
GHF

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RESTRICTED HARTREE FOCK, BY MEANS OF SCF PROCEDURE

This class is used to calculate the RHF energy of a given molecule and the number of electrons. The molecule has to be created in pySCF: molecule = gto.M(atom = geometry, spin = diff. in alpha and beta electrons, basis = basis set)

class hf.RHF.**RHF** (molecule, number_of_electrons, int_method='pyscf')

Input is a molecule and the number of electrons.

Molecules are made in pySCF and calculations are performed as follows, eg.: The following snippet prints and returns RHF energy of h_2 and the number of iterations needed to get this value.

```
>>> h_2 = gto.M(atom = 'h 0 0 0; h 0 0 1', spin = 0, basis = 'cc-pvdz')
>>> x = RHF(h_2, 2)
>>> x.get_scf_solution()
Number of iterations: 6
Converged SCF energy in Hartree: -1.0661086493179357 (RHF)
```

diis (convergence=1e-12, complex_method=False)

When needed, DIIS can be used to speed up the RHF calculations by reducing the needed iterations.

Parameters

- **convergence** – Set the convergence criterion. If none is given, 1e-12 is used.
- **complex_method** – Specify whether or not you want to work in the complex space. Default is real.

Returns scf energy, number of iterations, mo coefficients, last density matrix, last fock matrix

get_last_dens ()

Returns the last density matrix of the converged solution.

Returns The last density matrix.

get_last_fock ()

Returns the last fock matrix of the converged solution.

Returns The last Fock matrix.

get_mo_coeff ()

Returns mo coefficients of the converged solution.

Returns The mo coefficients

get_one_e ()

Returns The one electron integral matrix: T + V

get_ovlp ()

Returns The overlap matrix

get_scf_solution (*convergence=1e-12, complex_method=False*)

Prints the number of iterations and the converged scf energy.

Parameters

- **convergence** – Set the convergence criterion. If none is given, 1e-12 is used.
- **complex_method** – Specify whether or not you want to work in the complex space. Default is real.

Returns the converged energy

get_scf_solution_diis (*convergence=1e-12, complex_method=False*)

Prints the number of iterations and the converged DIIS energy. The number of iterations will be lower than with a normal scf, but the energy value will be the same. Example:

```
>>> h2 = gto.M(atom = 'h 0 0 0; h 1 0 0', basis = 'cc-pvdz')
>>> x = RHF(h2, 2)
>>> x.get_scf_solution_diis()
Number of iterations: 5
Converged SCF energy in Hartree: -1.100153764878446 (RHF)
```

Parameters

- **convergence** – Set the convergence criterion. If none is given, 1e-12 is used.
- **complex_method** – Specify whether or not you want to work in the complex space. Default is real.

Returns The converged scf energy, using DIIS.

get_two_e ()

Returns The electron repulsion interaction tensor

nuc_rep ()

Returns The nuclear repulsion value

scf (*convergence=1e-12, complex_method=False*)

Performs a self consistent field calculation to find the lowest RHF energy.

Parameters

- **convergence** – Convergence criterion. If none is specified, 1e-12 is used.
- **complex_method** – Specify whether or not you want to work in the complex space. Default is real.

Returns number of iterations, scf energy, mo coefficients, last density matrix, last fock matrix

UNRESTRICTED HARTREE FOCK, BY MEANS OF SCF PROCEDURE

This class is used to calculate the UHF energy for a given molecule and the number of electrons of that molecule. Several options are available to make sure you get the lowest energy from your calculation, as well as some useful functions to get intermediate values such as MO coefficients, density and fock matrices.

class hf.UHF.UHF (molecule, number_of_electrons, int_method='pyscf')

Input is a molecule and the number of electrons.

Molecules are made in pySCF and calculations are performed as follows, eg.: The following snippet prints and returns UHF energy of h₃ and the number of iterations needed to get this value.

For a normal scf calculation your input looks like the following example:

```
>>> h3 = gto.M(atom = 'h 0 0 0; h 0 0.86602540378 0.5; h 0 0 1', spin = 1, basis_
↳= 'cc-pvdz')
>>> x = UHF(h3, 3)
>>> x.get_scf_solution()
Number of iterations: 47
Converged SCF energy in Hartree: -1.506274320261134 (UHF)
<S^2> = 0.7735672504295973, <S_z> = 0.5, Multiplicity = 2.023430009098014
```

diis (initial_guess=None, convergence=1e-12, complex_method=False)

When needed, DIIS can be used to speed up the UHF calculations by reducing the needed iterations.

