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CHAPTER 2

Many Electron Wave Functions and Operators

2.1 The Electron Problem

2.1.1 Atomic Units

2.1.2 The Born-Oppenheimer Approximation

2.1.3 The Antisymmetry or Pauli Exclusion Principle

2.2 Orbitals, Slater Determinants, and Basis Functions

2.2.1 Spin Orbitals and Spatial Orbitals

Exercise 2.1

Given a set of K orthonormal spatial functions, $\{\psi_i^\alpha(\mathbf{r})\}$, and another set of K orthonormal functions, $\{\psi_i^\beta(\mathbf{r})\}$, such that the first set is not orthogonal to the second set, i.e.,

$$\int d\mathbf{r} \psi_i^{\alpha*}(\mathbf{r}) \psi_j^\beta(\mathbf{r}) = S_{ij}$$

where \mathbf{S} is an overlap matrix, show that the set $\{\chi_i\}$ of $2K$ spin orbitals, formed by multiplying $\psi_i^\alpha(\mathbf{r})$ by the α spin function and $\psi_i^\beta(\mathbf{r})$ by the β spin function, i.e.,

$$\left. \begin{aligned} \chi_{2i-1}(\mathbf{x}) &= \psi_i^\alpha(\mathbf{r})\alpha(\omega) \\ \chi_{2i}(\mathbf{x}) &= \psi_i^\beta(\mathbf{r})\beta(\omega) \end{aligned} \right\} i = 1, 2, \dots, K$$

is an orthonormal set.

Solution 2.1

It is easy to verify the normalization of any χ_{2i-1} or χ_{2j} , where $i = 1, 2, \dots, K$ and $j = 1, 2, \dots, K$,

$$\langle \chi_{2i-1} | \chi_{2j-1} \rangle = \int d\mathbf{x} \chi_{2i-1}^*(\mathbf{x}) \chi_{2j-1}(\mathbf{x}) = \int d\mathbf{r} \psi_i^{\alpha*}(\mathbf{r}) \psi_j^\alpha(\mathbf{r}) \int d\omega \alpha^*(\omega) \alpha(\omega) = \delta_{ij} \times 1 = \delta_{ij},$$

$$\langle \chi_{2i} | \chi_{2j} \rangle = \int d\mathbf{x} \chi_{2i}^*(\mathbf{x}) \chi_{2j}(\mathbf{x}) = \int d\mathbf{r} \psi_i^{\beta*}(\mathbf{r}) \psi_j^\beta(\mathbf{r}) \int d\omega \beta^*(\omega) \beta(\omega) = \delta_{ij} \times 1 = \delta_{ij}.$$

and the orthogonality between χ_{2i-1} and χ_{2j} , where $i = 1, 2, \dots, K$ and $j = 1, 2, \dots, K$,

$$\langle \chi_{2i-1} | \chi_{2j} \rangle = \int d\mathbf{x} \chi_{2i-1}^*(\mathbf{x}) \chi_{2j}(\mathbf{x}) = \int d\mathbf{r} \psi_i^{\alpha*}(\mathbf{r}) \psi_j^\beta(\mathbf{r}) \int d\omega \alpha^*(\omega) \beta(\omega) = S_{ij} \times 0 = 0,$$

$$\langle \chi_{2i} | \chi_{2j-1} \rangle = \int d\mathbf{x} \chi_{2i}^*(\mathbf{x}) \chi_{2j-1}(\mathbf{x}) = \int d\mathbf{r} \psi_i^{\beta*}(\mathbf{r}) \psi_j^\alpha(\mathbf{r}) \int d\omega \beta^*(\omega) \alpha(\omega) = S_{ji}^* \times 0 = 0.$$

Thus, we can that the set $\{\chi_i\}$ of $2K$ spin orbitals is an orthonormal set.

2.2.2 Hartree Products

Exercise 2.2

Show that the Hartree product of (2.30) is an eigenfunction of $\mathcal{H} = \sum_{i=1}^N h(i)$ with an eigenvalue given by (2.32).

Solution 2.2

The verification is easy. With (2.29), we find that

$$\mathcal{H}\Psi^{\text{HP}} = \left(\sum_{i=1}^N h(i) \right) \left[\prod_{j=1}^N \chi_{j'}(\mathbf{x}_j) \right] = \sum_{i=1}^N \prod_{j=1}^N h(i) \chi_{j'}(\mathbf{x}_j) = \sum_{i=1}^N \prod_{j=1}^N \varepsilon_i \chi_{j'}(\mathbf{x}_j) = \sum_{i=1}^N \varepsilon_i \prod_{j=1}^N \chi_{j'}(\mathbf{x}_j). \quad (2.2-1)$$

2.2.3 Slater Determinants

Exercise 2.3

Show that $\Psi(\mathbf{x}_1, \mathbf{x}_2)$ of Eq.(2.34) is normalized.

Solution 2.3

The verification is direct, viz.,

$$\begin{aligned} \langle \Psi | \Psi \rangle &= \int d\vec{x} \langle \Psi | \vec{x} \rangle \langle \vec{x} | \Psi \rangle \\ &= \int d\mathbf{x}_1 \int d\mathbf{x}_2 \frac{1}{\sqrt{2}} [\chi_i(\mathbf{x}_1) \chi_j(\mathbf{x}_2) - \chi_j(\mathbf{x}_1) \chi_i(\mathbf{x}_2)]^* \frac{1}{\sqrt{2}} [\chi_i(\mathbf{x}_1) \chi_j(\mathbf{x}_2) - \chi_j(\mathbf{x}_1) \chi_i(\mathbf{x}_2)] \\ &= \frac{1}{2} \left(\int d\mathbf{x}_1 \int d\mathbf{x}_2 \chi_i^*(\mathbf{x}_1) \chi_j^*(\mathbf{x}_2) \chi_i(\mathbf{x}_1) \chi_j(\mathbf{x}_2) - \int d\mathbf{x}_1 \int d\mathbf{x}_2 \chi_i^*(\mathbf{x}_1) \chi_j^*(\mathbf{x}_2) \chi_j(\mathbf{x}_1) \chi_i(\mathbf{x}_2) \right. \\ &\quad \left. - \int d\mathbf{x}_1 \int d\mathbf{x}_2 \chi_j^*(\mathbf{x}_1) \chi_i^*(\mathbf{x}_2) \chi_i(\mathbf{x}_1) \chi_j(\mathbf{x}_2) + \int d\mathbf{x}_1 \int d\mathbf{x}_2 \chi_j^*(\mathbf{x}_1) \chi_i^*(\mathbf{x}_2) \chi_j(\mathbf{x}_1) \chi_i(\mathbf{x}_2) \right) \\ &= \frac{1}{2} (1 - 0 - 0 + 1) = 1. \end{aligned}$$

Exercise 2.4

Suppose the spin orbitals χ_i and χ_j are eigenfunctions of a one-electron operator h with eigenvalues ε_i and ε_j as in Eq.(2.29). Show that the Hartree products in Eqs.(2.33a, b) and the antisymmetrized wave function in Eq.(2.34) are eigenfunctions of the independent-particle Hamiltonian $\mathcal{H} = h(1) + h(2)$ (c.f. Eq.(2.28)) and have the same eigenvalue namely, $\varepsilon_i + \varepsilon_j$.

Solution 2.4

Firstly, we check the Hartree products of χ_i and χ_j . With the conclusion of Exercise 2.2, we get that

$$\begin{aligned} \mathcal{H}|\Psi_{12}^{\text{HP}}\rangle &= (\varepsilon_i + \varepsilon_j)|\Psi_{12}^{\text{HP}}\rangle, \\ \mathcal{H}|\Psi_{21}^{\text{HP}}\rangle &= (\varepsilon_j + \varepsilon_i)|\Psi_{21}^{\text{HP}}\rangle = (\varepsilon_i + \varepsilon_j)|\Psi_{21}^{\text{HP}}\rangle. \end{aligned}$$

Thus, the eigenvalue of the Hartree product of χ_i and χ_j is irrelevant to their order. Note that

$$\Psi = \frac{1}{\sqrt{2}} [\chi_i(\mathbf{x}_1) \chi_j(\mathbf{x}_2) - \chi_j(\mathbf{x}_1) \chi_i(\mathbf{x}_2)] = \frac{1}{\sqrt{2}} (\Psi_{12}^{\text{HP}} - \Psi_{21}^{\text{HP}}),$$

we find that

$$\mathcal{H}|\Psi\rangle = \mathcal{H} \frac{1}{\sqrt{2}} (|\Psi_{12}^{\text{HP}}\rangle - |\Psi_{21}^{\text{HP}}\rangle) = \frac{1}{\sqrt{2}} (\mathcal{H}|\Psi_{12}^{\text{HP}}\rangle - \mathcal{H}|\Psi_{21}^{\text{HP}}\rangle)$$

$$\begin{aligned}
&= \frac{1}{\sqrt{2}} [(\varepsilon_i + \varepsilon_j) |\Psi_{12}^{\text{HP}}\rangle - (\varepsilon_i + \varepsilon_j) |\Psi_{21}^{\text{HP}}\rangle] \\
&= (\varepsilon_i + \varepsilon_j) \frac{1}{\sqrt{2}} (|\Psi_{12}^{\text{HP}}\rangle - |\Psi_{21}^{\text{HP}}\rangle) = (\varepsilon_i + \varepsilon_j) |\Psi\rangle.
\end{aligned}$$

Thus, we have proved that the Hartree products in Eqs.(2.33a, b) and the antisymmetrized wave function in Eq.(2.34) are eigenfunctions of the independent-particle Hamiltonian $\mathcal{H} = h(1) + h(2)$ and have the same eigenvalue $\varepsilon_i + \varepsilon_j$.

Exercise 2.5

Consider the Slater determinants

$$|K\rangle = |\chi_i \chi_j\rangle, \quad |L\rangle = |\chi_k \chi_l\rangle.$$

Show that

$$\langle K|L\rangle = \delta_{ik}\delta_{jl} - \delta_{il}\delta_{jk}.$$

Note that the overlap is zero unless: 1) $k = i$ and $l = j$, in which case $|L\rangle = |K\rangle$ and the overlap is unity and 2) $k = j$ and $l = i$ in which case $|L\rangle = |\chi_j \chi_i\rangle = -|K\rangle$ and the overlap is minus one.

Solution 2.5

We calculate the inner product firstly,

$$\begin{aligned}
\langle K|L\rangle &= \int d\vec{x} \langle K|\vec{x}\rangle \langle \vec{x}|L\rangle \\
&= \int d\mathbf{x}_1 \int d\mathbf{x}_2 \frac{1}{\sqrt{2}} [\chi_i(\mathbf{x}_1)\chi_j(\mathbf{x}_2) - \chi_j(\mathbf{x}_1)\chi_i(\mathbf{x}_2)]^* \frac{1}{\sqrt{2}} [\chi_k(\mathbf{x}_1)\chi_l(\mathbf{x}_2) - \chi_l(\mathbf{x}_1)\chi_k(\mathbf{x}_2)] \\
&= \frac{1}{2} \left[\int d\mathbf{x}_1 \int d\mathbf{x}_2 \chi_i^*(\mathbf{x}_1)\chi_j^*(\mathbf{x}_2)\chi_k(\mathbf{x}_1)\chi_l(\mathbf{x}_2) - \int d\mathbf{x}_1 \int d\mathbf{x}_2 \chi_i^*(\mathbf{x}_1)\chi_j^*(\mathbf{x}_2)\chi_l(\mathbf{x}_1)\chi_k(\mathbf{x}_2) \right. \\
&\quad \left. - \int d\mathbf{x}_1 \int d\mathbf{x}_2 \chi_j^*(\mathbf{x}_1)\chi_i^*(\mathbf{x}_2)\chi_k(\mathbf{x}_1)\chi_l(\mathbf{x}_2) + \int d\mathbf{x}_1 \int d\mathbf{x}_2 \chi_j^*(\mathbf{x}_1)\chi_i^*(\mathbf{x}_2)\chi_l(\mathbf{x}_1)\chi_k(\mathbf{x}_2) \right] \\
&= \frac{1}{2} (\delta_{ik}\delta_{jl} - \delta_{il}\delta_{jk} - \delta_{jk}\delta_{il} + \delta_{jl}\delta_{ik}) = \delta_{ik}\delta_{jl} - \delta_{jk}\delta_{il}.
\end{aligned}$$

The conclusion is obvious.

- When $k = i$ and $l = j$, in which case $|L\rangle = |K\rangle$ and the overlap is 1.
- When $k = j$ and $l = i$ in which case $|L\rangle = |\chi_k \chi_l\rangle = |\chi_j \chi_i\rangle = -|K\rangle$ and the overlap is -1 .
- Otherwise, the overlap is 0.

2.2.4 The Hartree-Fock Approximation

2.2.5 The Minimal Basis H_2 Model

Exercise 2.6

Show that ψ_1 and ψ_2 form an orthonormal set.

Solution 2.6

Similar to Exercise 2.1, verify the normalization of ψ_1 or ψ_2 , with $S_{12} = S_{21}$,

$$\begin{aligned}
\langle \psi_1|\psi_1\rangle &= \int d\mathbf{r} \psi_1^*(\mathbf{r})\psi_1(\mathbf{r}) = \int d\mathbf{r} \frac{1}{\sqrt{2(1+S_{12})}} (\phi_1(\mathbf{r}) + \phi_2(\mathbf{r}))^* \frac{1}{\sqrt{2(1+S_{12})}} (\phi_1(\mathbf{r}) + \phi_2(\mathbf{r})) \\
&= \frac{1}{2(1+S_{12})} (1 + S_{12} + S_{21} + 1) = 1,
\end{aligned}$$

$$\begin{aligned}\langle\psi_2|\psi_2\rangle &= \int d\mathbf{r} \psi_2^*(\mathbf{r})\psi_2(\mathbf{r}) = \int d\mathbf{r} \frac{1}{\sqrt{2(1-S_{12})}} (\phi_1(\mathbf{r}) - \phi_2(\mathbf{r}))^* \frac{1}{\sqrt{2(1-S_{12})}} (\phi_1(\mathbf{r}) - \phi_2(\mathbf{r})) \\ &= \frac{1}{2(1-S_{12})} (1 - S_{12} - S_{21} + 1) = 1,\end{aligned}$$

and the orthogonality between ψ_1 and ψ_2 ,

$$\begin{aligned}\langle\psi_1|\psi_2\rangle &= \int d\mathbf{r} \psi_1^*(\mathbf{r})\psi_2(\mathbf{r}) = \int d\mathbf{r} \frac{1}{\sqrt{2(1+S_{12})}} (\phi_1(\mathbf{r}) + \phi_2(\mathbf{r}))^* \frac{1}{\sqrt{2(1-S_{12})}} (\phi_1(\mathbf{r}) - \phi_2(\mathbf{r})) \\ &= \frac{1}{2\sqrt{(1-(S_{12})^2)}} (1 - S_{12} + S_{21} - 1) = 0 = \langle\psi_2|\psi_1\rangle^*.\end{aligned}$$

Thus, we conclude that ψ_1 and ψ_2 form an orthonormal set.

2.2.6 Excited Determinants

2.2.7 Form of the Exact Wave Function and Configuration Interaction

Exercise 2.7

A minimal basis set for benzene consists of 72 spin orbitals. Calculate the size of the full CI matrix if it would be formed from determinants. How many singly excited determinants are there? How many doubly excited determinants are there?

Solution 2.7

To begin with, a benzene molecule consists of 6 carbon atoms, each contributing 6 electrons, and 6 hydrogen atoms, each contributing 1 electron. Consequently, the total number of electrons in a benzene molecule is calculated as $6 \times 6 + 6 \times 1 = 42$ electrons. Namely, $N = 42$.

Secondly, the minimal basis set of benzene includes 36 spatial orbitals. Each carbon atom provides its 1s, 2s, and three 2p orbitals (i.e., $2p_x$, $2p_y$, and $2p_z$), while each hydrogen atom contributes its 1s orbital. These 36 spatial orbitals can be used to construct 72 spin orbitals. Namely, $2K = 72$.

Thus, there are $\binom{72}{42} = 164307576757973059488$ determinants in full CI calculation. Besides, there are $\binom{42}{1}\binom{30}{1} = 1260$ singly excited determinants and $\binom{42}{2}\binom{30}{2} = 374535$ doubly excited determinants.

2.3 Operators and Matrix Elements

2.3.1 Minimal Basis H_2 Matrix Elements

Exercise 2.8

Show that

$$\langle\Psi_{12}^{34}|\mathcal{O}_1|\Psi_{12}^{34}\rangle = \langle 3|h|3\rangle + \langle 4|h|4\rangle$$

and

$$\langle\Psi_0|\mathcal{O}_1|\Psi_{12}^{34}\rangle = \langle\Psi_{12}^{34}|\mathcal{O}_1|\Psi_0\rangle = 0.$$

Solution 2.8

For $\langle\Psi_{12}^{34}|\mathcal{O}_1|\Psi_{12}^{34}\rangle$, it can be divided into two parts, too, viz.,

$$\langle\Psi_{12}^{34}|\mathcal{O}_1|\Psi_{12}^{34}\rangle = \langle\Psi_{12}^{34}|h(1)|\Psi_{12}^{34}\rangle + \langle\Psi_{12}^{34}|h(2)|\Psi_{12}^{34}\rangle.$$

Its first part is

$$\begin{aligned}&\langle\Psi_{12}^{34}|h(1)|\Psi_{12}^{34}\rangle \\ &= \int d\mathbf{x}_1 \int d\mathbf{x}_2 \frac{1}{\sqrt{2}} [\chi_3(1)\chi_4(2) - \chi_4(1)\chi_3(2)]^* h(1) \frac{1}{\sqrt{2}} [\chi_3(1)\chi_4(2) - \chi_4(1)\chi_3(2)]\end{aligned}$$

$$\begin{aligned}
&= \frac{1}{2} \left[\int d\mathbf{x}_1 \chi_3^*(1) h(1) \chi_3(1) \int d\mathbf{x}_2 \chi_4^*(2) \chi_4(2) - \int d\mathbf{x}_1 \chi_3^*(1) h(1) \chi_4(1) \int d\mathbf{x}_2 \chi_4^*(2) \chi_3(2) \right. \\
&\quad \left. - \int d\mathbf{x}_1 \chi_4^*(1) h(1) \chi_3(1) \int d\mathbf{x}_2 \chi_3^*(2) \chi_4(2) + \int d\mathbf{x}_1 \chi_4^*(1) h(1) \chi_4(1) \int d\mathbf{x}_2 \chi_3^*(2) \chi_3(2) \right] \\
&= \frac{1}{2} \left[\int d\mathbf{x}_1 \chi_3^*(1) h(1) \chi_3(1) + 0 + 0 + \int d\mathbf{x}_1 \chi_4^*(1) h(1) \chi_4(1) \right] = \frac{1}{2} (\langle 3|h|3 \rangle + \langle 4|h|4 \rangle).
\end{aligned}$$

Similarly, we can obtain that

$$\langle \Psi_{12}^{34} | h(2) | \psi_{12}^{34} \rangle = \frac{1}{2} (\langle 3|h|3 \rangle + \langle 4|h|4 \rangle).$$

Thus,

$$\langle \Psi_{12}^{34} | \mathcal{O}_1 | \Psi_{12}^{34} \rangle = \frac{1}{2} (\langle 3|h|3 \rangle + \langle 4|h|4 \rangle) + \frac{1}{2} (\langle 3|h|3 \rangle + \langle 4|h|4 \rangle) = \langle 3|h|3 \rangle + \langle 4|h|4 \rangle. \quad (2.8-1)$$

Besides, in the same way, we obtain that

$$\begin{aligned}
&\langle \Psi_{12}^{34} | h(1) | \Psi_0 \rangle \\
&= \int d\mathbf{x}_1 \int d\mathbf{x}_2 \frac{1}{\sqrt{2}} [\chi_3(1) \chi_4(2) - \chi_4(1) \chi_3(2)]^* h(1) \frac{1}{\sqrt{2}} [\chi_1(1) \chi_2(2) - \chi_2(1) \chi_1(2)] \\
&= \frac{1}{2} \left[\int d\mathbf{x}_1 \chi_3^*(1) h(1) \chi_1(1) \int d\mathbf{x}_2 \chi_4^*(2) \chi_2(2) - \int d\mathbf{x}_1 \chi_3^*(1) h(1) \chi_2(1) \int d\mathbf{x}_2 \chi_4^*(2) \chi_1(2) \right. \\
&\quad \left. - \int d\mathbf{x}_1 \chi_4^*(1) h(1) \chi_1(1) \int d\mathbf{x}_2 \chi_3^*(2) \chi_2(2) + \int d\mathbf{x}_1 \chi_4^*(1) h(1) \chi_2(1) \int d\mathbf{x}_2 \chi_3^*(2) \chi_1(2) \right] \\
&= \frac{1}{2} [0 - 0 - 0 + 0] = 0.
\end{aligned}$$

and

$$\langle \Psi_{12}^{34} | h(2) | \Psi_0 \rangle = 0,$$

Therefore,

$$\langle \Psi_{12}^{34} | h(1) | \Psi_{12}^{34} \rangle = \langle \Psi_{12}^{34} | h(1) | \Psi_0 \rangle + \langle \Psi_{12}^{34} | h(2) | \Psi_0 \rangle = 0 + 0 = 0, \quad (2.8-2)$$

and

$$\langle \Psi_0 | \mathcal{O}_1 | \Psi_{12}^{34} \rangle = \langle \Psi_{12}^{34} | \mathcal{O}_1 | \Psi_0 \rangle^* = 0^* = 0. \quad (2.8-3)$$

Exercise 2.9

Using the above approach, show that the full CI matrix for minimal basis H_2 is

$$\mathcal{H} = \begin{pmatrix} \langle 1|h|1 \rangle + \langle 2|h|2 \rangle + \langle 12|12 \rangle - \langle 12|21 \rangle & \langle 12|34 \rangle - \langle 12|43 \rangle \\ \langle 34|12 \rangle - \langle 34|21 \rangle & \langle 3|h|3 \rangle + \langle 4|h|4 \rangle + \langle 34|34 \rangle - \langle 34|43 \rangle \end{pmatrix}.$$

and that it is Hermitian.

Solution 2.9

From (2.92), we know that

$$\langle \Psi_0 | \mathcal{H} | \Psi_0 \rangle = \langle 1|h|1 \rangle + \langle 2|h|2 \rangle + \langle 12|12 \rangle - \langle 12|21 \rangle.$$

For $\langle \Psi_0 | \mathcal{H} | \Psi_{12}^{34} \rangle$, from Exercise 2.8, we know

$$\langle \Psi_0 | \mathcal{O}_1 | \Psi_{12}^{34} \rangle = 0.$$

Besides,

$$\begin{aligned}
&\langle \Psi_0 | \mathcal{O}_2 | \Psi_{12}^{34} \rangle \\
&= \int d\mathbf{x}_1 \int d\mathbf{x}_2 \frac{1}{\sqrt{2}} [\chi_1(1) \chi_2(2) - \chi_2(1) \chi_1(2)]^* r_{12}^{-1} \frac{1}{\sqrt{2}} [\chi_3(1) \chi_4(2) - \chi_4(1) \chi_3(2)]
\end{aligned}$$

$$\begin{aligned}
&= \frac{1}{2} \left[\int d\mathbf{x}_1 \int d\mathbf{x}_2 \chi_1^*(1) \chi_2^*(2) r_{12}^{-1} \chi_3(1) \chi_4(2) - \int d\mathbf{x}_1 \int d\mathbf{x}_2 \chi_1^*(1) \chi_2^*(2) r_{12}^{-1} \chi_4(1) \chi_3(2) \right. \\
&\quad \left. - \int d\mathbf{x}_1 \int d\mathbf{x}_2 \chi_2^*(1) \chi_1^*(2) r_{12}^{-1} \chi_3(1) \chi_4(2) + \int d\mathbf{x}_1 \int d\mathbf{x}_2 \chi_2^*(1) \chi_1^*(2) r_{12}^{-1} \chi_4(1) \chi_3(2) \right] \\
&= \frac{1}{2} (\langle 12|34 \rangle - \langle 12|43 \rangle - \langle 21|34 \rangle + \langle 21|43 \rangle) = \frac{1}{2} (\langle 12|34 \rangle - \langle 12|43 \rangle - \langle 12|43 \rangle + \langle 12|34 \rangle) \\
&= \langle 12|34 \rangle - \langle 12|43 \rangle.
\end{aligned}$$

Thus, we know that

$$\langle \Psi_0 | \mathcal{H} | \Psi_{12}^{34} \rangle = \langle \Psi_0 | \mathcal{O}_1 | \Psi_{12}^{34} \rangle + \langle \Psi_0 | \mathcal{O}_2 | \Psi_{12}^{34} \rangle = 0 + \langle 12|34 \rangle - \langle 12|43 \rangle = \langle 12|34 \rangle - \langle 12|43 \rangle, \quad (2.9-1)$$

and

$$\langle \Psi_{12}^{34} | \mathcal{H} | \Psi_0 \rangle = (\langle \Psi_0 | \mathcal{H} | \Psi_{12}^{34} \rangle)^* = \langle 34|12 \rangle - \langle 34|21 \rangle. \quad (2.9-2)$$

At last, for $\langle \Psi_{12}^{34} | \mathcal{H} | \Psi_{12}^{34} \rangle$, from Exercise 2.8,

$$\langle \Psi_{12}^{34} | \mathcal{O}_1 | \Psi_{12}^{34} \rangle = \langle 3|h|3 \rangle + \langle 4|h|4 \rangle.$$

Moreover,

$$\begin{aligned}
&\langle \Psi_{34}^{12} | \mathcal{O}_2 | \Psi_{12}^{34} \rangle \\
&= \int d\mathbf{x}_1 \int d\mathbf{x}_2 \frac{1}{\sqrt{2}} [\chi_3(1) \chi_4(2) - \chi_4(1) \chi_3(2)]^* r_{12}^{-1} \frac{1}{\sqrt{2}} [\chi_3(1) \chi_4(2) - \chi_4(1) \chi_3(2)] \\
&= \frac{1}{2} \left[\int d\mathbf{x}_1 \int d\mathbf{x}_2 \chi_3^*(1) \chi_4^*(2) r_{12}^{-1} \chi_3(1) \chi_4(2) - \int d\mathbf{x}_1 \int d\mathbf{x}_2 \chi_3^*(1) \chi_4^*(2) r_{12}^{-1} \chi_4(1) \chi_3(2) \right. \\
&\quad \left. - \int d\mathbf{x}_1 \int d\mathbf{x}_2 \chi_4^*(1) \chi_3^*(2) r_{12}^{-1} \chi_3(1) \chi_4(2) + \int d\mathbf{x}_1 \int d\mathbf{x}_2 \chi_4^*(1) \chi_3^*(2) r_{12}^{-1} \chi_4(1) \chi_3(2) \right] \\
&= \frac{1}{2} (\langle 34|34 \rangle - \langle 34|43 \rangle - \langle 43|34 \rangle + \langle 43|43 \rangle) = \frac{1}{2} (\langle 34|34 \rangle - \langle 34|43 \rangle - \langle 34|43 \rangle + \langle 34|34 \rangle) \\
&= \langle 34|34 \rangle - \langle 34|43 \rangle.
\end{aligned}$$

Hence,

$$\langle \Psi_{12}^{34} | \mathcal{H} | \Psi_{12}^{34} \rangle = \langle \Psi_{12}^{34} | \mathcal{O}_1 | \Psi_{12}^{34} \rangle + \langle \Psi_{12}^{34} | \mathcal{O}_2 | \Psi_{12}^{34} \rangle = \langle 3|h|3 \rangle + \langle 4|h|4 \rangle + \langle 34|34 \rangle - \langle 34|43 \rangle. \quad (2.9-3)$$

In conclusion, we have proved that

$$\mathcal{H} = \begin{pmatrix} \langle 1|h|1 \rangle + \langle 2|h|2 \rangle + \langle 12|12 \rangle - \langle 12|21 \rangle & \langle 12|34 \rangle - \langle 12|43 \rangle \\ \langle 34|12 \rangle - \langle 34|21 \rangle & \langle 3|h|3 \rangle + \langle 4|h|4 \rangle + \langle 34|34 \rangle - \langle 34|43 \rangle \end{pmatrix}. \quad (2.9-4)$$

Obviously, it is Hermitian.

2.3.2 Notations for One- and Two-Electron Integrals

2.3.3 General Rules for Matrix Elements

Exercise 2.10

Derive Eq.(2.110) from Eq.(2.107).

Solution 2.10

We derive Eqs.(2.109a, b) firstly, then derive other equations. Note that

$$\langle mm || mm \rangle = \langle mm | mm \rangle - \langle mm | mm \rangle = 0. \quad (2.10-1)$$

Thus (2.109a) has been proved. Besides, note that

$$\langle mn || mn \rangle = \langle mn | mn \rangle - \langle mn | nm \rangle = \langle nm | nm \rangle - \langle nm | mn \rangle = \langle nm || nm \rangle. \quad (2.10-2)$$

Thus, (2.109b) has been proved. From Eq.(2.107), with $\langle mm||mm\rangle = 0$, we find that

$$\begin{aligned}
 \langle K|\mathcal{H}|K\rangle &= \sum_m^N \langle m|h|m\rangle + \frac{1}{2} \sum_m^N \sum_n^N \langle mn||mn\rangle \\
 &= \sum_m^N \langle m|h|m\rangle + \frac{1}{2} \sum_m^N \sum_{n>m}^N \langle mn||mn\rangle + \frac{1}{2} \sum_n^N \sum_{m>n}^N \langle mn||mn\rangle \\
 &= \sum_m^N \langle m|h|m\rangle + \frac{1}{2} \sum_m^N \sum_{n>m}^N \langle mn||mn\rangle + \frac{1}{2} \sum_m^N \sum_{n>m}^N \langle nm||nm\rangle \\
 &= \sum_m^N \langle m|h|m\rangle + \sum_m^N \sum_{n>m}^N \langle mn||mn\rangle.
 \end{aligned}$$

The first line of (2.110) has been verified. Then,

$$\begin{aligned}
 \langle K|\mathcal{H}|K\rangle &= \sum_m^N \langle m|h|m\rangle + \sum_m^N \sum_{n>m}^N \langle mn||mn\rangle = \sum_m^N \langle m|h|m\rangle + \sum_m^N \sum_{n>m}^N \langle mn|mn\rangle - \langle mn|nm\rangle \\
 &= \sum_m^N [m|h|m] + \sum_m^N \sum_{n>m}^N [mm|nn] - [mn|nm].
 \end{aligned}$$

Exercise 2.11

If $|K\rangle = |\chi_1\chi_2\chi_3\rangle$ show that

$$\langle K|\mathcal{H}|K\rangle = \langle 1|h|1\rangle + \langle 2|h|2\rangle + \langle 3|h|3\rangle + \langle 12||12\rangle + \langle 13||13\rangle + \langle 23||23\rangle.$$

Solution 2.11

From the conclusion of Exercise 2.10, we immediately obtain that

$$\langle K|\mathcal{H}|K\rangle = \langle 1|h|1\rangle + \langle 2|h|2\rangle + \langle 3|h|3\rangle + \langle 12||12\rangle + \langle 13||13\rangle + \langle 23||23\rangle.$$

Exercise 2.12

Evaluate the matrix elements that occur in the minimal basis H_2 full CI matrix (Eq.(2.79)) using the rules. Compare with the result obtained in Exercise 2.9.

Solution 2.12

$\langle \Psi_0|\mathcal{H}|\Psi_0\rangle$ has been delivered by (2.111) and (2.114). Applying the rules, we get that

$$\begin{aligned}
 \langle \Psi_0|\mathcal{H}|\Psi_{12}^{34}\rangle &= \langle \Psi_0|\mathcal{O}_1|\Psi_{12}^{34}\rangle + \langle \Psi_0|\mathcal{O}_2|\Psi_{12}^{34}\rangle = 0 + \langle 12||34\rangle = \langle 12||34\rangle, \\
 \langle \Psi_{12}^{34}|\mathcal{H}|\Psi_0\rangle &= \langle \Psi_{12}^{34}|\mathcal{O}_1|\Psi_0\rangle + \langle \Psi_{12}^{34}|\mathcal{O}_2|\Psi_0\rangle = 0 + \langle 34||12\rangle = \langle 34||12\rangle, \\
 \langle \Psi_{12}^{34}|\mathcal{H}|\Psi_{12}^{34}\rangle &= \langle \Psi_{12}^{34}|\mathcal{O}_1|\Psi_{12}^{34}\rangle + \langle \Psi_{12}^{34}|\mathcal{O}_2|\Psi_{12}^{34}\rangle = \langle 3|h|3\rangle + \langle 4|h|4\rangle + \langle 34||34\rangle.
 \end{aligned}$$

This result equals that of Exercise 2.9.

Exercise 2.13

Show that

$$\langle \Psi_a^r|\mathcal{O}_1|\Psi_b^s\rangle = \begin{cases} 0, & \text{if } a \neq b, r \neq s; \\ \langle r|h|s\rangle, & \text{if } a = b, r \neq s; \\ -\langle b|h|a\rangle, & \text{if } a \neq b, r = s; \\ \sum_c^N \langle c|h|c\rangle - \langle a|h|a\rangle + \langle r|h|r\rangle & \text{if } a = b, r = s. \end{cases}$$

Solution 2.13

Note that this result is not correct until the two determinants are in maximum coincidence! We talk about the final result according to the different occupation of the bra and the ket.

- When $a \neq b$ and $r \neq s$, the two determinants differ by two spin orbitals, and thus

$$\langle \Psi_a^r | \mathcal{O}_1 | \Psi_b^s \rangle = 0.$$

- When $a = b$ and $r \neq s$, the two determinants differ by one spin orbital, and thus

$$\langle \Psi_a^r | \mathcal{O}_1 | \Psi_b^s \rangle = \langle \Psi_a^r | \mathcal{O}_1 | \Psi_a^s \rangle = \langle r | h | s \rangle.$$

- When $a \neq b$ and $r = s$, the two determinants differ by one spin orbital, and thus $\langle \Psi_a^r | \mathcal{O}_1 | \Psi_b^s \rangle = \langle \Psi_a^r | \mathcal{O}_1 | \Psi_b^r \rangle$. Note that

$$\langle \Psi_a^r | \mathcal{O}_1 | \Psi_b^r \rangle = \langle \cdots \chi_r \chi_b \cdots | \mathcal{O}_1 | \cdots \chi_a \chi_r \cdots \rangle = -\langle \cdots \chi_r \chi_b \cdots | \mathcal{O}_1 | \cdots \chi_r \chi_a \cdots \rangle = -\langle r | h | a \rangle.$$

- When $a = b$ and $r = s$, the two determinants are the same, and thus

$$\langle \Psi_a^r | \mathcal{O}_1 | \Psi_b^s \rangle = \langle \Psi_a^r | \mathcal{O}_1 | \Psi_a^r \rangle = \langle \cdots \chi_{a \rightarrow r} \cdots | \mathcal{O}_1 | \cdots \chi_{a \rightarrow r} \cdots \rangle = \sum_{c \neq a} \langle c | h | c \rangle + \langle r | h | r \rangle.$$

In other words,

$$\langle \Psi_a^r | \mathcal{O}_1 | \Psi_b^s \rangle = \sum_c \langle c | h | c \rangle - \langle a | h | a \rangle + \langle r | h | r \rangle.$$

In conclusion, we get that

$$\langle \Psi_a^r | \mathcal{O}_1 | \Psi_b^s \rangle = \begin{cases} 0, & \text{if } a \neq b, r \neq s; \\ \langle r | h | s \rangle, & \text{if } a = b, r \neq s; \\ -\langle b | h | a \rangle, & \text{if } a \neq b, r = s; \\ \sum_c^N \langle c | h | c \rangle - \langle a | h | a \rangle + \langle r | h | r \rangle & \text{if } a = b, r = s. \end{cases} \quad (2.13-1)$$

Exercise 2.14

The Hartree-Fock ground state energy for an N -electron system is ${}^N E_0 = \langle {}^N \Psi_0 | \mathcal{H} | {}^N \Psi_0 \rangle$. Consider a state of the ionized system (in which an electron has been removed from spin orbital χ_a) with energy ${}^{N-1} E_a = \langle {}^{N-1} \Psi_a | \mathcal{H} | {}^{N-1} \Psi_a \rangle$, where $|{}^{N-1} \Psi_a\rangle$ is a single determinant with all spin orbitals but χ_a occupied,

$$|{}^{N-1} \Psi_a\rangle = |\chi_1 \chi_2 \cdots \chi_{a-1} \chi_{a+1} \cdots \chi_N\rangle.$$

Show, using the rules in the tables, that the energy required for this ionization process is

$${}^N E_0 - {}^{N-1} E_a = \langle a | h | a \rangle + \sum_b^N \langle ab | ab \rangle.$$

Solution 2.14

The verification is direct.

$$\begin{aligned} {}^N E_0 - {}^{N-1} E_0 &= \left[\sum_b^N \langle b | h | b \rangle + \frac{1}{2} \sum_c^N \sum_d^N \langle cd | cd \rangle \right] - \left[\sum_{b \neq a}^N \langle b | h | b \rangle + \frac{1}{2} \sum_{c \neq a}^N \sum_{d \neq a}^N \langle cd | cd \rangle \right] \\ &= \left[\sum_b^N \langle b | h | b \rangle - \sum_{b \neq a}^N \langle b | h | b \rangle \right] + \frac{1}{2} \left[\sum_c^N \sum_d^N \langle cd | cd \rangle - \sum_{c \neq a}^N \sum_{d \neq a}^N \langle cd | cd \rangle \right] \\ &= \langle a | h | a \rangle + \frac{1}{2} \left[\sum_d^N \langle ad | ad \rangle + \sum_c^N \langle ca | ca \rangle - \langle aa | aa \rangle \right] \end{aligned}$$

$$= \langle a|h|a \rangle + \frac{1}{2} \left[\sum_b^N \langle ab||ab \rangle + \sum_b^N \langle ab||ab \rangle \right] = \langle a|h|a \rangle + \sum_b^N \langle ab||ab \rangle.$$

2.3.4 Derivation of the Rules for Matrix Elements

Exercise 2.15

Generalize the result of Exercise 2.4 to N -electron Slater determinants. Show that the Slater determinant $|\chi_i \chi_j \cdots \chi_k\rangle$ formed from spin orbitals, which are eigenfunctions of the one-electron operator h as in Eq.(2.29), is an eigenfunction of the independent-electron Hamiltonian (2.28), $\mathcal{H} = \sum_{i=1}^N h(i)$, with an eigenvalue $\varepsilon_i + \varepsilon_j + \cdots + \varepsilon_k$. *Hint:* Since \mathcal{H} is invariant to permutations of the electron labels, it commutes with the permutation operator \mathcal{P}_n .

Solution 2.15

Note that a Slater determinant is a linear combination of $N!$ Hartree product. With the conclusion of Exercise 2.2 and the truth that \mathcal{H} commutes with the permutation operator \mathcal{P}_n , we find that

$$\begin{aligned} \mathcal{H}\Psi &= \mathcal{H} \frac{1}{\sqrt{N!}} \sum_n^{N!} (-1)^{p_n} \mathcal{P}_n \prod_{j=1}^N \chi_{j'}(\mathbf{x}_j) = \frac{1}{\sqrt{N!}} \sum_n^{N!} (-1)^{p_n} \mathcal{P}_n \mathcal{H} \prod_{j=1}^N \chi_{j'}(\mathbf{x}_j) \\ &= \frac{1}{\sqrt{N!}} \sum_n^{N!} (-1)^{p_n} \mathcal{P}_n \sum_{i=1}^N \varepsilon_i \prod_{j=1}^N \chi_{j'}(\mathbf{x}_j) = \sum_{i=1}^N \varepsilon_i \frac{1}{\sqrt{N!}} \sum_n^{N!} (-1)^{p_n} \mathcal{P}_n \prod_{j=1}^N \chi_{j'}(\mathbf{x}_j) = \sum_{i=1}^N \varepsilon_i \Psi. \end{aligned}$$

Exercise 2.16

A different procedure for deriving the above matrix elements uses the theorem that $\langle K|\mathcal{H}|L\rangle = (N!)^{1/2} \langle K^{\text{HP}}|\mathcal{H}|L\rangle$ where $|K^{\text{HP}}\rangle$ is the Hartree product corresponding to the determinant $|K\rangle$, i.e.,

$$|K\rangle = |\chi_m(\mathbf{x}_1)\chi_n(\mathbf{x}_2)\cdots\rangle$$

and

$$|K^{\text{HP}}\rangle = \chi_m(\mathbf{x}_1)\chi_n(\mathbf{x}_2)\cdots.$$

Prove this theorem. Use it to derive the matrix elements of a sum of one-electron operators.

Solution 2.16

Firstly, note that for a general permutation operator \mathcal{P}_k and a general ket $|L\rangle$, due to the antisymmetry of the ket, we have that

$$(-1)^{p_k} \mathcal{P}_k \langle K^{\text{HP}}|\mathcal{H}|L\rangle = \langle K^{\text{HP}}|\mathcal{H}|L\rangle.$$

Thus, we find that

$$\begin{aligned} \langle K|\mathcal{H}|L\rangle &= \frac{1}{\sqrt{N!}} \sum_{k=1}^{N!} (-1)^{p_k} \mathcal{P}_k \langle K^{\text{HP}}|\mathcal{H}|L\rangle = \frac{1}{\sqrt{N!}} \sum_{k=1}^{N!} \langle K^{\text{HP}}|\mathcal{H}|L\rangle \\ &= \frac{1}{\sqrt{N!}} N! \langle K^{\text{HP}}|\mathcal{H}|L\rangle = \sqrt{N!} \langle K^{\text{HP}}|\mathcal{H}|L\rangle. \end{aligned}$$

Hence, for the operator \mathcal{O}_1 , which is a sum of one-electron operators o , we get that

$$\begin{aligned} \langle K|\mathcal{O}_1|L\rangle &= \sqrt{N!} \langle K^{\text{HP}}|\mathcal{O}_1|L\rangle \\ &= \frac{\sqrt{N!}}{\sqrt{N!}} \sum_j (-1)^{p_j} \int d\mathbf{x}_1 \int d\mathbf{x}_2 \cdots \int d\mathbf{x}_N \{ \chi_m^*(1)\chi_n^*(2)\cdots\chi_l^*(N) \} \sum_i o(i) \mathcal{P}_j \{ \chi_p(1)\chi_q(2)\cdots\chi_r(N) \} \\ &= \sum_i \sum_j (-1)^{p_j} \int d\mathbf{x}_1 \int d\mathbf{x}_2 \cdots \int d\mathbf{x}_N \{ \chi_m^*(1)\chi_n^*(2)\cdots\chi_l^*(N) \} o(i) \mathcal{P}_j \{ \chi_p(1)\chi_q(2)\cdots\chi_r(N) \}. \end{aligned}$$

Here, for a given Hartree product K^{HP} , the maximum coincidence is dependent of the number of spin orbitals which $|K\rangle$ and $|L\rangle$ differ.

- When $|K\rangle$ and $|L\rangle$ are the same, at this time, there are N ways for the maximum coincidence and all of them are using an even permutation, thus,

$$\langle K|\mathcal{O}_1|L\rangle = \sum_i \int d\mathbf{x}_i \chi_m^*(i) o(i) \chi_m(i) \int d\mathbf{x}_1 \chi_n^*(1) \chi_n(1) \cdots \int d\mathbf{x}_N \chi_l^*(N) \chi_l(N) = \sum_i \langle i|o|i\rangle.$$

- When $|K\rangle$ and $|L\rangle$ differ by one spin orbital, at this time, there is only one way for the maximum coincidence and it is using an even permutation, viz.,

$$\langle K|\mathcal{O}_1|L\rangle = \sum_i \int d\mathbf{x}_i \chi_m^*(i) o(i) \chi_p(i) \int d\mathbf{x}_1 \chi_n^*(1) \chi_n(1) \cdots \int d\mathbf{x}_N \chi_l^*(N) \chi_l(N) = \langle m|o|p\rangle.$$

- When $|K\rangle$ and $|L\rangle$ differ by more than one spin orbital, at this time, there is no way for the maximum coincidence and thus,

$$\langle K|\mathcal{O}_1|L\rangle = 0.$$

2.3.5 Transition from Spin Orbitals to Spatial Orbitals

Exercise 2.17

By integrating out spin, show that the full CI matrix for minimal basis H_2 (see Exercise 2.9) is

$$\mathbf{H} = \begin{pmatrix} 2(1|h|1) + (11|11) & (12|12) \\ (21|21) & 2(2|h|2) + (22|22) \end{pmatrix}.$$

Solution 2.17

Note that the general reduction using physicists' notation is

$$\langle ij|kl\rangle = \langle \bar{i}\bar{j}|\bar{k}\bar{l}\rangle = \langle i\bar{j}|k\bar{l}\rangle = \langle \bar{i}\bar{j}|\bar{k}l\rangle = (ik|jl). \quad (2.17-1)$$

From the conclusion of Exercise 2.9, we find that

$$\begin{aligned} H_{11} &= \langle 1|h|1\rangle + \langle \bar{1}|h|\bar{1}\rangle + \langle 1\bar{1}|1\bar{1}\rangle - \langle 1\bar{1}|\bar{1}1\rangle = (1|h|1) + (1|h|1) + (11|11) - 0 = 2(1|h|1) + (11|11), \\ H_{12} &= \langle 1\bar{1}|2\bar{2}\rangle - \langle 1\bar{1}|\bar{2}2\rangle = (12|12) - 0 = (12|12), \\ H_{21} &= \langle 2\bar{2}|1\bar{1}\rangle - \langle 2\bar{2}|\bar{1}1\rangle = (21|21) - 0 = (21|21), \\ H_{22} &= \langle 2|h|2\rangle + \langle \bar{2}|h|\bar{2}\rangle + \langle 2\bar{2}|2\bar{2}\rangle - \langle 2\bar{2}|\bar{2}2\rangle = (2|h|2) + (2|h|2) + (22|22) - 0 = 2(2|h|2) + (22|22). \end{aligned}$$

Exercise 2.18

In Chapter 6, where we consider perturbation theory, we show that the leading correction to the Hartree-Fock ground state energy is

$$E_0^{[2]} = \frac{1}{4} \sum_{abrs} \frac{|\langle ab||rs\rangle|^2}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s}.$$

Show that for a closed-shell system (where $\varepsilon_i = \varepsilon_{\bar{i}}$) this becomes

$$E_0^{[2]} = \sum_{a,b=1}^{N/2} \sum_{r,s=(N/2+1)}^K \frac{\langle ab||rs\rangle (2\langle rs|ab\rangle - \langle rs|ba\rangle)}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s}.$$

Solution 2.18

The simplification is direct.

$$E_0^{(2)} = \frac{1}{4} \sum_{abrs} \frac{|\langle ab||rs\rangle|^2}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} = \frac{1}{4} \sum_{abrs} \frac{\langle ab||rs\rangle \langle rs||ab\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} = \frac{1}{4} \sum_{abrs} \frac{(\langle ab||rs\rangle - \langle ab||sr\rangle)(\langle rs||ab\rangle - \langle rs||ba\rangle)}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s}$$

$$\begin{aligned}
&= \frac{1}{4} \sum_{abrs} \frac{\langle ab|rs\rangle\langle rs|ab\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} - \frac{\langle ab|rs\rangle\langle rs|ba\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} - \frac{\langle ab|sr\rangle\langle rs|ab\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} + \frac{\langle ab|sr\rangle\langle rs|ba\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} \\
&= \frac{1}{4} \sum_{abrs} \frac{\langle ab|rs\rangle\langle rs|ab\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} - \frac{\langle ab|rs\rangle\langle rs|ba\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} - \frac{\langle ab|rs\rangle\langle sr|ab\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_s - \varepsilon_r} + \frac{\langle ab|rs\rangle\langle sr|ba\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_s - \varepsilon_r} \\
&= \frac{1}{4} \sum_{abrs} \frac{2\langle ab|rs\rangle\langle rs|ab\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} - \frac{2\langle ab|rs\rangle\langle rs|ba\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} = \frac{1}{2} \sum_{abrs} \frac{\langle ab|rs\rangle\langle rs|ab\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} - \frac{\langle ab|rs\rangle\langle rs|ba\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s}.
\end{aligned}$$

Note that for a closed-shell system, only six terms in the numerators can be nonzero, viz.,

$$\begin{aligned}
E_0^{(2)} &= \frac{1}{2} \sum_{abrs} \left(\frac{\langle ab|rs\rangle\langle rs|ab\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} - \frac{\langle ab|rs\rangle\langle rs|ba\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} \right) \\
&= \frac{1}{2} \left(\sum_{abrs}^{N/2} \frac{\langle ab|rs\rangle\langle rs|ab\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} - \frac{\langle ab|rs\rangle\langle rs|ba\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} + \sum_{\bar{a}\bar{b}\bar{r}\bar{s}}^{N/2} \frac{\langle \bar{a}\bar{b}|\bar{r}\bar{s}\rangle\langle \bar{r}\bar{s}|\bar{a}\bar{b}\rangle}{\varepsilon_{\bar{a}} + \varepsilon_{\bar{b}} - \varepsilon_{\bar{r}} - \varepsilon_{\bar{s}}} - \frac{\langle \bar{a}\bar{b}|\bar{r}\bar{s}\rangle\langle \bar{r}\bar{s}|\bar{b}\bar{a}\rangle}{\varepsilon_{\bar{a}} + \varepsilon_{\bar{b}} - \varepsilon_{\bar{r}} - \varepsilon_{\bar{s}}} \right. \\
&\quad + \sum_{\bar{a}\bar{b}\bar{r}\bar{s}}^{N/2} \frac{\langle \bar{a}\bar{b}|\bar{r}\bar{s}\rangle\langle \bar{r}\bar{s}|\bar{a}\bar{b}\rangle}{\varepsilon_{\bar{a}} + \varepsilon_{\bar{b}} - \varepsilon_{\bar{r}} - \varepsilon_{\bar{s}}} - \frac{\langle \bar{a}\bar{b}|\bar{r}\bar{s}\rangle\langle \bar{r}\bar{s}|\bar{b}\bar{a}\rangle}{\varepsilon_{\bar{a}} + \varepsilon_{\bar{b}} - \varepsilon_{\bar{r}} - \varepsilon_{\bar{s}}} + \sum_{\bar{a}\bar{b}\bar{r}\bar{s}}^{N/2} \frac{\langle \bar{a}\bar{b}|\bar{r}\bar{s}\rangle\langle \bar{r}\bar{s}|\bar{a}\bar{b}\rangle}{\varepsilon_{\bar{a}} + \varepsilon_{\bar{b}} - \varepsilon_{\bar{r}} - \varepsilon_{\bar{s}}} - \frac{\langle \bar{a}\bar{b}|\bar{r}\bar{s}\rangle\langle \bar{r}\bar{s}|\bar{b}\bar{a}\rangle}{\varepsilon_{\bar{a}} + \varepsilon_{\bar{b}} - \varepsilon_{\bar{r}} - \varepsilon_{\bar{s}}} \Big) \\
&= \frac{1}{2} \left(\sum_{abrs}^{N/2} \frac{\langle ab|rs\rangle\langle rs|ab\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} - \frac{\langle ab|rs\rangle\langle rs|ba\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} + \sum_{abrs}^{N/2} \frac{\langle ab|rs\rangle\langle rs|ab\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} - 0 + 0 - 0 \right. \\
&\quad \left. + 0 - 0 + \sum_{abrs}^{N/2} \frac{\langle ab|rs\rangle\langle rs|ab\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} - 0 + \sum_{abrs}^{N/2} \frac{\langle ab|rs\rangle\langle rs|ab\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} - \frac{\langle ab|rs\rangle\langle rs|ba\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} \right) \\
&= 2 \sum_{abrs}^{N/2} \frac{\langle ab|rs\rangle\langle rs|ab\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} - \frac{\langle ab|rs\rangle\langle rs|ba\rangle}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} = \sum_{abrs}^{N/2} \frac{\langle ab|rs\rangle(2\langle rs|ab\rangle - \langle rs|ba\rangle)}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s},
\end{aligned}$$

where

$$\sum_{abrs}^{N/2} \equiv \sum_{a,b=1}^{N/2} \sum_{r,s=(N/2+1)}^K.$$

2.3.6 Coulomb and Exchange Integrals

Exercise 2.19

Prove the following properties of coulomb and exchange integrals

$$J_{ii} = K_{ii}, \quad J_{ij}^* = J_{ij}, \quad K_{ij}^* = K_{ij}, \quad J_{ij} = J_{ji}, \quad K_{ij} = K_{ji}.$$

Solution 2.19

Using (2.182) and (2.184), we get that

$$\begin{aligned}
J_{ii} &= (ii|ii) = K_{ii}, \\
J_{ij}^* &= (ii|jj)^* = (ij|ij) = J_{ij}, \quad K_{ij}^* = (ij|ji)^* = (ji|ij) = (ij|ji) = K_{ij}, \\
J_{ij} &= (ii|jj) = (jj|ii) = J_{ji}, \quad K_{ij} = (ij|ji) = (ji|ij) = K_{ji}.
\end{aligned}$$

Exercise 2.20

Show that for *real* spatial orbitals

$$K_{ij} = (ij|ij) = (ji|ji) = \langle ii|jj\rangle = \langle jj|ii\rangle.$$

Solution 2.20

With real spatial orbitals, we find that

$$\begin{aligned} K_{ij} &= (ij|ji) = (ij|ij) = \langle ii|jj \rangle \\ &= (ji|ji) = \langle jj|ii \rangle. \end{aligned}$$

Exercise 2.21

Show that the full CI matrix for minimal basis H_2 (see Exercise 2.17) is

$$\mathbf{H} = \begin{pmatrix} 2h_{11} + J_{11} & K_{12} \\ K_{12} & 2h_{22} + J_{22} \end{pmatrix}.$$

The spatial molecular orbitals of this model are real because they were constructed as linear combinations of real atomic orbitals (see Eqs.(2.54), (2.55), (2.57), and (2.58)).

