |i) \cong \sum_{\varepsilon} H_{\varepsilon i} |\varepsilon)$$

Note that this is done just for the sake of factorizing the triple integral and computing the OEP matrix for the incomplete auxiliary basis. Therefore, the intermediate basis set is used just for a while during density fitting and is no longer necessary later on. By inserting the above identity to the triple integral one can transform it into a sum of the two-electron integrals that are much easier to evaluate. This leads to equations given in the beginning of this section.

18.39.3 Member Function Documentation

18.39.3.1 compute()

```
std::shared_ptr< psi::Matrix > DoubleGeneralizedDensityFit::compute (
    void ) [virtual]
```

Returns

The OEP coefficients $G_{\xi i}$

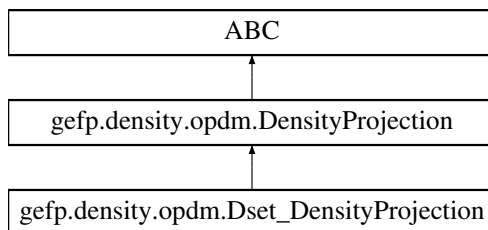
Implements [oepdev::GeneralizedDensityFit](#).

The documentation for this class was generated from the following files:

- [oepdev/liboep/oep_gdf.h](#)
- [oepdev/liboep/oep_gdf.cc](#)

18.40 gefp.density.opdm.Dset_DensityProjection Class Reference

Inheritance diagram for gefp.density.opdm.Dset_DensityProjection:



Public Member Functions

- `def __init__(self, np, S)`

Additional Inherited Members

18.40.1 Detailed Description

\

Gradient Projection Algorithm on D-sets.
 Ref.: Pernal, Cancès, J. Chem. Phys. 2005

Notes:

- Appropriate only for HF functional.

The documentation for this class was generated from the following file:

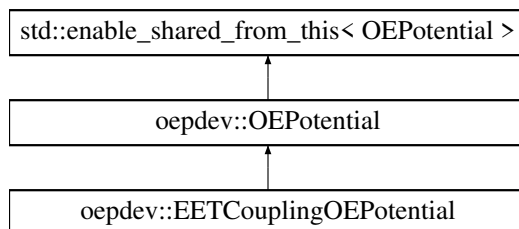
- `gefp/gefp/density/opdm.py`

18.41 oepdev::EETCouplingOEPotential Class Reference

Generalized One-Electron Potential for EET coupling calculations.

```
#include <oep.h>
```

Inheritance diagram for `oepdev::EETCouplingOEPotential`:



Public Member Functions

- **EETCouplingOEPotential** (SharedWavefunction [wfn](#), SharedBasisSet auxiliary, Shared-BasisSet intermediate, Options &options)

- **EETCouplingOEPotential** (SharedWavefunction [wfn](#), Options &options)
- virtual void [compute](#) (const std::string &oepType) override
Compute matrix forms of all OEP's within a specified OEP type.
- virtual void [compute_3D](#) (const std::string &oepType, const double &x, const double &y, const double &z, std::shared_ptr< psi::Vector > &v) override
Compute value of potential in point x, y, z and save at v.
- virtual void [print_header](#) () const override
Header information.

Additional Inherited Members

18.41.1 Detailed Description

Contains the following OEP types:

- `Fujimoto.GDF` - Joint OEP type for ET(L), ET(HL), HT(H) and HT(HL)
- `Fujimoto.CIS` - CIS data
- `Fujimoto.EXCH` - Pure-exchange coupling matrix $G_{\mu\nu} \equiv (\mu\mu|v\nu)$
- `Fujimoto.CTM` - ($HH|LL$) integral for the H_34 Hamiltonian matrix elements (CT)

The documentation for this class was generated from the following files:

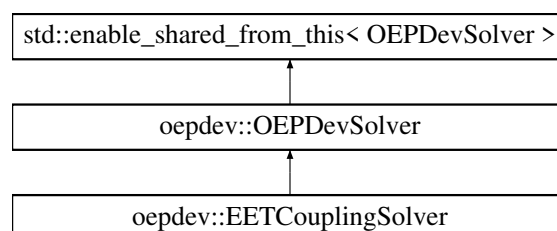
- `oepdev/liboep/oep.h`
- `oepdev/liboep/oep_coupling_eet.cc`

18.42 oepdev::EETCouplingSolver Class Reference

Compute the EET coupling energy between unperturbed wavefunctions.

```
#include <solver.h>
```

Inheritance diagram for oepdev::EETCouplingSolver:



Public Member Functions

- **EETCouplingSolver** (SharedWavefunctionUnion wfn_union)
- virtual double `compute_oep_based` (const std::string &method="DEFAULT")
Compute property by using OEP's.
- virtual double `compute_benchmark` (const std::string &method="DEFAULT")
Compute property by using benchmark method.

Additional Inherited Members

18.42.1 Detailed Description

The implemented methods are shown below

Table 18.29: Methods available in the Solver

Keyword	Method Description
Benchmark Methods	
FUJIMOTO_TI_CIS	<i>Default.</i> EET Coupling by Fujimoto JPC 2012.
OEP-Based Methods	
FUJIMOTO_TI_CIS	<i>Default.</i> OEP-based TI/CIS expressions.

In order to construct this solver, **always** use the `OEPDevSolver::build` static factory method.

Below the detailed description of the implemented equations is given for each of the above provided methods. In the formulae across, it is assumed that the orbitals are real. The Coulomb notation for electron repulsion integrals (ERI's) is adopted; i.e.,

$$(ac|bd) = \iint d\mathbf{r}_1 d\mathbf{r}_2 \phi_a(\mathbf{r}_1) \phi_c(\mathbf{r}_1) \frac{1}{r_{12}} \phi_b(\mathbf{r}_2) \phi_d(\mathbf{r}_2)$$

Greek subscripts denote basis set orbitals whereas *Italic* subscripts denote the occupied molecular orbitals.

Benchmark Methods

TI/CIS Method (Fujimoto JPC 2012).

In the simplest version of TI/CIS approach, the Hamiltonian of the molecular aggregate (dimer) is constructed from the CIS approximation and 4 basis functions constructed as follows:

$$\begin{aligned} |\Phi_1\rangle &= |\Psi_A^{(e)} \otimes \Psi_B^{(g)}\rangle \\ |\Phi_2\rangle &= |\Psi_A^{(g)} \otimes \Psi_B^{(e)}\rangle \\ |\Phi_3\rangle &= |\Psi_A^{(+)} \otimes \Psi_B^{(-)}\rangle \\ |\Phi_4\rangle &= |\Psi_A^{(-)} \otimes \Psi_B^{(+)}\rangle \end{aligned}$$

where g and e superscripts denote the ground and excited state of a molecule, $+$ and $-$ label the cationic and anionic state, respectively, whereas $|\Psi_X \otimes \Psi_Y\rangle$ denotes the antisymmetrized Hartree product of the monomer wavefunctions. The associated diagonal Hamiltonian matrix elements can be defined as

$$\begin{aligned} \langle \Phi_1 | \mathcal{H} - E_0 | \Phi_1 \rangle &\equiv E_1 = E_{e \rightarrow g}^A + \sum_{\mu\nu \in A} (P_{\nu\mu}^{A(e)} - P_{\nu\mu}^{A(g)}) \times \left\{ V_{\mu\nu}^{B(\text{nuc})} + \sum_{\lambda\sigma \in B} P_{\lambda\sigma}^{B(g)} \left[(\mu\nu|\sigma\lambda) - \frac{1}{2}(\mu\lambda|\sigma\nu) \right] \right\} \\ \langle \Phi_2 | \mathcal{H} - E_0 | \Phi_2 \rangle &\equiv E_2 = E_{e \rightarrow g}^B + \sum_{\mu\nu \in B} (P_{\nu\mu}^{B(e)} - P_{\nu\mu}^{B(g)}) \times \left\{ V_{\mu\nu}^{A(\text{nuc})} + \sum_{\lambda\sigma \in A} P_{\lambda\sigma}^{A(g)} \left[(\mu\nu|\sigma\lambda) - \frac{1}{2}(\mu\lambda|\sigma\nu) \right] \right\} \\ \langle \Phi_3 | \mathcal{H} - E_0 | \Phi_3 \rangle &\equiv E_3 = -\varepsilon_H^A + \varepsilon_L^B - (H^A H^A | L^B L^B) \\ \langle \Phi_4 | \mathcal{H} - E_0 | \Phi_4 \rangle &\equiv E_4 = \varepsilon_L^A - \varepsilon_H^B - (L^A L^A | H^B H^B) \end{aligned}$$

The associated off-diagonal Hamiltonian matrix elements can be defined as

$$\begin{aligned} \langle \Phi_1 | \mathcal{H} | \Phi_2 \rangle &\equiv V^{\text{Coul}} + V^{\text{Exch}} + V^{\text{Ovrl}} \\ \langle \Phi_1 | \mathcal{H} | \Phi_3 \rangle &\equiv V^{\text{ET1}} \\ \langle \Phi_2 | \mathcal{H} | \Phi_4 \rangle &\equiv V^{\text{ET2}} \\ \langle \Phi_1 | \mathcal{H} | \Phi_4 \rangle &\equiv V^{\text{HT1}} \\ \langle \Phi_2 | \mathcal{H} | \Phi_3 \rangle &\equiv V^{\text{HT2}} \\ \langle \Phi_3 | \mathcal{H} | \Phi_4 \rangle &\equiv V^{\text{CT}} \end{aligned}$$

where the Forster-type Coulombic (Coul), Dexter-type exchange (Exch), remaining overlap correction (Ovrl), as well as the electron, hole and charge (ET, HT, CT) transfer contributions are defined. The exchange-Coulomb coupling takes the form

$$\begin{aligned} V^{\text{Coul}} &= \frac{V^{\text{Coul},(0)}}{1 - S_{12}^2} \\ V^{\text{Exch}} &= \frac{V^{\text{Exch},(0)}}{1 - S_{12}^2} \\ V^{\text{Ovrl}} &= -\frac{(E_1 + E_2)S_{12}}{2(1 - S_{12}^2)} \end{aligned}$$

The overlap-corrected ET, HT and CT matrix elements read

$$\begin{aligned}
 V^{\text{ET1}} &= [1 - S_{13}^2]^{-1} \left\{ V^{\text{ET1},(0)} - \frac{1}{2}(E_1 + E_2)S_{13} \right\} \\
 V^{\text{ET2}} &= [1 - S_{24}^2]^{-1} \left\{ V^{\text{ET2},(0)} - \frac{1}{2}(E_1 + E_2)S_{24} \right\} \\
 V^{\text{HT1}} &= [1 - S_{14}^2]^{-1} \left\{ V^{\text{HT1},(0)} - \frac{1}{2}(E_1 + E_2)S_{14} \right\} \\
 V^{\text{HT2}} &= [1 - S_{23}^2]^{-1} \left\{ V^{\text{HT2},(0)} - \frac{1}{2}(E_1 + E_2)S_{23} \right\} \\
 V^{\text{CT}} &= [1 - S_{34}^2]^{-1} \left\{ V^{\text{CT},(0)} - \frac{1}{2}(E_1 + E_2)S_{34} \right\}
 \end{aligned}$$

In the above equations, the superscript (0) denotes that the matrix elements are not affected by the overlap between molecular wavefunctions, and are given by

$$\begin{aligned}
 V^{\text{Coul},(0)} &= \sum_{\mu\nu \in A} \sum_{\lambda\sigma \in B} P_{\nu\mu}^{g \rightarrow e(A)} P_{\lambda\sigma}^{g \rightarrow e(B)} (\mu\nu|\sigma\lambda) \\
 V^{\text{Exch},(0)} &= -\frac{1}{2} \sum_{\mu\nu \in A} \sum_{\lambda\sigma \in B} P_{\nu\mu}^{g \rightarrow e(A)} P_{\lambda\sigma}^{g \rightarrow e(B)} (\mu\lambda|\sigma\nu) \\
 V^{\text{ET1},(0)} &= t_{H \rightarrow L}^A \left\{ (L^A|\mathcal{F}|L^B) + 2(L^A H^A|H^A L^B) - (L^A L^B|H^A H^A) \right\} \\
 V^{\text{ET2},(0)} &= t_{H \rightarrow L}^B \left\{ (L^A|\mathcal{F}|L^B) + 2(L^A H^B|H^B L^B) - (L^A L^B|H^B H^B) \right\} \\
 V^{\text{HT1},(0)} &= t_{H \rightarrow L}^A \left\{ -(H^A|\mathcal{F}|H^B) + 2(H^A L^A|L^A H^B) - (H^A H^B|L^A L^A) \right\} \\
 V^{\text{HT2},(0)} &= t_{H \rightarrow L}^B \left\{ -(H^A|\mathcal{F}|H^B) + 2(H^A L^B|L^B H^B) - (H^A H^B|L^B L^B) \right\} \\
 V^{\text{CT},(0)} &= 2(H^A L^B|L^A H^B) - (H^A H^B|L^A L^B)
 \end{aligned}$$

In the above, \mathcal{F} is the Fock operator whereas H and L denote the HOMO and LUMO orbitals, respectively. The overlap integrals between the basis states are approximated by

$$\begin{aligned}
 S_{12} &\equiv (\Phi_1|\Phi_2) \cong -\frac{1}{N_{el}^{AB}} \text{Tr} \left[\mathbf{P}^{g \rightarrow e(A)} \mathbf{s}^{AB} \mathbf{P}^{g \rightarrow e(B)} \mathbf{s}^{BA} \right] \\
 S_{13} &\equiv (\Phi_1|\Phi_3) \cong -\frac{t_{H \rightarrow L}^A}{N_{el}^{AB}} S_{LL}^{AB} \\
 S_{14} &\equiv (\Phi_1|\Phi_4) \cong +\frac{t_{H \rightarrow L}^A}{N_{el}^{AB}} S_{HH}^{AB} \\
 S_{24} &\equiv (\Phi_2|\Phi_4) \cong -\frac{t_{H \rightarrow L}^B}{N_{el}^{AB}} S_{LL}^{AB} \\
 S_{23} &\equiv (\Phi_2|\Phi_3) \cong +\frac{t_{H \rightarrow L}^B}{N_{el}^{AB}} S_{HH}^{AB} \\
 S_{34} &\equiv (\Phi_3|\Phi_4) \cong -\frac{1}{N_{el}^{AB}} S_{HH}^{AB} S_{LL}^{AB}
 \end{aligned}$$

where the overlap between molecular orbitals U and W is given by

$$S_{UW}^{AB} \equiv \mathbf{s}^{AB} : \mathbf{c}_U^A \otimes \mathbf{c}_W^B$$

and s^{AB} is the AO overlap matrix between molecule A and B atomic basis functions.

For a closed-shell system, the EET coupling constant for two electronic transitions can be given approximately by

$$V \approx V^{\text{Direct}} + V^{\text{Indirect}}$$

where the overlap-corrected direct and indirect coupling constants are

$$V^{\text{Direct}} = V^{\text{Coul}} + V^{\text{Exch}} + V^{\text{Ovrl}}$$

$$V^{\text{Indirect}} = V^{\text{TI-2}} + V^{\text{TI-3}}$$

with

$$V^{\text{TI-2}} = -\frac{V^{\text{ET1}}V^{\text{HT2}}}{E_3 - E_1} - \frac{V^{\text{ET2}}V^{\text{HT1}}}{E_4 - E_1}$$

$$V^{\text{TI-3}} = \frac{V^{\text{CT}}(V^{\text{ET1}}V^{\text{ET2}} + V^{\text{HT1}}V^{\text{HT2}})}{(E_3 - E_1)(E_4 - E_1)}$$

Fock matrix in AB space

In the current implementation, Fock matrix in the AB space, that is necessary to evaluate ET and HT matrix elements, can be defined as

1. the AB block of full Hartree-Fock SCF Fock matrix for entire system;
2. the zeroth-order Fock matrix that is composed of monomer's unperturbed ground-state 1-particle density matrices.

In the latter case, the Fock matrix in AO representation is given by:

$$F_{\alpha \in A, \beta \in B}^{AB} \approx T_{\alpha\beta} + V_{\alpha\beta}^{A(\text{nuc})} + V_{\alpha\beta}^{B(\text{nuc})} + \sum_{\mu \nu \in A} P_{\nu\mu}^{A(g)} G_{\alpha\beta, \mu\nu} + \sum_{\sigma \lambda \in B} P_{\lambda\sigma}^{B(g)} G_{\alpha\beta, \sigma\lambda}$$

where

$$G_{\alpha\beta, \gamma\delta} \equiv (\alpha\beta|\gamma\delta) - \frac{1}{2}(\alpha\delta|\gamma\beta)$$

Mulliken approximated exchange-like contributions.

Exchange and CT contributions require ERI's of type (AB,AB). It is instructive to approximate these contributions in terms of the Coulomb-like ERI's for the sake of testing of OEP-based approximations which are given in the next Section.

Application of the Mullipen approximation

$$(ij|kl) \approx \frac{1}{4} S_{ij} S_{kl} [(ii|kk) + (jj|kk) + (ii|ll) + (jj|ll)]$$

results in the following approximations to the exchange-like terms

$$V^{\text{Exch},(0)} \approx -\frac{1}{8} \sum_{\mu \nu \in A} \sum_{\lambda \sigma \in B} P_{\nu\mu}^{g \rightarrow e(A)} P_{\lambda\sigma}^{g \rightarrow e(B)} S_{\mu\lambda} S_{\sigma\nu} [(\mu\mu|\sigma\sigma) + (\lambda\lambda|\nu\nu) + (\mu\mu|\nu\nu) + (\lambda\lambda|\sigma\sigma)]$$

$$V^{\text{CT},(0)} \approx \frac{1}{2} S_{HL}^{AB} S_{LH}^{AB} [r_{HL}^A + r_{HL}^B + \rho_H^A \odot \rho_H^B + \rho_L^A \odot \rho_L^B]$$

$$- \frac{1}{4} S_{HH}^{AB} S_{LL}^{AB} [r_{HL}^A + r_{HL}^B + \rho_H^A \odot \rho_L^B + \rho_L^A \odot \rho_H^B]$$

The former can be rewritten in a more convenient to implement formula:

$$V^{\text{Exch},(0)} \approx -\frac{1}{4} \sum_{\mu \in A} \sum_{\nu \in B} (\mu\mu|\sigma\sigma) [\mathbf{P}^A \mathbf{s}^{AB}]_{\mu\sigma} [\mathbf{P}^B \mathbf{s}^{BA}]_{\sigma\mu} - \frac{1}{8} \sum_{\mu\nu \in A} P_{\nu\mu}^A (\mu\mu|\nu\nu) [\mathbf{s}^{AB} \mathbf{P}^B \mathbf{s}^{BA}]_{\mu\nu} - \frac{1}{8} \sum_{\sigma\lambda \in B} P_{\lambda\sigma}^B (\lambda\lambda|\sigma\sigma)$$

In the CT term,

$$\begin{aligned} r_{HL}^A &\equiv \rho_H^A \odot \rho_L^A \\ r_{HL}^B &\equiv \rho_H^B \odot \rho_L^B \end{aligned}$$

where the effective Coulombic interaction energies are defined by

$$\rho_U^A \odot \rho_W^B \equiv (U^A U^A | W^A W^A)$$

OEP-Based Methods

TODO

OEP-Based TI/CIS theory

After introducing OEP's, the original TI/CIS theory by Fujimoto is reformulated *without* approximation as TODO

18.42.2 Member Function Documentation

18.42.2.1 compute_benchmark()

```
double EETCouplingSolver::compute_benchmark (
    const std::string & method = "DEFAULT" ) [virtual]
```

Each solver object has one `DEFAULT` benchmark method

Parameters

<i>method</i>	- benchmark method
---------------	--------------------

Implements [oepdev::OEPDevSolver](#).

18.42.2.2 compute_oep_based()

```
double EETCouplingSolver::compute_oep_based (
```



```
const std::string & method = "DEFAULT" ) [virtual]
```

Each solver object has one `DEFAULT` OEP-based method.

Parameters

<i>method</i>	- flavour of OEP model
---------------	------------------------

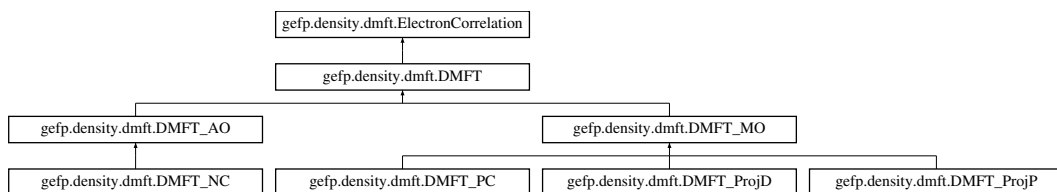
Implements [oepdev::OEPDevSolver](#).

The documentation for this class was generated from the following files:

- [oepdev/libsolver/solver.h](#)
- [oepdev/libsolver/solver_coupling_eet.cc](#)

18.43 gefp.density.dmft.ElectronCorrelation Class Reference

Inheritance diagram for `gefp.density.dmft.ElectronCorrelation`:



Static Public Member Functions

- def **degree_of_correlation** (dmft, scalar=True)

18.43.1 Detailed Description

\

The Electron Correlation: Dynamic and Non-dynamic Correlation.

The documentation for this class was generated from the following file:

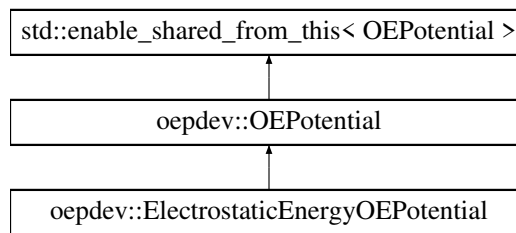
- [gefp/gefp/density/dmft.py](#)

18.44 oepdev::ElectrostaticEnergyOEPotential Class Reference

Generalized One-Electron Potential for Electrostatic Energy.

```
#include <oep.h>
```

Inheritance diagram for `oepdev::ElectrostaticEnergyOEPotential`:



Public Member Functions

- [ElectrostaticEnergyOEPotential](#) (SharedWavefunction [wfn](#), Options &options)
Only ESP-based potential is worth implementing.
- virtual void [compute](#) (const std::string &oepType) override
Compute matrix forms of all OEP's within a specified OEP type.
- virtual void [compute_3D](#) (const std::string &oepType, const double &x, const double &y, const double &z, std::shared_ptr< psi::Vector > &v) override
Compute value of potential in point x, y, z and save at v.
- virtual void [print_header](#) () const override
Header information.

Additional Inherited Members

18.44.1 Detailed Description

Contains the following OEP types:

- V

The documentation for this class was generated from the following files:

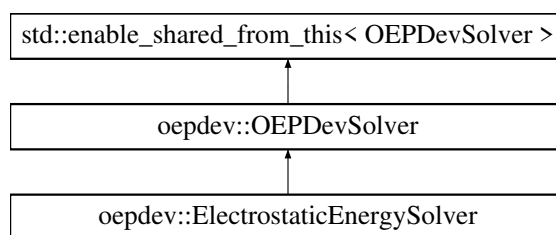
- oepdev/liboep/[oep.h](#)
- oepdev/liboep/oep_energy_coul.cc

18.45 oepdev::ElectrostaticEnergySolver Class Reference

Compute the Coulombic interaction energy between unperturbed wavefunctions.

```
#include <solver.h>
```

Inheritance diagram for oepdev::ElectrostaticEnergySolver:



Public Member Functions

- **ElectrostaticEnergySolver** (SharedWavefunctionUnion wfn_union)
- virtual double [compute_oep_based](#) (const std::string &method="DEFAULT")
Compute property by using OEP's.
- virtual double [compute_benchmark](#) (const std::string &method="DEFAULT")
Compute property by using benchmark method.

Additional Inherited Members

18.45.1 Detailed Description

The implemented methods are shown in below

Table 18.32: Methods available in the Solver

Keyword	Method Description
Benchmark Methods	
AO_EXPANDED	<i>Default.</i> Exact Coulombic energy from atomic orbital expansions.
MO_EXPANDED	Exact Coulombic energy from molecular orbital expansions
OEP-Based Methods	
ESP_SYMMETRIZED	<i>Default.</i> Coulombic energy from ESP charges interacting with nuclei and electronic density. Symmetrized with respect to monomers.
CAMM	Coulombic energy from CAMM distributions.

Below the detailed description of the above methods is given.

Benchmark Methods

Exact Coulombic energy from atomic orbital expansions.

The Coulombic interaction energy is given by

$$E^{\text{Coul}} = E^{\text{Nuc-Nuc}} + E^{\text{Nuc-El}} + E^{\text{El-El}}$$

where the nuclear-nuclear repulsion energy is

$$E^{\text{Nuc-Nuc}} = \sum_{x \in A} \sum_{y \in B} \frac{Z_x Z_y}{|\mathbf{r}_x - \mathbf{r}_y|}$$

the nuclear-electronic attraction energy is

$$E^{\text{Nuc-El}} = \sum_{x \in A} \sum_{\lambda \sigma \in B} Z_x V_{\lambda \sigma}^{(x)} \left(D_{\lambda \sigma}^{(\alpha)} + D_{\lambda \sigma}^{(\beta)} \right) + \sum_{y \in B} \sum_{\mu \nu \in A} Z_y V_{\mu \nu}^{(y)} \left(D_{\mu \nu}^{(\alpha)} + D_{\mu \nu}^{(\beta)} \right)$$

and the electron-electron repulsion energy is

$$E^{\text{El-El}} = \sum_{\mu \nu \in A} \sum_{\lambda \sigma \in B} \left\{ D_{\mu \nu}^{(\alpha)} + D_{\mu \nu}^{(\beta)} \right\} \left\{ D_{\lambda \sigma}^{(\alpha)} + D_{\lambda \sigma}^{(\beta)} \right\} (\mu \nu | \lambda \sigma)$$

In the above equations,

$$V_{\lambda \sigma}^{(x)} \equiv \int \frac{\phi_{\lambda}^*(\mathbf{r}) \phi_{\sigma}(\mathbf{r})}{|\mathbf{r} - \mathbf{r}_x|} d\mathbf{r}$$

Exact Coulombic energy from molecular orbital expansion.

This approach is fully equivalent to the atomic orbital expansion shown above. For the closed shell case, the Coulombic interaction energy is given by

$$E^{\text{Coul}} = E^{\text{Nuc-Nuc}} + E^{\text{Nuc-El}} + E^{\text{El-El}}$$

where the nuclear-nuclear repulsion energy is

$$E^{\text{Nuc-Nuc}} = \sum_{x \in A} \sum_{y \in B} \frac{Z_x Z_y}{|\mathbf{r}_x - \mathbf{r}_y|}$$

the nuclear-electronic attraction energy is

$$E^{\text{Nuc-El}} = 2 \sum_{i \in A} \sum_{y \in B} V_{ii}^{(y)} + 2 \sum_{j \in B} \sum_{x \in A} V_{jj}^{(x)}$$

and the electron-electron repulsion energy is

$$E^{\text{El-El}} = 4 \sum_{i \in A} \sum_{j \in B} (ii | jj)$$

OEP-Based Methods

Coulombic energy from ESP charges interacting with nuclei and electronic density.

In this approach, nuclear and electronic density of either species is approximated by ESP charges. In order to achieve symmetric expression, the interaction is computed twice (ESP of A interacting with density matrix and nuclear charges of B and vice versa) and then divided by 2. Thus,

$$E^{\text{Coul}} \approx \frac{1}{2} \left[\sum_{x \in A} \sum_{y \in B} \frac{Z_x q_y}{|\mathbf{r}_x - \mathbf{r}_y|} + \sum_{y \in B} \sum_{\mu \nu \in A} q_y V_{\mu \nu}^{(y)} \left(D_{\mu \nu}^{(\alpha)} + D_{\mu \nu}^{(\beta)} \right) + \sum_{y \in B} \sum_{x \in A} \frac{q_x Z_y}{|\mathbf{r}_x - \mathbf{r}_y|} + \sum_{x \in A} \sum_{\lambda \sigma \in B} q_x V_{\lambda \sigma}^{(x)} \left(D_{\lambda \sigma}^{(\alpha)} + D_{\lambda \sigma}^{(\beta)} \right) \right]$$

If the basis set is large and the number of ESP centres $q_{x(y)}$ is sufficient, the sum of first two contributions equals the sum of the latter two contributions.

Notes:

- This solver also computes and prints the ESP-ESP point charge interaction energy,

$$E^{\text{Coul,ESP}} \approx \sum_{x \in A} \sum_{y \in B} \frac{q_x q_y}{|\mathbf{r}_x - \mathbf{r}_y|}$$

for reference purposes.

- In order to construct this solver, **always** use the [OEPDevSolver::build](#) static factory method.

18.45.2 Member Function Documentation

18.45.2.1 compute_benchmark()

```
double ElectrostaticEnergySolver::compute_benchmark (
    const std::string & method = "DEFAULT" ) [virtual]
```

Each solver object has one DEFAULT benchmark method

Parameters

<i>method</i>	- benchmark method
---------------	--------------------

Implements [oepdev::OEPDevSolver](#).

18.45.2.2 compute_oep_based()

```
double ElectrostaticEnergySolver::compute_oep_based (
```

```
const std::string & method = "DEFAULT" ) [virtual]
```

Each solver object has one `DEFAULT` OEP-based method.

Parameters

<i>method</i>	- flavour of OEP model
---------------	------------------------

Implements [oepdev::OEPDevSolver](#).

The documentation for this class was generated from the following files:

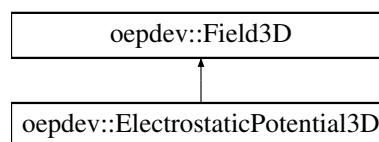
- [oepdev/libsolver/solver.h](#)
- [oepdev/libsolver/solver_energy_coul.cc](#)

18.46 oepdev::ElectrostaticPotential3D Class Reference

Electrostatic potential of a molecule.

```
#include <space3d.h>
```

Inheritance diagram for `oepdev::ElectrostaticPotential3D`:



Public Member Functions

- **ElectrostaticPotential3D** (const int &np, const double &padding, psi::SharedWavefunction [wfn](#), psi::Options &options)
- **ElectrostaticPotential3D** (const int &nx, const int &ny, const int &nz, const double &px, const double &py, const double &pz, psi::SharedWavefunction [wfn](#), psi::Options &options)
- virtual std::shared_ptr< psi::Vector > [compute_xyz](#) (const double &x, const double &y, const double &z)

Compute a value of 3D field at point (x, y, z)

- virtual void [print](#) () const

Print information of the object to Psi4 output.

Additional Inherited Members

18.46.1 Detailed Description

Computes the electrostatic potential of a molecule directly from the wavefunction. The electrostatic potential $v(\mathbf{r})$ at point \mathbf{r} is computed from the following formula:

$$v(\mathbf{r}) = v_{\text{nuc}}(\mathbf{r}) + v_{\text{el}}(\mathbf{r})$$

where the nuclear and electronic contributions are defined accordingly as

$$v_{\text{nuc}}(\mathbf{r}) = \sum_x \frac{Z_x}{|\mathbf{r} - \mathbf{r}_x|}$$

$$v_{\text{el}}(\mathbf{r}) = \sum_{\mu\nu} \left\{ D_{\mu\nu}^{(\alpha)} + D_{\mu\nu}^{(\beta)} \right\} V_{\nu\mu}(\mathbf{r})$$

In the above equations, Z_x denotes the charge of x th nucleus, $D_{\mu\nu}^{(\omega)}$ is the one-particle (relaxed) density matrix element in AO basis associated with the ω electron spin, and $V_{\mu\nu}(\mathbf{r})$ is the potential one-electron integral defined by

$$V_{\nu\mu}(\mathbf{r}) \equiv \int d\mathbf{r}' \phi_{\nu}^*(\mathbf{r}') \frac{1}{|\mathbf{r} - \mathbf{r}'|} \phi_{\mu}(\mathbf{r}')$$

The documentation for this class was generated from the following files:

- oepdev/lib3d/space3d.h
- oepdev/lib3d/space3d.cc

18.47 gefp.core.driver.Entry Class Reference

Public Member Functions

- `def __init__ (self, pade, description_short, description_full)`

Public Attributes

- `pade`
- `description_short`
- `description_full`

The documentation for this class was generated from the following file:

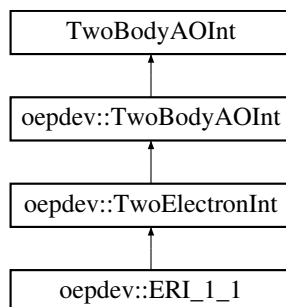
- `gefp/gefp/core/driver.py`

18.48 oepdev::ERI_1_1 Class Reference

2-centre ERI of the form $(a|O(2)|b)$ where $O(2) = 1/r_{12}$.

```
#include <eri.h>
```

Inheritance diagram for oepdev::ERI_1_1:



Public Member Functions

- [ERI_1_1](#) (const [IntegralFactory](#) *integral, int deriv=0, bool use_shell_pairs=false)
Constructor. Use [oepdev::IntegralFactory](#) to generate this object.
- [~ERI_1_1](#) ()
Destructor.

Protected Member Functions

- size_t [compute_doublet](#) (int, int)
Compute ERI's between 2 shells.

Protected Attributes

- double * [mdh_buffer_1_](#)
Buffer for McMurchie-Davidson-Hermite coefficients for monomial expansion (shell 1)
- double * [mdh_buffer_2_](#)
Buffer for McMurchie-Davidson-Hermite coefficients for monomial expansion (shell 2)

18.48.1 Detailed Description

ERI's are computed for a shell doublet $(P|Q)$ and stored in the `target_full_buffer`, accessible through `buffer()` method:

For each $(n_1, l_1, m_1) \in P$:
 For each $(n_2, l_2, m_2) \in Q$:

$$\text{ERI} = (A|B)[\{\alpha\}, \mathbf{n}, \mathbf{l}, \mathbf{m}]$$

For detailed description of the McMurchie-Davidson scheme, refer to [The Integral Package Library](#).

18.48.2 Implementation

A set of ERI's in a shell is decontracted as

$$(A|B)[\{\alpha\}, \mathbf{n}, \mathbf{l}, \mathbf{m}] = \sum_{ij} c_i(\alpha_1) c_j(\alpha_2) (i|j)[\{\alpha\}, \mathbf{n}, \mathbf{l}, \mathbf{m}]$$

where the primitive ERI is given by

$$(i|j)[\{\alpha\}, \mathbf{n}, \mathbf{l}, \mathbf{m}] = \sum_{N_1=0}^{n_1} \sum_{L_1=0}^{l_1} \sum_{M_1=0}^{m_1} \sum_{N_2=0}^{n_2} \sum_{L_2=0}^{l_2} \sum_{M_2=0}^{m_2} d_{N_1}^{n_1} d_{L_1}^{l_1} d_{M_1}^{m_1} d_{N_2}^{n_2} d_{L_2}^{l_2} d_{M_2}^{m_2} [N_1 L_1 M_1 | N_2 L_2 M_2]$$

The documentation for this class was generated from the following files:

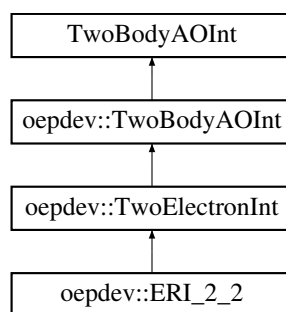
- oepdev/libints/[eri.h](#)
- oepdev/libints/eri.cc

18.49 oepdev::ERI_2_2 Class Reference

4-centre ERI of the form (ab|O(2)|cd) where O(2) = 1/r¹².

```
#include <eri.h>
```

Inheritance diagram for oepdev::ERI_2_2:



Public Member Functions

- [ERI_2_2](#) (const [IntegralFactory](#) *integral, int deriv=0, bool use_shell_pairs=false)
Constructor. Use [oepdev::IntegralFactory](#) to generate this object.
- [~ERI_2_2](#) ()
Destructor.

Protected Member Functions

- `size_t compute_quartet (int, int, int, int)`
Compute ERI's between 4 shells.

Protected Attributes

- `double * mdh_buffer_12_`
Buffer for McMurchie-Davidson-Hermite coefficients for binomial expansion (shells 1 and 2)
- `double * mdh_buffer_34_`
Buffer for McMurchie-Davidson-Hermite coefficients for binomial expansion (shells 3 and 4)

18.49.1 Detailed Description

ERI's are computed for a shell quartet (PQ|RS) and stored in the `target_full_buffer`, accessible through `buffer()` method:

For each $(n_1, l_1, m_1) \in P$:
 For each $(n_2, l_2, m_2) \in Q$:
 For each $(n_3, l_3, m_3) \in R$:
 For each $(n_4, l_4, m_4) \in S$:
 $\text{ERI} = (AB|CD)[\{\alpha\}, \mathbf{n}, \mathbf{l}, \mathbf{m}]$

For detailed description of the McMurchie-Davidson scheme, refer to [The Integral Package Library](#).

18.49.2 Implementation

A set of ERI's in a shell is decontracted as

$$(AB|CD)[\{\alpha\}, \mathbf{n}, \mathbf{l}, \mathbf{m}] = \sum_{ijkl} c_i(\alpha_1) c_j(\alpha_2) c_k(\alpha_3) c_l(\alpha_4) (ij|kl)[\{\alpha\}, \mathbf{n}, \mathbf{l}, \mathbf{m}]$$

where the primitive ERI is given by

$$(ij|kl)[\{\alpha\}, \mathbf{n}, \mathbf{l}, \mathbf{m}] = E_{ij}(\alpha_1, \alpha_2) E_{kl}(\alpha_3, \alpha_4) \\ \times \sum_{N_1=0}^{n_1+n_2} \sum_{L_1=0}^{l_1+l_2} \sum_{M_1=0}^{m_1+m_2} \sum_{N_2=0}^{n_3+n_4} \sum_{L_2=0}^{l_3+l_4} \sum_{M_2=0}^{m_3+m_4} d_{N_1}^{n_1 n_2} d_{L_1}^{l_1 l_2} d_{M_1}^{m_1 m_2} d_{N_2}^{n_3 n_4} d_{L_2}^{l_3 l_4} d_{M_2}^{m_3 m_4} [N_1 L_1 M_1 | N_2 L_2 M_2]$$

In the above equation, the multiplicative constants are given as

$$E_{ij}(\alpha_1, \alpha_2) = \exp \left[-\frac{\alpha_1 \alpha_2}{\alpha_1 + \alpha_2} |\mathbf{A} - \mathbf{B}|^2 \right] \\ E_{kl}(\alpha_3, \alpha_4) = \exp \left[-\frac{\alpha_3 \alpha_4}{\alpha_3 + \alpha_4} |\mathbf{C} - \mathbf{D}|^2 \right]$$

The documentation for this class was generated from the following files:

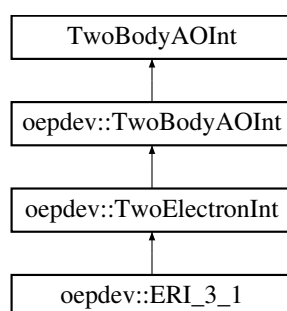
- oepdev/libints/[eri.h](#)
- oepdev/libints/eri.cc

18.50 oepdev::ERI_3_1 Class Reference

4-centre ERI of the form $(abc|O(2)|d)$ where $O(2) = 1/r^{12}$.

```
#include <eri.h>
```

Inheritance diagram for oepdev::ERI_3_1:



Public Member Functions

- [ERI_3_1](#) (const [IntegralFactory](#) *integral, int deriv=0, bool use_shell_pairs=false)
Constructor. Use [oepdev::IntegralFactory](#) to generate this object.
- [~ERI_3_1](#) ()
Destructor.

Protected Member Functions

- size_t [compute_quartet](#) (int, int, int, int)
Compute ERI's between 4 shells.

Protected Attributes

- double * [mdh_buffer_123_](#)
Buffer for McMurchie-Davidson-Hermite coefficients for trinomial expansion (shells 1, 2 and 3)
- double * [mdh_buffer_4_](#)
Buffer for McMurchie-Davidson-Hermite coefficients for monomial expansion (shell 4)

18.50.1 Detailed Description

ERI's are computed for a shell quartet (PQR|S) and stored in the `target_full_buffer`, accessible through `buffer()` method:

For each $(n_1, l_1, m_1) \in P$:
 For each $(n_2, l_2, m_2) \in Q$:
 For each $(n_3, l_3, m_3) \in R$:
 For each $(n_4, l_4, m_4) \in S$:
 $\text{ERI} = (ABC|D)[\{\alpha\}, \mathbf{n}, \mathbf{l}, \mathbf{m}]$

For detailed description of the McMurchie-Davidson scheme, refer to [The Integral Package Library](#).

