## Quantum Mechanics 2

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## Chapter 1

## Orbital Angular Momentum

## 1.1 Basics of QM

Let us recall some basic facts of quantum mechanics.

The expectation value of an observable A with an associated operator  $\hat{A}$  is given by,

$$\langle \hat{A} \rangle = \langle \Psi | \hat{A} | \Psi \rangle = \int \Psi^* \hat{A} \Psi \, d\mathbf{r} \,.$$
 (1.1)

The fundamental position, momentum, and angular momentum operators are defined as follows,

#### **Definition 1: Fundamental Operators**

$$\hat{\mathbf{r}} = x\hat{\mathbf{x}} + y\hat{\mathbf{y}} + z\hat{\mathbf{z}} \tag{1.2}$$

$$\hat{p} = -i\hbar \nabla \tag{1.3}$$

$$\hat{L}_i = \varepsilon_{ijk} \hat{r}_j \hat{p}_k \tag{1.4}$$

The Hamiltonian is defined,

#### Definition 2: Hamiltonian

$$\hat{H} = \hat{T} + \hat{V} = -\frac{\hbar^2}{2m} \nabla^2 + V(\mathbf{r}, t). \tag{1.5}$$

We obtain the wavefunction  $\Psi$  by solving the TDSE,

#### **Definition 3: Time Dependent Schrodinger Equation**

$$i\hbar \frac{\partial \Psi(\mathbf{r},t)}{\partial t} = \hat{H}\Psi(\mathbf{r},t).$$
 (1.6)

For the static case, this reduces to the TISE,

$$\hat{H}\Psi = E\Psi. \tag{1.7}$$

If  $\Psi(\mathbf{r},0)$  is written in the energy eigenbasis, i.e.,  $\Psi(\mathbf{r},0) = \sum_{i} c_{i} |E_{i}\rangle$ , then the time-dependent solution is trivial,

$$\Psi(\mathbf{r},t) = \sum_{i} c_i |E_i\rangle \exp\left(\frac{-iE_it}{\hbar}\right). \tag{1.8}$$

#### 1.1.1 The Simple Harmonic Oscillator

The SHO has a Hamiltonian,

$$\hat{H} = -\frac{\hbar^2}{2m} \frac{\mathrm{d}^2}{\mathrm{d}x^2} + \frac{1}{2} m\omega^2 x^2 \tag{1.9}$$

with energy eigenvalues,

$$E_n = \left(n + \frac{1}{2}\right)\hbar\omega\tag{1.10}$$

and has normalised Eigenfunctions,

$$\psi_n(x) = \left(\frac{1}{n!2^n a\sqrt{\pi}}\right) H_n\left(\frac{x}{a}\right) \exp\left(-\frac{x^2}{2a^2}\right)$$
(1.11)

where  $a = \sqrt{\hbar/m\omega}$  and  $H_n(x/a)$  is a Hermite polynomial.

#### 1.1.2 Simple Perturbation Theory

In simple perturbation theory, we write the Hamiltonian as,

$$\hat{H} = \hat{H}_0 + \hat{V} \tag{1.12}$$

where the Hamiltonian  $\hat{H}_0$  is trivial and for which we already have obtained its eigenfunction  $\psi$  and eigenvalues  $E_n^{(0)}$ . We then use this to find the expectation value of the total Hamiltonian,

$$\langle \hat{H} \rangle = \langle \psi | \hat{H}_0 + \hat{V} | \psi \rangle = E_n^{(0)} + \Delta E.$$
 (1.13)

Writing this more explicitly,

#### Definition 4: First Order Perturbation Theory

$$E_n = E_n^{(0)} + \langle \psi | \hat{V} | \psi \rangle \tag{1.14}$$

### 1.2 Particle in 2D SHO

The Hamiltonian of the 2D SHO is given by,

$$\hat{H}\psi(x,y) = -\frac{\hbar^2}{2m} \left( \frac{\partial^2}{\partial x^2} + \frac{\partial^2}{\partial y^2} \right) + \frac{1}{2} m\omega(x^2 + y^2)\psi(x,y) = E\psi(x,y)$$
(1.15)

We can separate this Hamiltonian into its x and y components,

$$\hat{H}_x = -\frac{\hbar^2}{2m} \frac{\partial^2}{\partial x^2} + \frac{1}{2} m\omega x^2 \qquad \qquad \hat{H}_y - \frac{\hbar^2}{2m} \frac{\partial^2}{\partial y^2} + \frac{1}{2} m\omega y^2. \tag{1.16}$$

We know the solution to the 1D SHO, as by eq. (1.10). We can intuit that the total solution of the 2D Hamiltonian will be a product of the two 1D wavefunctions. This comes from the fact that to add probabilities, we multiply the probability densities. So, we write,

$$\hat{H}\psi_{n_x}(x)\psi_{n_y}(y) = \left(\hat{H}_x + \hat{H}_y\right)\psi_{n_x}(x)\psi_{n_y}(y) 
= \left(\hat{H}_x\psi_{n_x}(x)\right)\psi_{n_y}(y) + \psi_{n_x}(x)\left(\hat{H}_y\psi_{n_y}(y)\right) 
= \left(n_x + \frac{1}{2}\right)\hbar\omega\psi_{n_y}(y) + \left(n_y + \frac{1}{2}\right)\hbar\omega\psi_{n_x}(x) 
= (n_x + n_y + 1)\hbar\omega\psi_{n_x}(x)\psi_{n_y}(y) 
\Longrightarrow E_{n_x,n_y} = (n_x + n_y + 1)\hbar\omega.$$
(1.17)

#### 1.2.1 Degeneracy

This is when there is more than one state with the same energy. The degeneracy D is the number of energy states that share the same energy. Non-degenerate states are those with D = 1.

