## Week 5

## Approximate Solutions of the Schrödinger Equation

## 5.1 Approximation Methods

10/25:

- The **variational method** and **perturbation theory** are two methods of approximating solutions to Schrödinger equations describing systems more complex than the hydrogen atom.
- To begin our investigation of the variational method, we will look at the particle in a box.
  - Consider a Hamiltonian for an electron in a box of length L=2 a.u. centered around x=0.
    - Note that we take the electron as the fundamental mass,  $\hbar$  as the fundamental unit of energy time, and the charge of the electron as the fundamental unit of charge, and the Bohr radius as the fundamental unit of length.
  - Our Hamiltonian is

$$H\psi(x) = -\frac{\hbar^2}{2m} \frac{\mathrm{d}^2}{\mathrm{d}x^2} \psi(x)$$

or, in atomic units,

$$H\psi(x) = -\frac{1}{2}\frac{\mathrm{d}^2}{\mathrm{d}x^2}\psi(x)$$

• Variational theorem: The expectation value of our Hamiltonian with respect to a trial wave function produces an approximate energy. Moreover<sup>[1]</sup>,

$$E_{\rm approx} \ge E_{\rm gr. st.}$$

- Variational method: Take  $\psi_{\text{trial}} = \sum_n a_n |\psi_n\rangle$  where  $\psi_n$  is a trial wave function and the  $a_j$ 's are parameters of the wave function which we want to optimize to lower  $E_{\text{trial}}$ .
  - Dirac's ket describes an abstract state of the particle (possibly position, possibly its Fourier transform, momentum).
- Back to the particle in a box:
  - A possible trial wave function (that satisfies the boundary conditions) is

$$\psi_{\rm tr} = (1+x)(1-x) = 1-x^2$$

<sup>&</sup>lt;sup>1</sup>We will prove that the approximate energy is an upper bound on the ground state energy in the homework.

- The energy of  $\psi_{\rm tr}$  may be evaluated as follows.

$$E = \frac{\int \psi_{\text{tr}}^*(x) \hat{H} \psi_{\text{tr}}(x) \, dx}{\int \psi_{\text{tr}}^*(x) \psi_{\text{tr}}(x) \, dx}$$

$$= \frac{\int_{-1}^{1} (1 - x^2) \left( -\frac{1}{2} \frac{d^2}{dx^2} \right) (1 - x^2) \, dx}{\int_{-1}^{1} (1 - x^2) (1 - x^2) \, dx}$$

$$= \frac{\int_{-1}^{1} (1 - x^2) \, dx}{\int_{-1}^{1} (1 - x^2) (1 - x^2) \, dx}$$

$$= \frac{4/3}{16/15}$$

$$= \frac{5}{4}$$

$$= 1.25 \text{ a.u.}$$

- From the exact solution to the particle in a box

$$E_1 = 1.23370055 < 1.25 = E_{\text{trial}}$$

so the variational theorem is satisfied.

- Next step: Trial wave function as a linear combination is  $\psi_{\rm tr}(x) = c_1 \psi_1(x) + c_2 \psi_2(x)$ .
- Plugging this into the SE yields

$$c_1(\hat{H} - E)\psi_1(x) + c_2(\hat{H} - E)\psi_2(x) = 0$$

- $\blacksquare \psi_1, \psi_2$  span the (Hilbert) space of solutions.
- To solve the above equation, multiply by  $\psi_1(x)$  and integrate to obtain

$$c_1 \int_{-1}^{1} \psi_1^*(x)(\hat{H} - E)\psi_1(x) dx + c_2 \int_{-1}^{1} \psi_1^*(x)(\hat{H} - E)\psi_2(x) dx = 0$$

and multiply by  $\psi_2(x)$  an integrate to obtain

$$c_1 \int_{-1}^{1} \psi_2^*(x)(\hat{H} - E)\psi_1(x) \, dx + c_2 \int_{-1}^{1} \psi_2^*(x)(\hat{H} - E)\psi_2(x) \, dx = 0$$

- Substituting, we have

$$c_1(H_{11} - ES_{11}) + c_2(H_{12} - ES_{12}) = 0$$
  $c_1(H_{21} - ES_{21}) + c_2(H_{22} - ES_{22}) = 0$ 

- In matrix form, the above two equations become

$$\begin{pmatrix} H_{11} & H_{12} \\ H_{21} & H_{22} \end{pmatrix} \begin{pmatrix} c_1 \\ c_2 \end{pmatrix} - E \begin{pmatrix} S_{11} & S_{12} \\ S_{21} & S_{22} \end{pmatrix} \begin{pmatrix} c_1 \\ c_2 \end{pmatrix} = \begin{pmatrix} 0 \\ 0 \end{pmatrix}$$
$$\mathbb{H}\vec{c} - E \mathbb{S}\vec{c} = 0$$

- We get a matrix that the same dimension as the size of the expansion (in the first case, we had a  $1 \times 1$  matrix).
- S is the overlap matrix because the wave functions aren't normalized.