18.50.2 Implementation

A set of ERI's in a shell is decontracted as

$$(ABC|D)[\{\alpha\}, \mathbf{n}, \mathbf{l}, \mathbf{m}] = \sum_{ijkl} c_i(\alpha_1) c_j(\alpha_2) c_k(\alpha_3) c_l(\alpha_4) (ijk|l)[\{\alpha\}, \mathbf{n}, \mathbf{l}, \mathbf{m}]$$

where the primitive ERI is given by

$$(ijk|l)[\{\alpha\}, \mathbf{n}, \mathbf{l}, \mathbf{m}] = E_{ijk}(\alpha_1, \alpha_2, \alpha_3) \times \sum_{N_1=0}^{n_1+n_2+n_3} \sum_{L_1=0}^{l_1+l_2+l_3} \sum_{M_1=0}^{m_1+m_2+m_3} \sum_{N_2=0}^{n_4} \sum_{L_2=0}^{l_4} \sum_{M_2=0}^{m_4} d_{N_1}^{n_1 n_2 n_3} d_{L_1}^{l_1 l_2 l_3} d_{M_1}^{m_1 m_2 m_3} d_{N_2}^{n_4} d_{L_2}^{l_4} d_{M_2}^{m_4} [N_1 L_1 M_1 | N_2 L_2 M_2]$$

In the above equation, the multiplicative constants are given as

$$E_{ijk}(\alpha_1, \alpha_2, \alpha_3) = \exp \left[-\frac{\alpha_1 \alpha_2}{\alpha_1 + \alpha_2} |\mathbf{A} - \mathbf{B}|^2 \right] \exp \left[-\frac{(\alpha_1 + \alpha_2) \alpha_3}{\alpha_1 + \alpha_2 + \alpha_3} |\mathbf{P} - \mathbf{C}|^2 \right]$$

The documentation for this class was generated from the following files:

- [oepdev/libints/eri.h](#)
- [oepdev/libints/eri.cc](#)

18.51 oepdev::ESPSolver Class Reference

Charges from Electrostatic Potential (ESP). A solver-type class.

```
#include <esp.h>
```

Public Member Functions

- [ESPSolver](#) (SharedField3D field)

- Construct from 3D vector field.
- [ESPSolver](#) (SharedField3D field, psi::SharedMatrix [centres](#))
Construct from 3D vector field.
- virtual [~ESPSolver](#) ()
Destructor.
- virtual psi::SharedMatrix [charges](#) () const
Get the (fit) charges.
- virtual psi::SharedMatrix [centres](#) () const
Get the charge distribution centres.
- virtual void [set_charge_sums](#) (psi::SharedVector s)
Set the charge sums Q_p .
- virtual void [set_charge_sums](#) (const double &s)
Set the charge sums Q_p (equal to all fields)
- virtual void [compute](#) ()
Perform fitting of effective charges.

Protected Attributes

- const int [nCentres_](#)
Number of fit centres.
- const int [nFields_](#)
Number of fields to fit.
- SharedField3D [field_](#)
Scalar field.
- psi::SharedMatrix [charges_](#)
Charges to be fit.
- psi::SharedMatrix [centres_](#)
Centres, at which fit charges will reside.
- psi::SharedVector [charge_sums_](#)
Vector of sums of partial charges.

18.51.1 Detailed Description

Solves the least-squares problem to fit the generalized charges $q_{m;p}$, that reproduce the reference generalized potential $v_p^{\text{ref}}(\mathbf{r})$ supplied by the [Field3D](#) object:

$$\int d\mathbf{r}' \left[v_p^{\text{ref}}(\mathbf{r}') - \sum_m \frac{q_{m;p}}{|\mathbf{r}' - \mathbf{r}_m|} \right]^2 \rightarrow \text{minimize}$$

The charges are subject to the following constraint:

$$\sum_m q_{m;p} = Q_p \text{ for all } p$$

Method description.

M generalized charges is found by solving the matrix equation

$$\begin{pmatrix} \mathbf{A} & 1 \\ 1 & 0 \end{pmatrix}^{-1} \cdot \begin{pmatrix} \mathbf{b}_p \\ Q_p \end{pmatrix} = \begin{pmatrix} \mathbf{q}_p \\ \lambda \end{pmatrix}$$

where the \mathbf{A} matrix of dimension $(M+1) \times (M+1)$ and \mathbf{b}_p vector of length $M+1$ are given as

$$A_{mn} = \sum_i \frac{1}{r_{im} r_{in}}$$

$$b_{m;p} = \sum_i \frac{v_p^{\text{ref}}(\mathbf{r}_m)}{r_{im}}$$

In the above equation, summations run over all sample points, at which reference potential is known. The solution is stored in the $M \times N$ matrix, where N is the dimensionality of the 3D vector field (i.e., the number of potentials supplied, p_{max}). As a default, $Q_p = 0$ for all potentials. This can be set by `oepdev::ESPSolver::set_charge_sums` method.

Note

Useful options:

- `ESP_PAD_SPHERE` - Padding spherical radius for random points selection. Default: 10.0 [A.U.]
- `ESP_NPOINTS_PER_ATOM` - Number of random points per atom in a molecule. Default: 1500
- `ESP_VDW_RADIUS_C` - The vdW radius for carbon atom. Default: 3.0 [A.U.]
- `ESP_VDW_RADIUS_H` - The vdW radius for hydrogen atom. Default: 4.0 [A.U.]
- `ESP_VDW_RADIUS_N` - The vdW radius for nitrogen atom. Default: 2.4 [A.U.]
- `ESP_VDW_RADIUS_O` - The vdW radius for oxygen atom. Default: 5.6 [A.U.]
- `ESP_VDW_RADIUS_F` - The vdW radius for fluorine atom. Default: 2.3 [A.U.]
- `ESP_VDW_RADIUS_CL` - The vdW radius for chlorine atom. Default: 2.9 [A.U.]

18.51.2 Constructor & Destructor Documentation**18.51.2.1 ESPSolver() [1/2]**

```
oepdev::ESPSolver::ESPSolver (
    SharedField3D field )
```

Assume that the centres are on atoms associated with the 3D vector field.

Parameters

<i>field</i>	- oepdev 3D vector field object
--------------	---------------------------------

18.51.2.2 ESPSolver() [2/2]

```
oepdev::ESPSolver::ESPSolver (
    SharedField3D field,
    psi::SharedMatrix centres )
```

Solve ESP equations for a custom set of charge distribution centres.

Parameters

<i>field</i>	- oepdev 3D vector field object
<i>centres</i>	- matrix with coordinates of charge distribution centres

The documentation for this class was generated from the following files:

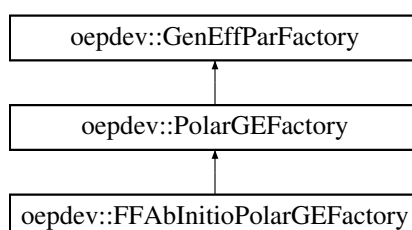
- oepdev/lib3d/[esp.h](#)
- oepdev/lib3d/[esp.cc](#)

18.52 oepdev::FFAbInitioPolarGEFactory Class Reference

Polarization GEFP Factory from First Principles: Finite-Difference Model. Arbitrary level of theory.

```
#include <gefp.h>
```

Inheritance diagram for oepdev::FFAbInitioPolarGEFactory:



Public Member Functions

- **FFAbInitioPolarGEFactory** (std::shared_ptr< psi::Wavefunction > [wfn](#), psi::Options &opt)
- virtual std::shared_ptr< [GenEffPar](#) > [compute](#) (void)
Compute the density matrix susceptibility tensors.

Additional Inherited Members

18.52.1 Detailed Description

Implements creation of the density matrix susceptibility tensors. Does not guarantee the idempotency of the density matrix in LCAO-MO variation, but for weak electric fields the idempotency is to be expected up to first order. The density matrix susceptibility tensor is represented by:

$$\delta D_{\alpha\beta} = \mathbf{B}_{\alpha\beta}^{(1)} \cdot \mathbf{F} + \mathbf{B}_{\alpha\beta}^{(2)} : \mathbf{F} \otimes \mathbf{F}$$

where $\mathbf{B}_{\alpha\beta}^{(1)}$ is the density matrix dipole polarizability defined as

$$\mathbf{B}_{\alpha\beta}^{(1)} = \left. \frac{\partial D_{\alpha\beta}}{\partial \mathbf{F}} \right|_{\mathbf{F}=\mathbf{0}}$$

whereas $\mathbf{B}_{\alpha\beta}^{(2)}$ is the density matrix dipole-dipole hyperpolarizability,

$$\mathbf{B}_{\alpha\beta}^{(2)} = \left. \frac{1}{2} \frac{\partial^2 D_{\alpha\beta}}{\partial \mathbf{F} \otimes \partial \mathbf{F}} \right|_{\mathbf{F}=\mathbf{0}}$$

The first derivative is evaluated numerically from central finite-field 3-point formula,

$$f' = \frac{f(h) - f(-h)}{2h} + \mathcal{O}(h^2)$$

where h is the differentiation step. Second derivatives are evaluated from the following formulae:

$$f_{uu} = \frac{f(h) + f(-h) - 2f(0)}{h^2} + \mathcal{O}(h^2)$$

$$f_{uw} = \frac{f(h, h) + f(-h, -h) + 2f(0) - f(h, 0) - f(-h, 0) - f(0, h) - f(0, -h)}{2h^2} + \mathcal{O}(h^2)$$

As long as the second-order susceptibility is considered, this susceptibility model works well for uniform weak, moderate and strong electric fields.

The documentation for this class was generated from the following files:

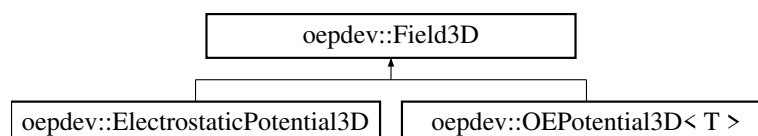
- oepdev/libgefp/gefp.h
- oepdev/libgefp/gefp_polar_ffabinitio.cc

18.53 oepdev::Field3D Class Reference

General Vector Field in 3D Space. Abstract base.

```
#include <space3d.h>
```

Inheritance diagram for oepdev::Field3D:



Public Member Functions

- **Field3D** (const int &ndim, const int &np, const double &pad, psi::SharedWavefunction **wfn**, psi::Options &options)
Construct potential on random grid by providing wavefunction. Excludes space within vdW volume.
- **Field3D** (const int &ndim, const int &nx, const int &ny, const int &nz, const double &px, const double &py, const double &pz, std::shared_ptr< psi::Wavefunction > **wfn**, psi::Options &options)
Construct potential on cube grid by providing wavefunction.
- virtual **~Field3D** ()
Destructor.
- virtual int **npoints** () const
Get the number of points at which the 3D field is defined.
- virtual std::shared_ptr< **PointsCollection3D** > **points_collection** () const
Get the collection of points.
- virtual std::shared_ptr< psi::Matrix > **data** () const
Get the data matrix in a form $\{ [x, y, z, f_1(x, y, z), f_2(x, y, z), \dots, f_n(x, y, z)] \}$ where $n = \text{ndim}$.
- virtual std::shared_ptr< psi::Wavefunction > **wfn** () const
Get the wavefunction.
- virtual bool **is_computed** () const
Get the information if data is already computed or not.
- int **dimension** () const
Get the number of fields.
- virtual void **compute** ()
Compute the 3D field in each point from the point collection.
- virtual std::shared_ptr< psi::Vector > **compute_xyz** (const double &x, const double &y, const double &z)=0
Compute a value of 3D field at point (x, y, z)
- virtual void **write_cube_file** (const std::string &name)
Write the cube file (only for Cube collections, otherwise does nothing)
- virtual void **print** () const =0
Print information of the object to Psi4 output.

Static Public Member Functions

- static shared_ptr< **Field3D** > **build** (const std::string &type, const int &np, const double &pad, psi::SharedWavefunction **wfn**, psi::Options &options, const int &ndim=1)
Build 3D field of random points. vdW volume is excluded.
- static shared_ptr< **Field3D** > **build** (const std::string &type, const int &nx, const int &ny, const int &nz, const double &px, const double &py, const double &pz, psi::SharedWavefunction **wfn**, psi::Options &options, const int &ndim=1)
Build 3D field of points on a g09-cube grid.

Protected Attributes

- `std::shared_ptr< PointsCollection3D > pointsCollection_`
Collection of points at which the 3D field is to be computed.
- `std::shared_ptr< psi::Matrix > data_`
The data matrix in a form $\{ [x, y, z, f_1(x, y, z), f_2(x, y, z), \dots, f_n(x, y, z)] \}$ where $n = nDim_$.
- `std::shared_ptr< psi::Wavefunction > wfn_`
Wavefunction.
- `psi::Matrix geom_`
Geometry of a molecule.
- `std::shared_ptr< psi::IntegralFactory > fact_`
Integral factory.
- `std::shared_ptr< psi::Matrix > pot_`
Matrix of potential one-electron integrals.
- `std::shared_ptr< psi::OneBodyAOInt > oneInt_`
One-electron integral shared pointer.
- `std::shared_ptr< PotentialInt > potInt_`
One-electron potential shared pointer.
- `std::shared_ptr< psi::BasisSet > primary_`
Basis set.
- `int nbf_`
Number of basis functions.
- `int nDim_`
Dimensionality of the 3D field (1: scalar field, 2>: vector field)
- `bool isComputed_`
Has data already computed?

18.53.1 Detailed Description

Create vector field defined at points distributed randomly or as an ordered g09 cube-like collection. Currently implemented fields are:

- Electrostatic potential - computes electrostatic potential (requires wavefunction)
- Template of generic classes - compute custom vector fields (requires generic object that is able to compute the field in 3D space)

Note: Always create instances by using static factory methods `build`. The following types of 3D vector fields are currently implemented:

- ELECTROSTATIC POTENTIAL

18.53.2 Constructor & Destructor Documentation

18.53.2.1 Field3D()

```
oepdev::Field3D::Field3D (
    const int & ndim,
    const int & nx,
    const int & ny,
    const int & nz,
    const double & px,
    const double & py,
    const double & pz,
    std::shared_ptr< psi::Wavefunction > wfn,
    psi::Options & options )
```

Construct potential on random grid by providing molecule. Excludes space within vdW volume Field3D(const int& ndim, const int& np, const double& pad, psi::SharedMolecule mol, psi::Options& options);

18.53.3 Member Function Documentation

18.53.3.1 build() [1/2]

```
std::shared_ptr< Field3D > oepdev::Field3D::build (
    const std::string & type,
    const int & np,
    const double & pad,
    psi::SharedWavefunction wfn,
    psi::Options & options,
    const int & ndim = 1 ) [static]
```

Parameters

<i>ndim</i>	- dimensionality of 3D field (1: scalar field, >2: vector field)
<i>type</i>	- type of 3D field
<i>np</i>	- number of points
<i>pad</i>	- radius padding of a minimal sphere enclosing the molecule
<i>wfn</i>	- Psi4 Wavefunction containing the molecule
<i>options</i>	- Psi4 options

18.53.3.2 build() [2/2]

```
std::shared_ptr< Field3D > oepdev::Field3D::build (
    const std::string & type,
    const int & nx,
    const int & ny,
    const int & nz,
    const double & px,
    const double & py,
    const double & pz,
    psi::SharedWavefunction wfn,
    psi::Options & options,
    const int & ndim = 1 ) [static]
```

Parameters

<i>ndim</i>	- dimensionality of 3D field (1: scalar field, >2: vector field)
<i>type</i>	- type of 3D field
<i>nx</i>	- number of points along x direction
<i>ny</i>	- number of points along y direction
<i>nz</i>	- number of points along z direction
<i>px</i>	- padding distance along x direction
<i>py</i>	- padding distance along y direction
<i>pz</i>	- padding distance along z direction
<i>wfn</i>	- Psi4 Wavefunction containing the molecule
<i>options</i>	- Psi4 options

The documentation for this class was generated from the following files:

- oepdev/lib3d/space3d.h
- oepdev/lib3d/space3d.cc

18.54 oepdev::Fourier9 Struct Reference

Simple structure to hold the Fourier series expansion coefficients for $N=4$.

```
#include <unitary_optimizer.h>
```

Public Attributes

- double **a0**
- double **a1**
- double **a2**
- double **a3**
- double **a4**

- double **b1**
- double **b2**
- double **b3**
- double **b4**

18.54.1 Detailed Description

The documentation for this struct was generated from the following file:

- oepdev/libutil/[unitary_optimizer.h](#)

18.55 oepdev::GenEffFrag Class Reference

Generalized Effective Fragment. Container Class.

```
#include <gefp.h>
```

Public Member Functions

- [GenEffFrag](#) ()
Initialize with default name of GEFP (Default)
- [GenEffFrag](#) (std::string name)
Initialize with custom name of GEFP.
- [~GenEffFrag](#) ()
Destruct.
- void [rotate](#) (std::shared_ptr< psi::Matrix > R)
Rotate.
- void [translate](#) (std::shared_ptr< psi::Vector > T)
Translate.
- void [superimpose](#) (std::shared_ptr< psi::Matrix > targetXYZ, std::vector< int > supList)
Superimpose.
- void [set_gefp_polarization](#) (const std::shared_ptr< [GenEffPar](#) > &par)
Set the Density Matrix Susceptibility Tensor Object.
- void [set_dmat_dipole_polarizability](#) (const std::vector< std::vector< std::shared_ptr< psi::Matrix >>> &susc)
Set the Density Matrix Dipole Polarizability.
- void [set_dmat_dipole_dipole_hyperpolarizability](#) (const std::vector< std::vector< std::shared_ptr< psi::Matrix >>> &susc)
Set the Density Matrix Dipole-Dipole Hyperpolarizability.
- void [set_dmat_quadrupole_polarizability](#) (const std::vector< std::vector< std::shared_ptr< psi::Matrix >>> &susc)

Set the Density Matrix Quadrupole Polarizability.

- `std::shared_ptr< psi::Matrix > susceptibility` (int fieldRank, int fieldGradientRank, int i, int x) const

Grab the Density Matrix Susceptibility.

- `std::vector< std::shared_ptr< psi::Matrix > > susceptibility` (int fieldRank, int fieldGradientRank, int i) const

Grab the Density Matrix Susceptibility.

- `std::vector< std::vector< std::shared_ptr< psi::Matrix > > > susceptibility` (int fieldRank, int fieldGradientRank) const

Grab the Density Matrix Susceptibility.

Public Attributes

- `std::map< std::string, std::shared_ptr< GenEffPar > > parameters`

Dictionary of All GEF Parameters.

Protected Attributes

- `std::string name_`

Name of GEFP.

- `std::shared_ptr< GenEffPar > densityMatrixSusceptibilityGEF_`

Density Matrix Susceptibility Tensor.

- `std::shared_ptr< GenEffPar > electrostaticEnergyGEF_`

Electrostatic Energy Effective One-Electron Potential.

- `std::shared_ptr< GenEffPar > repulsionEnergyGEF_`

Exchange-Repulsion Effective One-Electron Potential.

- `std::shared_ptr< GenEffPar > chargeTransferEnergyGEF_`

Charge-Transfer Effective One-Electron Potential.

- `std::shared_ptr< GenEffPar > EETCouplingConstantGEF_`

EET Coupling Effective One-Electron Potential.

18.55.1 Detailed Description

Describes the GEFP fragment that is in principle designed to work at correlated levels of theory.

18.55.2 Member Function Documentation

18.55.2.1 susceptibility() [1/3]

```
std::shared_ptr<psi::Matrix> oepdev::GenEffFrag::susceptibility (
    int fieldRank,
    int fieldGradientRank,
    int i,
    int x ) const [inline]
```

Parameters

<i>fieldRank</i>	- power dependency with respect to the electric field
<i>fieldGradientRank</i>	- power dependency with respect to the electric field gradient
<i>i</i>	- id of the distributed site
<i>x</i>	- id of the composite Cartesian component

18.55.2.2 susceptibility() [2/3]

```
std::vector<std::shared_ptr<psi::Matrix> > oepdev::GenEffFrag::susceptibility
(
    int fieldRank,
    int fieldGradientRank,
    int i ) const [inline]
```

Parameters

<i>fieldRank</i>	- power dependency with respect to the electric field
<i>fieldGradientRank</i>	- power dependency with respect to the electric field gradient
<i>i</i>	- id of the distributed site

18.55.2.3 susceptibility() [3/3]

```
std::vector<std::vector<std::shared_ptr<psi::Matrix> > > oepdev::GenEffFrag::susceptibility
(
    int fieldRank,
    int fieldGradientRank ) const [inline]
```

Parameters

<i>fieldRank</i>	- power dependency with respect to the electric field
<i>fieldGradientRank</i>	- power dependency with respect to the electric field gradient

The documentation for this class was generated from the following files:

- [oepdev/libgefp/gefp.h](#)
- [oepdev/libgefp/gefp.cc](#)

18.56 oepdev::GenEffPar Class Reference

Generalized Effective Fragment Parameters. Container Class.

```
#include <gefp.h>
```

Public Member Functions

- [GenEffPar](#) (std::string name)
Create with name of this parameter type.
- [~GenEffPar](#) ()
Destruct.
- void [set_susceptibility](#) (int fieldRank, int fieldGradientRank, const std::vector< std::vector< std::shared_ptr< psi::Matrix >>> &susc)
Set the Density Matrix Susceptibility.
- void [set_dipole_polarizability](#) (const std::vector< std::vector< std::shared_ptr< psi::Matrix >>> &susc)
Set The Density Matrix Dipole Polarizability.
- void [set_dipole_dipole_hyperpolarizability](#) (const std::vector< std::vector< std::shared_ptr< psi::Matrix >>> &susc)
Set The Density Matrix Dipole-Dipole Hyperpolarizability.
- void [set_quadropole_polarizability](#) (const std::vector< std::vector< std::shared_ptr< psi::Matrix >>> &susc)
Set The Density Matrix Quadrupole Polarizability.
- void [set_centres](#) (const std::vector< std::shared_ptr< psi::Vector >> ¢res)
Set the distributed centres' positions.
- void [allocate](#) (int fieldRank, int fieldGradientRank, int nsites, int nbf)
Allocate the Density Matrix Susceptibility.
- void [allocate_dipole_polarizability](#) (int nsites, int nbf)
Allocate The Density Matrix Dipole Polarizability.
- void [allocate_dipole_dipole_hyperpolarizability](#) (int nsites, int nbf)
Allocate The Density Matrix Dipole-Dipole Hyperpolarizability.
- void [allocate_quadropole_polarizability](#) (int nsites, int nbf)
Allocate The Density Matrix Quadrupole Polarizability.
- bool **hasDensityMatrixDipolePolarizability** () const
- bool **hasDensityMatrixDipoleDipoleHyperpolarizability** () const
- bool **hasDensityMatrixQuadrupolePolarizability** () const
- std::shared_ptr< psi::Matrix > [susceptibility](#) (int fieldRank, int fieldGradientRank, int i, int x) const

Grab the Density Matrix Susceptibility.

- `std::vector< std::shared_ptr< psi::Matrix > > susceptibility` (int fieldRank, int fieldGradientRank, int i) const

Grab the Density Matrix Susceptibility.

- `std::vector< std::vector< std::shared_ptr< psi::Matrix > > > susceptibility` (int fieldRank, int fieldGradientRank) const

Grab the Density Matrix Susceptibility.

- `std::vector< std::vector< std::shared_ptr< psi::Matrix > > > dipole_polarizability` () const

Grab the density matrix dipole polarizability tensor.

- `std::vector< std::shared_ptr< psi::Matrix > > dipole_polarizability` (int i) const

Grab the density matrix dipole polarizability tensor's x-th component.

- `std::shared_ptr< psi::Matrix > dipole_polarizability` (int i, int x) const

Grab the density matrix dipole polarizability tensor's x-th component of the i-th distributed site.

- `std::vector< std::vector< std::shared_ptr< psi::Matrix > > > dipole_dipole_hyperpolarizability` () const

Grab the density matrix dipole-dipole hyperpolarizability tensor.

- `std::vector< std::shared_ptr< psi::Matrix > > dipole_dipole_hyperpolarizability` (int i) const

Grab the density matrix dipole-dipole hyperpolarizability tensor's x-th component.

- `std::shared_ptr< psi::Matrix > dipole_dipole_hyperpolarizability` (int i, int x) const

Grab the density matrix dipole-dipole hyperpolarizability tensor's x-th component of the i-th distributed site.

- `std::vector< std::vector< std::shared_ptr< psi::Matrix > > > quadrupole_polarizability` () const

Grab the density matrix quadrupole polarizability tensor.

- `std::vector< std::shared_ptr< psi::Matrix > > quadrupole_polarizability` (int i) const

Grab the density matrix quadrupole polarizability tensor's x-th component.

- `std::shared_ptr< psi::Matrix > quadrupole_polarizability` (int i, int x) const

Grab the density matrix quadrupole polarizability tensor's x-th component of the i-th distributed site.

- `std::vector< std::shared_ptr< psi::Vector > > centres` () const

Grab the centres' positions.

- `std::shared_ptr< psi::Vector > centre` (int i) const

Grab the position of the i-th distributed site.

- `std::shared_ptr< psi::Matrix > compute_density_matrix` (std::shared_ptr< psi::Vector > field)

Compute the density matrix due to the uniform electric field perturbation.

- `std::shared_ptr< psi::Matrix > compute_density_matrix` (double fx, double fy, double fz)

Compute the density matrix due to the uniform electric field perturbation.

- `std::shared_ptr< psi::Matrix > compute_density_matrix` (std::vector< std::shared_ptr< psi::Vector > > fields)

Compute the density matrix due to the non-uniform electric field perturbation.

- `std::shared_ptr< psi::Matrix > compute_density_matrix` (`std::vector< std::shared_ptr< psi::Vector >> fields`, `std::vector< std::shared_ptr< psi::Matrix >> grads`)

Compute the density matrix due to the non-uniform electric field perturbation.

Protected Attributes

- `std::string name_`
The Name of Parameter Type.
- `std::vector< std::vector< std::shared_ptr< psi::Matrix > > > densityMatrixDipolePolarizability_`
The Density Matrix Dipole Polarizability.
- `std::vector< std::vector< std::shared_ptr< psi::Matrix > > > densityMatrixDipoleDipoleHyperpolarizability_`
The Density Matrix Dipole-Dipole Hyperpolarizability.
- `std::vector< std::vector< std::shared_ptr< psi::Matrix > > > densityMatrixQuadrupolePolarizability_`
The Density Matrix Quadrupole Polarizability.
- `std::vector< std::shared_ptr< psi::Vector > > distributedCentres_`
The Positions of the Distributed Centres.
- `bool hasDensityMatrixDipolePolarizability_`
- `bool hasDensityMatrixDipoleDipoleHyperpolarizability_`
- `bool hasDensityMatrixQuadrupolePolarizability_`

18.56.1 Detailed Description

18.56.2 Member Function Documentation

18.56.2.1 allocate()

```
void oepdev::GenEffPar::allocate (
    int fieldRank,
    int fieldGradientRank,
    int nsites,
    int nbf ) [inline]
```

Parameters

<i>fieldRank</i>	- power dependency with respect to the electric field \mathbf{F}
<i>fieldGradientRank</i>	- power dependency with respect to the electric field gradient $\nabla \otimes \mathbf{F}$
<i>nsites</i>	- number of distributed sites
<i>nbf</i>	- number of basis functions in the basis set

The following susceptibilities are supported (fieldRank, fieldGradientRank):

- (1, 0) - dipole polarizability, interacts with \mathbf{F}
- (2, 0) - dipole-dipole hyperpolarizability, interacts with $\mathbf{F} \otimes \mathbf{F}$
- (0, 1) - quadrupole polarizability, interacts with $\nabla \otimes \mathbf{F}$

18.56.2.2 compute_density_matrix() [1/4]

```
std::shared_ptr< psi::Matrix > oepdev::GenEffPar::compute_density_matrix (
    std::shared_ptr< psi::Vector > field )
```

Parameters

<i>field</i>	- the uniform electric field vector (A.U.)
--------------	--

18.56.2.3 compute_density_matrix() [2/4]

```
std::shared_ptr< psi::Matrix > oepdev::GenEffPar::compute_density_matrix (
    double fx,
    double fy,
    double fz )
```

Parameters

<i>fx</i>	- x-th Cartesian component of the uniform electric field vector (A.U.)
<i>fy</i>	- y-th Cartesian component of the uniform electric field vector (A.U.)
<i>fz</i>	- z-th Cartesian component of the uniform electric field vector (A.U.)

18.56.2.4 compute_density_matrix() [3/4]

```
std::shared_ptr< psi::Matrix > oepdev::GenEffPar::compute_density_matrix (
    std::vector< std::shared_ptr< psi::Vector >> fields )
```

Parameters

<i>fields</i>	- the list of non-uniform electric field vector (A.U.) evaluated at the distributed DMatPol sites
---------------	---

18.56.2.5 compute_density_matrix() [4/4]

```
std::shared_ptr< psi::Matrix > oepdev::GenEffPar::compute_density_matrix (
    std::vector< std::shared_ptr< psi::Vector >> fields,
    std::vector< std::shared_ptr< psi::Matrix >> grads )
```

Parameters

<i>fields</i>	- the list of electric field vectors (A.U.) evaluated at the distributed DMatPol sites
<i>grads</i>	- the list of electric field gradient matrices (A.U.) evaluated at the distributed DMatPol sites

18.56.2.6 set_susceptibility()

```
void oepdev::GenEffPar::set_susceptibility (
    int fieldRank,
    int fieldGradientRank,
    const std::vector< std::vector< std::shared_ptr< psi::Matrix >>>
    & susc ) [inline]
```

Parameters

<i>fieldRank</i>	- power dependency with respect to the electric field \mathbf{F}
<i>fieldGradientRank</i>	- power dependency with respect to the electric field gradient $\nabla \otimes \mathbf{F}$
<i>susc</i>	- the susceptibility tensor

The following susceptibilities are supported (fieldRank, fieldGradientRank):

- (1, 0) - dipole polarizability, interacts with \mathbf{F}
- (2, 0) - dipole-dipole hyperpolarizability, interacts with $\mathbf{F} \otimes \mathbf{F}$
- (0, 1) - quadrupole polarizability, interacts with $\nabla \otimes \mathbf{F}$

18.56.2.7 susceptibility() [1/3]

```
std::shared_ptr<psi::Matrix> oepdev::GenEffPar::susceptibility (
    int fieldRank,
    int fieldGradientRank,
    int i,
    int x ) const [inline]
```

Parameters

<i>fieldRank</i>	- power dependency with respect to the electric field
<i>fieldGradientRank</i>	- power dependency with respect to the electric field gradient
<i>i</i>	- id of the distributed site
<i>x</i>	- id of the composite Cartesian component

The following susceptibilities are supported (fieldRank, fieldGradientRank):

- (1, 0) - dipole polarizability, interacts with \mathbf{F}
- (2, 0) - dipole-dipole hyperpolarizability, interacts with $\mathbf{F} \otimes \mathbf{F}$
- (0, 1) - quadrupole polarizability, interacts with $\nabla \otimes \mathbf{F}$

The distributed sites are assumed to be atomic sites or molecular orbital centroids (depending on the polarization factory used). For the electric field, the composite Cartesian index is just an ordinary Cartesian index. For the electric field gradient and electric field squared, the composite Cartesian index is given as

$$I(x,y) = 3x + y$$

where the values of 0, 1 and 2 correspond to x , y and z Cartesian components, respectively. Therefore, in the latter case, there is 9 distinct composite Cartesian components.

18.56.2.8 susceptibility() [2/3]

```
std::vector<std::shared_ptr<psi::Matrix> > oepdev::GenEffPar::susceptibility
(
    int fieldRank,
    int fieldGradientRank,
    int i ) const [inline]
```

Parameters

<i>fieldRank</i>	- power dependency with respect to the electric field
<i>fieldGradientRank</i>	- power dependency with respect to the electric field gradient
<i>i</i>	- id of the distributed site

The following susceptibilities are supported (fieldRank, fieldGradientRank):

- (1, 0) - dipole polarizability, interacts with \mathbf{F}
- (2, 0) - dipole-dipole hyperpolarizability, interacts with $\mathbf{F} \otimes \mathbf{F}$
- (0, 1) - quadrupole polarizability, interacts with $\nabla \otimes \mathbf{F}$

The distributed sites are assumed to be atomic sites or molecular orbital centroids (depending on the polarization factory used).

18.56.2.9 susceptibility() [3 / 3]

```
std::vector<std::vector<std::shared_ptr<psi::Matrix> > > oepdev::GenEffPar::susceptibility(
(
    int fieldRank,
    int fieldGradientRank ) const [inline]
```

Parameters

<i>fieldRank</i>	- power dependency with respect to the electric field
<i>fieldGradientRank</i>	- power dependency with respect to the electric field gradient

The following susceptibilities are supported (fieldRank, fieldGradientRank):

- (1, 0) - dipole polarizability, interacts with \mathbf{F}
- (2, 0) - dipole-dipole hyperpolarizability, interacts with $\mathbf{F} \otimes \mathbf{F}$
- (0, 1) - quadrupole polarizability, interacts with $\nabla \otimes \mathbf{F}$

The documentation for this class was generated from the following files:

- oepdev/libgefp/gefp.h
- oepdev/libgefp/gefp.cc

18.57 oepdev::GenEffParFactory Class Reference

Generalized Effective Fragment Factory. Abstract Base.

```
#include <gefp.h>
```

Inheritance diagram for oepdev::GenEffParFactory:



Public Member Functions

- [GenEffParFactory](#) (std::shared_ptr< psi::Wavefunction > [wfn](#), psi::Options &opt)
Construct from wavefunction and Psi4 options.
- virtual [~GenEffParFactory](#) ()
Destruct.
- virtual std::shared_ptr< [GenEffPar](#) > [compute](#) (void)=0
Compute the fragment parameters.
- virtual std::shared_ptr< psi::Wavefunction > [wfn](#) (void) const

Grab wavefunction.

- virtual psi::Options & [options](#) (void) const

Grab options.

- std::shared_ptr< [CPHF](#) > [cphf_solver](#) () const

Grab the [CPHF](#) object.

Static Public Member Functions

- static std::shared_ptr< [GenEffParFactory](#) > [build](#) (const std::string &type, std::shared_ptr< psi::Wavefunction > [wfn](#), psi::Options &opt)

Build Density Matrix Susceptibility Generalized Factory.

Protected Member Functions

- virtual double [random_double](#) ()

Draw random number.

- virtual std::shared_ptr< psi::Vector > [draw_random_point](#) ()

Draw random point in 3D space, excluding the vdW region.

- virtual bool [is_in_vdWsphere](#) (double x, double y, double z) const

Is the point inside a vdW region?

Protected Attributes

- std::shared_ptr< psi::Wavefunction > [wfn_](#)

Wavefunction.

- psi::Options & [options_](#)

Psi4 Options.

- std::default_random_engine [randomNumberGenerator_](#)

Random number generators.

- std::uniform_real_distribution< double > [randomDistribution_](#)

- std::shared_ptr< psi::Matrix > [excludeSpheres_](#)

Matrix with vdW sphere information.

- std::map< std::string, double > [vdwRadius_](#)

Map with vdW radii.

- double [cx_](#)

Centre-of-mass coordinates.

- double [cy_](#)

- double [cz_](#)

- double [radius_](#)

Radius of padding sphere around the molecule.

- `const int nbf_`
Number of basis functions.
- `std::shared_ptr< CPHF > cphfSolver_`
The CPHF object.
- `std::shared_ptr< oepdev::GenEffParFactory > abInitioPolarizationSusceptibilitiesFactory_`
Ab initio polarization susceptibility factory.

18.57.1 Detailed Description

Describes the GEFP fragment that is in principle designed to work at correlated levels of theory.

18.57.2 Member Function Documentation

18.57.2.1 build()

```
std::shared_ptr< oepdev::GenEffParFactory > oepdev::GenEffParFactory::build
(
    const std::string & type,
    std::shared_ptr< psi::Wavefunction > wfn,
    psi::Options & opt ) [static]
```

Parameters

<i>type</i>	- Type of factory
<i>wfn</i>	- Psi4 wavefunction
<i>opt</i>	- Psi4 options

Available factory types:

- POLARIZATION - creates the polarization generalized effective fragment parameters' factory Factory subtype is specified in Psi4 options (input file).

Note

Useful options:

- POLARIZATION factory type:
 - DMATPOL_TRAINING_MODE - training mode. Default: EFIELD
 - DMATPOL_NSAMPLES - number of random samples (field or test charges sets). Default: 30
 - DMATPOL_FIELD_SCALE - electric field scale factor (relevant if training mode is EFIELD). Default: 0.01 [au]

- DMATPOL_NTEST_CHARGE - number of test charges per sample (relevant if training mode is CHARGES). Default: 1
- DMATPOL_TEST_CHARGE - test charge value (relevant if training mode is CHARGES). Default: 0.001 [au]
- DMATPOL_FIELD_RANK - electric field rank. Default: 1
- DMATPOL_GRADIENT_RANK - electric field gradient rank. Default: 0
- DMATPOL_TEST_FIELD_X - test electric field in X direction. Default: 0.000 [au]
- DMATPOL_TEST_FIELD_Y - test electric field in Y direction. Default: 0.000 [au]
- DMATPOL_TEST_FIELD_Z - test electric field in Z direction. Default: 0.008 [au]
- DMATPOL_OUT_STATS - output file name for statistical evaluation results. Default: dmatpol.stats.dat
- DMATPOL_DO_AB_INITIO - compute ab initio susceptibilities and evaluate statistics for it. Default: false
- DMATPOL_OUT_STATS_AB_INITIO - output file name for statistical evaluation results of ab initio model. Default: dmatpol.stats.abinitio.dat

The documentation for this class was generated from the following files:

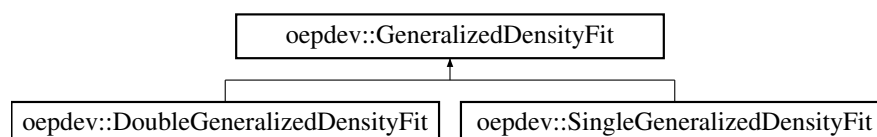
- oepdev/libgefp/gefp.h
- oepdev/libgefp/gefp.cc

18.58 oepdev::GeneralizedDensityFit Class Reference

Generalized Density Fitting Scheme. Abstract Base.

```
#include <oep_gdf.h>
```

Inheritance diagram for oepdev::GeneralizedDensityFit:



Public Member Functions

- [GeneralizedDensityFit](#) ()
Constructor. Initializes the pointers.
- virtual [~GeneralizedDensityFit](#) ()
Destructor.
- virtual std::shared_ptr< psi::Matrix > [compute](#) (void)=0

Perform the generalized density fit.

- `std::shared_ptr< psi::Matrix > G (void) const`

Extract the G_{ξ_i} coefficients.

Static Public Member Functions

- `static std::shared_ptr< GeneralizedDensityFit > build (std::shared_ptr< psi::BasisSet > bs_auxiliary, std::shared_ptr< psi::Matrix > v_vector)`

Factory for Single GDF Computer.

- `static std::shared_ptr< GeneralizedDensityFit > build (std::shared_ptr< psi::BasisSet > bs_auxiliary, std::shared_ptr< psi::BasisSet > bs_intermediate, std::shared_ptr< psi::Matrix > v_vector)`

Factory for Double GDF Computer.

Protected Member Functions

- `void invert_matrix (std::shared_ptr< psi::Matrix > &M)`
Invert a square matrix and check if the inverse is acceptable.

Protected Attributes

- `std::shared_ptr< psi::Matrix > G_`
The OEP coefficients G_{ξ_i} .
- `std::shared_ptr< psi::Matrix > H_`
The intermediate DF coefficients for $\hat{v}|i\rangle$.
- `std::shared_ptr< psi::Matrix > V_`
The V matrix ($\xi|\hat{v}i$).
- `int n_a_`
Number of auxiliary basis set functions.
- `int n_i_`
Number of intermediate basis set functions.
- `int n_o_`
Number of OEP's.
- `std::shared_ptr< psi::BasisSet > bs_a_`
Basis set: auxiliary.
- `std::shared_ptr< psi::BasisSet > bs_i_`
Basis set: intermediate.
- `std::shared_ptr< oepdev::IntegralFactory > ints_aa_`
Integral factory: aux - aux.
- `std::shared_ptr< oepdev::IntegralFactory > ints_ai_`
Integral factory: aux - int.

- `std::shared_ptr< oepdev::IntegralFactory > ints_ii_`
Integral factory: int - int.

18.58.1 Detailed Description

Performs the following map:

$$\hat{v}|i\rangle \cong \sum_{\eta} G_{\eta i} |\eta\rangle$$

where \hat{v} is the effective one-electron potential (OEP) operator, $|i\rangle$ is an arbitrary state vector and $|\eta\rangle$ is an auxiliary basis vector. The coefficients $G_{\eta i}$ are stored and define the OEP acting on the state i . The mapping onto the auxiliary space can be done in two ways:

- **Single Density Fit.** [This method](#) requires the auxiliary basis set to be nearly complete.
- **Double Density Fit.** [This method](#) can be used to arbitrary auxiliary basis sets.

18.58.2 Member Function Documentation

18.58.2.1 `build()` [1/2]

```
std::shared_ptr< GeneralizedDensityFit > GeneralizedDensityFit::build (
    std::shared_ptr< psi::BasisSet > bs_auxiliary,
    std::shared_ptr< psi::Matrix > v_vector ) [static]
```

Parameters

<i>bs_auxiliary</i>	- auxiliary basis set
<i>v_vector</i>	- the matrix with $V_{\xi i}$ elements

Returns

Generalized Density Fit Computer.

18.58.2.2 `build()` [2/2]

```
std::shared_ptr< GeneralizedDensityFit > GeneralizedDensityFit::build (
    std::shared_ptr< psi::BasisSet > bs_auxiliary,
    std::shared_ptr< psi::BasisSet > bs_intermediate,
    std::shared_ptr< psi::Matrix > v_vector ) [static]
```

Parameters

<i>bs_auxiliary</i>	- auxiliary basis set
<i>bs_intermediate</i>	- intermediate basis set
<i>v_vector</i>	- the matrix with V_{ei} elements

Returns

Generalized Density Fit Computer.