Parameters

- **initial_guess** – Initial guess for the scf procedure. None specified: core Hamiltonian.
- **convergence** – Set the convergence criterion. If none is given, 1e-12 is used.
- **complex_method** – Specify whether or not you want to work in the complex space. Default is real.

Returns scf energy, number of iterations, mo coefficients, last density matrix, last fock matrix

extra_electron_guess ()

This method adds two electrons to the system in order to get coefficients that can be used as a better guess for the scf procedure. This essentially forces the system into its $\langle S_z \rangle = 0$ state.

!!!IMPORTANT!!! Only supported with pyscf.

To perform a calculation with this method, you will have to work as follows:

```
>>> h4 = gto.M(atom = 'h 0 0 0; h 1 0 0; h 0 1 0; h 1 1 0' , spin = 2, basis_
↳= 'cc-pvdz')
>>> x = UHF(h4, 4)
>>> guess = x.extra_electron_guess()
```

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```
>>> x.get_scf_solution(guess)
Number of iterations: 60
Converged SCF energy in Hartree: -2.0210882477030547 (UHF)
<S^2> = 1.0565277001056579, <S_z> = 0.0, Multiplicity = 2.2860688529487976
```

Returns A new guess matrix to use for the scf procedure.

get_hessian()

Get the Hessian matrix after performing a stability analysis. :return: The hessian matrix

get_last_dens()

Gets the last density matrix of the converged solution. Alpha density in the first matrix, beta density in the second.

Returns The last density matrix.

get_last_fock()

Gets the last fock matrix of the converged solution. Alpha Fock matrix first, beta Fock matrix second.

Returns The last Fock matrix.

get_mo_coeff()

Gets the mo coefficients of the converged solution. Alpha coefficients in the first matrix, beta coefficients in the second.

Returns The mo coefficients

get_one_e()

Returns The one electron integral matrix: T + V

get_ovlp()

Returns The overlap matrix

get_scf_solution (*guess=None, convergence=1e-12, complex_method=False*)

Prints the number of iterations and the converged scf energy. Also prints the expectation value of S_z , S^2 and the multiplicity.

Parameters

- **guess** – The initial guess for the scf procedure. If none is given: core Hamiltonian.
- **convergence** – Set the convergence criterion. If none is given, 1e-12 is used.
- **complex_method** – Specify whether or not you want to work in the complex space. Default is real.

Returns The converged scf energy.

get_scf_solution_diis (*guess=None, convergence=1e-12, complex_method=False*)

Prints the number of iterations and the converged diis energy. Also prints the expectation value of S_z , S^2 and the multiplicity.

Parameters

- **guess** – The initial guess. If none is specified, core Hamiltonian.
- **convergence** – Set the convergence criterion. If none is given, 1e-12 is used.
- **complex_method** – Specify whether or not you want to work in the complex space. Default is real.

Returns The converged diis energy.

get_two_e()

Returns The electron repulsion interaction tensor

nuc_rep()

Returns The nuclear repulsion value

scf (*initial_guess=None, convergence=1e-12, complex_method=False*)

Performs a self consistent field calculation to find the lowest UHF energy.

Parameters

- **initial_guess** – A tuple of an alpha and beta guess matrix. If none, the core hamiltonian will be used.
- **convergence** – Set the convergence criterion. If none is given, 1e-12 is used.
- **complex_method** – Specify whether or not you want to work in the complex space. Default is real.

Returns The scf energy, number of iterations, the mo coefficients, the last density and the last fock matrices

stability()

Performing a stability analysis checks whether or not the wave function is stable, by checking the lowest eigen- value of the Hessian matrix. If there's an instability, the MO's will be rotated in the direction of the lowest eigenvalue. These new MO's can then be used to start a new scf procedure.

To perform a stability analysis, use the following syntax:

```
>>> h4 = gto.M(atom = 'h 0 0 0; h 1 0 0; h 0 1 0; h 1 1 0' , spin = 2, basis_
↳= 'cc-pvdz')
>>> x = UHF(h4, 4)
>>> guess = x.stability()
>>> x.get_scf_solution(guess)
There is an internal instability in the UHF wave function.
Number of iterations: 66
Converged SCF energy in Hartree: -2.0210882477030716 (UHF)
<S^2> = 1.056527700105677, <S_z> = 0.0, Multiplicity = 2.2860688529488145
```

Returns New and improved MO's.

stability_analysis (*method, step_size=1e-05*)

Internal stability analysis to verify whether the wave function is stable within the space of the used method.
: param method: Indicate whether you want to check the internal or external stability of the wave function.
Can be internal or external. : param step_size: Step size for orbital rotation. standard is 1e-5. : return: In case of internal stability analysis, it returns a new set of coefficients.

CONSTRAINED UNRESTRICTED HARTREE FOCK BY SCUSERIA

This class is used to calculate the ROHF energy for a given molecule and the number of electrons of that molecule, using a constrained version of unrestricted Hartree Fock, according to Scuseria.. Several options are available to make sure you get the lowest energy from your calculation, as well as some useful functions to get intermediate values such as MO coefficients, density and fock matrices.

class hf.cUHF_s.CUHF (*molecule, number_of_electrons, int_method='pyscf'*)

Input is a molecule and the number of electrons.

Molecules are made in pySCF/psi4 and calculations are performed as follows, eg.: The following snippet prints and returns UHF energy of h₃ and the number of iterations needed to get this value.

For a normal scf calculation your input looks like the following example:

```
>>> h3 = gto.M(atom = 'h 0 0 0; h 0 0.86602540378 0.5; h 0 0 1', spin = 1, basis_
↳= 'cc-pvdz')
>>> x = CUHF(h3, 3)
>>> x.get_scf_solution()
```

get_dens (*i=-1*)

Gets the last density matrix of the converged solution. Alpha density in the first matrix, beta density in the second.

Returns The last density matrix.

get_fock (*i=-1*)

Gets the last fock matrix of the converged solution. Alpha Fock matrix first, beta Fock matrix second.

Returns The last Fock matrix.

get_mo_coeff (*i=-1*)

Gets the mo coefficients of the converged solution. Alpha coefficients in the first matrix, beta coefficients in the second.

Returns The mo coefficients

get_one_e ()

Returns The one electron integral matrix: T + V

get_ovlp ()

Returns The overlap matrix

get_scf_solution (*convergence=1e-12, diis=True*)

Prints the number of iterations and the converged scf energy. Also prints the expectation value of S_z, S² and the multiplicity.

Parameters

- **convergence** – Set the convergence criterion. If none is given, 1e-12 is used.
- **diis** – Accelerates the convergence, default is true.

Returns The converged scf energy.

get_two_e()

Returns The electron repulsion interaction tensor

nuc_rep()

Returns The nuclear repulsion value

scf (*convergence=1e-12, diis=True*)

Performs a self consistent field calculation to find the lowest UHF energy.

Parameters

- **convergence** – Set the convergence criterion. If none is given, 1e-12 is used.
- **diis** – Accelerates the convergence, default is true.

Returns The scf energy, number of iterations, the mo coefficients, the last density and the last fock matrices

CONSTRAINED UNRESTRICTED HARTREE FOCK BY P. BULTINCK

This class is used to calculate the ROHF energy for a given molecule and the number of electrons of that molecule, using a constrained version of unrestricted Hartree Fock. This constraint is an idea from professor P. Bultinck, where the alpha and beta MO's are made equal for the closed shell part of the system. Several options are available to make sure you get the lowest energy from your calculation, as well as some useful functions to get intermediate values such as MO coefficients, density and fock matrices.

class hf.cUHF_b.CUHF (molecule, number_of_electrons, int_method='pyscf')

Input is a molecule and the number of electrons.

Molecules are made in pySCF/psi4 and calculations are performed as follows, eg.: The following snippet prints and returns UHF energy of h₃ and the number of iterations needed to get this value.

For a normal scf calculation your input looks like the following example:

```
>>> h3 = gto.M(atom = 'h 0 0 0; h 0 0.86602540378 0.5; h 0 0 1', spin = 1, basis_
↳= 'cc-pvdz')
>>> x = CUHF(h3, 3)
>>> x.get_scf_solution()
```

diis (initial_guess=None, convergence=1e-12)

When needed, DIIS can be used to speed up the UHF calculations by reducing the needed iterations.

Parameters

- **initial_guess** – Initial guess for the scf procedure. None specified: core Hamiltonian.
- **convergence** – Set the convergence criterion. If none is given, 1e-12 is used.

Returns scf energy, number of iterations, mo coefficients, last density matrix, last fock matrix

get_last_dens ()

Gets the last density matrix of the converged solution. Alpha density in the first matrix, beta density in the second.

Returns The last density matrix.

get_last_fock ()

Gets the last fock matrix of the converged solution. Alpha Fock matrix first, beta Fock matrix second.

Returns The last Fock matrix.

get_mo_coeff ()

Gets the mo coefficients of the converged solution. Alpha coefficients in the first matrix, beta coefficients in the second.

Returns The mo coefficients

get_one_e ()

Returns The one electron integral matrix: $T + V$

get_ovlp()

Returns The overlap matrix

get_scf_solution (*guess=None, convergence=1e-12*)

Prints the number of iterations and the converged scf energy. Also prints the expectation value of S_z , S^2 and the multiplicity.

Parameters

- **guess** – Initial scf guess
- **convergence** – Set the convergence criterion. If none is given, 1e-12 is used.

Returns The converged scf energy.

get_scf_solution_diis (*guess=None, convergence=1e-12*)

Prints the number of iterations and the converged diis energy. Also prints the expectation value of S_z , S^2 and the multiplicity.

Parameters

- **guess** – The initial guess. If none is specified, core Hamiltonian.
- **convergence** – Set the convergence criterion. If none is given, 1e-12 is used.

Returns The converged diis energy.

get_two_e()

Returns The electron repulsion interaction tensor

nuc_rep()

Returns The nuclear repulsion value

random_guess()

A function that creates a matrix with random values that can be used as an initial guess for the SCF calculations.

To use this guess:

```
>>> h3 = gto.M(atom = 'h 0 0 0; h 0 0.86602540378 0.5; h 0 0 1', spin = 1,   
↳basis = 'cc-pvdz')  
>>> x = CUHF(h3, 3)  
>>> guess = x.random_guess()  
>>> x.get_scf_solution(guess)
```

Returns A random hermitian matrix.

scf (*initial_guess=None, convergence=1e-12*)

Performs a self consistent field calculation to find the lowest UHF energy.

Parameters

- **initial_guess** – Set the convergence criterion. If none is given, 1e-12 is used.
- **convergence** – Set the convergence criterion. If none is given, 1e-12 is used.

Returns The scf energy, number of iterations, the mo coefficients, the last density and the last fock matrices

GENERALISED HARTREE FOCK, BY MEANS OF SCF PROCEDURE

This class creates a generalised Hartree-Fock object which can be used for scf calculations. Different initial guesses are provided as well as the option to perform a stability analysis. The molecule has to be created in pySCF: molecule = gto.M(atom = geometry, spin = diff. in alpha and beta electrons, basis = basis set)

class hf.GHF.GHF (molecule, number_of_electrons, int_method='pyscf')

Input is a molecule and the number of electrons.

Molecules are made in pySCF and calculations are performed as follows, eg.: The following snippet prints and returns UHF energy of h3 and the number of iterations needed to get this value.

For a normal scf calculation your input looks like the following example:

```
>>> h3 = gto.M(atom = 'h 0 0 0; h 0 0.86602540378 0.5; h 0 0 1', spin = 1, basis_
↳= 'cc-pvdz')
>>> x = GHF(h3, 3)
>>> x.get_scf_solution()
Number of iterations: 81
Converged SCF energy in Hartree: -1.5062743202607725 (Real GHF)
```

diis (guess=None, convergence=1e-12, complex_method=False)

The DIIS method is an alternative to the standard scf procedure. It reduces the number of iterations needed to find a solution. The same guesses can be used as for a standard scf calculation. Stability analysis can be done as well.

Parameters

- **convergence** – Set the convergence criterion. If none is given, 1e-12 is used.
- **guess** – The initial guess matrix, if none is specified, the spin blocked core Hamiltonian is used.
- **complex_method** – Specify whether or not you want to work in the complex space. Default is real.

Returns scf_energy, iterations, mo coefficients, last density matrix & last Fock matrix

get_hessian ()

After stability analysis is performed, the hessian is stored and can be used for further studying. :return: The Hessian matrix

get_last_dens ()

Gets the last density matrix of the converged solution.

Returns The last density matrix.

get_last_fock ()

Gets the last fock matrix of the converged solution.

Returns The last Fock matrix.

get_mo_coeff()

Gets the mo coefficients of the converged solution.

Returns The mo coefficients

get_one_e()

Returns The one electron integral matrix: $T + V$

get_ovlp()

Returns The overlap matrix

get_scf_solution (*guess=None, convergence=1e-12, complex_method=False*)

Prints the number of iterations and the converged scf energy.

Parameters

- **guess** – Initial guess for scf. If none is specified: expanded core Hamiltonian.
- **convergence** – Set the convergence criterion. If none is given, 1e-12 is used.
- **complex_method** – Specify whether or not you want to work in the complex space. Default is real.

Returns The converged scf energy.

get_scf_solution_diis (*guess=None, convergence=1e-12, complex_method=False*)

Prints the number of iterations and the converged energy after a diis calculation. Guesses can also be specified just like with a normal scf calculation.

Example:

```
>>> h3 = gto.M(atom = 'h 0 0 0; h 0 0.86602540378 0.5; h 0 0 1', spin = 1,
↳basis = 'cc-pvdz')
>>> x = GHF(h3, 3)
>>> guess = x.random_guess()
>>> x.get_scf_solution_diis(guess)
Number of iterations: 23
Converged SCF energy in Hartree: -1.5062743202915496 (Real GHF)
```

Without DIIS, 81 iterations are needed to find this solution.

Parameters

- **guess** – Initial guess for scf. None specified: expanded core Hamiltonian
- **convergence** – Set the convergence criterion. If none is given, 1e-12 is used.
- **complex_method** – Specify whether or not you want to work in the complex space. Default is real.cd

Returns The converged scf energy.

get_two_e()

Returns The electron repulsion interaction tensor

nuc_rep()

Returns The nuclear repulsion value

random_guess()

A function that creates a matrix with random values that can be used as an initial guess for the SCF calculations.

To use this guess:

```
>>> h3 = gto.M(atom = 'h 0 0 0; h 0 0.86602540378 0.5; h 0 0 1', spin = 1,
↳basis = 'cc-pvdz')
>>> x = GHF(h3, 3)
>>> guess = x.random_guess()
>>> x.get_scf_solution(guess)
```

Returns A random hermitian matrix.

scf (*guess=None, convergence=1e-12, complex_method=False*)

This function performs the SCF calculation by using the generalised Hartree-Fock formulas. Since we're working in the real class, all values throughout are real. For complex, see the "complex_GHF" class.

Parameters

- **guess** – Initial guess to start SCF. If none is given, core hamiltonian will be used.
- **convergence** – Set the convergence criterion. If none is given, 1e-12 is used.
- **complex_method** – Specify whether or not you want to work in the complex space. Default is real.

Returns scf_energy, iterations, mo coefficients, last density matrix & last Fock matrix

stability_analysis (*method, step_size=0.0001*)

Internal stability analysis to verify whether the wave function is stable within the space of the used method.
:param method: Indicate whether you want to check the internal or external stability of the wave function. Can be internal or external. :param step_size: Step size for orbital rotation. standard is 1e-4. :return: In case of internal stability analysis, it returns a new set of coefficients.

unitary_rotation_guess (*init=None*)

A function that creates an initial guess matrix by performing a unitary transformation on the core Hamiltonian matrix.

To use this guess:

```
>>> h3 = gto.M(atom = 'h 0 0 0; h 0 0.86602540378 0.5; h 0 0 1', spin = 1,
↳basis = 'cc-pvdz')
>>> x = GHF(h3, 3)
>>> guess = x.unitary_rotation_guess()
>>> x.get_scf_solution(guess)
```

Returns A rotated guess matrix.

USEFUL FUNCTIONS FOR SCF PROCEDURE

A number of functions used throughout the UHF and RHF calculations are summarised here.

`hf.SCF_functions.density_matrix(f_matrix, occ, trans)`

- `density()` creates a density matrix from a fock matrix and the number of occupied orbitals.
- Input is a fock matrix, the number of occupied orbitals, which can be separate for alpha and beta in case of UHF. And a transformation matrix X.

`hf.SCF_functions.eri_ao_to_mo(eri, coeff, complexity=False)`

Transform the two electron tensor to MO basis. Scales as N^5 . :param eri: Electron repulsion interaction, in ao notation, tensor in spinor basis :param coeff: coefficient matrix in spinor basis :param complexity: specify whether you are working with real or complex tensors. Default is real. :return: Electron repulsion interaction, in mo notation, tensor in spinor basis

`hf.SCF_functions.expand_matrix(matrix)`

Parameters matrix –

Returns a matrix double the size, where blocks of zero's are added top right and bottom left.

`hf.SCF_functions.expand_tensor(tensor, complexity=False)`

Expand every matrix within the tensor, in the same way the expand matrix function works. :param tensor: The tensor, usually eri, that you wish to expand. :param complexity: Is your tensor complex or not? Default is false. :return: a tensor where each dimension is doubled.

`hf.SCF_functions.get_integrals_psi4(mol)`

A function to calculate your integrals & nuclear repulsion with psi4. :param mol: Psi4 instance :return: overlap, core hamiltonian, eri tensor and nuclear repulsion

`hf.SCF_functions.get_integrals_pyscf(molecule)`

A function to calculate your integrals & nuclear repulsion with pyscf.

`hf.SCF_functions.ghf_spin(coeff, n_e, trans)`

A function used to calculate the spin expectation values in the generalised hartree fock formalism. :param coeff: The generalised MO coefficients :param n_e: number of electrons :param trans: transformation matrix, eg.: $S^{(-1/2)}$:return: The expectation values of S_z , S^2 and the multiplicity ($2S+1$)

`hf.SCF_functions.spin(occ_a, occ_b, coeff_a, coeff_b, overlap)`

Parameters

- **occ_a** – number of occupied alpha orbitals
- **occ_b** – number of occupied beta orbitals
- **coeff_a** – MO coefficients of alpha orbitals
- **coeff_b** – MO coefficients of beta orbitals

- **overlap** – overlap matrix of the molecule

Returns S^2 , S_z and spin multiplicity

`hf.SCF_functions.spin_blocked(block_1, block_2, block_3, block_4)`

When creating the blocks of the density or fock matrix separately, this function is used to add them together, and create the total density or Fock matrix in spin Blocked notation. :return: a density matrix in the spin-blocked notation

`hf.SCF_functions.trans_matrix(overlap)`

- Define a transformation matrix X, used to orthogonalize different matrices throughout the calculation.
- Input should be an overlap matrix.

`hf.SCF_functions.uhf_fock_matrix(density_matrix_1, density_matrix_2, one_electron, two_electron)`

- calculate a fock matrix from a given alpha and beta density matrix
- fock alpha if 1 = alpha and 2 = beta and vice versa
- input is the density matrix for alpha and beta, a one electron matrix and a two electron tensor.

`hf.SCF_functions.uhf_scf_energy(density_matrix_a, density_matrix_b, fock_a, fock_b, one_electron)`

- calculate the scf energy value from a given density matrix and a given fock matrix for both alpha and beta, so 4 matrices in total.
- then calculate the initial electronic energy and put it into an array
- input is the density matrices for alpha and beta, the fock matrices for alpha and beta and lastly a one electron matrix.

TESTING THE RHF AND UHF METHODS

Simple tests to check whether or not the functions return the correct value.

`hf.tests.test_auth.test_diis_real_ghf()`

Test whether diis gives the same energy in fewer iterations.

`hf.tests.test_auth.test_diis_rhf()`

Test whether diis gives the same energy in fewer iterations.

`hf.tests.test_auth.test_diis_uhf()`

Test whether diis gives the same energy in fewer iterations.

`hf.tests.test_auth.test_extra_e()`

`test_extra_e` will test the UHF method, with the added option of first adding 2 electrons to the system and using those coefficients for the actual system, by checking whether or not it returns the expected result. The accuracy is 10^{-6} .

`hf.tests.test_auth.test_one_e()`

Test whether or not psi4 and pyscf give the same core Hamiltonian integrals. :return:

`hf.tests.test_auth.test_overlap()`

Test whether or not psi4 and pyscf give the same overlap integrals.

`hf.tests.test_auth.test_pyscf_vs_psi4()`

Test a pyscf energy calculation vs a psi4 energy calculation.

`hf.tests.test_auth.test_rhf()`

`test_RHF` will test whether or not the RHF method returns the wanted result. The accuracy is 10^{-11} .

`hf.tests.test_auth.test_stability()`

`test_stability` will test the UHF method, with stability analysis, by checking whether or not it returns the expected result. The accuracy is 10^{-6} .

`hf.tests.test_auth.test_two_e()`

Test whether or not psi4 and pyscf give the same two electron integrals.

`hf.tests.test_auth.test_uhf()`

`test_UHF` will test the regular UHF method, by checking whether or not it returns the expected result. The accuracy is 10^{-6} .

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