### 1.3 Orbital Angular Momentum

The angular momentum in given direction in a classical system is given by,

$$L_i = \varepsilon_{ijk} r_j p_k. \tag{1.18}$$

The angular momentum operator in quantum mechanics is thus,

$$\hat{L}_i = \varepsilon_{ijk} \hat{r}_i \hat{p}_k. \tag{1.19}$$

We are particularly interested in the case where i=z, in which case the operator becomes,

$$\hat{L}_z = \hat{x}\hat{p}_y - \hat{y}\hat{p}_x = -i\hbar \left( x \frac{\partial}{\partial y} - y \frac{\partial}{\partial x} \right). \tag{1.20}$$

Let us consider this operator in plane polar coordinates,  $(r, \theta)$ . We have,

$$x = r\cos\theta \qquad \qquad y = r\sin\theta \tag{1.21}$$

Let us consider the following,

$$\frac{\partial}{\partial \theta} = \frac{\partial x}{\partial \theta} \frac{\partial}{\partial x} + \frac{\partial y}{\partial \theta} \frac{\partial}{\partial y} = -r \sin \theta \frac{\partial}{\partial x} + r \cos \theta \frac{\partial}{\partial y} 
= -y \frac{\partial}{\partial x} + x \frac{\partial}{\partial y}.$$
(1.22)

So, in plane polars,

#### Definition 5: Angular Momentum Operator in Z

$$\hat{L}_z = -i\hbar \frac{\partial}{\partial \theta}.\tag{1.23}$$

## 1.3.1 Eigenfunctions and Eigenvalues of $\hat{L}_z$

We wish to consider the following,

$$\hat{L}_z \Theta(\theta) = L_z \Theta(\theta). \tag{1.24}$$

So,

$$-i\hbar \frac{\mathrm{d}\Theta}{\mathrm{d}\theta} = L_z\Theta \tag{1.25}$$

which we can solve trivially,

$$\Theta(\theta) = Ae^{\frac{L_z \theta}{\hbar}} \tag{1.26}$$

where  $A = \frac{1}{\sqrt{2\pi}}$  is a normalisation constant. We require a cyclic boundary condition, such that  $\Theta(\theta) = \Theta(\theta + 2\pi)$ . So,

$$Ae^{\frac{iL_z(\theta+2\pi)}{\hbar}} = Ae^{\frac{iL_z\theta}{\hbar}}$$

$$e^{\frac{iL_z^2\pi}{\hbar}} = 1.$$
(1.27)

Not all values of  $L_z$  satisfy the eq. (1.27), so we have to impose the following restriction,

$$L_z = \hbar m, \quad m \in \mathbb{Z}$$
 (1.28)

and thus, we can write the angular momentum eigenfunction as,

#### Definition 6: Angular Momentum Eigenfunction

$$\Theta_m(\theta) = \frac{1}{\sqrt{2\pi}} e^{im\theta} \tag{1.29}$$

### 1.3.2 Angular Momentum of the 2D SHO

We wish to express eigenfunctions of the 2D SHO as eigenfunctions of angular momentum. we will find that we require a combination of all degenerate eigenfunctions for a givevn D in order to represent angular momentum eigenfunction. Observing the ground state,

$$\Psi_{00}(x,y) = e^{-\frac{x^2}{2a^2}} \cdot e^{-\frac{y^2}{2a^2}} = e^{-\frac{r^2}{2a^2}}, \qquad a^2 = \frac{\hbar}{2m}.$$
 (1.30)

Applying the angular momentum operator we find,

$$\hat{L}_z \Psi_{00} = 0 \cdot \Psi_{00} \tag{1.31}$$

which holds, as 0 is an allowed value of m. The first excited states of D=2 are given by,

$$\Psi_{10} = xe^{-\frac{x^2}{2a^2}} \cdot e^{-\frac{y^2}{2a^2}} \qquad \qquad \Psi_{01} = e^{-\frac{x^2}{2a^2}} \cdot ye^{-\frac{y^2}{2a^2}}$$
(1.32)

which we combine to form,

$$\Psi_{\pm} = \Psi_{10} \pm i\Psi_{01}$$

$$= [r\cos\theta \pm ir\sin\theta] e^{-\frac{r^2}{2a^2}} = re^{\pm i\theta} e^{-\frac{r^2}{2a^2}}.$$
(1.33)

Applying  $\hat{L}_z$  to eq. (1.33),

$$\hat{L}_z \Psi_{\pm} = \pm \hbar \Psi_{\pm} \tag{1.34}$$

 $\Longrightarrow \Psi_{\pm}$  is an eigenfunction of  $\hat{L}_z$  with eigenvalues  $\pm \hbar$ . Furthermore,  $\Psi_{\pm}$  is an eigenfunction of  $\hat{H}$ , so  $\hat{H}$  and  $\hat{L}_z$  commute. This allows for the 2D SHO to be described by *good quantum numbers*. These satisfy the following,

- 1. Can be known simultaneously,
- 2. Fully and uniquely specify the state of a system.

For the 2D SHO, its good quantum numbers are (n, m), where  $n = n_x + n_y$ . n specifies the energy of the system (as by  $E_n = (n+1)\hbar\omega$ ), and m specifies the angular momentum of the system (as by  $L_z = m\hbar$ ).