18.58.2.3 compute()

```
std::shared_ptr< psi::Matrix > GeneralizedDensityFit::compute (
    void ) [pure virtual]
```

Returns

The OEP coefficients $G_{\xi i}$

Implemented in [oepdev::DoubleGeneralizedDensityFit](#), and [oepdev::SingleGeneralizedDensityFit](#).

The documentation for this class was generated from the following files:

- [oepdev/liboep/oep_gdf.h](#)
- [oepdev/liboep/oep_gdf.cc](#)

18.59 oepdev::GeneralizedPolarGEFactory Class Reference

Polarization GEFP Factory with Least-Squares Parameterization.

```
#include <gefp.h>
```

Inheritance diagram for oepdev::GeneralizedPolarGEFactory:



Classes

- struct [StatisticalSet](#)

A structure to handle statistical data.

Public Member Functions

- [GeneralizedPolarGEFactory](#) (std::shared_ptr< psi::Wavefunction > [wfn](#), psi::Options &opt)
Construct from Psi4 wavefunction and options.
- virtual [~GeneralizedPolarGEFactory](#) ()
Destruct.
- virtual std::shared_ptr< [GenEffPar](#) > [compute](#) (void)
Perform Least-Squares Fit.
- bool [has_dipole_polarizability](#) () const
Dipole Polarizability (interacting with \mathbf{F})
- bool [has_dipole_dipole_hyperpolarizability](#) () const
Dipole-Dipole Hyperpolarizability (interacting with \mathbf{F}^2)
- bool [has_quadrupole_polarizability](#) () const
Quadrupole Polarizability (interacting with $\nabla \otimes \mathbf{F}$)
- bool [has_ab_initio_dipole_polarizability](#) () const
Ab Initio Dipole Polarizability (interacting with \mathbf{F})
- double [Zinit](#) () const
Grab initial summaric Z value.
- double [Z](#) () const
Grab final summaric Z value.

Protected Member Functions

- void [allocate](#) (void)
Allocate memory.
- void [invert_hessian](#) (void)
Invert Hessian (do also the identity test)
- void [compute_electric_field_sums](#) (void)
Compute electric field sum set.
- void [compute_electric_field_gradient_sums](#) (void)
Compute electric field gradient sum set.
- void [compute_statistics](#) (void)
Run the statistical evaluation of results.
- void [set_distributed_centres](#) (void)
Set the distributed centres.
- void [compute_parameters](#) (void)
Compute the parameters.
- void [fit](#) (void)
Perform least-squares fit.
- void [compute_ab_initio](#) (void)

- Compute ab initio parameters.*

 - void [save](#) (int i, int j)

Save susceptibility tensors associated with the i-th and j-th basis set function.
- virtual void [compute_samples](#) (void)=0

Compute samples of density matrices and select electric field distributions.
- virtual void [compute_gradient](#) (int i, int j)=0

Compute Gradient vector associated with the i-th and j-th basis set function.
- virtual void [compute_hessian](#) (void)=0

Compute Hessian matrix (independent on the parameters)

Protected Attributes

- int [nBlocks_](#)
- Number of parameter blocks.*
- int [nSites_](#)
- Number of distributed sites.*
- int [nSitesAbInitio_](#)
- Number of distributed sites of Ab Initio model (FF - single site (com); distributed: LMO sites)*
- int [nParameters_](#)
- Dimensionality of entire parameter space.*
- std::vector< int > [nParametersBlock_](#)
- Dimensionality of parameter space per block.*
- const int [nSamples_](#)
- Number of statistical samples.*
- const double [symmetryNumber_](#) [6]
- Symmetry number for matrix susceptibilities.*
- std::shared_ptr< psi::Matrix > [Gradient_](#)
- Gradient.*
- std::shared_ptr< psi::Matrix > [Hessian_](#)
- Hessian.*
- std::shared_ptr< psi::Matrix > [Parameters_](#)
- Parameters.*
- std::shared_ptr< oepdev::GenEffPar > [PolarizationSusceptibilities_](#)
- Density Matrix Susceptibility Tensors Object.*
- std::shared_ptr< oepdev::GenEffPar > [abInitioPolarizationSusceptibilities_](#)
- Density Matrix Susceptibility Tensors Object for Ab Initio Model.*
- bool [hasDipolePolarizability_](#)
- Has Dipole Polarizability?*
- bool [hasDipoleDipoleHyperpolarizability_](#)
- Has Dipole-Dipole Hyperpolarizability?*

- bool [hasQuadrupolePolarizability_](#)
Has Quadrupole Polarizability?
- bool [hasAbInitioDipolePolarizability_](#)
Has Ab Initio Dipole Polarizability?
- [StatisticalSet](#) [referenceStatisticalSet_](#)
Reference statistical data.
- [StatisticalSet](#) [referenceDpolStatisticalSet_](#)
Multipole reference statistical data.
- [StatisticalSet](#) [modelStatisticalSet_](#)
Model statistical data.
- [StatisticalSet](#) [abInitioModelStatisticalSet_](#)
Ab Initio Model statistical data.
- `std::vector< std::shared_ptr< psi::Matrix > >` [VMatrixSet_](#)
Potential matrix set.
- `std::vector< std::vector< std::shared_ptr< Vector > > >` [electricFieldSet_](#)
Electric field set.
- `std::vector< std::vector< std::shared_ptr< Matrix > > >` [electricFieldGradientSet_](#)
Electric field gradient set.
- `std::vector< std::vector< double > >` [electricFieldSumSet_](#)
Electric field sum set.
- `std::vector< std::vector< std::shared_ptr< psi::Vector > > >` [electricFieldGradientSumSet_](#)
Electric field gradient sum set.
- `std::vector< std::vector< std::shared_ptr< Vector > > >` [abInitioModelElectricFieldSet_](#)
Electric field set for Ab Initio Model (LMO-distributed)
- const double [mField_](#)
Level shifters for Hessian blocks.
- double [Zinit_](#)
Initial summaric Z value.
- double [Z_](#)
Final summaric Z value.
- `std::shared_ptr< psi::JK >` [jk_](#)
Computer of generalized JK objects.

Additional Inherited Members

18.59.1 Detailed Description

Implements a general class of methods for the density matrix susceptibility tensors represented by:

$$\delta D_{\alpha\beta} = \sum_i \left\{ \mathbf{B}_{i;\alpha\beta}^{(10)} \cdot \mathbf{F}(\mathbf{r}_i) + \mathbf{B}_{i;\alpha\beta}^{(20)} : \mathbf{F}(\mathbf{r}_i) \otimes \mathbf{F}(\mathbf{r}_i) + \mathbf{B}_{i;\alpha\beta}^{(01)} : \nabla_i \otimes \mathbf{F}(\mathbf{r}_i) + \dots \right\}$$

where:

- $\mathbf{B}_{i;\alpha\beta}^{(10)}$ is the density matrix dipole polarizability
- $\mathbf{B}_{i;\alpha\beta}^{(20)}$ is the density matrix dipole-dipole hyperpolarizability
- $\mathbf{B}_{i;\alpha\beta}^{(01)}$ is the density matrix quadrupole polarizability

all defined for the generalized distributed site at \mathbf{r}_i .

Available models:

1. Training against uniform electric fields

- [oepdev::LinearUniformEFieldPolarGEFactory](#) - linear with respect to electric field
- [oepdev::QuadraticUniformEFieldPolarGEFactory](#) - quadratic with respect to electric field

2. Training against non-uniform electric fields

- [oepdev::LinearNonUniformEFieldPolarGEFactory](#) - linear with respect to electric field, distributed site model
- [oepdev::QuadraticNonUniformEFieldPolarGEFactory](#) - quadratic with respect to electric field, distributed site model
- [oepdev::LinearGradientNonUniformEFieldPolarGEFactory](#) - linear with respect to electric field and linear with respect to electric field gradient, distributed site model. This model does not function now.
- [oepdev::QuadraticGradientNonUniformEFieldPolarGEFactory](#) - linear with respect to electric field and linear with respect to electric field gradient, distributed site model. This model does not function now.

For the non-linear field training, a set of point charges in each training sample is assumed. Distributed models use atomic centers as expansion points.

Determination of the generalized susceptibilities

Let $\{\mathbf{F}^{(1)}(\mathbf{r}), \mathbf{F}^{(2)}(\mathbf{r}), \dots, \mathbf{F}^{(N)}(\mathbf{r}), \dots\}$ be a set of N_{\max} distinct and randomly sampled spatial distributions of electric field. It is assumed that the exact difference one-particle density matrices (with respect to the unperturbed state) defined as

$$\delta\overline{\mathbf{D}}^{(N)} \equiv \overline{\mathbf{D}}^{(N)} - \overline{\mathbf{D}}^{(0)}$$

are known for each sample (overline symbolizes the exact estimate). Now, for each pair of the AO indices the following parameterization is constructed:

$$\delta D^{(N)} = \sum_i^M \left\{ \sum_u^{x,y,z} s_{iu}^{[1]} F_{iu}^{(N)} + \sum_u^{x,y,z} \sum_{w < u} r_{uw} s_{iww}^{[2]} F_{iu}^{(N)} F_{iw}^{(N)} + \dots \right\}$$

(the Greek subscripts were omitted here for notational simplicity). In the above equation, $B_u^{(i;1)} = s_{iu}^{[1]}$ and $B_{uw}^{(i;2)} = r_{uw}s_{iww}^{[2]}$, where r_{uw} is the symmetry factor equal to 1 for diagonal elements and 2 for off-diagonal elements of $B_{uw}^{(i;2)}$. The multiple parameter blocks ($s^{[1]}$, $s^{[2]}$ and so on) appear in the first power, allowing for linear least-squares regression. The square bracket superscripts denote the block of the parameter space.

To determine the optimum set, $\mathbf{s} = (s^{[1]} \ s^{[2]} \ \dots)^T$, a loss function Z that is subject to the least-squares minimization, is defined as

$$Z(\mathbf{s}) = \sum_N^{N_{\max}} \left(\delta D^{(N)} - \delta \bar{D}^{(N)} \right)^2.$$

The Hessian of Z computed with respect to the parameters is parameter-independent (constant) and generally non-singular as long as the electric fields on all distributed sites are different. Therefore, the exact solution for the optimal parameters is given by the Newton equation

$$\mathbf{s} = -\mathbf{H}^{-1} \cdot \mathbf{g},$$

where \mathbf{g} and \mathbf{H} are the gradient vector and the Hessian matrix, respectively. Note that in this case the dimensions of parameter space for the block 1 and 2 are equal to $3M$ and $6M$, respectively. The explicit forms of the gradient and Hessian up to second-order are given in the next section.

Explicit Formulae for Gradient and Hessian Blocks in Linear Regression DMS Model

The gradient vector \mathbf{g} and the Hessian matrix \mathbf{H} are built from blocks associated with a particular type of parameters, i.e.,

$$\mathbf{g} = \begin{pmatrix} \mathbf{g}^{[1]} \\ \mathbf{g}^{[2]} \end{pmatrix}, \quad \mathbf{H} = \begin{pmatrix} \mathbf{H}^{[11]} & \mathbf{H}^{[12]} \\ \mathbf{H}^{[21]} & \mathbf{H}^{[22]} \end{pmatrix},$$

where the block indices 1 and 2 correspond to the first- and second-order susceptibilities, respectively. Note that the second derivatives of $\delta D^{(N)}$ with respect to the adjustable parameters vanish due to the linear functional form of the parameterization formula given in the previous section. Thus, the gradient element of the r -th block and Hessian element of the (rs) -th block read

$$g^{[r]} \equiv \frac{\partial Z}{\partial s^{[r]}} = -2 \sum_N \bar{\delta D}^{(N)} \frac{\partial [\delta D^{(N)}]}{\partial s^{[r]}},$$

$$H^{[rs]} \equiv \frac{\partial^2 Z}{\partial s^{[r]} \partial s^{[s]}} = 2 \sum_N \frac{\partial [\delta D^{(N)}]}{\partial s^{[r]}} \frac{\partial [\delta D^{(N)}]}{\partial s^{[s]}}.$$

The explicit formulae for the gradient are

$$g_{ku}^{[1]} = -2 \sum_N \bar{\delta D}^{(N)} F_{ku}^{(N)},$$

$$g_{kuw}^{[2]} = -2 r_{uw} \sum_N \bar{\delta D}^{(N)} F_{ku}^{(N)} F_{kw}^{(N)}.$$

The Hessian subsequently follows to be %

$$H_{ku,lw}^{[11]} = 2 \sum_N F_{ku}^{(N)} F_{lw}^{(N)} ,$$

$$H_{ku,lu'w'}^{[12]} = 2 r_{u'w'} \sum_N F_{ku}^{(N)} F_{lu'}^{(N)} F_{lw'}^{(N)} ,$$

$$H_{kuw,lu'w'}^{[22]} = 2 r_{uw} r_{u'w'} \sum_N F_{ku}^{(N)} F_{kw}^{(N)} F_{lu'}^{(N)} F_{lw'}^{(N)} .$$

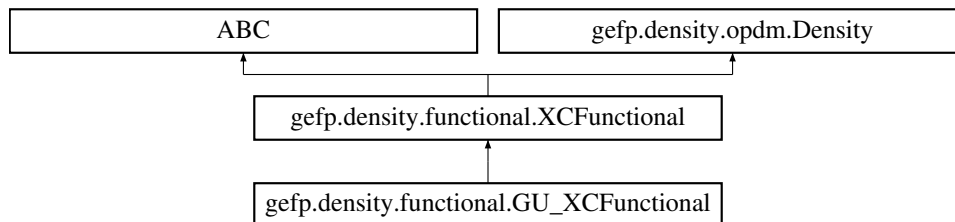
Note that due to the symmetry of the Hessian matrix, the block 21 is a transpose of the block 12. The composite indices ku and kuw are constructed from the distributed site index k and the appropriate symmetry-adapted ($w < u$) Cartesian component of a particular DMS tensor: u for the first-order, and uw for the second-order susceptibility tensor, respectively. The method described above can be easily extended to third and higher orders.

The documentation for this class was generated from the following files:

- oepdev/libgefp/gefp.h
- oepdev/libgefp/gefp_polar_base.cc

18.60 gefp.density.functional.GU_XCFunctional Class Reference

Inheritance diagram for gefp.density.functional.GU_XCFunctional:



Public Member Functions

- def **__init__** (self)
- def **abbr** (self)
- def **gradient_P_approximate_old** (self, x)
- def **gradient_P_approximate** (self, x)

Static Public Member Functions

- def **name** ()
- def **fij** (n)
- def **fij_1** (n, m)

Additional Inherited Members

18.60.1 Detailed Description

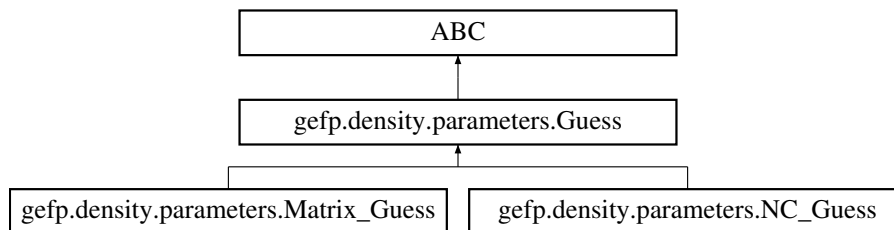
The Goedecker-Urmigar Exchange-Correlation Functional.

The documentation for this class was generated from the following file:

- `gefp/gefp/density/functional.py`

18.61 gefp.density.parameters.Guess Class Reference

Inheritance diagram for `gefp.density.parameters.Guess`:



Public Member Functions

- `def __init__ (self, n=None, c=None, matrix=None)`
- `def create (cls, n=None, c=None, matrix=None, t='matrix')`
- `def update (self, S=None, C=None)`
- `def matrix (self)`
- `def copy (self)`
- `def pack (self)`
- `def unpack (self)`
- `def __add__ (self, other)`
- `def __sub__ (self, other)`
- `def __rmul__ (self, other)`

18.61.1 Detailed Description

\

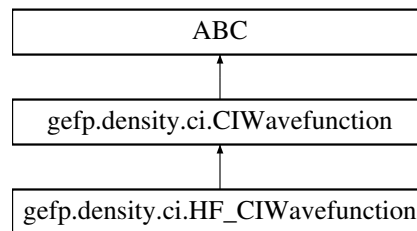
Container for handling density matrix guesses for DMFT calculations.
Contains functionalities for working with occupation numbers,
natural orbitals and density matrices.

The documentation for this class was generated from the following file:

- `gefp/gefp/density/parameters.py`

18.62 gefp.density.ci.HF_CIWavefunction Class Reference

Inheritance diagram for gefp.density.ci.HF_CIWavefunction:



Public Member Functions

- `def __init__ (self, ref.wfn, E)`
- `def make_ci_l (self)`

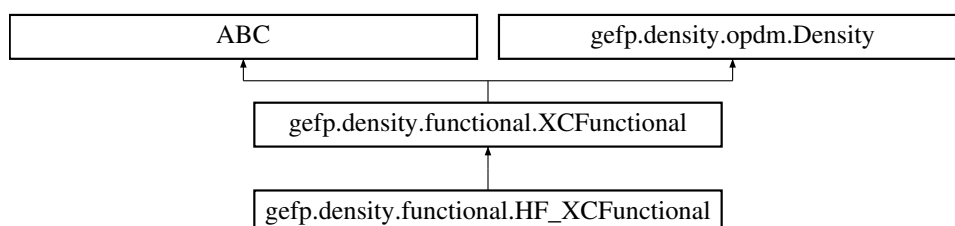
Additional Inherited Members

The documentation for this class was generated from the following file:

- `gefp/gefp/density/ci.py`

18.63 gefp.density.functional.HF_XCFunctional Class Reference

Inheritance diagram for gefp.density.functional.HF_XCFunctional:



Public Member Functions

- `def __init__ (self)`
- `def abbr (self)`
- `def energy_D (self, x, mode='scf-mo')`
- `def gradient_D (self, x)`
- `def gradient_nc (self, x)`

Static Public Member Functions

- def **name** ()
- def **fij** (n)
- def **fij_1** (n, m)

Additional Inherited Members

18.63.1 Detailed Description

The Hartree-Fock Exchange-Correlation Functional

The documentation for this class was generated from the following file:

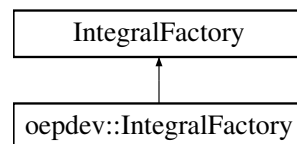
- gefp/gefp/density/functional.py

18.64 oepdev::IntegralFactory Class Reference

Extended [IntegralFactory](#) for computing integrals.

```
#include <integral.h>
```

Inheritance diagram for oepdev::IntegralFactory:



Public Member Functions

- [IntegralFactory](#) (std::shared_ptr< psi::BasisSet > bs1, std::shared_ptr< psi::BasisSet > bs2, std::shared_ptr< psi::BasisSet > bs3, std::shared_ptr< psi::BasisSet > bs4)
Initialize integral factory given a BasisSet for each center. Becomes (bs1 bs2 | bs3 bs4).
- [IntegralFactory](#) (std::shared_ptr< psi::BasisSet > bs1, std::shared_ptr< psi::BasisSet > bs2)
Initialize integral factory given a BasisSet for two centres. Becomes (bs1 bs2 | bs1 bs2).
- [IntegralFactory](#) (std::shared_ptr< psi::BasisSet > bs1)
Initialize integral factory given a BasisSet for two centres. Becomes (bs1 bs1 | bs1 bs1).
- virtual [~IntegralFactory](#) ()
Destructor.
- virtual [oepdev::TwoBodyAOInt * eri_1_1](#) (int deriv=0, bool use_shell_pairs=false)
Returns an [ERI_1_1](#) integral object.

- virtual `oepdev::TwoBodyAOInt * eri_2_1` (int deriv=0, bool use_shell_pairs=false)
Returns an *ERI_2_1* integral object.
- virtual `oepdev::TwoBodyAOInt * eri_2_2` (int deriv=0, bool use_shell_pairs=false)
Returns an *ERI_2_2* integral object.
- virtual `oepdev::TwoBodyAOInt * eri_3_1` (int deriv=0, bool use_shell_pairs=false)
Returns an *ERI_3_1* integral object.

18.64.1 Detailed Description

In addition to integrals available in Psi4, `oepdev::IntegralFactory` enables to compute also:

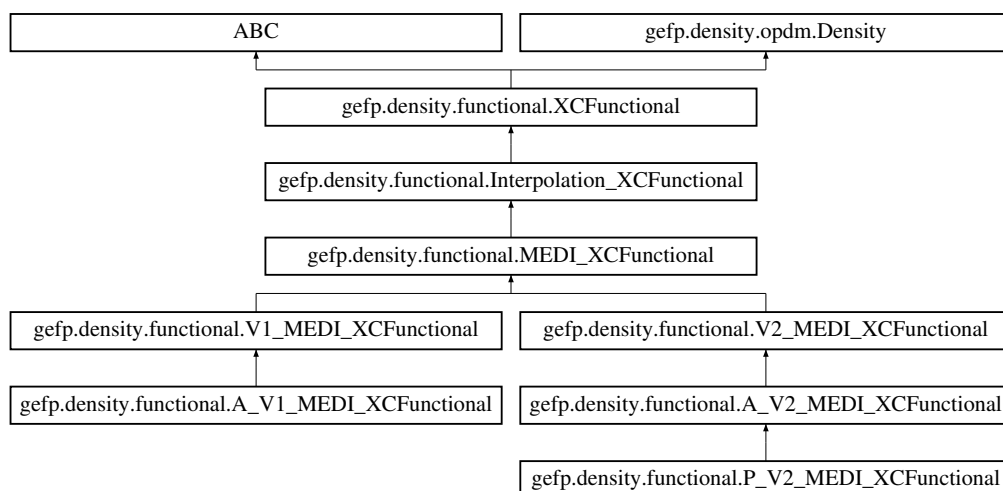
- OEI's:
 - none at that moment
- ERI's:
 - integrals of type (a|b) - `oepdev::ERI_1_1`
 - integrals of type (ab|c) - `oepdev::ERI_2_1`
 - integrals of type (abc|d) - `oepdev::ERI_3_1`
 - integrals of type (ab|cd) - `oepdev::ERI_2_2` (also in Psi4 as `psi::ERI`)

The documentation for this class was generated from the following files:

- `oepdev/libpsi/integral.h`
- `oepdev/libpsi/integral.cc`

18.65 `gefp.density.functional.Interpolation_XCFunctional` Class Reference

Inheritance diagram for `gefp.density.functional.Interpolation_XCFunctional`:



Public Member Functions

- def **__init__** (self, coeff, kmax=10)
- def **compute_a0** (self, n)
- def **compute_ak** (self, k, t)
- def **fij** (self, n)
- def **gradient_P** (self, x)

Additional Inherited Members

18.65.1 Detailed Description

The New Class of Exchange-Correlation Functionals: Interpolation Functionals

They differ in the model for the interpolation decay.
Each functional has to provide its own coefficients,
handled by 'coeff' dictionary in the constructor.

Eg.: coeff = {'coefficient_name': coefficient_object}

The documentation for this class was generated from the following file:

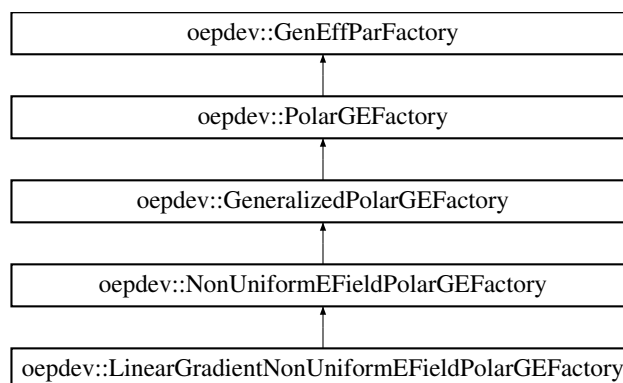
- gefp/gefp/density/functional.py

18.66 oepdev::LinearGradientNonUniformEFieldPolarGEFactory Class Reference

Polarization GEFP Factory with Least-Squares Parameterization.

```
#include <gefp.h>
```

Inheritance diagram for oepdev::LinearGradientNonUniformEFieldPolarGEFactory:



Public Member Functions

- **LinearGradientNonUniformEFieldPolarGEFactory** (std::shared_ptr< psi::Wavefunction > [wfn](#), psi::Options &opt)
- void [compute_gradient](#) (int i, int j)
Compute Gradient vector associated with the i-th and j-th basis set function.
- void [compute_hessian](#) (void)
Compute Hessian matrix (independent on the parameters)

Additional Inherited Members

18.66.1 Detailed Description

Implements the generalized density matrix susceptibility model of the form

$$\delta D_{\alpha\beta} \approx \sum_i \left\{ \mathbf{B}_{i;\alpha\beta}^{(10)} \cdot \mathbf{F}(\mathbf{r}_i) + \mathbf{B}_{i;\alpha\beta}^{(01)} : \nabla_i \otimes \mathbf{F}(\mathbf{r}_i) \right\}$$

where:

- $\mathbf{B}_{i;\alpha\beta}^{(10)}$ is the density matrix dipole polarizability
- $\mathbf{B}_{i;\alpha\beta}^{(01)}$ is the density matrix quadrupole polarizability all defined for the distributed site at \mathbf{r}_i .

Note

This model is not available now and probably will be deprecated in the future.

The documentation for this class was generated from the following files:

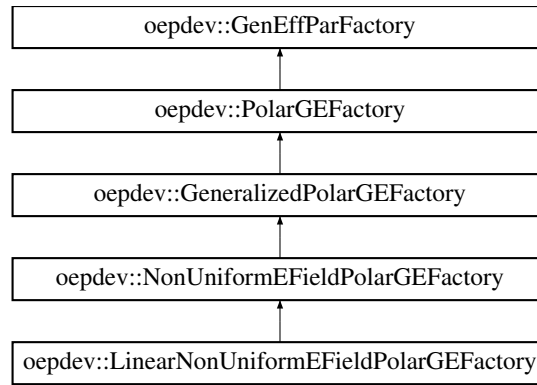
- oepdev/libgefp/[gefp.h](#)
- oepdev/libgefp/gefp_polar_nonuniform_field_1_grad_1.cc

18.67 oepdev::LinearNonUniformEFieldPolarGEFactory Class Reference

Polarization GEFP Factory with Least-Squares Parameterization.

```
#include <gefp.h>
```

Inheritance diagram for oepdev::LinearNonUniformEFieldPolarGEFactory:



Public Member Functions

- **LinearNonUniformEFieldPolarGEFactory** (std::shared_ptr< psi::Wavefunction > wfn, psi::Options &opt)
- void **compute_gradient** (int i, int j)
Compute Gradient vector associated with the i-th and j-th basis set function.
- void **compute_hessian** (void)
Compute Hessian matrix (independent on the parameters)

Additional Inherited Members

18.67.1 Detailed Description

Implements the generalized density matrix susceptibility model of the form

$$\delta D_{\alpha\beta} \approx \sum_i \mathbf{B}_{i;\alpha\beta}^{(10)} \cdot \mathbf{F}(\mathbf{r}_i)$$

where:

- $\mathbf{B}_{i;\alpha\beta}^{(10)}$ is the density matrix dipole polarizability defined for the distributed site at \mathbf{r}_i .

The documentation for this class was generated from the following files:

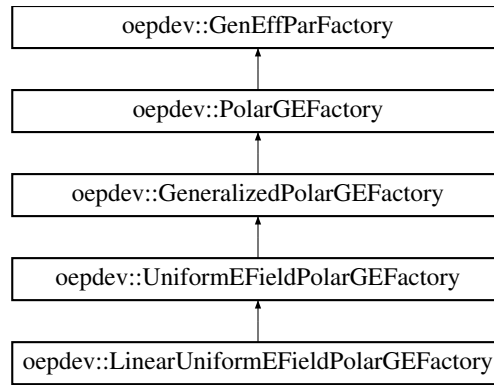
- oepdev/libgefp/gefp.h
- oepdev/libgefp/gefp_polar_nonuniform_field_1.cc

18.68 oepdev::LinearUniformEFieldPolarGEFactory Class Reference

Polarization GEFP Factory with Least-Squares Parameterization.

```
#include <gefp.h>
```

Inheritance diagram for oepdev::LinearUniformEFieldPolarGEFactory:



Public Member Functions

- **LinearUniformEFieldPolarGEFactory** (std::shared_ptr< psi::Wavefunction > [wfn](#), psi::Options &opt)
- void [compute_gradient](#) (int i, int j)
Compute Gradient vector associated with the i-th and j-th basis set function.
- void [compute_hessian](#) (void)
Compute Hessian matrix (independent on the parameters)

Additional Inherited Members

18.68.1 Detailed Description

Implements the generalized density matrix susceptibility model of the form

$$\delta D_{\alpha\beta} \approx \mathbf{B}_{\alpha\beta}^{(10)} \cdot \mathbf{F}$$

where:

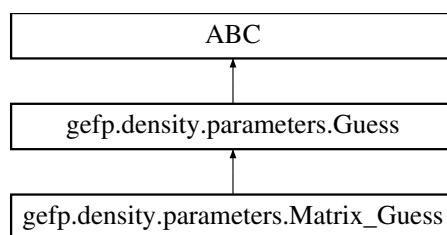
- $\mathbf{B}_{\alpha\beta}^{(10)}$ is the density matrix dipole polarizability

The documentation for this class was generated from the following files:

- oepdev/libgefp/[gefp.h](#)
- oepdev/libgefp/gefp_polar_uniform_field_1.cc

18.69 gefp.density.parameters.Matrix_Guess Class Reference

Inheritance diagram for gefp.density.parameters.Matrix_Guess:



Public Member Functions

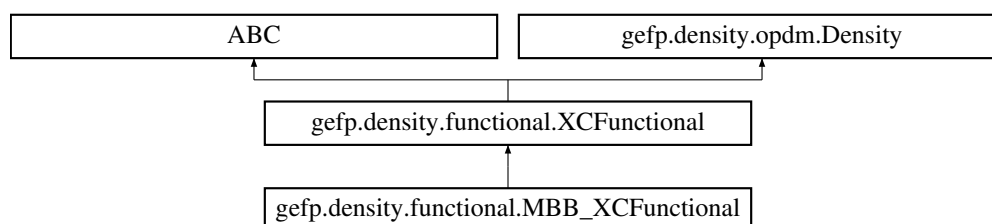
- def **__init__** (self, n=None, c=None, matrix=None)
- def **pack** (self)

The documentation for this class was generated from the following file:

- gefp/gefp/density/parameters.py

18.70 gefp.density.functional.MBB_XCFunctional Class Reference

Inheritance diagram for gefp.density.functional.MBB_XCFunctional:



Public Member Functions

- def **__init__** (self)
- def **abbr** (self)
- def **energy_P** (self, x)
- def **energy_pc** (self, x)
- def **gradient_P** (self, x)
- def **gradient_pc** (self, x)
- def **energy_D** (self, x, mode='scf-mo')

Static Public Member Functions

- def **name** ()
- def **fij** (n)

Additional Inherited Members

18.70.1 Detailed Description

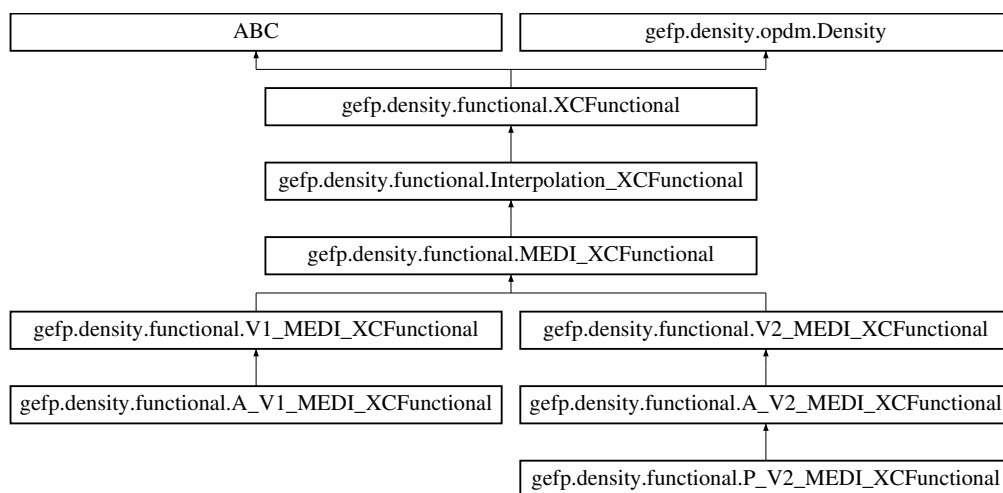
The Muller-Buijse-Baerends Exchange-Correlation Functional.

The documentation for this class was generated from the following file:

- `gefp/gefp/density/functional.py`

18.71 `gefp.density.functional.MEDI_XCFunctional` Class Reference

Inheritance diagram for `gefp.density.functional.MEDI_XCFunctional`:



Public Member Functions

- `def __init__ (self, coeff, kmax)`
- `def compute_t (self, n=None, c=None)`
- `def fij (self, n)`

Additional Inherited Members

18.71.1 Detailed Description

The New Class of Exchange-Correlation Functionals:
Interpolation Functionals with Monotonous Exponential Decay.

The decay in the interpolates is modelled by the monotonous decay

$$a_k = a_0 \exp(k \ln\{1 - a_0\})$$

Functional coefficients:

- o 'a0' - first term in the interpolates coefficient
(the one that multiplies MBB functional term - for k=0).
Parameter 'a0' has to be between 0.0 and 1.0. The smaller 'a0',
the more terms need to be taken (larger kmax).

The documentation for this class was generated from the following file:

- `gefp/gefp/density/functional.py`

18.72 oepdev::MultipoleConvergence Class Reference

Multipole Convergence.

```
#include <dmtip.h>
```

Public Types

- enum [ConvergenceLevel](#) {
 R1, R2, R3, R4,
 R5 }
- enum [Property](#) { **Energy, Potential** }

Public Member Functions

- [MultipoleConvergence](#) (std::shared_ptr< [DMTPole](#) > dmtip1, std::shared_ptr< [DMTPole](#) > dmtip2, [ConvergenceLevel](#) max_clevel=R5)
Construct from two shared [DMTPole](#) objects.
- virtual [~MultipoleConvergence](#) ()
Destructor.
- void [compute](#) ([Property](#) property=Energy)
- std::shared_ptr< psi::Matrix > [level](#) ([ConvergenceLevel](#) clevel=R5)

Protected Member Functions

- void [compute_energy](#) ()
Compute the generalized energy.
- void [compute_potential](#) ()
Void compute the generalized potential.

Protected Attributes

- [ConvergenceLevel max_clevel_](#)
Maximum allowed convergence level.
- `std::shared_ptr< DMTPole > dmtip_1_`
First DMTP set.
- `std::shared_ptr< DMTPole > dmtip_2_`
Second DMTP set.
- `std::map< std::string, std::shared_ptr< psi::Matrix > > convergenceList_`
Dictionary of available convergence level results.

18.72.1 Detailed Description

Handles the convergence of the distributed multipole expansions up to hexadecapole. Takes shared pointers to existing [DMTPole](#) objects and computes the generalized property:

- energy
- potential from the DMTP sets. The results are stored in matrix of size (N1, N2) where N1 and N2 are equal to the number of DMTP's in a set described by according [DMTPole](#) object given.

Note

Useful options:

- DMTP_CONVER - level of multipole series convergence (available: R1, R2, R3, R4 and R5). Default: R5.

See also

[DMTPole](#)

18.72.2 Member Enumeration Documentation

18.72.2.1 ConvergenceLevel

enum [oepdev::MultipoleConvergence::ConvergenceLevel](#)

Convergence level of the multipole expansion:

Parameters

<i>R1</i>	- qq term
-----------	-----------

Parameters

<i>R2</i>	- qd and sum of the above
<i>R3</i>	- qQ, dd and sum of the above
<i>R4</i>	- qO, dQ and sum of the above
<i>R5</i>	- qH, dO, QQ and sum of the above

18.72.2.2 Property

enum `oepdev::MultipoleConvergence::Property`

Property to be evaluated from interacting DMTP's:

Parameters

<i>Energy</i>	- generalized energy
<i>Potential</i>	- generalized potential

18.72.3 Constructor & Destructor Documentation

18.72.3.1 MultipoleConvergence()

```
oepdev::MultipoleConvergence::MultipoleConvergence (
    std::shared_ptr< DMTPole > dmtpl,
    std::shared_ptr< DMTPole > dmtpl2,
    MultipoleConvergence::ConvergenceLevel max_clevel = R5 )
```

Parameters

<i>dmtpl</i>	- first <code>DMTPole</code> object
<i>dmtpl2</i>	- second <code>DMTPole</code> object
<i>max_clevel</i>	- maximul allowed convergence level

18.72.4 Member Function Documentation

18.72.4.1 compute()

```
void oepdev::MultipoleConvergence::compute (
    MultipoleConvergence::Property property = Energy )
```

Compute the generalized property

Parameters

<i>property</i>	- generalized Property
-----------------	------------------------

18.72.4.2 level()

```
std::shared_ptr< psi::Matrix > oepdev::MultipoleConvergence::level (
    MultipoleConvergence::ConvergenceLevel clevel = R5 )
```

Grab the generalized property at specified level of convergence

Parameters

<i>clevel</i>	- ConvergenceLevel
---------------	--------------------

Returns

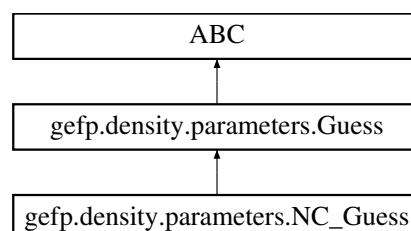
vector of results (each element corresponds to each DMTP pair in a set)

The documentation for this class was generated from the following files:

- oepdev/lib3d/[dmtplib.h](#)
- oepdev/lib3d/dmtp_base.cc

18.73 gefp.density.parameters.NC_Guess Class Reference

Inheritance diagram for gefp.density.parameters.NC_Guess:



Public Member Functions

- `def __init__ (self, n=None, c=None, matrix=None)`
- `def pack (self)`

The documentation for this class was generated from the following file:

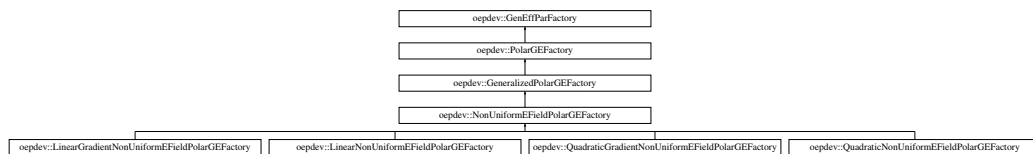
- `gefp/gefp/density/parameters.py`

18.74 oepdev::NonUniformEFieldPolarGEFactory Class Reference

Polarization GEFP Factory with Least-Squares Parameterization.

```
#include <gefp.h>
```

Inheritance diagram for oepdev::NonUniformEFieldPolarGEFactory:



Public Member Functions

- **NonUniformEFieldPolarGEFactory** (`std::shared_ptr< psi::Wavefunction > wfn`, `psi::Options &opt`)
- `void compute_samples (void)`
Compute samples of density matrices and select electric field distributions.
- `virtual void compute_gradient (int i, int j)=0`
Compute Gradient vector associated with the i-th and j-th basis set function.
- `virtual void compute_hessian (void)=0`
Compute Hessian matrix (independent on the parameters)

Additional Inherited Members

18.74.1 Detailed Description

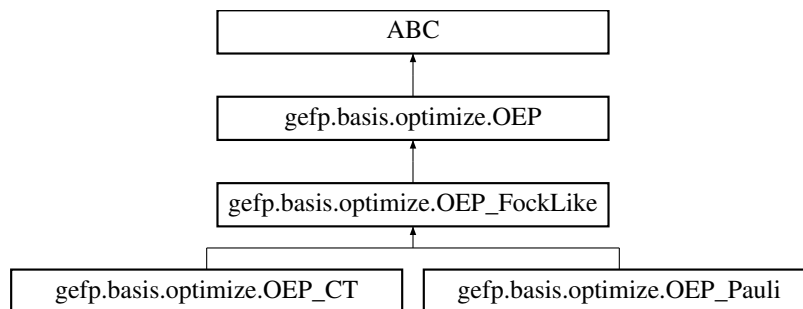
Implements a class of density matrix susceptibility models for parameterization in the non-uniform electric field generated by point charges.

The documentation for this class was generated from the following files:

- `oepdev/libgefp/gefp.h`
- `oepdev/libgefp/gefp_polar_nonuniform_base.cc`

18.75 gefp.basis.optimize.OEP Class Reference

Inheritance diagram for gefp.basis.optimize.OEP:



Public Member Functions

- `def __init__ (self, wfn, dfbasis)`
- `def create (cls, name, wfn, dfbasis)`

Public Attributes

- `wfn`
- `dfbasis`
- `mints`
- `basis_test`
- `basis_prim`
- `basis_aux`
- `eri_pppt`
- `V`

18.75.1 Detailed Description

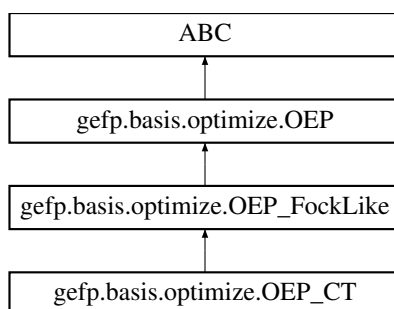
OEP object that defines the V matrix necessary for GDF.

