

PHSX 462: HW05

William Jardee

March 21, 2022

Griffiths 5.1

a) Let's start with what we are provided:

$$\vec{r} \equiv \vec{r}_1 - \vec{r}_2 \quad \vec{R} \equiv \frac{m_1 \vec{r}_1 + m_2 \vec{r}_2}{m_1 + m_2} \quad \mu \equiv \frac{m_1 m_2}{m_1 + m_2}$$

then, you know, do the thing:

$$\begin{aligned} \vec{R} &= \frac{(m_1(\vec{r} + \vec{r}_2) + m_2 \vec{r}_2)}{m_1 + m_2} & \vec{R} &= \frac{(m_2(\vec{r}_1 - \vec{r}) + \vec{r}_1 m_1)}{m_1 + m_2} \\ &= \vec{r}_2 + \frac{m_1}{m_2 + m_1} \vec{r} & &= \vec{r}_1 - \frac{m_2}{m_1 + m_2} \vec{r} \\ \vec{r}_2 &= \vec{R} - \frac{\mu}{m_2} \vec{r} & \vec{r}_1 &= \vec{R} + \frac{\mu}{m_1} \vec{r} \end{aligned} \quad \checkmark$$

$$\begin{aligned} \nabla_{\vec{r}_1} &= (\nabla_{\vec{r}_1} \cdot \vec{R}) \nabla_{\vec{R}} + (\nabla_{\vec{r}_1} \cdot \vec{r}) \nabla_{\vec{r}} & \nabla_{\vec{r}_2} &= (\nabla_{\vec{r}_2} \cdot \vec{R}) \nabla_{\vec{R}} + (\nabla_{\vec{r}_2} \cdot \vec{r}) \nabla_{\vec{r}} \\ &= \left(\frac{m_1}{m_1 + m_2} \right) \nabla_{\vec{R}} + \nabla_{\vec{r}} & &= \left(\frac{m_2}{m_1 + m_2} \right) \nabla_{\vec{R}} - \nabla_{\vec{r}} \\ &= \frac{\mu}{m_2} \nabla_{\vec{R}} + \nabla_{\vec{r}} & &= \frac{\mu}{m_1} \nabla_{\vec{R}} - \nabla_{\vec{r}} \end{aligned} \quad \checkmark$$

b)

$$\begin{aligned} (E - V)\Psi &= -\frac{\hbar^2}{2m_1} (\nabla_1)^2 \Psi - \frac{\hbar^2}{2m_2} (\nabla_2)^2 \Psi \\ &= -\frac{\hbar^2}{2m_1} \left(\frac{\mu}{m_2} \nabla_R + \nabla_r \right)^2 \Psi - \frac{\hbar^2}{2m_2} \left(\frac{\mu}{m_1} \nabla_R - \nabla_r \right)^2 \Psi \\ &= -\frac{\hbar^2}{2m_1} \left(\left(\frac{\mu}{m_2} \right)^2 \nabla_R^2 + \frac{\mu}{m_2} \nabla_R \nabla_r + \nabla_r^2 \right) \Psi \\ &\quad - \frac{\hbar^2}{2m_2} \left(\left(\frac{\mu}{m_1} \right)^2 \nabla_R^2 - \frac{\mu}{m_1} \nabla_R \nabla_r + \nabla_r^2 \right) \Psi \\ &= \left(\left[-\frac{\hbar^2}{2m_1} \left(\frac{\mu}{m_2} \right)^2 - \frac{\hbar^2}{2m_2} \left(\frac{\mu}{m_1} \right)^2 \right] \nabla_R^2 + \left[-\frac{\hbar^2}{2m_1} - \frac{\hbar^2}{2m_2} \right] \nabla_r^2 \right) \Psi \\ &= -\frac{\hbar^2}{2} \left[\frac{m_1 + m_2}{(m_1 + m_2)^2} \nabla_R^2 + \frac{m_1 + m_2}{m_1 m_2} \nabla_r^2 \right] \Psi \\ &= -\frac{\hbar^2}{2(m_1 + m_2)} \nabla_R^2 \Psi - \frac{\hbar^2}{2\mu} \nabla_r^2 \Psi \end{aligned} \quad \checkmark$$

- c) Not quite sure what they want us to do exactly here, I think it is just a statement about what the next step to solving would be.