Solution 2.21

From the conclusion of Exercise 2.17, we find that

$$\begin{aligned} H_{11} &= 2(1|h|1) + (11|11) = 2h_{11} + J_{11}, \\ H_{12} &= (12|12) = K_{12}, \\ H_{21} &= (21|21) = K_{21} = K_{12}, \\ H_{22} &= 2(2|h|2) + (22|22) = 2h_{22} + J_{22}. \end{aligned}$$

Exercise 2.22

Show that the energies of the Hartree products

$$\Psi_{\uparrow\downarrow}^{\text{HP}} = \psi_1(\mathbf{r}_1)\alpha(\omega_1)\psi_2(\mathbf{r}_2)\beta(\omega_2)$$

and

$$\Psi_{\downarrow\downarrow}^{\text{HP}} = \psi_1(\mathbf{r}_1)\beta(\omega_1)\psi_2(\mathbf{r}_2)\beta(\omega_2)$$

are the same and equal to $E(\uparrow\downarrow)$ as to be expected since the motion of electrons with parallel spin in not correlated within the Hartree product approximation to the wave function.

Solution 2.22

The energy of the Hartree product $\Psi_{\uparrow\downarrow}^{\text{HP}}$ is

$$\begin{aligned} E(\Psi_{\uparrow\downarrow}^{\text{HP}}) &= \langle \Psi_{\uparrow\downarrow}^{\text{HP}} | h(1) + h(2) + r_{12}^{-1} | \Psi_{\uparrow\downarrow}^{\text{HP}} \rangle = \int d\mathbf{x}_1 \int d\mathbf{x}_2 \langle \Psi_{\uparrow\downarrow}^{\text{HP}} | \mathbf{x}_1 \rangle \langle \mathbf{x}_1 | h(1) + h(2) + r_{12}^{-1} | \mathbf{x}_2 \rangle \langle \mathbf{x}_2 | \Psi_{\uparrow\downarrow}^{\text{HP}} \rangle \\ &= \int d\mathbf{r}_1 \psi_1^*(\mathbf{r}_1) h(1) \psi_1(\mathbf{r}_1) \int d\mathbf{r}_2 \psi_2^*(\mathbf{r}_2) \psi_2(\mathbf{r}_2) \int d\omega_1 \alpha^*(\omega_1) \alpha(\omega_1) \int d\omega_2 \beta^*(\omega_2) \beta(\omega_2) \\ &\quad + \int d\mathbf{r}_1 \psi_1^*(\mathbf{r}_1) \psi_1(\mathbf{r}_1) \int d\mathbf{r}_2 \psi_2^*(\mathbf{r}_2) h(2) \psi_2(\mathbf{r}_2) \int d\omega_1 \alpha^*(\omega_1) \alpha(\omega_1) \int d\omega_2 \beta^*(\omega_2) \beta(\omega_2) \\ &\quad + \int d\mathbf{r}_1 \int d\mathbf{r}_2 \psi_1^*(\mathbf{r}_1) \psi_1(\mathbf{r}_1) r_{12}^{-1} \psi_2^*(\mathbf{r}_2) \psi_2(\mathbf{r}_2) \int d\omega_1 \alpha^*(\omega_1) \alpha(\omega_1) \int d\omega_2 \beta^*(\omega_2) \beta(\omega_2) \\ &= h_{11} + h_{22} + J_{12}, \end{aligned}$$

while the energy of the Hartree product $\Psi_{\downarrow\downarrow}^{\text{HP}}$ is

$$\begin{aligned} E(\Psi_{\downarrow\downarrow}^{\text{HP}}) &= \langle \Psi_{\downarrow\downarrow}^{\text{HP}} | h(1) + h(2) + r_{12}^{-1} | \Psi_{\downarrow\downarrow}^{\text{HP}} \rangle = \int d\mathbf{x}_1 \int d\mathbf{x}_2 \langle \Psi_{\downarrow\downarrow}^{\text{HP}} | \mathbf{x}_1 \rangle \langle \mathbf{x}_1 | h(1) + h(2) + r_{12}^{-1} | \mathbf{x}_2 \rangle \langle \mathbf{x}_2 | \Psi_{\downarrow\downarrow}^{\text{HP}} \rangle \\ &= \int d\mathbf{r}_1 \psi_1^*(\mathbf{r}_1) h(1) \psi_1(\mathbf{r}_1) \int d\mathbf{r}_2 \psi_2^*(\mathbf{r}_2) \psi_2(\mathbf{r}_2) \int d\omega_1 \beta^*(\omega_1) \beta(\omega_1) \int d\omega_2 \beta^*(\omega_2) \beta(\omega_2) \\ &\quad + \int d\mathbf{r}_1 \psi_1^*(\mathbf{r}_1) \psi_1(\mathbf{r}_1) \int d\mathbf{r}_2 \psi_2^*(\mathbf{r}_2) h(2) \psi_2(\mathbf{r}_2) \int d\omega_1 \beta^*(\omega_1) \beta(\omega_1) \int d\omega_2 \beta^*(\omega_2) \beta(\omega_2) \\ &\quad + \int d\mathbf{r}_1 \int d\mathbf{r}_2 \psi_1^*(\mathbf{r}_1) \psi_1(\mathbf{r}_1) r_{12}^{-1} \psi_2^*(\mathbf{r}_2) \psi_2(\mathbf{r}_2) \int d\omega_1 \beta^*(\omega_1) \beta(\omega_1) \int d\omega_2 \beta^*(\omega_2) \beta(\omega_2) \end{aligned}$$

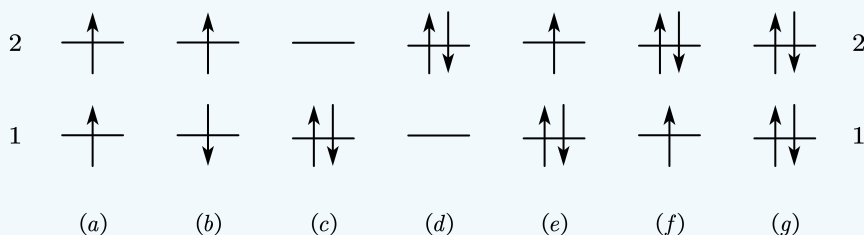
$$= h_{11} + h_{22} + J_{12}.$$

It is evident that they are the same and equal to $E(\uparrow\downarrow)$, namely, (2.186), as to be expected since the motion of electrons with parallel spin is not correlated within the Hartree product approximation to the wave function.

2.3.7 Pseudo-Classical Interpretation of Determinantal Energies

Exercise 2.23

Verify the energies of the following determinants by inspection.



- a. $h_{11} + h_{22} + J_{12} - K_{12}$.
- b. $h_{11} + h_{22} + J_{12}$.
- c. $2h_{11} + J_{11}$.
- d. $2h_{22} + J_{22}$.
- e. $2h_{11} + h_{22} + J_{11} + 2J_{12} - K_{12}$.
- f. $2h_{22} + h_{11} + J_{22} + 2J_{12} - K_{12}$.
- g. $2h_{11} + 2h_{22} + J_{11} + J_{22} + 4J_{12} - 2K_{12}$.

Solution 2.23

- a. There are 2 electrons in spatial orbital 1 and 2, contributing the terms h_{11} and h_{22} . There is only 1 unique pair of electrons in spatial orbitals 1 and 2, contributing the term J_{12} . There is only 1 unique pair of electrons with parallel spins in spatial orbitals 1 and 2, contributing the term $-K_{12}$. Thus,

$$E_a = h_{11} + h_{22} + J_{12} - K_{12}.$$

- b. There are 2 electrons in spatial orbital 1 and 2, contributing the terms h_{11} and h_{22} . There is only 1 unique pair of electrons in spatial orbitals 1 and 2, contributing the term J_{12} . However, there is no unique pair of electrons with parallel spins. Thus,

$$E_b = h_{11} + h_{22} + J_{12}.$$

- c. There are 2 electrons both in spatial orbital 1, contributing the terms $2h_{11}$. There is only 1 unique pair of electrons in spatial orbitals 1, contributing the term J_{11} . However, there is no unique pair of electrons with parallel spins. Thus,

$$E_c = 2h_{11} + J_{11}.$$

- d. There are 2 electrons both in spatial orbital 2, contributing the terms $2h_{22}$. There is only 1 unique pair of electrons in spatial orbitals 1, contributing the term J_{22} . However, there is no unique pair of electrons with parallel spins. Thus,

$$E_d = 2h_{22} + J_{22}.$$

- e. There are 3 electrons in spatial orbital 1, 1 and 2, contributing the terms h_{11} , h_{11} and h_{22} . There are 3 unique pairs of electrons in spatial orbitals 1 and 1, 1 and 2, 1 and 2, contributing the term

J_{11} , J_{12} and J_{12} . There is only 1 unique pair of electrons with parallel spins in spatial orbitals 1 and 2, contributing the term $-K_{12}$. Thus,

$$E_e = h_{11} + h_{11} + h_{22} + J_{11} + J_{12} + J_{12} - K_{12} = 2h_{11} + h_{22} + J_{11} + 2J_{12} - K_{12}.$$

- f. There are 3 electrons in spatial orbital 1, 2 and 2, contributing the terms h_{11} , h_{22} and h_{22} . There are 3 unique pairs of electrons in spatial orbitals 1 and 2, 1 and 2, 2 and 2, contributing the term J_{12} , J_{12} and J_{22} . There is only 1 unique pair of electrons with parallel spins in spatial orbitals 1 and 2, contributing the term $-K_{12}$. Thus,

$$E_f = h_{11} + h_{22} + h_{22} + J_{12} + J_{12} + J_{22} - K_{12} = 2h_{22} + h_{11} + J_{22} + 2J_{12} - K_{12}.$$

- g. There are 4 electrons in spatial orbital 1, 1, 2 and 2, contributing the terms h_{11} , h_{11} , h_{22} and h_{22} . There are 6 unique pairs of electrons in spatial orbitals 1 and 1, 1 and 2, 1 and 2, 1 and 2, 2 and 2, contributing the term J_{11} , J_{12} , J_{12} , J_{12} , J_{12} , and J_{22} . There are only 2 unique pair of electrons with parallel spins in spatial orbitals 1 and 2, 1 and 2, contributing the term $-K_{12}$ and $-K_{12}$. Thus,

$$\begin{aligned} E_g &= h_{11} + h_{11} + h_{22} + h_{22} + J_{11} + J_{12} + J_{12} + J_{12} + J_{12} + J_{22} - K_{12} - K_{12} \\ &= 2h_{11} + 2h_{22} + J_{11} + J_{22} + 4J_{12} - 2K_{12}. \end{aligned}$$

2.4 Second Quantization

2.4.1 Creation and Annihilation Operators and Their Anticommutation Relations

Exercise 2.24

Show, using the properties of determinants, that

$$(a_1^\dagger a_2^\dagger + a_2^\dagger a_1^\dagger)|K\rangle = 0$$

for every $|K\rangle$ in the set $\{|\chi_1\chi_2\rangle, |\chi_1\chi_3\rangle, |\chi_1\chi_4\rangle, |\chi_2\chi_3\rangle, |\chi_2\chi_4\rangle, |\chi_3\chi_4\rangle\}$.

Solution 2.24

Using (2.194), the conclusion can be obtained at once.

Exercise 2.25

Show, using the properties of determinants, that

$$\begin{aligned} (a_1 a_2^\dagger + a_2^\dagger a_1)|K\rangle &= 0, \\ (a_1 a_1^\dagger + a_1^\dagger a_1)|K\rangle &= |K\rangle \end{aligned}$$

for every $|K\rangle$ in the set $\{|\chi_1\chi_2\rangle, |\chi_1\chi_3\rangle, |\chi_1\chi_4\rangle, |\chi_2\chi_3\rangle, |\chi_2\chi_4\rangle, |\chi_3\chi_4\rangle\}$.

Solution 2.25

Using (2.217), the conclusion can be obtained at once.

Exercise 2.26

Show using second quantization that $\langle\chi_i|\chi_j\rangle = \delta_{ij}$.

Solution 2.26

The proof is direct.

$$\langle\chi_i|\chi_j\rangle = \langle|a_i a_j^\dagger|\rangle = \langle|\delta_{ij} - a_j^\dagger a_i|\rangle = \delta_{ij}\langle| \rangle - \langle|a_j^\dagger a_i|\rangle = \delta_{ij}.$$

Exercise 2.27

Given a state

$$|K\rangle = |\chi_1 \chi_2 \cdots \chi_N\rangle = a_1^\dagger a_2^\dagger \cdots a_N^\dagger | \rangle,$$

show that $\langle K | a_i^\dagger a_j | K \rangle = 1$ if $i = j$ and $i \in \{1, 2, \dots, N\}$, but is zero otherwise.

Solution 2.27

The proof is direct, too.

$$\langle K | a_i^\dagger a_j | K \rangle = \langle K | \delta_{ij} - a_j a_i^\dagger | K \rangle = \delta_{ij} \langle K | K \rangle - \langle K | a_j a_i^\dagger | K \rangle = \delta_{ij} - \langle K | a_j a_i^\dagger | K \rangle.$$

It is evident that $\langle K | a_i^\dagger a_j | K \rangle$ will be one if and only if $i = j$ and $i \in \{1, 2, \dots, N\}$. Otherwise, it will be zero.

Exercise 2.28

Let $|\Psi_0\rangle = |\chi_1 \cdots \chi_a \chi_b \cdots \chi_N\rangle$ be the Hartree-Fock ground state wave function. Show that

- $a_r |\Psi_0\rangle = 0 = \langle \Psi_0 | a_r^\dagger$.
- $a_a^\dagger |\Psi_0\rangle = 0 = \langle \Psi_0 | a_a$.
- $|\Psi_a^r\rangle = a_r^\dagger a_a |\Psi_0\rangle$.
- $\langle \Psi_a^r | = \langle \Psi_0 | a_a^\dagger a_r$.
- $|\Psi_{ab}^{rs}\rangle = a_s^\dagger a_b a_r^\dagger a_a |\Psi_0\rangle = a_r^\dagger a_s^\dagger a_b a_a |\Psi_0\rangle$.
- $\langle \Psi_{ab}^{rs} | = \langle \Psi_0 | a_a^\dagger a_r a_b^\dagger a_s = \langle \Psi_0 | a_a^\dagger a_b^\dagger a_s a_r$.

Solution 2.28

Assume that a, b are labels of occupied spin orbitals while r, s are labels of unoccupied spin orbitals.

- It is evident because there exists no electron occupying spin orbital r .
- It is also evident because there is an electron occupying spin orbital a .
- The verification is direct.

$$\begin{aligned} |\Psi_a^r\rangle &= |\chi_1 \cdots \chi_r \chi_b \cdots \chi_N\rangle = -|\chi_r \cdots \chi_1 \chi_b \cdots \chi_N\rangle = -a_r^\dagger |\cdots \chi_1 \chi_b \cdots \chi_N\rangle \\ &= -a_r^\dagger a_a |\chi_a \cdots \chi_1 \chi_b \cdots \chi_N\rangle = a_r^\dagger a_a |\chi_1 \cdots \chi_a \chi_b \cdots \chi_N\rangle = a_r^\dagger a_a |\Psi_0\rangle. \end{aligned}$$

- The verification is direct, too.

$$\begin{aligned} \langle \Psi_a^r | &= \langle \chi_1 \cdots \chi_r \chi_b \cdots \chi_N | = -\langle \chi_1 \cdots \chi_N \chi_b \cdots \chi_r | = -\langle \chi_1 \cdots \chi_N \chi_b \cdots | a_r \\ &= -\langle \chi_1 \cdots \chi_N \chi_b \cdots \chi_a | a_a^\dagger a_r = \langle \chi_1 \cdots \chi_a \chi_b \cdots \chi_N | a_a^\dagger a_r = \langle \Psi_0 | a_a^\dagger a_r. \end{aligned}$$

- The verification is direct, too.

$$\begin{aligned} |\Psi_{ab}^{rs}\rangle &= |\chi_1 \chi_2 \cdots \chi_r \chi_s \cdots \chi_N\rangle = (-1)^2 |\chi_r \chi_s \cdots \chi_1 \chi_2 \cdots \chi_N\rangle = a_r^\dagger |\chi_s \cdots \chi_1 \chi_2 \cdots \chi_N\rangle \\ &= a_r^\dagger a_s^\dagger |\cdots \chi_1 \chi_2 \cdots \chi_N\rangle = a_r^\dagger a_s^\dagger a_b |\chi_b \cdots \chi_1 \chi_2 \cdots \chi_N\rangle = -a_r^\dagger a_s^\dagger a_b |\chi_2 \cdots \chi_1 \chi_b \cdots \chi_N\rangle \\ &= -a_r^\dagger a_s^\dagger a_b a_a |\chi_a \chi_2 \cdots \chi_1 \chi_b \cdots \chi_N\rangle = a_r^\dagger a_s^\dagger a_b a_a |\chi_1 \chi_2 \cdots \chi_a \chi_b \cdots \chi_N\rangle = a_r^\dagger a_s^\dagger a_b a_a |\Psi_0\rangle. \end{aligned}$$

Note that with $\delta_{ar} = \delta_{br} = \delta_{as} = \delta_{bs} = 0$,

$$a_r^\dagger a_s^\dagger a_b = -a_s^\dagger a_r^\dagger a_b = -a_s^\dagger (\delta_{br} - a_b a_r^\dagger) = a_s^\dagger a_b a_r^\dagger.$$

We obtain that

$$|\Psi_{ab}^{rs}\rangle = a_r^\dagger a_s^\dagger a_b a_a |\Psi_0\rangle = a_s^\dagger a_b a_r^\dagger a_a |\Psi_0\rangle.$$

- The verification is direct, too.

$$\langle \Psi_{ab}^{rs} | = \langle \chi_1 \cdots \chi_r \chi_s \cdots \chi_{N-1} \chi_N | = (-1)^2 \langle \chi_1 \cdots \chi_N \chi_{N-1} \cdots \chi_s \chi_r | = \langle \chi_1 \cdots \chi_N \chi_{N-1} \cdots \chi_s | a_r$$

$$\begin{aligned}
&= \langle \chi_1 \cdots \chi_N \chi_{N-1} \cdots | a_s a_r = \langle \chi_1 \cdots \chi_N \chi_{N-1} \cdots \chi_b | a_b^\dagger a_s a_r = -\langle \chi_1 \cdots \chi_N \chi_b \cdots \chi_{N-1} | a_b^\dagger a_s a_r \\
&= -\langle \chi_1 \cdots \chi_N \chi_b \cdots \chi_{N-1} \chi_a | a_a^\dagger a_b^\dagger a_s a_r = \langle \chi_1 \cdots \chi_a \chi_b \cdots \chi_{N-1} \chi_N | a_a^\dagger a_b^\dagger a_s a_r = \langle \Psi_0 | a_a^\dagger a_b^\dagger a_s a_r.
\end{aligned}$$

In the same way, we can find that

$$a_b^\dagger a_s a_r = -a_b^\dagger a_r a_s = -(\delta_{br} - a_r a_b^\dagger) a_s = a_r a_b^\dagger a_s.$$

Thus,

$$\langle \Psi_{ab}^{rs} | = \langle \Psi_0 | a_a^\dagger a_b^\dagger a_s a_r = \langle \Psi_0 | a_a^\dagger a_r a_b^\dagger a_s.$$

2.4.2 Second-Quantized Operators and Their Matrix Elements

Exercise 2.29

Let $|\Psi_0\rangle = |\chi_1 \chi_2\rangle = a_1^\dagger a_2^\dagger | \rangle$ be the Hartree-Fock wave function for minimal basis H_2 . Show using second quantization that

$$\langle \Psi_0 | \mathcal{O}_1 | \Psi_0 \rangle = \sum_{ij} \langle i | h | j \rangle \langle | a_2 a_1 a_i^\dagger a_j a_1^\dagger a_2^\dagger | \rangle = \langle 1 | h | 1 \rangle + \langle 2 | h | 2 \rangle.$$

Solution 2.29

Using (2.231) and the conclusion of Exercise 2.28, we find that

$$\begin{aligned}
\langle \Psi_0 | \mathcal{O}_1 | \Psi_0 \rangle &= \langle \chi_1 \chi_2 | \sum_{ij} \langle i | h | j \rangle a_i^\dagger a_j | \chi_1 \chi_2 \rangle = \sum_{ij} \langle i | h | j \rangle \langle | a_2 a_1 a_i^\dagger a_j a_1^\dagger a_2^\dagger | \rangle \\
&= \sum_{ij} \langle i | h | j \rangle \langle | a_2 (\delta_{1i} - a_i^\dagger a_1) (\delta_{j1} - a_1^\dagger a_j) a_2^\dagger | \rangle \\
&= \sum_{ij} \langle i | h | j \rangle \left[\delta_{1i} \delta_{1j} \langle | a_2 a_2^\dagger | \rangle - \delta_{1i} \langle | a_2 a_1^\dagger a_j a_2^\dagger | \rangle - \delta_{j1} \langle | a_2 a_i^\dagger a_1 a_2^\dagger | \rangle + \langle | a_2 a_i^\dagger a_1 a_1^\dagger a_j a_2^\dagger | \rangle \right] \\
&= \langle 1 | h | 1 \rangle \langle | a_2 a_2^\dagger | \rangle - \sum_j \langle 1 | h | j \rangle \langle | a_2 a_1^\dagger a_j a_2^\dagger | \rangle - \sum_i \langle i | h | 1 \rangle \langle | a_2 a_i^\dagger a_1 a_2^\dagger | \rangle + \sum_{ij} \langle i | h | j \rangle \langle | a_2 a_i^\dagger a_1 a_1^\dagger a_j a_2^\dagger | \rangle \\
&= \langle 1 | h | 1 \rangle \left(\langle | \delta_{22} - a_2^\dagger a_2 | \rangle \right) - \sum_j \langle 1 | h | j \rangle \langle | (\delta_{12} - a_1^\dagger a_2) a_j a_2^\dagger | \rangle - \sum_i \langle i | h | 1 \rangle \langle | a_2 a_i^\dagger (\delta_{12} - a_2^\dagger a_1) | \rangle \\
&\quad + \sum_{ij} \langle i | h | j \rangle \langle | (\delta_{2i} - a_i^\dagger a_2) a_1 a_1^\dagger (\delta_{2j} - a_2^\dagger a_j) | \rangle \\
&= \langle 1 | h | 1 \rangle (1 - 0) + \sum_j \langle 1 | h | j \rangle \langle | a_1^\dagger a_2 a_j a_2^\dagger | \rangle + \sum_i \langle i | h | 1 \rangle \langle | a_2 a_i^\dagger a_2^\dagger a_1 | \rangle \\
&\quad + \sum_{ij} \langle i | h | j \rangle \left[\delta_{2i} \delta_{2j} \langle | a_1 a_1^\dagger | \rangle - \delta_{2i} \langle | a_1 a_1^\dagger a_2^\dagger a_j | \rangle - \delta_{2j} \langle | a_i^\dagger a_2 a_1 a_1^\dagger | \rangle + \langle | a_i^\dagger a_2 a_1 a_1^\dagger a_2^\dagger a_j | \rangle \right] \\
&= \langle 1 | h | 1 \rangle + 0 + 0 + \langle 2 | h | 2 \rangle (\delta_{11} - \langle | a_1 a_1^\dagger | \rangle) - 0 - 0 + 0 = \langle 1 | h | 1 \rangle + \langle 2 | h | 2 \rangle (1 - 0) = \langle 1 | h | 1 \rangle + \langle 2 | h | 2 \rangle.
\end{aligned}$$

Exercise 2.30

Show that

$$\langle \Psi_a^r | \mathcal{O}_1 | \Psi_0 \rangle = \sum_{ij} \langle i | h | j \rangle \langle \Psi_0 | a_a^\dagger a_r a_i^\dagger a_j | \Psi_0 \rangle = \langle r | h | a \rangle$$

by moving a_a^\dagger and a_r to the right.

Solution 2.30

Using (2.231) and the conclusion of Exercise 2.28, we find that

$$\langle \Psi_a^r | \mathcal{O}_1 | \Psi_0 \rangle = \langle \Psi_0 | a_a^\dagger a_r \sum_{ij} \langle i | h | j \rangle a_i^\dagger a_j | \Psi_0 \rangle = \sum_{ij} \langle i | h | j \rangle \langle \Psi_0 | a_a^\dagger a_r a_i^\dagger a_j | \Psi_0 \rangle$$

$$\begin{aligned}
&= \sum_{ij} \langle i|h|j \rangle \langle \Psi_0 | a_a^\dagger (\delta_{ri} - a_i^\dagger a_r) a_j | \Psi_0 \rangle = \sum_j \langle r|h|j \rangle \langle \Psi_0 | a_a^\dagger a_j | \Psi_0 \rangle - \sum_{ij} \langle i|h|j \rangle \langle \Psi_0 | a_a^\dagger a_i^\dagger a_r a_j | \Psi_0 \rangle \\
&= \sum_j \langle r|h|j \rangle \langle \Psi_0 | (\delta_{aj} - a_j a_a^\dagger) | \Psi_0 \rangle + \sum_{ij} \langle i|h|j \rangle \langle \Psi_0 | a_a^\dagger a_i^\dagger a_j a_r | \Psi_0 \rangle \\
&= \sum_j \langle r|h|j \rangle \delta_{aj} - \sum_j \langle r|h|j \rangle \langle \Psi_0 | a_j a_a^\dagger | \Psi_0 \rangle + 0 = \langle r|h|a \rangle + 0 = \langle r|h|a \rangle.
\end{aligned}$$

Exercise 2.31

Show that

$$\langle \Psi_a^r | \mathcal{O}_2 | \Psi_0 \rangle = \sum_b^N \langle rb | ab \rangle.$$

Hint: first show that

$$\begin{aligned}
\langle \Psi_0 | a_a^\dagger a_r a_i^\dagger a_j^\dagger a_l a_k | \Psi_0 \rangle &= \delta_{rj} \delta_{al} \langle \Psi_0 | a_i^\dagger a_k | \Psi_0 \rangle - \delta_{rj} \delta_{ak} \langle \Psi_0 | a_i^\dagger a_l | \Psi_0 \rangle \\
&\quad + \delta_{ri} \delta_{ak} \langle \Psi_0 | a_j^\dagger a_l | \Psi_0 \rangle - \delta_{ri} \delta_{al} \langle \Psi_0 | a_j^\dagger a_k | \Psi_0 \rangle
\end{aligned}$$

then refer to Exercise 2.27.

Solution 2.31

Similar to Exercise 2.30, we find that

$$\begin{aligned}
&\langle \Psi_0 | a_a^\dagger a_r a_i^\dagger a_j^\dagger a_l a_k | \Psi_0 \rangle \\
&= \langle \Psi_0 | a_a^\dagger (\delta_{ri} - a_i^\dagger a_r) a_j^\dagger a_l a_k | \Psi_0 \rangle = \delta_{ri} \langle \Psi_0 | a_a^\dagger a_j^\dagger a_l a_k | \Psi_0 \rangle - \langle \Psi_0 | a_a^\dagger a_i^\dagger a_r a_j^\dagger a_l a_k | \Psi_0 \rangle \\
&= -\delta_{ri} \langle \Psi_0 | a_j^\dagger a_a^\dagger a_l a_k | \Psi_0 \rangle - \langle \Psi_0 | a_a^\dagger a_i^\dagger (\delta_{rj} - a_j^\dagger a_r) a_l a_k | \Psi_0 \rangle \\
&= -\delta_{ri} \langle \Psi_0 | a_j^\dagger (\delta_{al} - a_l a_a^\dagger) a_k | \Psi_0 \rangle - \delta_{rj} \langle \Psi_0 | a_a^\dagger a_i^\dagger a_l a_k | \Psi_0 \rangle + \langle \Psi_0 | a_a^\dagger a_i^\dagger a_j^\dagger a_r a_l a_k | \Psi_0 \rangle \\
&= -\delta_{ri} \delta_{al} \langle \Psi_0 | a_j^\dagger a_k | \Psi_0 \rangle + \delta_{ri} \langle \Psi_0 | a_j^\dagger a_l a_a^\dagger a_k | \Psi_0 \rangle + \delta_{rj} \langle \Psi_0 | a_i^\dagger a_a^\dagger a_l a_k | \Psi_0 \rangle - \langle \Psi_0 | a_a^\dagger a_i^\dagger a_j^\dagger a_l a_r a_k | \Psi_0 \rangle \\
&= -\delta_{ri} \delta_{al} \langle \Psi_0 | a_j^\dagger a_k | \Psi_0 \rangle + \delta_{ri} \langle \Psi_0 | a_j^\dagger a_l (\delta_{ak} - a_k a_a^\dagger) | \Psi_0 \rangle \\
&\quad + \delta_{rj} \langle \Psi_0 | a_i^\dagger (\delta_{al} - a_l a_a^\dagger) a_k | \Psi_0 \rangle + \langle \Psi_0 | a_a^\dagger a_i^\dagger a_j^\dagger a_l a_r a_k | \Psi_0 \rangle \\
&= -\delta_{ri} \delta_{al} \langle \Psi_0 | a_j^\dagger a_k | \Psi_0 \rangle + \delta_{ri} \delta_{ak} \langle \Psi_0 | a_j^\dagger a_l | \Psi_0 \rangle - \delta_{ri} \langle \Psi_0 | a_j^\dagger a_l a_k a_a^\dagger | \Psi_0 \rangle \\
&\quad + \delta_{rj} \delta_{al} \langle \Psi_0 | a_i^\dagger a_k | \Psi_0 \rangle - \delta_{rj} \langle \Psi_0 | a_i^\dagger a_l a_a^\dagger a_k | \Psi_0 \rangle + 0 \\
&= -\delta_{ri} \delta_{al} \langle \Psi_0 | a_j^\dagger a_k | \Psi_0 \rangle + \delta_{ri} \delta_{ak} \langle \Psi_0 | a_j^\dagger a_l | \Psi_0 \rangle - 0 + \delta_{rj} \delta_{al} \langle \Psi_0 | a_i^\dagger a_k | \Psi_0 \rangle - \delta_{rj} \langle \Psi_0 | a_i^\dagger a_l (\delta_{ak} - a_k a_a^\dagger) | \Psi_0 \rangle \\
&= -\delta_{ri} \delta_{al} \langle \Psi_0 | a_j^\dagger a_k | \Psi_0 \rangle + \delta_{ri} \delta_{ak} \langle \Psi_0 | a_j^\dagger a_l | \Psi_0 \rangle \\
&\quad + \delta_{rj} \delta_{al} \langle \Psi_0 | a_i^\dagger a_k | \Psi_0 \rangle - \delta_{rj} \delta_{ak} \langle \Psi_0 | a_i^\dagger a_l | \Psi_0 \rangle + \delta_{rj} \langle \Psi_0 | a_i^\dagger a_l a_k a_a^\dagger | \Psi_0 \rangle \\
&= \delta_{rj} \delta_{al} \langle \Psi_0 | a_i^\dagger a_k | \Psi_0 \rangle - \delta_{rj} \delta_{ak} \langle \Psi_0 | a_i^\dagger a_l | \Psi_0 \rangle + \delta_{ri} \delta_{ak} \langle \Psi_0 | a_j^\dagger a_l | \Psi_0 \rangle - \delta_{ri} \delta_{al} \langle \Psi_0 | a_j^\dagger a_k | \Psi_0 \rangle.
\end{aligned}$$

Note that $\langle \Psi_0 | a_i^\dagger a_j | \Psi_0 \rangle$ equals 1 if and only if i and j should be labels of occupied spin orbitals like a and b . Hence, we find that

$$\begin{aligned}
\langle \Psi_a^r | \mathcal{O}_2 | \Psi_0 \rangle &= \langle \Psi_0 | a_a^\dagger a_r \frac{1}{2} \sum_{ijkl} \langle ij|kl \rangle a_i^\dagger a_j^\dagger a_l a_k | \Psi_0 \rangle = \frac{1}{2} \sum_{ijkl} \langle ij|kl \rangle \langle \Psi_0 | a_a^\dagger a_r a_i^\dagger a_j^\dagger a_l a_k | \Psi_0 \rangle \\
&= \frac{1}{2} \sum_{ijkl} \langle ij|kl \rangle (\delta_{rj} \delta_{al} \langle \Psi_0 | a_i^\dagger a_k | \Psi_0 \rangle - \delta_{rj} \delta_{ak} \langle \Psi_0 | a_i^\dagger a_l | \Psi_0 \rangle + \delta_{ri} \delta_{ak} \langle \Psi_0 | a_j^\dagger a_l | \Psi_0 \rangle - \delta_{ri} \delta_{al} \langle \Psi_0 | a_j^\dagger a_k | \Psi_0 \rangle) \\
&= \frac{1}{2} \sum_{ik} \langle ir|ka \rangle \langle \Psi_0 | a_i^\dagger a_k | \Psi_0 \rangle - \frac{1}{2} \sum_{il} \langle ir|al \rangle \langle \Psi_0 | a_i^\dagger a_l | \Psi_0 \rangle \\
&\quad + \frac{1}{2} \sum_{jl} \langle rj|al \rangle \langle \Psi_0 | a_j^\dagger a_l | \Psi_0 \rangle - \frac{1}{2} \sum_{jk} \langle rj|ka \rangle \langle \Psi_0 | a_j^\dagger a_k | \Psi_0 \rangle \\
&= \frac{1}{2} \sum_{ij} \langle ir|ja \rangle \langle \Psi_0 | a_i^\dagger a_j | \Psi_0 \rangle - \frac{1}{2} \sum_{ij} \langle ir|aj \rangle \langle \Psi_0 | a_i^\dagger a_j | \Psi_0 \rangle
\end{aligned}$$

$$\begin{aligned}
& + \frac{1}{2} \sum_{ij} \langle ri|aj \rangle \langle \Psi_0 | a_i^\dagger a_j | \Psi_0 \rangle - \frac{1}{2} \sum_{ij} \langle ri|ja \rangle \langle \Psi_0 | a_i^\dagger a_j | \Psi_0 \rangle \\
& = \sum_{ij} \langle ir|ja \rangle \langle \Psi_0 | a_i^\dagger a_j | \Psi_0 \rangle - \langle ir|aj \rangle \langle \Psi_0 | a_i^\dagger a_j | \Psi_0 \rangle = \sum_{ij} \langle ir||ja \rangle \langle \Psi_0 | a_i^\dagger a_j | \Psi_0 \rangle \\
& = \sum_{ij} \langle ri||aj \rangle \langle \Psi_0 | a_i^\dagger a_j | \Psi_0 \rangle = \sum_b^N \langle rb||ab \rangle.
\end{aligned}$$

2.5 Spin-Adapted Configurations

2.5.1 Spin Operators

Exercise 2.32

a) Derive (2.247) from (2.245); b) Derive (2.248).

Solution 2.32

a) Using (2.246a) and (2.245c,d), we find that

$$\begin{aligned}
s_+|\alpha\rangle &= (s_x + is_y)|\alpha\rangle = s_x|\alpha\rangle + is_y|\alpha\rangle = \frac{1}{2}|\beta\rangle + i \times \frac{i}{2}|\beta\rangle = \left(\frac{1}{2} + i \times \frac{i}{2}\right)|\beta\rangle = 0, \\
s_+|\beta\rangle &= (s_x + is_y)|\beta\rangle = s_x|\beta\rangle + is_y|\beta\rangle = \frac{1}{2}|\alpha\rangle + i \times \frac{-i}{2}|\alpha\rangle = \left(\frac{1}{2} + i \times \frac{-i}{2}\right)|\alpha\rangle = |\alpha\rangle, \\
s_-|\alpha\rangle &= (s_x - is_y)|\alpha\rangle = s_x|\alpha\rangle - is_y|\alpha\rangle = \frac{1}{2}|\beta\rangle - i \times \frac{i}{2}|\beta\rangle = \left(\frac{1}{2} - i \times \frac{i}{2}\right)|\beta\rangle = |\beta\rangle, \\
s_-|\beta\rangle &= (s_x - is_y)|\beta\rangle = s_x|\beta\rangle - is_y|\beta\rangle = \frac{1}{2}|\alpha\rangle - i \times \frac{-i}{2}|\alpha\rangle = \left(\frac{1}{2} - i \times \frac{-i}{2}\right)|\alpha\rangle = 0.
\end{aligned}$$

b) From (2.246a,b) and the conclusion of Exercise 1.4(f), we know that

$$\begin{pmatrix} s_+ \\ s_- \end{pmatrix} = \begin{pmatrix} 1 & i \\ 1 & -i \end{pmatrix} \begin{pmatrix} s_x \\ s_y \end{pmatrix} \Leftrightarrow \begin{pmatrix} s_x \\ s_y \end{pmatrix} = \begin{pmatrix} 1 & i \\ 1 & -i \end{pmatrix}^{-1} \begin{pmatrix} s_+ \\ s_- \end{pmatrix} = \frac{1}{-2i} \begin{pmatrix} -i & -i \\ -1 & 1 \end{pmatrix} \begin{pmatrix} s_+ \\ s_- \end{pmatrix}.$$

In other words, we know that

$$s_x = \frac{-is_+ - is_-}{-2i} = \frac{s_+ + s_-}{2}, \quad s_y = \frac{-s_+ + s_-}{-2i} = \frac{s_+ - s_-}{2i}.$$

Besides, with (2.242), it is evident that

$$[s_+, s_-] = [s_x + is_y, s_x - is_y] = [s_x, s_x] - i[s_x, s_y] + i[s_y, s_x] + [s_y, s_y] = -2i[s_x, s_y] = -2i(is_z) = 2s_z.$$

From (2.241), we find that

$$\begin{aligned}
s^2 &= s_x^2 + s_y^2 + s_z^2 = \left(\frac{s_+ + s_-}{2}\right)^2 + \left(\frac{s_+ - s_-}{2i}\right)^2 + s_z^2 \\
&= \frac{1}{4}(s_+^2 + s_+s_- + s_-s_+ + s_-^2) - \frac{1}{4}(s_+^2 - s_+s_- - s_-s_+ + s_-^2) + s_z^2 \\
&= \frac{1}{2}(s_+s_- + s_-s_+) + s_z^2 = s_+s_- + \frac{1}{2}(s_-s_+ - s_+s_-) + s_z^2 = s_-s_+ + \frac{1}{2}(s_+s_- - s_-s_+) + s_z^2 \\
&= s_+s_- + \frac{1}{2}[s_-, s_+] + s_z^2 = s_+s_- - \frac{1}{2}[s_+, s_-] + s_z^2 = s_+s_- - \frac{1}{2}(2s_z) + s_z^2 = s_+s_- - s_z + s_z^2 \\
&= s_-s_+ + \frac{1}{2}[s_+, s_-] + s_z^2 = s_-s_+ + \frac{1}{2}(2s_z) + s_z^2 = s_-s_+ + s_z + s_z^2.
\end{aligned}$$

Exercise 2.33

Find the 2×2 matrix representations of s^2 , s_z , s_+ , and s_- in the basis $|\alpha\rangle$, $|\beta\rangle$. Verify the identities analogous to (2.248a,b) for these matrix representations.

Solution 2.33

From (2.245a,b,c,d), at once we get that

$$\begin{aligned} s^2(|\alpha\rangle, |\beta\rangle) &= (|\alpha\rangle, |\beta\rangle) \begin{pmatrix} \frac{3}{4} & 0 \\ 0 & \frac{3}{4} \end{pmatrix}, \\ s_z(|\alpha\rangle, |\beta\rangle) &= (|\alpha\rangle, |\beta\rangle) \begin{pmatrix} \frac{1}{2} & 0 \\ 0 & -\frac{1}{2} \end{pmatrix}, \\ s_x(|\alpha\rangle, |\beta\rangle) &= (|\alpha\rangle, |\beta\rangle) \begin{pmatrix} 0 & \frac{1}{2} \\ \frac{1}{2} & 0 \end{pmatrix}, \\ s_y(|\alpha\rangle, |\beta\rangle) &= (|\alpha\rangle, |\beta\rangle) \begin{pmatrix} 0 & -\frac{i}{2} \\ \frac{i}{2} & 0 \end{pmatrix}. \end{aligned}$$

Thus, we find that

$$\begin{aligned} s_+ (|\alpha\rangle, |\beta\rangle) &= (s_x + is_y)(|\alpha\rangle, |\beta\rangle) = s_x(|\alpha\rangle, |\beta\rangle) + is_y(|\alpha\rangle, |\beta\rangle) \\ &= (|\alpha\rangle, |\beta\rangle) \begin{pmatrix} 0 & \frac{1}{2} \\ \frac{1}{2} & 0 \end{pmatrix} + (|\alpha\rangle, |\beta\rangle) \left[i \begin{pmatrix} 0 & -\frac{i}{2} \\ \frac{i}{2} & 0 \end{pmatrix} \right] = (|\alpha\rangle, |\beta\rangle) \begin{pmatrix} 0 & 1 \\ 0 & 0 \end{pmatrix}. \\ s_- (|\alpha\rangle, |\beta\rangle) &= (s_x - is_y)(|\alpha\rangle, |\beta\rangle) = s_x(|\alpha\rangle, |\beta\rangle) - is_y(|\alpha\rangle, |\beta\rangle) \\ &= (|\alpha\rangle, |\beta\rangle) \begin{pmatrix} 0 & \frac{1}{2} \\ \frac{1}{2} & 0 \end{pmatrix} + (|\alpha\rangle, |\beta\rangle) \left[-i \begin{pmatrix} 0 & -\frac{i}{2} \\ \frac{i}{2} & 0 \end{pmatrix} \right] = (|\alpha\rangle, |\beta\rangle) \begin{pmatrix} 0 & 0 \\ 1 & 0 \end{pmatrix}. \end{aligned}$$

Thus, in the basis $|\alpha\rangle$, $|\beta\rangle$, the 2×2 matrix representations of s^2 , s_z , s_+ and s_- are

$$s^2 = \begin{pmatrix} \frac{3}{4} & 0 \\ 0 & \frac{3}{4} \end{pmatrix}, \quad s_z = \begin{pmatrix} \frac{1}{2} & 0 \\ 0 & -\frac{1}{2} \end{pmatrix}, \quad s_+ = \begin{pmatrix} 0 & 1 \\ 0 & 0 \end{pmatrix}, \quad s_- = \begin{pmatrix} 0 & 0 \\ 1 & 0 \end{pmatrix}. \quad (2.33-1)$$

Readers can verify that these matrices represent identities similar to (2.248a,b) by themselves.

Exercise 2.34

Using the commutation relations (2.242), show that $[s^2, s_z] = 0$.

Solution 2.34

Before verification, note that for any operators (matrices) A , B , C ,

$$\begin{aligned} [AB, C] &= ABC - CAB = ABC - ACB + ACB - CAB \\ &= A(BC - CB) + (AC - CA)B = A[B, C] + [A, C]B. \end{aligned}$$

From (2.241) and (2.242), we find that

$$\begin{aligned} [s^2, s_z] &= [s_x^2 + s_y^2 + s_z^2, s_z] = [s_x^2, s_z] + [s_y^2, s_z] + [s_z^2, s_z] \\ &= s_x[s_x, s_z] + [s_x, s_z]s_x + s_y[s_y, s_z] + [s_y, s_z]s_y + s_z[s_z, s_z] + [s_z, s_z]s_z \\ &= -s_x[s_z, s_x] - [s_z, s_x]s_x + s_y[s_y, s_z] + [s_y, s_z]s_y + 0 + 0 \\ &= -s_x(is_y) - (is_y)s_x + s_y(is_x) + (is_x)s_y = (-i + i)s_xs_y + (-i + i)s_ys_x = 0. \end{aligned}$$

Exercise 2.35

Consider an operator \mathcal{A} that commutes with the Hamiltonian. Suppose $|\Phi\rangle$ is an eigenfunction of \mathcal{H} with eigenvalue E . Show that $\mathcal{A}|\Phi\rangle$ is also an eigenfunction of \mathcal{H} with eigenvalue E . Thus if $|\Phi\rangle$ is (energetically) nondegenerate, then $\mathcal{A}|\Phi\rangle$ is at most a constant multiple of $|\Phi\rangle$ (i.e., $\mathcal{A}|\Phi\rangle = a|\Phi\rangle$) and hence $|\Phi\rangle$ is an eigenfunction of \mathcal{A} . In case of degeneracies, we can always construct appropriate linear combinations of the degenerate eigenfunctions of \mathcal{H} that are also eigenfunctions of \mathcal{A} .

Solution 2.35

The proof of that if an operator \mathcal{A} that commutes with the Hamiltonian and $|\Phi\rangle$ is an eigenfunction of \mathcal{H} with eigenvalue E , $\mathcal{A}|\Phi\rangle$ will be also an eigenfunction of \mathcal{H} with eigenvalue E , is easy.

$$\mathcal{H}\mathcal{A}|\Phi\rangle = \mathcal{A}\mathcal{H}|\Phi\rangle = \mathcal{A}E|\Phi\rangle = E\mathcal{A}|\Phi\rangle.$$

Exercise 2.36

Given two nondegenerate eigenfunctions of a hermitian operator \mathcal{A} that commutes with \mathcal{H} , i.e., $\mathcal{A}|\Psi_1\rangle = a_1|\Psi_1\rangle$, $\mathcal{A}|\Psi_2\rangle = a_2|\Psi_2\rangle$, $a_1 \neq a_2$, show that $\langle\Psi_1|\mathcal{H}|\Psi_2\rangle = 0$. Thus the matrix element of the Hamiltonian between, say, singlet and triplet spin-adapted configurations is zero.

Solution 2.36

As \mathcal{A} is a hermitian operator, a_1 and a_2 are real and

$$\langle\Psi_2|\mathcal{A} = a_2\langle\Psi_2|.$$

Hence,

$$a_1\langle\Psi_2|\mathcal{H}|\Psi_1\rangle = \langle\Psi_2|\mathcal{H}a_1|\Psi_1\rangle = \langle\Psi_2|\mathcal{H}\mathcal{A}|\Psi_1\rangle = \langle\Psi_2|\mathcal{A}\mathcal{H}|\Psi_1\rangle = a_2\langle\Psi_2|\mathcal{H}|\Psi_1\rangle.$$

In other words,

$$(a_1 - a_2)\langle\Psi_2|\mathcal{H}|\Psi_1\rangle = 0.$$

Due to $a_1 \neq a_2$, we find that

$$\langle\Psi_1|\mathcal{H}|\Psi_2\rangle = 0.$$

Exercise 2.37

Prove Eq.(2.254). *Hint:* Use expansion (2.115) for a Slater determinant and note that \mathcal{S}_z , since it is invariant to any permutation of the electron labels, commutes with \mathcal{P}_n .

Solution 2.37

Firstly, note that

$$s_z(i)\chi_j(\mathbf{x}_i) = \frac{\delta_{j\alpha} - \delta_{j\beta}}{2}\chi_j(\mathbf{x}_i).$$

The derivation is exactly the same as that of Exercise 2.15 if $h(i)$ is replaced by $s_z(i)$. And thus, $\sum_{i=1}^N \varepsilon_i$ should be replaced by

$$\sum_{i=1}^N \frac{\delta_{j\alpha} - \delta_{j\beta}}{2} = \frac{N^\alpha - N^\beta}{2}.$$

2.5.2 Restricted Determinants and Spin-Adapted Configurations**Exercise 2.38**

Prove Eq.(2.256). *Hints:* 1) $\mathcal{S}^2 = \mathcal{S}_-\mathcal{S}_+ + \mathcal{S}_z + \mathcal{S}_z^2$, 2) as a result of Eq.(2.254) it is sufficient to show $\mathcal{S}_+|\psi_i\bar{\psi}_i\cdots\rangle = 0$, 3) use expansion (2.115) for the determinant, and note the \mathcal{S}_+ commutes with the permutation operator, 4) $s_+\psi_\alpha = 0$, 5) finally, $s_+\psi_\beta = \psi_\alpha$, but the determinant vanishes because it has two identical columns.