## 5.2 Variational Method

10/27: • Approximating the ground state energy with some trial wave function and applying

$$E_{\text{approx}} = \frac{\int \psi_{\text{tr}}^* \hat{H} \psi_{\text{tr}} \, dx}{\int \psi_{\text{tr}}^* \psi_{\text{tr}} \, dx}$$

where

$$\psi_{\rm tr} = \sum_{n} c_n \psi_n(x)$$

- Example 2:
- For our second term, we need another even function (since the ground state wavefunction is even). Thus, choose

$$\psi_{\rm tr}(x) = c_1(1-x^2) + c_2(1-x^2)x^2$$

- Think about this in the context of power series we have  $(1-x^2)$  times an even power series expansion  $(c_1 + c_2x^2)$ .
- To find  $c_1, c_2$ , we could plug into the approximation integral and minimize.
- Alternatively, we can use matrices. We essentially project the Schrödinger equation onto the space of the two wave functions.
- Take  $\hat{H}\psi = E\psi$  and expand it to  $\hat{H}(c_1\psi_1 + c_2\psi_2) = E(c_1\psi_1 + c_2\psi_2)$ . In matrix form,  $\mathbb{H}\vec{c} = E\mathbb{S}\vec{c}$ .
- We have an overlap matrix S because our wave functions aren't normalized. If the basis is orthonormal, S collapses to the identity matrix.
  - Each  $s_{ij}$  equals

$$s_{ij} = \int \psi_i^* \psi_j \, \mathrm{d}x$$

- If  $\psi_1, \psi_2$  is orthonormal, then  $s_{ij} = \delta_{ij}$ .
- The elements of the Hamiltonian matrix:

$$H_{11} = \int \psi_1^*(x) \hat{H} \psi_1(x) dx \qquad H_{12} = \int \psi_1^*(x) \hat{H} \psi_2(x) dx$$
$$= \frac{4}{3} \qquad = \frac{8}{15}$$

$$H_{21} = \frac{8}{15} \qquad \qquad H_{22} = \frac{44}{105}$$

- Notice that  $\mathbb{H}$  is symmetric with  $H_{12} = H_{21}$ .
- $\bullet\,$  Elements of the overlap matrix:

$$S_{11} = \frac{16}{15} \qquad \qquad S_{12} = \frac{32}{105}$$

$$S_{21} = \frac{32}{105} \qquad \qquad S_{22} = \frac{16}{315}$$

- Notice that  $\mathbb{S}$  is symmetric with  $S_{12} = S_{21}$ .
- Note that there are multiple ways to solve  $\mathbb{H}\vec{c} = E\mathbb{S}\vec{c}$ ; McQuarrie and Simon (1997) only teaches one. Thus, you can get computers to do the math and solve far bigger systems than you could by hand.

- Solving  $\mathbb{H}\vec{c} = E\mathbb{S}\vec{c}$  with the textbook method:
  - Rewrite as  $(\mathbb{H} E\mathbb{S})\vec{c} = 0$ . Find the null space of  $\mathbb{H} E\mathbb{S}$ .
  - Since the determinant is the product of the eigenvalues,  $\det(\mathbb{H} E\mathbb{S}) = (E_1 E)(E_2 E)$ .
  - This determinant is equal to zero only when  $E = E_1$  or  $E = E_2$ .
    - The energy is becoming quantized because of the linear algebra!
  - Now taking  $det(\mathbb{H} E\mathbb{S})$  gives a characteristic polynomial in E.

$$0 = \begin{vmatrix} \frac{4}{3} - \frac{16}{15}E & \frac{4}{15} - \frac{16}{105}E \\ \frac{4}{15} - \frac{16}{105}E & \frac{44}{105} - \frac{16}{315}E \end{vmatrix}$$
$$= \frac{256}{525} - \frac{2048}{4725}E + \frac{1024}{33075}E^2$$

- Solving the quadratic gives us

$$E = 7 \pm \frac{\sqrt{133}}{2}$$

- Thus,

$$E_1 = 1.233718705 \,\mathrm{a.u.}$$

$$E_2 = 12.766 \,\mathrm{a.u.}$$

- Notice that the  $E_1$  we found is only *marginally* greater than the real value of  $E_1$ . Our value is accurate to four decimal places!
- Solving for  $\vec{c}$  with  $E_1$  gives us

$$\vec{c}_1 = -0.9764$$

$$\vec{c}_2 = 0.2156$$