$$\begin{aligned}
& -\frac{\hbar^2}{2(m_1 + m_2)} \nabla_R^2 (\Psi_R \Psi_r) - \frac{\hbar^2}{2\mu} \nabla_r^2 (\Psi_R \Psi_r) + V(\Psi_R \Psi_r) = (E_R + E_r)(\Psi_R \Psi_r) \\
\Psi_r \left(-\frac{\hbar^2}{2(m_1 + m_2)} \right) \nabla_R^2 \Psi_R + \Psi_r \left(-\frac{\hbar^2}{2\mu} \right) \nabla_r^2 \Psi_r + V(\Psi_R \Psi_r) &= (E_R + E_r)(\Psi_R \Psi_r)
\end{aligned}$$

Griffiths 5.2

a) I did a lot of this in python, so I will be just stating the results:

$$m_e \rightarrow 13.6056 \text{ eV}$$

$$\mu \rightarrow 13.5983 \text{ eV}$$

$$\boxed{\% \text{ error} = 0.055\%}$$

b)

$$\text{hydrogen} \rightarrow -1.8886 \text{ eV}$$

$$\text{deutromium} \rightarrow -1.8892 \text{ eV}$$

where the $\mu_{deu} = \frac{m_e(m_p + m_n)}{m_e + m_n + m_p}$. Using the wavelength equation $\lambda = \frac{hc}{E}$

$$\boxed{\Delta\lambda = 0.1788 \text{ nm}}$$

c) with $\frac{m_e(m_n + m_e)}{2m_e + m_n}$

$$\boxed{E_{pos} = -13.5983 \text{ eV}}$$

d) with $\frac{m_\mu m_p}{m_\mu + m_p}$

$$E_{0,\mu} = 2528.51 \text{ eV}$$

$$\Delta E = 1896.38 \text{ eV} \rightarrow \boxed{\lambda = 0.6338 \text{ nm}}$$

Griffiths 5.4

a)

$$\begin{aligned}
 |\Psi_a(\vec{r}_1)\Psi_b(\vec{r}_2)|^2 &= \langle \Psi_a \Psi_b | \Psi_a \Psi_b \rangle \\
 &= \langle \Psi_a | \Psi_a \rangle \langle \Psi_b | \Psi_b \rangle \\
 &= 1
 \end{aligned}$$

Since there are two orthogonal terms, $\left| \frac{1}{A} \Psi_{\pm} \right|^2 = 1 + 1 = 2$. So:

$$A = \frac{1}{\sqrt{2}}$$

b) this one we are going to be a little more clear with:

$$\begin{aligned}
 1 &= A^2 \int [\Psi_a^1 \Psi_a^2 + \Psi_a^2 \Psi_a^1]^* [\Psi_a^1 \Psi_a^2 + \Psi_a^2 \Psi_a^1] \\
 &= A^2 \int [2\Psi_a^1 \Psi_a^2]^* [2\Psi_a^1 \Psi_a^2] \\
 &= 4A^2 \int (\Psi_a^1)^* \Psi_a^1 \int (\Psi_a^2)^* \Psi_a^2 \\
 &= 4A^2
 \end{aligned}$$

$$A = \frac{1}{2}$$

Question 4

- a) The energy of the system can be described as $E_n = E_a + E_b$. Since these particles are distinguishable, they can be in the same state:

$$E_0 = \frac{3}{2}\hbar\omega + \frac{3}{2}\hbar\omega \boxed{= 3\hbar\omega}$$

- b) There are two setups that we can do:

$$\psi_1 = (\hat{a}_+\psi_0(x_1))(\psi_0(x_2)) \qquad \psi_1 = (\psi_0(x_1))(\hat{a}_+\psi_0(x_2))$$

$$\begin{aligned} E_1 &= \left[\frac{1}{2}\hbar\omega \right] 4 + \left[\frac{3}{2}\hbar\omega \right] 2 \\ &= 2\hbar\omega + 3\hbar\omega \\ &\boxed{= 5\hbar\omega} \end{aligned}$$

- c) Only one of the states need to be excited:

$$\begin{array}{ll} \psi_{100}(x_1) \psi_{000}(x_2) & \psi_{000}(x_1) \psi_{100}(x_2) \\ \psi_{010}(x_1) \psi_{000}(x_2) & \psi_{000}(x_1) \psi_{010}(x_2) \\ \psi_{001}(x_1) \psi_{000}(x_2) & \psi_{000}(x_1) \psi_{001}(x_2) \end{array}$$