Solution 2.38

Firstly, we verify (2.251). Similar to Exercise 2.32, we find that

$$\mathcal{S}^2 = \sum_{i=1}^N \sum_{j=1}^N \vec{s}(i) \cdot \vec{s}(j) = \sum_{i=1}^N \sum_{j=1}^N s_x(i)s_x(j) + s_y(i)s_y(j) + s_z(i)s_z(j)$$

$$\begin{aligned}
&= \sum_{i=1}^N \sum_{j=1}^N \left(\frac{s_+(i) + s_-(i)}{2} \right) \left(\frac{s_+(j) + s_-(j)}{2} \right) + \left(\frac{s_+(i) - s_-(i)}{2i} \right) \left(\frac{s_+(j) - s_-(j)}{2j} \right) + s_z(i)s_z(j) \\
&= \sum_{i=1}^N \sum_{j=1}^N \frac{1}{4} [s_+(i)s_+(j) + s_+(i)s_-(j) + s_-(i)s_+(j) + s_-(i)s_-(j)] \\
&\quad - \frac{1}{4} [s_+(i)s_+(j) - s_+(i)s_-(j) - s_-(i)s_+(j) + s_-(i)s_-(j)] + s_z(i)s_z(j) \\
&= \sum_{i=1}^N \sum_{j=1}^N \frac{1}{2} [s_+(i)s_-(j) + s_-(i)s_+(j)] + s_z(i)s_z(j) \\
&= \sum_{i=1}^N \sum_{j=1}^N s_+(i)s_-(j) + \frac{1}{2} [s_-(i)s_+(j) - s_+(i)s_-(j)] + s_z(i)s_z(j) \\
&= \sum_{i=1}^N \sum_{j=1}^N s_-(i)s_+(j) + \frac{1}{2} [s_+(i)s_-(j) - s_-(i)s_+(j)] + s_z(i)s_z(j).
\end{aligned}$$

The first result can be further simplified to

$$\begin{aligned}
&\sum_{i=1}^N \sum_{j=1}^N s_+(i)s_-(j) + \frac{1}{2} [s_-(i)s_+(j) - s_+(i)s_-(j)] + s_z(i)s_z(j) \\
&= \sum_{i=1}^N s_+(i) \sum_{j=1}^N s_-(j) + \frac{1}{2} \sum_{i=1}^N \sum_{j=1}^N s_-(i)s_+(j) - \frac{1}{2} \sum_{i=1}^N \sum_{j=1}^N s_+(i)s_-(j) + \sum_{i=1}^N \sum_{j=1}^N s_z(i)s_z(j) \\
&= \sum_{i=1}^N s_+(i) \sum_{j=1}^N s_-(j) + \frac{1}{2} \sum_{i=1}^N \sum_{j=1}^N [s_-(i), s_+(j)] + s_+(j)s_-(i) - \frac{1}{2} \sum_{i=1}^N \sum_{j=1}^N s_+(j)s_-(i) + \sum_{i=1}^N \sum_{j=1}^N s_z(i)s_z(j) \\
&= \sum_{i=1}^N s_+(i) \sum_{j=1}^N s_-(j) + \frac{1}{2} \sum_{i=1}^N [s_-(i), s_+(i)] + \frac{1}{2} \sum_{i=1}^N \sum_{\substack{j=1 \\ j \neq i}}^N [s_-(i), s_+(j)] \\
&\quad + \frac{1}{2} \sum_{i=1}^N \sum_{j=1}^N s_+(j)s_-(i) - \frac{1}{2} \sum_{i=1}^N \sum_{j=1}^N s_+(j)s_-(i) + \sum_{i=1}^N \sum_{j=1}^N s_z(i)s_z(j) \\
&= \sum_{i=1}^N s_+(i) \sum_{j=1}^N s_-(j) + \frac{1}{2} \sum_{i=1}^N -2s_z(i) + 0 + \sum_{i=1}^N \sum_{j=1}^N s_z(i)s_z(j) \\
&= \sum_{i=1}^N s_+(i) \sum_{j=1}^N s_-(j) - \sum_{i=1}^N s_z(i) + \sum_{i=1}^N s_z(i) \sum_{j=1}^N s_z(j) \\
&= \mathcal{S}_+ \mathcal{S}_- - \mathcal{S}_z + \mathcal{S}_z^2.
\end{aligned}$$

Thus,

$$\mathcal{S}^2 = \mathcal{S}_+ \mathcal{S}_- - \mathcal{S}_z + \mathcal{S}_z^2. \quad (2.38-1)$$

The second result can be further simplified to

$$\begin{aligned}
&\sum_{i=1}^N \sum_{j=1}^N s_-(i)s_+(j) + \frac{1}{2} [s_+(i)s_-(j) - s_-(i)s_+(j)] + s_z(i)s_z(j) \\
&= \sum_{i=1}^N s_-(i) \sum_{j=1}^N s_+(j) + \frac{1}{2} \sum_{i=1}^N \sum_{j=1}^N s_+(i)s_-(j) - \frac{1}{2} \sum_{i=1}^N \sum_{j=1}^N s_-(i)s_+(j) + \sum_{i=1}^N \sum_{j=1}^N s_z(i)s_z(j) \\
&= \sum_{i=1}^N s_-(i) \sum_{j=1}^N s_+(j) + \frac{1}{2} \sum_{i=1}^N \sum_{j=1}^N [s_+(i), s_-(j)] + s_-(j)s_+(i) - \frac{1}{2} \sum_{i=1}^N \sum_{j=1}^N s_-(j)s_+(i) + \sum_{i=1}^N \sum_{j=1}^N s_z(i)s_z(j) \\
&= \sum_{i=1}^N s_-(i) \sum_{j=1}^N s_+(j) + \frac{1}{2} \sum_{i=1}^N [s_+(i), s_-(i)] + \frac{1}{2} \sum_{i=1}^N \sum_{\substack{j=1 \\ j \neq i}}^N [s_+(i), s_-(j)]
\end{aligned}$$

$$\begin{aligned}
& + \frac{1}{2} \sum_{i=1}^N \sum_{j=1}^N s_-(j) s_+(i) - \frac{1}{2} \sum_{i=1}^N \sum_{j=1}^N s_-(j) s_+(i) + \sum_{i=1}^N \sum_{j=1}^N s_z(i) s_z(j) \\
& = \sum_{i=1}^N s_-(i) \sum_{j=1}^N s_+(j) + \frac{1}{2} \sum_{i=1}^N 2s_z(i) + 0 + \sum_{i=1}^N \sum_{j=1}^N s_z(i) s_z(j) \\
& = \sum_{i=1}^N s_-(i) \sum_{j=1}^N s_+(j) + \sum_{i=1}^N s_z(i) + \sum_{i=1}^N s_z(i) \sum_{j=1}^N s_z(j) \\
& = \mathcal{S}_- \mathcal{S}_+ + \mathcal{S}_z + \mathcal{S}_z^2.
\end{aligned}$$

Thus,

$$\mathcal{S}^2 = \mathcal{S}_- \mathcal{S}_+ + \mathcal{S}_z + \mathcal{S}_z^2. \quad (2.38-2)$$

Secondly, we find that with $N = 2n$,

$$\begin{aligned}
\mathcal{S}_+ |\psi_1 \bar{\psi}_1 \cdots \psi_n \bar{\psi}_n\rangle & = \sum_{i=1}^{2n} s_+(i) |\psi_1 \bar{\psi}_1 \cdots \psi_n \bar{\psi}_n\rangle \\
& = \sum_{i=1}^n s_+(2i-1) |\psi_1 \bar{\psi}_1 \cdots \psi_n \bar{\psi}_n\rangle + \sum_{i=1}^n s_+(2i) |\psi_1 \bar{\psi}_1 \cdots \psi_n \bar{\psi}_n\rangle.
\end{aligned}$$

Note that

$$\begin{aligned}
& s_+(2i-1) |\psi_1 \bar{\psi}_1 \cdots \psi_n \bar{\psi}_n\rangle \\
& = s_+(2i-1) \frac{1}{\sqrt{N!}} \sum_k (-1)^k \mathcal{P}_k \psi_1(1) \bar{\psi}_1(2) \cdots \psi_j(2i-1) \bar{\psi}_j(2i) \cdots \psi_l(2n-1) \bar{\psi}_l(2n) \\
& = \frac{1}{\sqrt{N!}} \sum_k (-1)^k \mathcal{P}_k \psi_1(1) \bar{\psi}_1(2) \cdots [s_+(2i-1) \psi_j(2i-1)] \bar{\psi}_j(2i) \cdots \psi_l(2n-1) \bar{\psi}_l(2n) \\
& = \frac{1}{\sqrt{N!}} \sum_k (-1)^k \mathcal{P}_k \psi_1(1) \bar{\psi}_1(2) \cdots 0 \bar{\psi}_j(2i) \cdots \psi_l(2n-1) \bar{\psi}_l(2n) = 0, \\
& s_+(2i) |\psi_1 \bar{\psi}_1 \cdots \psi_n \bar{\psi}_n\rangle \\
& = s_+(2i) \frac{1}{\sqrt{N!}} \sum_k (-1)^k \mathcal{P}_k \psi_1(1) \bar{\psi}_1(2) \cdots \psi_j(2i-1) \bar{\psi}_j(2i) \cdots \psi_l(2n-1) \bar{\psi}_l(2n) \\
& = \frac{1}{\sqrt{N!}} \sum_k (-1)^k \mathcal{P}_k \psi_1(1) \bar{\psi}_1(2) \cdots \psi_j(2i-1) [s_+(2i) \bar{\psi}_j(2i)] \cdots \psi_l(2n-1) \bar{\psi}_l(2n) \\
& = \frac{1}{\sqrt{N!}} \sum_k (-1)^k \mathcal{P}_k \psi_1(1) \bar{\psi}_1(2) \cdots \psi_j(2i-1) \psi_j(2i) \cdots \psi_l(2n-1) \bar{\psi}_l(2n) \\
& = |\psi_1 \bar{\psi}_1 \cdots \psi_j \psi_j \psi_n \bar{\psi}_n\rangle = 0.
\end{aligned}$$

Thus,

$$\mathcal{S}_+ |\psi_1 \bar{\psi}_1 \cdots \psi_n \bar{\psi}_n\rangle = \sum_{i=1}^n s_+(2i-1) |\psi_1 \bar{\psi}_1 \cdots \psi_n \bar{\psi}_n\rangle + \sum_{i=1}^n s_+(2i) |\psi_1 \bar{\psi}_1 \cdots \psi_n \bar{\psi}_n\rangle = 0 + 0 = 0,$$

and using (2.254),

$$\begin{aligned}
\mathcal{S}^2 |\psi_1 \bar{\psi}_1 \cdots \psi_n \bar{\psi}_n\rangle & = (\mathcal{S}_- \mathcal{S}_+ + \mathcal{S}_z + \mathcal{S}_z^2) |\psi_1 \bar{\psi}_1 \cdots \psi_n \bar{\psi}_n\rangle \\
& = \mathcal{S}_- \mathcal{S}_+ |\psi_1 \bar{\psi}_1 \cdots \psi_n \bar{\psi}_n\rangle + \mathcal{S}_z |\psi_1 \bar{\psi}_1 \cdots \psi_n \bar{\psi}_n\rangle + \mathcal{S}_z \mathcal{S}_z |\psi_1 \bar{\psi}_1 \cdots \psi_n \bar{\psi}_n\rangle = 0.
\end{aligned}$$

In conclusion, (2.256) has been proved.

Exercise 2.39

Using $\mathcal{S}^2 = \mathcal{S}_- \mathcal{S}_+ + \mathcal{S}_z + \mathcal{S}_z^2$, show that $|\Psi_1^2\rangle$ is a singlet while $|\Psi_1^2\rangle$, $|\Psi_1^2\rangle$ and $|\Psi_1^2\rangle$ are triplets.

Solution 2.39

Note that spin operators like \mathcal{S}^2 only inspect the spin properties. Note that

$$\begin{aligned}
 \mathcal{S}_z \alpha(1)\alpha(2) &= (s_z(1) + s_z(2))\alpha(1)\alpha(2) = s_z(1)\alpha(1)\alpha(2) + s_z(2)\alpha(1)\alpha(2) \\
 &= s_z(1)\alpha(1)\alpha(2) + \alpha(1)s_z(2)\alpha(2) = \frac{1}{2}\alpha(1)\alpha(2) + \alpha(1)\frac{1}{2}\alpha(2) = \alpha(1)\alpha(2), \\
 \mathcal{S}_+ \alpha(1)\alpha(2) &= (s_+(1) + s_+(2))\alpha(1)\alpha(2) = s_+(1)\alpha(1)\alpha(2) + s_+(2)\alpha(1)\alpha(2) \\
 &= s_+(1)\alpha(1)\alpha(2) + \alpha(1)s_+(2)\alpha(2) = 0\alpha(2) + \alpha(1)0 = 0, \\
 \mathcal{S}^2 \alpha(1)\alpha(2) &= (\mathcal{S}_- \mathcal{S}_+ + \mathcal{S}_z + \mathcal{S}_z^2)\alpha(1)\alpha(2) \\
 &= \mathcal{S}_- \mathcal{S}_+ \alpha(1)\alpha(2) + \mathcal{S}_z \alpha(1)\alpha(2) + \mathcal{S}_z^2 \alpha(1)\alpha(2) \\
 &= 0 + \alpha(1)\alpha(2) + \mathcal{S}_z \alpha(1)\alpha(2) = \alpha(1)\alpha(2) + \alpha(1)\alpha(2) = 2\alpha(1)\alpha(2) = 1(1+1)\alpha(1)\alpha(2).
 \end{aligned}$$

Thus, we conclude that $|\Psi_1^2\rangle$ is a triplet. In the same way, note that

$$\begin{aligned}
 \mathcal{S}_z \beta(1)\beta(2) &= -\beta(1)\beta(2), \\
 \mathcal{S}_- \beta(1)\beta(2) &= 0, \\
 \mathcal{S}^2 \beta(1)\beta(2) &= (\mathcal{S}_+ \mathcal{S}_- - \mathcal{S}_z + \mathcal{S}_z^2)\beta(1)\beta(2) = 2\beta(1)\beta(2) = 1(1+1)\beta(1)\beta(2).
 \end{aligned}$$

Hence, $|\Psi_1^2\rangle$ is also a triplet.

And similarly, our calculation is listed below.

$$\begin{aligned}
 \mathcal{S}_z(\alpha(1)\beta(2) - \beta(1)\alpha(2)) &= 0, \\
 \mathcal{S}_-(\alpha(1)\beta(2) - \beta(1)\alpha(2)) &= \beta(1)\beta(2) - \beta(1)\beta(2) = 0, \\
 \mathcal{S}^2(\alpha(1)\beta(2) - \beta(1)\alpha(2)) &= 0 = 0(0+1)(\alpha(1)\beta(2) - \beta(1)\alpha(2)), \\
 \mathcal{S}_z(\alpha(1)\beta(2) + \beta(1)\alpha(2)) &= 0, \\
 \mathcal{S}_-(\alpha(1)\beta(2) + \beta(1)\alpha(2)) &= \beta(1)\beta(2) + \beta(1)\beta(2) = 2\beta(1)\beta(2), \\
 \mathcal{S}_+(\beta(1)\beta(2) + \alpha(1)\alpha(2)) &= \alpha(1)\beta(2) + \beta(1)\alpha(2), \\
 \mathcal{S}^2(\alpha(1)\beta(2) + \beta(1)\alpha(2)) &= 2\mathcal{S}_+(\beta(1)\beta(2) + \alpha(1)\alpha(2)) = 1(1+1)(\alpha(1)\beta(2) + \beta(1)\alpha(2)).
 \end{aligned}$$

Therefore, we conclude that $|\Psi_1^2\rangle$ is a triplet while $|\Psi_1^2\rangle$ is a singlet.

Exercise 2.40

Show that

$$\begin{aligned}
 \langle \Psi_1^2 | \mathcal{H} | \Psi_1^2 \rangle &= h_{11} + h_{22} + J_{12} + K_{12}, \\
 \langle \Psi_1^2 | \mathcal{H} | \Psi_1^2 \rangle &= h_{11} + h_{22} + J_{12} - K_{12}.
 \end{aligned}$$

Note that the energy of the triplet is lower than the energy of the singlet. Why is this to be expected from the space parts of the two wave functions?

Solution 2.40

The energy of the singlet and triplet can be calculated directly.

$$\begin{aligned}
 \langle \Psi_1^2 | \mathcal{H} | \Psi_1^2 \rangle &= \left(\frac{1}{\sqrt{2}} \langle 1\bar{2} | + \frac{1}{\sqrt{2}} \langle 2\bar{1} | \right) \mathcal{H} \left(\frac{1}{\sqrt{2}} | 1\bar{2} \rangle + \frac{1}{\sqrt{2}} | 2\bar{1} \rangle \right) \\
 &= \frac{1}{2} (\langle 1\bar{2} | \mathcal{H} | 1\bar{2} \rangle + \langle 1\bar{2} | \mathcal{H} | 2\bar{1} \rangle + \langle 2\bar{1} | \mathcal{H} | 1\bar{2} \rangle + \langle 2\bar{1} | \mathcal{H} | 2\bar{1} \rangle) \\
 &= \frac{1}{2} (h_{11} + h_{22} + J_{12} + K_{12} + K_{12} + h_{11} + h_{22} + J_{12}) \\
 &= h_{11} + h_{22} + J_{12} + K_{12}, \\
 \langle \Psi_1^2 | \mathcal{H} | \Psi_1^2 \rangle &= \left(\frac{1}{\sqrt{2}} \langle 1\bar{2} | - \frac{1}{\sqrt{2}} \langle 2\bar{1} | \right) \mathcal{H} \left(\frac{1}{\sqrt{2}} | 1\bar{2} \rangle - \frac{1}{\sqrt{2}} | 2\bar{1} \rangle \right) \\
 &= \frac{1}{2} (\langle 1\bar{2} | \mathcal{H} | 1\bar{2} \rangle - \langle 1\bar{2} | \mathcal{H} | 2\bar{1} \rangle - \langle 2\bar{1} | \mathcal{H} | 1\bar{2} \rangle + \langle 2\bar{1} | \mathcal{H} | 2\bar{1} \rangle)
 \end{aligned}$$

$$\begin{aligned}
&= \frac{1}{2} (h_{11} + h_{22} + J_{12} - K_{12} - K_{12} + h_{11} + h_{22} + J_{12}) \\
&= h_{11} + h_{22} + J_{12} - K_{12}.
\end{aligned}$$

In fact, we can find that

$$\langle {}^1\Psi_1^2 | \mathcal{H} | {}^1\Psi_1^2 \rangle - \langle {}^3\Psi_1^2 | \mathcal{H} | {}^3\Psi_1^2 \rangle = 2K_{12} > 0.$$

The energy difference between the singlet and the triplet results is a two-electron term. The only two-electron term in the Hamiltonian \mathcal{H} is the Coulomb interaction. From the expression of the singlet and the triplet, we find that in the triplet, two electrons are more concentrated while two electrons are diffused. It is evident that the former has a lower electrostatic potential. Thus, from the perspective of classical physics, the triplet state should have lower energy.

2.5.3 Unrestricted Determinants

Exercise 2.41

Consider the determinant $|K\rangle = |\psi_1^\alpha \bar{\psi}_1^\beta\rangle$ formed from *nonorthogonal* spatial orbitals, $\langle \psi_1^\alpha | \psi_1^\beta \rangle = S_{11}^{\alpha\beta}$.

- Show that $|K\rangle$ is an eigenfunction of \mathcal{S}^2 only if $\psi_1^\alpha = \psi_1^\beta$.
- Show that $\langle K | \mathcal{S}^2 | K \rangle = 1 - |S_{11}^{\alpha\beta}|^2$ in agreement with Eq.(2.271).

Solution 2.41

- For any $|K\rangle = |\psi_1^\alpha \bar{\psi}_1^\beta\rangle$, $\mathcal{S}_z |K\rangle = 0$. Thus, $|K\rangle$ is an eigenfunction of \mathcal{S}^2 if and only if it is also an eigenfunction of $\mathcal{S}_- \mathcal{S}_+$. However, we can find that

$$\begin{aligned}
\mathcal{S}_- \mathcal{S}_+ |K\rangle &= \mathcal{S}_- (s_+(1) + s_+(2)) \frac{1}{\sqrt{2}} \left(\psi_1^\alpha(1) \alpha(1) \psi_1^\beta(2) \beta(2) - \psi_1^\beta(1) \beta(1) \psi_1^\alpha(2) \alpha(2) \right) \\
&= \frac{1}{\sqrt{2}} \mathcal{S}_- \left(\psi_1^\alpha(1) \alpha(1) \psi_1^\beta(2) \alpha(2) - \psi_1^\beta(1) \alpha(1) \psi_1^\alpha(2) \alpha(2) \right) \\
&= \frac{1}{\sqrt{2}} (s_-(1) + s_-(2)) \left(\psi_1^\alpha(1) \alpha(1) \psi_1^\beta(2) \alpha(2) - \psi_1^\beta(1) \alpha(1) \psi_1^\alpha(2) \alpha(2) \right) \\
&= \frac{1}{\sqrt{2}} \left(\psi_1^\alpha(1) \beta(1) \psi_1^\beta(2) \alpha(2) + \psi_1^\alpha(1) \alpha(1) \psi_1^\beta(2) \beta(2) \right. \\
&\quad \left. - \psi_1^\beta(1) \beta(1) \psi_1^\alpha(2) \alpha(2) - \psi_1^\beta(1) \alpha(1) \psi_1^\alpha(2) \beta(2) \right) \\
&= |\psi_1^\alpha \bar{\psi}_1^\beta\rangle + |\bar{\psi}_1^\alpha \psi_1^\beta\rangle = |K\rangle + |\bar{\psi}_1^\alpha \psi_1^\beta\rangle.
\end{aligned}$$

Thus, $|\bar{\psi}_1^\alpha \psi_1^\beta\rangle$ is also an eigenfunction of \mathcal{S}^2 , belonging to the same eigenvalue as $|K\rangle$. Here, we apply the contrapositive of a lemma that if an operator \mathcal{A} has two eigenvector a_1 and a_2 which belong to different eigenvalues, for any real numbers λ_1 and λ_2 , $\lambda_1 a_1 + \lambda_2 a_2$ cannot be a eigenvector of \mathcal{A} .

It is evident that $|\bar{\psi}_1^\alpha \psi_1^\beta\rangle$ should be either a triplet or a singlet. Note that

$$\mathcal{S}_z |\bar{\psi}_1^\alpha \psi_1^\beta\rangle = 0, \quad \mathcal{S}_z |K\rangle = 0.$$

- If it is a triplet ($S = 1$), $|K\rangle$ will be a triplet, too. However, we find that

$$\begin{aligned}
2|K\rangle &= 1(1+1)|K\rangle = \mathcal{S}^2 |K\rangle = (\mathcal{S}_- \mathcal{S}_+ + \mathcal{S}_z + \mathcal{S}_z^2) |K\rangle \\
&= \mathcal{S}_- \mathcal{S}_+ |K\rangle + \mathcal{S}_z |K\rangle + \mathcal{S}_z^2 |K\rangle = \mathcal{S}_- \mathcal{S}_+ |K\rangle + 0 + 0 \\
&= |K\rangle + |\bar{\psi}_1^\alpha \psi_1^\beta\rangle,
\end{aligned}$$

which requires

$$|K\rangle \equiv |\psi_1^\alpha \bar{\psi}_1^\beta\rangle = |\bar{\psi}_1^\alpha \psi_1^\beta\rangle = -|\psi_1^\beta \bar{\psi}_1^\alpha\rangle.$$

Nevertheless, it is evident that this is forbidden.

– If it is a singlet ($S = 0$), $|K\rangle$ will be a singlet, too. We can find that

$$\begin{aligned} 0|K\rangle &= 0(0+1)|K\rangle = \mathcal{S}^2|K\rangle = (\mathcal{S}_-\mathcal{S}_+ + \mathcal{S}_z + \mathcal{S}_z^2)|K\rangle \\ &= \mathcal{S}_-\mathcal{S}_+|K\rangle + \mathcal{S}_z|K\rangle + \mathcal{S}_z^2|K\rangle = \mathcal{S}_-\mathcal{S}_+|K\rangle \\ &= |K\rangle + |\bar{\psi}_1^\alpha \psi_1^\beta\rangle, \end{aligned}$$

which requires

$$|K\rangle \equiv |\psi_1^\alpha \bar{\psi}_1^\beta\rangle = -|\bar{\psi}_1^\alpha \psi_1^\beta\rangle = |\psi_1^\beta \bar{\psi}_1^\alpha\rangle \Leftrightarrow \psi_1^\alpha = \psi_1^\beta.$$

Thus, we conclude that $|K\rangle$ is an eigenfunction of \mathcal{S}^2 if and only if $\psi_1^\alpha = \psi_1^\beta$.

b. Now we know that

$$\mathcal{S}_-\mathcal{S}_+|K\rangle = |K\rangle + |\bar{\psi}_1^\alpha \psi_1^\beta\rangle.$$

Thus,

$$\langle K|\mathcal{S}^2|K\rangle = \langle K|\mathcal{S}_-\mathcal{S}_+ + \mathcal{S}_z + \mathcal{S}_z^2|K\rangle = \langle K|\mathcal{S}_-\mathcal{S}_+|K\rangle = \langle K(|K\rangle + |\bar{\psi}_1^\alpha \psi_1^\beta\rangle) = 1 + \langle K|\bar{\psi}_1^\alpha \psi_1^\beta\rangle.$$

While

$$\begin{aligned} &\langle K|\bar{\psi}_1^\alpha \psi_1^\beta\rangle \\ &= \int d\mathbf{r}_1 \int d\mathbf{r}_2 \int d\omega_1 \int d\omega_2 \frac{1}{\sqrt{2}} \left(\psi_1^\alpha(1)\alpha(1)\psi_1^\beta(2)\beta(2) - \psi_1^\beta(1)\beta(1)\psi_1^\alpha(2)\alpha(2) \right)^* \\ &\quad \frac{1}{\sqrt{2}} \left(\psi_1^\alpha(1)\beta(1)\psi_1^\beta(2)\alpha(2) - \psi_1^\beta(1)\alpha(1)\psi_1^\alpha(2)\beta(2) \right) \\ &= \frac{1}{2} \left[\int d\mathbf{r}_1 \psi_1^{\alpha*}(1)\psi_1^\alpha(1) \int d\mathbf{r}_2 \psi_1^{\beta*}(2)\psi_1^\beta(2) \int d\omega_1 \alpha^*(1)\beta(1) \int d\omega_2 \beta^*(2)\alpha(2) \right. \\ &\quad - \int d\mathbf{r}_1 \psi_1^{\alpha*}(1)\psi_1^\beta(1) \int d\mathbf{r}_2 \psi_1^{\beta*}(2)\psi_1^\alpha(2) \int d\omega_1 \alpha^*(1)\alpha(1) \int d\omega_2 \beta^*(2)\beta(2) \\ &\quad - \int d\mathbf{r}_1 \psi_1^{\beta*}(1)\psi_1^\alpha(1) \int d\mathbf{r}_2 \psi_1^{\alpha*}(2)\psi_1^\beta(2) \int d\omega_1 \beta^*(1)\beta(1) \int d\omega_2 \alpha^*(2)\alpha(2) \\ &\quad \left. + \int d\mathbf{r}_1 \psi_1^{\beta*}(1)\psi_1^\beta(1) \int d\mathbf{r}_2 \psi_1^{\alpha*}(2)\psi_1^\alpha(2) \int d\omega_1 \beta^*(1)\alpha(1) \int d\omega_2 \alpha^*(2)\beta(2) \right] \\ &= \frac{1}{2} \left(0 - |S_{11}^{\alpha\beta}|^2 - |S_{11}^{\alpha\beta}|^2 + 0 \right) = -|S_{11}^{\alpha\beta}|^2. \end{aligned}$$

Therefore, we arrive at

$$\langle K|\mathcal{S}^2|K\rangle = 1 + \langle K|\bar{\psi}_1^\alpha \psi_1^\beta\rangle = 1 - |S_{11}^{\alpha\beta}|^2. \quad (2.41-1)$$

Moreover, when restricted spin orbitals are used, namely $\psi_1^\alpha = \psi_1^\beta$, at this time,

$$S_{11}^{\alpha\beta} = 1,$$

at once we get

$$\langle K|\mathcal{S}^2|K\rangle = 1 - 1^2 = 1 - 1 = 0,$$

which means that $|K\rangle$ is a singlet, indeed. Hence $|K\rangle$ will not be a singlet unless $\psi_1^\alpha = \psi_1^\beta$.

CHAPTER 4

Configuration Interaction

4.1 Multiconfigurational Wave Functions and the Structure of the Full CI Matrix

4.1.1 Intermediate Normalization and an Expression for the Correlation Energy

Exercise 4.1

Obtain Eq.(4.12) from Eq.(4.11). It will prove convenient to use unrestricted summations.

Solution 4.1

Note that the index r must be included in the set $\{t, u, v\}$ and the index a must be included in the set $\{c, d, e\}$ for a matrix element of $\langle \Psi_a^r | \mathcal{H} | \Psi_{cde}^{tuv} \rangle$. Therefore, we find that

$$\begin{aligned} \sum_{\substack{c < d < e \\ t < u < v}} \langle \Psi_a^r | \mathcal{H} | \Psi_{cde}^{tuv} \rangle c_{cde}^{tuv} &= \frac{1}{(3!)^2} \sum_{\substack{cde \\ tuv}} \langle \Psi_a^r | \mathcal{H} | \Psi_{cde}^{tuv} \rangle c_{cde}^{tuv} \\ &= \frac{1}{(3!)^2} \left[\sum_{\substack{de \\ uv}} \langle \Psi_a^r | \mathcal{H} | \Psi_{ade}^{ruv} \rangle c_{ade}^{ruv} + \sum_{\substack{de \\ tv}} \langle \Psi_a^r | \mathcal{H} | \Psi_{ade}^{trv} \rangle c_{ade}^{trv} + \sum_{\substack{de \\ tu}} \langle \Psi_a^r | \mathcal{H} | \Psi_{ade}^{tur} \rangle c_{ade}^{tur} \right. \\ &\quad + \sum_{\substack{ce \\ uv}} \langle \Psi_a^r | \mathcal{H} | \Psi_{cae}^{ruv} \rangle c_{cae}^{ruv} + \sum_{\substack{ce \\ tv}} \langle \Psi_a^r | \mathcal{H} | \Psi_{cae}^{trv} \rangle c_{cae}^{trv} + \sum_{\substack{ce \\ tu}} \langle \Psi_a^r | \mathcal{H} | \Psi_{cae}^{tur} \rangle c_{cae}^{tur} \\ &\quad \left. + \sum_{\substack{cd \\ uv}} \langle \Psi_a^r | \mathcal{H} | \Psi_{cda}^{ruv} \rangle c_{cda}^{ruv} + \sum_{\substack{cd \\ tv}} \langle \Psi_a^r | \mathcal{H} | \Psi_{cda}^{trv} \rangle c_{cda}^{trv} + \sum_{\substack{cd \\ tu}} \langle \Psi_a^r | \mathcal{H} | \Psi_{cda}^{tur} \rangle c_{cda}^{tur} \right]. \end{aligned}$$

Then, these dummy indices should be converted into the same one, viz.,

$$\begin{aligned} \sum_{\substack{c < d < e \\ t < u < v}} \langle \Psi_a^r | \mathcal{H} | \Psi_{cde}^{tuv} \rangle c_{cde}^{tuv} &= \frac{1}{(3!)^2} \left[\sum_{\substack{cd \\ tu}} \langle \Psi_a^r | \mathcal{H} | \Psi_{acd}^{rtu} \rangle c_{acd}^{rtu} + \sum_{\substack{cd \\ tu}} \langle \Psi_a^r | \mathcal{H} | \Psi_{acd}^{tru} \rangle c_{acd}^{tru} + \sum_{\substack{cd \\ tu}} \langle \Psi_a^r | \mathcal{H} | \Psi_{acd}^{tur} \rangle c_{acd}^{tur} \right. \\ &\quad + \sum_{\substack{cd \\ tu}} \langle \Psi_a^r | \mathcal{H} | \Psi_{cad}^{rtu} \rangle c_{cad}^{rtu} + \sum_{\substack{cd \\ tu}} \langle \Psi_a^r | \mathcal{H} | \Psi_{cad}^{tru} \rangle c_{cad}^{tru} + \sum_{\substack{cd \\ tu}} \langle \Psi_a^r | \mathcal{H} | \Psi_{cad}^{tur} \rangle c_{cad}^{tur} \\ &\quad \left. + \sum_{\substack{cd \\ tu}} \langle \Psi_a^r | \mathcal{H} | \Psi_{cda}^{rtu} \rangle c_{cda}^{rtu} + \sum_{\substack{cd \\ tu}} \langle \Psi_a^r | \mathcal{H} | \Psi_{cda}^{tru} \rangle c_{cda}^{tru} + \sum_{\substack{cd \\ tu}} \langle \Psi_a^r | \mathcal{H} | \Psi_{cda}^{tur} \rangle c_{cda}^{tur} \right] \\ &= \frac{1}{(3!)^2} \times 9 \sum_{\substack{cd \\ tu}} \langle \Psi_a^r | \mathcal{H} | \Psi_{acd}^{rtu} \rangle c_{acd}^{rtu} = \frac{1}{(2!)^2} \sum_{\substack{cd \\ tu}} \langle \Psi_a^r | \mathcal{H} | \Psi_{acd}^{rtu} \rangle c_{acd}^{rtu} = \sum_{\substack{c < d \\ t < u}} \langle \Psi_a^r | \mathcal{H} | \Psi_{acd}^{rtu} \rangle c_{acd}^{rtu}. \end{aligned}$$

Thus, we have proved that

$$\sum_{\substack{c < d < e \\ t < u < v}} \langle \Psi_a^r | \mathcal{H} | \Psi_{cde}^{tuv} \rangle c_{cde}^{tuv} = \sum_{\substack{c < d \\ t < u}} \langle \Psi_a^r | \mathcal{H} | \Psi_{acd}^{rtu} \rangle c_{acd}^{rtu}. \quad (4.1-1)$$

With this equation, it is clear that (4.12) can be obtained from (4.11).

Exercise 4.2

Using the secular determinant approach show that the lowest eigenvalue of the matrix

$$\begin{pmatrix} 0 & K_{12} \\ K_{12} & 2\Delta \end{pmatrix}$$

is given by Eq.(4.23).

Solution 4.2

The introduction of the secular determinant approach is demonstrated in the page 18. The matrix in the Exercise 4.2 is denoted as H , then

$$\det(H - \varepsilon \mathbf{1}) = \begin{vmatrix} -\varepsilon & K_{12} \\ K_{12} & 2\Delta - \varepsilon \end{vmatrix} = \varepsilon^2 - 2\Delta\varepsilon - K_{12}^2 = 0,$$

The discriminant Δ_E of this quadratic equation is

$$\Delta_E = 4\Delta^2 - 4 \times (-K_{12}^2) = 4(\Delta^2 + K_{12}^2)$$

Thus, the root are

$$E_1 = \Delta + \sqrt{\Delta^2 + K_{12}^2}, \quad E_2 = \Delta - \sqrt{\Delta^2 + K_{12}^2}.$$

Therefore, the lowest root is the correlation energy, viz.,

$$E_{\text{corr}} = \Delta - \sqrt{\Delta^2 + K_{12}^2}. \quad (4.2-1)$$

Exercise 4.3

Calculate the coefficient of the double excitation (c) in the intermediate normalized CI wave function at $R = 1.4$ a.u., using the STO-3G integrals given in Appendix D. Show analytically that as $R \rightarrow \infty$, $c \rightarrow -1$, and hence that at large distances the Hartree-Fock ground state and the doubly excited configuration have equal weight in the CI ground state. Finally, show that the CI wave function, when normalized to unity, becomes (at $R = \infty$)

$$\frac{1}{\sqrt{2}} (|\phi_1 \bar{\phi}_2\rangle + |\phi_2 \bar{\phi}_1\rangle)$$

where ϕ_1 and ϕ_2 are atomic orbitals on centers one and two, respectively.

Solution 4.3

When $R = 1.4$ a.u., we know that

$$\begin{aligned} \varepsilon_1 &= -0.5782 \text{ a.u.}, & \varepsilon_2 &= 0.6703 \text{ a.u.}, & J_{11} &= 0.6746 \text{ a.u.}, \\ J_{12} &= 0.6636 \text{ a.u.}, & J_{22} &= 0.6975 \text{ a.u.}, & K_{12} &= 0.1813 \text{ a.u.} \end{aligned}$$

Firstly, with (4.20), we calculate 2Δ at $R = 1.4$ a.u., viz.,

$$2\Delta = [2(\varepsilon_2 - \varepsilon_1) + J_{11} + J_{22} - 4J_{12} + 2K_{12}] = 1.5773 \text{ a.u.}$$

In other words, $\Delta = 0.78865$ a.u. Thus, the correlation energy E_{corr} at $R = 1.4$ a.u. is

$$E_{\text{corr}} = \Delta - \sqrt{\Delta^2 + K_{12}^2} = -0.02057 \text{ a.u.}$$

Therefore,

$$c = \frac{K_{12}}{E_{\text{corr}} - 2\Delta} = \frac{0.1813 \text{ a.u.}}{-0.02057 \text{ a.u.} - 1.5773 \text{ a.u.}} \approx -0.1135. \quad (4.3-1)$$

Indeed, we can find that

$$\Delta = \varepsilon_2 - \varepsilon_1 + \frac{1}{2}J_{11} + \frac{1}{2}J_{22} - 2J_{12} + K_{12} = h_{22} - h_{11} - \frac{1}{2}J_{11} + \frac{1}{2}J_{12}.$$

It is clear that

$$\lim_{R \rightarrow \infty} \Delta = \lim_{R \rightarrow \infty} \left[h_{22} - h_{11} + \frac{1}{2}J_{22} - \frac{1}{2}J_{11} \right] = E(\text{H}) - E(\text{H}) + \frac{1}{4}(\phi_1\phi_1|\phi_1\phi_1) - \frac{1}{4}(\phi_1\phi_1|\phi_1\phi_1) = 0.$$

Thus,

$$\begin{aligned} \lim_{R \rightarrow \infty} c &= \lim_{R \rightarrow \infty} \frac{K_{12}}{E_{\text{corr}} - 2\Delta} = \lim_{R \rightarrow \infty} \frac{K_{12}}{\Delta - \sqrt{\Delta^2 + K_{12}^2} - 2\Delta} = \lim_{R \rightarrow \infty} \frac{-K_{12}}{\Delta + \sqrt{\Delta^2 + K_{12}^2}} \\ &= - \lim_{\Delta \rightarrow 0} \frac{1}{\frac{\Delta}{K_{12}} + \sqrt{1 + \left(\frac{\Delta}{K_{12}}\right)^2}} = - \lim_{x \rightarrow 0} \frac{1}{x + \sqrt{1 + x^2}} = -1. \end{aligned}$$

This conclusion means that at large distances the Hartree-Fock ground state Ψ_0 and the doubly excited configuration Ψ_{11}^{22} have equal weight in the CI ground state Φ , viz.,

$$\lim_{R \rightarrow \infty} |\Phi\rangle = |\Psi_0\rangle - |\Psi_{11}^{22}\rangle = |\psi_1\bar{\psi}_1\rangle - |\psi_2\bar{\psi}_2\rangle.$$

Note that as $R \rightarrow \infty$, from (3.236) and (3.237), we find that

$$\lim_{R \rightarrow \infty} \psi_1 = \frac{1}{\sqrt{2}}(\phi_1 + \phi_2), \quad \lim_{R \rightarrow \infty} \psi_2 = \frac{1}{\sqrt{2}}(\phi_1 - \phi_2).$$

Thus,

$$\begin{aligned} \lim_{R \rightarrow \infty} |\psi_1\bar{\psi}_1\rangle &= \frac{1}{2}|(\phi_1 + \phi_2)(\bar{\phi}_1 + \bar{\phi}_2)\rangle = \frac{1}{2}(|\phi_1\bar{\phi}_1\rangle + |\phi_1\bar{\phi}_2\rangle + |\phi_2\bar{\phi}_1\rangle + |\phi_2\bar{\phi}_2\rangle), \\ \lim_{R \rightarrow \infty} |\psi_2\bar{\psi}_2\rangle &= \frac{1}{2}|(\phi_1 - \phi_2)(\bar{\phi}_1 - \bar{\phi}_2)\rangle = \frac{1}{2}(|\phi_1\bar{\phi}_1\rangle - |\phi_1\bar{\phi}_2\rangle - |\phi_2\bar{\phi}_1\rangle + |\phi_2\bar{\phi}_2\rangle), \end{aligned}$$

and then

$$\lim_{R \rightarrow \infty} |\Phi\rangle = \lim_{R \rightarrow \infty} |\psi_1\bar{\psi}_1\rangle - \lim_{R \rightarrow \infty} |\psi_2\bar{\psi}_2\rangle = |\phi_1\bar{\phi}_2\rangle + |\phi_2\bar{\phi}_1\rangle$$

Thus, at $R = \infty$, the normalized CI wave function is

$$\lim_{R \rightarrow \infty} |\Phi\rangle = \lim_{R \rightarrow \infty} \frac{1}{\langle \Phi_0 | \Phi_0 \rangle} |\Phi_0\rangle = \frac{1}{\sqrt{2}} (|\phi_1\bar{\phi}_2\rangle + |\phi_2\bar{\phi}_1\rangle). \quad (4.3-2)$$

We have proved two conclusions at $R = \infty$, the equal weight of the Hartree-Fock ground state Ψ_0 and the doubly excited configuration Ψ_{11}^{22} , and the form of normalized CI wave function.

4.2 Doubly Excited CI

4.3 Some Illustrative Calculations

4.4 Natural Orbitals and the One-Particle Reduced Density Matrix

Exercise 4.4

Show that γ is a Hermitian matrix.

Solution 4.4

Firstly, we find that $\gamma(\mathbf{x}_1, \mathbf{x}'_1)$ is Hermite, viz.,

$$\begin{aligned}\gamma^*(\mathbf{x}_1, \mathbf{x}'_1) &= \left(N \int_{\mathbb{R}^3} d\mathbf{x}_2 \cdots \int_{\mathbb{R}^3} d\mathbf{x}_N \Phi^*(\mathbf{x}_1, \mathbf{x}_2, \dots, \mathbf{x}_N) \Phi(\mathbf{x}'_1, \mathbf{x}_2, \dots, \mathbf{x}_N) \right)^* \\ &= N \int_{\mathbb{R}^3} d\mathbf{x}_2 \cdots \int_{\mathbb{R}^3} d\mathbf{x}_N \Phi^*(\mathbf{x}'_1, \mathbf{x}_2, \dots, \mathbf{x}_N) \Phi(\mathbf{x}_1, \mathbf{x}_2, \dots, \mathbf{x}_N) = \gamma(\mathbf{x}'_1, \mathbf{x}_1).\end{aligned}$$

Thus,

$$\begin{aligned}\gamma_{ji}^* &= \left(\int_{\mathbb{R}^3} d\mathbf{x}_1 \int_{\mathbb{R}^3} d\mathbf{x}'_1 \chi_j^*(\mathbf{x}_1) \gamma(\mathbf{x}_1, \mathbf{x}'_1) \chi_i(\mathbf{x}'_1) \right)^* = \int_{\mathbb{R}^3} d\mathbf{x}_1 \int_{\mathbb{R}^3} d\mathbf{x}'_1 \chi_i^*(\mathbf{x}'_1) \gamma^*(\mathbf{x}_1, \mathbf{x}'_1) \chi_j(\mathbf{x}_1) \\ &= \int_{\mathbb{R}^3} d\mathbf{x}_1 \int_{\mathbb{R}^3} d\mathbf{x}'_1 \chi_i^*(\mathbf{x}'_1) \gamma(\mathbf{x}'_1, \mathbf{x}_1) \chi_j(\mathbf{x}_1) = \int_{\mathbb{R}^3} d\mathbf{x}_1 \int_{\mathbb{R}^3} d\mathbf{x}_1 \chi_i^*(\mathbf{x}_1) \gamma(\mathbf{x}_1, \mathbf{x}'_1) \chi_j(\mathbf{x}'_1) = \gamma_{ij}.\end{aligned}$$

Hence we have proved that γ is a Hermitian matrix.

Exercise 4.5

Show that $\text{tr}(\gamma) = N$.

Solution 4.5

With (4.34), (4.36) and (4.37), we find that

$$\begin{aligned}N &= \int_{\mathbb{R}^3} d\mathbf{x}_1 \rho(\mathbf{x}_1) = \int_{\mathbb{R}^3} d\mathbf{x}_1 \gamma(\mathbf{x}_1, \mathbf{x}_1) = \int_{\mathbb{R}^3} d\mathbf{x}_1 \sum_{i=1}^N \sum_{j=1}^N \chi_i(\mathbf{x}_1) \gamma_{ij} \chi_j^*(\mathbf{x}_1) \\ &= \sum_{i=1}^N \sum_{j=1}^N \gamma_{ij} \int_{\mathbb{R}^3} d\mathbf{x}_1 \chi_i(\mathbf{x}_1) \chi_j^*(\mathbf{x}_1) = \sum_{i=1}^N \sum_{j=1}^N \gamma_{ij} \delta_{ij} = \sum_{i=1}^N \gamma_{ii} = \text{tr}(\gamma).\end{aligned}$$

Exercise 4.6

Consider the one-electron operator

$$\mathcal{O}_1 = \sum_{i=1}^N h(i).$$

a. Show that

$$\langle \Phi | \mathcal{O}_1 | \Phi \rangle = \int d\mathbf{x}_1 [h(\mathbf{x}_1) \gamma(\mathbf{x}_1, \mathbf{x}'_1)]_{\mathbf{x}'_1 = \mathbf{x}_1}$$

where the notation $[\]_{\mathbf{x}'_1 = \mathbf{x}_1}$ means that \mathbf{x}'_1 is set equal to \mathbf{x}_1 after $h(\mathbf{x}_1)$ has operated on $\gamma(\mathbf{x}_1, \mathbf{x}'_1)$.

b. Show that

$$\langle \Phi | \mathcal{O}_1 | \Phi \rangle = \text{tr}(\mathbf{h}\gamma)$$

where

$$h_{ij} = \langle i | h | j \rangle = \int d\mathbf{x}_1 \chi_i^*(\mathbf{x}_1) h(\mathbf{x}_1) \chi_j(\mathbf{x}_1).$$

Thus the expectation value of any one-electron operator can be expressed in terms of the one-matrix.

Solution 4.6

a. From the definition of \mathcal{O}_1 , we find that

$$\begin{aligned}\langle \Phi | \mathcal{O}_1 | \Phi \rangle &= \langle \Phi | \sum_{i=1}^N h(i) | \Phi \rangle \\ &= \sum_{i=1}^N \int_{\mathbb{R}^3} d\mathbf{x}_1 \int_{\mathbb{R}^3} d\mathbf{x}_2 \cdots \int_{\mathbb{R}^3} d\mathbf{x}_N \Phi^*(\mathbf{x}_1, \mathbf{x}_2, \dots, \mathbf{x}_N) h(\mathbf{x}_i) \Phi(\mathbf{x}_1, \mathbf{x}_2, \dots, \mathbf{x}_N)\end{aligned}$$

Considering that the different integral variables $d\mathbf{x}_1$ and $d\mathbf{x}_i$ ($i \neq 1$) have the same integral range, it is clear that

$$\begin{aligned}\langle \Phi | \mathcal{O}_1 | \Phi \rangle &= \langle \Phi | \sum_{i=1}^N h(i) | \Phi \rangle \\ &= N \int_{\mathbb{R}^3} d\mathbf{x}_1 \int_{\mathbb{R}^3} d\mathbf{x}_2 \cdots \int_{\mathbb{R}^3} d\mathbf{x}_N \Phi^*(\mathbf{x}_1, \mathbf{x}_2, \dots, \mathbf{x}_N) h(\mathbf{x}_1) \Phi(\mathbf{x}_1, \mathbf{x}_2, \dots, \mathbf{x}_N) \\ &= \int_{\mathbb{R}^3} d\mathbf{x}_1 h(\mathbf{x}_1) \times N \int_{\mathbb{R}^3} d\mathbf{x}_2 \cdots \int_{\mathbb{R}^3} d\mathbf{x}_N \Phi^*(\mathbf{x}_1, \mathbf{x}_2, \dots, \mathbf{x}_N) \Phi(\mathbf{x}_1, \mathbf{x}_2, \dots, \mathbf{x}_N) \\ &= \int_{\mathbb{R}^3} d\mathbf{x}_1 h(\mathbf{x}_1) \rho(\mathbf{x}_1) = \int_{\mathbb{R}^3} d\mathbf{x}_1 h(\mathbf{x}_1) \gamma(\mathbf{x}_1, \mathbf{x}_1) = \int_{\mathbb{R}^3} d\mathbf{x}_1 [h(\mathbf{x}_1) \gamma(\mathbf{x}_1, \mathbf{x}'_1)]_{\mathbf{x}'_1=\mathbf{x}_1}.\end{aligned}$$

b. From the former issue, we know that

$$\begin{aligned}\langle \Phi | \mathcal{O}_1 | \Phi \rangle &= \int_{\mathbb{R}^3} d\mathbf{x}_1 h(\mathbf{x}_1) \gamma(\mathbf{x}_1, \mathbf{x}_1) = \int_{\mathbb{R}^3} d\mathbf{x}_1 h(\mathbf{x}_1) \sum_{i=1}^N \sum_{j=1}^N \chi_i(\mathbf{x}_1) \gamma_{ij} \chi_j^*(\mathbf{x}_1) \\ &= \sum_{i=1}^N \sum_{j=1}^N \gamma_{ij} \int_{\mathbb{R}^3} d\mathbf{x}_1 \chi_i(\mathbf{x}_1) h(\mathbf{x}_1) \chi_j^*(\mathbf{x}_1) = \sum_{i=1}^N \sum_{j=1}^N \gamma_{ij} h_{ji} = \text{tr}(\gamma \mathbf{h}) = \text{tr}(\mathbf{h} \gamma).\end{aligned}$$

The last step uses the conclusion of Exercise 1.4(a).

Exercise 4.7

Recall that in second quantization a one-electron operator is

$$\mathcal{O}_1 = \sum_{ij} \langle i | h | j \rangle a_i^\dagger a_j.$$

a. Show that

$$\gamma_{ij} = \langle \Phi | a_j^\dagger a_i | \Phi \rangle.$$

b. Show that the matrix elements of γ^{HF} are given by Eq.(4.40).

Solution 4.7

a. We can use the conclusion in Exercise 4.6(b), viz.,

$$\text{tr}(\mathbf{h} \gamma) = \sum_{ij} h_{ij} \gamma_{ji} = \langle \Phi | \mathcal{O}_1 | \Phi \rangle.$$

With the second quantization form, we find that

$$\sum_{ij} h_{ij} \gamma_{ji} = \langle \Phi | \mathcal{O}_1 | \Phi \rangle = \sum_{ij} \langle \Phi | a_i^\dagger a_j | \Phi \rangle h_{ij} \Leftrightarrow \sum_{ij} [\langle \Phi | a_i^\dagger a_j | \Phi \rangle - \gamma_{ji}] h_{ij} = 0.$$

For any system, this equation holds true, which means that terms h_{ij} are linearly independent. Thus,

$$\gamma_{ji} = \langle \Phi | a_i^\dagger a_j | \Phi \rangle, \quad (4.7-1)$$

which equals $\gamma_{ij} = \langle \Phi | a_j^\dagger a_i | \Phi \rangle$.

b. If i belongs to unoccupied, $a_i | \Phi \rangle$ vanishes, and so does $\langle \Phi | a_j^\dagger$ if j belongs to unoccupied. If the indices i and j are occupied, viz., $i = a$ and $j = b$, with $a_a^\dagger | \Phi \rangle = 0$,

$$\gamma_{ab}^{\text{HF}} = \langle \Phi | a_a^\dagger a_b | \Phi \rangle = \langle \Phi | (\delta_{ab} - a_b a_a^\dagger) | \Phi \rangle = \delta_{ab} - \langle \Phi | a_b a_a^\dagger | \Phi \rangle = \delta_{ab} - 0 = \delta_{ab}.$$

Thus, we have proved

$$\gamma_{ij}^{\text{HF}} = \begin{cases} \delta_{ij}, & \text{iff } i, j \in \text{occupied}, \\ 0, & \text{otherwise.} \end{cases}$$

Exercise 4.8

For the special case of a two-electron system, the use of natural orbitals dramatically reduces the size of the full CI expansion. If ψ_1 is the occupied Hartree-Fock spatial orbital and ψ_r , $r = 2, 3, \dots, K$ are virtual spatial orbitals, the normalized full CI singlet wave function has the form

$$|^1\Phi_0\rangle = c_0|1\bar{1}\rangle + \sum_{r=2}^K c_1^r|^1\Psi_1^r\rangle + \frac{1}{2} \sum_{r=2}^K \sum_{s=2}^K c_{11}^{rs} |^1\Psi_{11}^{rs}\rangle$$

where the singly and doubly excited spin adapted configurations are defined in Subsection 2.5.2.

- a. Show that $|^1\Phi_0\rangle$ can be cast into the form

$$|^1\Phi_0\rangle = \sum_{i=1}^K \sum_{j=1}^K C_{ij} |\psi_i \bar{\psi}_j\rangle$$

where \mathbf{C} is a symmetric $K \times K$ matrix.

- b. Show that

$$\gamma(\mathbf{x}_1, \mathbf{x}'_1) = \sum_{ij} (\mathbf{C}\mathbf{C}^\dagger)_{ij} (\psi_i(1)\psi_j^*(1') + \bar{\psi}_i(1)\bar{\psi}_j^*(1')).$$

- c. Let \mathbf{U} be the unitary transformation which diagonalizes \mathbf{C}

$$\mathbf{U}^\dagger \mathbf{C} \mathbf{U} = \mathbf{d}$$

where $(\mathbf{d})_{ij} = d_i \delta_{ij}$. Show that

$$\mathbf{U}^\dagger \mathbf{C} \mathbf{C}^\dagger \mathbf{U} = \mathbf{d}^2.$$

- d. Show that

$$\gamma(\mathbf{x}_1, \mathbf{x}'_1) = \sum_i d_i^2 (\zeta_i(1)\zeta_i^*(1') + \bar{\zeta}_i(1)\bar{\zeta}_i^*(1'))$$

where

$$\zeta_i = \sum_k \psi_k U_{ki}.$$

Thus \mathbf{U} diagonalizes the one-matrix, and hence ζ_i are natural spatial orbitals for the two-electron system.

- e. Finally, since \mathbf{C} is symmetric, \mathbf{U} can be chosen as real. Show that in terms of the natural spatial orbitals, $|^1\Phi_0\rangle$ given in part (a) can be rewritten as

$$|^1\Phi_0\rangle = \sum_{i=1}^K d_i |\zeta_i \bar{\zeta}_i\rangle$$

and note that this expansion contains only K terms.