The documentation for this class was generated from the following file:

- `gefp/gefp/basis/optimize.py`

18.76 gefp.basis.optimize.OEP_CT Class Reference

Inheritance diagram for gefp.basis.optimize.OEP_CT:



Public Member Functions

- `def __init__ (self, wfn, dfbasis)`

Additional Inherited Members

18.76.1 Detailed Description

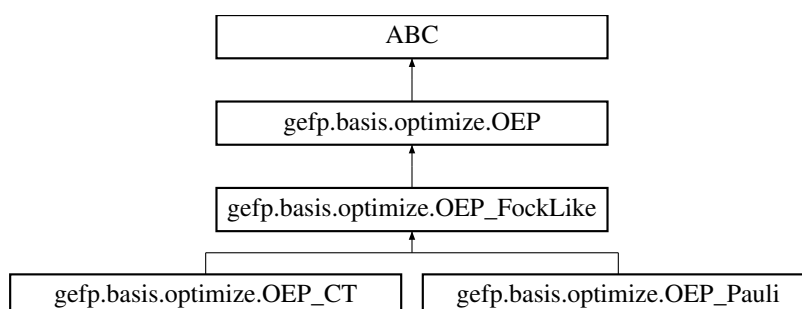
OEP for Group-(i) term of Otto-Ladik's theory of Charge-Transfer Energy

The documentation for this class was generated from the following file:

- `gefp/gefp/basis/optimize.py`

18.77 gefp.basis.optimize.OEP_FockLike Class Reference

Inheritance diagram for `gefp.basis.optimize.OEP_FockLike`:



Public Member Functions

- `def __init__ (self, wfn, dfbasis)`

Additional Inherited Members

18.77.1 Detailed Description

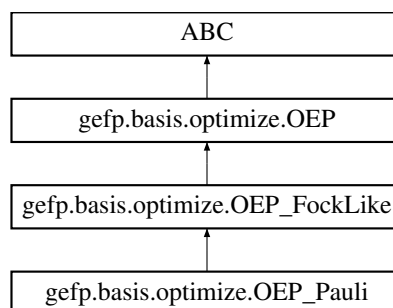
OEP for S1 term in Murrell et al.'s theory of Pauli repulsion

The documentation for this class was generated from the following file:

- `gefp/gefp/basis/optimize.py`

18.78 `gefp.basis.optimize.OEP_Pauli` Class Reference

Inheritance diagram for `gefp.basis.optimize.OEP_Pauli`:



Public Member Functions

- `def __init__ (self, wfn, dfbasis)`

Additional Inherited Members

18.78.1 Detailed Description

OEP for S1 term in Murrell et al.'s theory of Pauli repulsion

The documentation for this class was generated from the following file:

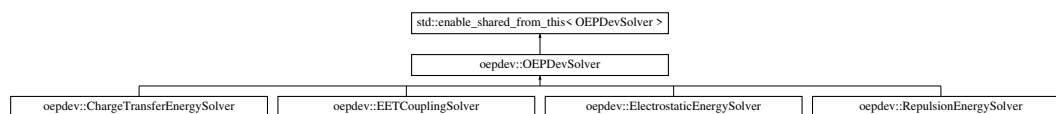
- `gefp/gefp/basis/optimize.py`

18.79 `oepdev::OEPDevSolver` Class Reference

Solver of properties of molecular aggregates. Abstract base.

```
#include <solver.h>
```

Inheritance diagram for oepdev::OEPDevSolver:



Public Member Functions

- [OEPDevSolver](#) (SharedWavefunctionUnion wfn_union)
Take wavefunction union and initialize the Solver.
- virtual [~OEPDevSolver](#) ()
Destructor.
- virtual double [compute_oep_based](#) (const std::string &method="DEFAULT")=0
Compute property by using OEP's.
- virtual double [compute_benchmark](#) (const std::string &method="DEFAULT")=0
Compute property by using benchmark method.

Static Public Member Functions

- static std::shared_ptr< [OEPDevSolver](#) > [build](#) (const std::string &target, SharedWavefunctionUnion wfn_union)
Build a solver of a particular property for given molecular cluster.

Protected Attributes

- SharedWavefunctionUnion [wfn_union_](#)
Wavefunction union.
- psi::Options & [options_](#)
Options.
- std::vector< std::string > [methods_oepBased_](#)
Names of all OEP-based methods implemented for a solver.
- std::vector< std::string > [methods_benchmark_](#)
Names of all benchmark methods implemented for a solver.

18.79.1 Detailed Description

Uses only a wavefunction union object to initialize.

Available solvers

- ELECTROSTATIC ENERGY
- REPULSION ENERGY
- CHARGE TRANSFER ENERGY
- EET COUPLING CONSTANT

Options

Interaction Property Method

- OEPDEV_SOLVER_EINT_COUL_AO - Coulombic energy: AO expanded
- OEPDEV_SOLVER_EINT_COUL_MO - Coulombic energy: MO expanded
- OEPDEV_SOLVER_EINT_COUL_ESP - Coulombic energy: ESP
- OEPDEV_SOLVER_EINT_COUL_CAMM - Coulombic energy: [CAMM](#)
- OEPDEV_SOLVER_EINT_REP_HS - Exchange-repulsion energy: Hayes-Stone
- OEPDEV_SOLVER_EINT_REP_DDS - Exchange-repulsion energy: DDS
- OEPDEV_SOLVER_EINT_REP_MRW - Exchange-repulsion energy: Murrell et al.
- OEPDEV_SOLVER_EINT_REP_OL - Exchange-repulsion energy: Otto-Ladik
- OEPDEV_SOLVER_EINT_REP_OEP1 - Exchange-repulsion energy: OEP (S1: GDF, S2: ESP)
- OEPDEV_SOLVER_EINT_REP_OEP2 - Exchange-repulsion energy: OEP (S1: GDF, S2: [CAMM](#))
- OEPDEV_SOLVER_EINT_REP_EFP2 - Exchange-repulsion energy: EFP2
- OEPDEV_SOLVER_EINT_CT_OL - Charge-transfer energy: Otto-Ladik
- OEPDEV_SOLVER_EINT_CT_OEP - Charge-transfer energy: OEP
- OEPDEV_SOLVER_EINT_CT_EFP2 - Charge-transfer energy: EFP2

Generalized density fitting (GDF) options:

- OEPDEV_DF_TYPE - type of the GDF. Default: DOUBLE. Other: SINGLE.
- DF_BASIS_OEP - auxiliary basis set. Default: sto-3g.
- DF_BASIS_INT - intermediate basis set. Relevant only if double GDF is used. Default: aug-cc-pVDZ-jkfit. Note that intermediate basis set should be nearly complete.

EFP2 Charge transfer energy options:

- `EFP2_CT_POTENTIAL_INTS` - Type of potential one-electron operator. Default: 'DMTP'. Other: 'ERI'.
- `EFP2_CT_NO_OCTUPOLES` - Ignore octupole moments from potential integrals? Default: `True`.

Excited States

- `EXCITED_STATE` - ID of state for all monomers to consider. If $-n$, then the n th bright state is taken. Default: -1 .
- `EXCITED_STATE_A` - ID of state for monomer A to consider. If $-n$, then the n th bright state is taken. Default: -1 .
- `EXCITED_STATE_B` - ID of state for monomer B to consider. If $-n$, then the n th bright state is taken. Default: -1 .
- `OSCILLATOR_STRENGTH_THRESHOLD` - Threshold for oscillator strength for bright states selection. Default: `0.01`.
- `TrCAMM_SYMMETRIZE` - Whether to use the 'symmetrized transition density' or not. Default: `true`.
- `TI_CIS_SCF_FOCK_MATRIX` - Whether to compute the full SCF Fock matrix for the dimer or approximate it from monomer OPDM's. Default: `false`.
- `TI_CIS_PRINT_FOCK_MATRIX` - Whether to print the Fock matrix (AB block in AO basis) or not. Default: `false`.

Environmental variables

One can easily access those variables from Python level by calling

```
psi4.get_variable("name of variable")
```

in your Python script.

Table 18.59: Environmental variables in the OEPDev solver.

Keyword	Description
Coulombic Interaction Energy	
<i>Distributed Multipole Series</i>	
EINT COUL CAMM R-1	CAMM charge-charge terms

Keyword	Description
EINT COUL CAMM R-2	CAMM charge-dipole terms + all above
EINT COUL CAMM R-3	CAMM charge-quadrupole, dipole-dipole + all above
EINT COUL CAMM R-4	CAMM charge-octupole, dipole-quadrupole + all above
EINT COUL CAMM R-5	CAMM charge-hexadecapole, dipole-octupole, quadrupole-quadrupole + all above
EINT COUL ESP	ESP charge-charge terms
Exact First-Order Perturbation Theory	
EINT COUL EXACT	MO or AO expanded Coulombic energy. Both give same results but MO is much faster.
Exchange-Repulsion Interaction Energy	
Density Decomposition Scheme	
EINT REP DDS KCAL	Pauli repulsion
EINT EXC DDS KCAL	DDS exchange
EINT EXR DDS KCAL	Sum of the above
Hayes-Stone model	
EINT REP HAYES-STONE KCAL	Pauli repulsion
EINT EXC HAYES-STONE KCAL	Pure exchange
EINT EXR HAYES-STONE KCAL	Sum of the above
Murrell et al. model	
EINT REP MURRELL-ETAL KCAL	Pauli repulsion
EINT EXC MURRELL-ETAL KCAL	Pure exchange (same as Hayes-Stone)
EINT EXR MURRELL-ETAL KCAL	Sum of the above
EINT REP MURRELL-ETAL:S1 KCAL	Pauli repulsion: S^{-1} term
EINT REP MURRELL-ETAL:S2 KCAL	Pauli repulsion: S^{-2} term
Otto-Ladik model	
EINT REP OTTO-LADIK KCAL	Pauli repulsion
EINT EXC OTTO-LADIK KCAL	Pure exchange (same as Hayes-Stone)
EINT EXR OTTO-LADIK KCAL	Sum of the above
EINT REP OTTO-LADIK:S1 KCAL	Pauli repulsion: S^{-1} term

Keyword	Description
EINT REP OTTO-LADIK:S2 KCAL	Pauli repulsion: S^{-2} term
EFP2 model	
EINT REP EFP2 KCAL	Pauli repulsion
EINT EXC EFP2 KCAL	Exchange: SGO approximation of Jensen
EINT EXR EFP2 KCAL	Sum of the above
EINT REP EFP2:S1 KCAL	Pauli repulsion: S^{-1} term
EINT REP EFP2:S2 KCAL	Pauli repulsion: S^{-2} term
OEP-based models	
EINT REP OEP-MURRELL-ETAL:S1-GDF KCAL	Pauli repulsion: S1 term using GDF, S2 term using CAMM
EINT REP OEP-MURRELL-ETAL:S1-GDF KCAL	S/S_{21} term of the above total term
EINT REP OEP-MURRELL-ETAL:S1-GDF KCAL	S/S_{22} term of the above total term
EINT REP OEP-MURRELL-ETAL:S1-GDF KCAL	Pauli repulsion: S1 term using GDF, S2 term using ESP
EINT REP OEP-MURRELL-ETAL:S1-GDF KCAL	S/S_{21} term of the above total term
EINT REP OEP-MURRELL-ETAL:S1-GDF KCAL	S/S_{22} term of the above total term
Charge-Transfer Interaction Energy	
EFP2 Model	
EINT CT EFP2 KCAL	Total charge-transfer energy (kcal/mole)
Otto-Ladik Model	
EINT CT OTTO-LADIK KCAL	Total charge-transfer energy (kcal/mole)
OEP-Based Otto-Ladik Model	
EINT CT OEP-OTTO-LADIK KCAL	Total charge-transfer energy (kcal/mole)
EET Coupling Constant	

Keyword	Description
<i>TrCamm Model</i>	
EET V0 TRCamm R1 CM-1	Overlap-uncorrected, converged to R1 (cm-1)
EET V TRCamm R1 CM-1	Overlap-corrected, converged to R1 (cm-1)
EET V0 TRCamm R2 CM-1	Overlap-uncorrected, converged to R2 (cm-1)
EET V TRCamm R2 CM-1	Overlap-corrected, converged to R2 (cm-1)
EET V0 TRCamm R3 CM-1	Overlap-uncorrected, converged to R3 (cm-1)
EET V TRCamm R3 CM-1	Overlap-corrected, converged to R3 (cm-1)
EET V0 TRCamm R4 CM-1	Overlap-uncorrected, converged to R4 (cm-1)
EET V TRCamm R4 CM-1	Overlap-corrected, converged to R4 (cm-1)
EET V0 TRCamm R5 CM-1	Overlap-uncorrected, converged to R5 (cm-1)
EET V TRCamm R5 CM-1	Overlap-corrected, converged to R5 (cm-1)
<i>Ti/CIS Model</i>	
EET V0 COUL CM-1	Overlap-uncorrected Coulomb (Forster) coupling (cm-1)
EET V0 EXCH CM-1	Overlap-uncorrected exchange (Dexter) coupling (cm-1)
EET V COUL CM-1	Overlap-corrected Coulomb (Forster) coupling (cm-1)
EET V EXCH CM-1	Overlap-corrected exchange (Dexter) coupling (cm-1)
EET V OVRL CM-1	Remaining overlap correction to direct coupling(cm-1)
EET V0 ET1 CM-1	Overlap-uncorrected H_13 matrix element (cm-1)
EET V0 ET2 CM-1	Overlap-uncorrected H_24 matrix element (cm-1)
EET V0 HT1 CM-1	Overlap-uncorrected H_14 matrix element (cm-1)
EET V0 HT2 CM-1	Overlap-uncorrected H_23 matrix element (cm-1)
EET V0 CT CM-1	Overlap-uncorrected H_34 matrix element (cm-1)
EET V ET1 CM-1	Overlap-corrected H_13 matrix element (cm-1)
EET V ET2 CM-1	Overlap-corrected H_24 matrix element (cm-1)
EET V HT1 CM-1	Overlap-corrected H_14 matrix element (cm-1)
EET V HT2 CM-1	Overlap-corrected H_23 matrix element (cm-1)

Keyword	Description
EET V CT CM-1	Overlap-corrected H_34 matrix element (cm-1)
EET V0 TI-2 CM-1	Approximate 2nd-order indirect coupling (cm-1)
EET V0 TI-3 CM-1	Approximate 3rd-order indirect coupling (cm-1)
EET V TI-2 CM-1	2nd-order indirect coupling (cm-1)
EET V TI-3 CM-1	3rd-order indirect coupling (cm-1)
EET V0 DIRECT CM-1	Approximate direct coupling (cm-1)
EET V0 INDIRECT CM-1	Approximate indirect coupling (cm-1)
EET V DIRECT CM-1	Direct coupling (cm-1)
EET V INDIRECT CM-1	Indirect coupling (cm-1)
EET V0 TI-CIS CM-1	Approximate total coupling (cm-1)
EET V TI-CIS CM-1	Total coupling (cm-1)
EET V0 EXCH-M CM-1	Overlap-uncorrected exchange (Dexter) coupling in Mulliken approximation (cm-1)
EET V EXCH-M CM-1	Overlap-corrected exchange (Dexter) coupling in Mulliken approximation (cm-1)
EET V0 CT-M CM-1	Overlap-uncorrected H_34 matrix element in Mulliken approximation (cm-1)
EET V CT-M CM-1	Overlap-corrected H_34 matrix element in Mulliken approximation (cm-1)
<i>OEP-Based TI/CIS Model</i>	
EET V OEP:COUL CM-1	Overlap-corrected Coulomb (Forster) coupling (TrCamm; cm-1)
EET V OEP:EXCH CM-1	Overlap-corrected exchange (Dexter) coupling (Mulliken approximation of AO ERI's; cm-1)
EET V OEP:OVRL CM-1	Remaining overlap correction to direct coupling (cm-1)
EET V0 OEP:ET1 CM-1	Overlap-uncorrected H_13 matrix element (cm-1)
EET V0 OEP:ET2 CM-1	Overlap-uncorrected H_24 matrix element (cm-1)
EET V0 OEP:HT1 CM-1	Overlap-uncorrected H_14 matrix element (cm-1)
EET V0 OEP:HT2 CM-1	Overlap-uncorrected H_23 matrix element (cm-1)
EET V0 OEP:CT:CAMM CM-1	Overlap-uncorrected H_34 matrix element: CAMM approximation of ionic interaction (cm-1)

Keyword	Description
EET V0 OEP:CT:CC CM-1	Overlap-uncorrected H ₃₄ matrix element: Point-charge approximation of ionic interaction (cm ⁻¹)
EET V OEP:ET1 CM-1	Overlap-corrected H ₁₃ matrix element (cm ⁻¹)
EET V OEP:ET2 CM-1	Overlap-corrected H ₂₄ matrix element (cm ⁻¹)
EET V OEP:HT1 CM-1	Overlap-corrected H ₁₄ matrix element (cm ⁻¹)
EET V OEP:HT2 CM-1	Overlap-corrected H ₂₃ matrix element (cm ⁻¹)
EET V OEP:CT:CAMM CM-1	Overlap-corrected H ₃₄ matrix element: CAMM approximation of ionic interaction (cm ⁻¹)
EET V OEP:CT:CC CM-1	Overlap-corrected H ₃₄ matrix element: Point-charge approximation of ionic interaction (cm ⁻¹)
EET V OEP:TI-2 CM-1	2nd-order indirect coupling (cm ⁻¹)
EET V OEP:TI-3:CAMM CM-1	3rd-order indirect coupling with CAMM approximation for V _{CT} (cm ⁻¹)
EET V OEP:TI-3:CC CM-1	3rd-order indirect coupling with point-charge approximation for V _{CT} (cm ⁻¹)
EET V OEP:DIRECT CM-1	Direct coupling (cm ⁻¹)
EET V OEP:INDIRECT:CAMM CM-1	Indirect coupling with CAMM approximation for V _{CT} (cm ⁻¹)
EET V OEP:INDIRECT:CC CM-1	Indirect coupling with point-charge approximation for V _{CT} (cm ⁻¹)
EET V OEP:TI-CIS:CAMM CM-1	Total coupling with CAMM approximation for V _{CT} (cm ⁻¹)
EET V OEP:TI-CIS:CC CM-1	Total coupling with point-charge approximation for V _{CT} (cm ⁻¹)

18.79.2 Constructor & Destructor Documentation

18.79.2.1 OEPDevSolver()

```
OEPDevSolver::OEPDevSolver (
    SharedWavefunctionUnion wfn_union )
```

Parameters

<i>wfn_union</i>	- wavefunction union of isolated molecular wavefunctions
------------------	--

18.79.3 Member Function Documentation

18.79.3.1 build()

```
std::shared_ptr< OEPDevSolver > OEPDevSolver::build (
    const std::string & target,
    SharedWavefunctionUnion wfn_union ) [static]
```

Parameters

<i>target</i>	- target property
<i>wfn_union</i>	- wavefunction union of isolated molecular wavefunctions

Implemented target properties:

- `ELECTROSTATIC_ENERGY` - Coulombic interaction energy between unperturbed wavefunctions.
- `REPULSION_ENERGY` - Pauli repulsion interaction energy between unperturbed wavefunctions.

See also

[ElectrostaticEnergySolver](#)

18.79.3.2 compute_benchmark()

```
double OEPDevSolver::compute_benchmark (
    const std::string & method = "DEFAULT" ) [pure virtual]
```

Each solver object has one `DEFAULT` benchmark method

Parameters

<i>method</i>	- benchmark method
---------------	--------------------

Implemented in [oepdev::EETCouplingSolver](#), [oepdev::ChargeTransferEnergySolver](#), [oepdev::RepulsionEnergySolver](#) and [oepdev::ElectrostaticEnergySolver](#).

18.79.3.3 compute_oep_based()

```
double OEPDevSolver::compute_oep_based (
    const std::string & method = "DEFAULT" ) [pure virtual]
```

Each solver object has one `DEFAULT` OEP-based method.

Parameters

<i>method</i>	- flavour of OEP model
---------------	------------------------

Implemented in [oepdev::EETCouplingSolver](#), [oepdev::ChargeTransferEnergySolver](#), [oepdev::RepulsionEnergySolver](#) and [oepdev::ElectrostaticEnergySolver](#).

The documentation for this class was generated from the following files:

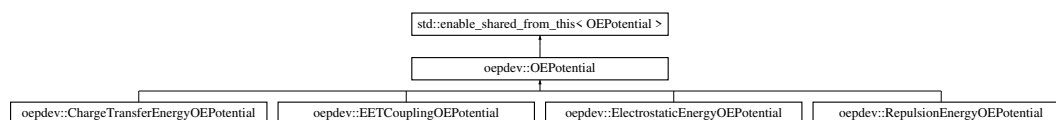
- [oepdev/libsolver/solver.h](#)
- [oepdev/libsolver/solver_base.cc](#)

18.80 oepdev::OEPotential Class Reference

Generalized One-Electron Potential: Abstract base.

```
#include <oep.h>
```

Inheritance diagram for `oepdev::OEPotential`:



Public Member Functions

- [OEPotential](#) (SharedWavefunction [wfn](#), Options &options)
Fully ESP-based OEP object.
- [OEPotential](#) (SharedWavefunction [wfn](#), SharedBasisSet auxiliary, SharedBasisSet intermediate, Options &options)
General OEP object.
- virtual [~OEPotential](#) ()
Destructor.
- virtual void [compute](#) (void)
Compute matrix forms of all OEP's within all OEP types.
- virtual void [compute](#) (const std::string &oepType)=0
Compute matrix forms of all OEP's within a specified OEP type.

- virtual void [compute_3D](#) (const std::string &oepType, const double &x, const double &y, const double &z, std::shared_ptr< psi::Vector > &v)=0
Compute value of potential in point x, y, z and save at v.
- std::shared_ptr< [OEPotential3D](#)< [OEPotential](#) > > [make_oeps3d](#) (const std::string &oepType)
Create 3D vector field with OEP.
- virtual void [write_cube](#) (const std::string &oepType, const std::string &fileName)
Write potential to a cube file.
- virtual void [localize](#) (void)
Localize Occupied MO's.
- virtual std::vector< psi::SharedVector > [mo_centroids](#) (psi::SharedMatrix C)
Compute MO centroids from LCAO-MO matrix.
- virtual void [rotate](#) (const Matrix &rotmat)
Rotate.
- virtual void [translate](#) (const Vector &trans)
Translate.
- virtual void [superimpose](#) (const Matrix &refGeometry, const std::vector< int > &supList, const std::vector< int > &reordList)
Superimpose.
- std::string [name](#) () const
Retrieve name of this OEP.
- [OEPTyp](#)e [oep](#) (const std::string &oepType) const
Retrieve the potentials.
- SharedMatrix [matrix](#) (const std::string &oepType) const
Retrieve the potentials of a particular OEP type in a matrix form.
- int [n](#) (const std::string &oepType) const
Retrieve the number of a particular OEP type.
- SharedWavefunction [wfn](#) () const
Retrieve wavefunction object.
- SharedMatrix [cOcc](#) () const
- SharedMatrix [cVir](#) () const
- SharedMatrix [IOcc](#) () const
- SharedLocalizer [localizer](#) () const
Retrieve MO Localizer.
- std::vector< std::shared_ptr< psi::Vector > > [lmoc](#) () const
Retrieve LMO Centroids.
- void [set_name](#) (const std::string &name)
Set the name of this OEP.
- virtual void [print_header](#) () const =0
Header information.
- void [print](#) () const
Print the contents (OEP data)

Static Public Member Functions

- static `std::shared_ptr< OEPotential > build` (const `std::string` &category, SharedWavefunction `wfn`, Options &options)
Build fully ESP-based OEP object.
- static `std::shared_ptr< OEPotential > build` (const `std::string` &category, SharedWavefunction `wfn`, SharedBasisSet auxiliary, SharedBasisSet intermediate, Options &options)
Build general OEP object.

Protected Attributes

- Options `options_`
Psi4 options.
- SharedWavefunction `wfn_`
Wavefunction.
- SharedBasisSet `primary_`
Promary Basis set.
- SharedBasisSet `auxiliary_`
Auxiliary Basis set.
- SharedBasisSet `intermediate_`
Intermediate Basis set.
- SharedLocalizer `localizer_`
Molecular Orbital Localizer.
- `std::string` `name_`
Name of this OEP;.
- `std::map< std::string, OEType > oepTypes_`
Types of OEP's within the scope of this object.
- `std::shared_ptr< psi::IntegralFactory > intsFactory_`
Integral factory.
- `std::shared_ptr< psi::Matrix > potMat_`
Matrix of potential one-electron integrals.
- `std::shared_ptr< psi::OneBodyAOInt > OEInt_`
One-electron integral shared pointer.
- `std::shared_ptr< oepdev::PotentialInt > potInt_`
One-electron potential shared pointer.
- `std::shared_ptr< psi::Matrix > cOcc_`
Occupied orbitals: Canonical (CMO)
- `std::shared_ptr< psi::Matrix > cVir_`
Virtual orbitals.
- `std::shared_ptr< psi::Matrix > lOcc_`
Occupied orbitals: Localized (LMO)
- `std::vector< std::shared_ptr< psi::Vector > > lmoc_`
LMO Centroids.

18.80.1 Detailed Description

Manages OEP's in matrix and 3D forms. Available OEP categories:

- ELECTROSTATIC ENERGY
- REPULSION ENERGY
- CHARGE TRANSFER ENERGY
- EET COUPLING CONSTANT

18.80.2 Constructor & Destructor Documentation

18.80.2.1 OEPotential() [1/2]

```
OEPotential::OEPotential (  
    SharedWavefunction wfn,  
    Options & options )
```

Parameters

<i>wfn</i>	- wavefunction
<i>options</i>	- Psi4 options

18.80.2.2 OEPotential() [2/2]

```
OEPotential::OEPotential (  
    SharedWavefunction wfn,  
    SharedBasisSet auxiliary,  
    SharedBasisSet intermediate,  
    Options & options )
```

Parameters

<i>wfn</i>	- wavefunction
<i>auxiliary</i>	- auxiliary basis set for density fitting of OEP's
<i>intermediate</i>	- intermediate basis set for density fitting of OEP's
<i>options</i>	- Psi4 options

18.80.3 Member Function Documentation

18.80.3.1 build() [1/2]

```
std::shared_ptr< OEPotential > OEPotential::build (
    const std::string & category,
    SharedWavefunction wfn,
    Options & options ) [static]
```

Parameters

<i>type</i>	- OEP category
<i>wfn</i>	- wavefunction
<i>options</i>	- Psi4 options

18.80.3.2 build() [2/2]

```
std::shared_ptr< OEPotential > OEPotential::build (
    const std::string & category,
    SharedWavefunction wfn,
    SharedBasisSet auxiliary,
    SharedBasisSet intermediate,
    Options & options ) [static]
```

Parameters

<i>type</i>	- OEP category
<i>wfn</i>	- wavefunction
<i>auxiliary</i>	- auxiliary basis set for density fitting of OEP's
<i>intermediate</i>	- intermediate basis set for density fitting of OEP's
<i>options</i>	- Psi4 options

18.80.3.3 make_oeps3d()

```
std::shared_ptr< OEPotential3D< OEPotential > > OEPotential::make_oeps3d (
    const std::string & oepType )
```

Parameters

<i>oepType</i>	- type of OEP. ESP-based OEP is assumed.
----------------	--

Returns

Vector field 3D with the OEP values.

The documentation for this class was generated from the following files:

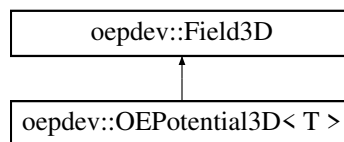
- oepdev/liboep/oep.h
- oepdev/liboep/oep_base.cc

18.81 oepdev::OEPotential3D< T > Class Template Reference

Class template for OEP 3D fields.

```
#include <space3d.h>
```

Inheritance diagram for oepdev::OEPotential3D< T >:

**Public Member Functions**

- [OEPotential3D](#) (const int &ndim, const int &np, const double &padding, std::shared_ptr< T > oep, const std::string &oepType)
Construct random spherical collection of 3D field of type T.
- [OEPotential3D](#) (const int &ndim, const int &nx, const int &ny, const int &nz, const double &px, const double &py, const double &pz, std::shared_ptr< T > oep, const std::string &oepType, psi::Options &options)
Construct ordered 3D collection of 3D field of type T.
- virtual [~OEPotential3D](#) ()
Destructor.
- virtual std::shared_ptr< psi::Vector > [compute_xyz](#) (const double &x, const double &y, const double &z)
Compute a value of 3D field at point (x, y, z)
- virtual void [print](#) () const
Print information of the object to Psi4 output.

Protected Attributes

- std::shared_ptr< T > [oep_](#)
Shared pointer to the instance of class T
- std::string [oepType_](#)
Descriptor of the 3D field type stored in instance of T

Additional Inherited Members

18.81.1 Detailed Description

`template<class T>`
`class oepdev::OEPotential3D< T >`

Used for special type of classes T that contain following public member functions:

```
class T : public std::enable_shared_from_this<T> {
public:
    void compute_3D(const std::string& descriptor,
                   const double& x, const double& y, const double& z,
                   std::shared_ptr<psi::Vector> &v);

    shared_ptr<psi::Wavefunction> wfn() const {return wfn-;}
};
```

with the `descriptor` of a certain 3D field type, `x`, `y`, `z` the points in 3D space in which the scalar or vector field has to be computed and stored at `v`. Instances of `T` should store shared pointer to wavefunction object. List of classes `T` that are compatible with this class template and are currently implemented in `oepdev` is given below:

- `oepdev::OEPotential` abstract base (do not use derived classes as `T`)

Template parameters:

Template Parameters

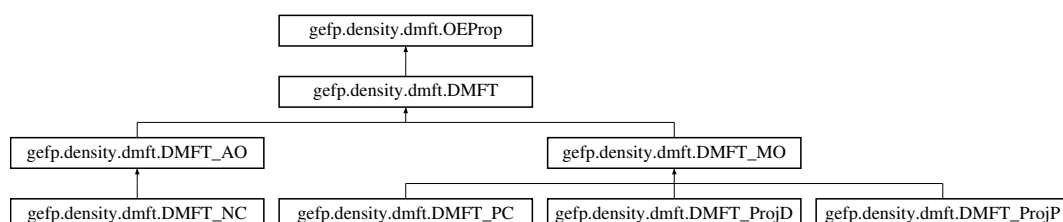
<code>T</code>	the compatible class (e.g. <code>oepdev::OEPotential</code>)
----------------	---

The documentation for this class was generated from the following file:

- `oepdev/lib3d/space3d.h`

18.82 gefp.density.dmft.OEProp Class Reference

Inheritance diagram for `gefp.density.dmft.OEProp`:



Static Public Member Functions

- def **dipole_moment** (dmft)
- def **quadrupole_moment** (dmft)

The documentation for this class was generated from the following file:

- `gefp/gefp/density/dmft.py`

18.83 oepdev::OEType Struct Reference

Container to handle the type of One-Electron Potentials.

```
#include <oep.h>
```

Public Attributes

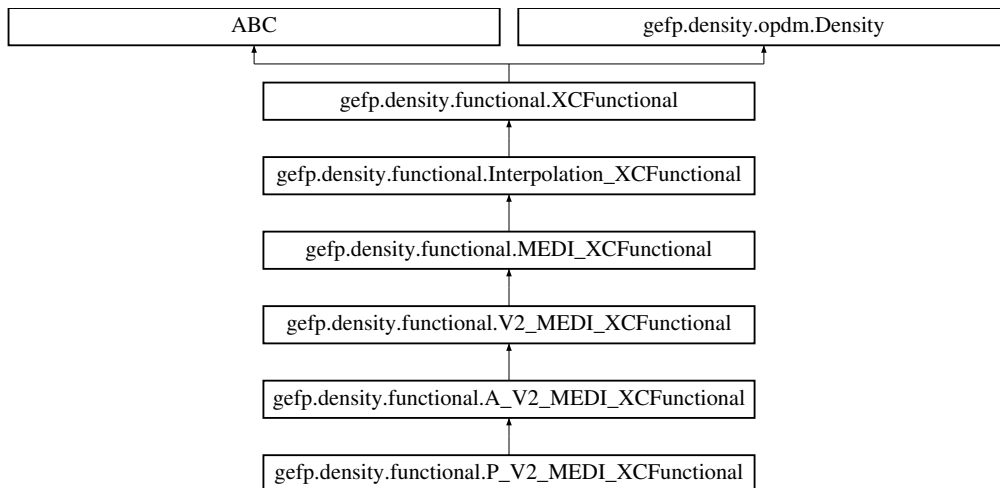
- std::string [name](#)
Name of this type of OEP.
- bool [is_density_fitted](#)
Is this OEP DF-based?
- int [n](#)
Number of OEP's within a type.
- SharedMatrix [matrix](#)
All OEP's of this type gathered in a matrix form.
- SharedDMTPole [dmtp](#)
Distributed Multipole Object.
- SharedCISData [cis_data](#)
CIS data.

The documentation for this struct was generated from the following file:

- `oepdev/liboep/oep.h`

18.84 gefp.density.functional.P_V2_MEDI_XCFunctional Class Reference

Inheritance diagram for gefp.density.functional.P_V2_MEDI_XCFunctional:



Public Member Functions

- `def __init__ (self, coeff, kmax)`
- `def abbr (self)`
- `def compute.t (self, n=None, c=None)`

Static Public Member Functions

- `def name ()`

Additional Inherited Members

18.84.1 Detailed Description

The New Class of Exchange-Correlation Functionals:

Interpolation Functionals with Monotonous Exponential Decay: 2D Pade Approximant for Un

Functional coefficients:

- `'pade_coefficients'` - 2D Pade Coefficients that fit the `'t'` parameter.

The documentation for this class was generated from the following file:

- `gefp/gefp/density/functional.py`

18.85 gefp.core.driver.PadeApproximant_2D Class Reference

Public Member Functions

- `def __init__ (self)`
- `def add_a (self, nx, ny, a)`

- def **add_b** (self, nx, ny, b)
- def **value** (self, x, y)

Public Attributes

- **a**
- **b**

18.85.1 Detailed Description

\

Pade Approximant of Function of 2 Variables:

$$f(x, y) = \frac{\sum_{n=0, m=0} a_{nm} x^n y^m}{1.0 + \sum_{n, m \neq (0, 0)} b_{nm} x^n y^m}$$

The documentation for this class was generated from the following file:

- gefp/gefp/core/driver.py

18.86 oepdev::PerturbCharges Struct Reference

Structure to hold perturbing charges.

```
#include <scf_perturb.h>
```

Public Attributes

- std::vector< double > [charges](#)
Vector of charge values.
- std::vector< std::shared_ptr< psi::Vector > > [positions](#)
Vector of charge position vectors.

18.86.1 Detailed Description

The documentation for this struct was generated from the following file:

- oepdev/libutil/[scf_perturb.h](#)

18.87 oepdev::Points3DIterator::Point Struct Reference

Public Attributes

- double **x**
- double **y**
- double **z**
- int **index**

The documentation for this struct was generated from the following file:

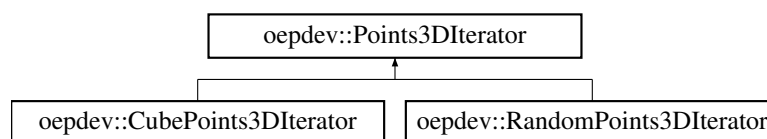
- oepdev/lib3d/space3d.h

18.88 oepdev::Points3DIterator Class Reference

Iterator over a collection of points in 3D space. Abstract base.

```
#include <space3d.h>
```

Inheritance diagram for oepdev::Points3DIterator:



Classes

- struct [Point](#)

Public Member Functions

- [Points3DIterator](#) (const int &np)
Plain constructor. Initializes the abstract features.
- virtual [~Points3DIterator](#) ()
Destructor.
- virtual bool [is_done](#) ()
Check if iteration is finished.
- virtual void [first](#) ()=0
Initialize first iteration.
- virtual void [next](#) ()=0
Step to next iteration.

- virtual void [rewind](#) ()
Rewind to the beginning.

- virtual double **x** () const
- virtual double **y** () const
- virtual double **z** () const
- virtual int **index** () const

Static Public Member Functions

- static shared_ptr< [Points3DIterator](#) > [build](#) (const int &nx, const int &ny, const int &nz, const double &dx, const double &dy, const double &dz, const double &ox, const double &oy, const double &oz)
Build G09 Cube collection iterator.
- static shared_ptr< [Points3DIterator](#) > [build](#) (const int &np, const double &radius, const double &cx, const double &cy, const double &cz)
Build random collection iterator.
- static shared_ptr< [Points3DIterator](#) > [build](#) (const int &np, const double &pad, psi::SharedMolecule mol)
Build random collection iterator.

Protected Attributes

- const int [np_](#)
Number of points.
- bool [done_](#)
Status of the iterator.
- int [index_](#)
Current index.
- [Point](#) [current_](#)

18.88.1 Detailed Description

Points3DIterators are constructed either as iterators over:

- a random collections or
- an ordered (g09 cube-like) collections. **Note:** Always create instances by using static factory methods.

18.88.2 Constructor & Destructor Documentation

18.88.2.1 Points3DIterator()

```
oepdev::Points3DIterator::Points3DIterator (
    const int & np )
```

Parameters

<i>np</i>	- number of points this iterator is constructed for
-----------	---

18.88.3 Member Function Documentation

18.88.3.1 build() [1/3]

```
std::shared_ptr< Points3DIterator > oepdev::Points3DIterator::build (
    const int & nx,
    const int & ny,
    const int & nz,
    const double & dx,
    const double & dy,
    const double & dz,
    const double & ox,
    const double & oy,
    const double & oz ) [static]
```

The points are generated according to Gaussian cube file format.

Parameters

<i>nx</i>	- number of points along x direction
<i>ny</i>	- number of points along y direction
<i>nz</i>	- number of points along z direction
<i>dx</i>	- spacing distance along x direction
<i>dy</i>	- spacing distance along y direction
<i>dz</i>	- spacing distance along y direction
<i>ox</i>	- coordinate x of cube origin
<i>oy</i>	- coordinate y of cube origin
<i>oz</i>	- coordinate z of cube origin

18.88.3.2 build() [2/3]

```
std::shared_ptr< Points3DIterator > oepdev::Points3DIterator::build (
    const int & np,
    const double & radius,
    const double & cx,
    const double & cy,
    const double & cz ) [static]
```

The points are drawn according to uniform distrinution in 3D space.

Parameters

<i>np</i>	- number of points to draw
<i>radius</i>	- sphere radius inside which points are to be drawn
<i>cx</i>	- coordinate x of sphere's centre
<i>cy</i>	- coordinate y of sphere's centre
<i>cz</i>	- coordinate z of sphere's centre

18.88.3.3 build() [3/3]

```
shared_ptr< Points3DIterator > oepdev::Points3DIterator::build (
    const int & np,
    const double & pad,
    psi::SharedMolecule mol ) [static]
```

The points are drawn according to uniform distrinution in 3D space enclosing a molecule given. All drawn points lie outside the van der Waals volume.

Parameters

<i>np</i>	- number of points to draw
<i>pad</i>	- radius padding of a minimal sphere enclosing the molecule
<i>mol</i>	- Psi4 molecule object

The documentation for this class was generated from the following files:

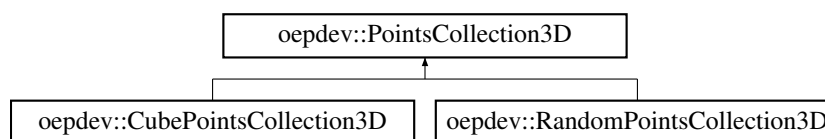
- oepdev/lib3d/space3d.h
- oepdev/lib3d/space3d.cc

18.89 oepdev::PointsCollection3D Class Reference

Collection of points in 3D space. Abstract base.

```
#include <space3d.h>
```

Inheritance diagram for oepdev::PointsCollection3D:



Public Types

- enum [Collection](#) { **Random**, **Cube** }
Public descriptor of collection type.

Public Member Functions

- [PointsCollection3D](#) ([Collection](#) collectionType, int &np)
Initialize abstract features.
- PointsCollection3D** ([Collection](#) collectionType, const int &np)
- virtual [~PointsCollection3D](#) ()
Destructor.
- virtual int [npoints](#) () const
Get the number of points.
- virtual shared_ptr< [Points3DIterator](#) > [points_iterator](#) () const
Get the iterator over this collection of points.
- virtual [Collection](#) [get_type](#) () const
Get the collection type.
- virtual void [print](#) () const =0
Print the information to Psi4 output file.

Static Public Member Functions

- static shared_ptr< [PointsCollection3D](#) > [build](#) (const int &[npoints](#), const double &radius, const double &cx=0.0, const double &cy=0.0, const double &cz=0.0)
Build random collection of points.
- static shared_ptr< [PointsCollection3D](#) > [build](#) (const int &[npoints](#), const double &padding, psi::SharedMolecule mol)
Build random collection of points.
- static shared_ptr< [PointsCollection3D](#) > [build](#) (const int &nx, const int &ny, const int &nz, const double &px, const double &py, const double &pz, psi::SharedBasisSet bs, psi::Options &options)
Build G09 Cube collection of points.

Protected Attributes

- const int `np_`
Number of points.
- `Collection` `collectionType_`
Collection type.
- `shared_ptr< Points3DIterator >` `pointsIterator_`
iterator over points collection

18.89.1 Detailed Description

Create random or ordered (g09 cube-like) collections of points in 3D space.

Note: Always create instances by using static factory methods.