Question 5

a)

$$\begin{aligned}
 1 &= A^2 \int \left[2 \sin\left(\frac{2\pi}{a}x_1\right) \sin\left(\frac{\pi}{a}x_2\right) \sin\left(\frac{\pi}{a}x_3\right) + 3 \sin\left(\frac{\pi}{a}x_1\right) \sin\left(\frac{2\pi}{a}x_2\right) \sin\left(\frac{5\pi}{a}x_3\right) \right]^* \\
 &\quad \times \left[2 \sin\left(\frac{2\pi}{a}x_1\right) \sin\left(\frac{\pi}{a}x_2\right) \sin\left(\frac{\pi}{a}x_3\right) + 3 \sin\left(\frac{\pi}{a}x_1\right) \sin\left(\frac{2\pi}{a}x_2\right) \sin\left(\frac{5\pi}{a}x_3\right) \right] \\
 &\text{recognizing that none of the states between the two state, so the cross terms are zero} \\
 &= A^2 \int \left[4 \sin^2\left(\frac{2\pi}{a}x_1\right) \sin^2\left(\frac{\pi}{a}x_2\right) \sin^2\left(\frac{\pi}{a}x_3\right) + 9 \sin^2\left(\frac{\pi}{a}x_1\right) \sin^2\left(\frac{2\pi}{a}x_2\right) \sin^2\left(\frac{5\pi}{a}x_3\right) \right] \\
 &\quad \text{remembering that } \int_0^{n\pi} \sin^2(x) = \frac{x}{2} \\
 &= A^2 \left[4 \left(\frac{a}{2}\right)^3 + 9 \left(\frac{a}{2}\right)^3 \right]
 \end{aligned}$$

$$A = \frac{1}{\sqrt{13}} \left(\frac{2}{a}\right)^{3/2}$$

This makes sense, since the second part is the normal coefficient, and the $\sqrt{13}$ accounts for normalizing the two different wavefunctions.

b) The wavefunction of interest is the second one, with the coefficient of 3. So, the probability of having E_3 with this energy is:

$$\left(\frac{3}{\sqrt{13}}\right)^2 = \frac{9}{13}$$

c) Independent of the energy, the average x value is the center of the well. So, we have a $\frac{4}{13}$ chance for $\frac{a}{2}$ and a $\frac{9}{13}$ chance for $\frac{a}{2}$, so $\langle x \rangle = \frac{a}{2}$.

d) We need to find the expectation value of each the energies independently, then add them all together:

$$\left| \frac{4}{13} \begin{bmatrix} 4 \\ 1 \\ 1 \end{bmatrix} + \frac{9}{13} \begin{bmatrix} 1 \\ 4 \\ 25 \end{bmatrix} \right| = \frac{1}{13} \left| \begin{bmatrix} 25 \\ 40 \\ 229 \end{bmatrix} \right| = \frac{294}{13}$$

$$\langle E \rangle \approx \frac{22.62 \hbar^2 \pi^2}{2ma^2}$$

Griffiths 5.8

b) I will do the next part, with the fermion, then remove all the negatives to take the permanent.

$$\begin{aligned}
 & \frac{1}{\sqrt{3!}} \begin{vmatrix} |\Psi_a\rangle_1 & |\Psi_b\rangle_1 & |\Psi_c\rangle_1 \\ |\Psi_a\rangle_2 & |\Psi_b\rangle_2 & |\Psi_c\rangle_2 \\ |\Psi_a\rangle_3 & |\Psi_b\rangle_3 & |\Psi_c\rangle_3 \end{vmatrix} \\
 &= \frac{1}{\sqrt{6}} \left[|\Psi_a\rangle_1 \begin{vmatrix} |\Psi_b\rangle_2 & |\Psi_c\rangle_2 \\ |\Psi_b\rangle_3 & |\Psi_c\rangle_3 \end{vmatrix} - |\Psi_a\rangle_2 \begin{vmatrix} |\Psi_b\rangle_1 & |\Psi_c\rangle_1 \\ |\Psi_b\rangle_3 & |\Psi_c\rangle_3 \end{vmatrix} + |\Psi_a\rangle_3 \begin{vmatrix} |\Psi_b\rangle_1 & |\Psi_c\rangle_1 \\ |\Psi_b\rangle_2 & |\Psi_c\rangle_2 \end{vmatrix} \right] \\
 &= \frac{1}{\sqrt{6}} [|\Psi_a\rangle_1 |\Psi_b\rangle_2 |\Psi_c\rangle_3 - |\Psi_a\rangle_1 |\Psi_c\rangle_2 |\Psi_b\rangle_3 - |\Psi_b\rangle_1 |\Psi_a\rangle_2 |\Psi_c\rangle_3 + |\Psi_c\rangle_1 |\Psi_a\rangle_2 |\Psi_b\rangle_3 \\
 &\quad + |\Psi_b\rangle_1 |\Psi_c\rangle_2 |\Psi_a\rangle_3 - |\Psi_c\rangle_1 |\Psi_b\rangle_2 |\Psi_a\rangle_3]
 \end{aligned}$$