Solution 4.8

- a. From (2.263) at the page 103 and Table 2.7 at the page 104, we know that

$$\begin{aligned} |^1\Psi_1^r\rangle &= \frac{1}{\sqrt{2}} (|\Psi_1^{\bar{r}}\rangle + |\Psi_1^r\rangle) = \frac{1}{\sqrt{2}} (|1\bar{r}\rangle + |r\bar{1}\rangle), \\ |^1\Psi_{11}^{rr}\rangle &= |\Psi_{11}^{r\bar{r}}\rangle = |r\bar{r}\rangle, \\ |^1\Psi_{11}^{rs}\rangle &= \frac{1}{\sqrt{2}} (|\Psi_{11}^{r\bar{s}}\rangle + |\Psi_{11}^{s\bar{r}}\rangle) = \frac{1}{\sqrt{2}} (|r\bar{s}\rangle + |s\bar{r}\rangle), \quad \forall r \neq s. \end{aligned}$$

Thus,

$$|^1\Phi_0\rangle = c_0|1\bar{1}\rangle + \sum_{r=2}^K \frac{c_1^r}{\sqrt{2}} (|1\bar{r}\rangle + |r\bar{1}\rangle) + \frac{1}{2} \sum_{r=2}^K c_{11}^{rr} |r\bar{r}\rangle + \sum_{r=2}^K \sum_{\substack{s=2 \\ r>s}}^K c_{11}^{rs} \frac{1}{\sqrt{2}} (|r\bar{s}\rangle + |s\bar{r}\rangle).$$

From this equation, we find that the coefficients are

$$\begin{aligned} C_{11} &= c_0, \\ C_{r1} &= C_{1r} = \frac{c_1^r}{\sqrt{2}}, \\ C_{rr} &= \frac{c_{11}^{rr}}{2}, \\ C_{rs} &= C_{sr} = \frac{c_{11}^{rs} + c_{11}^{sr}}{2}, \forall r \neq s, \end{aligned}$$

which equals

$$|^1\Phi_0\rangle = \sum_{i=1}^K \sum_{j=1}^K C_{ij} |\psi_i \bar{\psi}_j\rangle, \quad (4.8-1)$$

where

$$C_{ij} = C_{ji}, \forall i, j \in \{1, 2, \dots, K\}. \quad (4.8-2)$$

In other words, \mathbf{C} is a symmetric $K \times K$ matrix.

b. Note that there are two electrons in this system,

$$\begin{aligned} \gamma(\mathbf{x}_1, \mathbf{x}'_1) &= 2 \int_{\mathbb{R}^3} d\mathbf{x}_2 \Phi(\mathbf{x}_1, \mathbf{x}_2) \Phi^*(\mathbf{x}'_1, \mathbf{x}_2) \\ &= 2 \int_{\mathbb{R}^3} d\mathbf{x}_2 \frac{1}{\sqrt{2}} \sum_{i=1}^K \sum_{j=1}^K C_{ij} [\psi_i(\mathbf{x}_1) \bar{\psi}_j(\mathbf{x}_2) - \psi_i(\mathbf{x}_2) \bar{\psi}_j(\mathbf{x}_1)] \\ &\quad \times \frac{1}{\sqrt{2}} \sum_{k=1}^K \sum_{l=1}^K C_{kl}^* [\psi_k^*(\mathbf{x}'_1) \bar{\psi}_l^*(\mathbf{x}_2) - \psi_k^*(\mathbf{x}_2) \bar{\psi}_l^*(\mathbf{x}'_1)] \\ &= \sum_{i=1}^K \sum_{j=1}^K \sum_{k=1}^K \sum_{l=1}^K C_{ij} C_{kl}^* \\ &\quad \left[\psi_i(\mathbf{x}_1) \psi_k^*(\mathbf{x}'_1) \int_{\mathbb{R}^3} d\mathbf{x}_2 \bar{\psi}_j(\mathbf{x}_2) \bar{\psi}_l^*(\mathbf{x}_2) - \psi_i(\mathbf{x}_1) \bar{\psi}_l^*(\mathbf{x}'_1) \int_{\mathbb{R}^3} d\mathbf{x}_2 \bar{\psi}_j(\mathbf{x}_2) \psi_k^*(\mathbf{x}_2) \right. \\ &\quad \left. - \bar{\psi}_j(\mathbf{x}_1) \psi_k^*(\mathbf{x}'_1) \int_{\mathbb{R}^3} d\mathbf{x}_2 \psi_i(\mathbf{x}_2) \bar{\psi}_l^*(\mathbf{x}_2) + \bar{\psi}_j(\mathbf{x}_1) \bar{\psi}_l^*(\mathbf{x}'_1) \int_{\mathbb{R}^3} d\mathbf{x}_2 \psi_i(\mathbf{x}_2) \psi_k^*(\mathbf{x}_2) \right] \\ &= \sum_{i=1}^K \sum_{j=1}^K \sum_{k=1}^K \sum_{l=1}^K C_{ij} C_{kl}^* [\psi_i(\mathbf{x}_1) \psi_k^*(\mathbf{x}'_1) \delta_{jl} + \bar{\psi}_j(\mathbf{x}_1) \bar{\psi}_l^*(\mathbf{x}'_1) \delta_{ik}] \\ &= \sum_{i=1}^K \sum_{j=1}^K \sum_{k=1}^K C_{ij} C_{kj}^* \psi_i(\mathbf{x}_1) \psi_k^*(\mathbf{x}'_1) + \sum_{i=1}^K \sum_{j=1}^K \sum_{l=1}^K C_{ij} C_{il}^* \bar{\psi}_j(\mathbf{x}_1) \bar{\psi}_l^*(\mathbf{x}'_1) \\ &= \sum_{i=1}^K \sum_{j=1}^K \sum_{k=1}^K C_{ij} C_{jk}^\dagger \psi_i(\mathbf{x}_1) \psi_k^*(\mathbf{x}'_1) + \sum_{i=1}^K \sum_{j=1}^K \sum_{k=1}^K C_{ji} C_{ik}^\dagger \bar{\psi}_j(\mathbf{x}_1) \bar{\psi}_k^*(\mathbf{x}'_1) \\ &= \sum_{i=1}^K \sum_{j=1}^K \psi_i(\mathbf{x}_1) \psi_j^*(\mathbf{x}'_1) \left(\sum_{k=1}^K C_{ik} C_{kj}^\dagger \right) + \sum_{i=1}^K \sum_{j=1}^K \bar{\psi}_i(\mathbf{x}_1) \bar{\psi}_j^*(\mathbf{x}'_1) \left(\sum_{k=1}^K C_{ik} C_{kj}^\dagger \right) \\ &= \sum_{i=1}^K \sum_{j=1}^K (\mathbf{C} \mathbf{C}^\dagger)_{ij} (\psi_i(\mathbf{x}_1) \psi_j^*(\mathbf{x}'_1) + \bar{\psi}_i(\mathbf{x}_1) \bar{\psi}_j^*(\mathbf{x}'_1)). \end{aligned}$$

Thus, we have proved that

$$\gamma(\mathbf{x}_1, \mathbf{x}'_1) = \sum_{i=1}^K \sum_{j=1}^K (\mathbf{C} \mathbf{C}^\dagger)_{ij} (\psi_i(\mathbf{x}_1) \psi_j^*(\mathbf{x}'_1) + \bar{\psi}_i(\mathbf{x}_1) \bar{\psi}_j^*(\mathbf{x}'_1)). \quad (4.8-3)$$

c. In fact, all d_i are real as all eigenvalues of a real symmetric matrix are real, thus $\mathbf{d}^\dagger = \mathbf{d}$, and

$$\mathbf{d}^2 = \mathbf{d} \mathbf{d}^\dagger = \mathbf{U}^\dagger \mathbf{C} \mathbf{U} (\mathbf{U}^\dagger \mathbf{C} \mathbf{U})^\dagger = \mathbf{U}^\dagger \mathbf{C} \mathbf{U} \mathbf{U}^\dagger \mathbf{C}^\dagger \mathbf{U} = \mathbf{U}^\dagger \mathbf{C} \mathbf{C}^\dagger \mathbf{U}. \quad (4.8-4)$$

d. From the former issue, we know

$$\mathbf{C}\mathbf{C}^\dagger = \mathbf{U}\mathbf{d}^2\mathbf{U}^\dagger.$$

Thus, we find that

$$\begin{aligned}\gamma(\mathbf{x}_1, \mathbf{x}'_1) &= \sum_{i=1}^K \sum_{j=1}^K (\mathbf{C}\mathbf{C}^\dagger)_{ij} (\psi_i(\mathbf{x}_1)\psi_j^*(\mathbf{x}'_1) + \bar{\psi}_i(\mathbf{x}_1)\bar{\psi}_j^*(\mathbf{x}'_1)) \\ &= \sum_{i=1}^K \sum_{j=1}^K (\mathbf{U}\mathbf{d}^2\mathbf{U}^\dagger)_{ij} (\psi_i(\mathbf{x}_1)\psi_j^*(\mathbf{x}'_1) + \bar{\psi}_i(\mathbf{x}_1)\bar{\psi}_j^*(\mathbf{x}'_1)) \\ &= \sum_{i=1}^K \sum_{j=1}^K \sum_{k=1}^K \mathbf{U}_{ik}d_k^2\mathbf{U}_{kj}^\dagger (\psi_i(\mathbf{x}_1)\psi_j^*(\mathbf{x}'_1) + \bar{\psi}_i(\mathbf{x}_1)\bar{\psi}_j^*(\mathbf{x}'_1)) \\ &= \sum_{k=1}^K d_k^2 \left(\sum_{i=1}^K \psi_i(\mathbf{x}_1)\mathbf{U}_{ik} \sum_{j=1}^K \psi_j^*(\mathbf{x}'_1)\mathbf{U}_{jk}^* + \sum_{i=1}^K \bar{\psi}_i(\mathbf{x}_1)\mathbf{U}_{ik} \sum_{j=1}^K \bar{\psi}_j^*(\mathbf{x}'_1)\mathbf{U}_{jk}^* \right).\end{aligned}$$

Therefore, we define

$$\zeta_i = \sum_{k=1}^K \psi_k U_{ki},$$

and we obtain

$$\gamma(\mathbf{x}_1, \mathbf{x}'_1) = \sum_{k=1}^K d_k^2 (\zeta_k(\mathbf{x}_1)\zeta_k^*(\mathbf{x}'_1) + \bar{\zeta}_k(\mathbf{x}_1)\bar{\zeta}_k^*(\mathbf{x}'_1)) = \sum_{i=1}^K d_i^2 (\zeta_i(\mathbf{x}_1)\zeta_i^*(\mathbf{x}'_1) + \bar{\zeta}_i(\mathbf{x}_1)\bar{\zeta}_i^*(\mathbf{x}'_1)). \quad (4.8-5)$$

We conclude that U diagonalizes the one-matrix, and hence ζ_i are natural spatial orbitals for the two-electron system.

e. Now we convert \mathbf{C} firstly,

$$\mathbf{C} = \mathbf{U}\mathbf{d}\mathbf{U}^\dagger \Leftrightarrow C_{ij} = \sum_{k=1}^K \mathbf{U}_{ik}d_k\mathbf{U}_{kj}^\dagger = \sum_{k=1}^K \mathbf{U}_{ik}d_k\mathbf{U}_{jk}^*$$

Thus we arrive at

$$|^1\Phi_0\rangle = \sum_{i=1}^K \sum_{j=1}^K C_{ij} |\psi_i\bar{\psi}_j\rangle = \sum_{i=1}^K \sum_{j=1}^K \sum_{k=1}^K \mathbf{U}_{ik}d_k\mathbf{U}_{jk}^* |\psi_i\bar{\psi}_j\rangle = \sum_{k=1}^K d_k |\zeta_k\bar{\zeta}_k\rangle = \sum_{i=1}^K d_i |\zeta_i\bar{\zeta}_i\rangle. \quad (4.8-6)$$

We find that this expansion contains only K terms.

4.5 The Multiconfiguration Self-Consistent Field (MCSCF) and Generalized Valence Bond (GVB) Methods

Exercise 4.9

Consider the transformation

$$\begin{aligned}u &= \frac{1}{\sqrt{a^2+b^2}} (a\psi_A + b\psi_B), \\ v &= \frac{1}{\sqrt{a^2+b^2}} (a\psi_A - b\psi_B).\end{aligned}$$

a. Show that

$$\langle u|u\rangle = \langle v|v\rangle = 1$$

and

$$\langle u|v\rangle \equiv S = \frac{a^2 - b^2}{a^2 + b^2}.$$

b. Show that $|\Psi_{\text{GVB}}\rangle$ in Eq.(4.52) can be rewritten as

$$|\Psi_{\text{GVB}}\rangle = \frac{1}{\sqrt{a^4 + b^4}} [a^2 \psi_A(1) \psi_A(2) - b^2 \psi_B(1) \psi_B(2)] \frac{1}{\sqrt{2}} (\alpha(1)\beta(2) - \alpha(2)\beta(1))$$

and conclude that this is identical to $|\Psi_{\text{MCSCF}}\rangle$ in Eq.(4.48) if

$$c_A = \frac{a^2}{\sqrt{a^4 + b^4}},$$

$$c_B = -\frac{b^2}{\sqrt{a^4 + b^4}}.$$

Solution 4.9

a. Check $\langle u|u\rangle$, $\langle u|v\rangle$ and $\langle v|v\rangle$. Note that

$$\langle \psi_A | \psi_A \rangle = \langle \psi_B | \psi_B \rangle = 1, \quad \langle \psi_A | \psi_B \rangle = \langle \psi_B | \psi_A \rangle = 0.$$

My results are as follows.

$$\begin{aligned} \langle u|u \rangle &= \frac{1}{a^2 + b^2} [a^2 \langle \psi_A | \psi_A \rangle + ab \langle \psi_A | \psi_B \rangle + ab \langle \psi_B | \psi_A \rangle + b^2 \langle \psi_B | \psi_B \rangle] = 1, \\ \langle v|v \rangle &= \frac{1}{a^2 + b^2} [a^2 \langle \psi_A | \psi_A \rangle - ab \langle \psi_A | \psi_B \rangle - ab \langle \psi_B | \psi_A \rangle + b^2 \langle \psi_B | \psi_B \rangle] = 1, \\ S \equiv \langle u|v \rangle &= \frac{1}{a^2 + b^2} [a^2 \langle \psi_A | \psi_A \rangle - ab \langle \psi_A | \psi_B \rangle + ab \langle \psi_B | \psi_A \rangle - b^2 \langle \psi_B | \psi_B \rangle] = \frac{a^2 - b^2}{a^2 + b^2}. \end{aligned}$$

b. Substitute these results into (4.52), we find that

$$\begin{aligned} |\Psi_{\text{GVB}}\rangle &= \frac{1}{\sqrt{2 \left[1 + \left(\frac{a^2 - b^2}{a^2 + b^2} \right)^2 \right]}} \left[\frac{1}{a^2 + b^2} (a\psi_A(1) + b\psi_B(1))(a\psi_A(2) - b\psi_B(2)) \right. \\ &\quad \left. + \frac{1}{a^2 + b^2} (a\psi_A(2) + b\psi_B(2))(a\psi_A(1) - b\psi_B(1)) \right] \frac{1}{\sqrt{2}} [\alpha(1)\beta(2) - \alpha(2)\beta(1)] \\ &= \frac{a^2 + b^2}{\sqrt{2 [(a^2 + b^2)^2 + (a^2 - b^2)^2]}} \times \frac{1}{\sqrt{2} (a^2 + b^2)} [\alpha(1)\beta(2) - \alpha(2)\beta(1)] \\ &\quad [(a\psi_A(1) + b\psi_B(1))(a\psi_A(2) - b\psi_B(2)) + (a\psi_A(2) + b\psi_B(2))(a\psi_A(1) - b\psi_B(1))] \\ &= \frac{1}{2\sqrt{2a^4 + 2b^4}} [2a^2 \psi_A(1) \psi_A(2) - 2b^2 \psi_B(1) \psi_B(2)] [\alpha(1)\beta(2) - \alpha(2)\beta(1)] \\ &= \frac{1}{\sqrt{a^4 + b^4}} [a^2 \psi_A(1) \psi_A(2) - b^2 \psi_B(1) \psi_B(2)] \frac{1}{\sqrt{2}} [\alpha(1)\beta(2) - \alpha(2)\beta(1)]. \end{aligned}$$

Thus, we conclude that this is identical to $|\Psi_{\text{MCSCF}}\rangle$ in Eq.(4.48) if

$$c_A = \frac{a^2}{\sqrt{a^4 + b^4}}, \quad c_B = -\frac{b^2}{\sqrt{a^4 + b^4}}. \quad (4.9-1)$$

4.6 Truncated CI and the Size-Consistency Problem

Exercise 4.10

Show that $|1_1 \bar{1}_1 2_1 \bar{2}_1\rangle$ has a zero matrix element with any of the configurations in Eq.(4.55).

Solution 4.10

We only check whether $\langle \Psi_0 | \mathcal{H} | 1_1 \bar{1}_1 2_1 \bar{2}_1 \rangle$, $\langle \Psi_{1_1 \bar{1}_1}^{2_1 \bar{2}_1} | \mathcal{H} | 1_1 \bar{1}_1 2_1 \bar{2}_1 \rangle$ and $\langle \Psi_{1_2 \bar{1}_2}^{2_2 \bar{2}_2} | \mathcal{H} | 1_1 \bar{1}_1 2_1 \bar{2}_1 \rangle$ vanish. That is

enough.

$$\begin{aligned}\langle \Psi_0 | \mathcal{H} | 1_1 \bar{1}_1 2_1 \bar{2}_1 \rangle &= \langle 1_1 \bar{1}_1 1_2 \bar{1}_2 | \mathcal{H} | 1_1 \bar{1}_1 2_1 \bar{2}_1 \rangle = \langle 1_2 \bar{1}_2 | 2_1 \bar{2}_1 \rangle = (1_2 2_1 | 1_2 2_1) = 0, \\ \langle \Psi_{1_1 \bar{1}_1}^{2_1 \bar{2}_1} | \mathcal{H} | 1_1 \bar{1}_1 2_1 \bar{2}_1 \rangle &= \langle 2_1 \bar{2}_1 1_2 \bar{1}_2 | \mathcal{H} | 1_1 \bar{1}_1 2_1 \bar{2}_1 \rangle = 0, \\ \langle \Psi_{1_2 \bar{1}_2}^{2_2 \bar{2}_2} | \mathcal{H} | 1_1 \bar{1}_1 2_1 \bar{2}_1 \rangle &= \langle 1_1 \bar{1}_1 2_2 \bar{2}_2 | \mathcal{H} | 1_1 \bar{1}_1 2_1 \bar{2}_1 \rangle = \langle 2_2 \bar{2}_2 | 2_1 \bar{2}_1 \rangle = (2_2 2_1 | 2_2 2_1) = 0.\end{aligned}$$

Exercise 4.11

Use the integrals for STO-3G H₂ at $R = 1.4$ a.u., given in Appendix D, to calculate ${}^N E_{\text{corr}}(\text{DCI})/N$ for $N = 1, 10$, and 100 .

Solution 4.11

When $R = 1.4$ a.u., we know that

$$\begin{aligned}\varepsilon_1 &= -0.5782 \text{ a.u.}, & \varepsilon_2 &= 0.6703 \text{ a.u.}, & J_{11} &= 0.6746 \text{ a.u.}, \\ J_{12} &= 0.6636 \text{ a.u.}, & J_{22} &= 0.6975 \text{ a.u.}, & K_{12} &= 0.1813 \text{ a.u.}\end{aligned}$$

Similar to Exercise 4.3, we obtain $\Delta = 0.78865$ a.u. Thus, with (4.67), we calculate $\frac{{}^N E_{\text{corr}}(\text{DCI})}{N}$ at various N . These results are listed in Table 4.1. We conclude that as N increases, $\frac{{}^N E_{\text{corr}}(\text{DCI})}{N}$ vanishes.

Table 4.1: The table of $\frac{{}^N E_{\text{corr}}(\text{DCI})}{N}$ to N . Here $\Delta = 0.78865$ a.u.

N	1	10	100	1000	10000
$\frac{{}^N E_{\text{corr}}(\text{DCI})}{N}$	-0.02057	-0.01864	-0.01188	-0.00500	-0.00174

Exercise 4.12

Show that full CI is size consistent for a dimer of non-interacting minimal basis H₂ molecules. A full CI calculation includes, in addition to the excitations in Eq.(4.55), the quadruply excited state $|2_1 \bar{2}_1 2_2 \bar{2}_2\rangle = |\Psi_{1_1 \bar{1}_1 1_2 \bar{1}_2}^{2_1 \bar{2}_1 2_2 \bar{2}_2}\rangle$,

$$|\Phi_0\rangle = |\Psi_0\rangle + c_1 |2_1 \bar{2}_1 1_2 \bar{1}_2\rangle + c_2 |1_1 \bar{1}_1 2_2 \bar{2}_2\rangle + c_3 |2_1 \bar{2}_1 2_2 \bar{2}_2\rangle.$$

a. Show that the full CI matrix equation is

$$\begin{pmatrix} 0 & K_{12} & K_{12} & 0 \\ K_{12} & 2\Delta & 0 & K_{12} \\ K_{12} & 0 & 2\Delta & K_{12} \\ 0 & K_{12} & K_{12} & 4\Delta \end{pmatrix} \begin{pmatrix} 1 \\ c_1 \\ c_2 \\ c_3 \end{pmatrix} = {}^2 E_{\text{corr}} \begin{pmatrix} 1 \\ c_1 \\ c_2 \\ c_3 \end{pmatrix}.$$

Go directly to part (e). If you need help return to part (b).

b. Show that $c_1 = c_2$ and hence ${}^2 E_{\text{corr}} = 2K_{12}c_1$.

c. Show that

$$c_3 = \frac{{}^2 E_{\text{corr}}}{{}^2 E_{\text{corr}} - 4\Delta}.$$

d. Show that

$$c_1 = \frac{2K_{12}}{{}^2 E_{\text{corr}} - 4\Delta}.$$

e. Finally, show that

$${}^2 E_{\text{corr}} = 2 \left(\Delta - \sqrt{\Delta^2 + K_{12}^2} \right)$$

which is indeed exact for the model.

It is interesting to note that we can express the coefficient of the quadruple excitation (c_3) as the product of the coefficients of the double excitations:

$$c_3 = \frac{{}^2E_{\text{corr}}}{{}^2E_{\text{corr}} - 4\Delta} = \frac{2K_{12}c_1}{{}^2E_{\text{corr}} - 4\Delta} = (c_1)^2$$

where we have used results in parts (b), (c), and (d). This result is not true in general but is a consequence of the fact that the two monomers are independent. However, it suggests that it might be reasonable to *approximate* the coefficient of a quadruply excited configuration as a product of the coefficients of the double excitations that combine to give the quadruply excited configuration. This idea plays a central role in the next chapter.

Solution 4.12

a. Similar to the instance shown at the page 263, we obtain that

$$\begin{aligned} \langle \Psi_0 | \mathcal{H} - E_0 | \Psi_0 \rangle &= 0, \\ \langle \Psi_0 | \mathcal{H} - E_0 | 2_1 \bar{2}_1 1_2 \bar{1}_2 \rangle &= \langle 2_1 \bar{2}_1 1_2 \bar{1}_2 | \mathcal{H} - E_0 | \Psi_0 \rangle = \langle 1_1 \bar{1}_1 | 2_1 \bar{2}_1 \rangle = (1_1 2_1 | 1_1 2_1) = K_{12}, \\ \langle \Psi_0 | \mathcal{H} - E_0 | 1_1 \bar{1}_1 2_2 \bar{2}_2 \rangle &= \langle 1_1 \bar{1}_1 2_2 \bar{2}_2 | \mathcal{H} - E_0 | \Psi_0 \rangle = \langle 1_2 \bar{1}_2 | 2_2 \bar{2}_2 \rangle = (1_2 2_2 | 1_2 2_2) = K_{12}, \\ \langle \Psi_0 | \mathcal{H} - E_0 | 2_1 \bar{2}_1 2_2 \bar{2}_2 \rangle &= \langle 2_1 \bar{2}_1 2_2 \bar{2}_2 | \mathcal{H} - E_0 | \Psi_0 \rangle = \langle 1_1 \bar{1}_1 1_2 \bar{1}_2 | \mathcal{H} | 2_1 \bar{2}_1 2_2 \bar{2}_2 \rangle = 0, \\ \langle 2_1 \bar{2}_1 1_2 \bar{1}_2 | \mathcal{H} - E_0 | 2_1 \bar{2}_1 1_2 \bar{1}_2 \rangle &= (2h_{11} + 2h_{22} + J_{11} + J_{12}) - (4h_{11} + 2J_{11}) \\ &= 2(\varepsilon_1 + \varepsilon_2) - J_{11} + J_{22} - 4J_{12} + 2K_{12} - 2(2\varepsilon_1 - J_{11}) \\ &= 2(\varepsilon_2 - \varepsilon_1) + J_{11} + J_{22} + 2K_{12} - 4J_{12} = 2\Delta, \\ \langle 2_1 \bar{2}_1 1_2 \bar{1}_2 | \mathcal{H} - E_0 | 1_1 \bar{1}_1 2_2 \bar{2}_2 \rangle &= \langle 1_1 \bar{1}_1 2_2 \bar{2}_2 | \mathcal{H} - E_0 | 2_1 \bar{2}_1 1_2 \bar{1}_2 \rangle = \langle 1_1 \bar{1}_1 2_2 \bar{2}_2 | \mathcal{H} | 2_1 \bar{2}_1 1_2 \bar{1}_2 \rangle = 0, \\ \langle 2_1 \bar{2}_1 1_2 \bar{1}_2 | \mathcal{H} - E_0 | 2_1 \bar{2}_1 2_2 \bar{2}_2 \rangle &= \langle 2_1 \bar{2}_1 2_2 \bar{2}_2 | \mathcal{H} - E_0 | 2_1 \bar{2}_1 1_2 \bar{1}_2 \rangle = \langle 2_1 \bar{2}_1 2_2 \bar{2}_2 | \mathcal{H} | 2_1 \bar{2}_1 1_2 \bar{1}_2 \rangle \\ &= \langle 2_2 \bar{2}_2 | 1_2 \bar{1}_2 \rangle = (2_2 1_2 | 2_2 1_2) = K_{12}, \\ \langle 1_1 \bar{1}_1 2_2 \bar{2}_2 | \mathcal{H} - E_0 | 1_1 \bar{1}_1 2_2 \bar{2}_2 \rangle &= (2h_{11} + 2h_{22} + J_{11} + J_{22}) - (4h_{11} + 2J_{11}) \\ &= 2(\varepsilon_1 + \varepsilon_2) - J_{11} + J_{22} - 4J_{12} + 2K_{12} - 2(2\varepsilon_1 - J_{11}) \\ &= 2(\varepsilon_2 - \varepsilon_1) + J_{11} + J_{22} + 2K_{12} - 4J_{12} = 2\Delta, \\ \langle 1_1 \bar{1}_1 2_2 \bar{2}_2 | \mathcal{H} - E_0 | 2_1 \bar{2}_1 2_2 \bar{2}_2 \rangle &= \langle 2_1 \bar{2}_1 2_2 \bar{2}_2 | \mathcal{H} - E_0 | 1_1 \bar{1}_1 2_2 \bar{2}_2 \rangle = \langle 1_1 \bar{1}_1 2_2 \bar{2}_2 | \mathcal{H} | 2_1 \bar{2}_1 2_2 \bar{2}_2 \rangle \\ &= \langle 1_1 \bar{1}_1 | 2_1 \bar{2}_1 \rangle = (1_1 2_1 | 1_1 2_1) = K_{12}, \\ \langle 2_1 \bar{2}_1 2_2 \bar{2}_2 | \mathcal{H} - E_0 | 2_1 \bar{2}_1 2_2 \bar{2}_2 \rangle &= (4h_{22} + 2J_{22}) - (4h_{11} + 2J_{11}) \\ &= (4\varepsilon_2 - 8J_{12} + 4K_{12} + 2J_{12}) - (4\varepsilon_1 - 2J_{11}) \\ &= 4(\varepsilon_2 - \varepsilon_1) + 2J_{11} + 2J_{12} + 4K_{12} - 8J_{12} = 4\Delta. \end{aligned}$$

Here, note that the definition of Δ is located on page 240. Thus, we obtain the full CI matrix equation

$$\begin{pmatrix} 0 & K_{12} & K_{12} & 0 \\ K_{12} & 2\Delta & 0 & K_{12} \\ K_{12} & 0 & 2\Delta & K_{12} \\ 0 & K_{12} & K_{12} & 4\Delta \end{pmatrix} \begin{pmatrix} 1 \\ c_1 \\ c_2 \\ c_3 \end{pmatrix} = {}^2E_{\text{corr}} \begin{pmatrix} 1 \\ c_1 \\ c_2 \\ c_3 \end{pmatrix}. \quad (4.12-1)$$

It also equals

$$\begin{aligned} K_{12}c_1 + K_{12}c_2 &= {}^2E_{\text{corr}}, \\ K_{12} + 2\Delta c_1 + K_{12}c_3 &= {}^2E_{\text{corr}}c_1, \\ K_{12} + 2\Delta c_2 + K_{12}c_3 &= {}^2E_{\text{corr}}c_2, \\ K_{12}c_1 + K_{12}c_2 + 4\Delta c_3 &= {}^2E_{\text{corr}}c_3. \end{aligned}$$

b. From the second and the third equations, it is evident that $c_1 = c_2$. From the first equation,

$${}^2E_{\text{corr}} = K_{12}c_1 + K_{12}c_2 = 2K_{12}c_1. \quad (4.12-2)$$

c. The fourth equation can be subtracted by the first one, viz.,

$$4\Delta c_3 = {}^2E_{\text{corr}}c_3 - {}^2E_{\text{corr}},$$

which is equal to

$$c_3 = \frac{{}^2E_{\text{corr}}}{2E_{\text{corr}} - 4\Delta}. \quad (4.12-3)$$

d. By substituting c_3 into the second equation, we find that

$$\begin{aligned} c_1 &= \frac{K_{12} + K_{12}c_3}{2E_{\text{corr}} - 2\Delta} = \frac{K_{12}}{2E_{\text{corr}} - 2\Delta} \left(1 + \frac{{}^2E_{\text{corr}}}{2E_{\text{corr}} - 4\Delta} \right) \\ &= \frac{K_{12}(2E_{\text{corr}} - 4\Delta)}{(2E_{\text{corr}} - 2\Delta)(2E_{\text{corr}} - 4\Delta)} = \frac{2K_{12}}{2E_{\text{corr}} - 4\Delta}. \end{aligned}$$

e. Thus, by substituting c_1 into the first equation, we find that

$$2E_{\text{corr}} = 2K_{12}c_1 = \frac{4K_{12}^2}{2E_{\text{corr}} - 4\Delta} \Rightarrow (2E_{\text{corr}})^2 - 4\Delta(2E_{\text{corr}}) - 4K_{12}^2 = 0.$$

The discriminant Δ_E of this quadratic equation is

$$\Delta_E = (4\Delta)^2 - 4 \times 1 \times (-4K_{12}^2) = 16(\Delta^2 + K_{12}^2) > 0$$

Thus, the root are

$$E_1 = 2 \left(\Delta + \sqrt{\Delta^2 + K_{12}^2} \right), \quad E_2 = 2 \left(\Delta - \sqrt{\Delta^2 + K_{12}^2} \right).$$

In current problem, the correlation energy is the lowest root,

$${}^2E_{\text{corr}} = 2 \left(\Delta - \sqrt{\Delta^2 + K_{12}^2} \right), \quad (4.12-4)$$

which is indeed exact for the model.

Exercise 4.13

Consider the exact basis set correlation energy of minimal basis H_2 given in Eq.(4.23). Assuming that $K_{12}^2/\Delta^2 \ll 1$ show that

$${}^1E_{\text{corr}}(\text{exact}) \approx -\frac{K_{12}^2}{2\Delta}.$$

Hint: $(1+x)^{1/2} \approx 1 + \frac{1}{2}x$ when $x \ll 1$. This approximate result is the same as the simplest expression for the correlation energy obtained via a form of perturbation theory. Show that by expanding ${}^N E_{\text{corr}}(\text{DCI})$ in the same way, assuming that $NK_{12}^2/\Delta^2 \ll 1$, one obtains simply N times the above result. This approximation is equivalent to a perturbation result for the correlation energy of a supermolecule, and then form of the result is a reflection of the fact that, in contrast to truncated CI, perturbation theory is size consistent (see Chapter 6).

Solution 4.13

With $(1+x)^{\frac{1}{2}} \approx 1 + \frac{1}{2}x$ as $x \ll 1$, when $K_{12}^2/\Delta^2 \ll 1$, we find that

$${}^1E_{\text{corr}}(\text{exact}) = \Delta \left(1 - \sqrt{1 + \frac{K_{12}^2}{\Delta^2}} \right) \approx \Delta \left(1 - \left(1 + \frac{K_{12}^2}{2\Delta^2} \right) \right) = -\frac{K_{12}^2}{2\Delta}. \quad (4.13-1)$$

The conclusion has been proved.

Exercise 4.14

DCI calculations have become relatively routine; therefore, it is of interest to ask whether one can correct the DCI correlation energy so that it becomes approximately size consistent. A simple prescription for doing this, which is approximately valid *only* for relatively small systems, is to write

$$E_{\text{corr}} = E_{\text{corr}}(\text{DCI}) + \Delta E_{\text{Davidson}}$$

where the Davidson correction is given by

$$\Delta E_{\text{Davidson}} = (1 - c_0^2) E_{\text{corr}}(\text{DCI})$$

where c_0 is the coefficient of the Hartree-Fock wave function in the *normalized* DCI wave function. The Davidson correction can be computed without additional labor since c_0 is available in a DCI calculation. Moreover, there is numerical evidence that it leads to an improvement over DCI for relatively small molecules. For example, for H_2O , using the 39-STO basis and the Davidson correction, $\theta_e = 104.6^\circ$, $R_e = 1.809$ a.u., $f_{RR} = 8.54$, and $f_{\theta\theta} = 0.80$ (see Table 4.7). Also, the ionization potentials of N_2 , using the same basis as in Table 4.9 and the Davidson correction, are 0.575 a.u. ($3\sigma_g$) and 0.617 a.u. ($1\pi_u$). The purpose of this exercise is to explore the nature of the Davidson correction.

- a. For the model of N independent minimal basis H_2 molecules, assume that N is large, yet small enough that NK_{12}^2/Δ^2 is still less than unity. In addition, remember that $\Delta \gg K_{12}$. Using the identity $(1+x)^{1/2} \approx 1 + \frac{1}{2}x - \frac{1}{8}x^2 + \dots$ show that

$${}^N E_{\text{corr}}(\text{DCI}) = -\frac{NK_{12}^2}{2\Delta} + \frac{N^2 K_{12}^4}{8\Delta^3} + \dots$$

The term proportional to N^2 is spurious and is not present in the similar expansion of ${}^N E_{\text{corr}}(\text{exact})$.

- b. Show that

$$1 - c_0^2 = \frac{Nc_1^2}{1 + Nc_1^2}.$$

- c. Show that

$$c_1 = -\frac{K_{12}}{2\Delta} + \dots$$

- d. Show that

$$\Delta E_{\text{Davidson}} = -\frac{N^2 K_{12}^4}{8\Delta^3} + \dots$$

Finally, note that the Davidson correction exactly cancels the term proportional to N^2 in the expansion of ${}^N E_{\text{corr}}(\text{DCI})$. However, spurious terms containing higher powers of N still remain. For large N , the whole analysis breaks down since NK_{12}^2/Δ^2 eventually becomes greater than unity. For $N = 1$, $E_{\text{corr}}(\text{DCI})$ is exact within the model, yet $\Delta E_{\text{Davidson}}$ is not zero.

- e. Numerically investigate the range of validity of the Davidson correction for N independent H_2 molecules. Calculate ${}^N E_{\text{corr}}(\text{DCI})/{}^N E_{\text{corr}}(\text{exact})$ and $({}^N E_{\text{corr}}(\text{DCI}) + \Delta E_{\text{Davidson}})/{}^N E_{\text{corr}}(\text{exact})$ for $N = 1, \dots, 20$ using the values of the two-electron integrals for $R = 1.4$ a.u. given in Appendix D. You will find that the correlation energy calculated using the Davidson correction is within 1% of the exact value for $3 < N < 11$. The DCI correlation energy, on the other hand, errs by 2.5% for $N = 3$ and by 10% for $N = 11$. For $N = 100$, the correlation energy with and without the Davidson correction errs by 25% and 42%, respectively. *Hint:* show that

$$\Delta E_{\text{Davidson}} = \frac{({}^N E_{\text{corr}}(\text{DCI}))^3}{NK_{12}^2 + ({}^N E_{\text{corr}}(\text{DCI}))^2}.$$

- f. Calculate the Davidson-corrected correlation energy of H_2O using $E_{\text{corr}}(\text{DCI})$ and c_0 given by Saxe et al. (see Further Reading). Compare your result with $E_{\text{corr}}(\text{DQCI})$ and the exact basis set correlation energy.

Solution 4.14

- a. From (4.67), we find that

$$\frac{{}^N E_{\text{corr}}(\text{DCI})}{\Delta} = 1 - \sqrt{1 + \frac{NK_{12}^2}{\Delta^2}} = 1 - \left(1 + \frac{NK_{12}^2}{2\Delta^2} - \frac{N^2 K_{12}^4}{8\Delta^4} + \dots\right) = -\frac{NK_{12}^2}{2\Delta^2} + \frac{N^2 K_{12}^4}{8\Delta^4} + \dots$$

It equals

$${}^N E_{\text{corr}}(\text{DCI}) = -\frac{NK_{12}^2}{2\Delta} + \frac{N^2 K_{12}^4}{8\Delta^3} + \dots \quad (4.14-1)$$

b. From (4.55), we know the norm of $|\Phi_0\rangle$ of the two independent minimal basis, viz.,

$$\langle \Phi_0 | \Phi_0 \rangle = \left(\langle \Psi_0 | + c_1^* \langle \Psi_{1_1 \bar{1}_1}^{2_1 \bar{2}_1} | + c_2^* \langle \Psi_{1_2 \bar{1}_2}^{2_2 \bar{2}_2} | \right) \left(|\Psi_0\rangle + c_1 |\Psi_{1_1 \bar{1}_1}^{2_1 \bar{2}_1}\rangle + c_2 |\Psi_{1_2 \bar{1}_2}^{2_2 \bar{2}_2}\rangle \right) = 1 + c_1^2 + c_2^2 = 1 + 2c_1^2.$$

Similarly, we can derive that the norm of $|\Phi_0\rangle$ of N independent minimal basis, viz.,

$$\langle \Phi_0 | \Phi_0 \rangle = 1 + Nc_1^2.$$

Therefore, we can obtain the coefficient of $|\Psi_0\rangle$,

$$c_0 = \frac{1}{\sqrt{\langle \Phi_0 | \Phi_0 \rangle}} = \frac{1}{\sqrt{1 + Nc_1^2}}$$

and

$$1 - c_0^2 = 1 - \left(\frac{1}{\sqrt{1 + Nc_1^2}} \right)^2 = \frac{Nc_1^2}{1 + Nc_1^2}. \quad (4.14-2)$$

c. From (4.64) and (4.67), we obtain

$$c_1 = \frac{K_{12}}{{}^N E_{\text{corr}}(\text{DCI}) - 2\Delta} = \frac{K_{12}}{\Delta - \sqrt{\Delta^2 + NK_{12}^2} - 2\Delta} = -\frac{\frac{K_{12}}{\Delta}}{1 + \sqrt{1 + N\left(\frac{K_{12}}{\Delta}\right)^2}} \approx -\frac{K_{12}}{2\Delta}, \quad (4.14-3)$$

where $N\left(\frac{K_{12}}{\Delta}\right)^2 \approx 0$. Note that as $N\left(\frac{K_{12}}{\Delta}\right)^2 \approx 0$,

$$1 + Nc_1^2 \approx 1 + N\left(-\frac{K_{12}}{2\Delta}\right)^2 \approx 1.$$

d. From (4.65), we find that

$$\Delta E_{\text{Davidson}} = \frac{Nc_1^2}{1 + Nc_1^2} NK_{12}c_1 \approx N^2 K_{12}c_1^3 \approx N^2 K_{12} \left(-\frac{K_{12}}{2\Delta}\right)^3 = -\frac{N^2 K_{12}^4}{8\Delta^3}.$$

Higher terms are omitted. In other words,

$$\Delta E_{\text{Davidson}} = -\frac{N^2 K_{12}^4}{8\Delta^3} + \dots \quad (4.14-4)$$

e. Substitute (4.64) into the equation of $\Delta E_{\text{Davidson}}$, we find that

$$\begin{aligned} \Delta E_{\text{Davidson}} &= \frac{Nc_1^2}{1 + Nc_1^2} {}^N E_{\text{corr}}(\text{DCI}) \\ &= \frac{N\left(\frac{{}^N E_{\text{corr}}(\text{DCI})}{NK_{12}}\right)^2}{{}^N E_{\text{corr}}(\text{DCI})^2} {}^N E_{\text{corr}}(\text{DCI}) = \frac{({}^N E_{\text{corr}}(\text{DCI}))^3}{NK_{12}^2 + ({}^N E_{\text{corr}}(\text{DCI}))^2}. \end{aligned}$$

A program named `14.cc` in the directory `./scripts/4.14` can be found. It is designed to calculate the correlation energy for $N = 1, 2, \dots, 19, 20, 100$. Relevant results are listed in Table 4.2.

f. The paper by Saxe et al. can be found in the directory “references”. From this paper, we know

$$\begin{aligned} c_0(\text{DCI}) &= 0.97938, \\ {}^N E_{\text{corr}}(\text{DCI}) &= -0.139340 \text{ a.u.} \\ {}^N E_{\text{corr}}(\text{DQCI}) &= -0.145859 \text{ a.u.} \\ {}^N E_{\text{corr}}(\text{FCI}) &= -0.148028 \text{ a.u.} \end{aligned}$$

Thus,

$$\Delta E_{\text{Davidson}} = (1 - c_0^2(\text{DCI}))^N E_{\text{corr}}(\text{DCI}) = -0.005687 \text{ a.u.}$$

and

$$^N E_{\text{corr}}(\text{DCI}) + \Delta E_{\text{Davidson}} = -0.145027 \text{ a.u.}$$

Compared to the exact basis set correlation energy $^N E_{\text{corr}}(\text{FCI}) = -0.148028 \text{ a.u.}$, the error of DCI with the Davidson correction is about 2.03% while the error of QDCI is about 1.47%.

Table 4.2: The table of the correlation energy to N .

N	$^N E_{\text{corr}}(\text{DCI})$	error	$^N E_{\text{corr}}(\text{DCI}) + \Delta E_{\text{Davidson}}$	error	$^N E_{\text{corr}}(\text{exact})$
1	-0.020571	-0.00%	-0.020832	1.27%	-0.020571
2	-0.040632	-1.24%	-0.041627	1.18%	-0.041142
3	-0.060219	-2.42%	-0.062355	1.04%	-0.061713
4	-0.079364	-3.55%	-0.082992	0.86%	-0.082284
5	-0.098095	-4.63%	-0.103521	0.65%	-0.102855
6	-0.116439	-5.66%	-0.123929	0.41%	-0.123426
7	-0.134419	-6.65%	-0.144206	0.16%	-0.143997
8	-0.152055	-7.60%	-0.164344	-0.14%	-0.164567
9	-0.169367	-8.52%	-0.184338	-0.43%	-0.185138
10	-0.186371	-9.40%	-0.204183	-0.74%	-0.205709
11	-0.203084	-10.25%	-0.223877	-1.06%	-0.226280
12	-0.219519	-11.07%	-0.243418	-1.39%	-0.246851
13	-0.235691	-11.87%	-0.262806	-1.73%	-0.267422
14	-0.251612	-12.63%	-0.282041	-2.07%	-0.287993
15	-0.267292	-13.38%	-0.301122	-2.41%	-0.308564
16	-0.282743	-14.09%	-0.320052	-2.76%	-0.329135
17	-0.297975	-14.79%	-0.338830	-3.11%	-0.349706
18	-0.312996	-15.47%	-0.357459	-3.46%	-0.370277
19	-0.327814	-16.13%	-0.375941	-3.81%	-0.390848
20	-0.342439	-16.77%	-0.394276	-4.17%	-0.411419
100	-1.188453	-42.23%	-1.545648	-24.86%	-2.057093

Exercise 4.15

The normalized full CI wave function for a minimal basis H_2 molecule is

$$|\Phi_0\rangle = \frac{1}{\sqrt{1+c^2}}|1\bar{1}\rangle + \frac{c}{\sqrt{1+c^2}}|2\bar{2}\rangle$$

where $c = {}^1E_{\text{corr}}/K_{12}$. Show that for N independent minimal H_2 molecules, the overlap between the Hartree-Fock wave functions, $|\Psi_0\rangle$, and the exact *normalized* ground state wave function is

$$\langle\Psi_0|\Phi_0\rangle = \frac{1}{(1+c^2)^{\frac{N}{2}}}.$$

Using the values of the two-electron integrals for $R=1.4$ a.u., given in Appendix D, calculate $\langle\Psi_0|\Phi_0\rangle$ for $N=1, 10$, and 100 . Note that this overlap decreases quickly (in fact, exponentially) as N increases. Thus the overlap between the Hartree-Fock and the exact wave functions of the system exponentially approaches zero as the size of the system increases, even though the Hartree-Fock energy is size consistent. *Hint:* Because the N independent H_2 molecules are infinitely separated we can, for all intents and purposes, ignore the requirement that the wave function of this system be antisymmetric with respect to the interchange of electrons which belong to different H_2 molecules. Thus we can write

$$|\Phi_0\rangle \sim \prod_{i=1}^N \left(\frac{1}{\sqrt{1+c^2}}|1_i\bar{1}_i\rangle + \frac{c}{\sqrt{1+c^2}}|2_i\bar{2}_i\rangle \right)$$

and

$$|\Psi_0\rangle \sim \prod_{i=1}^N |1_i\bar{1}_i\rangle.$$

Solution 4.15

With the wave functions supplied in hints, we find that

$$\langle \Psi_0 | \Phi_0 \rangle = \frac{1}{(1+c^2)^{N/2}} \prod_{i=1}^N \prod_{j=1}^N [\langle 1_i \bar{1}_i | 1_j \bar{1}_j \rangle + \langle 1_i \bar{1}_i | 2_j \bar{2}_j \rangle] = \frac{1}{(1+c^2)^{N/2}} \prod_{i=1}^N \prod_{j=1}^N \delta_{ij} = \frac{1}{(1+c^2)^{N/2}}. \quad (4.15-1)$$

As $R = 1.4$ a.u., $\Delta = 0.78865$ a.u. and

$$c = \frac{{}^1E_{\text{corr}}}{K_{12}} = \frac{\Delta - \sqrt{\Delta^2 + K_{12}^2}}{K_{12}} = -0.1135.$$

The table of $\langle \Psi_0 | \Phi_0 \rangle$ and $\ln \langle \Psi_0 | \Phi_0 \rangle$ to N can be seen below. We conclude that this overlap decreases exponentially as N increases.

Table 4.3: The table of $\langle \Psi_0 | \Phi_0 \rangle$ and $\ln \langle \Psi_0 | \Phi_0 \rangle$ to N .

N	$\langle \Psi_0 \Phi_0 \rangle$	$\ln \langle \Psi_0 \Phi_0 \rangle$
1	0.9936	-0.00642
10	0.9380	-0.06401
100	0.5273	-0.63999
1000	0.0017	-6.37713

CHAPTER 5

Pair and Coupled-Pair Theories

5.1 The Independent Electron Pair Approximation (IEPA)

Exercise 5.1

The application of pair theory to minimal basis H_2 is trivial since we are dealing with a two-electron system for which the IEPA is exact, i.e., it gives the full CI result obtained in the last chapter, viz.

$${}^1E_{\text{corr}} = \Delta - (\Delta^2 + K_{12}^2)^{1/2}$$

where (see Eq.(4.20))

$$\Delta = (\varepsilon_2 - \varepsilon_1) + \frac{1}{2}(J_{11} + J_{22} - 4J_{12} + 2K_{12}).$$

- a. Calculate the correlation energy using first-order pairs. Remember that the summations in Eq.(5.19) go over spin orbitals (i.e., $a = 1, \bar{1}$, and $r = 2, \bar{2}$). Show that

$${}^1E_{\text{corr}}(\text{FO}) = \frac{K_{12}^2}{2(\varepsilon_1 - \varepsilon_2)}.$$

- b. Approximate Δ in the exact correlation energy by $\varepsilon_2 - \varepsilon_1$ and recover the first-order pair correlation energy by expanding the exact answer to first order using the relation $(1 + x)^{1/2} \approx 1 + x/2$.

Solution 5.1

- a. At this time, there is only one pair of electrons and one pair of virtual spin orbitals. Thus,

$$E_{\text{corr}}(\text{FO}) = \sum_{\substack{a < b \\ r < s}} \frac{|\langle \Psi_0 | \mathcal{H} | \Psi_{ab}^{rs} \rangle|^2}{\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s} = \frac{|\langle 1\bar{1} | 2\bar{2} \rangle|^2}{\varepsilon_1 + \varepsilon_{\bar{1}} - \varepsilon_2 - \varepsilon_{\bar{2}}} = -\frac{K_{12}^2}{2(\varepsilon_2 - \varepsilon_1)}. \quad (5.1-1)$$

- b. As $K_{12} \ll \Delta$, we find that

$${}^1E_{\text{corr}} = \Delta \left[1 - \sqrt{1 + \frac{K_{12}^2}{\Delta^2}} \right] = \Delta \left[1 - \left(1 + \frac{K_{12}^2}{2\Delta^2} + \dots \right) \right] = -\frac{K_{12}^2}{2\Delta} + \dots$$

Here, the truth that when $|x| \ll 1$,

$$(1 + x)^{\frac{1}{2}} \approx 1 + \frac{x}{2},$$

has been used.

After substitute $\Delta = \varepsilon_2 - \varepsilon_1$, we obtain

$${}^1E_{\text{corr}}(\text{FO}) = -\frac{K_{12}^2}{2(\varepsilon_2 - \varepsilon_1)} = \frac{K_{12}^2}{2(\varepsilon_1 - \varepsilon_2)}. \quad (5.1-2)$$

Exercise 5.2

Derive Eqs.(5.22a) and (5.22b).

Solution 5.2

From (5.9a), with $\langle \Psi_{1_i \bar{1}_i}^{2_i \bar{2}_i} | \mathcal{H} | \Psi_0 \rangle = \langle \Psi_0 | \mathcal{H} | \Psi_{1_i \bar{1}_i}^{2_i \bar{2}_i} \rangle = \langle 1_i \bar{1}_i | 2_i \bar{2}_i \rangle = K_{12}$, we obtain

$$K_{12} c_{1_i \bar{1}_i}^{2_i \bar{2}_i} = e_{1_i \bar{1}_i}, \quad (5.2-1)$$

which is (5.22a). Similarly, with (4.20), we obtain

$$K_{12} + \langle \Psi_{1_i \bar{1}_i}^{2_i \bar{2}_i} | \mathcal{H} - E_0 | \Psi_{1_i \bar{1}_i}^{2_i \bar{2}_i} \rangle = e_{1_i \bar{1}_i} c_{1_i \bar{1}_i}^{2_i \bar{2}_i} \quad (5.2-2)$$

which is (5.22b).

Exercise 5.3

Calculate the total first-order pair correlation energy for the dinner using Eq.(5.19) and show that it is twice the result obtained in Exercise 5.1.

Solution 5.3

Note that only $|2_1 \bar{2}_1 1_2 \bar{1}_2\rangle$ and $|1_1 \bar{1}_1 2_2 \bar{2}_2\rangle$ can interact with $\langle \Phi_0 |$ via the Hamiltonian \mathcal{H} , thus

$$E_{\text{corr}}(\text{FO}, 2\text{H}_2) = \frac{\langle \Psi_0 | \mathcal{H} | 2_1 \bar{2}_1 1_2 \bar{1}_2 \rangle}{\varepsilon_1 + \varepsilon_{\bar{1}} - \varepsilon_2 - \varepsilon_{\bar{2}}} + \frac{\langle \Psi_0 | \mathcal{H} | 1_1 \bar{1}_1 2_2 \bar{2}_2 \rangle}{\varepsilon_1 + \varepsilon_{\bar{1}} - \varepsilon_2 - \varepsilon_{\bar{2}}} = -\frac{K_{12}^2}{2(\varepsilon_1 - \varepsilon_2)} - \frac{K_{12}^2}{2(\varepsilon_1 - \varepsilon_2)} = 2E_{\text{corr}}(\text{FO}, \text{H}_2). \quad (5.3-1)$$

It is twice the result obtained in Exercise 5.1.

5.1.1 Invariance under Unitary Transformations: An Example**Exercise 5.4**

Show that $|a\bar{a}b\bar{b}\rangle = |1_1 \bar{1}_1 1_2 \bar{1}_2\rangle$. *Hint:* use Eq.(1.40) repeatedly. Eq.(1.40) for Slater determinants is

$$|\chi_1 \chi_2 \cdots \left(\sum_k c_k \chi'_k \right) \cdots \chi_N\rangle = \sum_k c_k |\chi_1 \chi_2 \cdots \chi'_k \cdots \chi_N\rangle.$$

Solution 5.4

Note that if any two rows (or columns) of a determinant are equal, the value of the determinant is zero. Therefore, we find that

$$\begin{aligned} |a\bar{a}b\bar{b}\rangle &= \frac{1}{\sqrt{2}} (|1_1 \bar{a} b \bar{b}\rangle + |1_2 \bar{a} b \bar{b}\rangle) \\ &= \frac{1}{\sqrt{2}} \left(\frac{1}{\sqrt{2}} |1_1 \bar{a} 1_1 \bar{b}\rangle - \frac{1}{\sqrt{2}} |1_1 \bar{a} 1_2 \bar{b}\rangle \right) + \frac{1}{\sqrt{2}} \left(\frac{1}{\sqrt{2}} |1_2 \bar{a} 1_1 \bar{b}\rangle - \frac{1}{\sqrt{2}} |1_2 \bar{a} 1_2 \bar{b}\rangle \right) \\ &= -\frac{1}{2} |1_1 \bar{a} 1_2 \bar{b}\rangle + \frac{1}{2} |1_2 \bar{a} 1_1 \bar{b}\rangle = -|1_1 \bar{a} 1_2 \bar{b}\rangle \\ &= -\frac{1}{\sqrt{2}} (|1_1 \bar{1}_1 1_2 \bar{b}\rangle + |1_1 \bar{1}_2 1_2 \bar{b}\rangle) \\ &= -\frac{1}{\sqrt{2}} \left(\frac{1}{\sqrt{2}} |1_1 \bar{1}_1 1_2 \bar{1}_1\rangle - \frac{1}{\sqrt{2}} |1_1 \bar{1}_1 1_2 \bar{1}_2\rangle \right) - \frac{1}{\sqrt{2}} \left(\frac{1}{\sqrt{2}} |1_1 \bar{1}_2 1_2 \bar{1}_1\rangle - \frac{1}{\sqrt{2}} |1_1 \bar{1}_2 1_2 \bar{1}_2\rangle \right) \\ &= \frac{1}{2} |1_1 \bar{1}_1 1_2 \bar{1}_2\rangle - \frac{1}{2} |1_1 \bar{1}_2 1_2 \bar{1}_1\rangle = |1_1 \bar{1}_1 1_2 \bar{1}_2\rangle. \end{aligned}$$

Concisely,

$$|a\bar{a}b\bar{b}\rangle = |1_1 \bar{1}_1 1_2 \bar{1}_2\rangle. \quad (5.4-1)$$

Exercise 5.5

Derive Eqs.(5.31a) and (5.31b).