18.89.2 Constructor & Destructor Documentation

18.89.2.1 PointsCollection3D()

```
oepdev::PointsCollection3D::PointsCollection3D (
    Collection collectionType,
    int & np )
```

Parameters

<i>np</i>	- number of points to be created
-----------	----------------------------------

18.89.3 Member Function Documentation

18.89.3.1 build() [1/3]

```
std::shared_ptr< PointsCollection3D > oepdev::PointsCollection3D::build (
    const int & npoints,
    const double & radius,
    const double & cx = 0.0,
    const double & cy = 0.0,
    const double & cz = 0.0 ) [static]
```

Points uniformly span a sphere.

Parameters

<i>npoints</i>	- number of points to draw
<i>radius</i>	- sphere radius inside which points are to be drawn
<i>cx</i>	- coordinate x of sphere's centre
<i>cy</i>	- coordinate y of sphere's centre
<i>cz</i>	- coordinate z of sphere's centre

18.89.3.2 build() [2/3]

```
std::shared_ptr< PointsCollection3D > oepdev::PointsCollection3D::build (
    const int & npoints,
    const double & padding,
    psi::SharedMolecule mol ) [static]
```

Points uniformly span space inside a sphere enclosing a molecule. excluding the van der Waals volume.

Parameters

<i>np</i>	- number of points to draw
<i>padding</i>	- radius padding of a minimal sphere enclosing the molecule
<i>mol</i>	- Psi4 molecule object

18.89.3.3 build() [3/3]

```
std::shared_ptr< PointsCollection3D > oepdev::PointsCollection3D::build (
    const int & nx,
    const int & ny,
    const int & nz,
    const double & px,
    const double & py,
    const double & pz,
    psi::SharedBasisSet bs,
    psi::Options & options ) [static]
```

The points span a parallelepiped according to Gaussian cube file format.

Parameters

<i>nx</i>	- number of points along x direction
<i>ny</i>	- number of points along y direction
<i>nz</i>	- number of points along z direction

Parameters

<i>px</i>	- padding distance along x direction
<i>py</i>	- padding distance along y direction
<i>pz</i>	- padding distance along z direction
<i>bs</i>	- Psi4 basis set object
<i>options</i>	- Psi4 options object

The documentation for this class was generated from the following files:

- oepdev/lib3d/space3d.h
- oepdev/lib3d/space3d.cc

18.90 oepdev::PolarGEFactory Class Reference

Polarization GEFP Factory. Abstract Base.

```
#include <gefp.h>
```

Inheritance diagram for oepdev::PolarGEFactory:



Public Member Functions

- [PolarGEFactory](#) (std::shared_ptr< psi::Wavefunction > [wfn](#), psi::Options &opt)
Construct from Psi4 options.
- virtual [~PolarGEFactory](#) ()
Destruct.
- virtual std::shared_ptr< [GenEffPar](#) > [compute](#) (void)=0
Compute the density matrix susceptibility tensors.

Protected Member Functions

- std::shared_ptr< psi::Vector > [draw_field](#) ()
Randomly draw electric field value.
- double [draw_charge](#) ()
Randomly draw charge value.
- std::shared_ptr< [oepdev::RHFPerturbed](#) > [perturbed_state](#) (const std::shared_ptr< psi::Vector > &field)
Solve SCF equations to find perturbed state due to uniform electric field.

- `std::shared_ptr< oepdev::RHFPerturbed > perturbed_state` (const `std::shared_ptr< psi::Vector > &pos`, const double &charge)
Solve SCF equations to find perturbed state due to point charge.
- `std::shared_ptr< oepdev::RHFPerturbed > perturbed_state` (const `std::shared_ptr< psi::Matrix > &charges`)
Solve SCF equations to find perturbed state due to set of point charges.
- `std::shared_ptr< psi::Vector > field_due_to_charges` (const `std::shared_ptr< psi::Matrix > &charges`, const double &x, const double &y, const double &z)
Evaluate electric field at point (x,y,z) due to point charges.
- `std::shared_ptr< psi::Vector > field_due_to_charges` (const `std::shared_ptr< psi::Matrix > &charges`, const `std::shared_ptr< psi::Vector > &pos`)
- `std::shared_ptr< psi::Matrix > field_gradient_due_to_charges` (const `std::shared_ptr< psi::Matrix > &charges`, const double &x, const double &y, const double &z)
Evaluate electric field gradient at point (x,y,z) due to point charges.
- `std::shared_ptr< psi::Matrix > field_gradient_due_to_charges` (const `std::shared_ptr< psi::Matrix > &charges`, const `std::shared_ptr< psi::Vector > &pos`)

Additional Inherited Members

18.90.1 Detailed Description

Basic interface for the polarization density matrix susceptibility parameters.

The documentation for this class was generated from the following files:

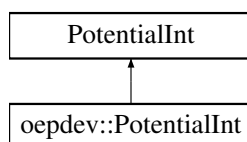
- `oepdev/libgefp/gefp.h`
- `oepdev/libgefp/gefp_polar_base.cc`

18.91 oepdev::PotentialInt Class Reference

Computes potential integrals.

```
#include <potential.h>
```

Inheritance diagram for `oepdev::PotentialInt`:



Public Member Functions

- `PotentialInt` (`std::vector< psi::SphericalTransform > &st`, `std::shared_ptr< psi::BasisSet > bs1`, `std::shared_ptr< psi::BasisSet > bs2`, int deriv=0)

Constructor. Initialize identically like in psi::Potentillnt.

- [PotentialInt](#) (std::vector< psi::SphericalTransform > &st, std::shared_ptr< psi::BasisSet > bs1, std::shared_ptr< psi::BasisSet > bs2, std::shared_ptr< psi::Matrix > Qxyz, int deriv=0)

Constructor. Takes an arbitrary collection of charges.

- [PotentialInt](#) (std::vector< psi::SphericalTransform > &, std::shared_ptr< psi::BasisSet >, std::shared_ptr< psi::BasisSet >, const double &x, const double &y, const double &z, const double &q=1.0, int deriv=0)

Constructor. Computes potential for one point x, y, z for a test particle of charge q.

- void [set_charge_field](#) (const double &x, const double &y, const double &z, const double &q=1.0)

Mutator. Set the charge field to be a x, y, z point of charge q.

18.91.1 Constructor & Destructor Documentation

18.91.1.1 PotentialInt() [1/3]

```
oepdev::PotentialInt::PotentialInt (
    std::vector< psi::SphericalTransform > & st,
    std::shared_ptr< psi::BasisSet > bs1,
    std::shared_ptr< psi::BasisSet > bs2,
    int deriv = 0 )
```

Parameters

<i>st</i>	- Spherical transform object
<i>bs1</i>	- basis set for first space
<i>bs2</i>	- basis set for second space
<i>deriv</i>	- derivative level

18.91.1.2 PotentialInt() [2/3]

```
oepdev::PotentialInt::PotentialInt (
    std::vector< psi::SphericalTransform > & st,
    std::shared_ptr< psi::BasisSet > bs1,
    std::shared_ptr< psi::BasisSet > bs2,
    std::shared_ptr< psi::Matrix > Qxyz,
    int deriv = 0 )
```

Parameters

<i>st</i>	- Spherical transform object
<i>bs1</i>	- basis set for first space
<i>bs2</i>	- basis set for second space
<i>Qxyz</i>	- matrix with charges and their positions
<i>deriv</i>	- derivative level

18.91.1.3 PotentialInt() [3/3]

```
oepdev::PotentialInt::PotentialInt (
    std::vector< psi::SphericalTransform > & st,
    std::shared_ptr< psi::BasisSet > bs1,
    std::shared_ptr< psi::BasisSet > bs2,
    const double & x,
    const double & y,
    const double & z,
    const double & q = 1.0,
    int deriv = 0 )
```

Parameters

<i>st</i>	- Spherical transform object
<i>bs1</i>	- basis set for first space
<i>bs2</i>	- basis set for second space
<i>x</i>	- x coordinate of q
<i>y</i>	- y coordinate of q
<i>z</i>	- z coordinate of q
<i>q</i>	- value of the probe charge
<i>deriv</i>	- derivative level

18.91.2 Member Function Documentation**18.91.2.1 set_charge_field()**

```
void oepdev::PotentialInt::set_charge_field (
    const double & x,
    const double & y,
    const double & z,
    const double & q = 1.0 )
```

Parameters

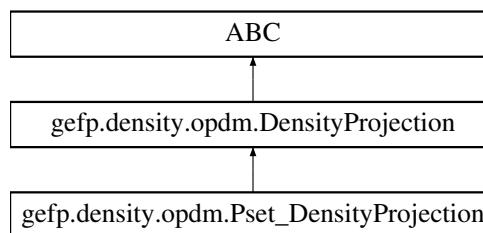
x	- x coordinate of q
y	- y coordinate of q
z	- z coordinate of q
q	- value of the probe charge

The documentation for this class was generated from the following files:

- oepdev/libpsi/[potential.h](#)
- oepdev/libpsi/potential.cc

18.92 gefp.density.opdm.Pset_DensityProjection Class Reference

Inheritance diagram for gefp.density.opdm.Pset_DensityProjection:

**Public Member Functions**

- `def __init__(self, np, S)`

Additional Inherited Members**18.92.1 Detailed Description**

\
 Gradient Projection Algorithm on P-sets.
 Ref.: Pernal, Cancès, J. Chem. Phys. 2005

Notes:

- Appropriate for any DMFT functional.

The documentation for this class was generated from the following file:

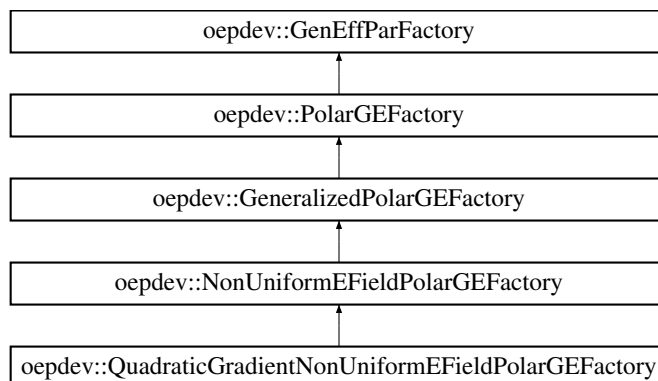
- `gefp/gefp/density/opdm.py`

18.93 oepdev::QuadraticGradientNonUniformEFieldPolarGEFactory Class Reference

Polarization GEFP Factory with Least-Squares Parameterization.

```
#include <gefp.h>
```

Inheritance diagram for oepdev::QuadraticGradientNonUniformEFieldPolarGEFactory:



Public Member Functions

- **QuadraticGradientNonUniformEFieldPolarGEFactory** (`std::shared_ptr< psi::Wavefunction > wfn`, `psi::Options &opt`)
- void **compute_gradient** (`int i`, `int j`)
Compute Gradient vector associated with the i-th and j-th basis set function.
- void **compute_hessian** (`void`)
Compute Hessian matrix (independent on the parameters)

Additional Inherited Members

18.93.1 Detailed Description

Implements the generalized density matrix susceptibility model of the form

$$\delta D_{\alpha\beta} \approx \sum_i \left\{ \mathbf{B}_{i;\alpha\beta}^{(10)} \cdot \mathbf{F}(\mathbf{r}_i) + \mathbf{B}_{i;\alpha\beta}^{(20)} : \mathbf{F} \otimes \mathbf{F}(\mathbf{r}_i) + \mathbf{B}_{i;\alpha\beta}^{(01)} : \nabla_i \otimes \mathbf{F}(\mathbf{r}_i) \right\}$$

where:

- $\mathbf{B}_{i;\alpha\beta}^{(10)}$ is the density matrix dipole polarizability
- $\mathbf{B}_{i;\alpha\beta}^{(20)}$ is the density matrix dipole-dipole hyperpolarizability
- $\mathbf{B}_{i;\alpha\beta}^{(01)}$ is the density matrix quadrupole polarizability all defined for the distributed site at \mathbf{r}_i .

Note

This model is not available now and probably will be deprecated in the future.

The documentation for this class was generated from the following files:

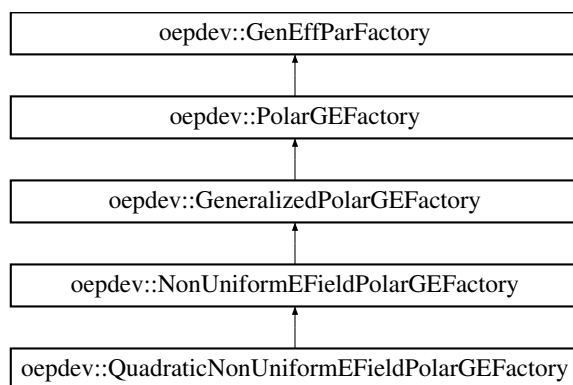
- oepdev/libgefp/gefp.h
- oepdev/libgefp/gefp_polar_nonuniform_field_2_grad_1.cc

18.94 oepdev::QuadraticNonUniformEFieldPolarGEFactory Class Reference

Polarization GEFP Factory with Least-Squares Parameterization.

```
#include <gefp.h>
```

Inheritance diagram for oepdev::QuadraticNonUniformEFieldPolarGEFactory:



Public Member Functions

- **QuadraticNonUniformEFieldPolarGEFactory** (std::shared_ptr< psi::Wavefunction > wfn, psi::Options &opt)
- void **compute_gradient** (int i, int j)
Compute Gradient vector associated with the i-th and j-th basis set function.
- void **compute_hessian** (void)
Compute Hessian matrix (independent on the parameters)

Additional Inherited Members

18.94.1 Detailed Description

Implements the generalized density matrix susceptibility model of the form

$$\delta D_{\alpha\beta} \approx \sum_i \left\{ \mathbf{B}_{i;\alpha\beta}^{(10)} \cdot \mathbf{F}(\mathbf{r}_i) + \mathbf{B}_{i;\alpha\beta}^{(20)} : \mathbf{F}(\mathbf{r}_i) \otimes \mathbf{F}(\mathbf{r}_i) \right\}$$

where:

- $\mathbf{B}_{i;\alpha\beta}^{(10)}$ is the density matrix dipole polarizability
- $\mathbf{B}_{i;\alpha\beta}^{(20)}$ is the density matrix dipole-dipole hyperpolarizability all defined for the distributed site at \mathbf{r}_i .

The documentation for this class was generated from the following files:

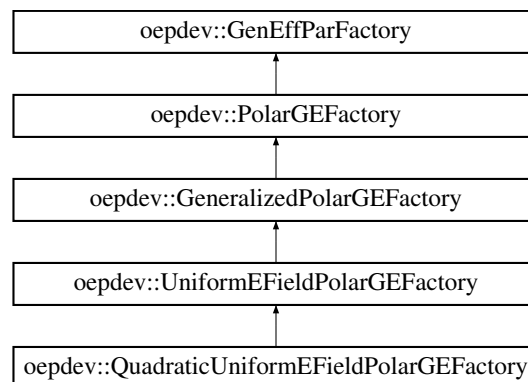
- oepdev/libgefp/gefp.h
- oepdev/libgefp/gefp_polar_nonuniform_field_2.cc

18.95 oepdev::QuadraticUniformEFieldPolarGEFactory Class Reference

Polarization GEFP Factory with Least-Squares Parameterization.

```
#include <gefp.h>
```

Inheritance diagram for oepdev::QuadraticUniformEFieldPolarGEFactory:



Public Member Functions

- **QuadraticUniformEFieldPolarGEFactory** (std::shared_ptr< psi::Wavefunction > wfn, psi::Options &opt)
- void **compute_gradient** (int i, int j)
Compute Gradient vector associated with the i-th and j-th basis set function.
- void **compute_hessian** (void)
Compute Hessian matrix (independent on the parameters)

Additional Inherited Members

18.95.1 Detailed Description

Implements the generalized density matrix susceptibility model of the form

$$\delta D_{\alpha\beta} \approx \mathbf{B}_{\alpha\beta}^{(10)} \cdot \mathbf{F} + \mathbf{B}_{\alpha\beta}^{(20)} : \mathbf{F} \otimes \mathbf{F}$$

where:

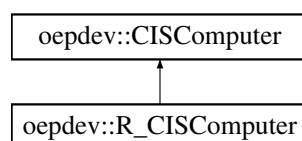
- $\mathbf{B}_{\alpha\beta}^{(10)}$ is the density matrix dipole polarizability
- $\mathbf{B}_{\alpha\beta}^{(20)}$ is the density matrix dipole-dipole hyperpolarizability

The documentation for this class was generated from the following files:

- oepdev/libgefp/[gefp.h](#)
- oepdev/libgefp/gefp_polar_uniform_field_2.cc

18.96 oepdev::R_CISComputer Class Reference

Inheritance diagram for oepdev::R_CISComputer:



Public Member Functions

- **R_CISComputer** (std::shared_ptr< psi::Wavefunction > wfn, psi::Options &opt)

Protected Member Functions

- virtual void **set_beta_** (void)
- virtual void **build_hamiltonian_** (void)

Additional Inherited Members

The documentation for this class was generated from the following files:

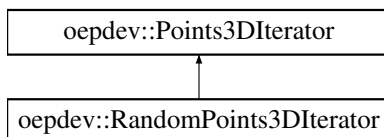
- oepdev/libutil/[cis.h](#)
- oepdev/libutil/cis_rhf.cc

18.97 oepdev::RandomPoints3DIterator Class Reference

Iterator over a collection of points in 3D space. Random collection.

```
#include <space3d.h>
```

Inheritance diagram for oepdev::RandomPoints3DIterator:



Public Member Functions

- **RandomPoints3DIterator** (const int &np, const double &radius, const double &cx, const double &cy, const double &cz)
- **RandomPoints3DIterator** (const int &np, const double &pad, psi::SharedMolecule mol)
- virtual void [first](#) ()
Initialize first iteration.
- virtual void [next](#) ()
Step to next iteration.

Protected Member Functions

- virtual double **random_double** ()
- virtual void **draw_random_point** ()
- virtual bool **is_in_vdWsphere** (double x, double y, double z) const

Protected Attributes

- double **cx_**
- double **cy_**
- double **cz_**
- double **radius_**
- double **r_**
- double **phi_**
- double **theta_**
- double **x_**
- double **y_**
- double **z_**
- psi::SharedMatrix **excludeSpheres_**
- std::map< std::string, double > **vdwRadius_**
- std::default_random_engine **randomNumberGenerator_**
- std::uniform_real_distribution< double > **randomDistribution_**

Additional Inherited Members

18.97.1 Detailed Description

Note: Always create instances by using static factory method from [Points3DIterator](#). Do not use constructors of this class.

The documentation for this class was generated from the following files:

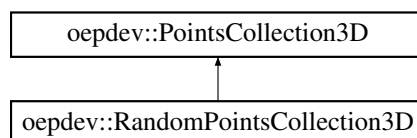
- oepdev/lib3d/space3d.h
- oepdev/lib3d/space3d.cc

18.98 oepdev::RandomPointsCollection3D Class Reference

Collection of random points in 3D space.

```
#include <space3d.h>
```

Inheritance diagram for oepdev::RandomPointsCollection3D:



Public Member Functions

- **RandomPointsCollection3D** ([Collection](#) collectionType, const int &npoints, const double &radius, const double &cx, const double &cy, const double &cz)
- **RandomPointsCollection3D** ([Collection](#) collectionType, const int &npoints, const double &padding, psi::SharedMolecule mol)
- virtual void [print](#) () const

Print the information to Psi4 output file.

Additional Inherited Members

18.98.1 Detailed Description

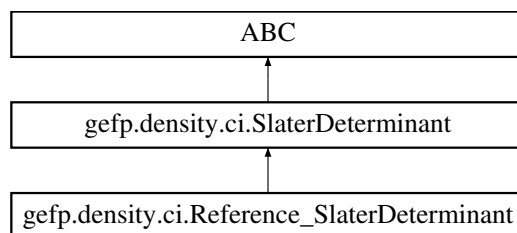
Note: Do not use constructors of this class explicitly. Instead, use static factory methods of the superclass to create instances.

The documentation for this class was generated from the following files:

- oepdev/lib3d/space3d.h
- oepdev/lib3d/space3d.cc

18.99 gefp.density.ci.Reference_SlaterDeterminant Class Reference

Inheritance diagram for gefp.density.ci.Reference_SlaterDeterminant:



Public Member Functions

- `def __init__ (self, nao, nbo, nmo)`

Public Attributes

- **`is_reference`**

The documentation for this class was generated from the following file:

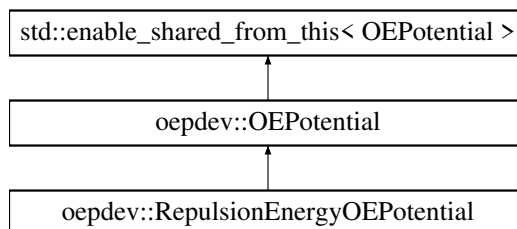
- `gefp/gefp/density/ci.py`

18.100 oepdev::RepulsionEnergyOEPotential Class Reference

Generalized One-Electron Potential for Pauli Repulsion Energy.

```
#include <oep.h>
```

Inheritance diagram for oepdev::RepulsionEnergyOEPotential:



Public Member Functions

- **`RepulsionEnergyOEPotential`** (SharedWavefunction [wfn](#), SharedBasisSet auxiliary, SharedBasisSet intermediate, Options &options)
- **`RepulsionEnergyOEPotential`** (SharedWavefunction [wfn](#), Options &options)

- virtual void [compute](#) (const std::string &oepType) override
Compute matrix forms of all OEP's within a specified OEP type.
- virtual void [compute_3D](#) (const std::string &oepType, const double &x, const double &y, const double &z, std::shared_ptr< psi::Vector > &v) override
Compute value of potential in point x, y, z and save at v.
- virtual void [print_header](#) () const override
Header information.

Additional Inherited Members

18.100.1 Detailed Description

Contains the following OEP types:

- Murrell-etal.S1
- Otto-Ladik.S2

The documentation for this class was generated from the following files:

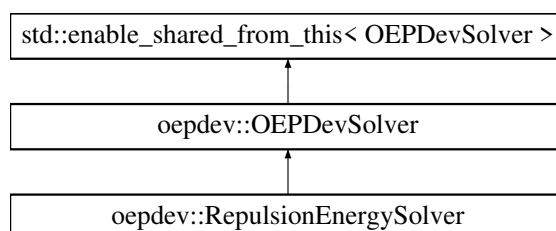
- oepdev/liboep/[oep.h](#)
- oepdev/liboep/oep_energy_pauli.cc

18.101 oepdev::RepulsionEnergySolver Class Reference

Compute the Pauli-Repulsion interaction energy between unperturbed wavefunctions.

```
#include <solver.h>
```

Inheritance diagram for oepdev::RepulsionEnergySolver:



Public Member Functions

- **RepulsionEnergySolver** (SharedWavefunctionUnion wfn_union)
- virtual double [compute_oep_based](#) (const std::string &method="DEFAULT")
Compute property by using OEP's.
- virtual double [compute_benchmark](#) (const std::string &method="DEFAULT")
Compute property by using benchmark method.

Additional Inherited Members

18.101.1 Detailed Description

The implemented methods are shown below

Table 18.82: Methods available in the Solver

Keyword	Method Description
Benchmark Methods	
HAYES_STONE	<i>Default.</i> Pauli Repulsion energy at HF level from Hayes and Stone (1984).
DDS	Pauli Repulsion energy at HF level from Mandado and Hermida-Ramon (2012).
MURRELL_ETAL	Approximate Pauli Repulsion energy at HF level from Murrell et al (1967).
OTTO_LADIK	Approximate Pauli Repulsion energy at HF level from Otto and Ladik (1975).
EFP2	Approximate Pauli Repulsion energy at HF level from EFP2 model.
OEP-Based Methods	
MURRELL_ETAL_GDF_ESP	<i>Default.</i> OEP-Murrell et al's: S1 term via DF-OEP, S2 term via ESP-OEP.
MURRELL_ETAL_GDF_CAMM	OEP-Murrell et al's: S1 term via DF-OEP, S2 term via CAMM-OEP.
MURRELL_ETAL_ESP	OEP-Murrell et al's: S1 and S2 via ESP-OEP (not implemented)

Note:

- This solver also computes and prints the exchange energy at HF level (formulae are given below) for reference purposes.
- In order to construct this solver, **always** use the `OEPDevSolver::build` static factory method.

Below the detailed description of the implemented equations is given for each of the above provided methods. In the formulae across, it is assumed that the orbitals are real. The Coulomb notation for electron repulsion integrals (ERI's) is adopted; i.e.,

$$(ac|bd) = \iint d\mathbf{r}_1 d\mathbf{r}_2 \phi_a(\mathbf{r}_1) \phi_c(\mathbf{r}_1) \frac{1}{r_{12}} \phi_b(\mathbf{r}_2) \phi_d(\mathbf{r}_2)$$

Greek subscripts denote basis set orbitals whereas Italic subscripts denote the occupied molecular orbitals.

Benchmark Methods

Pauli Repulsion energy at HF level by Hayes and Stone (1984).

For a closed-shell system, equation of Hayes and Stone (1984) becomes

$$E^{\text{Rep}} = 2 \sum_{kl} \left(V_{kl}^A + V_{kl}^B + T_{kl} \right) \left[[\mathbf{S}^{-1}]_{lk} - \delta_{lk} \right] + \sum_{klmn} (kl|mn) \left\{ 2[\mathbf{S}^{-1}]_{kl}[\mathbf{S}^{-1}]_{mn} - [\mathbf{S}^{-1}]_{kn}[\mathbf{S}^{-1}]_{lm} - 2\delta_{kl}\delta_{mn} + \delta_{kn}\delta_{lm} \right\}$$

where \mathbf{S} is the overlap matrix between the doubly-occupied orbitals. The exact, pure exchange energy is for a closed shell case given as

$$E^{\text{Ex,pure}} = -2 \sum_{a \in A} \sum_{b \in B} (ab|ba)$$

Similarity transformation of molecular orbitals does not affect the resulting energies. The overall exchange-repulsion interaction energy is then (always net repulsive)

$$E^{\text{Ex-Rep}} = E^{\text{Ex,pure}} + E^{\text{Rep}}$$

Repulsion energy of Mandado and Hermida-Ramon (2011)

At the Hartree-Fock level, the exchange-repulsion energy from the density-based scheme of Mandado and Hermida-Ramon (2011) is fully equivalent to the method by Hayes and Stone (1984). However, density-based method enables to compute exchange-repulsion energy at any level of theory. It is derived based on the Pauli deformation density matrix,

$$\Delta \mathbf{D}^{\text{Pauli}} \equiv \mathbf{D}^{oo} - \mathbf{D}$$

where \mathbf{D}^{oo} and \mathbf{D} are the density matrix formed from mutually orthogonal sets of molecular orbitals within the entire aggregate (formed by symmetric orthogonalization of MO's) and the density matrix of the unperturbed system (that can be understood as a Hadamard sum $\mathbf{D} \equiv \mathbf{D}^A \oplus \mathbf{D}^B$).

At HF level, the Pauli deformation density matrix is given by

$$\Delta \mathbf{D}^{\text{Pauli}} = \mathbf{C} [\mathbf{S}^{-1} - \mathbf{1}] \mathbf{C}^\dagger$$

whereas the density matrix constructed from mutually orthogonal orbitals is

$$\mathbf{D}^{oo} = \mathbf{C} \mathbf{S}^{-1} \mathbf{C}^\dagger$$

In the above equations, \mathbf{S} is the overlap matrix between doubly occupied molecular orbitals of the entire aggregate.

Here, the expressions for the exchange-repulsion energy at any level of theory are shown for the case of open-shell system. The net repulsive energy is given as

$$E^{\text{Ex-Rep}} = E^{\text{Rep,1}} + E^{\text{Rep,2}} + E^{\text{Ex}}$$

where the one- and two-electron part of the repulsion energy is

$$\begin{aligned} E^{\text{Rep,1}} &= E^{\text{Rep,Kin}} + E^{\text{Rep,Nuc}} \\ E^{\text{Rep,2}} &= E^{\text{Rep,el-}\Delta} + E^{\text{Rep,}\Delta-\Delta} \end{aligned}$$

The kinetic and nuclear contributions are

$$E^{\text{Rep,Kin}} = 2 \sum_{\alpha\beta \in A,B} \Delta D_{\alpha\beta}^{\text{Pauli}} T_{\alpha\beta}$$

$$E^{\text{Rep,Nuc}} = 2 \sum_{\alpha\beta \in A,B} \Delta D_{\alpha\beta}^{\text{Pauli}} \sum_{z \in A,B} V_{\alpha\beta}^{(z)}$$

whereas the electron-deformation and deformation-deformation interaction contributions are

$$E^{\text{Rep,el-}\Delta} = 4 \sum_{\alpha\beta\gamma\delta \in A,B} \Delta D_{\alpha\beta}^{\text{Pauli}} D_{\gamma\delta}(\alpha\beta|\gamma\delta)$$

$$E^{\text{Rep,}\Delta-\Delta} = 2 \sum_{\alpha\beta\gamma\delta \in A,B} \Delta D_{\alpha\beta}^{\text{Pauli}} \Delta D_{\gamma\delta}^{\text{Pauli}}(\alpha\beta|\gamma\delta)$$

The associated exchange energy is given by

$$E^{\text{Ex}} = - \sum_{\alpha\beta\gamma\delta \in A,B} \left[D_{\alpha\delta}^{oo} D_{\beta\gamma}^{oo} - D_{\alpha\delta}^A D_{\beta\gamma}^A - D_{\alpha\delta}^B D_{\beta\gamma}^B \right] (\alpha\beta|\gamma\delta)$$

It is important to emphasise that, although, at HF level, the particular 'repulsive' and 'exchange' energies computed by using either Hayes and Stone or Mandado and Hermida-Ramon methods are not equal to each other, they sum up to exactly the same exchange-repulsion energy, $E^{\text{Ex-Rep}}$. Therefore, these methods at HF level are fully equivalent but the nature of partitioning of repulsive and exchange parts is different. It is also noted that the orbital localization does *not* affect the resulting energies, as opposed to the few approximate methods described below (Otto-Ladik and EFP2 methods).

Approximate Pauli Repulsion energy at HF level from Murrell et al.

By expanding the overlap matrix in a Taylor series one can show that the Pauli repulsion energy is approximately given as

$$E^{\text{Rep}} = E^{\text{Rep}}(\mathcal{O}(S)) + E^{\text{Rep}}(\mathcal{O}(S^2))$$

where the first-order term is

$$E^{\text{Rep}}(\mathcal{O}(S)) = -2 \sum_{a \in A} \sum_{b \in B} S_{ab} \left\{ V_{ab}^A + \sum_{c \in A} [2(ab|cc) - (ac|bc)] + V_{ab}^B + \sum_{d \in B} [2(ab|dd) - (ad|bd)] \right\}$$

whereas the second-order term is

$$E^{\text{Rep}}(\mathcal{O}(S^2)) = 2 \sum_{a \in A} \sum_{b \in B} S_{ab} \left\{ \sum_{c \in A} S_{bc} \left[V_{ac}^B + 2 \sum_{d \in B} (ac|dd) \right] + \sum_{d \in B} S_{ad} \left[V_{bd}^A + 2 \sum_{x \in A} (bd|cc) \right] - \sum_{c \in A} \sum_{d \in B} S_{cd} (ac|bd) \right\}$$

Thus derived repulsion energy is invariant with respect to transformation of molecular orbitals, similarly as Hayes-Stone's method and density-based method. By using OEP technique, the above theory can be exactly re-cast *without* any further approximations.

Approximate Pauli Repulsion energy at HF level from Otto and Ladik (1975).

The Pauli repulsion energy is approximately given as

$$E^{\text{Rep}} = E^{\text{Rep}}(\mathcal{O}(S)) + E^{\text{Rep}}(\mathcal{O}(S^2))$$

where the first-order term is

$$E^{\text{Rep}}(\mathcal{O}(S)) = -2 \sum_{a \in A} \sum_{b \in B} S_{ab} \left\{ V_{ab}^A + 2 \sum_{c \in A} (ab|cc) - (ab|aa) + V_{ab}^B + 2 \sum_{d \in B} (ab|dd) - (ab|bb) \right\}$$

whereas the second-order term is

$$E^{\text{Rep}}(\mathcal{O}(S^2)) = 2 \sum_{a \in A} \sum_{b \in B} S_{ab}^2 \left\{ V_{aa}^B + V_{bb}^A + 2 \sum_{c \in A} (cc|bb) + 2 \sum_{d \in B} (aa|dd) - (aa|bb) \right\}$$

Thus derived repulsion energy is *not* invariant with respect to transformation of molecular orbitals, in contrast to Hayes-Stone's method and density-based method. It was shown that good results are obtained when using localized molecular orbitals, whereas using canonical molecular orbitals brings poor results. By using OEP technique, the above theory can be exactly re-cast *without* any further approximations.

Approximate Pauli Repulsion energy at HF level from Jensen and Gordon (1996).

The Pauli repulsion energy used within the EFP2 approach is approximately given as

$$E^{\text{Rep}} = E^{\text{Rep}}(\mathcal{O}(S)) + E^{\text{Rep}}(\mathcal{O}(S^2))$$

where the first-order term is

$$E^{\text{Rep}}(\mathcal{O}(S)) = -2 \sum_{a \in A} \sum_{b \in B} S_{ab} \left\{ \sum_{c \in A} F_{ac}^A S_{cb} + \sum_{d \in B} F_{bd}^B S_{da} - 2T_{ab} \right\}$$

whereas the second-order term is

$$E^{\text{Rep}}(\mathcal{O}(S^2)) = 2 \sum_{a \in A} \sum_{b \in B} S_{ab}^2 \left\{ \sum_{x \in A} \frac{-Z_x}{R_{xb}} + \sum_{y \in B} \frac{-Z_y}{R_{ya}} + \sum_{c \in A} \frac{2}{R_{bc}} + \sum_{d \in B} \frac{2}{R_{ad}} - \frac{1}{R_{ab}} \right\}$$

Thus derived repulsion energy is *not* invariant with respect to transformation of molecular orbitals, in contrast to Hayes-Stone's method and density-based method. It was shown that good results are obtained when using localized molecular orbitals, whereas using canonical molecular orbitals brings poor results.

In EFP2, exchange energy is approximated by spherical Gaussian approximation (SGO). The result of this is the following formula for the exchange energy:

$$E^{\text{Ex}} \approx -4 \sum_{a \in A} \sum_{b \in B} \sqrt{\frac{-2 \ln |S_{ab}|}{\pi}} \frac{S_{ab}^2}{R_{ab}}$$

In all the above formulas, R_{ij} are distances between position vectors of i -th and j -th point. The LMO centroids are defined by

$$\mathbf{r}_a = (a|\mathbf{r}|a)$$

where a denotes the occupied molecular orbital.

OEP-Based Methods

The Murrell et al's theory of Pauli repulsion for S-1 term and the Otto-Ladik's theory for S-2 term is here re-cast by introducing OEP's. The S-1 term is expressed via DF-OEP, whereas the S-2 term via ESP-OEP.

S-1 term (Murrell et al.)

The OEP reduction without any approximations leads to the following formula

$$E^{\text{Rep}}(\mathcal{O}(S^1)) = -2 \sum_{a \in A} \sum_{b \in B} S_{ab} \left\{ \sum_{\xi \in A} S_{b\xi} G_{\xi a}^A + \sum_{\eta \in B} S_{a\eta} G_{\eta b}^B \right\}$$

where the OEP matrices are given as

$$G_{\xi a}^A = \sum_{\xi' \in A} [\mathbf{S}^{-1}]_{\xi\xi'} \sum_{\alpha \in A} \left\{ C_{\alpha a} V_{\alpha\xi'}^A + \sum_{\mu \nu \in A} [2C_{\alpha a} D_{\mu\nu} - C_{\nu a} D_{\alpha\mu}] (\alpha\xi' | \mu\nu) \right\}$$

and analogously for molecule *B*. Here, the nuclear attraction integrals are denoted by $V_{\alpha\xi'}^A$.

S-2 term (Otto-Ladik)

After the OEP reduction, this contribution under Otto-Ladik approximation has the following form:

$$E^{\text{Rep}}(\mathcal{O}(S^2)) = 2 \sum_{a \in A} \sum_{b \in B} S_{ab}^2 \left\{ \sum_{x \in A} q_{xa} V_{bb}^{(x)} + \sum_{y \in B} q_{yb} V_{aa}^{(y)} \right\}$$

where the ESP charges associated with each occupied molecular orbital reproduce the *effective potential* of molecule in question, i.e.,

$$\sum_{x \in A} \frac{q_{xa}}{|\mathbf{r} - \mathbf{r}_x|} \cong v_a^A(\mathbf{r})$$

where the potential is given by

$$v_a^A(\mathbf{r}) = \sum_{x \in A} \frac{-Z_x}{|\mathbf{r} - \mathbf{r}_x|} + 2 \sum_{c \in A} \int \frac{\phi_c(\mathbf{r}') \phi_c(\mathbf{r}')}{|\mathbf{r} - \mathbf{r}'|} d\mathbf{r}' - \frac{1}{2} \int \frac{\phi_a(\mathbf{r}') \phi_a(\mathbf{r}')}{|\mathbf{r} - \mathbf{r}'|} d\mathbf{r}'$$

18.101.2 Member Function Documentation

18.101.2.1 compute_benchmark()

```
double RepulsionEnergySolver::compute_benchmark (
    const std::string & method = "DEFAULT" ) [virtual]
```

Each solver object has one `DEFAULT` benchmark method

Parameters

<i>method</i>	- benchmark method
---------------	--------------------

Implements [oepdev::OEPDevSolver](#).

18.101.2.2 compute_oep_based()

```
double RepulsionEnergySolver::compute_oep_based (
    const std::string & method = "DEFAULT" ) [virtual]
```

Each solver object has one DEFAULT OEP-based method.

Parameters

<i>method</i>	- flavour of OEP model
---------------	------------------------

Implements [oepdev::OEPDevSolver](#).

The documentation for this class was generated from the following files:

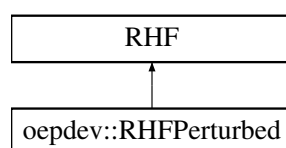
- oepdev/libsolver/[solver.h](#)
- oepdev/libsolver/solver_energy_pauli.cc

18.102 oepdev::RHFPerturbed Class Reference

RHF theory under electrostatic perturbation.

```
#include <scf_perturb.h>
```

Inheritance diagram for oepdev::RHFPerturbed:



Public Member Functions

- [RHFPerturbed](#) (std::shared_ptr< psi::Wavefunction > ref_wfn, std::shared_ptr< psi::SuperFunctional > functional)
Build from wavefunction and superfunctional.
- [RHFPerturbed](#) (std::shared_ptr< psi::Wavefunction > ref_wfn, std::shared_ptr< psi::SuperFunctional > functional, psi::Options &options, std::shared_ptr< psi::PSIO > psio)

Build from wavefunction and superfunctional + options and psio.

- virtual `~RHFPerturbed ()`

Clear memory.

- virtual double `compute_energy ()`

Compute total energy.

- virtual void `set_perturbation (std::shared_ptr< psi::Vector > field)`

Perturb the system with external electric field.

- virtual void `set_perturbation (const double &fx, const double &fy, const double &fz)`

This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

- virtual void `set_perturbation (std::shared_ptr< psi::Vector > position, const double &charge)`

Perturb the system with a point charge.

- virtual void `set_perturbation (const double &rx, const double &ry, const double &rz, const double &charge)`

This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

- `std::shared_ptr< psi::Matrix > Vpert () const`

Get a copy of the perturbation potential one-electron matrix.

- double `nuclear_interaction_energy () const`

Get the interaction energy of the nuclei with the perturbing potential.

Protected Member Functions

- virtual void `perturb_Hcore ()`

Add the electrostatic perturbation to the Hcore matrix.

Protected Attributes

- `std::shared_ptr< psi::Vector > perturbField_`

Perturbing electric field.

- `std::shared_ptr< PerturbCharges > perturbCharges_`

Perturbing charges.

- `std::shared_ptr< psi::Matrix > Vpert_`

Perturbation potential one-electron matrix.

- double `nuclearInteractionEnergy_`

Electrostatic interaction energy due to nuclei.

18.102.1 Detailed Description

Compute RHF wavefunction under the following conditions:

- external uniform electric field
- set of point charges The mixed conditions can also be used.

Theory

The electrostatic perturbation is here understood as a distribution of external (generally non-uniform) electric field. It is assumed that this perturbation is one-electron in nature. Therefore, the one-electron Hamiltonian is changed according to the following

$$\mathbf{H}^{\text{core}} \rightarrow \mathbf{H}^{\text{core}} + \sum_n q_n \mathbf{V}^{(n)} - \mathbb{M} \cdot \mathbf{F}$$

where q_n is the external classical point charge, $\mathbf{V}^{(n)}$ is the associated matrix of potential integrals, \mathbb{M} is the vector of dipole integrals and \mathbf{F} is an external uniform electric field. The total energy is then computed by performing an SCF procedure on the above one-electron Hamiltonian. The contribution due to nuclei is included, i.e.,

$$E_{\text{Nuc}} \rightarrow E_{\text{Nuc-Nuc}} + \sum_{In} \frac{q_n Z_I}{r_{In}} - \mu_{\text{Nuc}} \cdot \mathbf{F}$$

where μ_{Nuc} is the nuclear dipole moment and Z_I is the atomic number of the I th nucleus. It is added in the nuclear repulsion energy $E_{\text{Nuc-Nuc}}$ (note that the resulting energy can be negative as well depending on the electric field direction and configuration of point charges).