So, for a bosonic system:

$$\begin{aligned}
 & \frac{1}{\sqrt{6}} [|\Psi_a\rangle_1 |\Psi_b\rangle_2 |\Psi_c\rangle_3 + |\Psi_a\rangle_1 |\Psi_c\rangle_2 |\Psi_b\rangle_3 + |\Psi_b\rangle_1 |\Psi_a\rangle_2 |\Psi_c\rangle_3 + |\Psi_c\rangle_1 |\Psi_a\rangle_2 |\Psi_b\rangle_3 \\
 &\quad + |\Psi_b\rangle_1 |\Psi_c\rangle_2 |\Psi_a\rangle_3 + |\Psi_c\rangle_1 |\Psi_b\rangle_2 |\Psi_a\rangle_3]
 \end{aligned}$$

c) Stealing the answer from the last part, before changing the signs:

$$\begin{aligned}
 & \frac{1}{\sqrt{6}} [|\Psi_a\rangle_1 |\Psi_b\rangle_2 |\Psi_c\rangle_3 - |\Psi_a\rangle_1 |\Psi_c\rangle_2 |\Psi_b\rangle_3 - |\Psi_b\rangle_1 |\Psi_a\rangle_2 |\Psi_c\rangle_3 + |\Psi_c\rangle_1 |\Psi_a\rangle_2 |\Psi_b\rangle_3 \\
 &\quad + |\Psi_b\rangle_1 |\Psi_c\rangle_2 |\Psi_a\rangle_3 - |\Psi_c\rangle_1 |\Psi_b\rangle_2 |\Psi_a\rangle_3]
 \end{aligned}$$

Griffiths 5.9

a) I just brute forced these bad bois to find all of them.

For the first energy: $E_0 = 2K$, degeneracy 1

$$\Psi_1(x_1)\Psi_1(x_2) |\uparrow\downarrow\rangle - \Psi_1(x_1)\Psi_1(x_2) |\downarrow\uparrow\rangle$$

for the second energy: $E_1 = 5K$, degeneracy 4

$$\begin{aligned} & \Psi_1(x_1)\Psi_2(x_2) |\downarrow\downarrow\rangle - \Psi_2(x_1)\Psi_1(x_2) |\downarrow\downarrow\rangle \\ & (\Psi_1(x_1)\Psi_2(x_2)) \frac{1}{\sqrt{2}}(|\uparrow\downarrow\rangle + |\downarrow\uparrow\rangle) - (\Psi_2(x_1)\Psi_1(x_2)) \frac{1}{\sqrt{2}}(|\uparrow\downarrow\rangle + |\downarrow\uparrow\rangle) \\ & \Psi_1(x_1)\Psi_2(x_2) |\uparrow\uparrow\rangle - \Psi_2(x_1)\Psi_1(x_2) |\uparrow\uparrow\rangle \\ & (\Psi_1(x_1)\Psi_2(x_2)) \frac{1}{\sqrt{2}}(|\uparrow\downarrow\rangle - |\downarrow\uparrow\rangle) + (\Psi_2(x_1)\Psi_1(x_2)) \frac{1}{\sqrt{2}}(|\uparrow\downarrow\rangle - |\downarrow\uparrow\rangle) \end{aligned}$$

for the third energy: $E_2 = 8k$, degeneracy 1

$$\Psi_2(x_1)\Psi_2(x_2) |\uparrow\downarrow\rangle - \Psi_2(x_1)\Psi_2(x_2) |\downarrow\uparrow\rangle$$

for the second energy: $E_3 = 10K$, degeneracy 4

$$\begin{aligned} & \Psi_1(x_1)\Psi_3(x_2) |\downarrow\downarrow\rangle - \Psi_3(x_1)\Psi_1(x_2) |\downarrow\downarrow\rangle \\ & (\Psi_1(x_1)\Psi_3(x_2)) \frac{1}{\sqrt{2}}(|\uparrow\downarrow\rangle + |\downarrow\uparrow\rangle) - (\Psi_3(x_1)\Psi_1(x_2)) \frac{1}{\sqrt{2}}(|\uparrow\downarrow\rangle + |\downarrow\uparrow\rangle) \\ & \Psi_1(x_1)\Psi_3(x_2) |\uparrow\uparrow\rangle - \Psi_3(x_1)\Psi_1(x_2) |\uparrow\uparrow\rangle \\ & (\Psi_1(x_1)\Psi_3(x_2)) \frac{1}{\sqrt{2}}(|\uparrow\downarrow\rangle - |\downarrow\uparrow\rangle) + (\Psi_3(x_1)\Psi_1(x_2)) \frac{1}{\sqrt{2}}(|\uparrow\downarrow\rangle - |\downarrow\uparrow\rangle) \end{aligned}$$