Solution 5.5

With (5.28a), (5.28b), and (5.29), we find that

$$\begin{aligned}\langle \Psi_0 | \mathcal{H} | \Psi_{aa}^{**} \rangle &= \langle \Psi_0 | \mathcal{H} | \left(\frac{1}{\sqrt{2}} |\Psi_{aa}^{r\bar{r}}\rangle + \frac{1}{\sqrt{2}} |\Psi_{aa}^{s\bar{s}}\rangle \right) \rangle = \frac{1}{\sqrt{2}} \langle \Psi_0 | \mathcal{H} | \Psi_{aa}^{r\bar{r}} \rangle + \frac{1}{\sqrt{2}} \langle \Psi_0 | \mathcal{H} | \Psi_{aa}^{s\bar{s}} \rangle \\ &= \frac{1}{\sqrt{2}} \frac{K_{12}}{2} + \frac{1}{\sqrt{2}} \frac{K_{12}}{2} = \frac{K_{12}}{\sqrt{2}}.\end{aligned}$$

This is (5.31a). From (4.17a) and (4.17d), we know that

$$E_0 = \langle \Psi_0 | \mathcal{H} | \Psi_0 \rangle = \langle \Psi_0 | \mathcal{O}_1 | \Psi_0 \rangle + \langle \Psi_0 | \mathcal{O}_2 | \Psi_0 \rangle = 2(2h_{11} + J_{11}) = 4\varepsilon_1 - 2J_{11}.$$

And it is evident that

$$\begin{aligned}\langle \Psi_{aa}^{**} | \mathcal{H} | \Psi_{aa}^{**} \rangle &= \left(\frac{1}{\sqrt{2}} \langle \Psi_{aa}^{r\bar{r}} | + \frac{1}{\sqrt{2}} \langle \Psi_{aa}^{s\bar{s}} | \right) \mathcal{H} \left(\frac{1}{\sqrt{2}} |\Psi_{aa}^{r\bar{r}}\rangle + \frac{1}{\sqrt{2}} |\Psi_{aa}^{s\bar{s}}\rangle \right) \\ &= \frac{1}{2} \langle \Psi_{aa}^{r\bar{r}} | \mathcal{H} | \Psi_{aa}^{r\bar{r}} \rangle + \frac{1}{2} \langle \Psi_{aa}^{r\bar{r}} | \mathcal{H} | \Psi_{aa}^{s\bar{s}} \rangle + \frac{1}{2} \langle \Psi_{aa}^{s\bar{s}} | \mathcal{H} | \Psi_{aa}^{r\bar{r}} \rangle + \frac{1}{2} \langle \Psi_{aa}^{s\bar{s}} | \mathcal{H} | \Psi_{aa}^{s\bar{s}} \rangle.\end{aligned}$$

We have to calculate $\langle \Psi_{aa}^{r\bar{r}} | \mathcal{H} | \Psi_{aa}^{r\bar{r}} \rangle$, $\langle \Psi_{aa}^{r\bar{r}} | \mathcal{H} | \Psi_{aa}^{s\bar{s}} \rangle$ and $\langle \Psi_{aa}^{s\bar{s}} | \mathcal{H} | \Psi_{aa}^{s\bar{s}} \rangle$. Before their formal derivation, note that

$$h_{rr} = h_{ss} = h_{22}, \quad h_{aa} = h_{bb} = h_{11}.$$

With (5.26a)-(5.26d), we obtain

$$\begin{aligned}\langle \Psi_{aa}^{r\bar{r}} | \mathcal{H} | \Psi_{aa}^{r\bar{r}} \rangle &= \langle \Psi_{aa}^{r\bar{r}} | \mathcal{O}_1 | \Psi_{aa}^{r\bar{r}} \rangle + \langle \Psi_{aa}^{r\bar{r}} | \mathcal{O}_2 | \Psi_{aa}^{r\bar{r}} \rangle \\ &= \langle r|h|r \rangle + \langle \bar{r}|h|\bar{r} \rangle + \langle b|h|b \rangle + \langle \bar{b}|h|\bar{b} \rangle \\ &\quad + \langle r\bar{r}||r\bar{r} \rangle + \langle r\bar{b}||r\bar{b} \rangle + \langle r\bar{b}||r\bar{b} \rangle + \langle \bar{r}b||\bar{r}b \rangle + \langle \bar{r}\bar{b}||\bar{r}\bar{b} \rangle + \langle b\bar{b}||b\bar{b} \rangle \\ &= 2h_{rr} + 2h_{bb} + J_{rr} + (J_{rb} - K_{rb}) + J_{rb} + J_{rb} + (J_{rb} - K_{rb}) + J_{bb} \\ &= 2h_{11} + 2h_{22} + \frac{1}{2}J_{11} + \frac{1}{2}J_{22} + 2J_{12} - K_{12} \\ &= 2\varepsilon_1 + 2\varepsilon_2 - \frac{3}{2}J_{11} + \frac{1}{2}J_{22} - 2J_{12} + K_{12}.\end{aligned}$$

Similarly, we obtain that

$$\langle \Psi_{aa}^{r\bar{r}} | \mathcal{H} | \Psi_{aa}^{s\bar{s}} \rangle = \langle r\bar{r}||s\bar{s} \rangle = J_{rs} = \frac{1}{2}J_{22},$$

and

$$\begin{aligned}\langle \Psi_{aa}^{s\bar{s}} | \mathcal{H} | \Psi_{aa}^{s\bar{s}} \rangle &= 2h_{ss} + 2h_{aa} + J_{ss} + J_{sb} - K_{sb} + J_{sb} + J_{sb} + J_{sb} - K_{sb} + J_{bb} \\ &= 2h_{11} + 2h_{22} + \frac{1}{2}J_{11} + \frac{1}{2}J_{22} + 2J_{12} - K_{12}, \\ &= 2\varepsilon_1 + 2\varepsilon_2 - \frac{3}{2}J_{11} + \frac{1}{2}J_{22} - 2J_{12} + K_{12}.\end{aligned}$$

Hence,

$$\begin{aligned}\langle \Psi_{aa}^{**} | \mathcal{H} | \Psi_{aa}^{**} \rangle &= 2\varepsilon_1 + 2\varepsilon_2 - \frac{3}{2}J_{11} + \frac{1}{2}J_{22} - 2J_{12} + K_{12} + \frac{1}{2}J_{22} \\ &= 2\varepsilon_1 + 2\varepsilon_2 - \frac{3}{2}J_{11} + J_{22} - 2J_{12} + K_{12}, \\ \langle \Psi_{aa}^{**} | \mathcal{H} - E_0 | \Psi_{aa}^{**} \rangle &= 2\varepsilon_1 + 2\varepsilon_2 - \frac{3}{2}J_{11} + J_{22} - 2J_{12} + K_{12} - (4\varepsilon_1 - 2J_{11}) \\ &= 2(\varepsilon_2 - \varepsilon_1) + \frac{1}{2}J_{11} + J_{22} - 2J_{12} + K_{12}.\end{aligned}$$

In conclusion, we obtain

$$\langle \Psi_{aa}^{**} | \mathcal{H} - E_0 | \Psi_{aa}^{**} \rangle = 2(\varepsilon_2 - \varepsilon_1) + J_{22} + \frac{1}{2}(J_{11} - 4J_{12} + 2K_{12}) \equiv 2\Delta'. \quad (5.5-1)$$

This is (5.31b).

Exercise 5.6

Show that $e_{a\bar{b}} = e_{\bar{a}b} = e_{a\bar{a}}$.

Solution 5.6

The key point is to prove that the equations which determine $e_{a\bar{b}}$ are identical to that of $e_{a\bar{a}}$. Note that similar to Exercise 5.5, we can obtain

$$\begin{aligned}\langle \Psi_0 | \mathcal{H} | \Psi_{ab}^{**} \rangle &= \frac{K_{12}}{\sqrt{2}}, \\ \langle \Psi_{ab}^{**} | \mathcal{H} - E_0 | \Psi_{ab}^{**} \rangle &= 2\Delta'.\end{aligned}$$

Thus the equations of $e_{a\bar{b}}$ are

$$\begin{aligned}\frac{K_{12}}{\sqrt{2}}c &= e_{a\bar{b}}, \\ \frac{K_{12}}{\sqrt{2}} + 2\Delta'c &= e_{a\bar{b}}c.\end{aligned}$$

They are identical to that of $e_{a\bar{a}}$, thus $e_{a\bar{b}} = e_{a\bar{a}}$ and so does $e_{\bar{a}b}$.

Exercise 5.7

Show that DCI is invariant to unitary transformations for the above model.

a. The DCI wave function is

$$|\Psi_{\text{DCI}}\rangle = |\Psi_0\rangle + c_1|\Psi_{a\bar{a}}^{**}\rangle + c_2|\Psi_{b\bar{b}}^{**}\rangle + c_3|\Psi_{a\bar{b}}^{**}\rangle + c_4|\Psi_{\bar{a}b}^{**}\rangle.$$

Show that the corresponding eigenvalue problem which determines the DCI correlation energy of the dimer (${}^2E_{\text{corr}}(\text{DCI})$) is

$$\begin{pmatrix} 0 & \frac{1}{\sqrt{2}}K_{12} & \frac{1}{\sqrt{2}}K_{12} & \frac{1}{\sqrt{2}}K_{12} & \frac{1}{\sqrt{2}}K_{12} \\ \frac{1}{\sqrt{2}}K_{12} & 2\Delta' & \frac{1}{2}J_{11} & \frac{1}{2}K_{12} - J_{12} & \frac{1}{2}K_{12} - J_{12} \\ \frac{1}{\sqrt{2}}K_{12} & \frac{1}{2}J_{11} & 2\Delta' & \frac{1}{2}K_{12} - J_{12} & \frac{1}{2}K_{12} - J_{12} \\ \frac{1}{\sqrt{2}}K_{12} & \frac{1}{2}K_{12} - J_{12} & \frac{1}{2}K_{12} - J_{12} & 2\Delta' & \frac{1}{2}J_{11} \\ \frac{1}{\sqrt{2}}K_{12} & \frac{1}{2}K_{12} - J_{12} & \frac{1}{2}K_{12} - J_{12} & \frac{1}{2}J_{11} & 2\Delta' \end{pmatrix} \begin{pmatrix} 1 \\ c_1 \\ c_2 \\ c_3 \\ c_4 \end{pmatrix} = {}^2E_{\text{corr}}(\text{DCI}) \begin{pmatrix} 1 \\ c_1 \\ c_2 \\ c_3 \\ c_4 \end{pmatrix}.$$

b. Show that $c_1 = c_2 = c_3 = c_4 = c$ and then solve the equations to show

$${}^2E_{\text{corr}}(\text{DCI}) = \Delta - (\Delta^2 + 2K_{12}^2)^{1/2},$$

which is the same result as found in the last chapter (see Eq.(4.60)).

Solution 5.7

a. Some matrix elements have been solved in Exercise 5.5 and Exercise 5.6, and they are listed as follows.

$$\begin{aligned}\langle \Psi_0 | \mathcal{H} - E_0 | \Psi_0 \rangle &= 0, \\ \langle \Psi_0 | \mathcal{H} | \Psi_{a\bar{a}}^{**} \rangle &= \langle \Psi_{a\bar{a}}^{**} | \mathcal{H} | \Psi_0 \rangle = \frac{1}{\sqrt{2}}K_{12}, \\ \langle \Psi_0 | \mathcal{H} | \Psi_{b\bar{b}}^{**} \rangle &= \langle \Psi_{b\bar{b}}^{**} | \mathcal{H} | \Psi_0 \rangle = \frac{1}{\sqrt{2}}K_{12}, \\ \langle \Psi_0 | \mathcal{H} | \Psi_{a\bar{b}}^{**} \rangle &= \langle \Psi_{a\bar{b}}^{**} | \mathcal{H} | \Psi_0 \rangle = \frac{1}{\sqrt{2}}K_{12}, \\ \langle \Psi_0 | \mathcal{H} | \Psi_{\bar{a}b}^{**} \rangle &= \langle \Psi_{\bar{a}b}^{**} | \mathcal{H} | \Psi_0 \rangle = \frac{1}{\sqrt{2}}K_{12}, \\ \langle \Psi_{a\bar{a}}^{**} | \mathcal{H} - E_0 | \Psi_{a\bar{a}}^{**} \rangle &= 2\Delta', \\ \langle \Psi_{b\bar{b}}^{**} | \mathcal{H} - E_0 | \Psi_{b\bar{b}}^{**} \rangle &= 2\Delta',\end{aligned}$$

$$\begin{aligned}\langle \Psi_{a\bar{b}}^{**} | \mathcal{H} - E_0 | \Psi_{a\bar{b}}^{**} \rangle &= 2\Delta', \\ \langle \Psi_{\bar{a}b}^{**} | \mathcal{H} - E_0 | \Psi_{\bar{a}b}^{**} \rangle &= 2\Delta'.\end{aligned}$$

Now we pay attention to calculate other matrix elements.

$$\begin{aligned}\langle \Psi_{a\bar{a}}^{**} | \mathcal{H} | \Psi_{b\bar{b}}^{**} \rangle &= \left(\frac{1}{\sqrt{2}} \langle \Psi_{a\bar{a}}^{r\bar{r}} | + \frac{1}{\sqrt{2}} \langle \Psi_{a\bar{a}}^{s\bar{s}} | \right) \mathcal{H} \left(\frac{1}{\sqrt{2}} | \Psi_{b\bar{b}}^{r\bar{r}} \rangle + \frac{1}{\sqrt{2}} | \Psi_{b\bar{b}}^{s\bar{s}} \rangle \right) \\ &= \frac{1}{2} \langle \Psi_{a\bar{a}}^{r\bar{r}} | \mathcal{H} | \Psi_{b\bar{b}}^{r\bar{r}} \rangle + \frac{1}{2} \langle \Psi_{a\bar{a}}^{r\bar{r}} | \mathcal{H} | \Psi_{b\bar{b}}^{s\bar{s}} \rangle + \frac{1}{2} \langle \Psi_{a\bar{a}}^{s\bar{s}} | \mathcal{H} | \Psi_{b\bar{b}}^{r\bar{r}} \rangle + \frac{1}{2} \langle \Psi_{a\bar{a}}^{s\bar{s}} | \mathcal{H} | \Psi_{b\bar{b}}^{s\bar{s}} \rangle \\ &= \frac{1}{2} (\langle b\bar{b} | a\bar{a} \rangle + 0 + 0 + \langle b\bar{b} | a\bar{a} \rangle) = \frac{1}{2} (K_{ab} + K_{ab}) = K_{ab} = \frac{1}{2} J_{11}, \\ \langle \Psi_{b\bar{b}}^{**} | \mathcal{H} | \Psi_{a\bar{a}}^{**} \rangle &= (\langle \Psi_{a\bar{a}}^{**} | \mathcal{H} | \Psi_{b\bar{b}}^{**} \rangle)^* = \frac{1}{2} J_{11}, \\ \langle \Psi_{a\bar{a}}^{**} | \mathcal{H} | \Psi_{a\bar{b}}^{**} \rangle &= \left(\frac{1}{\sqrt{2}} \langle \Psi_{a\bar{a}}^{r\bar{r}} | + \frac{1}{\sqrt{2}} \langle \Psi_{a\bar{a}}^{s\bar{s}} | \right) \mathcal{H} \left(\frac{1}{\sqrt{2}} | \Psi_{a\bar{b}}^{r\bar{s}} \rangle + \frac{1}{\sqrt{2}} | \Psi_{a\bar{b}}^{s\bar{r}} \rangle \right) \\ &= \frac{1}{2} \langle \Psi_{a\bar{a}}^{r\bar{r}} | \mathcal{H} | \Psi_{a\bar{b}}^{r\bar{s}} \rangle + \frac{1}{2} \langle \Psi_{a\bar{a}}^{r\bar{r}} | \mathcal{H} | \Psi_{a\bar{b}}^{s\bar{r}} \rangle + \frac{1}{2} \langle \Psi_{a\bar{a}}^{s\bar{s}} | \mathcal{H} | \Psi_{a\bar{b}}^{r\bar{s}} \rangle + \frac{1}{2} \langle \Psi_{a\bar{a}}^{s\bar{s}} | \mathcal{H} | \Psi_{a\bar{b}}^{s\bar{r}} \rangle \\ &= \frac{1}{2} (\langle \bar{r}\bar{b} | \bar{a}\bar{s} \rangle - \langle \bar{r}\bar{b} | s\bar{a} \rangle - \langle \bar{s}\bar{b} | r\bar{a} \rangle + \langle \bar{s}\bar{b} | \bar{a}\bar{r} \rangle) \\ &= \frac{1}{2} ((ra|bs) - (rs|ba) - (rs|ba) - (sr|ba) + (sa|br) - (sr|ba)) \\ &= \frac{1}{2} \left(\frac{K_{12}}{2} - \frac{J_{12}}{2} - \frac{J_{12}}{2} - \frac{J_{12}}{2} + \frac{K_{12}}{2} - \frac{J_{12}}{2} \right) = \frac{1}{2} K_{12} - J_{12}, \\ \langle \Psi_{a\bar{b}}^{**} | \mathcal{H} | \Psi_{a\bar{a}}^{**} \rangle &= (\langle \Psi_{a\bar{a}}^{**} | \mathcal{H} | \Psi_{a\bar{b}}^{**} \rangle)^* = \frac{1}{2} K_{12} - J_{12}, \\ \langle \Psi_{a\bar{a}}^{**} | \mathcal{H} | \Psi_{\bar{a}b}^{**} \rangle &= \left(\frac{1}{\sqrt{2}} \langle \Psi_{a\bar{a}}^{r\bar{r}} | + \frac{1}{\sqrt{2}} \langle \Psi_{a\bar{a}}^{s\bar{s}} | \right) \mathcal{H} \left(\frac{1}{\sqrt{2}} | \Psi_{\bar{a}b}^{s\bar{r}} \rangle + \frac{1}{\sqrt{2}} | \Psi_{\bar{a}b}^{r\bar{s}} \rangle \right) \\ &= \frac{1}{2} \langle \Psi_{a\bar{a}}^{r\bar{r}} | \mathcal{H} | \Psi_{\bar{a}b}^{s\bar{r}} \rangle + \frac{1}{2} \langle \Psi_{a\bar{a}}^{r\bar{r}} | \mathcal{H} | \Psi_{\bar{a}b}^{r\bar{s}} \rangle + \frac{1}{2} \langle \Psi_{a\bar{a}}^{s\bar{s}} | \mathcal{H} | \Psi_{\bar{a}b}^{s\bar{r}} \rangle + \frac{1}{2} \langle \Psi_{a\bar{a}}^{s\bar{s}} | \mathcal{H} | \Psi_{\bar{a}b}^{r\bar{s}} \rangle \\ &= \frac{1}{2} (-\langle \bar{r}b | \bar{s}a \rangle + \langle \bar{r}b | as \rangle + \langle \bar{s}b | ar \rangle - \langle \bar{s}b | \bar{r}a \rangle) \\ &= \frac{1}{2} (-(rs|ba) + (ra|bs) - (rs|ba) + (sa|br) - (sr|ba) - (sr|ba)) \\ &= \frac{1}{2} \left(-\frac{J_{12}}{2} + \frac{K_{12}}{2} - \frac{J_{12}}{2} + \frac{K_{12}}{2} - \frac{J_{12}}{2} - \frac{J_{12}}{2} \right) = \frac{1}{2} K_{12} - J_{12}, \\ \langle \Psi_{\bar{a}b}^{**} | \mathcal{H} | \Psi_{a\bar{a}}^{**} \rangle &= (\langle \Psi_{a\bar{a}}^{**} | \mathcal{H} | \Psi_{\bar{a}b}^{**} \rangle)^* = \frac{1}{2} K_{12} - J_{12}, \\ \langle \Psi_{b\bar{b}}^{**} | \mathcal{H} | \Psi_{a\bar{b}}^{**} \rangle &= \left(\frac{1}{\sqrt{2}} \langle \Psi_{b\bar{b}}^{r\bar{r}} | + \frac{1}{\sqrt{2}} \langle \Psi_{b\bar{b}}^{s\bar{s}} | \right) \mathcal{H} \left(\frac{1}{\sqrt{2}} | \Psi_{a\bar{b}}^{r\bar{s}} \rangle + \frac{1}{\sqrt{2}} | \Psi_{a\bar{b}}^{s\bar{r}} \rangle \right) \\ &= \frac{1}{2} \langle \Psi_{b\bar{b}}^{r\bar{r}} | \mathcal{H} | \Psi_{a\bar{b}}^{r\bar{s}} \rangle + \frac{1}{2} \langle \Psi_{b\bar{b}}^{r\bar{r}} | \mathcal{H} | \Psi_{a\bar{b}}^{s\bar{r}} \rangle + \frac{1}{2} \langle \Psi_{b\bar{b}}^{s\bar{s}} | \mathcal{H} | \Psi_{a\bar{b}}^{r\bar{s}} \rangle + \frac{1}{2} \langle \Psi_{b\bar{b}}^{s\bar{s}} | \mathcal{H} | \Psi_{a\bar{b}}^{s\bar{r}} \rangle \\ &= \frac{1}{2} (-\langle a\bar{r} | b\bar{s} \rangle + \langle ar | sb \rangle + \langle as | rb \rangle - \langle a\bar{s} | b\bar{r} \rangle) \\ &= \frac{1}{2} (-(ab|rs) + (as|rb) - (ab|rs) + (ar|sb) - (ab|sr) - (ab|sr)) \\ &= \frac{1}{2} \left(-\frac{J_{12}}{2} + \frac{K_{12}}{2} - \frac{J_{12}}{2} + \frac{K_{12}}{2} - \frac{J_{12}}{2} - \frac{J_{12}}{2} \right) = \frac{1}{2} K_{12} - J_{12}, \\ \langle \Psi_{a\bar{b}}^{**} | \mathcal{H} | \Psi_{b\bar{b}}^{**} \rangle &= (\langle \Psi_{b\bar{b}}^{**} | \mathcal{H} | \Psi_{a\bar{b}}^{**} \rangle)^* = \frac{1}{2} K_{12} - J_{12}, \\ \langle \Psi_{b\bar{b}}^{**} | \mathcal{H} | \Psi_{\bar{a}b}^{**} \rangle &= \left(\frac{1}{\sqrt{2}} \langle \Psi_{b\bar{b}}^{r\bar{r}} | + \frac{1}{\sqrt{2}} \langle \Psi_{b\bar{b}}^{s\bar{s}} | \right) \mathcal{H} \left(\frac{1}{\sqrt{2}} | \Psi_{\bar{a}b}^{s\bar{r}} \rangle + \frac{1}{\sqrt{2}} | \Psi_{\bar{a}b}^{r\bar{s}} \rangle \right) \\ &= \frac{1}{2} \langle \Psi_{b\bar{b}}^{r\bar{r}} | \mathcal{H} | \Psi_{\bar{a}b}^{s\bar{r}} \rangle + \frac{1}{2} \langle \Psi_{b\bar{b}}^{r\bar{r}} | \mathcal{H} | \Psi_{\bar{a}b}^{r\bar{s}} \rangle + \frac{1}{2} \langle \Psi_{b\bar{b}}^{s\bar{s}} | \mathcal{H} | \Psi_{\bar{a}b}^{s\bar{r}} \rangle + \frac{1}{2} \langle \Psi_{b\bar{b}}^{s\bar{s}} | \mathcal{H} | \Psi_{\bar{a}b}^{r\bar{s}} \rangle \\ &= \frac{1}{2} (\langle \bar{a}\bar{r} | \bar{s}\bar{b} \rangle - \langle \bar{a}\bar{r} | \bar{b}s \rangle - \langle \bar{a}s | \bar{b}r \rangle + \langle \bar{a}\bar{s} | \bar{r}\bar{b} \rangle) \\ &= \frac{1}{2} ((as|rb) - (ab|rs) - (ab|rs) - (ab|sr) + (ar|sb) - (ab|sr))\end{aligned}$$

$$\begin{aligned}
&= \frac{1}{2} \left(\frac{K_{12}}{2} - \frac{J_{12}}{2} - \frac{J_{12}}{2} - \frac{J_{12}}{2} + \frac{K_{12}}{2} - \frac{J_{12}}{2} \right) = \frac{1}{2} K_{12} - J_{12}, \\
\langle \Psi_{ab}^{**} | \mathcal{H} | \Psi_{bb}^{**} \rangle &= (\langle \Psi_{bb}^{**} | \mathcal{H} | \Psi_{ab}^{**} \rangle)^* = \frac{1}{2} K_{12} - J_{12}, \\
\langle \Psi_{ab}^{**} | \mathcal{H} | \Psi_{ab}^{**} \rangle &= \left(\frac{1}{\sqrt{2}} \langle \Psi_{ab}^{r\bar{s}} | + \frac{1}{\sqrt{2}} \langle \Psi_{ab}^{s\bar{r}} | \right) \mathcal{H} \left(\frac{1}{\sqrt{2}} | \Psi_{ab}^{r\bar{s}} \rangle + \frac{1}{\sqrt{2}} | \Psi_{ab}^{s\bar{r}} \rangle \right) \\
&= \frac{1}{2} \langle \Psi_{ab}^{r\bar{s}} | \mathcal{H} | \Psi_{ab}^{r\bar{s}} \rangle + \frac{1}{2} \langle \Psi_{ab}^{r\bar{s}} | \mathcal{H} | \Psi_{ab}^{s\bar{r}} \rangle + \frac{1}{2} \langle \Psi_{ab}^{s\bar{r}} | \mathcal{H} | \Psi_{ab}^{r\bar{s}} \rangle + \frac{1}{2} \langle \Psi_{ab}^{s\bar{r}} | \mathcal{H} | \Psi_{ab}^{s\bar{r}} \rangle \\
&= \frac{1}{2} (\langle \bar{a}b | \bar{b}a \rangle + 0 + 0 + \langle \bar{a}b | \bar{b}a \rangle) = \frac{1}{2} (K_{ab} + K_{ab}) = \frac{1}{2} J_{11}, \\
\langle \Psi_{ab}^{**} | \mathcal{H} | \Psi_{ab}^{**} \rangle &= (\langle \Psi_{ab}^{**} | \mathcal{H} | \Psi_{ab}^{**} \rangle)^* = \frac{1}{2} J_{11}.
\end{aligned}$$

Thus, the corresponding DCI eigenvalue problem is

$$\begin{pmatrix} 0 & \frac{1}{\sqrt{2}} K_{12} & \frac{1}{\sqrt{2}} K_{12} & \frac{1}{\sqrt{2}} K_{12} & \frac{1}{\sqrt{2}} K_{12} \\ \frac{1}{\sqrt{2}} K_{12} & 2\Delta' & \frac{1}{2} J_{11} & \frac{1}{2} K_{12} - J_{12} & \frac{1}{2} K_{12} - J_{12} \\ \frac{1}{\sqrt{2}} K_{12} & \frac{1}{2} J_{11} & 2\Delta' & \frac{1}{2} K_{12} - J_{12} & \frac{1}{2} K_{12} - J_{12} \\ \frac{1}{\sqrt{2}} K_{12} & \frac{1}{2} K_{12} - J_{12} & \frac{1}{2} K_{12} - J_{12} & 2\Delta' & \frac{1}{2} J_{11} \\ \frac{1}{\sqrt{2}} K_{12} & \frac{1}{2} K_{12} - J_{12} & \frac{1}{2} K_{12} - J_{12} & \frac{1}{2} J_{11} & 2\Delta' \end{pmatrix} \begin{pmatrix} 1 \\ c_1 \\ c_2 \\ c_3 \\ c_4 \end{pmatrix} = {}^2E_{\text{corr}}(\text{DCI}) \begin{pmatrix} 1 \\ c_1 \\ c_2 \\ c_3 \\ c_4 \end{pmatrix}. \quad (5.7-1)$$

b. From (5.7-1), we know that

$$\begin{aligned}
\frac{K_{12}}{\sqrt{2}} + 2\Delta' c_1 + \frac{J_{11}}{2} c_2 + \frac{K_{12} - J_{12}}{2} c_3 + \frac{K_{12} - J_{12}}{2} c_4 &= {}^2E_{\text{corr}}(\text{DCI}) c_1, \\
\frac{K_{12}}{\sqrt{2}} + \frac{J_{11}}{2} c_1 + 2\Delta' c_2 + \frac{K_{12} - J_{12}}{2} c_3 + \frac{K_{12} - J_{12}}{2} c_4 &= {}^2E_{\text{corr}}(\text{DCI}) c_2.
\end{aligned}$$

The first equation can be subtracted by the second one, viz.,

$$(2\Delta' - \frac{J_{11}}{2} - {}^2E_{\text{corr}}(\text{DCI}))(c_1 - c_2) = 0.$$

Assume that ${}^2E_{\text{corr}}(\text{DCI}) \neq 2\Delta' - \frac{J_{11}}{2}$ (in fact, this holds true), we find that

$$c_1 = c_2.$$

In this way, we can prove $c_1 = c_2 = c_3 = c_4$. In fact, we can permute 1 and 2 of the second equation of (5.7-1), then we will find that the third equation of (5.7-1) is obtained, and vice versa. Thus $c_1 = c_2$. This method is suitable for not only c_1 and c_2 , but also c_1 and c_3 , c_1 and c_4 . Hence we can also conclude that $c_1 = c_2 = c_3 = c_4$.

Thus, we set $c_1 = c$, thus

$$\begin{aligned}
{}^2E_{\text{corr}}(\text{DCI}) &= 2\sqrt{2}K_{12}c, \\
{}^2E_{\text{corr}}(\text{DCI})c &= \frac{K_{12}}{\sqrt{2}} + (2\Delta' + \frac{1}{2}J_{11} + \frac{1}{2}K_{12} - J_{12} + \frac{1}{2}K_{12} - J_{12})c = \frac{K_{12}}{\sqrt{2}} + 2\Delta c.
\end{aligned}$$

In fact, they can be converted to a quadratic equation,

$$({}^2E_{\text{corr}}(\text{DCI}))^2 - 2\Delta({}^2E_{\text{corr}}(\text{DCI})) - 2K_{12}^2 = 0,$$

The discriminant Δ_E of this quadratic equation is

$$\Delta_E = (-2\Delta)^2 - 4 \times 1 \times (-2K_{12}^2) = 4(\Delta^2 + 2K_{12}^2) > 0,$$

and the root are

$$E_1 = \Delta + \sqrt{\Delta^2 + 2K_{12}^2}, \quad E_2 = \Delta - \sqrt{\Delta^2 + 2K_{12}^2}.$$

Therefore, the lowest root is the correlation energy, viz.,

$${}^2E_{\text{corr}} = \Delta - \sqrt{\Delta^2 + 2K_{12}^2}. \quad (5.7-2)$$

which is the same result as found in the last chapter (see Eq.(4.60)).

Exercise 5.8

Show directly from Eq.(5.19) using delocalized orbitals and the two-electron integrals in Eq.(5.26) that the total first-order pair correlation energy (which is the same as the many-body second-order perturbation energy) of the dimer is given by Eq.(5.46).

Solution 5.8

In fact, from Eq.(5.18) using delocalized orbitals and the two-electron integrals in Eq.(5.26), it is also easy to derive the the total first-order pair correlation energy. However, I think this method is much better because it shows the absence of e_{aa}^{FO} and e_{bb}^{FO} as their corresponding configurations are *ungerade* while Ψ_0 is *gerade*. Thus I will firstly show e_{ab}^{FO} , $e_{a\bar{b}}^{\text{FO}}$, $e_{b\bar{a}}^{\text{FO}}$, $e_{b\bar{b}}^{\text{FO}}$ and then calculate ${}^2E_{\text{corr}}(\text{FO}(D))$. Note that the integrals, which have three *gerade* orbitals and one *ungerade* orbital, or three *ungerade* orbitals and one *gerade* orbital, vanish. Thus,

$$\begin{aligned} e_{a\bar{a}}^{\text{FO}} &= \frac{|\langle a\bar{a}||r\bar{r}\rangle|^2}{\varepsilon_a + \varepsilon_{\bar{a}} - \varepsilon_r - \varepsilon_{\bar{r}}} + \frac{|\langle a\bar{a}||s\bar{s}\rangle|^2}{\varepsilon_a + \varepsilon_{\bar{a}} - \varepsilon_s - \varepsilon_{\bar{s}}} + \frac{|\langle a\bar{a}||r\bar{s}\rangle|^2}{\varepsilon_a + \varepsilon_{\bar{a}} - \varepsilon_r - \varepsilon_{\bar{s}}} + \frac{|\langle a\bar{a}||s\bar{r}\rangle|^2}{\varepsilon_a + \varepsilon_{\bar{a}} - \varepsilon_s - \varepsilon_{\bar{r}}} \\ &= \frac{1}{2(\varepsilon_1 - \varepsilon_2)} \left(\frac{1}{4}K_{12}^2 + \frac{1}{4}K_{12}^2 + 0 + 0 \right) = \frac{K_{12}^2}{4(\varepsilon_1 - \varepsilon_2)}. \end{aligned}$$

Similarly, we obtain

$$\begin{aligned} e_{a\bar{b}}^{\text{FO}} &= \frac{|\langle a\bar{b}||r\bar{r}\rangle|^2}{\varepsilon_a + \varepsilon_{\bar{b}} - \varepsilon_r - \varepsilon_{\bar{r}}} + \frac{|\langle a\bar{b}||s\bar{s}\rangle|^2}{\varepsilon_a + \varepsilon_{\bar{b}} - \varepsilon_s - \varepsilon_{\bar{s}}} + \frac{|\langle a\bar{b}||r\bar{s}\rangle|^2}{\varepsilon_a + \varepsilon_{\bar{b}} - \varepsilon_r - \varepsilon_{\bar{s}}} + \frac{|\langle a\bar{b}||s\bar{r}\rangle|^2}{\varepsilon_a + \varepsilon_{\bar{b}} - \varepsilon_s - \varepsilon_{\bar{r}}} = \frac{K_{12}^2}{4(\varepsilon_1 - \varepsilon_2)}, \\ e_{b\bar{a}}^{\text{FO}} &= \frac{|\langle b\bar{a}||r\bar{r}\rangle|^2}{\varepsilon_b + \varepsilon_{\bar{a}} - \varepsilon_r - \varepsilon_{\bar{r}}} + \frac{|\langle b\bar{a}||s\bar{s}\rangle|^2}{\varepsilon_b + \varepsilon_{\bar{a}} - \varepsilon_s - \varepsilon_{\bar{s}}} + \frac{|\langle b\bar{a}||r\bar{s}\rangle|^2}{\varepsilon_b + \varepsilon_{\bar{a}} - \varepsilon_r - \varepsilon_{\bar{s}}} + \frac{|\langle b\bar{a}||s\bar{r}\rangle|^2}{\varepsilon_b + \varepsilon_{\bar{a}} - \varepsilon_s - \varepsilon_{\bar{r}}} = \frac{K_{12}^2}{4(\varepsilon_1 - \varepsilon_2)}, \\ e_{b\bar{b}}^{\text{FO}} &= \frac{|\langle b\bar{b}||r\bar{r}\rangle|^2}{\varepsilon_b + \varepsilon_{\bar{b}} - \varepsilon_r - \varepsilon_{\bar{r}}} + \frac{|\langle b\bar{b}||s\bar{s}\rangle|^2}{\varepsilon_b + \varepsilon_{\bar{b}} - \varepsilon_s - \varepsilon_{\bar{s}}} + \frac{|\langle b\bar{b}||r\bar{s}\rangle|^2}{\varepsilon_b + \varepsilon_{\bar{b}} - \varepsilon_r - \varepsilon_{\bar{s}}} + \frac{|\langle b\bar{b}||s\bar{r}\rangle|^2}{\varepsilon_b + \varepsilon_{\bar{b}} - \varepsilon_s - \varepsilon_{\bar{r}}} = \frac{K_{12}^2}{4(\varepsilon_1 - \varepsilon_2)}. \end{aligned}$$

Thus

$${}^2E_{\text{corr}}(\text{FO}(D)) = e_{a\bar{a}}^{\text{FO}} + e_{a\bar{b}}^{\text{FO}} + e_{b\bar{a}}^{\text{FO}} + e_{b\bar{b}}^{\text{FO}} = \frac{K_{12}^2}{\varepsilon_1 - \varepsilon_2} = 2 \left(\frac{K_{12}^2}{2(\varepsilon_1 - \varepsilon_2)} \right). \quad (5.8-1)$$

Exercise 5.9

Show that the total correlation energy obtained using Epstein-Nesbet pairs is not invariant to unitary transformations.

a. Show, using localized orbitals, that

$${}^2E_{\text{corr}}(\text{EN}(L)) = -\frac{K_{12}^2}{\Delta}.$$

b. Show, using delocalized spin-orbital pairs, that

$${}^2E_{\text{corr}}(\text{EN}(D)) = -\frac{K_{12}^2}{\Delta'}.$$

c. Show, using delocalized spin-adapted pairs, that

$${}^2E_{\text{corr}}^{\text{singlet}}(\text{EN}(D)) = -\frac{K_{12}^2}{2\Delta'} - \frac{K_{12}^2}{2\Delta''}.$$

d. Using the STO-3G integrals for H_2 in Appendix D compare the numerical values of the above expressions at $R = 1.4$ a.u..

Solution 5.9

a. From Exercise 5.2, we know

$$\begin{aligned} \langle \Psi_0 | \mathcal{H} | \Psi_{1_1 \bar{1}_1}^{2_1 \bar{2}_1} \rangle &= \langle \Psi_0 | \mathcal{H} | \Psi_{1_2 \bar{1}_2}^{2_2 \bar{2}_2} \rangle = K_{12}, \\ \langle \Psi_{1_1 \bar{1}_1}^{2_1 \bar{2}_1} | \mathcal{H} - E_0 | \Psi_{1_1 \bar{1}_1}^{2_1 \bar{2}_1} \rangle &= \langle \Psi_{1_2 \bar{1}_2}^{2_2 \bar{2}_2} | \mathcal{H} - E_0 | \Psi_{1_2 \bar{1}_2}^{2_2 \bar{2}_2} \rangle = 2\Delta, \end{aligned}$$

thus we obtain that

$$e_{1_1\bar{1}_1}^{\text{EN}} = -\frac{|\langle \Psi_0 | \mathcal{H} | \Psi_{1_1\bar{1}_1}^{2_1\bar{2}_1} \rangle|^2}{\langle \Psi_{1_1\bar{1}_1}^{2_1\bar{2}_1} | \mathcal{H} - E_0 | \Psi_{1_1\bar{1}_1}^{2_1\bar{2}_1} \rangle} = -\frac{K_{12}^2}{2\Delta},$$

$$e_{1_2\bar{1}_2}^{\text{EN}} = -\frac{|\langle \Psi_0 | \mathcal{H} | \Psi_{1_2\bar{1}_2}^{2_2\bar{2}_2} \rangle|^2}{\langle \Psi_{1_2\bar{1}_2}^{2_2\bar{2}_2} | \mathcal{H} - E_0 | \Psi_{1_2\bar{1}_2}^{2_2\bar{2}_2} \rangle} = -\frac{K_{12}^2}{2\Delta}.$$

Therefore, with (5.16), we find that

$${}^2E_{\text{corr}}(\text{EN}(L)) = e_{1_1\bar{1}_1}^{\text{EN}} + e_{1_2\bar{1}_2}^{\text{EN}} = -\frac{K_{12}^2}{\Delta}. \quad (5.9-1)$$

b. From Exercise 5.5, we know

$$\langle \Psi_0 | \mathcal{H} | \Psi_{a\bar{a}}^{**} \rangle = \langle \Psi_{a\bar{a}}^{**} | \mathcal{H} | \Psi_0 \rangle = \frac{K_{12}}{\sqrt{2}},$$

$$\langle \Psi_{a\bar{a}}^{**} | \mathcal{H} - E_0 | \Psi_{a\bar{a}}^{**} \rangle = 2(\varepsilon_2 - \varepsilon_1) + J_{22} + \frac{1}{2}(J_{11} - 4J_{12} + 2K_{12}) \equiv 2\Delta',$$

thus we obtain that

$$e_{a\bar{a}}^{\text{EN}} = -\frac{|\langle \Psi_0 | \mathcal{H} | \Psi_{a\bar{a}}^{**} \rangle|^2}{\langle \Psi_{a\bar{a}}^{**} | \mathcal{H} - E_0 | \Psi_{a\bar{a}}^{**} \rangle} = -\frac{K_{12}^2}{4\Delta'}.$$

As described in the textbook, $e_{b\bar{b}}^{\text{EN}}$ equals $e_{a\bar{a}}^{\text{EN}}$ due to the high symmetry of this problem. Besides, in Exercise 5.6, we know that

$$\langle \Psi_0 | \mathcal{H} | \Psi_{a\bar{b}}^{**} \rangle = \langle \Psi_{a\bar{b}}^{**} | \mathcal{H} | \Psi_0 \rangle = \frac{K_{12}}{\sqrt{2}},$$

$$\langle \Psi_{a\bar{b}}^{**} | \mathcal{H} - E_0 | \Psi_{a\bar{b}}^{**} \rangle = 2\Delta',$$

and thus

$$e_{a\bar{b}}^{\text{EN}} = -\frac{|\langle \Psi_0 | \mathcal{H} | \Psi_{a\bar{b}}^{**} \rangle|^2}{\langle \Psi_{a\bar{b}}^{**} | \mathcal{H} - E_0 | \Psi_{a\bar{b}}^{**} \rangle} = -\frac{K_{12}^2}{4\Delta'}.$$

Similarly, we know $e_{\bar{a}b}^{\text{EN}} = e_{a\bar{b}}^{\text{EN}}$. Hence, we find that

$${}^2E_{\text{corr}}(\text{EN}(D)) = e_{a\bar{a}}^{\text{EN}} + e_{a\bar{b}}^{\text{EN}} + e_{\bar{a}b}^{\text{EN}} + e_{b\bar{b}}^{\text{EN}} = -\frac{K_{12}^2}{\Delta'}. \quad (5.9-2)$$

c. From (5.42a) and (5.42b), we know that

$$\langle \Psi_0 | \mathcal{H} | {}^B\Psi_{a\bar{b}}^{rs} \rangle = \langle {}^B\Psi_{a\bar{b}}^{rs} | \mathcal{H} | \Psi_0 \rangle = K_{12},$$

$$\langle {}^B\Psi_{a\bar{b}}^{rs} | \mathcal{H} - E_0 | {}^B\Psi_{a\bar{b}}^{rs} \rangle = 2(\varepsilon_2 - \varepsilon_1) + J_{11} + J_{22} - 2J_{12} + K_{12} \equiv 2\Delta'',$$

and thus,

$$e_{a\bar{b}}^{\text{singlet,EN}} = -\frac{|\langle \Psi_0 | \mathcal{H} | {}^B\Psi_{a\bar{b}}^{rs} \rangle|^2}{\langle {}^B\Psi_{a\bar{b}}^{rs} | \mathcal{H} - E_0 | {}^B\Psi_{a\bar{b}}^{rs} \rangle} = -\frac{K_{12}^2}{2\Delta''}.$$

Therefore, we obtain that

$${}^2E_{\text{corr}}^{\text{singlet}}(\text{EN}(D)) = e_{a\bar{a}}^{\text{EN}} + e_{b\bar{b}}^{\text{EN}} + e_{a\bar{b}}^{\text{singlet,EN}} = -\frac{K_{12}^2}{2\Delta'} - \frac{K_{12}^2}{2\Delta''}. \quad (5.9-3)$$

d. As $R = 1.4$ a.u., we know that

$$\begin{aligned} \varepsilon_1 &= -0.5782 \text{ a.u.}, & \varepsilon_2 &= 0.6703 \text{ a.u.}, & J_{11} &= 0.6746 \text{ a.u.}, \\ J_{12} &= 0.6636 \text{ a.u.}, & J_{22} &= 0.6975 \text{ a.u.}, & K_{12} &= 0.1813 \text{ a.u.} \end{aligned}$$

Thus, we obtain that

$$\begin{aligned}\Delta &= \varepsilon_2 - \varepsilon_1 + \frac{1}{2}J_{11} + \frac{1}{2}J_{22} - 2J_{12} + K_{12} = 0.78865 \text{ a.u.}, \\ \Delta' &= \varepsilon_2 - \varepsilon_1 + \frac{1}{4}J_{11} + \frac{1}{2}J_{22} - J_{12} + \frac{1}{2}K_{12} = 1.19295 \text{ a.u.}, \\ \Delta'' &= \varepsilon_2 - \varepsilon_1 + \frac{1}{2}J_{11} + \frac{1}{2}J_{22} - J_{12} + \frac{1}{2}K_{12} = 1.3616 \text{ a.u.},\end{aligned}$$

Finally, we get that

$$\begin{aligned}{}^2E_{\text{corr}}(\text{EN}(L)) &= -0.04168 \text{ a.u.}, \\ {}^2E_{\text{corr}}(\text{EN}(D)) &= -0.02755 \text{ a.u.}, \\ {}^2E_{\text{corr}}^{\text{singlet}}(\text{EN}(D)) &= -0.02585 \text{ a.u.}\end{aligned}$$

Now we conclude that the Epstein-Nesbet pair theory is not only variant to unitary transformations of degenerate HF orbitals, but also gives different answers depending on whether one uses spin-orbital or spin-adapted pair functions.

Exercise 5.10

The DCI wave function for the H_2 dimer using spin-adapted configurations is

$$|\Psi_{\text{DCI}}\rangle = |\Psi_0\rangle + c_1|\Psi_{a\bar{a}}^{**}\rangle + c_2|\Psi_{b\bar{b}}^{**}\rangle + c_3|{}^B\Psi_{ab}^{rs}\rangle.$$

Show that the corresponding DCI eigenvalue problem is

$$\begin{pmatrix} 0 & \frac{1}{\sqrt{2}}K_{12} & \frac{1}{\sqrt{2}}K_{12} & K_{12} \\ \frac{1}{\sqrt{2}}K_{12} & 2\Delta' & \frac{1}{2}J_{11} & \frac{1}{\sqrt{2}}(K_{12} - 2J_{12}) \\ \frac{1}{\sqrt{2}}K_{12} & \frac{1}{2}J_{11} & 2\Delta' & \frac{1}{\sqrt{2}}(K_{12} - 2J_{12}) \\ K_{12} & \frac{1}{\sqrt{2}}(K_{12} - 2J_{12}) & \frac{1}{\sqrt{2}}(K_{12} - 2J_{12}) & 2\Delta'' \end{pmatrix} \begin{pmatrix} 1 \\ c_1 \\ c_2 \\ c_3 \end{pmatrix} = {}^2E_{\text{corr}}(\text{DCI}) \begin{pmatrix} 1 \\ c_1 \\ c_2 \\ c_3 \end{pmatrix}.$$

and then solve the resulting equations to show that

$${}^2E_{\text{corr}}(\text{DCI}) = \Delta - (\Delta^2 + 2K_{12}^2)^{1/2}.$$

Solution 5.10

From Exercise 5.7 and Exercise 5.9, we know that

$$\begin{aligned}\langle \Psi_0 | \mathcal{H} - E_0 | \Psi_0 \rangle &= 0, \\ \langle \Psi_{a\bar{a}}^{**} | \mathcal{H} - E_0 | \Psi_{a\bar{a}}^{**} \rangle &= 2\Delta', \\ \langle \Psi_{b\bar{b}}^{**} | \mathcal{H} - E_0 | \Psi_{b\bar{b}}^{**} \rangle &= 2\Delta', \\ \langle {}^B\Psi_{ab}^{rs} | \mathcal{H} - E_0 | {}^B\Psi_{ab}^{rs} \rangle &= 2(\varepsilon_2 - \varepsilon_1) + J_{11} + J_{22} - 2J_{12} + K_{12} = 2\Delta'', \\ \langle \Psi_0 | \mathcal{H} | \Psi_{a\bar{a}}^{**} \rangle &= \langle \Psi_{a\bar{a}}^{**} | \mathcal{H} | \Psi_0 \rangle = \frac{1}{\sqrt{2}}K_{12}, \\ \langle \Psi_0 | \mathcal{H} | \Psi_{b\bar{b}}^{**} \rangle &= \langle \Psi_{b\bar{b}}^{**} | \mathcal{H} | \Psi_0 \rangle = \frac{1}{\sqrt{2}}K_{12}, \\ \langle \Psi_0 | \mathcal{H} | {}^B\Psi_{ab}^{rs} \rangle &= \langle {}^B\Psi_{ab}^{rs} | \mathcal{H} | \Psi_0 \rangle = K_{12}.\end{aligned}$$

Thus, we pay attention to calculating $\langle \Psi_{a\bar{a}}^{**} | \mathcal{H} | \Psi_{b\bar{b}}^{**} \rangle$, $\langle \Psi_{a\bar{a}}^{**} | \mathcal{H} | {}^B\Psi_{ab}^{rs} \rangle$.

$$\begin{aligned}\langle \Psi_{a\bar{a}}^{**} | \mathcal{H} | \Psi_{b\bar{b}}^{**} \rangle &= \left(\frac{1}{\sqrt{2}}\langle \Psi_{a\bar{a}}^{r\bar{r}} | + \frac{1}{\sqrt{2}}\langle \Psi_{a\bar{a}}^{s\bar{s}} | \right) \mathcal{H} \left(\frac{1}{\sqrt{2}}| \Psi_{b\bar{b}}^{r\bar{r}} \rangle + \frac{1}{\sqrt{2}}| \Psi_{b\bar{b}}^{s\bar{s}} \rangle \right) \\ &= \frac{1}{2}\langle \Psi_{a\bar{a}}^{r\bar{r}} | \mathcal{H} | \Psi_{b\bar{b}}^{r\bar{r}} \rangle + \frac{1}{2}\langle \Psi_{a\bar{a}}^{r\bar{r}} | \mathcal{H} | \Psi_{b\bar{b}}^{s\bar{s}} \rangle + \frac{1}{2}\langle \Psi_{a\bar{a}}^{s\bar{s}} | \mathcal{H} | \Psi_{b\bar{b}}^{r\bar{r}} \rangle + \frac{1}{2}\langle \Psi_{a\bar{a}}^{s\bar{s}} | \mathcal{H} | \Psi_{b\bar{b}}^{s\bar{s}} \rangle \\ &= \frac{1}{2}(\langle b\bar{b} | a\bar{a} \rangle + 0 + 0 + \langle b\bar{b} | a\bar{a} \rangle) = \frac{1}{2}(K_{ba} + K_{ba}) = K_{ab} = \frac{1}{2}J_{11},\end{aligned}$$

$$\begin{aligned}
\langle \Psi_{aa}^{**} | \mathcal{H} |^B \Psi_{ab}^{rs} \rangle &= \left(\frac{1}{\sqrt{2}} \langle \Psi_{aa}^{r\bar{r}} | + \frac{1}{\sqrt{2}} \langle \Psi_{aa}^{s\bar{s}} | \right) \mathcal{H} \left(\frac{1}{2} | \Psi_{ab}^{s\bar{r}} \rangle + \frac{1}{2} | \Psi_{ab}^{\bar{r}s} \rangle + \frac{1}{2} | \Psi_{ab}^{r\bar{s}} \rangle + \frac{1}{2} | \Psi_{ab}^{s\bar{s}} \rangle \right) \\
&= \frac{1}{2\sqrt{2}} [\langle \Psi_{aa}^{r\bar{r}} | \mathcal{H} | \Psi_{ab}^{s\bar{r}} \rangle + \langle \Psi_{aa}^{r\bar{r}} | \mathcal{H} | \Psi_{ab}^{\bar{r}s} \rangle + \langle \Psi_{aa}^{r\bar{r}} | \mathcal{H} | \Psi_{ab}^{r\bar{s}} \rangle + \langle \Psi_{aa}^{r\bar{r}} | \mathcal{H} | \Psi_{ab}^{s\bar{s}} \rangle \\
&\quad + \langle \Psi_{aa}^{s\bar{s}} | \mathcal{H} | \Psi_{ab}^{s\bar{r}} \rangle + \langle \Psi_{aa}^{s\bar{s}} | \mathcal{H} | \Psi_{ab}^{\bar{r}s} \rangle + \langle \Psi_{aa}^{s\bar{s}} | \mathcal{H} | \Psi_{ab}^{r\bar{s}} \rangle + \langle \Psi_{aa}^{s\bar{s}} | \mathcal{H} | \Psi_{ab}^{s\bar{s}} \rangle] \\
&= \frac{1}{2\sqrt{2}} (-\langle \bar{r}b | | \bar{s}a \rangle + \langle rb | | as \rangle + \langle \bar{r}\bar{b} | | \bar{a}\bar{s} \rangle - \langle r\bar{b} | | s\bar{a} \rangle \\
&\quad + \langle sb | | ar \rangle - \langle \bar{s}b | | \bar{r}a \rangle - \langle s\bar{b} | | r\bar{a} \rangle + \langle \bar{s}\bar{b} | | \bar{a}\bar{r} \rangle) \\
&= \frac{1}{2\sqrt{2}} (-\langle rs | ba \rangle + \langle ra | bs \rangle - \langle rs | ba \rangle + \langle ra | bs \rangle - \langle rs | ba \rangle - \langle rs | ba \rangle \\
&\quad + \langle sa | br \rangle - \langle sr | ba \rangle - \langle sr | ba \rangle - \langle sr | ba \rangle + \langle sa | br \rangle - \langle sr | ba \rangle) \\
&= \frac{1}{2\sqrt{2}} \left(-8 \times \frac{J_{12}}{2} + 4 \times \frac{K_{12}}{2} \right) = -\sqrt{2}J_{12} + \frac{1}{\sqrt{2}}K_{12} = \frac{1}{\sqrt{2}}(K_{12} - 2J_{12}).
\end{aligned}$$

Due to the high symmetry of this problem, we find that

$$\langle \Psi_{bb}^{**} | \mathcal{H} |^B \Psi_{ab}^{rs} \rangle = \langle \Psi_{aa}^{**} | \mathcal{H} |^B \Psi_{ab}^{rs} \rangle = \frac{1}{\sqrt{2}}(K_{12} - 2J_{12}).$$

and

$$\begin{aligned}
\langle \Psi_{bb}^{**} | \mathcal{H} | \Psi_{aa}^{**} \rangle &= (\langle \Psi_{aa}^{**} | \mathcal{H} | \Psi_{bb}^{**} \rangle)^* = \frac{1}{2}J_{11}, \\
\langle {}^B \Psi_{ab}^{rs} | \mathcal{H} | \Psi_{aa}^{**} \rangle &= (\langle \Psi_{aa}^{**} | \mathcal{H} |^B \Psi_{ab}^{rs} \rangle)^* = \frac{1}{\sqrt{2}}(K_{12} - 2J_{12}). \\
\langle {}^B \Psi_{ab}^{rs} | \mathcal{H} | \Psi_{bb}^{**} \rangle &= (\langle \Psi_{bb}^{**} | \mathcal{H} |^B \Psi_{ab}^{rs} \rangle)^* = \frac{1}{\sqrt{2}}(K_{12} - 2J_{12}).
\end{aligned}$$

Thus, we obtain the corresponding DCI eigenvalue problem is

$$\begin{pmatrix} 0 & \frac{1}{\sqrt{2}}K_{12} & \frac{1}{\sqrt{2}}K_{12} & K_{12} \\ \frac{1}{\sqrt{2}}K_{12} & 2\Delta' & \frac{1}{2}J_{11} & \frac{1}{\sqrt{2}}(K_{12} - 2J_{12}) \\ \frac{1}{\sqrt{2}}K_{12} & \frac{1}{2}J_{11} & 2\Delta' & \frac{1}{\sqrt{2}}(K_{12} - 2J_{12}) \\ K_{12} & \frac{1}{\sqrt{2}}(K_{12} - 2J_{12}) & \frac{1}{\sqrt{2}}(K_{12} - 2J_{12}) & 2\Delta'' \end{pmatrix} \begin{pmatrix} 1 \\ c_1 \\ c_2 \\ c_3 \end{pmatrix} = {}^2E_{\text{corr}}(\text{DCI}) \begin{pmatrix} 1 \\ c_1 \\ c_2 \\ c_3 \end{pmatrix}.$$

In the same way, it is evident that $c_1 = c_2$. Let $c_1 = c$ and then, the matrix can be converted into

$$\begin{pmatrix} \sqrt{2}K_{12} & K_{12} \\ 2\Delta - {}^2E_{\text{corr}}(\text{DCI}) & \frac{1}{\sqrt{2}}(2\Delta - {}^2E_{\text{corr}}(\text{DCI})) \\ \sqrt{2}(K_{12} - 2J_{12}) & 2\Delta'' - {}^2E_{\text{corr}}(\text{DCI}) \end{pmatrix} \begin{pmatrix} c \\ c_3 \end{pmatrix} = \begin{pmatrix} {}^2E_{\text{corr}}(\text{DCI}) \\ -\sqrt{2}K_{12} \\ -K_{12} \end{pmatrix}.$$

Note that a minor of the coefficient matrix is zero, viz.,

$$\begin{vmatrix} \sqrt{2}K_{12} & K_{12} \\ 2\Delta - {}^2E_{\text{corr}}(\text{DCI}) & \frac{1}{\sqrt{2}}(2\Delta - {}^2E_{\text{corr}}(\text{DCI})) \end{vmatrix} = 0,$$

and we conclude that the first two equations are linearly dependent if c and c_3 exist. Thus,

$$\frac{{}^2E_{\text{corr}}(\text{DCI})}{\sqrt{2}K_{12}} = -\frac{\sqrt{2}K_{12}}{2\Delta - {}^2E_{\text{corr}}(\text{DCI})} \Leftrightarrow ({}^2E_{\text{corr}}(\text{DCI}))^2 - 2\Delta({}^2E_{\text{corr}}(\text{DCI})) - 2K_{12}^2 = 0.$$

The discriminant Δ_E of this quadratic equation is

$$\Delta_E = (-2\Delta)^2 - 4 \times 1 \times (-2K_{12}^2) = 4(\Delta^2 + 2K_{12}^2) > 0$$

and the lowest root is

$${}^2E_{\text{corr}}(\text{DCI}) = \Delta - \sqrt{\Delta^2 + 2K_{12}^2}. \quad (5.10-1)$$

As the textbook mentioned, the DCI gives the same answer regardless of using spin-orbital or spin-adapted pair functions.