The documentation for this class was generated from the following files:

- oepdev/libutil/[scf_perturb.h](#)
- oepdev/libutil/scf_perturb.cc

18.103 gefp.density.dfi.SCF Class Reference

Public Member Functions

- `def __init__(self, mol, bfs=None)`
- `def run(self, maxit=30, conv=1.0e-7, guess=None, damp=0.01, ndamp=10, verbose=True, v_ext=None)`

Public Attributes

- [e_nuc](#)
Accessors nuclear repulsion energy.
- **E**

- **D**
- **Co**
- **C**
- **F**
- **eps**
- **H**
- **S**
- **X**

18.103.1 Detailed Description

Self-Consistent Field (SCF) Procedure for Hartree-Fock Model

Demo for RHF-SCF method (closed shells). Implements SCF algorithm with primitive damping of the AO Fock matrix.

Usage:

```
scf = SCF(molecule)
scf.run(maxit=30, conv=1.0e-7, guess=None, damp=0.01, ndamp=10, verbose=True)
```

The above example runs SCF on 'molecule' psi4.core.Molecule object starting from core Hamiltonian as guess (guess=None) and convergence 1.0E-7 A.U. in total energy with 30 maximum iterations (10 of which are performed by damping of the Fock matrix with damping coefficient of 0.1). The SCF iterations are printed to standard output (verbose=True).

Last Revision: Gund

The documentation for this class was generated from the following file:

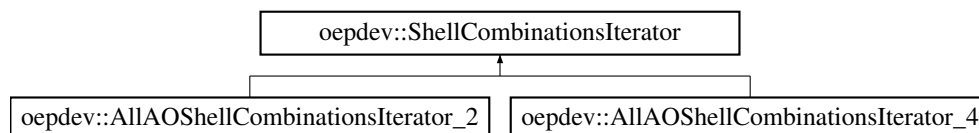
- gefp/gefp/density/dfi.py

18.104 oepdev::ShellCombinationsIterator Class Reference

Iterator for Shell Combinations. Abstract Base.

```
#include <integrals_iter.h>
```

Inheritance diagram for oepdev::ShellCombinationsIterator:



Public Member Functions

- [ShellCombinationsIterator](#) (int nshell)
Constructor.
- virtual [~ShellCombinationsIterator](#) ()
Destructor.
- virtual void [first](#) (void)=0
First iteration.
- virtual void [next](#) (void)=0
Next iteration.
- virtual std::shared_ptr< psi::BasisSet > [bs_1](#) (void) const
Grab the basis set of axis 1.
- virtual std::shared_ptr< psi::BasisSet > [bs_2](#) (void) const
Grab the basis set of axis 2.
- virtual std::shared_ptr< psi::BasisSet > [bs_3](#) (void) const
Grab the basis set of axis 3.
- virtual std::shared_ptr< psi::BasisSet > [bs_4](#) (void) const
Grab the basis set of axis 4.
- virtual int [P](#) (void) const
Grab the current shell P index.
- virtual int [Q](#) (void) const
Grab the current shell Q index.
- virtual int [R](#) (void) const
Grab the current shell R index.
- virtual int [S](#) (void) const
Grab the current shell S index.
- virtual bool [is_done](#) (void)
Return status of an iterator.
- virtual const int [nshell](#) (void) const
Return number of shells this iterator is for.
- virtual std::shared_ptr< [AOIntegralsIterator](#) > [ao_iterator](#) (std::string mode="ALL") const

- virtual void [compute_shell](#) (std::shared_ptr< oepdev::TwoBodyAOInt > tei) const =0
- virtual void [compute_shell](#) (std::shared_ptr< psi::TwoBodyAOInt > tei) const =0

Static Public Member Functions

- static std::shared_ptr< [ShellCombinationsIterator](#) > [build](#) (const [IntegralFactory](#) &ints, std::string mode="ALL", int [nshell](#)=4)

Build shell iterator from [oepdev::IntegralFactory](#).

- static std::shared_ptr< [ShellCombinationsIterator](#) > [build](#) (std::shared_ptr< [IntegralFactory](#) > ints, std::string mode="ALL", int [nshell](#)=4)

This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

- static std::shared_ptr< [ShellCombinationsIterator](#) > [build](#) (const psi::IntegralFactory &ints, std::string mode="ALL", int [nshell](#)=4)

Build shell iterator from [psi::IntegralFactory](#).

- static std::shared_ptr< [ShellCombinationsIterator](#) > [build](#) (std::shared_ptr< psi::IntegralFactory > ints, std::string mode="ALL", int [nshell](#)=4)

This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

Protected Attributes

- SharedBasisSet [bs_1_](#)

Basis set of axis 1.

- SharedBasisSet [bs_2_](#)

Basis set of axis 2.

- SharedBasisSet [bs_3_](#)

Basis set of axis 3.

- SharedBasisSet [bs_4_](#)

Basis set of axis 4.

- const int [nshell_](#)

Number of shells this iterator is for.

- bool [done](#)

Status of an iterator.

18.104.1 Detailed Description

Date

2018/03/01 17:22:00

18.104.2 Constructor & Destructor Documentation

18.104.2.1 ShellCombinationsIterator()

```
ShellCombinationsIterator::ShellCombinationsIterator (
    int nshell )
```

Parameters

<i>nshell</i>	- number of shells this iterator is for
---------------	---

18.104.3 Member Function Documentation

18.104.3.1 ao_iterator()

```
std::shared_ptr< AOIntegralsIterator > ShellCombinationsIterator::ao_iterator
(
    std::string mode = "ALL" ) const [virtual]
```

Make an AO integral iterator based on current shell

Parameters

<i>mode</i>	- either "ALL" or "UNIQUE" (iterate over all or unique integrals)
-------------	---

Returns

iterator over AO integrals

18.104.3.2 build() [1/2]

```
std::shared_ptr< ShellCombinationsIterator > ShellCombinationsIterator::build
(
    const IntegralFactory & ints,
    std::string mode = "ALL",
    int nshell = 4 ) [static]
```

Parameters

<i>ints</i>	- integral factory
<i>mode</i>	- mode of iteration (either ALL or UNIQUE)
<i>nshell</i>	- number of shells to iterate through

Returns

shell iterator

Examples:

[example_integrals_iter.cc](#).

18.104.3.3 build() [2/2]

```
std::shared_ptr< ShellCombinationsIterator > ShellCombinationsIterator::build
(
    const psi::IntegralFactory & ints,
    std::string mode = "ALL",
    int nshell = 4 ) [static]
```

Parameters

<i>ints</i>	- integral factory
<i>mode</i>	- mode of iteration (either ALL or UNIQUE)
<i>nshell</i>	- number of shells to iterate through

Returns

shell iterator

18.104.3.4 compute_shell() [1/2]

```
void ShellCombinationsIterator::compute_shell (
    std::shared_ptr< oepdev::TwoBodyAOInt > tei ) const [pure virtual]
```

Compute integrals in a current shell. Works both for [oepdev::TwoBodyAOInt](#) and [psi::TwoBodyAOInt](#)

Parameters

<i>tei</i>	- two body integral object
------------	----------------------------

Implemented in [oepdev::AllAOShellCombinationsIterator_2](#), and [oepdev::AllAOShellCombinationsIterator_4](#).

18.104.3.5 compute_shell() [2/2]

```
void ShellCombinationsIterator::compute_shell (
```



```
std::shared_ptr< psi ::TwoBodyAOInt > tei ) const [pure virtual]
```

Compute integrals in a current shell. Works both for [oepdev::TwoBodyAOInt](#) and [psi::TwoBodyAOInt](#)

Parameters

<i>tei</i>	- two body integral object
------------	----------------------------

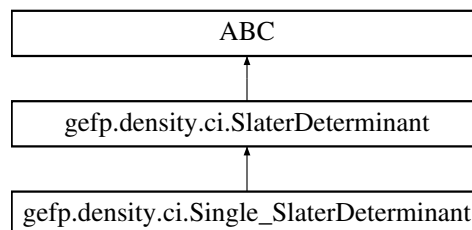
Implemented in [oepdev::AllAOShellCombinationsIterator_4](#).

The documentation for this class was generated from the following files:

- [oepdev/libutil/integrals_iter.h](#)
- [oepdev/libutil/integrals_iter.cc](#)

18.105 gefp.density.ci.Single_SlaterDeterminant Class Reference

Inheritance diagram for [gefp.density.ci.Single_SlaterDeterminant](#):



Public Member Functions

- `def __init__ (self, nao, nbo, nmo, rule)`

Public Attributes

- `is_single`
- `change_alpha`

The documentation for this class was generated from the following file:

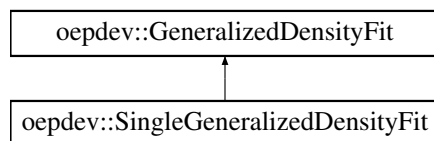
- [gefp/gefp/density/ci.py](#)

18.106 oepdev::SingleGeneralizedDensityFit Class Reference

Generalized Density Fitting Scheme - Single Fit.

```
#include <oep_gdf.h>
```

Inheritance diagram for oepdev::SingleGeneralizedDensityFit:



Public Member Functions

- **SingleGeneralizedDensityFit** (std::shared_ptr< psi::BasisSet > bs_auxiliary, std::shared_ptr< psi::Matrix > v_vector)
- std::shared_ptr< psi::Matrix > **compute** (void)

Perform the generalized density fit.

Additional Inherited Members

18.106.1 Detailed Description

The density fitting map projects the OEP onto the auxiliary, nearly complete basis set space through application of the resolution of identity. Refer to [density fitting in complete space](#) for more details.

18.106.2 Determination of the OEP matrix

Coefficients \mathbf{G} are computed by using the following relation

$$\mathbf{G}^{(i)} = \mathbf{v}^{(i)} \cdot \mathbf{S}^{-1}$$

where

$$S_{\xi\eta} = (\xi|\eta)$$

$$v_{\xi}^{(i)} = (\xi|\hat{v}i)$$

In the above, $|\cdot\rangle$ denotes the single integration over electron coordinate, i.e.,

$$(a|b) \equiv \int d\mathbf{r} \phi_a^*(\mathbf{r}) \phi_b(\mathbf{r})$$

whereas the spatial form of the potential operator \hat{v} can be expressed by

$$v(\mathbf{r}) \equiv \int d\mathbf{r}' \frac{\rho(\mathbf{r}')}{|\mathbf{r}' - \mathbf{r}|}$$

with $\rho(\mathbf{r})$ being the effective one-electron density associated with \hat{v} .

18.106.3 Member Function Documentation

18.106.3.1 compute()

```
std::shared_ptr< psi::Matrix > SingleGeneralizedDensityFit::compute (
    void ) [virtual]
```

Returns

The OEP coefficients G_{ξ_i}

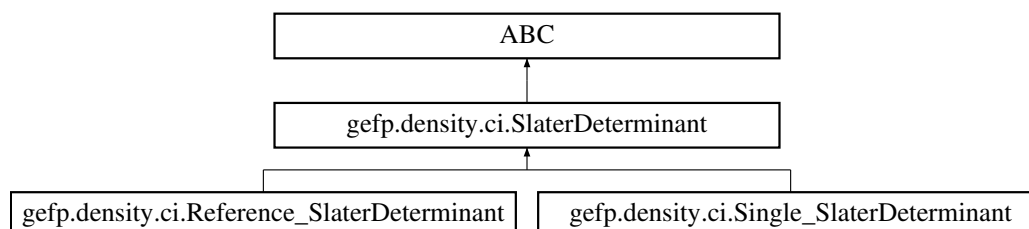
Implements [oepdev::GeneralizedDensityFit](#).

The documentation for this class was generated from the following files:

- oepdev/liboep/oep_gdf.h
- oepdev/liboep/oep_gdf.cc

18.107 gefp.density.ci.SlaterDeterminant Class Reference

Inheritance diagram for gefp.density.ci.SlaterDeterminant:



Public Member Functions

- `def __init__ (self, nao, nbo, nmo, rule)`

Public Attributes

- `is_reference`
- `is_single`
- `is_double`
- `is_triple`
- `rule`
- `nao`
- `nbo`

- **nav**
- **nbv**
- **nmo**

The documentation for this class was generated from the following file:

- [gefp/gefp/density/ci.py](#)

18.108 oepdev::GeneralizedPolarGEFactory::StatisticalSet Struct Reference

A structure to handle statistical data.

```
#include <gefp.h>
```

Public Attributes

- `std::vector< double >` [InducedInteractionEnergySet](#)
Interaction energy set.
- `std::vector< std::shared_ptr< psi::Matrix > >` [DensityMatrixSet](#)
Density matrix set.
- `std::vector< std::shared_ptr< psi::Vector > >` [InducedDipoleSet](#)
Induced dipole moment set.
- `std::vector< std::shared_ptr< psi::Vector > >` [InducedQuadrupoleSet](#)
Induced quadrupole moment set.
- `std::vector< std::shared_ptr< psi::Matrix > >` [JKMatrixSet](#)
Sum of J and K matrix set.

The documentation for this struct was generated from the following file:

- [oepdev/libgefp/gefp.h](#)

18.109 gefp.math.matrix.Superimposer Class Reference

Public Member Functions

- `def __init__ (self)`
SVDSuperimposer from BIOPYTHON PACKAGE Copyright (C) 2002, Thomas Hamelryck (thamelry@vub.ac.be) This code is part of the Biopython distribution and governed by its license.
- `def set (self, reference_coords, coords)`

- def **run** (self)
- def **get_transformed** (self)
- def **get_rotran** (self)
- def **get_init_rms** (self)
- def **get_rms** (self)

Public Attributes

- **reference_coords**
- **coords**
- **transformed_coords**
- **rot**
- **tran**
- **rms**
- **init_rms**
- **n**

18.109.1 Detailed Description

\

SVDSuperimposer finds the best rotation and translation to put two point sets on top of each other (minimizing the RMSD). This is eg. useful to superimpose crystal structures.

SVD stands for Singular Value Decomposition, which is used to calculate the superposition.

Reference:

Matrix computations, 2nd ed. Golub, G. & Van Loan, CF., The Johns Hopkins University Press, Baltimore, 1989

18.109.2 Constructor & Destructor Documentation

18.109.2.1 __init__()

```
def gefp.math.matrix.Superimposer.__init__ (
    self )
```

Please see the LICENSE file that should have been included as part of this package.

18.109.3 Member Function Documentation

18.109.3.1 set()

```
def gefp.math.matrix.Superimposer.set (
    self,
    reference_coords,
    coords )
```

Set the coordinates to be superimposed.
 coords will be put on top of reference_coords.

o reference_coords: an NxDIM array
 o coords: an NxDIM array

DIM is the dimension of the points, N is the number
 of points to be superimposed.

The documentation for this class was generated from the following file:

- gefp/gefp/math/matrix.py

18.110 oepdev::test::Test Class Reference

Manages test routines.

```
#include <test.h>
```

Public Member Functions

- [Test](#) (std::shared_ptr< psi::Wavefunction > wfn, psi::Options &options)
Construct the tester.
- [~Test](#) ()
Destructor.
- double [run](#) (void)
Peform the test.

Protected Member Functions

- double [test_basic](#) (void)
Test the basic functionalities of OEPEv.
- double [test_cis_rhf](#) (void)
Test the CIS(RHF) method.
- double [test_cis_uhf](#) (void)
Test the CIS(UHF) method.
- double [test_cphf](#) (void)

- *Test the CPHF method.*
- double `test_dmatPol` (void)
- *Test the density matrix susceptibility ($X = 1$)*
- double `test_dmatPolX` (void)
- *Test the density matrix susceptibility.*
- double `test_eri_1_1` (void)
- *Test the oepdev::ERI_1_1 class against psi::ERI.*
- double `test_eri_2_2` (void)
- *Test the oepdev::ERI_2_2 class against psi::ERI.*
- double `test_eri_3_1` (void)
- *Test the oepdev::ERI_3_1 class against psi::ERI.*
- double `test_unitaryOptimizer` (void)
- *Test the oepdev::UnitaryOptimizer class.*
- double `test_unitaryOptimizer_4_2` (void)
- *Test the oepdev::UnitaryOptimizer_4_2 class.*
- double `test_scf_perturb` (void)
- *Test the oepdev::RHFPerturbed class.*
- double `test_camm` (void)
- *Test the oepdev::CAMM class.*
- double `test_dmt energy` (void)
- *Test the oepdev::DMTP class for energy calculations.*
- double `test_esp_solver` (void)
- *Test the oepdev::ESPSolver.*
- double `test_points_collection3d` (void)
- *Test the cube file generation (oepdev::Field3D electrostatic potential and oepdev::Points3DIterator for cube collection)*
- double `test_ct_energy_benchmark_ol` (void)
- *Test the Charge-transfer Energy Solver (benchmark method Otto-Ladik)*
- double `test_rep_energy_benchmark_hs` (void)
- *Test the Repulsion Energy Solver: (benchmark method Hayes-Stone)*
- double `test_rep_energy_benchmark_dds` (void)
- *Test the Repulsion Energy Solver: (benchmark method Density-Based - DDS/HF)*
- double `test_rep_energy_benchmark_murrell_etal` (void)
- *Test the Repulsion Energy Solver: (benchmark method Murrell-etal)*
- double `test_rep_energy_oep_based_murrell_etal` (void)
- *Test the Repulsion Energy Solver: (OEP-based method Murrell-etal)*
- double `test_rep_energy_benchmark_ol` (void)
- *Test the Repulsion Energy Solver: (benchmark method Otto-Ladik)*
- double `test_rep_energy_benchmark_efp2` (void)
- *Test the Repulsion Energy Solver: (benchmark method EFP2)*
- double `test_custom` (void)
- *Test the custom code (to be deprecated)*

Protected Attributes

- `std::shared_ptr< psi::Wavefunction > wfn_`
Wavefunction object.
- `psi::Options & options_`
Psi4 Options.

The documentation for this class was generated from the following files:

- `oepdev/libtest/test.h`
- `oepdev/libtest/basic.cc`
- `oepdev/libtest/camm.cc`
- `oepdev/libtest/cis_rhf.cc`
- `oepdev/libtest/cis_uhf.cc`
- `oepdev/libtest/cphf.cc`
- `oepdev/libtest/ct_energy_benchmark_ol.cc`
- `oepdev/libtest/dmatpol.cc`
- `oepdev/libtest/dmatpolX.cc`
- `oepdev/libtest/dmtp_energy.cc`
- `oepdev/libtest/eri_1_1.cc`
- `oepdev/libtest/eri_2_2.cc`
- `oepdev/libtest/eri_3_1.cc`
- `oepdev/libtest/esp_solver.cc`
- `oepdev/libtest/points_collection3d.cc`
- `oepdev/libtest/rep_energy_benchmark_dds.cc`
- `oepdev/libtest/rep_energy_benchmark_efp2.cc`
- `oepdev/libtest/rep_energy_benchmark_hs.cc`
- `oepdev/libtest/rep_energy_benchmark_murrell_etal.cc`
- `oepdev/libtest/rep_energy_benchmark_ol.cc`
- `oepdev/libtest/rep_energy_oep_based_murrell_etal.cc`
- `oepdev/libtest/scf_perturb.cc`
- `oepdev/libtest/test.cc`
- `oepdev/libtest/test_custom.cc`
- `oepdev/libtest/unitary_optimizer.cc`
- `oepdev/libtest/unitary_optimizer_4_2.cc`

18.111 oepdev::TIData Class Reference

Transfer Integral EET Data.

```
#include <ti_data.h>
```


Public Member Functions

- [TIData](#) ()
Constructor.
- virtual [~TIData](#) ()
Destructor.
- void [set_s](#) (double, double, double, double, double, double)
Set the overlap integrals between basis states, S_{ij} , for $ij=12,13,14,23,24,34$.
- void [set_e](#) (double, double, double, double)
Set the diagonal exciton Hamiltonian matrix elements E_n for $n=1,2,3,4$.
- void [set_de](#) (double, double)
Set environmental corrections ΔE_1 and ΔE_2 .
- void [set_trcamm_coupling](#) (oepdev::SharedDMTPConvergence)
Set the convergence object for TrCamm-based $V^{\text{Coul},(0)}$.
- virtual double [coupling_trcamm](#) (const std::string &rn)
- virtual double [coupling_direct](#) (void)
- virtual double [coupling_direct_coul](#) (void)
- virtual double [coupling_direct_exch](#) (void)
- virtual double [coupling_indirect](#) (void)
- virtual double [coupling_indirect_ti2](#) (void)
- virtual double [coupling_indirect_ti3](#) (void)
- virtual double [coupling_total](#) (void)
- virtual double [overlap_corrected](#) (const std::string &type)
- virtual double [overlap_corrected_direct](#) (void)
- virtual double [overlap_corrected_direct](#) (double v)
- virtual double [overlap_corrected_indirect](#) (double v, double s)

Public Attributes

- [oepdev::MultipoleConvergence::ConvergenceLevel trcamm_convergence](#)
Convergence object for Coulombic coupling under TrCamm approximation.
- bool [diagonal_correction](#)
Environmental correction activated?
- bool [mulliken_approximation](#)
Mulliken approximation activated?
- bool [overlap_correction](#)
Overlap correction activated?
- bool [trcamm_approximation](#)
TrCamm approximation activated?
- std::map< std::string, double > [v0](#)
- oepdev::SharedDMTPConvergence [v0_trcamm](#)

V0_Coul multipole convergence.

- double [s12](#)
Overlap matrix element between basis functions.
- double [s13](#)
Overlap matrix element between basis functions.
- double [s14](#)
Overlap matrix element between basis functions.
- double [s23](#)
Overlap matrix element between basis functions.
- double [s24](#)
Overlap matrix element between basis functions.
- double [s34](#)
Overlap matrix element between basis functions.

- double [e1](#)
Diagonal Hamiltonian matrix element.
- double [e2](#)
Diagonal Hamiltonian matrix element.
- double [e3](#)
Diagonal Hamiltonian matrix element.
- double [e4](#)
Diagonal Hamiltonian matrix element.

- double [de1](#)
Environmental correction to the E_n for $n=1,2$.
- double [de2](#)
Environmental correction to the E_n for $n=1,2$.

Protected Attributes

- double [c_](#)
Conversion factor (unused now)

18.111.1 Detailed Description

Container for storing and managing TI data for EET coupling calculations, according to Fujimoto JCP 2012:

- exciton Hamiltonian matrix elements
- overlap integrals between basis states
- TrCamm EET coupling convergence object

Contains useful methods to process exciton Hamiltonian matrix elements:

- compute direct and indirect EET coupling constants
- compute overlap-corrected exciton Hamiltonian off-diagonal matrix elements
- include or exclude environmental correction in the diagonal exciton Hamiltonian
- activate TrCamm approximation of $V^{\text{Coul},(0)}$
- activate Mulliken approximation for $V^{\text{Exch},(0)}$ and $V^{\text{CT},(0)}$

To activate/deactivate the various approximations and corrections listed above, set the following attributes

- `diagonal_correction`
- `mulliken_approximation`
- `overlap_correction`
- `trcamm_approximation`

to `true/false`, according to your need.

Example of usage.

```
{c++}
// Set up exciton Hamiltonian
TIData data = TIData();
data.set_s(S12, S13, S14, S32, S42, S34);
data.set_e(E1, E2, E3, E4);
data.set_de(E1 - E01, E2 - E02);
data.v0["COUL"] = V0_Coul;
data.v0["EXCH"] = V0_Exch;
data.v0["ET1"] = V0_ET1;
data.v0["ET2"] = V0_ET2;
data.v0["HT1"] = V0_HT1;
data.v0["HT2"] = V0_HT2;
data.v0["CT"] = V0_CT;
data.v0["EXCH.M"] = V0_Exch.M;
data.v0["CT.M"] = V0_CT.M;

// Set up approximations and corrections
data.diagonal_correction = true;
data.mulliken_approximation = false;
```

```

data.trcamm_approximation = false;
data.overlap_correction = true;

// Compute overlap-corrected indirect coupling matrix elements
double V_ET1 = data.overlap_corrected("ET1");
double V_ET2 = data.overlap_corrected("ET2");
double V_HT1 = data.overlap_corrected("HT1");
double V_HT2 = data.overlap_corrected("HT2");
double V_CT = data.overlap_corrected("CT");
double V_CTM = data.overlap_corrected("CTM");

// Compute final coupling contributions
double V_Coul = data.overlap_corrected("COUL");
double V_Exch = data.overlap_corrected("EXCH");
double V_Ovrl = data.overlap_corrected("OVRL");

double V_ExchM = data.overlap_corrected("EXCH.M");

double V_TI_2 = data.coupling_indirect_ti2();
double V_TI_3 = data.coupling_indirect_ti3();

data.diagonal_correction = false;
double V0_TI_2 = data.coupling_indirect_ti2();
double V0_TI_3 = data.coupling_indirect_ti3();

data.mulliken_approximation = true;
double V0_TI_3_M = data.coupling_indirect_ti3();
data.diagonal_correction = true;
double V_TI_3_M = data.coupling_indirect_ti3();

double V_direct = V_Coul + V_Exch + V_Ovrl;
double V_indirect = V_TI_2 + V_TI_3;

```

See also

[oepdev::EETCouplingSolver](#)

18.111.2 Member Function Documentation

18.111.2.1 coupling_direct()

```
double TIData::coupling_direct (
    void ) [virtual]
```

Compute the direct EET coupling constant.

Returns

$$V^{\text{Dir}} = V^{\text{Coul}} + V^{\text{Exch}} + V^{\text{Ovrl}}$$

Overlap and diagonal corrections as well as TrCamm and Mulliken approximations for Coulomb and pure exchange parts can be set.

18.111.2.2 coupling_direct_coul()

```
double TIData::coupling_direct_coul (
    void ) [virtual]
```

Compute the direct EET coupling constant in Forster limit (Coulombic approximation)

Returns

$$V^{\text{Dir}} = V^{\text{Coul}}$$

Overlap correction as well as TrCamm approximation for Coulomb coupling can be set.

18.111.2.3 coupling_direct_exch()

```
double TIData::coupling_direct_exch (
    void ) [virtual]
```

Compute the direct EET coupling constant due to pure exchange.

Returns

$$V^{\text{Dir}} = V^{\text{Exch}}$$

Overlap correction as well Mulliken approximation for pure exchange coupling can be set.

18.111.2.4 coupling_indirect()

```
double TIData::coupling_indirect (
    void ) [virtual]
```

Compute the indirect EET coupling constant.

Returns

$$V^{\text{Indir}} = V^{\text{TI},(2)} + V^{\text{TI},(3)}$$

Overlap and diagonal corrections as well as Mulliken approximations for $V^{\text{CT},(0)}$ can be set.

18.111.2.5 coupling_indirect_ti2()

```
double TIData::coupling_indirect_ti2 (
    void ) [virtual]
```

Compute the indirect EET coupling constant in second-order with respect to TI.

Returns

$$V^{\text{TI},(2)} = -\frac{V^{\text{ET1}}V^{\text{HT2}}}{E_3-E_1} - \frac{V^{\text{ET2}}V^{\text{HT1}}}{E_4-E_1}$$

Overlap and diagonal corrections can be set.

18.111.2.6 coupling_indirect_ti3()

```
double TIData::coupling_indirect_ti3 (
    void ) [virtual]
```

Compute the indirect EET coupling constant in third-order with respect to TI.

Returns

$$V^{\text{TI},(3)} = \frac{V^{\text{CT}}(V^{\text{ET}1}V^{\text{ET}2} + V^{\text{HT}1}V^{\text{HT}2})}{(E_3 - E_1)(E_4 - E_1)}$$

Overlap and diagonal corrections as well as Mulliken approximations for $V^{\text{CT},(0)}$ can be set.

18.111.2.7 coupling_total()

```
double TIData::coupling_total (
    void ) [virtual]
```

Compute the *total* EET coupling constant.

Returns

$$V^{\text{Total}} = V^{\text{Dir}} + V^{\text{Indir}}$$

Overlap and diagonal corrections, TrCamm approximation for Coulomb coupling, and Mulliken approximations for pure exchange coupling and $V^{\text{CT},(0)}$ can be set.

18.111.2.8 coupling_trcamm()

```
double TIData::coupling_trcamm (
    const std::string & rn ) [virtual]
```

Compute Coulombic coupling approximated by TrCamm.

Parameters

<i>rn</i>	- convergence of TrCamm coupling. Can be from R1 to R5, which corresponds to the R^{-n} series expansion of distributed multipoles.
-----------	---

Returns

$$V^{\text{Coul},(0)} \approx V^{\text{TrCamm},(0)}(R^{-n})$$

18.111.2.9 overlap_corrected()

```
double TIData::overlap_corrected (
    const std::string & type ) [virtual]
```

Compute overlap corrected matrix elements.

Parameters

<i>type</i>	<p>- matrix element $V^{\text{type},(0)}$ subject to overlap correction, where type is one of the following:</p> <ul style="list-style-type: none"> • COUL - $V^{\text{Coul},(0)}$, • EXCH - $V^{\text{Exch},(0)}$, • TrCamm_R1 - $V^{\text{TrCamm},(0)}(R^{-1})$, • TrCamm_R2 - $V^{\text{TrCamm},(0)}(R^{-2})$, • TrCamm_R3 - $V^{\text{TrCamm},(0)}(R^{-3})$, • TrCamm_R4 - $V^{\text{TrCamm},(0)}(R^{-4})$, • TrCamm_R5 - $V^{\text{TrCamm},(0)}(R^{-5})$, • ET1 - $V^{\text{ET1},(0)}$, • ET2 - $V^{\text{ET2},(0)}$, • HT1 - $V^{\text{HT1},(0)}$, • HT2 - $V^{\text{HT2},(0)}$, • CT - $V^{\text{CT},(0)}$, • CT_M - Mulliken-approximated $V^{\text{CT},(0)}$, • EXCH_M - Mulliken-approximated $V^{\text{Exch},(0)}$.
-------------	---

If type = OVRL, the overlap-correction to the direct EET coupling constant is returned, V^{Ovrl} .

Returns

overlap-corrected exciton Hamiltonian matrix element contribution of selected type

Diagonal correction can be set.

18.111.2.10 overlap_corrected_direct() [1/2]

```
double TIData::overlap_corrected_direct (
    void ) [virtual]
```

Compute overlap-corrected direct EET coupling constant.

Returns

$$V^{\text{Dir}} = V^{\text{Coul}} + V^{\text{Exch}} + V^{\text{Ovr}}$$

Diagonal correction, TrCAMM approximation and Mulliken approximation can be set.

18.111.2.11 overlap_corrected_direct() [2/2]

```
double TIData::overlap_corrected_direct (
    double v ) [virtual]
```

Compute overlap-corrected direct EET coupling constant from value v .

Returns

$$\frac{v}{1-s_{12}^2}$$

18.111.2.12 overlap_corrected_indirect()

```
double TIData::overlap_corrected_indirect (
    double v,
    double s ) [virtual]
```

Compute overlap-corrected coupling constant from value v and associated overlap integral s .

Returns

$$\frac{1}{1-s^2} \left(v - \frac{(E_1+E_2)s}{2} \right)$$

Diagonal correction can be set.

18.111.3 Member Data Documentation**18.111.3.1 v0**

```
std::map<std::string, double> oepdev::TIData::v0
```

Dictionary of all zeroth-order off-diagonal matrix elements.

Use only the following keywords:

- COUL - $V^{\text{Coul},(0)}$,
- EXCH - $V^{\text{Exch},(0)}$,
- TrCAMM_R1 - $V^{\text{TrCAMM},(0)}(R^{-1})$,

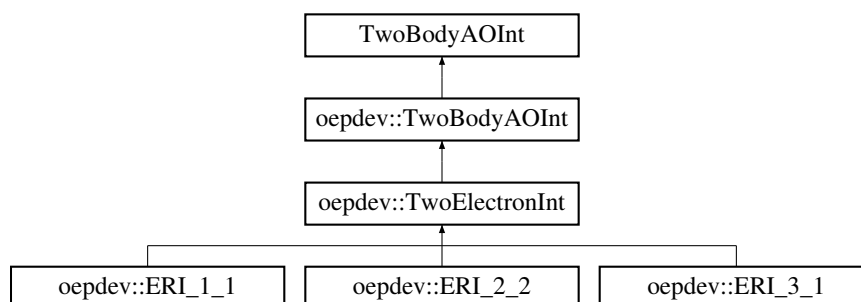
- $\text{TrCAMM_R2} - V^{\text{TrCAMM},(0)}(R^{-2})$,
- $\text{TrCAMM_R3} - V^{\text{TrCAMM},(0)}(R^{-3})$,
- $\text{TrCAMM_R4} - V^{\text{TrCAMM},(0)}(R^{-4})$,
- $\text{TrCAMM_R5} - V^{\text{TrCAMM},(0)}(R^{-5})$,
- $\text{ET1} - V^{\text{ET1},(0)}$,
- $\text{ET2} - V^{\text{ET2},(0)}$,
- $\text{HT1} - V^{\text{HT1},(0)}$,
- $\text{HT2} - V^{\text{HT2},(0)}$,
- $\text{CT} - V^{\text{CT},(0)}$,
- CT_M - Mulliken-approximated $V^{\text{CT},(0)}$,
- EXCH_M - Mulliken-approximated $V^{\text{Exch},(0)}$,
- OVR - V^{Ovr} .

The documentation for this class was generated from the following files:

- oepdev/libsolver/ti_data.h
- oepdev/libsolver/ti_data.cc

18.112 oepdev::TwoBodyAOInt Class Reference

Inheritance diagram for oepdev::TwoBodyAOInt:



Public Member Functions

- virtual void [compute](#) (std::shared_ptr< psi::Matrix > &result, int ibs1=0, int ibs2=2)
Compute two-body two-centre integral and put it into matrix.
- virtual void [compute](#) (psi::Matrix &result, int ibs1=0, int ibs2=2)
- virtual size_t **compute_shell** (int, int, int, int)=0

- virtual size_t **compute_shell** (int, int, int)=0
- virtual size_t **compute_shell** (int, int)=0
- virtual size_t **compute_shell_deriv1** (int, int, int, int)=0
- virtual size_t **compute_shell_deriv2** (int, int, int, int)=0
- virtual size_t **compute_shell_deriv1** (int, int, int)=0
- virtual size_t **compute_shell_deriv2** (int, int, int)=0
- virtual size_t **compute_shell_deriv1** (int, int)=0
- virtual size_t **compute_shell_deriv2** (int, int)=0

Protected Member Functions

- **TwoBodyAOInt** (const [IntegralFactory](#) *intsfactory, int deriv=0)
- **TwoBodyAOInt** (const [TwoBodyAOInt](#) &rhs)

18.112.1 Member Function Documentation

18.112.1.1 compute() [1/2]

```
void oepdev::TwoBodyAOInt::compute (
    std::shared_ptr< psi::Matrix > & result,
    int ibs1 = 0,
    int ibs2 = 2 ) [virtual]
```

Parameters

<i>result</i>	- matrix where to store (i j) two-body integrals
<i>ibs1</i>	- first basis set axis
<i>ibs2</i>	- second basis set axis

18.112.1.2 compute() [2/2]

```
void oepdev::TwoBodyAOInt::compute (
    psi::Matrix & result,
    int ibs1 = 0,
    int ibs2 = 2 ) [virtual]
```

This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

The documentation for this class was generated from the following files:

- oepdev/libpsi/[integral.h](#)

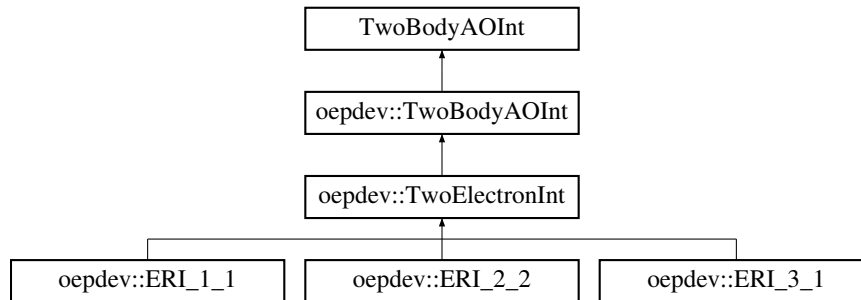
- oepdev/libpsi/integral.cc

18.113 oepdev::TwoElectronInt Class Reference

General Two Electron Integral.

```
#include <eri.h>
```

Inheritance diagram for oepdev::TwoElectronInt:



Public Member Functions

- **TwoElectronInt** (const [IntegralFactory](#) *integral, int deriv, bool use_shell_pairs)
- virtual size_t [compute_shell](#) (int, int)
Compute ERI's between 2 shells. Result is stored in buffer.
- virtual size_t [compute_shell](#) (int, int, int)
Compute ERI's between 3 shells. Result is stored in buffer.
- virtual size_t [compute_shell](#) (int, int, int, int)
Compute ERI's between 4 shells. Result is stored in buffer.
- virtual size_t [compute_shell](#) (const psi::AOShellCombinationsIterator &)
- virtual size_t [compute_shell_deriv1](#) (int, int)
Compute first derivatives of ERI's between 2 shells.
- virtual size_t [compute_shell_deriv2](#) (int, int)
Compute second derivatives of ERI's between 2 shells.
- virtual size_t [compute_shell_deriv1](#) (int, int, int)
Compute first derivatives of ERI's between 3 shells.
- virtual size_t [compute_shell_deriv2](#) (int, int, int)
Compute second derivatives of ERI's between 3 shells.
- virtual size_t [compute_shell_deriv1](#) (int, int, int, int)
Compute first derivatives of ERI's between 4 shells.
- virtual size_t [compute_shell_deriv2](#) (int, int, int, int)
Compute second derivatives of ERI's between 4 shells.

Protected Member Functions

- int [get_cart_am](#) (int am, int n, int x)
Get the angular momentum per Cartesian component.
- double [get_R](#) (int N, int L, int M)
Get the (N,L,M)th McMurchie-Davidson coefficient.
- virtual size_t [compute_doublet](#) (int, int)
Computes the ERI's between three shells.
- virtual size_t [compute_triplet](#) (int, int, int)
Computes the ERI's between three shells.
- virtual size_t [compute_quartet](#) (int, int, int, int)
Computes the ERI's between four shells.

Protected Attributes

- const int [max_am_](#)
Maximum angular momentum.
- const int [n_max_am_](#)
Maximum number of angular momentum functions.
- psi::Fjt * [fjt_](#)
Computes the fundamental: Boys function value at T for degree v.
- bool [use_shell_pairs_](#)
Should we use shell pair information?
- const double [cartMap_](#) [60]
Map of Cartesian components per each am.
- const double [df_](#) [8]
Double factorial array.
- double * [mdh_buffer_R_](#)
Buffer for the McMurchie-Davidson-Hermite R coefficients.

18.113.1 Detailed Description

Implements the McMurchie-Davidson recursive scheme for all integral types. The integral can be defined for any number of Gaussian centres, thus it is not limited to 2-by-2 four-centre ERI. Currently implemented subtypes are:

- [oepdev::ERI_1_1](#) - 2-centre electron-repulsion integral (i|j)
- [oepdev::ERI_2_2](#) - 4-centre electron-repulsion integral (ij|kl)
- [oepdev::ERI_3_1](#) - 4-centre electron-repulsion integral (ijk|l)

See also

[The Integral Package Library](#)

18.113.2 Member Function Documentation

18.113.2.1 compute_shell()

```
size_t oepdev::TwoElectronInt::compute_shell (
    const psi::AOShellCombinationsIterator & shellIter ) [virtual]
```

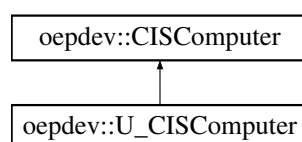
Compute ERIs between 4 shells. Result is stored in buffer. Only for use with [ERI.2.2](#) and the same basis sets, otherwise shell pairs won't be compatible.

The documentation for this class was generated from the following files:

- oepdev/libints/[eri.h](#)
- oepdev/libints/eri.cc

18.114 oepdev::U_CISComputer Class Reference

Inheritance diagram for oepdev::U_CISComputer:



Public Member Functions

- **U_CISComputer** (std::shared_ptr< psi::Wavefunction > wfn, psi::Options &opt)

Protected Member Functions

- virtual void **set_beta_** (void)
- virtual void **build_hamiltonian_** (void)

Additional Inherited Members

The documentation for this class was generated from the following files:

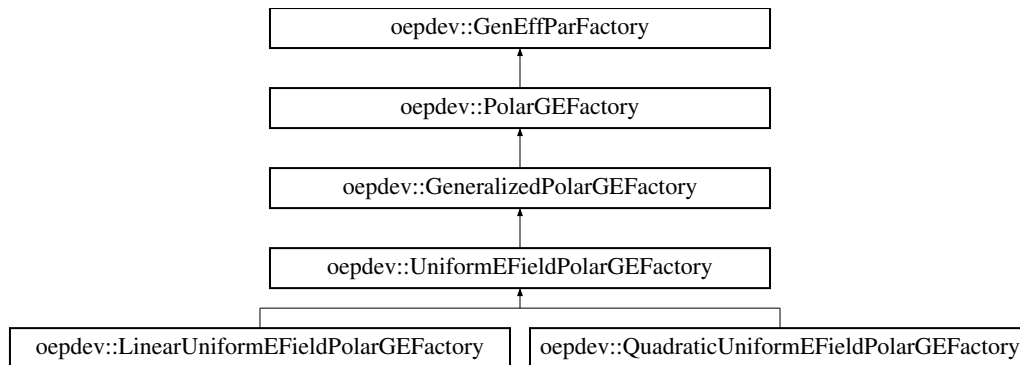
- oepdev/libutil/[cis.h](#)
- oepdev/libutil/cis_uhf.cc

18.115 oepdev::UniformEFieldPolarGEFactory Class Reference

Polarization GEFP Factory with Least-Squares Parameterization.

```
#include <gefp.h>
```

Inheritance diagram for oepdev::UniformEFieldPolarGEFactory:



Public Member Functions

- **UniformEFieldPolarGEFactory** (std::shared_ptr< psi::Wavefunction > [wfn](#), psi::Options &opt)
- void [compute_samples](#) (void)
Compute samples of density matrices and select electric field distributions.
- virtual void [compute_gradient](#) (int i, int j)=0
Compute Gradient vector associated with the i-th and j-th basis set function.
- virtual void [compute_hessian](#) (void)=0
Compute Hessian matrix (independent on the parameters)

Additional Inherited Members

18.115.1 Detailed Description

Implements a class of density matrix susceptibility models for parameterization in the uniform electric field.