5.1.2 Some Illustrative Calculations

5.2 Coupled-Pair Theories

5.2.1 The Coupled Cluster Approximation (CCA)

5.2.2 The Cluster Expansion of the Wave Function

Exercise 5.11

Show that the wave function two independent H₂ molecules in Eqs.(5.49) and (5.50) can be written as

$$|\Phi\rangle = \exp(c_{1_1\bar{1}_1}^{2_1\bar{2}_1} a_{2_1}^\dagger a_{\bar{2}_1}^\dagger a_{\bar{1}_1} a_{1_1} + c_{1_2\bar{1}_2}^{2_2\bar{2}_2} a_{2_2}^\dagger a_{\bar{2}_2}^\dagger a_{\bar{1}_2} a_{1_2}) |1_1\bar{1}_1 1_2\bar{1}_2\rangle.$$

Solution 5.11

It is evident that

$$\begin{aligned} |\Phi\rangle &= \exp(c_{1_1\bar{1}_1}^{2_1\bar{2}_1} a_{2_1}^\dagger a_{\bar{2}_1}^\dagger a_{\bar{1}_1} a_{1_1} + c_{1_2\bar{1}_2}^{2_2\bar{2}_2} a_{2_2}^\dagger a_{\bar{2}_2}^\dagger a_{\bar{1}_2} a_{1_2}) |1_1\bar{1}_1 1_2\bar{1}_2\rangle \\ &= (1 + c_{1_1\bar{1}_1}^{2_1\bar{2}_1} a_{2_1}^\dagger a_{\bar{2}_1}^\dagger a_{\bar{1}_1} a_{1_1} + c_{1_2\bar{1}_2}^{2_2\bar{2}_2} a_{2_2}^\dagger a_{\bar{2}_2}^\dagger a_{\bar{1}_2} a_{1_2} + c_{1_1\bar{1}_1}^{2_1\bar{2}_1} c_{1_2\bar{1}_2}^{2_2\bar{2}_2} a_{2_1}^\dagger a_{\bar{2}_1}^\dagger a_{\bar{1}_1} a_{1_1} a_{2_2}^\dagger a_{\bar{2}_2}^\dagger a_{\bar{1}_2} a_{1_2}) |1_1\bar{1}_1 1_2\bar{1}_2\rangle \\ &= |1_1\bar{1}_1 1_2\bar{1}_2\rangle + c_{1_1\bar{1}_1}^{2_1\bar{2}_1} |2_1\bar{2}_1 1_2\bar{1}_2\rangle + c_{1_2\bar{1}_2}^{2_2\bar{2}_2} |1_1\bar{1}_1 2_2\bar{2}_2\rangle + c_{1_1\bar{1}_1}^{2_1\bar{2}_1} c_{1_2\bar{1}_2}^{2_2\bar{2}_2} |2_1\bar{2}_1 2_2\bar{2}_2\rangle, \end{aligned}$$

which equals (5.49) and (5.50).

5.2.3 Linear CCA and the Coupled Electron Pair Approximation (CEPA)

Exercise 5.12

- Show that if the matrix \mathbf{D} is approximated by its diagonal elements, the L-CCA correlation energy is identical to the result obtained using Epstein-Nesbet pairs (i.e., Eqs.(5.15) and (5.16)).
- Show that linear CCA is invariant under unitary transformations for the problem of two independent H₂ molecules. First show that for this model the correlation energy of the dimer using localized orbitals is the same as that obtained in Exercise 5.9a. Then show using delocalized spin orbitals that, in contrast to the results of Exercise 5.9, one gets the same correlation energy. You will find the DCI matrix given in Exercise 5.7 useful.

Solution 5.12

- If the matrix \mathbf{D} is approximated by its diagonal elements, their mathematical expression will be

$$(\mathbf{D})_{rasb,tcud} = \delta_{rs,tu} \delta_{ab,cd} \langle \Psi_{ab}^{rs} | \mathcal{H} - E_0 | \Psi_{ab}^{rs} \rangle.$$

It is evident that the mathematical expression of matrix elements of \mathbf{D}^{-1} is

$$(\mathbf{D}^{-1})_{rasb,tcud} = \frac{\delta_{rs,tu} \delta_{ab,cd}}{\langle \Psi_{ab}^{rs} | \mathcal{H} - E_0 | \Psi_{ab}^{rs} \rangle}.$$

Thus, at this time, the L-CCA correlation energy is

$$\begin{aligned} E_{\text{corr}}(\text{L-CCA}(\text{diagonal } \mathbf{D})) &= - \sum_{\substack{a < b \\ r < s}} \sum_{\substack{c < d \\ t < u}} \mathbf{B}_{rasb}^\dagger (\mathbf{D}^{-1})_{rasb,tcud} \mathbf{B}_{tcud} \\ &= - \sum_{\substack{a < b \\ r < s}} \sum_{\substack{c < d \\ t < u}} \langle \Psi_0 | \mathcal{H} | \Psi_{ab}^{rs} \rangle \frac{\delta_{rs,tu} \delta_{ab,cd}}{\langle \Psi_{ab}^{rs} | \mathcal{H} - E_0 | \Psi_{ab}^{rs} \rangle} \langle \Psi_{cd}^{tu} | \mathcal{H} | \Psi_0 \rangle \\ &= - \sum_{\substack{a < b \\ r < s}} \frac{\langle \Psi_0 | \mathcal{H} | \Psi_{ab}^{rs} \rangle \langle \Psi_{ab}^{rs} | \mathcal{H} | \Psi_0 \rangle}{\langle \Psi_{ab}^{rs} | \mathcal{H} - E_0 | \Psi_{ab}^{rs} \rangle}. \end{aligned}$$

However, the (5.15) and (5.16) gives

$$E_{\text{corr}}(\text{EN}) = \sum_{a < b} e_{ab}^{\text{EN}} = \sum_{a < b} \left(- \sum_{r < s} \frac{|\langle \Psi_0 | \mathcal{H} | \Psi_{ab}^{rs} \rangle|^2}{\langle \Psi_{ab}^{rs} | \mathcal{H} - E_0 | \Psi_{ab}^{rs} \rangle} \right) = - \sum_{\substack{a < b \\ r < s}} \frac{\langle \Psi_0 | \mathcal{H} | \Psi_{ab}^{rs} \rangle \langle \Psi_{ab}^{rs} | \mathcal{H} | \Psi_0 \rangle}{\langle \Psi_{ab}^{rs} | \mathcal{H} - E_0 | \Psi_{ab}^{rs} \rangle}.$$

In conclusion, if the matrix \mathbf{D} is approximated by its diagonal elements, the L-CCA correlation energy is identical to the result obtained using Epstein-Nesbet pairs.

- b. When localized orbitals are used, based on $|2_1\bar{2}_11_2\bar{1}_2\rangle$ and $|1_1\bar{1}_12_2\bar{2}_2\rangle$, its \mathbf{B} and \mathbf{D} are

$$\mathbf{B} = \begin{pmatrix} K_{12} \\ K_{12} \end{pmatrix}, \quad \mathbf{D} = \begin{pmatrix} 2\Delta & 0 \\ 0 & 2\Delta \end{pmatrix} \Rightarrow \mathbf{D}^{-1} = \begin{pmatrix} \frac{1}{2\Delta} & 0 \\ 0 & \frac{1}{2\Delta} \end{pmatrix}.$$

From (5.65), we obtain that

$$E_{\text{corr}}(\text{L-CCA}(L)) = - (K_{12} \quad K_{12}) \begin{pmatrix} \frac{1}{2\Delta} & 0 \\ 0 & \frac{1}{2\Delta} \end{pmatrix} \begin{pmatrix} K_{12} \\ K_{12} \end{pmatrix} = -\frac{K_{12}^2}{\Delta}. \quad (5.12-1)$$

When delocalized orbitals are used, based on $|\Psi_{a\bar{a}}^{**}\rangle$, $|\Psi_{b\bar{b}}^{**}\rangle$, $|\Psi_{a\bar{b}}^{**}\rangle$, $|\Psi_{\bar{a}b}^{**}\rangle$, its \mathbf{B} and \mathbf{D} are

$$\mathbf{B} = \begin{pmatrix} \frac{K_{12}}{\sqrt{2}} \\ \frac{K_{12}}{\sqrt{2}} \\ \frac{K_{12}}{\sqrt{2}} \\ \frac{K_{12}}{\sqrt{2}} \end{pmatrix}, \quad \mathbf{D} = \begin{pmatrix} 2\Delta' & \frac{1}{2}J_{11} & \frac{1}{2}K_{12} - J_{12} & \frac{1}{2}K_{12} - J_{12} \\ \frac{1}{2}J_{11} & 2\Delta' & \frac{1}{2}K_{12} - J_{12} & \frac{1}{2}K_{12} - J_{12} \\ \frac{1}{2}K_{12} - J_{12} & \frac{1}{2}K_{12} - J_{12} & 2\Delta' & \frac{1}{2}J_{11} \\ \frac{1}{2}K_{12} - J_{12} & \frac{1}{2}K_{12} - J_{12} & \frac{1}{2}J_{11} & 2\Delta' \end{pmatrix}.$$

In fact, it is difficult to calculate the inverse of \mathbf{D} directly. Here I will introduce how to calculate it. Notations are listed as follows.

$$a = 2\Delta' + \frac{1}{2}J_{11}, \quad c = \frac{1}{2}K_{12} - J_{12}, \quad A \equiv \begin{pmatrix} 2\Delta' & \frac{1}{2}J_{11} \\ \frac{1}{2}J_{11} & 2\Delta' \end{pmatrix}, \quad \delta \equiv \begin{pmatrix} 1 \\ 1 \end{pmatrix}.$$

Thus, the matrix \mathbf{D} can be expressed as

$$\mathbf{D} = \begin{pmatrix} A & c\delta\delta^T \\ c\delta\delta^T & A \end{pmatrix}.$$

Due to the symmetry of \mathbf{D} , its inverse \mathbf{D}^{-1} can be set in the form of

$$\begin{pmatrix} M & N \\ N & M \end{pmatrix},$$

where M and N are 2×2 matrices and M is symmetric. Then,

$$\mathbf{D}\mathbf{D}^{-1} = \begin{pmatrix} A & c\delta\delta^T \\ c\delta\delta^T & A \end{pmatrix} \begin{pmatrix} M & N \\ N & M \end{pmatrix} = \begin{pmatrix} \mathbf{1}_2 & 0_2 \\ 0_2 & \mathbf{1}_2 \end{pmatrix},$$

where $\mathbf{1}_2$ is a 2-order identity matrix and 0_2 is a 2×2 zero matrix. Thus, two matrix equations are obtained, viz.,

$$AM + c\delta\delta^T N = \mathbf{1}_2, \quad (a)$$

$$AN + c\delta\delta^T M = 0_2. \quad (b)$$

Note that

$$\delta^T A = (1, 1) \begin{pmatrix} 2\Delta' & \frac{1}{2}J_{11} \\ \frac{1}{2}J_{11} & 2\Delta' \end{pmatrix} = (a, a) = a\delta^T, \quad \delta^T \delta = (1, 1) \begin{pmatrix} 1 \\ 1 \end{pmatrix} = 2.$$

The equation (a) can be left-multiplied by δ^T ,

$$\delta^T = \delta^T \mathbf{1}_2 = \delta^T AM + c\delta^T \delta \delta^T N = a\delta^T M + 2c\delta^T N,$$

so does the equation (b),

$$0_2 = \delta^T 0_2 = \delta^T AN + c\delta^T \delta \delta^T M = a\delta^T N + 2c\delta^T M.$$

In other words, we obtain a new equation.

$$\begin{pmatrix} a & 2c \\ 2c & a \end{pmatrix} \begin{pmatrix} \delta^T M \\ \delta^T N \end{pmatrix} = \begin{pmatrix} \delta^T \\ \mathbf{0} \end{pmatrix}$$

The determinant of the coefficient matrix is

$$a^2 - 4c^2 = (a + 2c)(a - 2c) = 2\Delta(a - 2c) = 2\Delta \left[2(\varepsilon_2 - \varepsilon_1) + \frac{1}{2}J_{11} + J_{22} \right] \neq 0,$$

thus the coefficient matrix is reversible. With the truth that for a general 2×2 matrix \mathbf{C} , if its determinant $|\mathbf{C}| \neq 0$, its inverse

$$\mathbf{C}^{-1} = \frac{1}{|\mathbf{C}|} \begin{pmatrix} c_{22} & -c_{12} \\ -c_{21} & c_{11} \end{pmatrix},$$

we find that

$$\begin{pmatrix} \delta^T M \\ \delta^T N \end{pmatrix} = \frac{1}{a^2 - 4c^2} \begin{pmatrix} a & -2c \\ -2c & a \end{pmatrix} \begin{pmatrix} \delta^T \\ \mathbf{0} \end{pmatrix} = \frac{1}{a^2 - 4c^2} \begin{pmatrix} a\delta^T \\ -2c\delta^T \end{pmatrix}.$$

Now we know that

$$\begin{aligned} \delta^T M &= \frac{a}{a^2 - 4c^2} \delta^T, \\ \delta^T N &= -\frac{2c}{a^2 - 4c^2} \delta^T. \end{aligned}$$

Hence, we take it into the equation (a),

$$\begin{aligned} M &= A^{-1}(\mathbf{1}_2 - c\delta\delta^T N) \\ &= \frac{1}{(2\Delta')^2 - (\frac{1}{2}J_{11})^2} \begin{pmatrix} 2\Delta' & -\frac{1}{2}J_{11} \\ -\frac{1}{2}J_{11} & 2\Delta' \end{pmatrix} \left[\begin{pmatrix} 1 & 0 \\ 0 & 1 \end{pmatrix} - c \times \frac{-2c}{a^2 - 4c^2} \begin{pmatrix} 1 & 1 \\ 1 & 1 \end{pmatrix} \right] \\ &= \frac{1}{a(a - J_{11})} \begin{pmatrix} 2\Delta' & -\frac{1}{2}J_{11} \\ -\frac{1}{2}J_{11} & 2\Delta' \end{pmatrix} \times \frac{1}{2\Delta(a - 2c)} \begin{pmatrix} a^2 - 2c^2 & 2c^2 \\ 2c^2 & a^2 - 2c^2 \end{pmatrix} \\ &= \frac{1}{2\Delta(a - 2c)(a - J_{11})} \begin{pmatrix} 2(\Delta'a - c^2) & 2c^2 - \frac{1}{2}J_{11}a \\ 2c^2 - \frac{1}{2}J_{11}a & 2(\Delta'a - c^2) \end{pmatrix} \end{aligned}$$

and the equation (b),

$$N = -A^{-1}c\delta\delta^T M = -c(A^{-1}\delta)(\delta^T M) = -c \frac{a - J_{11}}{a(a - J_{11})} \delta \times \frac{a}{a^2 - 4c^2} \delta^T = -\frac{c}{a^2 - 4c^2} \begin{pmatrix} 1 & 1 \\ 1 & 1 \end{pmatrix}.$$

In conclusion, the inverse of \mathbf{D} is

$$\mathbf{D}^{-1} = \begin{pmatrix} M & N \\ N & M \end{pmatrix} = \frac{1}{a^2 - 4c^2} \begin{pmatrix} \frac{2(\Delta'a - c^2)}{a - J_{11}} & \frac{2c^2 - \frac{1}{2}J_{11}a}{a - J_{11}} & -c & -c \\ \frac{2c^2 - \frac{1}{2}J_{11}a}{a - J_{11}} & \frac{2(\Delta'a - c^2)}{a - J_{11}} & -c & -c \\ -c & -c & \frac{2(\Delta'a - c^2)}{a - J_{11}} & \frac{2c^2 - \frac{1}{2}J_{11}a}{a - J_{11}} \\ -c & -c & \frac{2c^2 - \frac{1}{2}J_{11}a}{a - J_{11}} & \frac{2(\Delta'a - c^2)}{a - J_{11}} \end{pmatrix}.$$

Now, it is turn to calculate the correlation energy, note that $a + 2c = \Delta$ and $2\Delta' - \frac{1}{2}J_{11} = a - J_{11}$, we find that

$$\begin{aligned} E_{\text{corr}}(\mathbf{L} - \text{CCA}(D)) &= -\mathbf{B}^\dagger \mathbf{D} \mathbf{B} \\ &= -\frac{1}{a^2 - 4c^2} \begin{pmatrix} \frac{K_{12}}{\sqrt{2}}, \frac{K_{12}}{\sqrt{2}}, \frac{K_{12}}{\sqrt{2}}, \frac{K_{12}}{\sqrt{2}} \end{pmatrix} \begin{pmatrix} \frac{2(\Delta'a - c^2)}{a - J_{11}} & \frac{2c^2 - \frac{1}{2}J_{11}a}{a - J_{11}} & -c & -c \\ \frac{2c^2 - \frac{1}{2}J_{11}a}{a - J_{11}} & \frac{2(\Delta'a - c^2)}{a - J_{11}} & -c & -c \\ -c & -c & \frac{2(\Delta'a - c^2)}{a - J_{11}} & \frac{2c^2 - \frac{1}{2}J_{11}a}{a - J_{11}} \\ -c & -c & \frac{2c^2 - \frac{1}{2}J_{11}a}{a - J_{11}} & \frac{2(\Delta'a - c^2)}{a - J_{11}} \end{pmatrix} \begin{pmatrix} \frac{K_{12}}{\sqrt{2}} \\ \frac{K_{12}}{\sqrt{2}} \\ \frac{K_{12}}{\sqrt{2}} \\ \frac{K_{12}}{\sqrt{2}} \end{pmatrix} \\ &= -\frac{1}{a^2 - 4c^2} \left(\frac{K_{12}}{\sqrt{2}} \right)^2 \left[-8c + 4 \times \frac{2(\Delta'a - c^2)}{a - J_{11}} + 4 \times \frac{2c^2 - \frac{1}{2}J_{11}a}{a - J_{11}} \right] \\ &= -\frac{1}{2\Delta(a - 2c)} \times \frac{K_{12}^2}{2} \left[-8c + 4 \times \frac{2\Delta'a - 2c^2 + 2c^2 - \frac{1}{2}J_{11}a}{a - J_{11}} \right] \\ &= -\frac{K_{12}^2}{4\Delta(a - 2c)} \times 4 \left(-2c + \frac{(2\Delta' - J_{11})a}{a - J_{11}} \right) \\ &= -\frac{K_{12}^2}{\Delta(a - 2c)} \times (a - 2c) = -\frac{K_{12}^2}{\Delta}. \end{aligned}$$

This states that one gets the same correlation energy using delocalized orbitals.

5.2.4 Some Illustrative Calculations

5.3 Many-Electron Theories with Single Particle Hamiltonians

Exercise 5.13

For the 2×2 matrix

$$\begin{pmatrix} \mathbf{H}_{AA} & \mathbf{H}_{AB} \\ \mathbf{H}_{BA} & \mathbf{H}_{BB} \end{pmatrix} \equiv \begin{pmatrix} H_{11} & H_{12} \\ H_{21} & H_{22} \end{pmatrix}$$

where $H_{11} < H_{22}$ and $H_{12} > 0$. Equation (5.96) simplifies to

$$E_R = \varepsilon_1 - H_{11} = H_{12}C$$

and

$$H_{21} + H_{22}C - CH_{11} - C^2H_{12} = 0.$$

Solve this quadratic equation for the lowest C and then show that ε_1 thus obtained is the lowest eigenvalue of the matrix.

Solution 5.13

The quadratic equation is

$$H_{12}C^2 + (H_{11} - H_{22})C - H_{21} = 0,$$

and its discriminant Δ_C equals

$$(H_{11} - H_{22})^2 - 4 \times H_{12} \times (-H_{21}) = (H_{11} - H_{22})^2 + 4H_{12}H_{21} > 0.$$

Thus, we obtain that the lowest root is

$$C = \frac{1}{2H_{12}} \left[H_{22} - H_{11} - \sqrt{(H_{11} - H_{22})^2 + 4H_{12}H_{21}} \right],$$

and the relaxation energy is

$$E_R = \varepsilon_1 - H_{11} = H_{12}C = \frac{1}{2} \left[H_{22} - H_{11} - \sqrt{(H_{11} - H_{22})^2 + 4H_{12}H_{21}} \right].$$

Thus, we obtain that

$$\varepsilon_1 = H_{11} + H_{12}C = \frac{1}{2} \left[H_{11} + H_{22} - \sqrt{(H_{11} - H_{22})^2 + 4H_{12}H_{21}} \right].$$

For the 2×2 matrix, its eigen polynomial is

$$\det(\mathbf{H} - \varepsilon \mathbf{1}) = \begin{vmatrix} H_{11} - \varepsilon & H_{12} \\ H_{21} & H_{22} - \varepsilon \end{vmatrix} = (\varepsilon - H_{11})(\varepsilon - H_{22}) - H_{12}H_{21},$$

which is a quadratic equation of ε , viz.,

$$\varepsilon^2 - (H_{11} + H_{22})\varepsilon + (H_{11}H_{22} - H_{12}H_{21}) = 0.$$

Its discriminant Δ_ε equals

$$\Delta_\varepsilon = (H_{11} + H_{22})^2 - 4 \times 1 \times (H_{11}H_{22} - H_{12}H_{21}) = (H_{11} - H_{22})^2 + 4H_{12}H_{21} > 0,$$

and the lowest root of is

$$\frac{1}{2} \left[(H_{11} + H_{22}) - \sqrt{(H_{11} - H_{22})^2 + 4H_{12}H_{21}} \right],$$

which equals ε_1 obtained by solving (5.96) and (5.98).

In fact, for a general square matrix \mathbf{A} , if it has an eigenvector a which belongs to the eigenvalue λ , for a general constant c , $\mathbf{A} + c\mathbf{1}$ will have the same eigenvector a which belongs to the eigenvalue $\lambda + c$, and vice versa, viz.,

$$\mathbf{A}a = \lambda a \Leftrightarrow (\mathbf{A} + c\mathbf{1})a = \mathbf{A}a + ca = \lambda a + ca = (\lambda + c)a,$$

Thus, the lowest eigenvalue of $\mathbf{A} + c\mathbf{1}$ is the sum of c and the lowest eigenvalue of \mathbf{A} .

5.3.1 The Relaxation Energy via CI, IEPA, CEPA, and CCA

Exercise 5.14

Show that

a. $\langle \Psi_0 | \mathcal{H} | \Psi_b^s \rangle = v_{bs}$.

b. $\langle \Psi_a^r | \mathcal{H} | \Psi_0 \rangle = v_{ra}$.

c. $\langle \Psi_a^r | \mathcal{H} - E_0 | \Psi_b^s \rangle = \begin{cases} 0, & \text{if } a \neq b, r \neq s, \\ v_{rs}, & \text{if } a = b, r \neq s, \\ -v_{ba}, & \text{if } a \neq b, r = s, \\ \varepsilon_r^{(0)} + v_{rr} - \varepsilon_a^{(0)} - v_{aa} & \text{if } a = b, r = s. \end{cases}$

d. $\langle \Psi_a^r | \mathcal{H} | \Psi_{ab}^{rs} \rangle = \begin{cases} v_{bs}, & \text{if } a \neq b, r \neq s, \\ 0, & \text{otherwise.} \end{cases}$

Solution 5.14

a. With (5.72) and (5.75),

$$\begin{aligned} \langle \Psi_0 | \mathcal{H} | \Psi_b^s \rangle &= \langle \Psi_0 | \mathcal{H}_0 | \Psi_b^s \rangle + \langle \Psi_0 | \mathcal{V} | \Psi_b^s \rangle \\ &= \left(\sum_a \varepsilon_a^{(0)} \right) \langle \Psi_0 | \Psi_b^s \rangle + \langle \Psi_0 | \sum_{i=1}^N v(i) | \Psi_b^s \rangle = 0 + \langle \chi_b^{(0)} | v | \chi_s^{(0)} \rangle = v_{bs}. \end{aligned}$$

Thus, we obtain that

$$\langle \Psi_0 | \mathcal{H} | \Psi_b^s \rangle = v_{bs}. \quad (5.14-1)$$

b. Similarly, we find that

$$\langle \Psi_a^r | \mathcal{H} | \Psi_0 \rangle = \langle \Psi_a^r | \Psi_0 \rangle \sum_a \varepsilon_a^{(0)} + \langle \Psi_a^r | \sum_{i=1}^N v(i) | \Psi_0 \rangle = 0 + \langle \chi_r^{(0)} | v | \chi_a^{(0)} \rangle = v_{ra}. \quad (5.14-2)$$

c. Firstly, we inspect $\langle \Psi_a^r | \mathcal{H} | \Psi_a^r \rangle$. With (5.73), (5.74), we know that

$$\begin{aligned} \langle \Psi_a^r | \mathcal{H} | \Psi_a^r \rangle &= \langle \Psi_a^r | \mathcal{H}_0 | \Psi_a^r \rangle + \langle \Psi_a^r | \mathcal{V} | \Psi_a^r \rangle = \left(\sum_{b \neq a} \varepsilon_b + \varepsilon_r \right) \langle \Psi_a^r | \Psi_a^r \rangle + \sum_{b \neq a} v_{bb} + v_{rr} \\ &= \sum_b \varepsilon_b + \varepsilon_r - \varepsilon_a + \sum_b v_{bb} - v_{aa} + v_{rr} = E_0 + \varepsilon_r^{(0)} + v_{rr} - \varepsilon_a^{(0)} - v_{aa}. \end{aligned}$$

Thus,

$$\langle \Psi_a^r | \mathcal{H} - E_0 | \Psi_a^r \rangle = \langle \Psi_a^r | \mathcal{H} | \Psi_a^r \rangle - E_0 = \varepsilon_r^{(0)} + v_{rr} - \varepsilon_a^{(0)} - v_{aa}.$$

Then, we inspect $\langle \Psi_a^r | \mathcal{H} | \Psi_a^s \rangle$. When $r \neq s$, we get that

$$\langle \Psi_a^r | \mathcal{H} | \Psi_a^s \rangle = \langle \Psi_a^r | \mathcal{H}_0 | \Psi_a^s \rangle + \langle \Psi_a^r | \mathcal{V} | \Psi_a^s \rangle = \left(\sum_{b \neq a} \varepsilon_b + \varepsilon_r \right) \langle \Psi_a^r | \Psi_a^s \rangle + v_{rs} = v_{rs}.$$

Similarly, when $a \neq b$, we find that

$$\langle \Psi_a^r | \mathcal{H} | \Psi_b^r \rangle = \langle \Psi_a^r | \mathcal{H}_0 | \Psi_b^r \rangle + \langle \Psi_a^r | \mathcal{V} | \Psi_b^r \rangle = \left(\sum_{c \neq a} \varepsilon_c + \varepsilon_r \right) \langle \Psi_a^r | \Psi_b^r \rangle - v_{ba} = -v_{ba}.$$

Otherwise, the matrix element of a Hamiltonian between Ψ_a^r and Ψ_b^s vanishes as the number of their different spin orbitals is more than one ($r \neq s$ and $a \neq b$). In conclusion, we find that

$$\langle \Psi_a^r | \mathcal{H} - E_0 | \Psi_b^s \rangle = \begin{cases} 0, & \text{if } a \neq b, r \neq s, \\ v_{rs}, & \text{if } a = b, r \neq s, \\ -v_{ba}, & \text{if } a \neq b, r = s, \\ \varepsilon_r^{(0)} + v_{rr} - \varepsilon_a^{(0)} - v_{aa} & \text{if } a = b, r = s. \end{cases} \quad (5.14-3)$$

This conclusion can be reached using the results of Exercise 2.13, too.

d . As $a = b$ or $r = s$, Ψ_{ab}^{rs} vanishes and thus $\langle \Psi_a^r | \mathcal{H} | \Psi_{ab}^{rs} \rangle = 0$. Otherwise, namely, $a \neq b$ and $r \neq s$, the number of their different spin orbitals is one, at this time, $\langle \Psi_a^r | \mathcal{H} | \Psi_{ab}^{rs} \rangle = \langle \Psi_0 | \mathcal{H} | \Psi_b^s \rangle = v_{bs}$. Finally, we get that

$$\langle \Psi_a^r | \mathcal{H} | \Psi_{ab}^{rs} \rangle = \begin{cases} v_{bs}, & \text{if } a \neq b, r \neq s, \\ 0, & \text{otherwise.} \end{cases} \quad (5.14-4)$$

Exercise 5.15

Repeat the above analysis using

$$\begin{aligned} |\chi_1\rangle &= a_1|\chi_1^{(0)}\rangle + a_2|\chi_2^{(0)}\rangle + a_3|\chi_3^{(0)}\rangle + a_4|\chi_4^{(0)}\rangle, \\ |\chi_2\rangle &= b_1|\chi_1^{(0)}\rangle + b_2|\chi_2^{(0)}\rangle + b_3|\chi_3^{(0)}\rangle + b_4|\chi_4^{(0)}\rangle \end{aligned}$$

instead of Eqs.(5.121a,b).

a. By repeated use of Eq.(1.40) show that

$$\begin{aligned} |\Phi_0\rangle &= (a_1b_2 - b_1a_2)|\chi_1^{(0)}\chi_2^{(0)}\rangle + (a_1b_3 - b_1a_3)|\chi_1^{(0)}\chi_3^{(0)}\rangle + (a_1b_4 - b_1a_4)|\chi_1^{(0)}\chi_4^{(0)}\rangle \\ &\quad + (a_3b_2 - b_3a_2)|\chi_3^{(0)}\chi_2^{(0)}\rangle + (a_4b_2 - b_4a_2)|\chi_4^{(0)}\chi_2^{(0)}\rangle + (a_3b_4 - b_3a_4)|\chi_3^{(0)}\chi_4^{(0)}\rangle. \end{aligned}$$

Intermediate normalize this wave function by dividing the right-hand side by $a_1b_2 - b_1a_2$ and then explicitly verify that Eq.(5.123) is satisfied.

b. To make contact with the general formalism, note that

$$\mathbf{U}_{AA} = \begin{pmatrix} a_1 & b_1 \\ a_2 & b_2 \end{pmatrix}, \quad \mathbf{U}_{BA} = \begin{pmatrix} a_3 & b_3 \\ a_4 & b_4 \end{pmatrix}.$$

Note that $|\mathbf{U}_{AA}| = a_1b_2 - b_1a_2$ as required to make Eq.(5.120a) consistent with the result obtained in part (a). Use the result of Exercise 1.4(f) to evaluate \mathbf{U}_{AA}^{-1} and then verify the general result given in Eq.(5.120b) by calculating

$$(\mathbf{U}_{BA}\mathbf{U}_{AA}^{-1})_{11} = c_1^3$$

and showing that it is identical to the coefficient of $|\chi_3^{(0)}\chi_2^{(0)}\rangle$ obtained in part (a).

Solution 5.15

a. Using (1.40) and $|\chi_i^{(0)}\chi_i^{(0)}\rangle = 0$ for $i = 1, 2, 3, 4$, we find that

$$\begin{aligned} |\Phi_0\rangle &= (a_1\chi_1^{(0)} + a_2\chi_2^{(0)} + a_3\chi_3^{(0)} + a_4\chi_4^{(0)})(b_1\chi_1^{(0)} + b_2\chi_2^{(0)} + b_3\chi_3^{(0)} + b_4\chi_4^{(0)}) \\ &= a_1b_2|\chi_1^{(0)}\chi_2^{(0)}\rangle + a_1b_3|\chi_1^{(0)}\chi_3^{(0)}\rangle + a_1b_4|\chi_1^{(0)}\chi_4^{(0)}\rangle + a_2b_1|\chi_2^{(0)}\chi_1^{(0)}\rangle \\ &\quad + a_2b_3|\chi_2^{(0)}\chi_3^{(0)}\rangle + a_2b_4|\chi_2^{(0)}\chi_4^{(0)}\rangle + a_3b_1|\chi_3^{(0)}\chi_1^{(0)}\rangle + a_3b_2|\chi_3^{(0)}\chi_2^{(0)}\rangle \\ &\quad + a_3b_4|\chi_3^{(0)}\chi_4^{(0)}\rangle + a_4b_1|\chi_4^{(0)}\chi_1^{(0)}\rangle + a_4b_2|\chi_4^{(0)}\chi_2^{(0)}\rangle + a_4b_3|\chi_4^{(0)}\chi_3^{(0)}\rangle \\ &= (a_1b_2 - a_2b_1)|\chi_1^{(0)}\chi_2^{(0)}\rangle + (a_1b_3 - a_3b_1)|\chi_1^{(0)}\chi_3^{(0)}\rangle + (a_1b_4 - a_4b_1)|\chi_1^{(0)}\chi_4^{(0)}\rangle \\ &\quad + (a_2b_3 - a_3b_2)|\chi_2^{(0)}\chi_3^{(0)}\rangle + (a_2b_4 - a_4b_2)|\chi_2^{(0)}\chi_4^{(0)}\rangle + (a_3b_4 - a_4b_3)|\chi_3^{(0)}\chi_4^{(0)}\rangle \\ &= (a_1b_2 - a_2b_1)|\Psi_0\rangle + (a_1b_3 - a_3b_1)|\Phi_2^3\rangle + (a_1b_4 - a_4b_1)|\Phi_2^4\rangle \\ &\quad - (a_2b_3 - a_3b_2)|\Psi_1^3\rangle - (a_2b_4 - a_4b_2)|\Psi_1^4\rangle + (a_3b_4 - a_4b_3)|\Psi_{12}^{34}\rangle. \end{aligned}$$

After the intermediate normalization (namely, $|\Phi_0\rangle$ divided by $a_1b_2 - a_2b_1$), we get that

$$\begin{aligned} |\Phi'_0\rangle &= \frac{1}{a_1b_2 - a_2b_1}|\Phi_0\rangle \\ &= |\Psi_0\rangle + \frac{1}{a_1b_2 - a_2b_1} [(a_1b_3 - a_3b_1)|\Phi_2^3\rangle + (a_1b_4 - a_4b_1)|\Phi_2^4\rangle \\ &\quad - (a_2b_3 - a_3b_2)|\Psi_1^3\rangle - (a_2b_4 - a_4b_2)|\Psi_1^4\rangle + (a_3b_4 - a_4b_3)|\Psi_{12}^{34}\rangle]. \end{aligned}$$

Thus,

$$\begin{aligned} c_2^3 &= \frac{a_1 b_3 - a_3 b_1}{a_1 b_2 - a_2 b_1}, & c_2^4 &= \frac{a_1 b_4 - a_4 b_1}{a_1 b_2 - a_2 b_1}, & c_1^3 &= -\frac{a_2 b_3 - a_3 b_2}{a_1 b_2 - a_2 b_1}, \\ c_1^4 &= -\frac{a_2 b_4 - a_4 b_2}{a_1 b_2 - a_2 b_1}, & c_{12}^{34} &= \frac{a_3 b_4 - a_4 b_3}{a_1 b_2 - a_2 b_1}. \end{aligned}$$

We can find that

$$\begin{aligned} c_1^3 c_2^4 - c_1^4 c_2^3 &= \left(-\frac{a_2 b_3 - a_3 b_2}{a_1 b_2 - a_2 b_1} \right) \left(\frac{a_1 b_4 - a_4 b_1}{a_1 b_2 - a_2 b_1} \right) - \left(-\frac{a_2 b_4 - a_4 b_2}{a_1 b_2 - a_2 b_1} \right) \left(\frac{a_1 b_3 - a_3 b_1}{a_1 b_2 - a_2 b_1} \right) \\ &= -\frac{a_2 a_3 b_1 b_4 + a_1 a_4 b_2 b_3 - a_2 a_4 b_1 b_3 - a_1 a_3 b_2 b_4}{(a_1 b_2 - a_2 b_1)^2} = -\frac{(a_1 b_2 - a_2 b_1)(a_4 b_3 - a_3 b_4)}{(a_1 b_2 - a_2 b_1)^2} \\ &= -\frac{a_4 b_3 - a_3 b_4}{a_1 b_2 - a_2 b_1} = \frac{a_3 b_4 - a_4 b_3}{a_1 b_2 - a_2 b_1} = c_{12}^{34}. \end{aligned}$$

b. The inverse of \mathbf{U}_{AA} is

$$\mathbf{U}_{AA}^{-1} = \frac{1}{|\mathbf{U}_{AA}|} \begin{pmatrix} b_2 & -b_1 \\ -a_2 & a_1 \end{pmatrix} = \frac{1}{a_1 b_2 - b_1 a_2} \begin{pmatrix} b_2 & -b_1 \\ -a_2 & a_1 \end{pmatrix},$$

and thus

$$\mathbf{U}_{BA} \mathbf{U}_{AA}^{-1} = \begin{pmatrix} a_3 & b_3 \\ a_4 & b_4 \end{pmatrix} \times \frac{1}{a_1 b_2 - b_1 a_2} \begin{pmatrix} b_2 & -b_1 \\ -a_2 & a_1 \end{pmatrix} = \frac{1}{a_1 b_2 - b_1 a_2} \begin{pmatrix} a_3 b_2 - a_2 b_3 & a_1 b_3 - a_3 b_1 \\ a_4 b_2 - a_2 b_4 & a_1 b_4 - a_4 b_1 \end{pmatrix}.$$

We know that

$$(\mathbf{U}_{BA} \mathbf{U}_{AA}^{-1})_{11} = \frac{a_3 b_2 - a_2 b_3}{a_1 b_2 - b_1 a_2} = c_1^3. \quad (5.15-1)$$

It is identical to the coefficient of $|\chi_3^{(0)} \chi_2^{(0)}\rangle$ obtained in part (a).

5.3.2 The Resonance Energy of Polyenes in Hückel Theory

Exercise 5.16

Set up the Hückel matrix for benzene and find its eigenvalues. Remember that if the carbon atoms are labeled clockwise from 1 to 6, then atoms 1 and 6 are nearest neighbors. Show that the six eigenvalues are identical to those given by Eq.(5.131). Find the total energy and compare it with the result given by Eq.(5.132).

Solution 5.16

The Hückel matrix for benzene is

$$\mathbf{H} = \begin{pmatrix} \alpha & \beta & 0 & 0 & 0 & \beta \\ \beta & \alpha & \beta & 0 & 0 & 0 \\ 0 & \beta & \alpha & \beta & 0 & 0 \\ 0 & 0 & \beta & \alpha & \beta & 0 \\ 0 & 0 & 0 & \beta & \alpha & \beta \\ \beta & 0 & 0 & 0 & \beta & \alpha \end{pmatrix},$$

whose eigen polynomial is a sixth-degree equation in one variable ε , viz.,

$$\det(\mathbf{H} - \varepsilon \mathbf{1}) = (\alpha - \varepsilon - 2\beta)(\alpha - \varepsilon - \beta)^2(\alpha - \varepsilon + \beta)^2(\alpha - \varepsilon + 2\beta) = 0.$$

This equation has six roots,

$$\varepsilon_1 = \alpha + 2\beta, \quad \varepsilon_2 = \varepsilon_3 = \alpha + \beta, \quad \varepsilon_4 = \varepsilon_5 = \alpha - \beta, \quad \varepsilon_6 = \alpha - 2\beta.$$

And thus

$$\mathcal{E}_0 = \sum_{i=1}^{6/2} \varepsilon_i = \varepsilon_1 + \varepsilon_2 + \varepsilon_3 = 6\alpha + 8\beta. \quad (5.16-1)$$

Note that the cosine function is an even function. Using Eq.(5.131), we obtain

$$\begin{aligned} \varepsilon_0 &= \alpha + 2\beta \cos \frac{0 \times \pi}{3} = \alpha + 2\beta \times 1 = \alpha + 2\beta, \\ \varepsilon_{\pm 1} &= \alpha + 2\beta \cos \frac{\pm 1 \times \pi}{3} = \alpha + 2\beta \times \frac{1}{2} = \alpha + \beta, \\ \varepsilon_{\pm 2} &= \alpha + 2\beta \cos \frac{\pm 2 \times \pi}{3} = \alpha + 2\beta \times \left(-\frac{1}{2}\right) = \alpha - \beta, \\ \varepsilon_3 &= \alpha + 2\beta \cos \frac{3 \times \pi}{3} = \alpha + 2\beta \times (-1) = \alpha - 2\beta, \end{aligned}$$

and

$$\mathcal{E}_0 = 6\alpha + \frac{4\beta}{\sin \frac{\pi}{6}} = 6\alpha + \frac{4\beta}{\frac{1}{2}} = 6\alpha + 8\beta.$$

It is evident that they are identical to my results before.

Exercise 5.17

Verify Eqs.(5.139) and (5.140).

Solution 5.17

The verification of all equations is as follows.

- For (5.139a):

$$\begin{aligned} \langle i|j \rangle &= \frac{1}{2} [\langle \phi_{2i-1} | \phi_{2j-1} \rangle + \langle \phi_{2i-1} | \phi_{2j} \rangle + \langle \phi_{2i} | \phi_{2j-1} \rangle + \langle \phi_{2i} | \phi_{2j} \rangle] = \frac{1}{2} [\delta_{ij} + 0 + 0 + \delta_{ij}] = \delta_{ij}, \\ \langle i^* | j^* \rangle &= \frac{1}{2} [\langle \phi_{2i-1} | \phi_{2j-1} \rangle - \langle \phi_{2i-1} | \phi_{2j} \rangle - \langle \phi_{2i} | \phi_{2j-1} \rangle + \langle \phi_{2i} | \phi_{2j} \rangle] = \frac{1}{2} [\delta_{ij} - 0 - 0 + \delta_{ij}] = \delta_{ij}. \end{aligned}$$

- For (5.139b):

$$\begin{aligned} \langle i | j^* \rangle &= \frac{1}{2} [\langle \phi_{2i-1} | \phi_{2j-1} \rangle - \langle \phi_{2i-1} | \phi_{2j} \rangle + \langle \phi_{2i} | \phi_{2j-1} \rangle - \langle \phi_{2i} | \phi_{2j} \rangle] = \frac{1}{2} [\delta_{ij} - 0 + 0 - \delta_{ij}] = 0, \\ \langle i^* | j \rangle &= (\langle i | j^* \rangle)^* = 0^* = 0. \end{aligned}$$

- For (5.140a):

$$\begin{aligned} \langle i | h_{\text{eff}} | i \rangle &= \frac{1}{2} [\langle \phi_{2i-1} | h_{\text{eff}} | \phi_{2i-1} \rangle + \langle \phi_{2i-1} | h_{\text{eff}} | \phi_{2i} \rangle + \langle \phi_{2i} | h_{\text{eff}} | \phi_{2i-1} \rangle + \langle \phi_{2i} | h_{\text{eff}} | \phi_{2i} \rangle] \\ &= \frac{1}{2} [\alpha + \beta + \beta + \alpha] = \alpha + \beta. \end{aligned}$$

- For (5.140b):

$$\begin{aligned} \langle i^* | h_{\text{eff}} | i^* \rangle &= \frac{1}{2} [\langle \phi_{2i-1} | h_{\text{eff}} | \phi_{2i-1} \rangle - \langle \phi_{2i-1} | h_{\text{eff}} | \phi_{2i} \rangle - \langle \phi_{2i} | h_{\text{eff}} | \phi_{2i-1} \rangle + \langle \phi_{2i} | h_{\text{eff}} | \phi_{2i} \rangle] \\ &= \frac{1}{2} [\alpha - \beta - \beta + \alpha] = \alpha - \beta. \end{aligned}$$

- For (5.140c):

$$\begin{aligned} \langle i | h_{\text{eff}} | i - 1 \rangle &= \frac{1}{2} [\langle \phi_{2i-1} | h_{\text{eff}} | \phi_{2i-3} \rangle + \langle \phi_{2i-1} | h_{\text{eff}} | \phi_{2i-2} \rangle + \langle \phi_{2i} | h_{\text{eff}} | \phi_{2i-3} \rangle + \langle \phi_{2i} | h_{\text{eff}} | \phi_{2i-2} \rangle] \\ &= \frac{1}{2} [0 + \beta + 0 + 0] = \frac{1}{2}\beta, \end{aligned}$$

$$\begin{aligned}\langle i|h_{\text{eff}}|i+1\rangle &= \frac{1}{2} [\langle \phi_{2i-1}|h_{\text{eff}}|\phi_{2i+1}\rangle + \langle \phi_{2i-1}|h_{\text{eff}}|\phi_{2i+2}\rangle + \langle \phi_{2i}|h_{\text{eff}}|\phi_{2i+1}\rangle + \langle \phi_{2i}|h_{\text{eff}}|\phi_{2i+2}\rangle] \\ &= \frac{1}{2} [0 + 0 + \beta + 0] = \frac{1}{2}\beta.\end{aligned}$$

Thus,

$$\langle i|h_{\text{eff}}|i\pm 1\rangle = \frac{\beta}{2}.$$

- For (5.140d):

$$\begin{aligned}\langle i^*|h_{\text{eff}}|(i-1)^*\rangle &= \frac{1}{2} [\langle \phi_{2i-1}|h_{\text{eff}}|\phi_{2i-3}\rangle - \langle \phi_{2i-1}|h_{\text{eff}}|\phi_{2i-2}\rangle - \langle \phi_{2i}|h_{\text{eff}}|\phi_{2i-3}\rangle + \langle \phi_{2i}|h_{\text{eff}}|\phi_{2i-2}\rangle] \\ &= \frac{1}{2} [0 - 0 - \beta + 0] = -\frac{1}{2}\beta, \\ \langle i^*|h_{\text{eff}}|(i+1)^*\rangle &= \frac{1}{2} [\langle \phi_{2i-1}|h_{\text{eff}}|\phi_{2i+1}\rangle - \langle \phi_{2i-1}|h_{\text{eff}}|\phi_{2i+2}\rangle - \langle \phi_{2i}|h_{\text{eff}}|\phi_{2i+1}\rangle + \langle \phi_{2i}|h_{\text{eff}}|\phi_{2i+2}\rangle] \\ &= \frac{1}{2} [0 - 0 - \beta + 0] = -\frac{1}{2}\beta.\end{aligned}$$

Thus,

$$\langle i^*|h_{\text{eff}}|(i\pm 1)^*\rangle = -\frac{\beta}{2}.$$

- For (5.140e):

$$\begin{aligned}\langle i|h_{\text{eff}}|(i-1)^*\rangle &= \frac{1}{2} [\langle \phi_{2i-1}|h_{\text{eff}}|\phi_{2i-3}\rangle - \langle \phi_{2i-1}|h_{\text{eff}}|\phi_{2i-2}\rangle + \langle \phi_{2i}|h_{\text{eff}}|\phi_{2i-3}\rangle - \langle \phi_{2i}|h_{\text{eff}}|\phi_{2i-2}\rangle] \\ &= \frac{1}{2} [0 - \beta + 0 - 0] = -\frac{1}{2}\beta, \\ \langle i|h_{\text{eff}}|(i+1)^*\rangle &= \frac{1}{2} [\langle \phi_{2i-1}|h_{\text{eff}}|\phi_{2i+1}\rangle - \langle \phi_{2i-1}|h_{\text{eff}}|\phi_{2i+2}\rangle + \langle \phi_{2i}|h_{\text{eff}}|\phi_{2i+1}\rangle - \langle \phi_{2i}|h_{\text{eff}}|\phi_{2i+2}\rangle] \\ &= \frac{1}{2} [0 - 0 + \beta - 0] = \frac{1}{2}\beta, \\ \langle (i-1)|h_{\text{eff}}|i^*\rangle &= \frac{1}{2} [\langle \phi_{2i-3}|h_{\text{eff}}|\phi_{2i-1}\rangle - \langle \phi_{2i-3}|h_{\text{eff}}|\phi_{2i}\rangle + \langle \phi_{2i-2}|h_{\text{eff}}|\phi_{2i-1}\rangle - \langle \phi_{2i-2}|h_{\text{eff}}|\phi_{2i}\rangle] \\ &= \frac{1}{2} [0 - 0 + \beta - 0] = \frac{1}{2}\beta, \\ \langle (i+1)|h_{\text{eff}}|i^*\rangle &= \frac{1}{2} [\langle \phi_{2i+1}|h_{\text{eff}}|\phi_{2i-1}\rangle - \langle \phi_{2i+1}|h_{\text{eff}}|\phi_{2i}\rangle + \langle \phi_{2i+2}|h_{\text{eff}}|\phi_{2i-1}\rangle - \langle \phi_{2i+2}|h_{\text{eff}}|\phi_{2i}\rangle] \\ &= \frac{1}{2} [0 - \beta + 0 - 0] = -\frac{1}{2}\beta.\end{aligned}$$

Thus,

$$\langle i|h_{\text{eff}}|(i\pm 1)^*\rangle = \langle (i\mp 1)|h_{\text{eff}}|i^*\rangle = \pm \frac{\beta}{2}.$$

Now we have proved all equations.

Exercise 5.18

Evaluate the matrix elements given in Eq.(5.149) and fill in the remaining steps leading to Eq.(5.150).

Solution 5.18

The verification of (5.149) is easy, viz.,

$$\langle \Psi_0|\mathcal{H}|_1^*\rangle = \langle \Psi_0|\mathcal{H}\left(\frac{1}{\sqrt{2}}|\Psi_1^{2*}\rangle - \frac{1}{\sqrt{2}}|\Psi_1^{3*}\rangle\right) = \frac{1}{\sqrt{2}}\langle \Psi_0|\mathcal{H}|\Psi_1^{2*}\rangle - \frac{1}{\sqrt{2}}\langle \Psi_0|\mathcal{H}|\Psi_1^{3*}\rangle = \frac{\beta}{\sqrt{2}}.$$

And now we pay attention to calculating $\langle {}_1^*|\mathcal{H}|_1^*\rangle$, which is

$$\langle {}_1^*|\mathcal{H}|_1^*\rangle = \left(\frac{1}{\sqrt{2}}\langle \Psi_1^{2*}| - \frac{1}{\sqrt{2}}\langle \Psi_1^{3*}| \right) \mathcal{H} \left(\frac{1}{\sqrt{2}}|\Psi_1^{2*}\rangle - \frac{1}{\sqrt{2}}|\Psi_1^{3*}\rangle\right)$$

$$\begin{aligned}
&= \frac{1}{2} (\langle \Psi_1^{2*} | \mathcal{H} | \Psi_1^{2*} \rangle - \langle \Psi_1^{2*} | \mathcal{H} | \Psi_1^{3*} \rangle - \langle \Psi_1^{3*} | \mathcal{H} | \Psi_1^{2*} \rangle + \langle \Psi_1^{3*} | \mathcal{H} | \Psi_1^{3*} \rangle) \\
&= \frac{1}{2} \left(E_0 + \varepsilon_{2*}^{(0)} - \varepsilon_1^{(0)} - \left(-\frac{\beta}{2} \right) - \left(-\frac{\beta}{2} \right) + E_0 + \varepsilon_{2*}^{(0)} - \varepsilon_1^{(0)} \right) \\
&= E_0 + \varepsilon_{2*}^{(0)} - \varepsilon_1^{(0)} + \frac{\beta}{2} = E_0 + (\alpha - \beta) - (\alpha + \beta) + \frac{\beta}{2} = E_0 - \frac{3\beta}{2}.
\end{aligned}$$

Thus, we find that

$$\langle 1^* | \mathcal{H} - E_0 | 1^* \rangle = \langle 1^* | \mathcal{H} | 1^* \rangle - E_0 = E_0 - \frac{3\beta}{2} - E_0 = -\frac{3\beta}{2}.$$

Finally, from (5.148), we obtain the corresponding “particle” equations, viz.,

$$\frac{\beta}{\sqrt{2}} c = e_1, \quad (a)$$

$$\frac{\beta}{\sqrt{2}} - \frac{3\beta}{2} c = e_1 c. \quad (b)$$

The equation (b) can be subtracted by the equation (a) multiplied by c , thus we obtain a quadratic equation of c , viz.,

$$\frac{\beta}{\sqrt{2}} c^2 + \frac{3\beta}{2} c - \frac{\beta}{\sqrt{2}} = 0.$$

Its discriminant is

$$\Delta_c = \left(\frac{3\beta}{2} \right)^2 - 4 \times \frac{\beta}{\sqrt{2}} \times \left(-\frac{\beta}{\sqrt{2}} \right) = \frac{17\beta^2}{4},$$

and the root of c are

$$c_{\pm} = \frac{-\frac{3\beta}{2} \mp \frac{\sqrt{17}}{2}\beta}{2 \times \frac{\beta}{\sqrt{2}}} = -\frac{3 \pm \sqrt{17}}{2\sqrt{2}}.$$

Here, note that β is negative, and thus $\sqrt{\beta^2} = -\beta!$ Thus, from the equation (a), we know that

$$e_{\pm} = \frac{\beta}{\sqrt{2}} c_{\pm} = -\frac{3 \pm \sqrt{17}}{4} \beta = \frac{\mp \sqrt{17} - 3}{4} \beta$$

In fact, only the minus root makes physical sense as the relaxation energy is negative, hence,

$$e_1 = e_- = \frac{-3 + \sqrt{17}}{4} \beta \approx 0.2808\beta,$$

and

$$E_R(\text{IEPA}) = 6e_1 = 6 \times 0.2808\beta = 1.6848\beta \approx 1.685\beta, \quad (5.18-1)$$

which is (5.150).

Exercise 5.19

- a) Extend the above analysis to calculate the IEPA resonance energy for a cyclic polyene with $N = 2n$ ($N > 6$) carbon atoms. As before, argue that all “particle” energies are the same so that

$$E_R = N e_1.$$

Consider only single excitations that mix with $|\Psi_0\rangle$. Show that the “particle” function $|\Psi_1\rangle$ is

$$|\Psi_1\rangle = |\Psi_0\rangle + c|1^*\rangle,$$

where $|1^*\rangle$ is obtained from Eq.(5.146),

$$|1^*\rangle = \frac{1}{\sqrt{2}} (|\Psi_1^{2*}\rangle - |\Psi_1^{n*}\rangle).$$

Now show that

$$\langle \Psi_0 | \mathcal{H} | 1^* \rangle = \frac{1}{\sqrt{2}} \beta$$

as before, but that here

$$\langle {}^*|_1 \mathcal{H} - E_0 |_1^* \rangle = -2\beta,$$

instead of the result in Eq.(5.149b). Why the difference? Finally, solve the resulting “particle” equations to show that

$$E_R(\text{IEPA}) = N \left(\sqrt{\frac{3}{2}} - 1 \right) \beta = 0.2247N\beta.$$

Note that the IEPA is indeed size consistent and that in the limit of large N it gives 82% of the exact resonance energy.

- b) The above result is not really exact within the IEPA. The reason for this is that there exist single excitations involving orbital $|1\rangle$ that do not mix with $|\Psi_0\rangle$ but do mix with $|_1^*\rangle$ and thus have some effect on the “particle” energy e_1 . These excitations are analogous to single excitations in CI for a real many-particle system in the sense that although single excitations do not mix with the HF wave function because of Brillouin’s theorem, they do mix indirectly through the double excitations. Investigate the effect of such excitations for the case $N = 10$. Show that the exact “particle” function $|\Psi_1\rangle$ is

$$|\Psi_1\rangle = |\Psi_0\rangle + c_1|_1^*\rangle + c_3|\Psi_1^{3*}\rangle + c_4|\Psi_1^{4*}\rangle.$$

Now show that

$$\begin{aligned} \langle \Psi_0 | \mathcal{H} | \Psi_1^{3*} \rangle &= \langle \Psi_0 | \mathcal{H} | \Psi_1^{4*} \rangle = 0, \\ \langle \Psi_1^{3*} | \mathcal{H} - E_0 | \Psi_1^{3*} \rangle &= \langle \Psi_1^{4*} | \mathcal{H} - E_0 | \Psi_1^{4*} \rangle = -2\beta, \\ \langle \Psi_1^{3*} | \mathcal{H} | \Psi_1^{4*} \rangle &= -\frac{1}{2}\beta, \\ \langle {}^*|_1 \mathcal{H} | \Psi_1^{3*} \rangle &= -\frac{1}{2\sqrt{2}}\beta, \\ \langle {}^*|_1 \mathcal{H} | \Psi_1^{4*} \rangle &= \frac{1}{2\sqrt{2}}\beta. \end{aligned}$$

Finally, show from the resulting “particle” equations that e_1 is the solution of

$$4e_1^3 + 14\beta e_1^2 + 9\beta^2 e_1 - 3\beta^3 = 0.$$

The cubic equation can be solved to yield $e_1 = 0.2387\beta$ so that the exact IEPA resonance energy for $N = 10$ is 2.387β , which is to be compared with the approximate result of 2.247β obtained in part (a), so that there is a 6% difference. The exact resonance energy found from Eq.(5.135) is 2.944β for this case.

Solution 5.19

- a) Similar to Exercise 5.18, we get

$$\begin{aligned} \langle \Psi_0 | \mathcal{H} |_1^* \rangle &= \langle \Psi_0 | \mathcal{H} \left(\frac{1}{\sqrt{2}} |\Psi_1^{2*}\rangle - \frac{1}{\sqrt{2}} |\Psi_1^{n*}\rangle \right) = \frac{1}{\sqrt{2}} \langle \Psi_0 | \mathcal{H} | \Psi_1^{2*} \rangle - \frac{1}{\sqrt{2}} \langle \Psi_0 | \mathcal{H} | \Psi_1^{n*} \rangle = \frac{\beta}{\sqrt{2}}. \\ \langle {}^*|_1 \mathcal{H} |_1^* \rangle &= \left(\frac{1}{\sqrt{2}} \langle \Psi_1^{2*} | - \frac{1}{\sqrt{2}} \langle \Psi_1^{n*} | \right) \mathcal{H} \left(\frac{1}{\sqrt{2}} |\Psi_1^{2*}\rangle - \frac{1}{\sqrt{2}} |\Psi_1^{n*}\rangle \right) \\ &= \frac{1}{2} (\langle \Psi_1^{2*} | \mathcal{H} | \Psi_1^{2*} \rangle - \langle \Psi_1^{2*} | \mathcal{H} | \Psi_1^{n*} \rangle - \langle \Psi_1^{n*} | \mathcal{H} | \Psi_1^{2*} \rangle + \langle \Psi_1^{n*} | \mathcal{H} | \Psi_1^{n*} \rangle) \\ &= \frac{1}{2} (E_0 + \varepsilon_{2^*}^{(0)} - \varepsilon_1^{(0)} - 0 - 0 + E_0 + \varepsilon_{2^*}^{(0)} - \varepsilon_1^{(0)}) \\ &= E_0 + \varepsilon_{2^*}^{(0)} - \varepsilon_1^{(0)} + \frac{\beta}{2} = E_0 + (\alpha - \beta) - (\alpha + \beta) = E_0 - 2\beta. \end{aligned}$$

Thus, we find that

$$\langle {}^*|_1 \mathcal{H} - E_0 |_1^* \rangle = \langle {}^*|_1 \mathcal{H} |_1^* \rangle - E_0 = E_0 - 2\beta - E_0 = -2\beta.$$

Hence, we obtain the corresponding “particle” equations, viz.,

$$\frac{\beta}{\sqrt{2}}c = e_1, \quad (a)$$

$$\frac{\beta}{\sqrt{2}} - 2\beta c = e_1 c. \quad (b)$$

The equation (b) can be subtracted by the equation (a) multiplied by c , thus we obtain a quadratic equation of c , viz.,

$$\frac{\beta}{\sqrt{2}}c^2 + 2\beta c - \frac{\beta}{\sqrt{2}} = 0.$$

Its discriminant is

$$\Delta_c = (2\beta)^2 - 4 \times \frac{\beta}{\sqrt{2}} \times \left(-\frac{\beta}{\sqrt{2}}\right) = 6\beta^2,$$

and the root of c are

$$c_{\pm} = \frac{-2\beta \mp \sqrt{6}\beta}{2 \times \frac{\beta}{\sqrt{2}}} = -\sqrt{2} \mp \sqrt{3}.$$

Thus, from the equation (a), we know that

$$e_{\pm} = \frac{\beta}{\sqrt{2}}c_{\pm} = \frac{-\sqrt{2} \mp \sqrt{3}}{\sqrt{2}}\beta = \left(\mp\sqrt{\frac{3}{2}} - 1\right)\beta.$$

In fact, only the minus root makes physical sense as the relaxation energy is negative, hence,

$$e_1 = e_- = \left(\sqrt{\frac{3}{2}} - 1\right)\beta \approx 0.2247\beta,$$

and

$$\lim_{N \rightarrow \infty} \frac{E_R(\text{IEPA})}{N} = e_1 = 0.2247\beta. \quad (5.19-1)$$

In the limit of large N , the propotion of relaxation energy IEPA delivers is

$$\lim_{N \rightarrow \infty} \frac{E_R(\text{IEPA})}{E_R} = \lim_{N \rightarrow \infty} \frac{E_R(\text{IEPA})}{N} \frac{N}{E_R(\text{IEPA})} = \frac{0.2247\beta}{0.2732\beta} = 0.822. \quad (5.19-2)$$

In other words, the IPEA is size consistent and that in the limit of large N it gives 82% of the exact resonance energy. In my opinion, the differences between Exercise 5.18 and Exercise 5.19(a) are that $\langle \Psi_1^{2*} | \mathcal{H} | \Psi_1^{n*} \rangle$ vanishes when $N > 6$. This corresponds the truth that in the Hückel theory, when the larger the system is, the less the proportion of other electrons, which interact with electrons in a given occupied orbital, in all electrons is, as $|\Psi_1\rangle = |\Psi_0\rangle + c|\Psi_1^*\rangle$ described.

- b) When $N = 10$, $n = 5$ and neither the orbital 3 nor the orbital 4 is the nearest-neighbor ones of the orbital 1. Hence,

$$\langle \Psi_0 | \mathcal{H} | \Psi_1^{3*} \rangle = \langle \Psi_0 | \mathcal{H} | \Psi_1^{4*} \rangle = 0.$$

Moreover, other matrix elements are

$$\begin{aligned} \langle \Psi_1^{3*} | \mathcal{H} - E_0 | \Psi_1^{3*} \rangle &= \langle \Psi_1^{3*} | \mathcal{H} | \Psi_1^{3*} \rangle - E_0 = E_0 + \varepsilon_{3*} - \varepsilon_1 - E_0 = \alpha - \beta - (\alpha + \beta) = -2\beta, \\ \langle \Psi_1^{4*} | \mathcal{H} - E_0 | \Psi_1^{4*} \rangle &= \langle \Psi_1^{4*} | \mathcal{H} | \Psi_1^{4*} \rangle - E_0 = E_0 + \varepsilon_{4*} - \varepsilon_1 - E_0 = \alpha - \beta - (\alpha + \beta) = -2\beta, \\ \langle \Psi_1^{3*} | \mathcal{H} | \Psi_1^{4*} \rangle &= \langle 3^* | v(1) | 4^* \rangle = -\frac{\beta}{2}, \\ \langle {}^*1 | \mathcal{H} | \Psi_1^{3*} \rangle &= \left(\frac{1}{\sqrt{2}} \langle \Psi_1^{2*} | - \frac{1}{\sqrt{2}} \langle \Psi_1^{5*} | \right) \mathcal{H} | \Psi_1^{3*} \rangle = \frac{1}{\sqrt{2}} \langle \Psi_1^{2*} | \mathcal{H} | \Psi_1^{3*} \rangle - \frac{1}{\sqrt{2}} \langle \Psi_1^{5*} | \mathcal{H} | \Psi_1^{3*} \rangle \\ &= \frac{1}{\sqrt{2}} \times \left(-\frac{\beta}{2}\right) - \frac{1}{\sqrt{2}} \times 0 = -\frac{\beta}{2\sqrt{2}}, \\ \langle {}^*1 | \mathcal{H} | \Psi_1^{4*} \rangle &= \left(\frac{1}{\sqrt{2}} \langle \Psi_1^{2*} | - \frac{1}{\sqrt{2}} \langle \Psi_1^{5*} | \right) \mathcal{H} | \Psi_1^{4*} \rangle = \frac{1}{\sqrt{2}} \langle \Psi_1^{2*} | \mathcal{H} | \Psi_1^{4*} \rangle - \frac{1}{\sqrt{2}} \langle \Psi_1^{5*} | \mathcal{H} | \Psi_1^{4*} \rangle \end{aligned}$$

$$= \frac{1}{\sqrt{2}} \times 0 - \frac{1}{\sqrt{2}} \times \left(-\frac{\beta}{2}\right) = \frac{\beta}{2\sqrt{2}}.$$

Thus, the corresponding matrix is

$$\begin{pmatrix} 0 & \frac{1}{\sqrt{2}}\beta & 0 & 0 \\ \frac{1}{\sqrt{2}}\beta & -2\beta & -\frac{1}{2\sqrt{2}}\beta & \frac{1}{2\sqrt{2}}\beta \\ 0 & -\frac{1}{2\sqrt{2}}\beta & -2\beta & -\frac{1}{2}\beta \\ 0 & \frac{1}{2\sqrt{2}}\beta & -\frac{1}{2}\beta & -2\beta \end{pmatrix} \begin{pmatrix} 1 \\ c_1 \\ c_2 \\ c_3 \end{pmatrix} = e_1 \begin{pmatrix} 1 \\ c_1 \\ c_2 \\ c_3 \end{pmatrix}.$$

The eigen equation is

$$\det(\mathbf{H} - e\mathbf{1}) = \frac{1}{8} (2e + 5\beta) (4e^3 + 14\beta e^2 + 9\beta^2 e - 3\beta^3) = 0,$$

whose four roots are

$$e_1 = -2.5\beta, \quad e_2 \approx -2.46271\beta, \quad e_3 \approx -1.27596\beta, \quad e_4 \approx 0.238676\beta.$$

Thus, $e = e_4 \approx 0.2387\beta$.

$$E'_R(\text{IEPA}) = 10e = 2.387\beta.$$

Compared with the approximate result of 2.247β obtained in part (a), so that there is a

$$\frac{2.387\beta - 2.247\beta}{2.247\beta} = 6.2\%$$

difference.

Exercise 5.20

Verify Eq.(5.152c), derive Eq.(5.153a,b), and solve them to obtain the result shown in Eq.(5.154).

Solution 5.20

The verification of (5.152c) is listed as following:

$$\begin{aligned} \langle {}^*_1 | \mathcal{H} | {}^*_2 \rangle &= \frac{1}{2} [\langle \Psi_1^{2*} | \mathcal{H} | \Psi_2^{3*} \rangle - \langle \Psi_1^{2*} | \mathcal{H} | \Psi_2^{1*} \rangle - \langle \Psi_1^{3*} | \mathcal{H} | \Psi_2^{3*} \rangle + \langle \Psi_1^{3*} | \mathcal{H} | \Psi_2^{1*} \rangle] \\ &= \frac{1}{2} [0 - 0 - (-v_{21}) + 0] = \frac{1}{2} v_{21} = \frac{1}{2} \frac{\beta}{2} = \frac{\beta}{4}, \\ \langle {}^*_1 | \mathcal{H} | {}^*_3 \rangle &= \frac{1}{2} [\langle \Psi_1^{2*} | \mathcal{H} | \Psi_3^{1*} \rangle - \langle \Psi_1^{2*} | \mathcal{H} | \Psi_3^{2*} \rangle - \langle \Psi_1^{3*} | \mathcal{H} | \Psi_3^{1*} \rangle + \langle \Psi_1^{3*} | \mathcal{H} | \Psi_3^{2*} \rangle] \\ &= \frac{1}{2} [0 - (-v_{31}) - 0 + 0] = \frac{1}{2} v_{31} = \frac{1}{2} \frac{\beta}{2} = \frac{\beta}{4}, \\ \langle {}^*_2 | \mathcal{H} | {}^*_3 \rangle &= \frac{1}{2} [\langle \Psi_2^{3*} | \mathcal{H} | \Psi_3^{1*} \rangle - \langle \Psi_2^{3*} | \mathcal{H} | \Psi_3^{2*} \rangle - \langle \Psi_2^{1*} | \mathcal{H} | \Psi_3^{1*} \rangle + \langle \Psi_2^{1*} | \mathcal{H} | \Psi_3^{2*} \rangle] \\ &= \frac{1}{2} [0 - 0 - (-v_{32}) + 0] = \frac{1}{2} v_{32} = \frac{1}{2} \frac{\beta}{2} = \frac{\beta}{4}. \end{aligned}$$

Thus the first two equations of the corresponding DCI eigenvalue problem are

$$\begin{aligned} \langle \Psi_0 | \mathcal{H} - E_0 | \Psi_0 \rangle + \sum_{i=1}^3 \langle \Psi_0 | \mathcal{H} | \Psi_i^* \rangle c_i + \sum_{i=1}^3 \langle \Psi_0 | \mathcal{H} | \Psi_i^{\bar{*}} \rangle \bar{c}_i &= E_R(\text{SCI}), \\ \langle \Psi_1^* | \mathcal{H} | \Psi_0 \rangle + \langle \Psi_1^* | \mathcal{H} - E_0 | \Psi_1^* \rangle c_1 + \sum_{i=2}^3 \langle \Psi_1^* | \mathcal{H} | \Psi_i^* \rangle c_i + \sum_{i=1}^3 \langle \Psi_1^* | \mathcal{H} | \Psi_i^{\bar{*}} \rangle \bar{c}_i &= E_R(\text{SCI})c_1. \end{aligned}$$

As $c_1 = c_2 = c_3 = \bar{c}_1 = \bar{c}_2 = \bar{c}_3 = c$, we get

$$\begin{aligned} 0 + \sqrt{2}\beta c + \sqrt{2}\beta c + \sqrt{2}\beta c + \sqrt{2}\beta c + \sqrt{2}\beta c + \sqrt{2}\beta c &= E_R(\text{SCI}), \\ \frac{1}{\sqrt{2}}\beta - \frac{3}{2}\beta c + \frac{\beta}{4}c + \frac{\beta}{4}c + 0 + 0 + 0 &= E_R(\text{SCI})c. \end{aligned}$$

These two equations are (5.153a) and (5.153b), which equals

$$\begin{pmatrix} 0 & 3\sqrt{2}\beta \\ \frac{1}{\sqrt{2}}\beta & -\beta \end{pmatrix} \begin{pmatrix} 1 \\ c \end{pmatrix} = E_R(\text{SCI}) \begin{pmatrix} 1 \\ c \end{pmatrix}.$$

The eigen equation is a quadratic equation of ε , viz.,

$$\det(\mathbf{H} - \varepsilon \mathbf{1}) = -\varepsilon(-\beta - \varepsilon) - 3\beta^2 = \varepsilon^2 + \beta\varepsilon - 3\beta^2 = 0.$$

The discriminant Δ_ε is

$$\beta^2 - 4 \times 1 \times (-3\beta^2) = 13\beta^2,$$

and the lowest root is

$$E_R(\text{SCI}) = \frac{1}{2} \left(-\beta + \sqrt{13}\beta \right) = \frac{\sqrt{13} - 1}{2} \beta \approx 1.3028\beta. \quad (5.20-1)$$

Exercise 5.21

Extend the above analysis to calculate the SCI resonance energy for a cyclic polyene with $N = 2n$ ($N > 6$) carbon atoms. If we restrict ourselves to only those configurations which interact with $|\Psi_0\rangle$, then the appropriate generalization of Eq.(5.151) is

$$|\Psi_{\text{SCI}}\rangle = |\Psi_0\rangle + \sum_{i=1}^n c_i |i^*\rangle + \sum_{i=1}^n \bar{c}_i |\bar{i}^*\rangle.$$

As discussed in Exercise 5.19b, this is not the complete SCI wave function because there exist additional singly excited configurations which, although they do not mix with $|\Psi_0\rangle$, they do mix with $|i^*\rangle$. The omission of these does not affect our qualitative conclusions. Show that the required matrix elements are

$$\begin{aligned} \langle \Psi_0 | \mathcal{H} | i^* \rangle &= \frac{1}{\sqrt{2}} \beta, \\ \langle i^* | \mathcal{H} - E_0 | j^* \rangle &= (-2\beta) \delta_{ij}. \end{aligned}$$

Why are Eqs.(5.152b) and Eqs.(5.152c) different? Using these matrix elements, show that the SCI equations are

$$\begin{aligned} E_R(\text{SCI}) &= \frac{1}{\sqrt{2}} N c \beta, \\ \frac{1}{\sqrt{2}} \beta - 2c\beta &= E_R(\text{SCI})c. \end{aligned}$$

Finally, solve them to obtain

$$E_R(\text{SCI}) = \left(\sqrt{1 + \frac{N}{2}} - 1 \right) \beta$$

which is proportional to $N^{1/2}$ as N becomes large.

Solution 5.21

Similar to Exercise 5.20, we find that

$$\langle \Psi_0 | \mathcal{H} | i^* \rangle = \frac{1}{\sqrt{2}} \langle \Psi_0 | \mathcal{H} | \Psi_i^{(i+1)*} \rangle - \frac{1}{\sqrt{2}} \langle \Psi_0 | \mathcal{H} | \Psi_i^{(i-1)*} \rangle = \frac{1}{\sqrt{2}} \left(\frac{\beta}{2} \right) - \frac{1}{\sqrt{2}} \left(-\frac{\beta}{2} \right) = \frac{\beta}{\sqrt{2}}.$$

However, note that though $i \neq j \Leftrightarrow i \pm 1 \neq j \neq 1$, $i \neq j$ cannot guarantee $i \pm 1 \neq j \mp 1$, and we find that

$$\begin{aligned} \langle \Psi_i^{(i+1)*} | \mathcal{H} | \Psi_j^{(j+1)*} \rangle &= \delta_{ij} (E_0 + \varepsilon_{(i+1)*} - \varepsilon_i) = \delta_{ij} [E_0 + (\alpha - \beta) - (\alpha + \beta)] = \delta_{ij} (E_0 - 2\beta), \\ \langle \Psi_i^{(i+1)*} | \mathcal{H} | \Psi_j^{(j-1)*} \rangle &= \delta_{i+2,j} \langle i | v(i) | (i+2) \rangle = 0, \\ \langle \Psi_i^{(i-1)*} | \mathcal{H} | \Psi_j^{(j+1)*} \rangle &= \delta_{i,j+2} \langle (j+2) | v(j) | j \rangle = 0, \end{aligned}$$

$$\langle \Psi_i^{(i-1)*} | \mathcal{H} | \Psi_j^{(j-1)*} \rangle = \delta_{ij} (E_0 + \varepsilon_{(i-1)*} - \varepsilon_i) = \delta_{ij} [E_0 + (\alpha - \beta) - (\alpha + \beta)] = \delta_{ij} (E_0 - 2\beta).$$

Thus,

$$\begin{aligned} \langle {}^*_i | \mathcal{H} - E_0 | {}^*_j \rangle &= \left(\frac{1}{\sqrt{2}} \langle \Psi_i^{(i+1)*} | - \frac{1}{\sqrt{2}} \langle \Psi_i^{(i-1)*} | \right) \mathcal{H} \left(\frac{1}{\sqrt{2}} | \Psi_j^{(j+1)*} \rangle - \frac{1}{\sqrt{2}} | \Psi_j^{(j-1)*} \rangle \right) - E_0 \\ &= \frac{1}{2} \left(\langle \Psi_i^{(i+1)*} | \mathcal{H} | \Psi_j^{(j+1)*} \rangle - \langle \Psi_i^{(i+1)*} | \mathcal{H} | \Psi_j^{(j-1)*} \rangle \right. \\ &\quad \left. - \langle \Psi_i^{(i-1)*} | \mathcal{H} | \Psi_j^{(j+1)*} \rangle + \langle \Psi_i^{(i-1)*} | \mathcal{H} | \Psi_j^{(j-1)*} \rangle \right) - E_0 \\ &= \frac{1}{2} (E_0 - 2\beta - 0 - 0 + E_0 - 2\beta) - E_0 = E_0 - 2\beta - E_0 = -2\beta, \end{aligned}$$

which is different from (5.152b) because when $N > 6$, there exist other occupied orbitals which are not located in nearest-neighbor carbon atoms. This is the same as my opinion in Exercise 5.19(a).

Similarly, due to the symmetry of benzene, it follows that $c_i = \bar{c}_i = c$. The first two equations of SCI equations are

$$\begin{aligned} 0 + \sum_{i=1}^n \frac{\beta}{\sqrt{2}} c + \sum_{i=1}^n \frac{\beta}{\sqrt{2}} c &= E_R(\text{SCI}), \\ \frac{\beta}{\sqrt{2}} + (-2\beta) + \sum_{j=2}^n (-2\beta) \delta_{1j} c + (-2\beta) + \sum_{j=1}^n 0 \times c &= E_R(\text{SCI})c, \end{aligned}$$

which can be simplified to

$$\begin{aligned} \frac{N\beta}{\sqrt{2}} c &= E_R(\text{SCI}), \\ \frac{\beta}{\sqrt{2}} - 2\beta c &= E_R(\text{SCI})c. \end{aligned}$$

In the same way, we obtain a quadratic equation of c ,

$$\frac{N\beta}{\sqrt{2}} c^2 + 2\beta c - \frac{\beta}{\sqrt{2}} = 0.$$

Its discriminant Δ_c is

$$(2\beta)^2 - 4 \times \left(\frac{N\beta}{\sqrt{2}} \right) \times \left(-\frac{\beta}{\sqrt{2}} \right) = (4 + 2N)\beta^2 = 4\left(1 + \frac{N}{2}\right)\beta^2,$$

and the lowest root is

$$c_{\min} = \frac{-2\beta + 2\sqrt{1 + \frac{N}{2}}\beta}{2 \times \frac{N\beta}{\sqrt{2}}} = \frac{1}{N} \left(\sqrt{2 + N} - \sqrt{2} \right).$$

Thus, the correlation energy of SCI is

$$E_R(\text{SCI}) = \frac{N\beta}{\sqrt{2}} \frac{1}{N} \left(\sqrt{2 + N} - \sqrt{2} \right) = \left(\sqrt{1 + \frac{N}{2}} - 1 \right) \beta,$$

which is proportional to $N^{1/2}$ as N becomes large.

CHAPTER 6

Many-Body Perturbation Theory

6.1 Rayleigh-Schrödinger (RS) Perturbation Theory

*6.2 Diagrammatic Representation of RS Perturbation Theory

6.2.1 Diagrammatic Perturbation Theory for Two States

Exercise 6.1

Write down and evaluate all fifth-order diagrams that have the property that an imaginary horizontal line crosses only one hole and one particle line. Show that the sum of such diagrams is

$$\frac{V_{12}V_{21}(V_{22} - V_{11})^3}{(E_1^{(0)} - E_2^{(0)})^4}$$

Hint: There are eight such diagrams, and they can be generated by adding three dots to the second-order diagram in all positive ways.

Solution 6.1

The final results are listed below firstly.

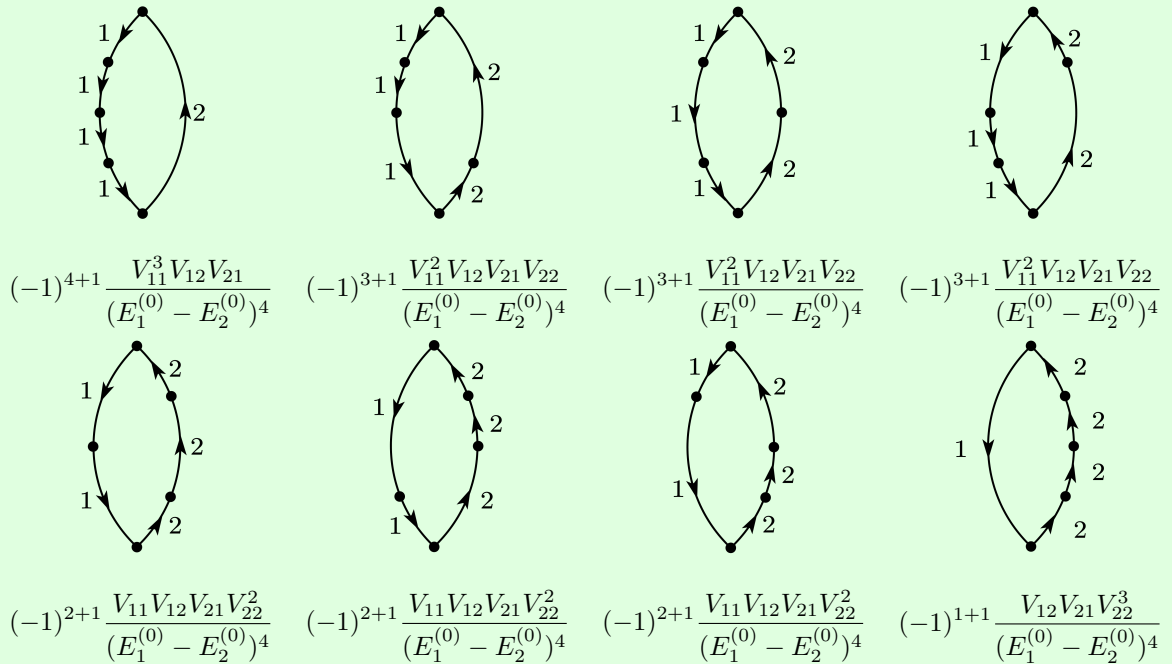


Figure 6.1: All fifth-order diagrams, which have the property that an imaginary horizontal line crosses only one hole and one particle line, and their mathematical expressions.

Note that the diagrams, which have the property that an imaginary horizontal line crosses only one hole

and one particle line, have no pair of hole/particle lines whose overlap is nonempty. For any pair of hole/particle lines which connect the dots m_1 and n_1 , m_2 and n_2 , where $n_1 > m_1$ and $n_2 > m_2$, their overlap will be

- $[\max\{m_1, m_2\}, \min\{n_1, n_2\}]$, if $\min\{n_1, n_2\} > \max\{m_1, m_2\}$;
- empty, otherwise.

For example, if there is a pair of hole lines, one connecting the dot 1 and 3, and the other connecting 2 and 5, their overlap will be $[2, 3]$. Take an another example, if there is a pair of particle lines, one connecting the dot 5 and 4, and the other connecting 2 and 1, their overlap will be empty.

Thus, there are four cases.

- Four hole lines and one particle line. There is only one method, hole lines are $(1, 2)$, $(2, 3)$, $(3, 4)$, $(4, 5)$ and the only particle line is $(5, 1)$, as the first subdiagram in Fig 6.1.
- Three hole lines and two particle lines. There are three methods as follows.
 - Hole lines are $(1, 2)$, $(2, 3)$, and $(3, 5)$ while particle lines are $(5, 4)$, $(4, 1)$.
 - Hole lines are $(1, 2)$, $(2, 4)$, and $(4, 5)$ while particle lines are $(5, 3)$, $(3, 1)$.
 - Hole lines are $(1, 3)$, $(3, 4)$, and $(4, 5)$ while particle lines are $(5, 2)$, $(2, 1)$.

They correspond to the second, third, fourth subdiagram in Fig 6.1.

- Two hole lines and three particle lines. There are three methods as follows.
 - Hole lines are $(1, 3)$, $(3, 5)$ while particle lines are $(5, 4)$, $(4, 2)$, and $(2, 1)$.
 - Hole lines are $(1, 4)$, $(4, 5)$ while particle lines are $(5, 3)$, $(3, 2)$, and $(2, 1)$.
 - Hole lines are $(1, 2)$, $(2, 5)$ while particle lines are $(5, 4)$, $(4, 3)$, and $(3, 1)$.

They correspond to the fifth, sixth, seventh subdiagram in Fig 6.1.

- One hole line and four particle lines. There is only one method, the only hole line is $(1, 5)$ and particle lines are $(5, 4)$, $(4, 3)$, $(3, 2)$, and $(2, 1)$, as the eighth subdiagram in Fig 6.1.

Thus, the sum of such diagrams is

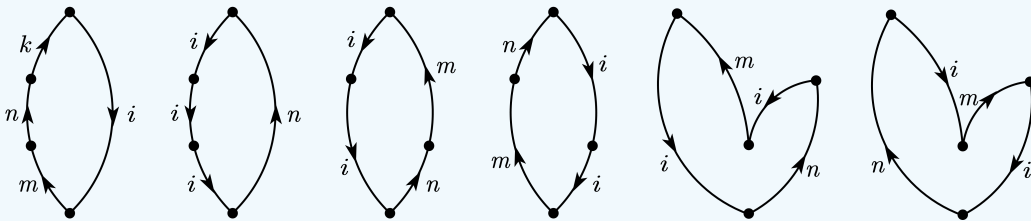
$$-\frac{V_{12}V_{21}(V_{11}^3 - 3V_{11}^2V_{22} + 3V_{11}V_{22}^2 - V_{22}^3)}{(E_1^{(0)} - E_2^{(0)})^4} = -\frac{V_{12}V_{21}(V_{11} - V_{22})^3}{(E_1^{(0)} - E_2^{(0)})^4} = \frac{V_{12}V_{21}(V_{22} - V_{11})^3}{(E_1^{(0)} - E_2^{(0)})^4}.$$

In fact, as the textbook says, these eight diagrams can be generated by adding three dots to the second-order diagram in all positive ways. In fact, any pair of hole/particle lines in them has also empty overlap. I think the calculation of the overlap is much direct than inspecting the property of lines.

6.2.2 Diagrammatic Perturbation Theory for N States

Exercise 6.2

Use diagrammatic techniques to obtain the fourth-order perturbation energy of a particular state (say, i) of an N -state system. That is, evaluate the diagrams



where the indices m, n, k, \dots exclude i . Using the approach of Section 6.1, obtain an algebraic expression for the fourth-order energy and compare it to the diagrammatic result.

Solution 6.2

Firstly, all fourth-order diagrams and their mathematical expressions are listed in Fig 6.2.

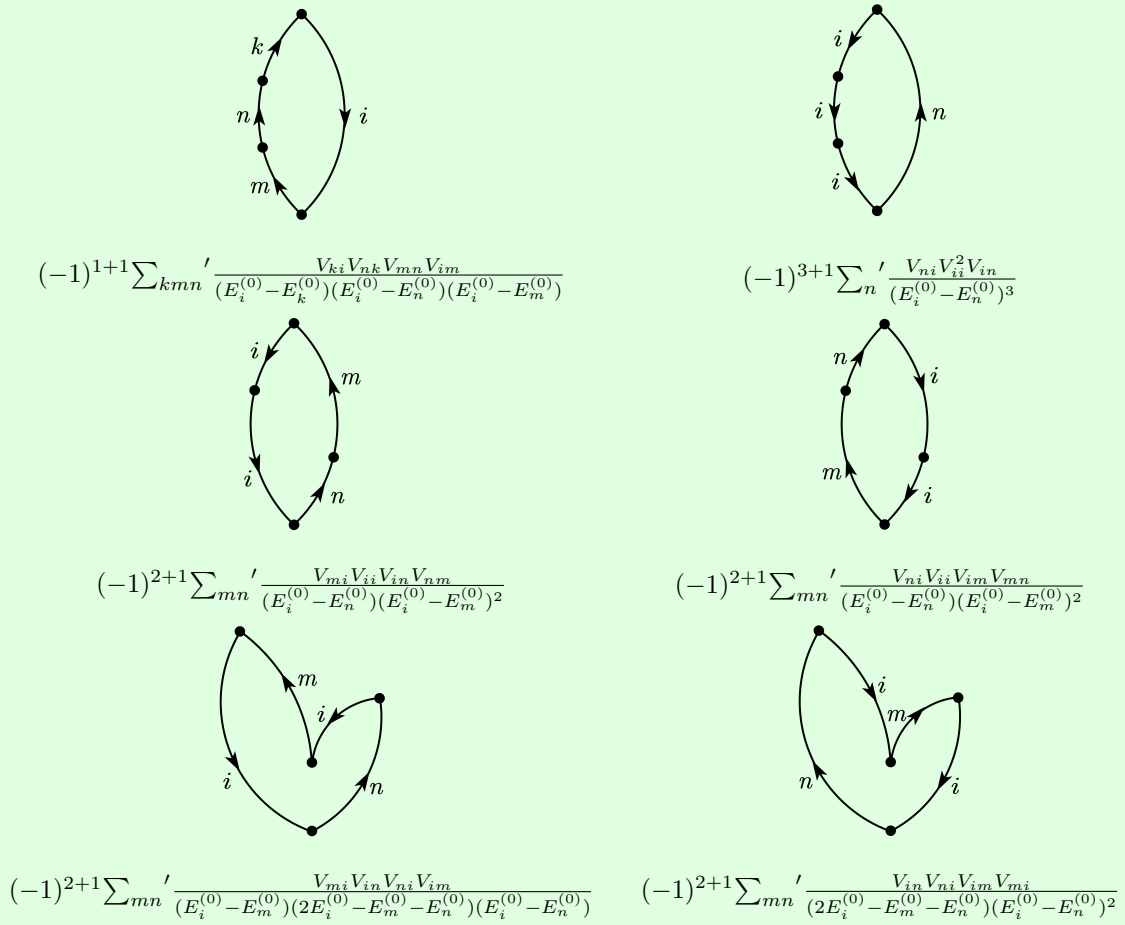


Figure 6.2: All fourth-order diagrams and their mathematical expressions.

Before the formal algebraic derivation, we should obtain some useful intermediate results. From (6.7d), multiplying by $\langle n|$, where $n \neq i$, we find that

$$E_n^{(0)} \langle n | \Psi_i^{(3)} \rangle + \langle n | \mathcal{V} | \Psi_i^{(2)} \rangle = E_i^{(0)} \langle n | \Psi_i^{(3)} \rangle + E_i^{(1)} \langle n | \Psi_i^{(2)} \rangle + E_i^{(1)} \langle n | \Psi_i^{(2)} \rangle,$$

and thus

$$\langle n | \Psi_i^{(3)} \rangle = \frac{1}{E_i^{(0)} - E_n^{(0)}} \left[\langle n | \mathcal{V} | \Psi_i^{(2)} \rangle - E_i^{(1)} \langle n | \Psi_i^{(2)} \rangle - E_i^{(1)} \langle n | \Psi_i^{(2)} \rangle \right].$$

Moreover, (6.8b), (6.10), (6.12), and (6.14) are used in the formal derivation. The fourth-order perturbation energy $E_i^{(4)}$ can be divided into 3 terms, viz.,

$$\begin{aligned} E_i^{(4)} &= \langle i | \mathcal{V} | \Psi_i^{(3)} \rangle = \sum_n ' \langle i | \mathcal{V} | n \rangle \langle n | \Psi_i^{(3)} \rangle = \sum_n ' V_{in} \frac{\langle n | \mathcal{V} | \Psi_i^{(2)} \rangle - E_i^{(1)} \langle n | \Psi_i^{(2)} \rangle - E_i^{(2)} \langle n | \Psi_i^{(1)} \rangle}{E_i^{(0)} - E_n^{(0)}} \\ &= \sum_n ' \frac{V_{in} \langle n | \mathcal{V} | \Psi_i^{(2)} \rangle}{E_i^{(0)} - E_n^{(0)}} - E_i^{(1)} \sum_n ' \frac{V_{in} \langle n | \Psi_i^{(2)} \rangle}{E_i^{(0)} - E_n^{(0)}} - E_i^{(2)} \sum_n ' \frac{V_{in} \langle n | \Psi_i^{(1)} \rangle}{E_i^{(0)} - E_n^{(0)}}. \end{aligned}$$

The first term is

$$\begin{aligned} \sum_n ' \frac{V_{in} \langle n | \mathcal{V} | \Psi_i^{(2)} \rangle}{E_i^{(0)} - E_n^{(0)}} &= \sum_{mn} ' \frac{V_{in} \langle n | \mathcal{V} | m \rangle \langle m | \Psi_i^{(2)} \rangle}{E_i^{(0)} - E_n^{(0)}} = \sum_{mn} ' \frac{V_{in} V_{nm}}{E_i^{(0)} - E_n^{(0)}} \langle m | \Psi_i^{(2)} \rangle \\ &= \sum_{mn} ' \frac{V_{in} V_{nm}}{E_i^{(0)} - E_n^{(0)}} \frac{\langle m | \mathcal{V} | \Psi_i^{(1)} \rangle - E_i^{(1)} \langle m | \Psi_i^{(1)} \rangle}{E_i^{(0)} - E_m^{(0)}} \\ &= \sum_{mn} ' \frac{V_{in} V_{nm}}{(E_i^{(0)} - E_n^{(0)})(E_i^{(0)} - E_m^{(0)})} \left[\langle m | \mathcal{V} | \Psi_i^{(1)} \rangle - E_i^{(1)} \langle m | \Psi_i^{(1)} \rangle \right] \end{aligned}$$

$$\begin{aligned}
&= \sum'_{mn} \frac{V_{in} V_{nm}}{(E_i^{(0)} - E_n^{(0)})(E_i^{(0)} - E_m^{(0)})} \left[\sum'_k \langle m | \mathcal{V} | k \rangle \langle k | \Psi_i^{(1)} \rangle - E_i^{(1)} \langle m | \Psi_i^{(1)} \rangle \right] \\
&= \sum'_{mnk} \frac{V_{in} V_{nm} V_{mk}}{(E_i^{(0)} - E_n^{(0)})(E_i^{(0)} - E_m^{(0)})} \langle k | \Psi_i^{(1)} \rangle - \sum'_{mn} \frac{V_{ii} V_{in} V_{nm}}{(E_i^{(0)} - E_n^{(0)})(E_i^{(0)} - E_m^{(0)})} \langle m | \Psi_i^{(1)} \rangle \\
&= \sum'_{mnk} \frac{V_{in} V_{nm} V_{mk}}{(E_i^{(0)} - E_n^{(0)})(E_i^{(0)} - E_m^{(0)})} \frac{V_{ki}}{E_i^{(0)} - E_k^{(0)}} - \sum'_{mn} \frac{V_{ii} V_{in} V_{nm}}{(E_i^{(0)} - E_n^{(0)})(E_i^{(0)} - E_m^{(0)})} \frac{V_{mi}}{E_i^{(0)} - E_m^{(0)}} \\
&= \sum'_{mnk} \frac{V_{in} V_{nm} V_{mk} V_{ki}}{(E_i^{(0)} - E_n^{(0)})(E_i^{(0)} - E_m^{(0)})(E_i^{(0)} - E_k^{(0)})} - \sum'_{mn} \frac{V_{ii} V_{in} V_{nm} V_{mi}}{(E_i^{(0)} - E_n^{(0)})(E_i^{(0)} - E_m^{(0)})^2} \\
&= \sum'_{mnk} \frac{V_{im} V_{mn} V_{nk} V_{ki}}{(E_i^{(0)} - E_n^{(0)})(E_i^{(0)} - E_m^{(0)})(E_i^{(0)} - E_k^{(0)})} - \sum'_{mn} \frac{V_{ii} V_{in} V_{nm} V_{mi}}{(E_i^{(0)} - E_n^{(0)})(E_i^{(0)} - E_m^{(0)})^2}.
\end{aligned}$$

It is evident that the first part of the first term correspond to the first subdiagram and the second part of the first term correspond to the third subdiagram.

The second term is

$$\begin{aligned}
&- E_i^{(1)} \sum'_n \frac{V_{in} \langle n | \Psi_i^{(2)} \rangle}{E_i^{(0)} - E_n^{(0)}} - \sum'_n \frac{V_{ii} V_{in}}{E_i^{(0)} - E_n^{(0)}} \langle n | \Psi_i^{(2)} \rangle \\
&= - \sum'_n \frac{V_{ii} V_{in}}{E_i^{(0)} - E_n^{(0)}} \frac{\langle n | \mathcal{V} | \Psi_i^{(1)} \rangle}{E_i^{(0)} - E_n^{(0)}} - E_i^{(1)} \sum'_n \frac{V_{ii} V_{in}}{(E_i^{(0)} - E_n^{(0)})^2} \langle n | \Psi_i^{(1)} \rangle \\
&= - \sum'_n \frac{V_{ii} V_{in}}{(E_i^{(0)} - E_n^{(0)})^2} \langle n | \mathcal{V} | \Psi_i^{(1)} \rangle + E_i^{(1)} \sum'_n \frac{V_{ii} V_{in}}{(E_i^{(0)} - E_n^{(0)})^2} \langle n | \Psi_i^{(1)} \rangle \\
&= - \sum'_n \frac{V_{ii} V_{in}}{(E_i^{(0)} - E_n^{(0)})^2} \sum'_m \langle n | \mathcal{V} | m \rangle \langle m | \Psi_i^{(1)} \rangle + \sum'_n \frac{V_{ii}^2 V_{in}}{(E_i^{(0)} - E_n^{(0)})^2} \langle n | \Psi_i^{(1)} \rangle \\
&= - \sum'_{mn} \frac{V_{ii} V_{in} V_{nm}}{(E_i^{(0)} - E_n^{(0)})^2} \langle m | \Psi_i^{(1)} \rangle + \sum'_n \frac{V_{ii}^2 V_{in}}{(E_i^{(0)} - E_n^{(0)})^2} \langle n | \Psi_i^{(1)} \rangle \\
&= - \sum'_{mn} \frac{V_{ii} V_{in} V_{nm}}{(E_i^{(0)} - E_n^{(0)})^2} \frac{\langle m | \mathcal{V} | i \rangle}{E_i^{(0)} - E_m^{(0)}} + \sum'_n \frac{V_{ii}^2 V_{in}}{(E_i^{(0)} - E_n^{(0)})^2} \frac{\langle n | \mathcal{V} | i \rangle}{E_i^{(0)} - E_n^{(0)}} \\
&= - \sum'_{mn} \frac{V_{ii} V_{in} V_{nm} V_{mi}}{(E_i^{(0)} - E_n^{(0)})^2 (E_i^{(0)} - E_m^{(0)})} + \sum'_n \frac{V_{ii}^2 V_{in} V_{ni}}{(E_i^{(0)} - E_n^{(0)})^3} \\
&= - \sum'_{mn} \frac{V_{ii} V_{im} V_{mn} V_{ni}}{(E_i^{(0)} - E_m^{(0)})^2 (E_i^{(0)} - E_n^{(0)})} + \sum'_n \frac{V_{ii}^2 V_{in} V_{ni}}{(E_i^{(0)} - E_n^{(0)})^3}.
\end{aligned}$$

It is evident that the first part of the second term correspond to the fourth subdiagram and the second part of the second term correspond to the second subdiagram.

The third term is

$$\begin{aligned}
&- E_i^{(2)} \sum'_n \frac{V_{in} \langle n | \Psi_i^{(1)} \rangle}{E_i^{(0)} - E_n^{(0)}} = - E_i^{(2)} \sum'_n \frac{V_{in}}{E_i^{(0)} - E_n^{(0)}} \frac{\langle n | \mathcal{V} | i \rangle}{E_i^{(0)} - E_n^{(0)}} = - E_i^{(2)} \sum'_n \frac{V_{in} V_{ni}}{(E_i^{(0)} - E_n^{(0)})^2} \\
&= - \left(\sum'_m \frac{V_{im} V_{mi}}{(E_i^{(0)} - E_m^{(0)})} \right) \sum'_n \frac{V_{in} V_{ni}}{(E_i^{(0)} - E_n^{(0)})^2} = - \sum'_{mn} \frac{V_{im} V_{mi} V_{in} V_{ni}}{(E_i^{(0)} - E_m^{(0)})(E_i^{(0)} - E_n^{(0)})^2}.
\end{aligned}$$

It seems that it does not directly correspond to any subdiagram in Fig 6.2. However, we can find that the sum of the mathematical expressions of the fifth and sixth subdiagram is

$$\begin{aligned}
&(-1)^{2+1} \sum'_{mn} \frac{V_{mi} V_{in} V_{ni} V_{im}}{(E_i^{(0)} - E_m^{(0)})(2E_i^{(0)} - E_m^{(0)} - E_n^{(0)})(E_i^{(0)} - E_n^{(0)})} \\
&\quad + (-1)^{2+1} \sum'_{mn} \frac{V_{in} V_{ni} V_{im} V_{mi}}{(2E_i^{(0)} - E_m^{(0)} - E_n^{(0)})(E_i^{(0)} - E_n^{(0)})^2} \\
&= - \sum'_{mn} \frac{V_{in} V_{ni} V_{im} V_{mi}}{(2E_i^{(0)} - E_m^{(0)} - E_n^{(0)})(E_i^{(0)} - E_n^{(0)})} \left[\frac{1}{E_i^{(0)} - E_m^{(0)}} + \frac{1}{E_i^{(0)} - E_n^{(0)}} \right]
\end{aligned}$$

$$\begin{aligned}
&= -\sum'_{mn} \frac{V_{in} V_{ni} V_{im} V_{mi}}{(2E_i^{(0)} - E_m^{(0)} - E_n^{(0)})(E_i^{(0)} - E_n^{(0)})} \frac{2E_i^{(0)} - E_m^{(0)} - E_n^{(0)}}{(E_i^{(0)} - E_m^{(0)})(E_i^{(0)} - E_n^{(0)})} \\
&= -\sum'_{mn} \frac{V_{im} V_{mi} V_{in} V_{ni}}{(E_i^{(0)} - E_m^{(0)})(E_i^{(0)} - E_n^{(0)})^2},
\end{aligned}$$

which is the third term exactly. Thus we can conclude that the results obtained by algebraic methods is the same as that by diagrammatic techniques. The mathematical expression of the fourth-order perturbation energy $E_i^{(4)}$ is

$$\begin{aligned}
E_i^{(4)} &= \sum'_{mnk} \frac{V_{im} V_{mn} V_{nk} V_{ki}}{(E_i^{(0)} - E_n^{(0)})(E_i^{(0)} - E_m^{(0)})(E_i^{(0)} - E_k^{(0)})} - \sum'_{mn} \frac{V_{ii} V_{in} V_{nm} V_{mi}}{(E_i^{(0)} - E_n^{(0)})(E_i^{(0)} - E_m^{(0)})^2} \\
&\quad - \sum'_{mn} \frac{V_{ii} V_{im} V_{mn} V_{ni}}{(E_i^{(0)} - E_m^{(0)})^2 (E_i^{(0)} - E_n^{(0)})} + \sum'_n \frac{V_{ii}^2 V_{in} V_{ni}}{(E_i^{(0)} - E_n^{(0)})^3} - \sum'_{mn} \frac{V_{im} V_{mi} V_{in} V_{ni}}{(E_i^{(0)} - E_m^{(0)})(E_i^{(0)} - E_n^{(0)})^2}.
\end{aligned}$$

6.2.3 Summation of Diagrams

6.3 Orbital Perturbation Theory: One-Particle Perturbations

Exercise 6.3

Derive

$$E_0^{(2)} = \sum_{ar} \frac{v_{ar} v_{ra}}{\varepsilon_a^{(0)} - \varepsilon_r^{(0)}}$$

starting with the general expression for the second-order energy (Eq.(6.12)) applied to an N -electron system,

$$E_0^{(2)} = \sum'_n \frac{\left| \langle \Psi_0 | \sum_i v(i) | n \rangle \right|^2}{E_0^{(0)} - E_n^{(0)}}$$

where the sum runs over all states of the system except the ground state.