The documentation for this class was generated from the following files:

- oepdev/libgefp/[gefp.h](#)
- oepdev/libgefp/gefp_polar_uniform_base.cc

18.116 oepdev::UnitaryOptimizer Class Reference

Find the optimum unitary matrix of quadratic matrix equation.

```
#include <unitary_optimizer.h>
```

Public Member Functions

- [UnitaryOptimizer](#) (double *R, double *P, int n, double conv=1.0e-6, int maxiter=100, bool verbose=true)
Create from R and P matrices and optimization options.
- [UnitaryOptimizer](#) (std::shared_ptr< psi::Matrix > R, std::shared_ptr< psi::Vector > P, double conv=1.0e-6, int maxiter=100, bool verbose=true)
Create from R and P matrices and optimization options.
- [~UnitaryOptimizer](#) ()
Clear memory.
- bool [maximize](#) ()
Run the minimization.
- bool [minimize](#) ()
Run the maximization.
- std::shared_ptr< psi::Matrix > [X](#) ()
Get the unitary matrix (solution)
- double * [get_X](#) () const
Get the unitary matrix (pointer to solution)
- double [Z](#) ()
Get the actual value of Z function.
- bool [success](#) () const
Get the status of the optimization.

Protected Member Functions

- [UnitaryOptimizer](#) (int n, double conv, int maxiter, bool verbose)
Initialize the basic memory.
- void [common_init_](#) ()
Prepare the optimizer.
- void [run_](#) (const std::string &opt)
Run the optimization (intermediate interface)
- void [optimize_](#) (const std::string &opt)
Run the optimization (inner interface)
- void [refresh_](#) ()
Restore the initial state of the optimizer.
- void [update_conv_](#) ()
Update the convergence.
- void [update_iter_](#) ()

- Update the iterates.*

 - void [update_Z_](#) ()

Update Z value.
- void [update_RP_](#) ()

Uptade R and P matrices.
- void [update_X_](#) ()

Update the solution matrix X.
- double [eval_Z_](#) (double *X, double *R, double *P)

Evaluate the objective Z function.
- double [eval_Z_](#) ()
- double [eval_dZ_](#) (double g, double *R, double *P, int i, int j)

Evaluate the change in Z.
- double [eval_Z_trial_](#) (int i, int j, double gamma)

Evaluate the trial Z value.
- void [form_X0_](#) ()

Create identity matrix.
- void [form_X_](#) (int i, int j, double gamma)

Form unitary matrix X (store in buffer Xnew.)
- void [form_next_X_](#) (const std::string &opt)

Form the next unitary matrix X.
- [ABCD](#) [get_ABCD_](#) (int i, int j)

Retrieve [ABCD](#) parameters for root search.
- void [find_roots_boyd_](#) (const [ABCD](#) &abcd)

Solve for all roots of equation $A\sin(g) + B*\cos(g) + C*\sin(2*g) + D*\cos(2*g) = 0$ -> implements Boyd's method.*
- double [find_root_halley_](#) (double x0, const [ABCD](#) &abcd)

Solve for root of equation $A\sin(g) + B*\cos(g) + C*\sin(2*g) + D*\cos(2*g) = 0$ -> implements Halley's method.*
- double [find_gamma_](#) (const [ABCD](#) &abcd, int i, int j, const std::string &opt)

Compute gamma from roots of base equations.
- bool [lt_](#) (double a, double b)

less-than function
- bool [gt_](#) (double a, double b)

greater-than function
- double [func_0_](#) (double g, const [ABCD](#) &abcd)

Function $f(\text{gamma}) = d(dZ)/d\text{gamma}$.
- double [func_1_](#) (double g, const [ABCD](#) &abcd)

Gradient of $f(\text{gamma})$
- double [func_2_](#) (double g, const [ABCD](#) &abcd)

Hessian of $f(\text{gamma})$ - used only for Halley method (not implemented since Boyd method is more suitable here)
- std::shared_ptr< psi::Matrix > [psi_X_](#) ()

Form the Psi4 matrix with the transformation matrix.

Protected Attributes

- const int `n_`
Dimension of the problem.
- const double `conv_`
Convergence.
- const int `maxiter_`
Maximum number of iterations.
- const bool `verbose_`
Verbose mode.
- double * `R_`
R matrix.
- double * `P_`
P vector.
- double * `R0_`
Reference R matrix.
- double * `P0_`
Reference P vector.
- double * `X_`
X Matrix (accumulated solution)
- double * `W_`
Work place.
- double * `Xold_`
Temporary X matrix.
- double * `Xnew_`
Temporary X matrix.
- int `niter_`
Current number of iterations.
- double `S_` [4]
Current solutions.
- double `Zinit_`
Initial Z value.
- double `Zold_`
Old Z value.
- double `Znew_`
New Z value.
- double `conv_current_`
Current convergence.
- bool `success_`
Status of optimization.

18.116.1 Detailed Description

The objective function of the orthogonal matrix \mathbf{X}

$$Z(\mathbf{X}) \equiv \sum_{ijkl} X_{ij} X_{kl} R_{jl} - \sum_{ij} X_{ij} P_j$$

is optimized by using the Jacobi iteration algorithm. In the above equation, \mathbf{R} is a square, general real matrix of size $N \times N$ whereas \mathbf{P} is a real vector of length N .

Algorithm.

Optimization of \mathbf{X} is factorized into a sequence of 2-dimensional rotations with one real parameter γ :

$$\mathbf{X}^{\text{New}} = \mathbf{U}(\gamma) \cdot \mathbf{X}^{\text{Old}}$$

where

$$\mathbf{U}(\gamma) \equiv \begin{pmatrix} \ddots & & & \\ & \cos(\gamma) & \cdots & \sin(\gamma) \\ & \vdots & \ddots & \vdots \\ & -\sin(\gamma) & \cdots & \cos(\gamma) \\ & & & \ddots \end{pmatrix}$$

is the Jacobi transformation matrix constructed for the I th and J th element from the entire N -dimensional set. For the sake of algorithmic simplicity, every iteration after $\mathbf{U}(\gamma)$ has been formed, \mathbf{X}^{Old} is for a while assumed to be an identity matrix and the \mathbf{R} matrix and \mathbf{P} vector are transformed according to the following formulae

$$\begin{aligned} \mathbf{R} &\rightarrow \mathbf{U} \mathbf{R} \mathbf{U}^T \\ \mathbf{P} &\rightarrow \mathbf{U} \mathbf{P} \end{aligned}$$

The full transformation matrix is accumulated in the memory buffer until convergence.

In each iteration, the optimum angle γ is found as follows: First, the roots of the finite Fourier series

$$A \sin(\gamma) + B \cos(\gamma) + C \sin(2\gamma) + D \cos(2\gamma) = 0$$

are found. In the above equations, the expansion coefficients are given as

$$\begin{aligned} A &= P_I + P_J - \sum_{k \neq I, J} (R_{Ik} + R_{Jk} + R_{kI} + R_{kJ}) \\ B &= P_I - P_J - \sum_{k \neq I, J} (R_{Ik} - R_{Jk} + R_{kI} - R_{kJ}) \\ C &= -2(R_{IJ} + R_{JI}) \\ D &= -2(R_{II} - R_{JJ}) \end{aligned}$$

and I, J are the chosen indices in the Jacobi iteration subspace. The roots are evaluated by applying the Boyd's method[1], in which they are given as

$$\gamma_n = \Re[-i \ln(\lambda_n)]$$

where λ_n is an eigenvalue of the following 4 by 4 complex matrix:

$$\begin{pmatrix} 0 & 1 & 0 & 0 \\ 0 & 0 & 1 & 0 \\ 0 & 0 & 0 & 1 \\ -\frac{D+iC}{D-iC} & -\frac{B+iC}{D-iC} & 0 & -\frac{B-iC}{D-iC} \end{pmatrix}$$

Once the four roots of the Fourier series equation are found, one solution out of four is chosen which satisfies the global optimum condition, i.e., the largest increase/decrease in the objective function given by

$$\delta Z = A(1 - \cos(\gamma)) + B \sin(\gamma) + C \sin^2(\gamma) + \frac{D}{2} \sin(2\gamma)$$

The discrimination between the minima/maxima is performed based on the evaluation of the Hessian of Z with respect to γ ,

$$\frac{\partial^2 Z}{\partial \gamma^2} = A \cos(\gamma) - B \sin(\gamma) + 2C \cos(2\gamma) - 2D \sin(2\gamma)$$

All the $N(N-1)/2$ unique pairs of molecular orbitals are checked and the optimal set of γ, I, J is chosen to construct \mathbf{X}^{New} .

References:

[1] Boyd, J.P.; J. Eng. Math. (2006) 56, pp. 203-219

18.116.2 Constructor & Destructor Documentation

18.116.2.1 UnitaryOptimizer() [1/3]

```
oepdev::UnitaryOptimizer::UnitaryOptimizer (
    double * R,
    double * P,
    int n,
    double conv = 1.0e-6,
    int maxiter = 100,
    bool verbose = true )
```

Parameters

<i>R</i>	- R matrix
<i>P</i>	- P vector
<i>n</i>	- dimensionality of the problem (<i>N</i>)
<i>conv</i>	- convergence in the <i>Z</i> function
<i>maxiter</i>	- maximum number of iterations
<i>verbose</i>	- whether print information of iteration process or not Sets up the optimizer.

18.116.2.2 UnitaryOptimizer() [2/3]

```
oepdev::UnitaryOptimizer::UnitaryOptimizer (
    std::shared_ptr< psi::Matrix > R,
    std::shared_ptr< psi::Vector > P,
    double conv = 1.0e-6,
    int maxiter = 100,
    bool verbose = true )
```

Parameters

<i>R</i>	- R matrix
<i>P</i>	- P vector
<i>conv</i>	- convergence in the <i>Z</i> function
<i>maxiter</i>	- maximum number of iterations
<i>verbose</i>	- whether print information of iteration process or not Sets up the optimizer.

18.116.2.3 UnitaryOptimizer() [3/3]

```
oepdev::UnitaryOptimizer::UnitaryOptimizer (
    int n,
    double conv,
    int maxiter,
    bool verbose ) [protected]
```

Parameters

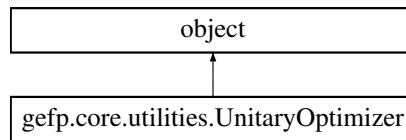
<i>n</i>	- dimensionality of the problem (<i>N</i>)
<i>conv</i>	- convergence in the <i>Z</i> function
<i>maxiter</i>	- maximum number of iterations
<i>verbose</i>	- whether print information of iteration process or not Sets up the optimizer.

The documentation for this class was generated from the following files:

- oepdev/libutil/[unitary_optimizer.h](#)
- oepdev/libutil/unitary_optimizer.cc

18.117 gefp.core.utilities.UnitaryOptimizer Class Reference

Inheritance diagram for gefp.core.utilities.UnitaryOptimizer:



Public Member Functions

- def `__init__` (self, R, P, conv=1.0e-8, maxiter=100, verbose=True)
- def `maximize` (self)
- def `minimize` (self)
- def `run` (self, opt='minimize')
- def `Z` (self)

Public Attributes

- **X**
- **conv**
- **maxiter**
- **verbose**

18.117.1 Detailed Description

Finds the unitary matrix X that optimizes the following function:

$$Z(X) = \sum_{ijkl} X_{ij} X_{kl} R_{jl} - \sum_{ij} X_{ij} P_{ij}$$

where

- * X is a square unitary matrix of size $N \times N$
- * R is a square, in general non-symmetric matrix of size $N \times N$
- * P is a vector of length N

Usage:

```

optimizer = UnitaryOptimizer(R, P, conv=1.0e-8, maxiter=100, verbose=True)
optimizer.maximize() #or minimize()
X = optimizer.X
Z = optimizer.Z
  
```

Last Revision: 25.03.2018

18.117.2 Constructor & Destructor Documentation

18.117.2.1 `__init__()`

```
def gefp.core.utilities.UnitaryOptimizer.__init__ (
    self,
    R,
    P,
    conv = 1.0e-8,
    maxiter = 100,
    verbose = True )
```

Initialize with R and P matrix, as well as optimization options

18.117.3 Member Function Documentation

18.117.3.1 `maximize()`

```
def gefp.core.utilities.UnitaryOptimizer.maximize (
    self )
```

Maximize the Z function under unitary constraint for X

18.117.3.2 `minimize()`

```
def gefp.core.utilities.UnitaryOptimizer.minimize (
    self )
```

Minimize the Z function under unitary constraint for X

18.117.3.3 `run()`

```
def gefp.core.utilities.UnitaryOptimizer.run (
    self,
    opt = 'minimize' )
```

Perform the optimization

18.117.3.4 Z()

```
def gefp.core.utilities.UnitaryOptimizer.Z (
    self )
```

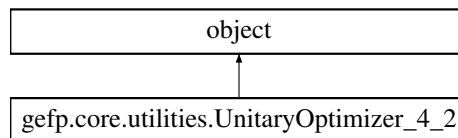
Return the current value of objective function

The documentation for this class was generated from the following file:

- gefp/gefp/core/utilities.py

18.118 gefp.core.utilities.UnitaryOptimizer_4_2 Class Reference

Inheritance diagram for gefp.core.utilities.UnitaryOptimizer_4_2:



Public Member Functions

- def `__init__` (self, R, P, conv=1.0e-8, maxiter=100, verbose=True)
- def `maximize` (self)
- def `minimize` (self)
- def `run` (self, opt='minimize')
- def `Z` (self)

Public Attributes

- **X**
- **conv**
- **maxiter**
- **verbose**

18.118.1 Detailed Description

Finds the unitary matrix X that optimizes the following function:

$$Z(X) = \sum_{ijklmn} X_{\{ki\}} X_{\{lj\}} X_{\{mi\}} X_{\{nj\}} R_{\{ijklmn\}} + \sum_{ijk} X_{\{ji\}} X_{\{ki\}} P_{\{ijk\}}$$

where

- * X is a square unitary matrix of size $N \times N$
- * R is a general real 6-th rank tensor of size N^6
- * P is a general real 3-rd rank tensor of size N^3

Usage:

```
optimizer = UnitaryOptimizer(R, P, conv=1.0e-8, maxiter=100, verbose=True)
optimizer.maximize() #or minimize()
X = optimizer.X
Z = optimizer.Z
```

Last Revision: 07.04.2018

18.118.2 Constructor & Destructor Documentation

18.118.2.1 `__init__()`

```
def gefp.core.utilities.UnitaryOptimizer_4.2.__init__ (
    self,
    R,
    P,
    conv = 1.0e-8,
    maxiter = 100,
    verbose = True )
```

Initialize with R and P matrix, as well as optimization options

18.118.3 Member Function Documentation

18.118.3.1 `maximize()`

```
def gefp.core.utilities.UnitaryOptimizer_4.2.maximize (
    self )
```

Maximize the Z function under unitary constraint for X

18.118.3.2 minimize()

```
def gefp.core.utilities.UnitaryOptimizer_4_2.minimize (
    self )
```

Minimize the Z function under unitary constraint for X

18.118.3.3 run()

```
def gefp.core.utilities.UnitaryOptimizer_4_2.run (
    self,
    opt = 'minimize' )
```

Perform the optimization

18.118.3.4 Z()

```
def gefp.core.utilities.UnitaryOptimizer_4_2.Z (
    self )
```

Return the current value of objective function

The documentation for this class was generated from the following file:

- `gefp/gefp/core/utilities.py`

18.119 oepdev::UnitaryOptimizer_4_2 Class Reference

Find the optimim unitary matrix for quartic-quadratic matrix equation with trace.

```
#include <unitary_optimizer.h>
```

Public Member Functions

- [UnitaryOptimizer_4_2](#) (double *R, double *P, int n, double conv=1.0e-6, int maxiter=100, bool verbose=true)
Create from R and P matrices and optimization options.
- [~UnitaryOptimizer_4_2](#) ()
Clear memory.
- bool [maximize](#) ()

- Run the minimization.*
 - bool `minimize` ()
- Run the maximization.*
 - std::shared_ptr< psi::Matrix > `X` ()
- Get the unitary matrix (solution)*
 - double * `get_X` () const
- Get the unitary matrix (pointer to solution)*
 - double `Z` ()
- Get the actual value of Z function.*
 - bool `success` () const
- Get the status of the optimization.*

Protected Member Functions

- `UnitaryOptimizer_4_2` (int n, double conv, int maxiter, bool verbose)
 - Initialize the basic memory.*
- void `common_init` ()
 - Prepare the optimizer.*
- void `run` (const std::string &opt)
 - Run the optimization (intermediate interface)*
- void `optimize` (const std::string &opt)
 - Run the optimization (inner interface)*
- void `refresh` ()
 - Restore the initial state of the optimizer.*
- void `update_conv` ()
 - Update the convergence.*
- void `update_iter` ()
 - Update the iterates.*
- void `update_Z` ()
 - Update Z value.*
- void `update_RP` ()
 - Uptade R and P matrices.*
- void `update_X` ()
 - Update the solution matrix X.*
- double `eval_Z` (double *X, double *R, double *P)
 - Evaluate the objective Z function.*
- double `eval_Z` ()
- double `eval_dZ` (double g, double *R, double *P, int I, int J)
 - Evaluate the change in Z.*
- double `eval_Z_trial` (int I, int J, double gamma)

- Evaluate the trial Z value.*

 - void [form_X0_](#) ()
- Create identity matrix.*

 - void [form_X_](#) (int I, int J, double gamma)

Form unitary matrix X (store in buffer Xnew_)
- void [form_next_X_](#) (const std::string &opt)

Form the next unitary matrix X.
- [Fourier9](#) [get_fourier_](#) (int I, int J)

Retrieve [ABCD](#) parameters for root search.
- void [find_roots_boyd_](#) (const [Fourier9](#) &abcd)

Solve for all roots of equation $A\sin(g) + B*\cos(g) + C*\sin(2*g) + D*\cos(2*g) = 0$ -> implements Boyd's method.*
- double [find_root_halley_](#) (double x0, const [Fourier9](#) &abcd)

Solve for root of equation $A\sin(g) + B*\cos(g) + C*\sin(2*g) + D*\cos(2*g) = 0$ -> implements Halley's method.*
- double [find_gamma_](#) (const [Fourier9](#) &abcd, int i, int j, const std::string &opt)

Compute gamma from roots of base equations.
- bool [lt_](#) (double a, double b)

less-than function
- bool [gt_](#) (double a, double b)

greater-than function
- std::shared_ptr< [psi::Matrix](#) > [psi_X_](#) ()

Form the Psi4 matrix with the transformation matrix.

Protected Attributes

- const int [n_](#)

Dimension of the problem.
- const double [conv_](#)

Convergence.
- const int [maxiter_](#)

Maximum number of iterations.
- const bool [verbose_](#)

Verbose mode.
- double * [R_](#)

R tensor.
- double * [P_](#)

P tensor.
- double * [R0_](#)

Reference R tensor.
- double * [P0_](#)

- *Reference P tensor.*
- double * [X_](#)
X Matrix (accumulated solution)
- double * [W_](#)
Work place.
- double * [Xold_](#)
Temporary X matrix.
- double * [Xnew_](#)
Temporary X matrix.
- int [niter_](#)
Current number of iterations.
- double [S_](#) [8]
Current solutions.
- double [Zinit_](#)
Initial Z value.
- double [Zold_](#)
Old Z value.
- double [Znew_](#)
New Z value.
- double [conv_current_](#)
Current convergence.
- bool [success_](#)
Status of optimization.

18.119.1 Detailed Description

The objective function of the orthogonal matrix \mathbf{X}

$$Z(\mathbf{X}) \equiv \sum_{ijklmn} X_{ki} X_{lj} X_{mi} X_{nj} R_{ijklmn} + \sum_{ijk} X_{ji} X_{ki} P_{ijk}$$

is optimized by using the Jacobi iteration algorithm. In the above equation, \mathbf{R} is a general real sixth-rank tensor of size N^6 whereas \mathbf{P} is a general real third-rank tensor of size N^3 .

Algorithm.

Optimization of \mathbf{X} is factorized into a sequence of 2-dimensional rotations with one real parameter γ :

$$\mathbf{X}^{\text{New}} = \mathbf{X}^{\text{Old}} \cdot \mathbf{U}(\gamma)$$

where

$$\mathbf{U}(\gamma) \equiv \begin{pmatrix} \ddots & & & & & & & \\ & \cos(\gamma) & \cdots & \sin(\gamma) & & & & \\ & \vdots & \ddots & \vdots & & & & \\ & -\sin(\gamma) & \cdots & \cos(\gamma) & & & & \\ & & & & \ddots & & & \end{pmatrix}$$

is the Jacobi transformation matrix constructed for the I th and J th element from the entire N -dimensional set. For the sake of algorithmic simplicity, every iteration after $\mathbf{U}(\gamma)$ has been formed, \mathbf{X}^{Old} is for a while assumed to be an identity matrix and the \mathbf{R} as well as \mathbf{P} tensors are transformed according to the following formulae

$$R_{ijklmn} \rightarrow \sum_{k'l'm'n'} R_{ijk'l'm'n'} X_{k'k} X_{l'l} X_{m'm} X_{n'n}$$

$$P_{ijk} \rightarrow \sum_{j'k'} P_{ij'k'} X_{j'j} X_{k'k}$$

The full transformation matrix is accumulated in the memory buffer until convergence.

In each iteration, the optimum angle γ is found as follows: First, the roots of the finite Fourier series

$$a_0 + \sum_{p=1}^4 \{a_p \cos(px) + b_p \sin(px)\} = 0$$

are found. In the above equations, the expansion coefficients are calculated analytically as a function of I, J - the chosen indices in the Jacobi iteration subspace. The roots are evaluated by applying the Boyd's method[1], in which they are given as

$$\gamma_n = \Re[-i \ln(\lambda_n)]$$

where λ_n is an eigenvalue of the following 8 by 8 complex matrix:

$$\begin{pmatrix} 0 & 1 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 1 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 1 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 1 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 & 1 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 & 0 & 1 & 0 \\ -\frac{a_4+ib_4}{a_4-ib_4} & -\frac{a_3+ib_3}{a_4-ib_4} & -\frac{a_2+ib_2}{a_4-ib_4} & -\frac{a_1+ib_1}{a_4-ib_4} & -\frac{2a_0}{a_4-ib_4} & -\frac{a_1-ib_1}{a_4-ib_4} & -\frac{a_2-ib_2}{a_4-ib_4} & -\frac{a_3-ib_3}{a_4-ib_4} \end{pmatrix}$$

Once the eight roots of the Fourier series equation are found, one solution out of eight is chosen which satisfies the global optimum condition, i.e., the largest increase/decrease in the objective function given by

$$\delta Z = Z(\mathbf{U}(\gamma)) - Z(\mathbf{1})$$

The Hessian is not computed. All the $N(N-1)/2$ unique pairs of molecular orbitals are checked and the optimal set of γ, I, J is chosen to construct \mathbf{X}^{New} .

References:

[1] Boyd, J.P.; J. Eng. Math. (2006) 56, pp. 203-219

18.119.2 Constructor & Destructor Documentation

18.119.2.1 UnitaryOptimizer_4_2() [1/2]

```
oepdev::UnitaryOptimizer_4_2::UnitaryOptimizer_4_2 (
    double * R,
    double * P,
    int n,
    double conv = 1.0e-6,
    int maxiter = 100,
    bool verbose = true )
```

Parameters

<i>R</i>	- R tensor (flattened row-wise)
<i>P</i>	- P tensor (flattened row-wise)
<i>n</i>	- dimensionality of the problem (<i>N</i>)
<i>conv</i>	- convergence in the <i>Z</i> function
<i>maxiter</i>	- maximum number of iterations
<i>verbose</i>	- whether print information of iteration process or not Sets up the optimizer.

18.119.2.2 UnitaryOptimizer_4_2() [2/2]

```
oepdev::UnitaryOptimizer_4_2::UnitaryOptimizer_4_2 (
    int n,
    double conv,
    int maxiter,
    bool verbose ) [protected]
```

Parameters

<i>n</i>	- dimensionality of the problem (<i>N</i>)
<i>conv</i>	- convergence in the <i>Z</i> function
<i>maxiter</i>	- maximum number of iterations
<i>verbose</i>	- whether print information of iteration process or not Sets up the optimizer.

The documentation for this class was generated from the following files:

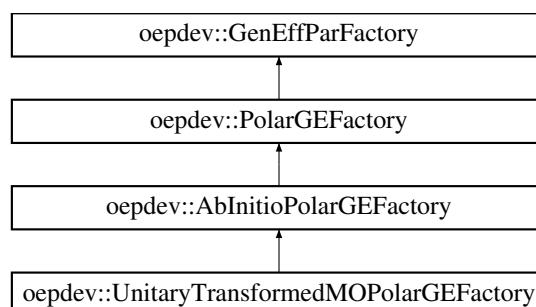
- [oepdev/libutil/unitary_optimizer.h](#)
- [oepdev/libutil/unitary_optimizer.cc](#)

18.120 oepdev::UnitaryTransformedMOPolarGEFactory Class Reference

Polarization GEFP Factory with Least-Squares Scaling of MO Space.

```
#include <gefp.h>
```

Inheritance diagram for oepdev::UnitaryTransformedMOPolarGEFactory:



Public Member Functions

- [UnitaryTransformedMOPolarGEFactory](#) (std::shared_ptr< psi::Wavefunction > wfn, psi::Options &opt)
Construct from CPHF object and Psi4 options.
- virtual [~UnitaryTransformedMOPolarGEFactory](#) ()
Destruct.
- std::shared_ptr< [GenEffPar](#) > [compute](#) (void)
Perform Least-Squares Fit.

Additional Inherited Members

18.120.1 Detailed Description

Implements creation of the density matrix susceptibility tensors for which $\mathbf{X} \neq \mathbf{1}$. Guarantees the idempotency of the density matrix up to first-order in LCAO-MO variation.

Note

This method does not give better results than the $X=1$ method and is extremely time and memory consuming. Therefore, it is placed here only for future reference about solving unitary optimization problem in case it occurs.

The documentation for this class was generated from the following files:

- oepdev/libgefp/[gefp.h](#)
- oepdev/libgefp/gefp_polar_abinitio.cc

18.121 gefp.core.driver.UniversalSurface Class Reference

Static Public Attributes

- string `par_descr_fci_sto3g_1`
- string `par_fulld_fci_sto3g_1`
- `par_pade_fci_sto3g_1` = [PadeApproximant_2D\(\)](#)
- string `par_descr_fci_sto3g_2`
- string `par_fulld_fci_sto3g_2`
- `par_pade_fci_sto3g_2` = [PadeApproximant_2D\(\)](#)
- `par_fci_sto3g_1` = [Entry](#)(`par_pade_fci_sto3g_1`, `par_descr_fci_sto3g_1`, `par_fulld_fci_sto3g_1`)
- `par_fci_sto3g_2` = [Entry](#)(`par_pade_fci_sto3g_2`, `par_descr_fci_sto3g_2`, `par_fulld_fci_sto3g_2`)

18.121.1 Member Data Documentation

18.121.1.1 `par_descr_fci_sto3g_1`

string gefp.core.driver.UniversalSurface.par_descr_fci_sto3g_1 [static]

Initial value:

```
= """\
Universal Surface at FCI/STO-3G level.
Systems: 2el (H2), 4el (H4), 10el (H2O)
"""
```

18.121.1.2 `par_descr_fci_sto3g_2`

string gefp.core.driver.UniversalSurface.par_descr_fci_sto3g_2 [static]

Initial value:

```
= """\
Universal Surface at FCI/STO-3G level.
Systems: 2el (H2), 4el (H4), 10el (H2O)
"""
```


18.121.1.3 par_fulld_fci_sto3g_1

```
string gefp.core.driver.UniversalSurface.par_fulld_fci_sto3g_1 [static]
```

Initial value:

```
= """\
function used for fitting: g(x,y)
g(x,y) = (a0 + a1*x + a2*y + a3*x*y + a4*y*y + a5*y*y*x + a6*y*y*y)/(1.0 + b1*x + b2*y + b3*x*y + b4*y*y
        + b5*y*y*x + b6*y*y*y)

degrees of freedom      (FIT_NDF)                : 47
rms of residuals        (FIT_STDFIT) = sqrt(WSSR/ndf) : 0.0217961
variance of residuals   (reduced chisquare) = WSSR/ndf : 0.00047507

Final set of parameters          Asymptotic Standard Error
=====
a0          = 0.173323          +/- 0.007845      (4.526%)
a1          = -6.85743          +/- 3.626          (52.87%)
a2          = 0.732897          +/- 0.5255          (71.7%)
a3          = -8.10918          +/- 3.951           (48.73%)
a4          = -0.272454          +/- 0.5683          (208.6%)
a5          = 1.73152           +/- 3.307           (191%)
a6          = -0.0591462         +/- 0.08416         (142.3%)
b1          = -68.04            +/- 38.64           (56.8%)
b2          = -3.57443           +/- 3.66           (102.4%)
b3          = 23.0961            +/- 43.09           (186.5%)
b4          = -3.80028           +/- 3.203           (84.29%)
b5          = 30.2229            +/- 27.85           (92.14%)
b6          = -0.269245          +/- 0.328           (121.8%)

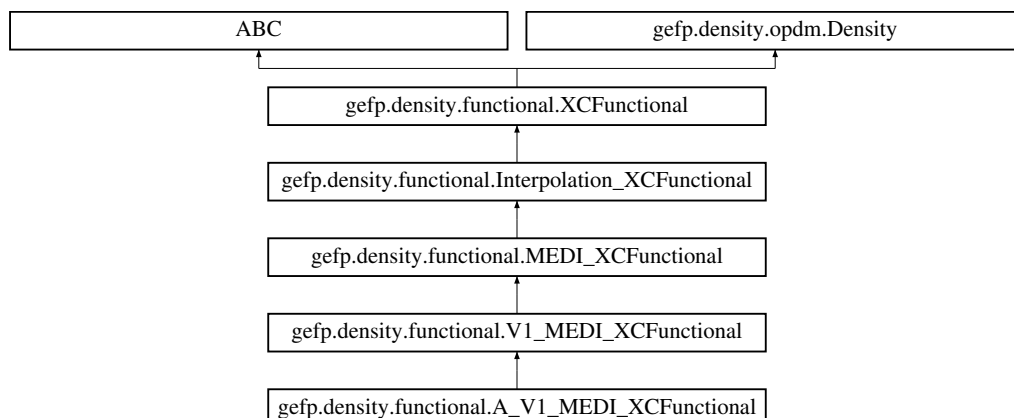
"""
```

The documentation for this class was generated from the following file:

- gefp/gefp/core/driver.py

18.122 gefp.density.functional.V1_MEDI_XCFunctional Class Reference

Inheritance diagram for gefp.density.functional.V1_MEDI_XCFunctional:

**Public Member Functions**

- def `__init__` (self, coeff, kmax)

- def **compute_ak** (self, k, t)
- def **compute_t** (self, n=None, c=None)

Additional Inherited Members

18.122.1 Detailed Description

\

Version 1 of MEDI functional. It is a proper functional of density matrix. However, it cannot be represented explicitly in terms of density matrix.

Functional coefficients:

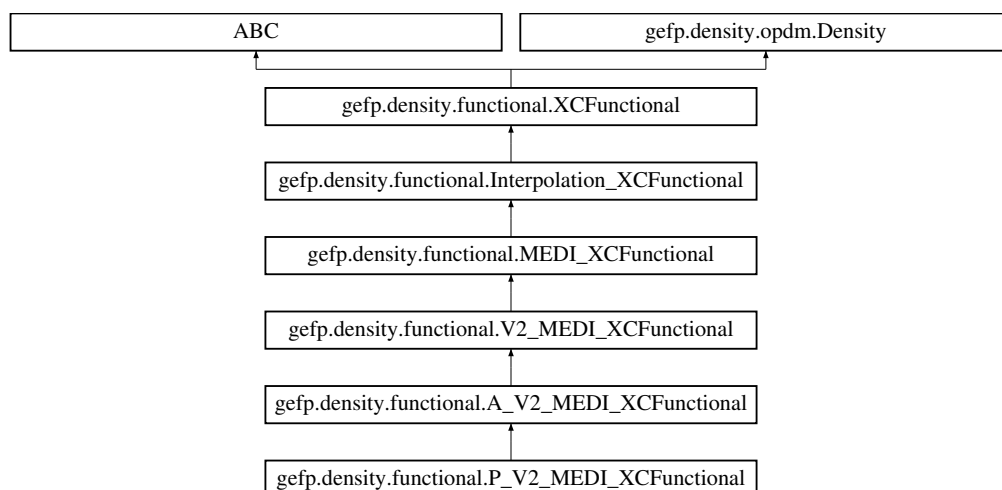
- 't' - t parameter in Gegenbauer Series. Parameter t to be between 0.0 and 1.0. The larger t, the more terms should be included (higher kmax).

The documentation for this class was generated from the following file:

- gefp/gefp/density/functional.py

18.123 gefp.density.functional.V2_MEDI_XCFunctional Class Reference

Inheritance diagram for gefp.density.functional.V2_MEDI_XCFunctional:



Public Member Functions

- def **__init__** (self, coeff, kmax)
- def **compute_ak** (self, k, t)
- def **compute_t** (self, n=None, c=None)
- def **compute_ak_derivative_t** (self, k, t)

Static Public Attributes

- float **parameter_A** = 100.0
- float **parameter_L** = -0.4

Additional Inherited Members

18.123.1 Detailed Description

\

Version 2 of MEDI functional. It is a proper functional of density matrix. Also, it can be represented explicitly in terms of density matrix.

Functional coefficients:

- 't' - t parameter in Gegenbauer Series. Parameter t to be between 0.0 and 1.0. The larger t, the more terms should be included (higher kmax).

The documentation for this class was generated from the following file:

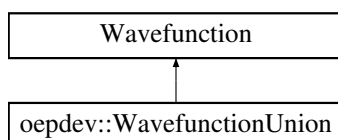
- gefp/gefp/density/functional.py

18.124 oepdev::WavefunctionUnion Class Reference

Union of two Wavefunction objects.

```
#include <wavefunction_union.h>
```

Inheritance diagram for oepdev::WavefunctionUnion:



Public Member Functions

- [WavefunctionUnion](#) (SharedWavefunction ref_wfn, Options &options)

Constructor.

- [WavefunctionUnion](#) (SharedMolecule dimer, SharedBasisSet [primary](#), SharedBasisSet auxiliary_df, SharedBasisSet primary_1, SharedBasisSet primary_2, SharedBasisSet auxiliary_1, SharedBasisSet auxiliary_2, SharedBasisSet auxiliary_df_1, SharedBasisSet auxiliary_df_2, SharedBasisSet intermediate_1, SharedBasisSet intermediate_2, SharedWavefunction wfn_1, SharedWavefunction wfn_2, Options &options)

Constructor.

- virtual [~WavefunctionUnion](#) ()

- Destructor.*
- virtual double `compute_energy` ()
Compute Energy (now blank)
- virtual double `nuclear_repulsion_interaction_energy` ()
Compute Nuclear Repulsion Energy between unions.
- void `localize_orbitals` ()
Localize Molecular Orbitals.
- void `transform_integrals` ()
Transform Integrals (2- and 4-index transformations)
- void `clear_dpd` ()
Close the DPD instance.
- int `L_nmo` (int n) const
*Get number of molecular orbitals of the *n*th fragment.*
- int `L_nso` (int n) const
*Get number of symmetry orbitals of the *n*th fragment.*
- int `L_ndocc` (int n) const
*Get number of doubly occupied orbitals of the *n*th fragment.*
- int `L_nvir` (int n) const
*Get number of virtual orbitals of the *n*th fragment.*
- int `L_nalpha` (int n) const
*Get the number of the alpha electrons of the *n*th fragment.*
- int `L_nbeta` (int n) const
*Get the number of the beta electrons of the *n*th fragment.*
- int `L_nbf` (int n) const
*Get number of basis functions of the *n*th fragment.*
- int `L_noffs_ao` (int n) const
*Get the basis set offset of the *n*th fragment.*
- double `L_energy` (int n) const
*Get the reference energy of the *n*th fragment.*
- SharedMolecule `L_molecule` (int n) const
*Get the molecule object of the *n*th fragment.*
- SharedBasisSet `L_primary` (int n) const
*Get the primary basis set object of the *n*th fragment.*
- SharedBasisSet `L_auxiliary` (int n) const
*Get the auxiliary basis set object of the *n*th fragment.*
- SharedBasisSet `L_intermediate` (int n) const
*Get the intermediate basis set object of the *n*th fragment.*
- SharedWavefunction `L_wfn` (int n) const
*Get the wavefunction object of the *n*th fragment.*
- SharedMOSpace `L_mospace` (int n, const std::string &label) const

- Get the MO space named `label` (either OCC or VIR) of the *n*th fragment.*

 - SharedLocalizer `l_localizer` (int n) const

*Get the orbital localizer object of the *n*th fragment.*
 - SharedIntegralTransform `integrals` (void) const

Get the integral transform object of the entire union.
 - bool `has_localized_orbitals` (void) const

If union got its molecular orbital localized or not.
 - SharedBasisSet `primary` (void) const

Get the primary basis set for the entire union.
 - SharedMOSpace `mospace` (const std::string &label) const

Get the MO space named `label` (either OCC or VIR)
 - SharedMatrix `Ca_subset` (const std::string &basis="SO", const std::string &subset="ALL")
 - SharedMatrix `Cb_subset` (const std::string &basis="SO", const std::string &subset="ALL")
 - SharedMatrix `C_subset_helper` (SharedMatrix C, const Dimension &noccpi, SharedVector epsilon, const std::string &basis, const std::string &subset)
- Helpers for Ca_ and Cb_ matrix transformers.*
- SharedVector `epsilon_subset_helper` (SharedVector epsilon, const Dimension &noccpi, const std::string &basis, const std::string &subset)
- Helper for epsilon transformer.*
- void `print_header` (void)
- Print information about this wavefunction union.*
- void `print_mo_integrals` (void)
- Print the MO ingegrals.*

Protected Attributes

- int `nisolatedMolecules_`
- Number of isolated molecules.*
- SharedWavefunction `dimer_wavefunction_`
- The wavefunction for a dimer (electrons relaxed in the field of monomers)*
- SharedIntegralTransform `integrals_`
- Integral transform object (2- and 4-index transformations)*
- bool `hasLocalizedOrbitals_`
- whether orbitals of the union were localized (or not)*
- std::map< const std::string, SharedMOSpace > `mospacesUnion_`
- Dictionary of MO spaces for the entire union (OCC and VIR)*
- std::vector< SharedMolecule > `l_molecule_`
- List of molecules.*
- std::vector< SharedBasisSet > `l_primary_`
- List of primary basis functions per molecule.*
- std::vector< SharedBasisSet > `l_auxiliary_`

List of auxiliary basis functions per molecule.

- `std::vector< SharedBasisSet >` [I.intermediate_](#)

List of intermediate basis functions per molecule.

- `std::vector< SharedWavefunction >` [I.wfn_](#)

List of original isolated wavefunctions (electrons unrelaxed)

- `std::vector< std::string >` [I.name_](#)

List of names of isolated wavefunctions.

- `std::vector< int >` [I.nbf_](#)

List of basis function numbers per molecule.

- `std::vector< int >` [I.nmo_](#)

List of numbers of molecular orbitals (MO's) per molecule.

- `std::vector< int >` [I.nso_](#)

List of numbers of SO's per molecule.

- `std::vector< int >` [I.ndocc_](#)

List of numbers of doubly occupied orbitals per molecule.

- `std::vector< int >` [I.nvir_](#)

List of numbers of virtual orbitals per molecule.

- `std::vector< int >` [I.noffs_ao_](#)

List of basis set offsets per molecule.

- `std::vector< double >` [I.energy_](#)

List of energies of isolated wavefunctions.

- `std::vector< double >` [I.efzc_](#)

List of frozen-core energies per isolated wavefunction.

- `std::vector< bool >` [I.density_fitted_](#)

List of information per wfn whether it was obtained using DF or not.

- `std::vector< int >` [I.nalpha_](#)

List of numbers of alpha electrons per isolated wavefunction.

- `std::vector< int >` [I.nbeta_](#)

List of numbers of beta electrons per isolated wavefunction.

- `std::vector< int >` [I.nfrzc_](#)

List of numbers of frozen-core orbitals per isolated molecule.

- `std::vector< SharedLocalizer >` [I.localizer_](#)

List of orbital localizers.

- `std::vector< std::map< const std::string, SharedMOSpace > >` [I.mospace_](#)

List of dictionaries of MO spaces.

- `std::shared_ptr< psi::OEProp >` [oeprop_](#)

One-Electron Property.