Hint: The states $|n\rangle$ must be single excitations of the type

$$|\Psi_a^r\rangle = |\chi_1^{(0)} \cdots \chi_{a-1}^{(0)} \chi_r^{(0)} \chi_{a+1}^{(0)} \cdots \chi_N^{(0)}\rangle.$$

Solution 6.3

Note that (6.31) states that $\mathcal{V} = \sum_i v(i)$ only connects two Slater determinants whose different occupied orbitals should be no more than one, thus we obtain that

$$E_0^{(2)} = \sum'_n \frac{|\langle \Psi_0 | \sum_i v(i) | n \rangle|^2}{E_0^{(0)} - E_n^{(0)}} = \sum_{ar} \frac{|\langle \Psi_0 | \mathcal{V} | \Psi_a^r \rangle|^2}{E_0^{(0)} - (E_0^{(0)} + \varepsilon_r^{(0)} - \varepsilon_a^{(0)})} = \sum_{ar} \frac{v_{ar} v_{ra}}{\varepsilon_a^{(0)} - \varepsilon_r^{(0)}}. \quad (6.3-1)$$

Exercise 6.4

Calculate the third-order energy $E_0^{(3)}$ using the general expression given in Eq.(6.15).

a. Show that

$$B_0^{(3)} = -E_0^{(1)} \sum'_n \frac{|\langle \Psi_0 | \mathcal{V} | n \rangle|^2}{(E_0^{(0)} - E_n^{(0)})^2} = -\sum_{abr} \frac{v_{aa} v_{rb} v_{br}}{(\varepsilon_b^{(0)} - \varepsilon_r^{(0)})^2}.$$

b. Show that

$$A_0^{(3)} = \sum'_{nm} \frac{\langle \Psi_0 | \mathcal{V} | n \rangle \langle n | \mathcal{V} | m \rangle \langle m | \mathcal{V} | \Psi_0 \rangle}{(E_0^{(0)} - E_n^{(0)})(E_0^{(0)} - E_m^{(0)})} = \sum_{abrs} \frac{v_{ar} v_{sb} \langle \Psi_a^r | \mathcal{V} | \Psi_b^s \rangle}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_b^{(0)} - \varepsilon_s^{(0)})}.$$

c. Show that

$$\begin{aligned}\langle \Psi_a^r | \mathcal{V} | \Psi_b^s \rangle &= v_{rs} && \text{if } a = b \quad r \neq s, \\ &= -v_{ba} && \text{if } a \neq b \quad r = s, \\ &= \sum_c v_{cc} - v_{aa} + v_{rr} && \text{if } a = b \quad r = s,\end{aligned}$$

and zero otherwise.

d. Finally, combine the two terms to obtain

$$E_0^{(3)} = A_0^{(3)} + B_0^{(3)} = \sum_{ars} \frac{v_{ar} v_{rs} v_{sa}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_a^{(0)} - \varepsilon_s^{(0)})} - \sum_{abr} \frac{v_{ra} v_{ab} v_{br}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_b^{(0)} - \varepsilon_r^{(0)})}.$$

e. Show that for a closed-shell system

$$E_0^{(3)} = 2 \sum_{ars}^{N/2} \frac{v_{ar} v_{rs} v_{sa}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_a^{(0)} - \varepsilon_s^{(0)})} - 2 \sum_{abr}^{N/2} \frac{v_{ra} v_{ab} v_{br}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_b^{(0)} - \varepsilon_r^{(0)})}.$$

Solution 6.4

a. Similar to Exercise 6.3, it is evident that

$$\begin{aligned}B_0^{(3)} &= -E_0^{(1)} \sum_n' \frac{|\langle \Psi_0 | \mathcal{V} | n \rangle|^2}{(E_0^{(0)} - E_n^{(0)})^2} = - \left(\sum_a v_{aa} \right) \sum_{br} \frac{v_{rb} v_{br}}{[E_0^{(0)} - (E_n^{(0)} + \varepsilon_r^{(0)} - \varepsilon_b^{(0)})]^2} \\ &= - \sum_{abr} \frac{v_{aa} v_{rb} v_{br}}{(\varepsilon_b^{(0)} - \varepsilon_r^{(0)})^2}.\end{aligned}$$

b. In the same way, we obtain that

$$\begin{aligned}A_0^{(3)} &= \sum_{nm}' \frac{\langle \Psi_0 | \mathcal{V} | n \rangle \langle n | \mathcal{V} | m \rangle \langle m | \mathcal{V} | \Psi_0 \rangle}{(E_0^{(0)} - E_n^{(0)})(E_0^{(0)} - E_m^{(0)})} \\ &= \sum_{ar} \sum_{bs} \frac{\langle \Psi_0 | \mathcal{V} | \Psi_a^r \rangle \langle \Psi_a^r | \mathcal{V} | \Psi_b^s \rangle \langle \Psi_b^s | \mathcal{V} | \Psi_0 \rangle}{[E_0^{(0)} - (E_0^{(0)} + \varepsilon_r^{(0)} - \varepsilon_a^{(0)})][E_0^{(0)} - (E_0^{(0)} + \varepsilon_s^{(0)} - \varepsilon_b^{(0)})]} \\ &= \sum_{abrs} \frac{v_{ar} v_{sb} \langle \Psi_a^r | \mathcal{V} | \Psi_b^s \rangle}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_b^{(0)} - \varepsilon_s^{(0)})}.\end{aligned}$$

c. Using the conclusion of Exercise 2.13, at once we get

$$\begin{aligned}\langle \Psi_a^r | \mathcal{V} | \Psi_b^s \rangle &= v_{rs} && \text{if } a = b \quad r \neq s, \\ &= -v_{ba} && \text{if } a \neq b \quad r = s, \\ &= \sum_c v_{cc} - v_{aa} + v_{rr} && \text{if } a = b \quad r = s,\end{aligned}$$

and zero otherwise.

d. Thus, we simplify $A_0^{(3)}$ as follows.

$$\begin{aligned}A_0^{(3)} &= \sum_{abrs} \frac{v_{ar} v_{sb} \langle \Psi_a^r | \mathcal{V} | \Psi_b^s \rangle}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_b^{(0)} - \varepsilon_s^{(0)})} \\ &= \sum_{ar} \frac{v_{ar} v_{ra}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})^2} \left(\sum_c v_{cc} - v_{aa} + v_{rr} \right) + \sum_{\substack{ar \\ s \neq r}} \frac{v_{ar} v_{sa} v_{rs}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_a^{(0)} - \varepsilon_s^{(0)})} \\ &\quad + \sum_{\substack{ar \\ b \neq a}} \frac{v_{ar} v_{rb} (-v_{ba})}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_b^{(0)} - \varepsilon_r^{(0)})}\end{aligned}$$

$$\begin{aligned}
&= \sum_{ar} \frac{v_{ar}v_{ra}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})^2} \left(\sum_b v_{bb} \right) - v_{aa} \sum_{ar} \frac{v_{ar}v_{ra}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})^2} + v_{rr} \sum_{ar} \frac{v_{ar}v_{ra}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})^2} \\
&\quad + \sum_{\substack{ar \\ s \neq r}} \frac{v_{ar}v_{sa}v_{rs}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_a^{(0)} - \varepsilon_s^{(0)})} - \sum_{\substack{ar \\ b \neq a}} \frac{v_{ar}v_{rb}v_{ba}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_b^{(0)} - \varepsilon_r^{(0)})} \\
&= \left(\sum_a v_{aa} \right) \sum_{br} \frac{v_{br}v_{rb}}{(\varepsilon_b^{(0)} - \varepsilon_r^{(0)})^2} + \sum_{ars} \frac{v_{ar}v_{sa}v_{rs}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_a^{(0)} - \varepsilon_s^{(0)})} - \sum_{abr} \frac{v_{ar}v_{rb}v_{ba}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_b^{(0)} - \varepsilon_r^{(0)})} \\
&= \sum_{abr} \frac{v_{aa}v_{br}v_{rb}}{(\varepsilon_b^{(0)} - \varepsilon_r^{(0)})^2} + \sum_{ars} \frac{v_{ar}v_{sa}v_{rs}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_a^{(0)} - \varepsilon_s^{(0)})} - \sum_{abr} \frac{v_{ar}v_{rb}v_{ba}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_b^{(0)} - \varepsilon_r^{(0)})}.
\end{aligned}$$

Finally,

$$\begin{aligned}
E_0^{(3)} &= A_0^{(3)} + B_0^{(3)} \\
&= \sum_{abr} \frac{v_{aa}v_{br}v_{rb}}{(\varepsilon_b^{(0)} - \varepsilon_r^{(0)})^2} + \sum_{ars} \frac{v_{ar}v_{sa}v_{rs}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_a^{(0)} - \varepsilon_s^{(0)})} \\
&\quad - \sum_{abr} \frac{v_{ar}v_{rb}v_{ba}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_b^{(0)} - \varepsilon_r^{(0)})} - \sum_{abr} \frac{v_{aa}v_{rb}v_{br}}{(\varepsilon_b^{(0)} - \varepsilon_r^{(0)})^2} \\
&= \sum_{ars} \frac{v_{ar}v_{sa}v_{rs}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_a^{(0)} - \varepsilon_s^{(0)})} - \sum_{abr} \frac{v_{br}v_{ra}v_{ab}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_b^{(0)} - \varepsilon_r^{(0)})}.
\end{aligned}$$

- e. Since a matrix element $v_{ij} = \langle i|v|j \rangle$ is nonzero only if both spin orbitals i and j have the same spin, thus only $v_{ar}v_{sa}v_{rs}$ and $v_{\bar{a}\bar{r}}v_{\bar{s}\bar{a}}v_{\bar{r}\bar{s}}$ contribute, and so $v_{br}v_{ra}v_{ab}$ and $v_{\bar{b}\bar{r}}v_{\bar{r}\bar{a}}v_{\bar{a}\bar{b}}$ do $v_{br}v_{ra}v_{ab}$. Besides, the denominators is invariant regardless of spin up or down. In this way, we obtain that

$$\begin{aligned}
E_0^{(3)} &= \sum_{ars}^{N/2} \frac{v_{ar}v_{sa}v_{rs}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_a^{(0)} - \varepsilon_s^{(0)})} + \sum_{\bar{a}\bar{r}\bar{s}}^{N/2} \frac{v_{\bar{a}\bar{r}}v_{\bar{s}\bar{a}}v_{\bar{r}\bar{s}}}{(\varepsilon_{\bar{a}}^{(0)} - \varepsilon_{\bar{r}}^{(0)})(\varepsilon_{\bar{a}}^{(0)} - \varepsilon_{\bar{s}}^{(0)})} \\
&\quad - \sum_{abr}^{N/2} \frac{v_{br}v_{ra}v_{ab}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_b^{(0)} - \varepsilon_r^{(0)})} - \sum_{\bar{a}\bar{b}\bar{r}}^{N/2} \frac{v_{\bar{b}\bar{r}}v_{\bar{r}\bar{a}}v_{\bar{a}\bar{b}}}{(\varepsilon_{\bar{a}}^{(0)} - \varepsilon_{\bar{r}}^{(0)})(\varepsilon_{\bar{b}}^{(0)} - \varepsilon_{\bar{r}}^{(0)})} \\
&= 2 \sum_{ars}^{N/2} \frac{v_{ar}v_{sa}v_{rs}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_a^{(0)} - \varepsilon_s^{(0)})} - 2 \sum_{abr}^{N/2} \frac{v_{br}v_{ra}v_{ab}}{(\varepsilon_a^{(0)} - \varepsilon_r^{(0)})(\varepsilon_b^{(0)} - \varepsilon_r^{(0)})}.
\end{aligned}$$

Exercise 6.5

Show that the second term in Eq.(6.52) is equal to $\frac{3}{8}\beta$ for benzene.

Solution 6.5

The pictorial representation of the second term in $E_0^{(3)}$ can be seen in Fig 6.3.

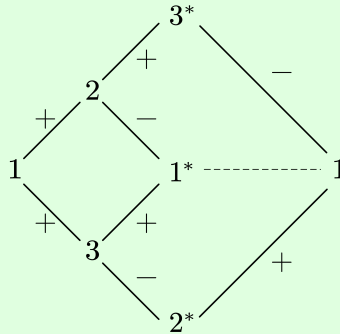


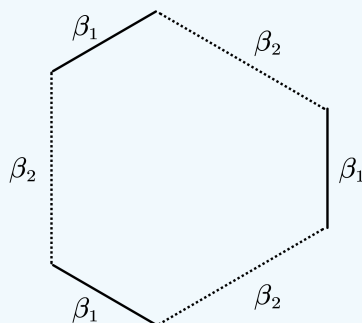
Figure 6.3: Pictorial representation of the second term in $E_0^{(3)}$, where the order is v_{ij} , v_{jk^*} and v_{k^*i} from the left to the right. The plus and minus signs indicate whether the matrix elements between the two orbitals is $\pm \frac{\beta}{2}$.

With the high symmetry, it is

$$-2 \times \frac{1}{(2\beta)^2} \sum_{i=1}^3 [v_{12}v_{23}^*v_{31} + v_{13}v_{32}^*v_{21}] = -\frac{3}{2\beta^2} \left[\frac{\beta}{2} \times \frac{\beta}{2} \times \left(-\frac{\beta}{2}\right) + \frac{\beta}{2} \times \left(-\frac{\beta}{2}\right) \times \frac{\beta}{2} \right] = \frac{3}{8\beta}. \quad (6.5-1)$$

Exercise 6.6

Consider a cyclic polyene with $N = 4\nu + 2$, $\nu = 1, 2, \dots$ carbons. Instead of assuming that all the bonds are identical, suppose they alternate in length. In the context of Hückel theory this means that the resonance integrals between adjacent carbons are not all equal to β but alternate between β_1 and β_2 . For example, for benzene we have



Now it can be shown that the exact energy for a cyclic polyene of this type is

$$\mathcal{E}_0 = N\alpha - 2 \sum_{j=-\nu}^{\nu} \left(\beta_1^2 + \beta_2^2 + 2\beta_1\beta_2 \cos \frac{2j\pi}{2\nu+1} \right)^{1/2}$$

(see for example, L. Salem, *Molecular Orbital Theory of Conjugated Systems*, Benjamin, New York, 1966, pp.498-500). Note that when $\beta_1 = \beta_2 = \beta$, since $2\cos^2\theta = (1 + \cos 2\theta)$ and β is negative, we recover

$$\mathcal{E}_0 = N\alpha + 4\beta \sum_{j=-\nu}^{\nu} \cos \frac{j\pi}{2\nu+1}$$

which is the result quoted in Subsection 5.3.2. Also note that when $\beta_1 = \beta$ but $\beta_2 = 0$, we have

$$\mathcal{E}_0 = N\alpha + N\beta$$

which is just the total energy of the polyene using the localized ethylenic description. The purpose of this exercise is to obtain the perturbation expansion for the resonance energy by expanding the exact result in powers of β_2/β_1 .

- Show that for benzene ($\nu = 1$) the exact ground state energy in the alternating short and long bond model is

$$\mathcal{E}_0 = 6\alpha + 2(\beta_1 + \beta_2) - 4(\beta_1^2 + \beta_2^2 - \beta_1\beta_2)^{1/2}$$

Do this first by using the general expression and then by setting up the Hückel matrix, diagonalizing it and adding up the occupied orbital energies. Note that when $\beta_1 = \beta_2 = \beta$ we recover our old result, $6\alpha + 8\beta$.

- Setting $\beta_1 = \beta$ and $\beta_2/\beta_1 = x$ show that the resonance energy of benzene can be written as

$$E_R = 4\beta \left(\frac{1}{2}x - 1 + (1 - x + x^2)^{1/2} \right)$$

Note that when $x = 0$, $E_R = 0$ and when $x = 1$, $E_R = 2\beta$ which is exact.

- Using the relation

$$(1+y)^{1/2} = 1 + \frac{1}{2}y - \frac{1}{8}y^2 + \frac{1}{16}y^3 - \frac{5}{128}y^4 + \dots, \quad |y| < 1$$

expand E_R to fourth order in x and thus show that

$$E_R = \beta \left(\frac{3}{2}x^2 + \frac{3}{4}x^3 + \frac{3}{32}x^4 + \cdots \right)$$

Identifying the coefficient of x^n with the n th-order perturbation result (i.e., $E_0^{(n)}$), we have

$$E_0^{(2)} = \frac{3}{2}\beta,$$

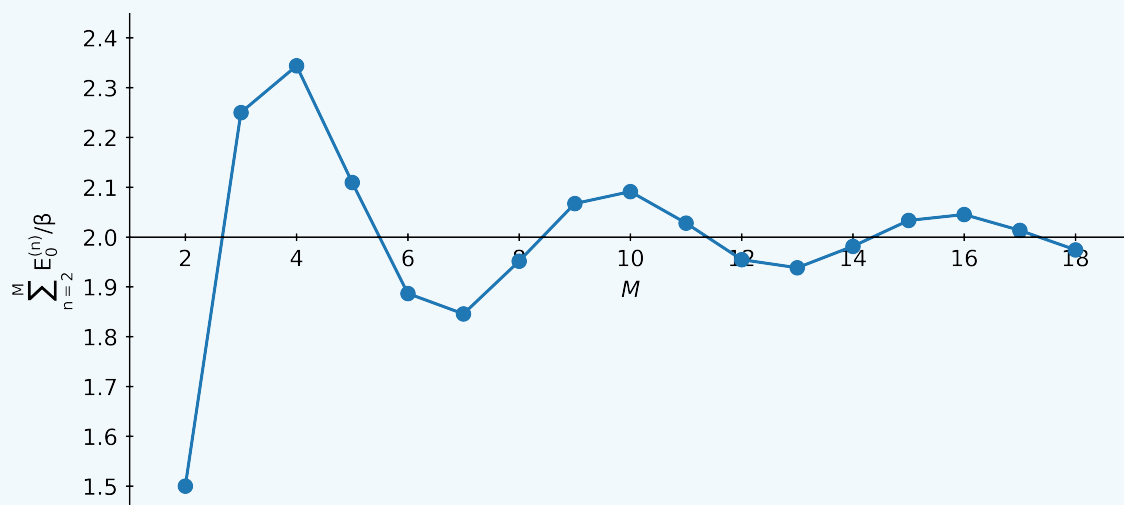
$$E_0^{(3)} = \frac{3}{4}\beta,$$

$$E_0^{(4)} = \frac{3}{32}\beta.$$

Note that $E_0^{(2)}$ and $E_0^{(3)}$ agree with our previously calculated values. This derivation provides some insight into the poor convergence of the perturbation expansion of the resonance energy of benzene. Basically, the perturbation expansion converges rapidly when x is small. However, for our problem x is equal to unity.

The resonance energy calculated up to M th-order as a function of M is shown below. Note the oscillatory convergence towards the exact value of 2β . The method used above to obtain $E_0^{(n)}$ for $n = 2, 3, 4$ becomes extremely laborious for larger n . The results below were calculated by first showing that $E_0^{(n)} = 4\beta C_n^{-1/2}(1/2)$, where $C_n^{-1/2}(x)$ is a Gegenbauer polynomial of degree n and order $-\frac{1}{2}$, and then using the recursive properties of these polynomials, to show that

$$(n+1)E_0^{(n+1)} = (n-1)E_0^{(n)} - (n-2)E_0^{(n-1)}.$$



Solution 6.6

a. Using the general expression and note that $\beta_1 \leq 0$ and $\beta_2 \leq 0$, we get that

$$\begin{aligned} \mathcal{E}_0(\text{benzene}) &= 6 \times \alpha - 2 \sum_{j=1}^1 \left(\beta_1^2 + \beta_2^2 + 2\beta_1\beta_2 \cos \frac{2j\pi}{2 \times 1 + 1} \right)^{1/2} \\ &= 6\alpha - 2 \left(\beta_1^2 + \beta_2^2 + 2\beta_1\beta_2 \cos \frac{-2\pi}{3} \right)^{1/2} \\ &\quad - 2 \left(\beta_1^2 + \beta_2^2 + 2\beta_1\beta_2 \cos \frac{0\pi}{3} \right)^{1/2} - 2 \left(\beta_1^2 + \beta_2^2 + 2\beta_1\beta_2 \cos \frac{2\pi}{3} \right)^{1/2} \\ &= 6\alpha - 2 (\beta_1^2 + \beta_2^2 - \beta_1\beta_2)^{1/2} + 2(\beta_1 + \beta_2) - 2 (\beta_1^2 + \beta_2^2 - \beta_1\beta_2)^{1/2}. \end{aligned}$$

In a nut shell,

$$\mathcal{E}_0(\text{benzene}) = 6\alpha + 2(\beta_1 + \beta_2) - 4(\beta_1^2 + \beta_2^2 - \beta_1\beta_2)^{1/2}. \quad (6.6-1)$$

At this time, we should remember that the Hückel matrix is

$$\mathbf{H} = \begin{pmatrix} \alpha & \beta_2 & 0 & 0 & 0 & \beta_1 \\ \beta_2 & \alpha & \beta_1 & 0 & 0 & 0 \\ 0 & \beta_1 & \alpha & \beta_2 & 0 & 0 \\ 0 & 0 & \beta_2 & \alpha & \beta_1 & 0 \\ 0 & 0 & 0 & \beta_1 & \alpha & \beta_2 \\ \beta_1 & 0 & 0 & 0 & \beta_2 & \alpha \end{pmatrix}.$$

Its eigen function is

$$\det(\mathbf{H} - \varepsilon \mathbf{1}) = (\beta_1^2 - \beta_1\beta_2 + \beta_2^2 - (\alpha - \varepsilon)^2)^2(\alpha - \beta_1 - \beta_2 - \varepsilon)(\alpha + \beta_1 + \beta_2 - \varepsilon) = 0,$$

and there are six roots:

$$\begin{aligned} \varepsilon_1 &= \alpha + \beta_1 + \beta_2, \\ \varepsilon_2 &= \varepsilon_3 = \alpha - \sqrt{\beta_1^2 - \beta_1\beta_2 + \beta_2^2}, \\ \varepsilon_4 &= \varepsilon_5 = \alpha + \sqrt{\beta_1^2 - \beta_1\beta_2 + \beta_2^2}, \\ \varepsilon_6 &= \alpha - \beta_1 - \beta_2. \end{aligned}$$

Thus,

$$\mathcal{E}'_0(\text{benzene}) = 2\varepsilon_1 + 2\varepsilon_2 + 2\varepsilon_3 = 6\alpha + 2(\beta_1 + \beta_2) - 4(\beta_1^2 + \beta_2^2 - \beta_1\beta_2)^{1/2}. \quad (6.6-2)$$

It equals the results obtained by using the general expression.

b. The verification is direct. With $\beta_1 = \beta$, $\beta_2 = x\beta_1 = x\beta$, we get that

$$\begin{aligned} E_R &= \mathcal{E}_0(\text{benzene}) - E_0 = 6\alpha + 2(\beta + x\beta) - 4(\beta^2 + (x\beta)^2 - x\beta^2)^{1/2} - (6\alpha + 6\beta) \\ &= -4\beta + 2x\beta + 4\beta(1 + x^2 - x)^{1/2} = 4\beta\left(\frac{x}{2} - 1 + (1 + x^2 - x)^{1/2}\right) \end{aligned}$$

c. When $0 \leq x \leq 1$, $-\frac{1}{4} \leq x^2 - x \leq 0$, thus we obtain that

$$\begin{aligned} E_R &= 4\beta\left(\frac{x}{2} - 1 + (1 + x^2 - x)^{1/2}\right) = 4\beta\left[\frac{x}{2} - 1 + (1 + (x^2 - x))^{1/2}\right] \\ &= 4\beta\left[\frac{x}{2} - 1 + 1 + \frac{1}{2}(x^2 - x) - \frac{1}{8}(x^2 - x)^2 + \frac{1}{16}(x^2 - x)^3 - \frac{5}{128}(x^2 - x)^4 + \dots\right] \\ &= 4\beta\left[\frac{x}{2} + \frac{1}{2}(-x + x^2) - \frac{1}{8}(x^2 - 2x^3 + x^4) \right. \\ &\quad \left. + \frac{1}{16}(-x^3 + 3x^4 - 3x^5 + x^6) - \frac{5}{128}(x^4 - 4x^5 + 6x^6 - 4x^7 + x^8) + \dots\right] \\ &= 4\beta\left[\left(\frac{1}{2} - \frac{1}{2}\right)x + \left(\frac{1}{2} - \frac{1}{8}\right)x^2 + \left(\frac{1}{4} - \frac{1}{16}\right)x^3 + \left(-\frac{1}{8} + \frac{3}{16} - \frac{5}{128}\right)x^4 + \dots\right] \\ &= \beta\left(\frac{3}{2}x^2 + \frac{3}{4}x^3 + \frac{3}{32}x^4 + \dots\right). \end{aligned}$$

In fact, the Gegenbauer polynomials $C_n^\lambda(x)$ can be generated by $(1 - 2xt + t^2)^{-\lambda}$, viz.,

$$(1 - 2xt + t^2)^{-\lambda} = \sum_{n=0}^{\infty} C_n^\lambda(x)t^n, \quad \forall t \in (-1, 1), x \in (-1, 1).$$

Let $\lambda = -\frac{1}{2}$ and $x = \frac{1}{2}$, we obtain that

$$(1 - t + t^2)^{\frac{1}{2}} = \sum_{n=0}^{\infty} C_n^{-\frac{1}{2}}\left(\frac{1}{2}\right)t^n.$$

Thus, with

$$C_0^\lambda(x) = 1, \quad C_1^\lambda(x) = 2\lambda x,$$

we find that

$$\sum_{n=0}^{\infty} C_n^{-\frac{1}{2}} \left(\frac{1}{2} \right) t^n = C_0^{-\frac{1}{2}} \left(\frac{1}{2} \right) + C_1^{-\frac{1}{2}} \left(\frac{1}{2} \right) t + \sum_{n=2}^{\infty} C_n^{-\frac{1}{2}} \left(\frac{1}{2} \right) t^n = 1 - \frac{t}{2} + \sum_{n=2}^{\infty} C_n^{-\frac{1}{2}} \left(\frac{1}{2} \right) t^n.$$

Hence, we get

$$E_R = 4\beta \left(\frac{x}{2} - 1 + (1 - x + x^2)^{1/2} \right) = 4\beta \sum_{n=2}^{\infty} C_n^{-\frac{1}{2}} \left(\frac{1}{2} \right) x^n. \quad (6.6-3)$$

It equals

$$E_0^{(n)} = 4\beta C_n^{-\frac{1}{2}} \left(\frac{1}{2} \right), \quad n = 2, 3, 4, \dots \quad (6.6-4)$$

Due to the recurrence relation of Gegenbauer polynomials,

$$(n+1)C_{n+1}^\lambda(x) = 2(n+\lambda)x C_n^\lambda(x) - (n+2\lambda-1)C_{n-1}^\lambda(x), \quad \forall n \geq 1,$$

we know that for $n = 2, 3, \dots$,

$$\begin{aligned} (n+1)E_0^{(n+1)} &= (n+1) \times 4\beta C_{n+1}^{-\frac{1}{2}} \left(\frac{1}{2} \right) = 4\beta(n+1)C_{n+1}^{-\frac{1}{2}} \left(\frac{1}{2} \right) \\ &= 4\beta \left[2 \left(n - \frac{1}{2} \right) \frac{1}{2} C_n^{-\frac{1}{2}} \left(\frac{1}{2} \right) - \left(n + 2 \times \left(-\frac{1}{2} \right) - 1 \right) C_{n-1}^{-\frac{1}{2}} \left(\frac{1}{2} \right) \right] \\ &= 4\beta \left[\left(n - \frac{1}{2} \right) C_n^{-\frac{1}{2}} \left(\frac{1}{2} \right) - (n-2)C_{n-1}^{-\frac{1}{2}} \left(\frac{1}{2} \right) \right] \\ &= \left(n - \frac{1}{2} \right) 4\beta C_n^{-\frac{1}{2}} \left(\frac{1}{2} \right) - (n-2)4\beta C_{n-1}^{-\frac{1}{2}} \left(\frac{1}{2} \right) \\ &= \left(n - \frac{1}{2} \right) E_0^{(n)} - (n-2)E_0^{(n-1)}. \end{aligned}$$

Remark: You can read some introduction about Gegenbauer polynomials from WIKIPEDIA, whose url is https://en.wikipedia.org/wiki/Gegenbauer_polynomials.

*6.4 Diagrammatic Representation of Orbital Perturbation Theory

Exercise 6.7

Find the fourth-order energy for a closed-shell cyclic polyene.

a. Show that

$$= \frac{N\beta}{64}$$

and

$$= -\frac{3N\beta}{128}$$

so that

$$E_0^{(4)} = \frac{N\beta}{64}$$

Thus the resonance energy calculated for a cyclic polyene with $N > 6$ up to fourth order is $(1/4 + 1/64)N\beta = 0.2656N\beta$, which compares very favorably with the asymptotically exact value of 0.2732β (i.e., 97%).

- b. For benzene, show that the diagrammatic result for the fourth-order energy agrees with the independently calculated result found in Exercise 6.6.

Solution 6.7

- a. Firstly, we list all diagrams of the fourth-order energy.

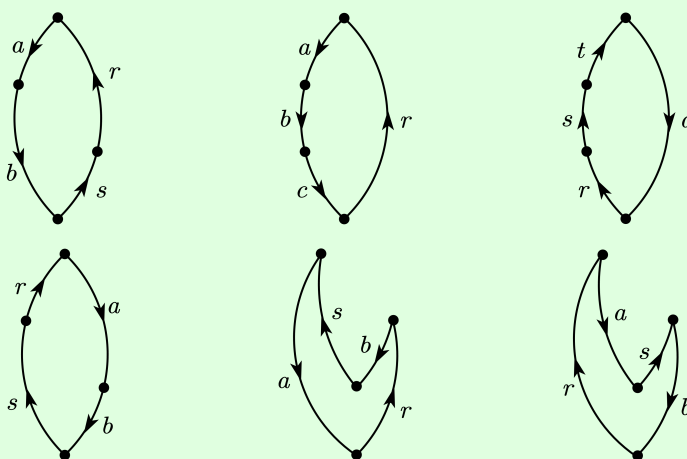


Figure 6.4: All fourth-order diagrams.

Note that when $N > 6$, $(i+3)^*$ cannot interact with i , and when $N = 6$, 1^* cannot interact with 1 , either. Thus, there is no essential difference in the shape and properties between the case of $N > 6$ and $N = 6$. It is enough to draw the pictorial representation of benzene ($N = 6$) in order to get the same result of $N > 6$. The pictorial representation of benzene can be seen in Fig 6.5.

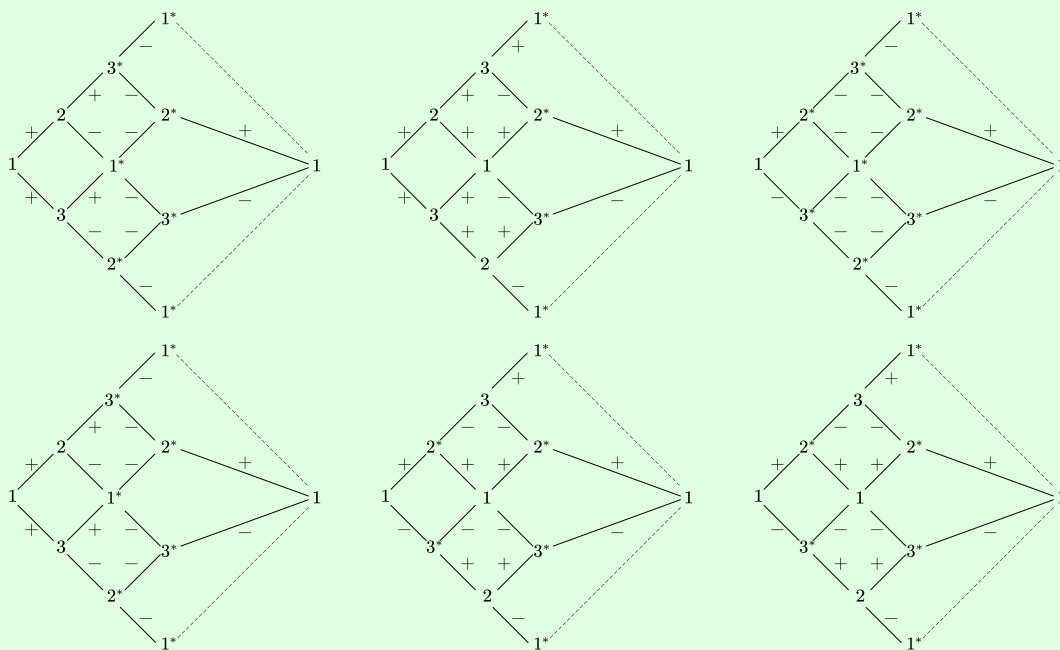


Figure 6.5: The pictorial representation of all fourth-order diagrams of the benzene.

Now we will calculate all terms according to their pictorial representation. Instead of lengthy calculation, we only care about the path from 1 to 1 in each diagram. For example, for the first subdiagram of Fig 6.5, we know that it has eight paths, six valid but two invalid. If a path has a segment with a plus/minus token, it will be marked as $+/-$, and zero otherwise. And thus, its eight paths are:

$$\begin{array}{cccc} (+, +, -, 0) & (+, +, -, +) & (+, -, -, +) & (+, -, -, -) \\ (+, +, -, +) & (+, +, -, -) & (+, -, -, -) & (+, -, -, 0) \end{array}$$

If a path has a 0, the corresponding term is zero. Otherwise, the corresponding term has a factor $(-1)^k$, where k is the number of its minus signs. Therefore, there are two positive and four negative terms, and the first term is

$$(-1)^{2+1} \frac{2 \times 3}{(2\beta)^3} \left[2 \times \left(\frac{\beta^4}{16} \right) + 4 \times \left(-\frac{\beta^4}{16} \right) \right] = \frac{6}{64} \beta = \frac{N}{64} \beta,$$

where the factor 2 occurs for the closed-shell structure, and the factor 3 is the number of occupied spin orbitals.

Similarly, we find that the second, third, ..., sixth term is $\frac{N}{64} \beta$, $\frac{N}{64} \beta$, $\frac{N}{64} \beta$, $-\frac{3N}{128} \beta$, $-\frac{3N}{128} \beta$. Thus, we obtain that

$$E_0^{(4)} = 4 \times \left(\frac{N}{64} \beta \right) + 2 \times \left(-\frac{3N}{128} \beta \right) = \frac{N}{64} \beta. \quad (6.7-1)$$

- b. As mentioned before, the general expression of the correlation energy with $N > 6$ is also suitable for benzene. Thus, the correlation energy of the benzene is

$$E_0^{(4)}(\text{benzene}) = \frac{6}{64} \beta = \frac{3}{32} \beta, \quad (6.7-2)$$

which agrees with the independently calculated result found in Exercise 6.6.

6.5 Perturbation Expansion of the Correlation Energy

Exercise 6.8

Derive Eqs.(6.73) and (6.74) starting with Eq.(6.72).

Solution 6.8

The detailed simplification can be seen in Exercise 2.18.

Exercise 6.9

Derive Eqs.(6.77) and (6.78) from Eq.(6.76).

Solution 6.9

We assume that $\Delta \approx (\varepsilon_2 - \varepsilon_1)$, thus

$$\Delta = (\varepsilon_2 - \varepsilon_1) + \frac{1}{2} J_{11} + \frac{1}{2} J_{22} - 2J_{12} + K_{12} = (\varepsilon_2 - \varepsilon_1) \left[1 + \frac{J_{11} + J_{22} - 4J_{12} + 2K_{12}}{2(\varepsilon_2 - \varepsilon_1)} \right].$$

Using binomial series and geometric series, we find that

$$\begin{aligned} E_{\text{corr}} &= \Delta - (\Delta^2 + K_{12}^2)^{\frac{1}{2}} = \Delta \left[1 - \left(1 + \frac{K_{12}^2}{\Delta^2} \right)^{\frac{1}{2}} \right] = \Delta \left[1 - \left(1 + \frac{K_{12}^2}{2\Delta^2} - \frac{K_{12}^4}{8\Delta^4} + \dots \right) \right] \\ &= -\frac{K_{12}^2}{2\Delta} + \frac{K_{12}^4}{8\Delta^3} + \dots \end{aligned}$$

$$\begin{aligned}
&= -\frac{K_{12}^2}{2(\varepsilon_2 - \varepsilon_1)} \left[1 + \frac{J_{11} + J_{22} - 4J_{12} + 2K_{12}}{2(\varepsilon_2 - \varepsilon_1)} \right] + \frac{K_{12}^4}{8(\varepsilon_2 - \varepsilon_1)^3} \left[1 + \frac{J_{11} + J_{22} - 4J_{12} + 2K_{12}}{2(\varepsilon_2 - \varepsilon_1)} \right]^3 + \dots \\
&= -\frac{K_{12}^2}{2(\varepsilon_2 - \varepsilon_1)} \left[1 - \frac{J_{11} + J_{22} - 4J_{12} + 2K_{12}}{2(\varepsilon_2 - \varepsilon_1)} + \left(\frac{J_{11} + J_{22} - 4J_{12} + 2K_{12}}{2(\varepsilon_2 - \varepsilon_1)} \right)^2 + \dots \right] + \dots \\
&= -\frac{K_{12}^2}{2(\varepsilon_2 - \varepsilon_1)} + \frac{K_{12}^2(J_{11} + J_{22} - 4J_{12} + 2K_{12})}{4(\varepsilon_2 - \varepsilon_1)^2} + o\left(\left(\frac{1}{\varepsilon_2 - \varepsilon_1}\right)^2\right).
\end{aligned}$$

Thus, we obtain (6.77) and (6.78) from (6.76).

6.6 The N -Dependence of the RS Perturbation Expansion

Exercise 6.10

Derive Eqs.(6.80b) and (6.90).

Solution 6.10

From (6.68), we find that with the truth that all two-electron integrals involving orbitals from different units are zero and $\langle mm||mm \rangle = 0$,

$$\begin{aligned}
E_0^{(1)} &= -\frac{1}{2} \sum_{i=1}^{2N} \sum_{j=1}^{2N} \langle 1_i 1_j || 1_i 1_j \rangle = -\frac{1}{2} \sum_{i=1}^{2N} \langle 1_i 1_i || 1_i 1_i \rangle \\
&= -\frac{1}{2} \sum_{i=1}^N \langle 1_i 1_i || 1_i 1_i \rangle + \langle 1_i \bar{1}_i || 1_i \bar{1}_i \rangle + \langle \bar{1}_i 1_i || \bar{1}_i 1_i \rangle + \langle \bar{1}_i \bar{1}_i || \bar{1}_i \bar{1}_i \rangle = -\frac{1}{2} \sum_{i=1}^N 2J_{11} = -\sum_{i=1}^N J_{11} = -NJ_{11}.
\end{aligned}$$

Besides, we obtain that

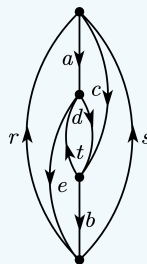
$$\begin{aligned}
\langle \Psi_{1_i \bar{1}_i}^{2_i \bar{2}_i} | \mathcal{V} | \Psi_{1_i \bar{1}_i}^{2_i \bar{2}_i} \rangle &= \langle \Psi_{1_i \bar{1}_i}^{2_i \bar{2}_i} | \mathcal{H} | \Psi_{1_i \bar{1}_i}^{2_i \bar{2}_i} \rangle - \langle \Psi_{1_i \bar{1}_i}^{2_i \bar{2}_i} | \mathcal{H}_0 | \Psi_{1_i \bar{1}_i}^{2_i \bar{2}_i} \rangle \\
&= \sum_{i=1}^{2N} h_{11} + 2h_{22} - 2h_{11} + \sum_{i=1}^N J_{11} + J_{22} - J_{11} - [(2N-2)\varepsilon_1 + 2\varepsilon_2] \\
&= (2N-2)h_{11} + 2h_{22} + (N-1)J_{11} + J_{22} - (2N-2)(h_{11} + J_{11}) - 2(h_{22} + 2J_{12} - K_{12}) \\
&= -(N-1)J_{11} + J_{22} - 4J_{12} + 2K_{12}.
\end{aligned}$$

*6.7 Diagrammatic Representation of the Perturbation Expansion of the Correlation Energy

6.7.1 Hugenholtz Diagrams

Exercise 6.11

Show that the fourth-order diagram



is equal to

$$-\frac{1}{2} \sum_{abcderst} \frac{\langle rs||ac\rangle\langle at||de\rangle\langle dc||tb\rangle\langle eb||rs\rangle}{(\varepsilon_a + \varepsilon_c - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d + \varepsilon_e - \varepsilon_r - \varepsilon_t - \varepsilon_s)(\varepsilon_b + \varepsilon_e - \varepsilon_r - \varepsilon_s)}.$$

Solution 6.11

Using the diagrammatic method, this term is made of four parts as follows.

1. There are 5 hole lines (a, b, c, d, e) and 3 particle lines (r, s, t). Thus we add a summation token with subscripts $abcderst$.
2. There are 2 loops, $r \rightarrow a \rightarrow d \rightarrow t \rightarrow e \rightarrow r$, $s \rightarrow c \rightarrow b \rightarrow s$, and 5 hole lines, thus we should add a $(-1)^{2+5} = (-1)^7$.
3. From the top to the bottom, 4 dots contribute $\langle rs||ac\rangle$, $\langle at||de\rangle$, $\langle dc||tb\rangle$, $\langle eb||rs\rangle$, respectively.
4. There are 3 imaginary horizontal lines.
 - The first one crosses a, c, r, s , contributing a factor $\varepsilon_a + \varepsilon_c - \varepsilon_r - \varepsilon_s$ to the denominator.
 - The second one crosses c, d, e, r, t, s contributing a factor $\varepsilon_c + \varepsilon_d + \varepsilon_e - \varepsilon_r - \varepsilon_t - \varepsilon_s$ to the denominator.
 - The third one crosses b, e, r, s , contributing a factor $\varepsilon_b + \varepsilon_e - \varepsilon_r - \varepsilon_s$ to the denominator.

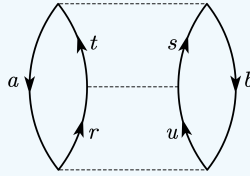
In conclusion, we know the mathematical expression of this diagram is

$$-\frac{1}{2} \sum_{abcderst} \frac{\langle rs||ac\rangle\langle at||de\rangle\langle dc||tb\rangle\langle eb||rs\rangle}{(\varepsilon_a + \varepsilon_c - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d + \varepsilon_e - \varepsilon_r - \varepsilon_t - \varepsilon_s)(\varepsilon_b + \varepsilon_e - \varepsilon_r - \varepsilon_s)}. \quad (6.11-1)$$

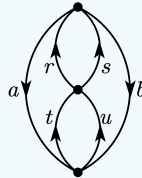
6.7.2 Goldstone Diagrams

Exercise 6.12

We stated that the Goldstone diagrams in Table 6.2 can be obtained by “pulling apart” the second- and third-order Hugenholtz diagrams. This is quite tricky to see but the converse is much easier. For example, if we push



together, it is clear that we obtain



Push all third-order diagrams in Table 6.2 together in a similar way and, thus, find which Goldstone diagram comes from which Hugenholtz diagram. For the above Hugenholtz diagram, verify that its mathematical value is indeed the sum of the values of the corresponding Goldstone diagrams.

Solution 6.12

As the textbook mentioned at the page 361, there are three third-order Hugenholtz diagrams, which are listed in Fig 6.6.

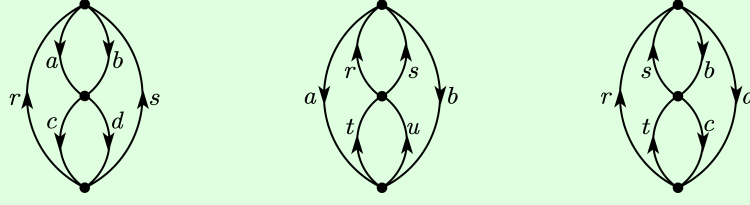


Figure 6.6: All third-order Hugenholtz diagrams.

In fact, the “pushing together” results of these third-order Goldstone diagrams in the table 6.2 can be summarized as follows.

- Pushing together the second or the seventh third-order Goldstone diagrams leads to the first Hugenholtz diagram.
- Pushing together the first and the eighth third-order Goldstone diagrams leads to the second Hugenholtz diagram.
- Pushing together the rest of third-order Goldstone diagrams leads to the third Hugenholtz diagram.

Firstly, we will verify that the sum of the second and the seventh third-order Goldstone diagrams is the first third-order Hugenholtz diagram. The sum of the second and the seventh third-order Goldstone diagrams is

$$\begin{aligned}
 & (-1)^{4+2} \left(\frac{1}{2} \right) \sum_{abcdrs} \frac{\langle ad|rs\rangle \langle cb|ad\rangle \langle rs|cb\rangle}{(\varepsilon_a + \varepsilon_d - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_b - \varepsilon_r - \varepsilon_s)} \\
 & + (-1)^{4+1} \left(\frac{1}{2} \right) \sum_{abcdrs} \frac{\langle ac|rs\rangle \langle db|ac\rangle \langle sr|db\rangle}{(\varepsilon_a + \varepsilon_c - \varepsilon_r - \varepsilon_s)(\varepsilon_d + \varepsilon_b - \varepsilon_r - \varepsilon_s)} \\
 & = \frac{1}{2} \sum_{abcdrs} \frac{\langle ad|rs\rangle \langle cb|ad\rangle \langle rs|cb\rangle}{(\varepsilon_a + \varepsilon_d - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_b - \varepsilon_r - \varepsilon_s)} - \frac{\langle ad|rs\rangle \langle cb|ad\rangle \langle sr|cb\rangle}{(\varepsilon_a + \varepsilon_d - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_b - \varepsilon_r - \varepsilon_s)} \\
 & = \frac{1}{2} \sum_{abcdrs} \frac{\langle ad|rs\rangle \langle cb|ad\rangle (\langle rs|cb\rangle - \langle sr|cb\rangle)}{(\varepsilon_a + \varepsilon_d - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_b - \varepsilon_r - \varepsilon_s)} = \frac{1}{2} \sum_{abcdrs} \frac{\langle ab|rs\rangle \langle cd|ab\rangle (\langle rs|cd\rangle - \langle sr|cd\rangle)}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)},
 \end{aligned}$$

and the first Hugenholtz diagram can be simplified to

$$\begin{aligned}
 & \frac{1}{8} \sum_{abcdrs} \frac{\langle ab|rs\rangle \langle rs|cd\rangle \langle cd|ab\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)} \\
 & = \frac{1}{8} \sum_{abcdrs} \frac{(\langle ab|rs\rangle - \langle ab|sr\rangle)(\langle rs|cd\rangle - \langle rs|dc\rangle)(\langle cd|ab\rangle - \langle cd|ba\rangle)}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)} \\
 & = \frac{1}{8} \sum_{abcdrs} \frac{\langle ab|rs\rangle \langle rs|cd\rangle \langle cd|ab\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)} - \frac{\langle ab|rs\rangle \langle rs|cd\rangle \langle cd|ba\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)} \\
 & \quad - \frac{\langle ab|rs\rangle \langle rs|dc\rangle \langle cd|ab\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)} + \frac{\langle ab|rs\rangle \langle rs|dc\rangle \langle cd|ba\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)} \\
 & \quad - \frac{\langle ab|sr\rangle \langle rs|cd\rangle \langle cd|ab\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)} + \frac{\langle ab|sr\rangle \langle rs|cd\rangle \langle cd|ba\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)} \\
 & \quad + \frac{\langle ab|sr\rangle \langle rs|dc\rangle \langle cd|ab\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)} - \frac{\langle ab|sr\rangle \langle rs|dc\rangle \langle cd|ba\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)} \\
 & = \frac{1}{8} \sum_{abcdrs} \frac{\langle ab|rs\rangle \langle cd|ab\rangle \langle rs|cd\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)} - \frac{\langle ab|rs\rangle \langle rs|dc\rangle \langle dc|ba\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)} \\
 & \quad - \frac{\langle ab|rs\rangle \langle sr|cd\rangle \langle cd|ab\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)} + \frac{\langle ab|rs\rangle \langle sr|dc\rangle \langle dc|ba\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)} \\
 & \quad - \frac{\langle ab|rs\rangle \langle sr|cd\rangle \langle cd|ab\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)} + \frac{\langle ab|rs\rangle \langle sr|dc\rangle \langle dc|ba\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)} \\
 & \quad + \frac{\langle ab|rs\rangle \langle sr|dc\rangle \langle cd|ab\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)} - \frac{\langle ab|rs\rangle \langle sr|cd\rangle \langle dc|ba\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_d - \varepsilon_r - \varepsilon_s)}
 \end{aligned}$$

$$\begin{aligned}
&= \sum_{abcrst} \frac{[\langle ab|rs\rangle - \langle ab|sr\rangle][\langle cs|tb\rangle - \langle cs|bt\rangle][\langle rt|ac\rangle - \langle rt|ca\rangle]}{(\varepsilon_a + \varepsilon_b - \varepsilon_s - \varepsilon_r)(\varepsilon_a + \varepsilon_c - \varepsilon_r - \varepsilon_t)} \\
&= \sum_{abcrst} \frac{\langle ab|rs\rangle\langle cs|tb\rangle\langle rt|ac\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_a + \varepsilon_c - \varepsilon_r - \varepsilon_t)} - \frac{\langle ab|rs\rangle\langle cs|tb\rangle\langle rt|ca\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_a + \varepsilon_c - \varepsilon_r - \varepsilon_t)} \\
&\quad - \frac{\langle ab|rs\rangle\langle cs|bt\rangle\langle rt|ac\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_a + \varepsilon_c - \varepsilon_r - \varepsilon_t)} + \frac{\langle ab|rs\rangle\langle cs|bt\rangle\langle rt|ca\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_a + \varepsilon_c - \varepsilon_r - \varepsilon_t)} \\
&\quad - \frac{\langle ab|sr\rangle\langle cs|tb\rangle\langle rt|ac\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_a + \varepsilon_c - \varepsilon_r - \varepsilon_t)} + \frac{\langle ab|sr\rangle\langle cs|tb\rangle\langle rt|ca\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_a + \varepsilon_c - \varepsilon_r - \varepsilon_t)} \\
&\quad + \frac{\langle ab|sr\rangle\langle cs|bt\rangle\langle rt|ac\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_a + \varepsilon_c - \varepsilon_r - \varepsilon_t)} - \frac{\langle ab|sr\rangle\langle cs|bt\rangle\langle rt|ca\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_a + \varepsilon_c - \varepsilon_r - \varepsilon_t)} \\
&= \sum_{abcrst} \frac{\langle ac|rt\rangle\langle bt|sc\rangle\langle rs|ab\rangle}{(\varepsilon_a + \varepsilon_c - \varepsilon_r - \varepsilon_t)(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)} - \frac{\langle bc|rt\rangle\langle at|sc\rangle\langle rs|ab\rangle}{(\varepsilon_b + \varepsilon_c - \varepsilon_t - \varepsilon_r)(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)} \\
&\quad - \sum_{abcrst} \frac{\langle bc|rt\rangle\langle ra|sb\rangle\langle st|ac\rangle}{(\varepsilon_b + \varepsilon_c - \varepsilon_r - \varepsilon_t)(\varepsilon_a + \varepsilon_c - \varepsilon_s - \varepsilon_t)} + \sum_{abcrst} \frac{\langle bc|rt\rangle\langle ar|bs\rangle\langle ts|ac\rangle}{(\varepsilon_c + \varepsilon_b - \varepsilon_r - \varepsilon_t)(\varepsilon_a + \varepsilon_c - \varepsilon_s - \varepsilon_t)} \\
&\quad - \sum_{abcrst} \frac{\langle ab|rs\rangle\langle sc|at\rangle\langle rt|bc\rangle}{(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)(\varepsilon_c + \varepsilon_b - \varepsilon_r - \varepsilon_t)} + \sum_{abcrst} \frac{\langle cb|rt\rangle\langle at|sc\rangle\langle rs|ab\rangle}{(\varepsilon_c + \varepsilon_b - \varepsilon_r - \varepsilon_t)(\varepsilon_a + \varepsilon_b - \varepsilon_r - \varepsilon_s)} \\
&\quad + \sum_{abcrst} \frac{\langle cb|rt\rangle\langle ra|sb\rangle\langle st|ac\rangle}{(\varepsilon_c + \varepsilon_b - \varepsilon_r - \varepsilon_t)(\varepsilon_a + \varepsilon_c - \varepsilon_s - \varepsilon_t)} - \sum_{abcrst} \frac{\langle ac|rt\rangle\langle rb|sc\rangle\langle st|ab\rangle}{(\varepsilon_a + \varepsilon_c - \varepsilon_r - \varepsilon_t)(\varepsilon_a + \varepsilon_b - \varepsilon_s - \varepsilon_t)}.
\end{aligned}$$

These eight terms correspond to the fifth, twelfth, fourth, ninth, eleventh, sixth, tenth, third third-order Goldstone diagrams, respectively.

In conclusion, we have verified that the first third-order Hugenholtz diagram corresponds the second and the seventh third-order Goldstone diagrams, the second third-order Hugenholtz diagram corresponds the first and the eighth third-order Goldstone diagrams, and the third third-order Hugenholtz diagram corresponds the third, fourth, fifth, sixth, ninth, tenth, eleventh, twelfth third-order Goldstone diagrams.

6.7.3 Summation of Diagrams

6.7.4 What Is the Linked Cluster Theorem?

Exercise 6.13

Calculate $E_0^{(3)}$ for a supermolecule consisting of N non-interacting minimal basis H_2 molecules by evaluating the Goldstone diagrams in Table 6.2. Compare your result with that of Eq.(6.92), which was obtained algebraically by explicitly cancelling terms proportional to N^2 .

Hint: Simply show that the value of each Goldstone diagram for the supermolecule is N times the result for a single molecule.

Solution 6.13

Note that in this model, all two-electron integrals involving orbitals from different units are zero, and all third-order terms in the table 6.2 will be zero unless their numerators are non-zero. Thus, each third-order term of this model is just the sum of that of each units, leading that the value of each Goldstone diagram for the supermolecule is N times the result for a single molecule.

6.8 Some Illustrative Calculations