18.124.1 Detailed Description

The [WavefunctionUnion](#) is the union of two unperturbed Wavefunctions.

Notes:

1. Works only for C1 symmetry! Therefore `this->nirrep() = 1`.
 2. Does not set `reference_wavefunction_`
 3. Sets `oeprop_` for the union of uncoupled molecules
-
1. Performs Hadamard sums on `H_`, `Fa_`, `Da_`, `Ca_` and `S_` based on uncoupled wavefunctions.
 2. Since it is based on shallow copy of the original Wavefunction, it **changes** contents of this wavefunction. Reallocate and copy if you want to keep the original wavefunction.

Warnings:

1. Gradients, Hessians and frequencies are not touched, hence they are **wrong**!
2. Lagrangian (if present) is not touched, hence its **wrong**!
3. `Ca/Cb` and `epsilon` subsets were reimplemented from `psi::Wavefunction` to remove sorting of orbitals. However, the corresponding member functions are not virtual in `psi::Wavefunction`. This could bring problems when upcasting.

The following variables are *shallow* copies of variables inside the Wavefunction object, that is created for the *whole* molecule cluster:

- `basissets_` (DF/RI/F12/etc basis sets)
- `basisset_` (ORBITAL basis set)
- `sobasisset_` (Primary basis set for SO integrals)
- `AO2SO_` (AO2SO conversion matrix (AO in rows, SO in cols))
- `molecule_` (Molecule that this wavefunction is run on)
- `options_` (Options object)
- `psio_` (PSI file access variables)
- `integral_` (Integral factory)
- `factory_` (Matrix factory for creating standard sized matrices)
- `memory_` (How much memory you have access to)
- `nalpha_, nbeta_` (Total alpha and beta electrons)
- `nfrzc_` (Total frozen core orbitals)

- `doccpi_` (Number of doubly occupied per irrep)
- `soccpi_` (Number of singly occupied per irrep)
- `frzcpi_` (Number of frozen core per irrep)
- `frzvpi_` (Number of frozen virtuals per irrep)
- `nalphapi_` (Number of alpha electrons per irrep)
- `nbetapi_` (Number of beta electrons per irrep)
- `nsopi_` (Number of so per irrep)
- `nmopi_` (Number of mo per irrep)
- `nso_` (Total number of SOs)
- `nmo_` (Total number of MOs)
- `nirrep_` (Number of irreps; must be equal to 1 due to symmetry reasons)
- `same_a_b_dens_` and `same_a_b_orbs_` The rest is altered so that the Wavefunction parameters reflect a cluster of non-interacting (uncoupled, isolated, unrelaxed) molecular electron densities.

18.124.2 Constructor & Destructor Documentation

18.124.2.1 WavefunctionUnion() [1/2]

```
oepdev::WavefunctionUnion::WavefunctionUnion (
    SharedWavefunction ref_wfn,
    Options & options )
```

Provide wavefunction with molecule containing at least 2 fragments.

Parameters

<i>ref_wfn</i>	- reference wavefunction
<i>options</i>	- Psi4 options

18.124.2.2 WavefunctionUnion() [2/2]

```
oepdev::WavefunctionUnion::WavefunctionUnion (
    SharedMolecule dimer,
    SharedBasisSet primary,
```



```

SharedBasisSet auxiliary_df,
SharedBasisSet primary_1,
SharedBasisSet primary_2,
SharedBasisSet auxiliary_1,
SharedBasisSet auxiliary_2,
SharedBasisSet auxiliary_df_1,
SharedBasisSet auxiliary_df_2,
SharedBasisSet intermediate_1,
SharedBasisSet intermediate_2,
SharedWavefunction wfn_1,
SharedWavefunction wfn_2,
Options & options )

```

Provide molecule dimer and all the required monomer basis sets and wavefunctions.

Parameters

<i>dimer</i>	- molecule object
<i>primary</i>	- basis set object
<i>auxiliary_df</i>	- basis set object (for DF SCF)
<i>primary_1</i>	- basis set object for 1st monomer
<i>primary_2</i>	- basis set object for 2nd monomer
<i>auxiliary_1</i>	- basis set object for 1st monomer
<i>auxiliary_2</i>	- basis set object for 2nd monomer
<i>auxiliary_df_1</i>	- basis set object for 1st monomer
<i>auxiliary_df_2</i>	- basis set object for 2nd monomer
<i>intermediate_1</i>	- basis set object for 1st monomer
<i>intermediate_2</i>	- basis set object for 2nd monomer
<i>wfn_1</i>	- unperturbed wavefunction object
<i>wfn_2</i>	- unperturbed wavefunction object
<i>options</i>	- Psi4 options

18.124.3 Member Function Documentation

18.124.3.1 Ca_subset()

```

SharedMatrix oepdev::WavefunctionUnion::Ca_subset (
    const std::string & basis = "SO",
    const std::string & subset = "ALL" )

```

Return a subset of the Ca matrix in a desired basis

Parameters

<i>basis</i>	the symmetry basis to use AO, SO
--------------	----------------------------------

Parameters

<i>subset</i>	the subset of orbitals to return ALL, ACTIVE, FROZEN, OCC, VIR, FROZEN_OCC, ACTIVE_OCC, ACTIVE_VIR, FROZEN_VIR
---------------	--

Returns

the matrix in Pitzer order in the desired basis

18.124.3.2 Cb_subset()

```
SharedMatrix oepdev::WavefunctionUnion::Cb_subset (
    const std::string & basis = "SO",
    const std::string & subset = "ALL" )
```

Return a subset of the Cb matrix in a desired basis

Parameters

<i>basis</i>	the symmetry basis to use AO, SO
<i>subset</i>	the subset of orbitals to return ALL, ACTIVE, FROZEN, OCC, VIR, FROZEN_OCC, ACTIVE_OCC, ACTIVE_VIR, FROZEN_VIR

Returns

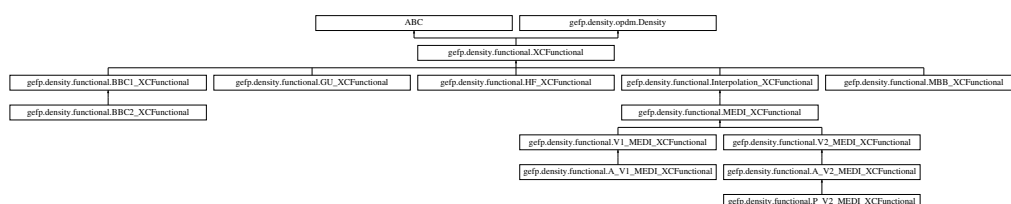
the matrix in Pitzer order in the desired basis

The documentation for this class was generated from the following files:

- oepdev/libutil/[wavefunction_union.h](#)
- oepdev/libutil/wavefunction_union.cc

18.125 gefp.density.functional.XCFunctional Class Reference

Inheritance diagram for gefp.density.functional.XCFunctional:



Public Member Functions

- def **__init__** (self, jk=None, wfn=None, ints=None)
- def **set_wfn** (self, wfn)
- def **set_jk** (self, jk)
- def **set_ints** (self, ints)
- def **create** (cls, name=default, kwargs)
- def **energy_D** (self, x, mode)
- def **energy_P** (self, x)
- def **energy_pc** (self, x)
- def **gradient_D** (self, x)
- def **gradient_P** (self, x)
- def **gradient_P_approximate** (self, x)
- def **gradient_P_numerical** (self, x)
- def **gradient_nc** (self, x)
- def **gradient_pc** (self, x)
- def **abbr** (self)

Static Public Member Functions

- def **name** ()
- def **fij** (n)
- def **fij_1** (n, m)

Static Public Attributes

- string **default** = 'hf'

18.125.1 Detailed Description

\

The Exchange-Correlation DMFT functional.

18.125.2 Member Function Documentation

18.125.2.1 create()

```
def gefp.density.functional.XCFunctional.create (  
    cls,  
    name = default,  
    kwargs )
```

\

Create a density matrix exchange-correlation functional.

Available functionals:

- o 'HF' - the Hartree-Fock functional (default)
- o 'MBB' - the Muller-Buijse-Baerends functional
- o 'GU' - the Goedecker-Urmigar functional
- o 'BBC1' - the BBC1 functional
- o 'BBC2' - the BBC2 functional
- o 'MEDI' - the monotonous exponential decay
of interpolates between MBB and
MBB with zero exchange.
- o 'OEDI' - the oscillatory exponential decay
of interpolates between MBB and
MBB with zero exchange.

Sets:

D, NC

P

P

P

P

P

P

Analytic Derivatives:

Yes

Yes

Approximate

No

No

No

No

The documentation for this class was generated from the following file:

- `gefp/gefp/density/functional.py`

Chapter 19

File Documentation

19.1 include/oepdev_files.h File Reference

Macros

- #define `OEPDEV_USE_PSI4_DIIS_MANAGER` 0
Use DIIS from Psi4 (1) or OEPDev (0)?
- #define `OEPDEV_MAX_AM` 8
L_max.
- #define `OEPDEV_N_MAX_AM` 17
2L_max+1
- #define `OEPDEV_CRIT_ERI` 1e-9
*ERI criterion for E12, E34, E123 and lambda*EXY coefficients.*
- #define `OEPDEV_SIZE_BUFFER_R` 250563
*Size of R buffer ($OEPDEV_N_MAX_AM * OEPDEV_N_MAX_AM * OEPDEV_N_MAX_AM * OEPDEV_N_MAX_AM * 3$)*
- #define `OEPDEV_SIZE_BUFFER_D2` 3264
*Size of D2 buffer ($3 * (OEPDEV_MAX_AM + 1) * (OEPDEV_MAX_AM + 1) * OEPDEV_N_MAX_AM$)*
- #define `OEPDEV_AU_KcalPerMole` 627.509
Energy converters.
- #define `OEPDEV_AU_CMRec` 219474.63
- #define `OEPDEV_AU_EV` 27.21138

19.2 include/oepdev_options.h File Reference

Namespaces

- `psi`
Psi4 package namespace.

Functions

- PSI_API int [psi::read_options](#) (std::string name, Options &options)

Options for the OEPCDev plugin.

19.3 main.cc File Reference

```
#include <string>
#include "include/oepdev_files.h"
#include "psi4/liboptions/liboptions.h"
#include "psi4/libmints/wavefunction.h"
#include "include/oepdev_options.h"
#include "oepdev/liboep/oep.h"
#include "oepdev/libgefp/gefp.h"
#include "oepdev/libsolver/solver.h"
#include "oepdev/libtest/test.h"
#include <pybind11/pybind11.h>
```

Namespaces

- [psi](#)

Psi4 package namespace.

Typedefs

- using **SharedWavefunction** = std::shared_ptr< [psi::Wavefunction](#) >
- using **SharedUnion** = std::shared_ptr< [oepdev::WavefunctionUnion](#) >
- using **SharedOEPotential** = std::shared_ptr< [oepdev::OEPotential](#) >
- using **SharedGEFPFactory** = std::shared_ptr< [oepdev::GenEffParFactory](#) >
- using **SharedGEFPParameters** = std::shared_ptr< [oepdev::GenEffPar](#) >

Functions

- void **psi::export_dmt** (py::module &)
- void **psi::export_cphf** (py::module &)
- void **psi::export_solver** (py::module &)
- void **psi::export_util** (py::module &)
- void **psi::export_oep** (py::module &)
- PSI_API SharedWavefunction [psi::oepdev](#) (SharedWavefunction ref_wfn, Options &options)

Main routine of the OEPCDev plugin.

- **psi::PYBIND11_MODULE** (oepdev, m)

19.4 oepdev/lib3d/dmtp.h File Reference

```
#include "psi4/libmints/wavefunction.h"
#include "psi4/libmints/basisset.h"
#include "psi4/libmints/molecule.h"
#include "psi4/libmints/matrix.h"
#include "psi4/libmints/vector.h"
```

Classes

- class [oepdev::MultipoleConvergence](#)
Multipole Convergence.
- class [oepdev::DMTPole](#)
Distributed Multipole Analysis Container and Computer. Abstract Base.
- class [oepdev::CAMM](#)
Cumulative Atomic Multipole Moments.

Namespaces

- [psi](#)
Psi4 package namespace.
- [oepdev](#)
OEPEv module namespace.

Typedefs

- using [psi::SharedBasisSet](#) = std::shared_ptr< BasisSet >

19.5 oepdev/lib3d/esp.h File Reference

```
#include "space3d.h"
```

Classes

- class [oepdev::ESPSolver](#)
Charges from Electrostatic Potential (ESP). A solver-type class.

Namespaces

- [oepdev](#)

OEPDev module namespace.

Typedefs

- using [oepdev::SharedField3D](#) = std::shared_ptr< [oepdev::Field3D](#) >

19.6 oepdev/libgefp/gefp.h File Reference

```
#include <vector>
#include <string>
#include <random>
#include <cmath>
#include "psi4/libpsi4util/PsiOutStream.h"
#include "psi4/libpsi4util/process.h"
#include "psi4/libmints/wavefunction.h"
#include "psi4/libmints/matrix.h"
#include "psi4/libmints/vector.h"
#include "psi4/libmints/vector3.h"
#include "../liboep/oep.h"
#include "../libutil/util.h"
#include "../libutil/cphf.h"
#include "../libutil/scf_perturb.h"
```

Classes

- class [oepdev::GenEffPar](#)
Generalized Effective Fragment Parameters. Container Class.
- class [oepdev::GenEffFrag](#)
Generalized Effective Fragment. Container Class.
- class [oepdev::GenEffParFactory](#)
Generalized Effective Fragment Factory. Abstract Base.
- class [oepdev::PolarGEFactory](#)
Polarization GEFP Factory. Abstract Base.
- class [oepdev::AbInitioPolarGEFactory](#)
Polarization GEFP Factory from First Principles. Hartree-Fock Approximation.
- class [oepdev::FFAbInitioPolarGEFactory](#)
Polarization GEFP Factory from First Principles: Finite-Difference Model. Arbitrary level of theory.
- class [oepdev::GeneralizedPolarGEFactory](#)

Polarization GEFP Factory with Least-Squares Parameterization.

- struct [oepdev::GeneralizedPolarGEFactory::StatisticalSet](#)

A structure to handle statistical data.

- class [oepdev::UniformEFieldPolarGEFactory](#)

Polarization GEFP Factory with Least-Squares Parameterization.

- class [oepdev::NonUniformEFieldPolarGEFactory](#)

Polarization GEFP Factory with Least-Squares Parameterization.

- class [oepdev::LinearUniformEFieldPolarGEFactory](#)

Polarization GEFP Factory with Least-Squares Parameterization.

- class [oepdev::QuadraticUniformEFieldPolarGEFactory](#)

Polarization GEFP Factory with Least-Squares Parameterization.

- class [oepdev::LinearNonUniformEFieldPolarGEFactory](#)

Polarization GEFP Factory with Least-Squares Parameterization.

- class [oepdev::QuadraticNonUniformEFieldPolarGEFactory](#)

Polarization GEFP Factory with Least-Squares Parameterization.

- class [oepdev::LinearGradientNonUniformEFieldPolarGEFactory](#)

Polarization GEFP Factory with Least-Squares Parameterization.

- class [oepdev::QuadraticGradientNonUniformEFieldPolarGEFactory](#)

Polarization GEFP Factory with Least-Squares Parameterization.

- class [oepdev::UnitaryTransformedMOPolarGEFactory](#)

Polarization GEFP Factory with Least-Squares Scaling of MO Space.

Namespaces

- [oepdev](#)

OEPEDev module namespace.

19.7 oepdev/libints/eri.h File Reference

```
#include "psi4/libpsi4util/exception.h"
#include "psi4/libmints/integral.h"
#include "psi4/libmints/basisset.h"
#include "psi4/libmints/fjt.h"
#include "../libpsi/integral.h"
#include "recurr.h"
```

Classes

- class [oepdev::TwoElectronInt](#)
General Two Electron Integral.
- class [oepdev::ERI_1_1](#)
2-centre ERI of the form $(a|O(2)|b)$ where $O(2) = 1/r^{12}$.
- class [oepdev::ERI_2_2](#)
4-centre ERI of the form $(ab|O(2)|cd)$ where $O(2) = 1/r^{12}$.
- class [oepdev::ERI_3_1](#)
4-centre ERI of the form $(abc|O(2)|d)$ where $O(2) = 1/r^{12}$.

Namespaces

- [oepdev](#)
OEPEv module namespace.

19.8 oepdev/libints/recurr.h File Reference

Namespaces

- [oepdev](#)
OEPEv module namespace.

Macros

- #define [D1_INDEX](#)(x, i, n) $((81*(x))+(9*(i))+(n))$
Get the index of McMurchie-Davidson-Hermite D1 coefficient stored in the `mdh_buffer_`, that is attributed to the x Cartesian coordinate from angular momentum i of function 1, and the Hermite index n.
- #define [D2_INDEX](#)(x, i, j, n) $((1377*(x))+(153*(i))+(17*(j))+(n))$
Get the index of McMurchie-Davidson-Hermite D2 coefficient stored in the `mdh_buffer_`, that is attributed to the x Cartesian coordinate from angular momenta i, j of function 1 and 2, and the Hermite index n.
- #define [D3_INDEX](#)(x, i, j, k, n) $((18225*(x))+(2025*(i))+(225*(j))+(25*(k))+(n))$
Get the index of McMurchie-Davidson-Hermite D3 coefficient stored in the `mdh_buffer_`, that is attributed to the x Cartesian coordinate from angular momenta i, j and k of function 1, 2 and 3, and the Hermite index n.
- #define [R_INDEX](#)(n, l, m, j) $((14739*(n))+(867*(l))+(51*(m))+(j))$
Get the index of McMurchie-Davidson R coefficient stored in the `mdh_buffer_R_` from angular momenta n, l and m and the Boys index j.

Functions

- double [oepdev::d_N_n1_n2](#) (int N, int n1, int n2, double PA, double PB, double aP)
Compute McMurchie-Davidson-Hermite (MDH) coefficient for binomial expansion.
- void [oepdev::make_mdh_D1_coeff](#) (int n1, double aPd, double *buffer)
Compute the McMurchie-Davidson-Hermite coefficients for monomial expansion.
- void [oepdev::make_mdh_D2_coeff](#) (int n1, int n2, double aPd, double *PA, double *PB, double *buffer)
Compute the McMurchie-Davidson-Hermite coefficients for binomial expansion.
- void [oepdev::make_mdh_D3_coeff](#) (int n1, int n2, int n3, double aPd, double *PA, double *PB, double *PC, double *buffer)
Compute the McMurchie-Davidson-Hermite coefficients for trinomial expansion.
- void [oepdev::make_mdh_D2_coeff_explicit_recursion](#) (int n1, int n2, double aP, double *PA, double *PB, double *buffer)
Compute the McMurchie-Davidson-Hermite coefficients for binomial expansion by explicit recursion. This function makes the same changes to buffers as [oepdev::make_mdh_D2_coeff](#), but implements it through explicit recursion by calls to [oepdev::d_N_n1_n2](#). Therefore, it is slightly slower. Here for debugging purposes.
- void [oepdev::make_mdh_R_coeff](#) (int N, int L, int M, double alpha, double a, double b, double c, double *F, double *buffer)
Compute the McMurchie-Davidson R coefficients.

19.9 oepdev/liboep/oep.h File Reference

```
#include <cstdio>
#include <string>
#include <vector>
#include <map>
#include "psi4/liboptions/liboptions.h"
#include "psi4/libpsi4util/PsiOutStream.h"
#include "psi4/libmints/wavefunction.h"
#include "psi4/libmints/basisset.h"
#include "psi4/libmints/matrix.h"
#include "psi4/libmints/vector.h"
#include "psi4/libmints/local.h"
#include "../libutil/cis.h"
#include "../libpsi/integral.h"
#include "../libpsi/potential.h"
#include "../lib3d/space3d.h"
#include "../lib3d/dmtp.h"
```

Classes

- struct [oepdev::OEType](#)

Container to handle the type of One-Electron Potentials.

- class [oepdev::OEPotential](#)

Generalized One-Electron Potential: Abstract base.

- class [oepdev::ElectrostaticEnergyOEPotential](#)

Generalized One-Electron Potential for Electrostatic Energy.

- class [oepdev::RepulsionEnergyOEPotential](#)

Generalized One-Electron Potential for Pauli Repulsion Energy.

- class [oepdev::ChargeTransferEnergyOEPotential](#)

Generalized One-Electron Potential for Charge-Transfer Interaction Energy.

- class [oepdev::EETCouplingOEPotential](#)

Generalized One-Electron Potential for EET coupling calculations.

Namespaces

- [oepdev](#)

OEPDev module namespace.

Typedefs

- using **oepdev::SharedWavefunction** = std::shared_ptr< Wavefunction >
- using **oepdev::SharedBasisSet** = std::shared_ptr< BasisSet >
- using **oepdev::SharedMatrix** = std::shared_ptr< Matrix >
- using **oepdev::SharedVector** = std::shared_ptr< Vector >
- using **oepdev::SharedDMTPole** = std::shared_ptr< DMTPole >
- using **oepdev::SharedLocalizer** = std::shared_ptr< Localizer >
- using **oepdev::SharedCISData** = std::shared_ptr< CISData >

19.10 oepdev/liboep/oep_gdf.h File Reference

```
#include <cstdio>
#include <string>
#include "psi4/liboptions/liboptions.h"
#include "psi4/libmints/wavefunction.h"
#include "psi4/libmints/basisset.h"
#include "psi4/libmints/matrix.h"
#include "../libpsi/integral.h"
```

Classes

- class [oepdev::GeneralizedDensityFit](#)
Generalized Density Fitting Scheme. Abstract Base.
- class [oepdev::SingleGeneralizedDensityFit](#)
Generalized Density Fitting Scheme - Single Fit.
- class [oepdev::DoubleGeneralizedDensityFit](#)
Generalized Density Fitting Scheme - Double Fit.

Namespaces

- [oepdev](#)
OEPEv module namespace.

19.11 oepdev/libpsi/integral.h File Reference

```
#include "psi4/libmints/integral.h"
#include "psi4/libmints/basisset.h"
#include "psi4/libmints/matrix.h"
```

Classes

- class [oepdev::TwoBodyAOInt](#)
- class [oepdev::IntegralFactory](#)
Extended [IntegralFactory](#) for computing integrals.

Namespaces

- [oepdev](#)
OEPEv module namespace.

19.12 oepdev/libpsi/potential.h File Reference

```
#include <vector>
#include "psi4/psi4-dec.h"
#include "psi4/liboptions/liboptions.h"
#include "psi4/libpsio/psio.h"
#include "psi4/libmints/typedefs.h"
#include "psi4/libmints/onebody.h"
#include "psi4/libmints/potential.h"
```

```
#include "psi4/libmints/sointegral_onebody.h"
#include "psi4/libmints/osrecur.h"
```

Classes

- class [oepdev::PotentialInt](#)
Computes potential integrals.

Namespaces

- [oepdev](#)
OEPEv module namespace.

19.13 oepdev/libsolver/solver.h File Reference

```
#include <cstdio>
#include <string>
#include <map>
#include "psi4/psi4-dec.h"
#include "psi4/liboptions/liboptions.h"
#include "psi4/libpsio/psio.h"
#include "psi4/libmints/potential.h"
#include "psi4/libmints/integral.h"
#include "psi4/libpsi4util/PsiOutputStream.h"
#include "../libutil/wavefunction_union.h"
#include "../libutil/integrals_iter.h"
#include "../libpsi/integral.h"
#include "../liboep/oep.h"
#include "../../include/oepdev_files.h"
```

Classes

- class [oepdev::OEPEvSolver](#)
Solver of properties of molecular aggregates. Abstract base.
- class [oepdev::ElectrostaticEnergySolver](#)
Compute the Coulombic interaction energy between unperturbed wavefunctions.
- class [oepdev::RepulsionEnergySolver](#)
Compute the Pauli-Repulsion interaction energy between unperturbed wavefunctions.
- class [oepdev::ChargeTransferEnergySolver](#)
Compute the Charge-Transfer interaction energy between unperturbed wavefunctions.

- class [oepdev::EETCouplingSolver](#)

Compute the EET coupling energy between unperturbed wavefunctions.

Namespaces

- [oepdev](#)

OEPEv module namespace.

Typedefs

- using **oepdev::SharedWavefunctionUnion** = std::shared_ptr< WavefunctionUnion >
- using **oepdev::SharedOEPotential** = std::shared_ptr< OEPotential >

19.14 oepdev/libsolver/ti_data.h File Reference

```
#include <cstdio>
#include <string>
#include <map>
#include "../lib3d/dmtp.h"
```

Classes

- class [oepdev::TIData](#)

Transfer Integral EET Data.

Namespaces

- [oepdev](#)

OEPEv module namespace.

Typedefs

- using **oepdev::SharedDMTPConvergence** = std::shared_ptr< [oepdev::MultipoleConvergence](#) >

19.15 oepdev/libtest/test.h File Reference

```
#include <vector>
#include "psi4/psi4-dec.h"
```

```
#include "psi4/liboptions/liboptions.h"
#include "psi4/libpsio/psio.h"
#include "psi4/libpsi4util/PsiOutStream.h"
#include "psi4/libmints/integral.h"
#include "psi4/libmints/matrix.h"
#include "psi4/libmints/wavefunction.h"
#include "psi4/libmints/basisset.h"
#include "psi4/libqt/qt.h"
#include "../libpsi/integral.h"
#include "../libutil/integrals_iter.h"
#include "../../include/oepdev_files.h"
```

Classes

- class [oepdev::test::Test](#)

Manages test routines.

Namespaces

- [oepdev](#)

OEPEv module namespace.

19.16 oepdev/libutil/cis.h File Reference

```
#include <string>
#include <utility>
#include "psi4/liboptions/liboptions.h"
#include "psi4/libpsio/psio.h"
#include "psi4/libpsio/psio.hpp"
#include "psi4/libmints/matrix.h"
#include "psi4/libmints/vector.h"
#include "psi4/libmints/basisset.h"
#include "psi4/libmints/molecule.h"
#include "psi4/libmints/wavefunction.h"
#include "psi4/libtrans/integraltransform.h"
#include "psi4/libtrans/mospace.h"
#include "psi4/libdpd/dpd.h"
#include "../lib3d/dmtp.h"
```

Classes

- struct [oepdev::CISData](#)

Container to handle the CIS wavefunction parameters.

- class [oepdev::CISComputer](#)
CISComputer.
- class [oepdev::R_CISComputer](#)
- class [oepdev::U_CISComputer](#)

Namespaces

- [oepdev](#)
OEPEv module namespace.

Typedefs

- using **[oepdev::SharedMolecule](#)** = std::shared_ptr< psi::Molecule >
- using **[oepdev::SharedMOSpace](#)** = std::shared_ptr< psi::MOSpace >
- using **[oepdev::SharedMOSpaceVector](#)** = std::vector< std::shared_ptr< psi::MOSpace > >
- using **[oepdev::SharedIntegralTransform](#)** = std::shared_ptr< psi::IntegralTransform >

19.17 oepdev/libutil/diis.h File Reference

```
#include <cstdio>
#include <string>
#include <vector>
#include "psi4/libciomr/libciomr.h"
#include "psi4/libmints/vector.h"
#include "psi4/libmints/matrix.h"
#include "psi4/libqt/qt.h"
#include "psi4/libpsi4util/PsiOutStream.h"
```

Classes

- class [oepdev::DIISManager](#)
DIIS manager.

Namespaces

- [oepdev](#)
OEPEv module namespace.

19.18 oepdev/libutil/integrals_iter.h File Reference

```
#include <cstdio>
#include "psi4/libmints/basisset.h"
#include "psi4/libmints/integral.h"
#include "../libpsi/integral.h"
```

Classes

- class [oepdev::ShellCombinationsIterator](#)
Iterator for Shell Combinations. Abstract Base.
- class [oepdev::AOIntegralsIterator](#)
Iterator for AO Integrals. Abstract Base.
- class [oepdev::AllAOShellCombinationsIterator_4](#)
Loop over all possible ERI shells in a shell quartet.
- class [oepdev::AllAOShellCombinationsIterator_2](#)
Loop over all possible ERI shells in a shell doublet.
- class [oepdev::AllAOIntegralsIterator_4](#)
Loop over all possible ERI within a particular shell quartet.
- class [oepdev::AllAOIntegralsIterator_2](#)
Loop over all possible ERI within a particular shell doublet.

Namespaces

- [oepdev](#)
OEPEv module namespace.

Typedefs

- using **[oepdev::SharedIntegralFactory](#)** = std::shared_ptr< IntegralFactory >
- using **[oepdev::SharedTwoBodyAOInt](#)** = std::shared_ptr< TwoBodyAOInt >
- using [oepdev::SharedShellsIterator](#) = std::shared_ptr< ShellCombinationsIterator >
Iterator over shells as shared pointer.
- using [oepdev::SharedAOIntsIterator](#) = std::shared_ptr< AOIntegralsIterator >
Iterator over AO integrals as shared pointer.

19.19 oepdev/libutil/scf_perturb.h File Reference

```
#include "psi4/libmints/vector.h"
#include "psi4/libmints/matrix.h"
#include "psi4/libscf_solver/rhf.h"
```

Classes

- struct [oepdev::PerturbCharges](#)
Structure to hold perturbing charges.
- class [oepdev::RHFPerturbed](#)
RHF theory under electrostatic perturbation.

Namespaces

- [oepdev](#)
OEPEv module namespace.

19.20 oepdev/libutil/unitary_optimizer.h File Reference

```
#include <string>
#include <complex>
#include "psi4/libmints/vector.h"
#include "psi4/libmints/matrix.h"
```

Classes

- struct [oepdev::ABCD](#)
Simple structure to hold the Fourier series expansion coefficients.
- struct [oepdev::Fourier9](#)
Simple structure to hold the Fourier series expansion coefficients for N=4.
- class [oepdev::UnitaryOptimizer](#)
Find the optimum unitary matrix of quadratic matrix equation.
- class [oepdev::UnitaryOptimizer_4_2](#)
Find the optimum unitary matrix for quartic-quadratic matrix equation with trace.

Namespaces

- [oepdev](#)
OEPEv module namespace.

Macros

- `#define IDX(i, j, n) ((n)*(i)+(j))`
- `#define IDX3(i, j, k) (n2_*(i)+n_*(j)+(k))`
- `#define IDX6(i, j, k, l, m, n) (n5_*(i)+n4_*(j)+n3_*(k)+n2_*(l)+n_*(m)+(n))`

Functions

- `constexpr std::complex< double > oepdev::operator""_i (unsigned long long d)`
- `constexpr std::complex< double > oepdev::operator""_i (long double d)`

19.21 oepdev/libutil/util.h File Reference

```
#include <cstdio>
#include <string>
#include <cmath>
#include <map>
#include <cassert>
#include "psi4/psi4-dec.h"
#include "psi4/liboptions/liboptions.h"
#include "psi4/libpsio/psio.h"
#include "psi4/libciomr/libciomr.h"
#include "psi4/libpsio/psio.hpp"
#include "psi4/libiwl/iwl.h"
#include "psi4/libqt/qt.h"
#include "psi4/libmints/molecule.h"
#include "psi4/libmints/writer.h"
#include "psi4/libmints/writer_file_prefix.h"
#include "psi4/libmints/wavefunction.h"
#include "psi4/libmints/basisset.h"
#include "psi4/libmints/vector.h"
#include "psi4/libmints/matrix.h"
#include "psi4/libmints/oeprop.h"
#include "psi4/libmints/local.h"
#include "psi4/libfunctional/superfunctional.h"
#include "psi4/libtrans/mospace.h"
#include "psi4/libtrans/integraltransform.h"
#include "psi4/libscf_solver/rhf.h"
#include "psi4/libdpd/dpd.h"
```

Namespaces

- [oepdev](#)
OEPEv module namespace.

Typedefs

- using **oepdev::SharedSuperFunctional** = std::shared_ptr< SuperFunctional >

Functions

- PSI.API void **oepdev::preamble** (void)
Print preamble for module OEPDEV.
- template<typename... Args>
std::string **oepdev::string_sprintf** (const char *format, Args... args)
Format string output. Example: std::string text = oepdev::string_sprintf("Test %3d, %13.5f", 5, -10.5425);
- PSI.API std::shared_ptr< SuperFunctional > **oepdev::create_superfunctional** (std::string name, Options &options)
Set up DFT functional.
- PSI.API std::shared_ptr< Molecule > **oepdev::extract_monomer** (std::shared_ptr< const Molecule > molecule_dimer, int id)
Extract molecule from dimer.
- PSI.API double **oepdev::compute_distance** (psi::SharedVector v1, psi::SharedVector v2)
Compute distance between two points in nD space.
- PSI.API std::shared_ptr< Wavefunction > **oepdev::solve_scf** (std::shared_ptr< Molecule > molecule, std::shared_ptr< BasisSet > primary, std::shared_ptr< BasisSet > auxiliary, std::shared_ptr< SuperFunctional > functional, Options &options, std::shared_ptr< PSIO > psio, bool compute_mints=false)
Solve RHF-SCF equations for a given molecule in a given basis set.
- PSI.API std::shared_ptr< Wavefunction > **oepdev::solve_scf_sad** (std::shared_ptr< Molecule > molecule, std::shared_ptr< BasisSet > primary, std::shared_ptr< BasisSet > auxiliary, std::vector< std::shared_ptr< BasisSet >> sad, std::vector< std::shared_ptr< BasisSet >> sad_fit, std::shared_ptr< SuperFunctional > functional, Options &options, std::shared_ptr< PSIO > psio, bool compute_mints=false)
Solve RHF-SCF equations for a given molecule in a given basis set.
- PSI.API double **oepdev::average_moment** (std::shared_ptr< psi::Vector > moment)
Compute the scalar magnitude of multipole moment.
- PSI.API std::vector< std::shared_ptr< psi::Matrix >> **oepdev::calculate_JK** (std::shared_ptr< psi::Wavefunction > wfn, std::shared_ptr< psi::Matrix > C)
Compute the Coulomb and exchange integral matrices in MO basis.
- PSI.API std::vector< std::shared_ptr< psi::Matrix >> **oepdev::calculate_JK_r** (std::shared_ptr< psi::Wavefunction > wfn, std::shared_ptr< psi::IntegralTransform > tr, std::shared_ptr< psi::Matrix > Dij)
Compute the Coulomb and exchange integral matrices in MO basis.
- PSI.API std::shared_ptr< psi::Matrix > **oepdev::calculate_der_D** (std::shared_ptr< psi::Wavefunction > wfn, std::shared_ptr< psi::IntegralTransform > tr, std::shared_ptr< psi::Matrix > C, std::vector< std::shared_ptr< psi::Matrix >> A)

Compute the derivative of exchange-correlation energy wrt the density matrix in MO-A basis.

- PSI_API double [oepdev::calculate_e_xc](#) (std::shared_ptr< psi::Wavefunction > wfn, std::shared_ptr< psi::IntegralTransform > tr, std::shared_ptr< psi::Matrix > f, std::shared_ptr< psi::Matrix > C)

Compute the exchange-correlation energy from ERI in MO-SCF basis.

- PSI_API std::shared_ptr< psi::Matrix > [oepdev::matrix_power_derivative](#) (std::shared_ptr< psi::Matrix > A, double g, double step)

Compute the contracted derivative of power of a square and symmetric matrix.

- std::shared_ptr< psi::Matrix > [oepdev::calculate_DFI_Vel](#) (std::shared_ptr< psi::IntegralFactory > f_aabb, std::shared_ptr< psi::IntegralFactory > f_abab, std::shared_ptr< psi::Matrix > d.b)

Compute the Effective DFI Potential Matrix Due To Electrons.

- PSI_API std::shared_ptr< psi::Matrix > [oepdev::calculate_DFI_Vel_JK](#) (std::shared_ptr< psi::IntegralFactory > f_aabb, std::shared_ptr< psi::IntegralFactory > f_abab, std::shared_ptr< psi::Matrix > d.b)

Compute the Effective DFI Coulomb+Exchange Potential Matrix Due To Electrons.

- PSI_API std::shared_ptr< psi::Matrix > [oepdev::calculate_DFI_Vel_J](#) (std::shared_ptr< psi::IntegralFactory > f_aabb, std::shared_ptr< psi::Matrix > d.b)

Compute the Effective DFI Coulomb Potential Matrix Due To Electrons.

19.22 oepdev/libutil/wavefunction_union.h File Reference

```
#include <cstdio>
#include <string>
#include <map>
#include "psi4/psi4-dec.h"
#include "psi4/liboptions/liboptions.h"
#include "psi4/libpsio/psio.h"
#include "psi4/libciomr/libciomr.h"
#include "psi4/libpsio/psio.hpp"
#include "psi4/libqt/qt.h"
#include "psi4/libmints/molecule.h"
#include "psi4/libmints/writer.h"
#include "psi4/libmints/writer_file_prefix.h"
#include "psi4/libmints/wavefunction.h"
#include "psi4/libmints/basisset.h"
#include "psi4/libmints/vector.h"
#include "psi4/libmints/matrix.h"
#include "psi4/libmints/oeprop.h"
#include "psi4/libmints/local.h"
#include "psi4/libfunctional/superfunctional.h"
#include "psi4/libtrans/mospace.h"
#include "psi4/libtrans/integraltransform.h"
#include "psi4/libscf_solver/rhf.h"
```

```
#include "psi4/libdpd/dpd.h"
```

Classes

- class [oepdev::WavefunctionUnion](#)
Union of two Wavefunction objects.

Namespaces

- [oepdev](#)
OEPEv module namespace.

Chapter 20

Example Documentation

20.1 example_cphf.cc

Shows how to use the `oepdev::CPHF` solver to compute molecular and LMO-distributed polarizabilities at RHF level of theory.

```
void example_cphf(std::shared_ptr<psi::Wavefunction> wfn, psi::Options& opt){  
    // build the solver  
    std::shared_ptr<oepdev::CPHF> solver = std::make_shared<oepdev::CPHF>(wfn, opt);  
  
    // run the solver to converge CPHF equations  
    solver->compute();  
  
    // print the LMO-distributed polarizabilities  
    for (int i=0; i<solver->nocc(); i++) {  
        solver->polarizability(i)->print();  
    }  
  
    // print the molecular polarizability  
    solver->polarizability()->print();  
  
    // grab 4th LMO-distributed polarizability and its associated LMO centroid  
    psi::SharedMatrix pol_4 = solver->polarizability(3);  
    psi::SharedVector rmo_4 = solver->lmo.centroid(3);  
};
```

20.2 example_integrals_iter.cc

Iterations over electron repulsion integrals in AO basis. This is an example of how to use

- the `oepdev::ShellCombinationsIterator` class
- the `oepdev::AOIntegralsIterator` class.

```
void iterate(std::shared_ptr<oepdev::IntegralFactory> ints)  
{  
    // Prepare for direct calculation of ERI's (shell by shell)  
    std::shared_ptr<psi::TwoBodyAOInt> tei(ints->eri());  
  
    // Grab the buffer where the integrals for a current shell will be placed  
    const double* buffer = tei->buffer();
```

```

// Create iterator to go through all shell quartet combinations
oepdev::SharedShellsIterator shellIter =
    oepdev::ShellCombinationsIterator::build(ints, "ALL", 4);

// Iterate over shells, and then over all integrals in each shell quartet
for (shellIter->first(); shellIter->is_done() == false; shellIter->next())
{
    // Compute all integrals between shells in the current quartet
    shellIter->compute_shell(tei);

    // Create iterator to go through all integrals within a shell quartet
    oepdev::SharedAOIntsIterator intsIter = shellIter->ao_iterator("ALL");

    for (intsIter->first(); intsIter->is_done() == false; intsIter->next())
    {
        // Grab current (ij|kl) indices here
        int i = intsIter->i();
        int j = intsIter->j();
        int k = intsIter->k();
        int l = intsIter->l();

        // Grab the (ij|kl) integral
        double integral = buffer[intsIter->index()];
    }
}
}

```

20.3 example_scf_perturb.cc

Perturb HF Hamiltonian with external electrostatic potential. This is an example of how to use the `oepdev::RHFPerturbed` class.

```

void scf_perturb(std::shared_ptr<psi::Wavefunction> wfn, psi::Options& opt)
{
    // Set up HF superfunctional
    std::shared_ptr<psi::SuperFunctional> func = oepdev::create_superfunctional
        ("HF", opt);

    // Initialize the perturbed wavefunction
    std::shared_ptr<oepdev::RHFPerturbed> scf = std::make_shared<oepdev::RHFPerturbed>(wfn, func, opt, wfn->
        psio());

    /* Perturb the system with the uniform electric field [Fx, Fy, Fz].
       Then, add two point charges of charge qi placed at [Rxi, Ryi, Rzi].
       Provide all these values in atomic units! */
    const double Fx = 0.04, Fy = 0.05, Fz = -0.09;
    const double Rx1= 0.00, Rx2= 1.30, Rx3= -1.00;
    const double Rx1= 0.10, Rx2=-0.30, Rx3= 3.50;
    const double q1 = 0.30, q2 =-0.09;

    scf->set_perturbation(Fx, Fy, Fz);          /* set it only once, setting it again will overwrite the
        field, not add */
    scf->set_perturbation(Rx1, Ry1, Rz1, q1);
    scf->set_perturbation(Rx2, Ry2, Rz2, q2); /* more charges can be added */

    // Solve perturbed SCF equations
    scf->compute_energy();

    // Grab some data
    double energy = scf->reference_energy();    // Total energy of the system
    std::shared_ptr<psi::Matrix> Da = scf->Da(); // One-particle density matrix

    /* Note that the external field and charges perturb only one-electron Hamiltonian.*/
}

```

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