

Lecture Notes: Quantum Mechanics

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Chapter 1

General principles of quantum theory

1.1 Classical-Mechanical Quantities Are Represented by Linear Operators in Quantum Mechanics

In quantum mechanics, operators play a fundamental role in describing physical observables and the dynamics of quantum systems. In quantum mechanics, physical quantities such as position, momentum, energy, and angular momentum are represented by mathematical operators. These operators act on the wave function of a quantum system and yield the observable values of these quantities when applied to a wave function.

Postulate 2 To every observable in classical mechanics there corresponds a linear operator in quantum mechanics.

An operator \hat{A} is a mathematical entity that transforms a function $f(x)$ into another function $g(x)$

$$\hat{A}f(x) = g(x).$$

We usually denote an operator by a capital letter with a carat over it (e.g., \hat{A}). The operator \hat{A} operates on $f(x)$ to give a new function $g(x)$.

An operator is a symbol that tells you to do something to whatever follows the symbol. For example, we can consider $\frac{df}{dx}$ to be the $\frac{d}{dx}$ operator operating on the function $f(x)$.

Classical-Mechanical Observables and Their Corresponding Quantum-Mechanical Operators

Observable		Operator	
Name	Symbol	Symbol	Operation
Position	x	\hat{X}	Multiply by x
	\mathbf{r}	$\hat{\mathbf{R}}$	Multiply by \mathbf{r}
Momentum	p_x	\hat{P}_x	$-i\hbar \frac{\partial}{\partial x}$
	\mathbf{p}	$\hat{\mathbf{P}}$	$-i\hbar(\mathbf{i} \frac{\partial}{\partial x} + \mathbf{j} \frac{\partial}{\partial y} + \mathbf{k} \frac{\partial}{\partial z})$
Kinetic energy	T_x	\hat{T}_x	$-\frac{\hbar^2}{2m} \frac{\partial^2}{\partial x^2}$
	T	\hat{T}	$-\frac{\hbar^2}{2m} (\frac{\partial^2}{\partial x^2} + \frac{\partial^2}{\partial y^2} + \frac{\partial^2}{\partial z^2})$
			$= -\frac{\hbar^2}{2m} \nabla^2$
Potential energy	$V(x)$	$\hat{V}(\hat{x})$	Multiply by $V(x)$
	$V(x, y, z)$	$\hat{V}(\hat{x}, \hat{y}, \hat{z})$	Multiply by $V(x, y, z)$
Total energy	E	\hat{H}	$-\frac{\hbar^2}{2m} (\frac{\partial^2}{\partial x^2} + \frac{\partial^2}{\partial y^2} + \frac{\partial^2}{\partial z^2}) + V(x, y, z)$
			$= -\frac{\hbar^2}{2m} \nabla^2 + V(x, y, z)$
	$l_x = yp_z - zp_y$	\hat{l}_x	$-i\hbar(y \frac{\partial}{\partial z} - z \frac{\partial}{\partial y})$
$l_y = zp_x - xp_z$	\hat{l}_y		$-i\hbar(z \frac{\partial}{\partial x} - x \frac{\partial}{\partial z})$
	$l_z = xp_y - yp_x$	\hat{l}_z	$-i\hbar(x \frac{\partial}{\partial y} - y \frac{\partial}{\partial x})$

Some other examples are SQR (square what follows), $\int_0^1 dx$ (integrate what follows from 0 to 1), 3 (multiply by 3).

HW1: Perform the following operations

- (a) $\hat{A}(2x)$, $\hat{A} = \frac{d^2}{dx^2}$
- (b) $\hat{A}(x^2)$, $\hat{A} = \frac{d^2}{dx^2} + 2\frac{d}{dx} + 3$
- (c) $\hat{A}(xy^3)$, $\hat{A} = \frac{\partial}{\partial y}$
- (d) $\hat{A}(e^{ikx})$, $\hat{A} = -i\hbar \frac{d}{dx}$

In quantum mechanics, we deal only with **linear operators**. The operator \hat{A} is linear if it satisfies the following criteria

$$\hat{A}[c_1f_1(x) + c_2f_2(x)] = c_1\hat{A}f_1(x) + c_2\hat{A}f_2(x) \quad (1.1.1)$$

where c_1 and c_2 are (possibly complex) constants. Clearly, the "differentiate" and "integrate" operators are linear because

$$\frac{d}{dx}[c_1f_1(x) + c_2f_2(x)] = c_1\frac{df_1}{dx} + c_2\frac{df_2}{dx}$$

and

$$\int [c_1f_1(x) + c_2f_2(x)] dx = c_1 \int f_1(x) dx + c_2 \int f_2(x) dx.$$

The "square" operator, SQR, on the other hand, is nonlinear because

$$\begin{aligned} \text{SQR}[c_1f_1(x) + c_2f_2(x)] &= c_1^2f_1^2(x) + c_2^2f_2^2(x) + 2c_1c_2f_1(x)f_2(x) \\ &\neq c_1f_1^2(x) + c_2f_2^2(x) \end{aligned}$$

and therefore it does not satisfy the definition given by Equation 1.1.1.

HW2: Determine whether the following operators are linear or nonlinear:

(a) $\hat{A}f(x) = \text{SQRT } f(x)$ (**take the square root**)

(b) $\hat{A}f(x) = x^2f(x)$

1.2 Eigenfunctions and eigenvalues

In general, the function $\phi(x)$ is obtained by the application of the operator \hat{A} on an arbitrary function $\psi(x)$. The function $\phi(x)$ is linearly independent of $\psi(x)$. However, for some particular function $\psi_1(x)$, it is possible that

$$\hat{A}\psi_1(x) = \alpha_1\psi_1(x)$$

where α_1 is a complex number. In such a case $\psi_1(x)$ is said to be an **eigenfunction** of \hat{A} and α_1 is the corresponding **eigenvalue**. The problem of determining $\psi_1(x)$ and α_1 for a given \hat{A} is called an **eigenvalue problem**.

HW3: (a) Show that $e^{\alpha x}$ is an eigenfunction of the operator $\frac{d^n}{dx^n}$. What is the eigenvalue?

(b) Show that e^{ikx} is an eigen function of the operator $\hat{P}_x = -i\hbar\frac{\partial}{\partial x}$. What is the eigenvalue?

For a given operator \hat{A} , many eigenfunctions may exist, so that

$$\hat{A}\psi_i(x) = \alpha_i\psi_i(x)$$

where $\psi_i(x)$ are the eigenfunctions, which may even be infinite in number, and α_i are the corresponding eigenvalues. Each eigenfunction of \hat{A} is unique, that is to say, is **linearly independent** of the other eigenfunctions.

If $f(x) \neq cg(x)$ for any real number $c \neq 0$, we say $f(x)$ and $g(x)$ are linearly independent functions. Suppose $f(x) = 2x$ and $g(x) = 3x$, then both the functions are said to be called linearly dependent because $f(x) = \frac{2}{3}g(x)$. The functions $f(x) = 2x$ and $g(x) = 3x^2$ are linearly independent as $f(x) \neq cg(x)$. **Moreover, wavefunctions of a particle confined in a one-dimensional box are linearly independent.**

Sometimes two or more eigenfunctions have the same eigenvalue. In that situation the eigenvalue is said to be **degenerate**. When two, three, ..., n eigenfunctions have the same eigenvalue, the

eigenvalue is doubly, triply, . . . , n -fold degenerate. When an eigenvalue corresponds only to a single eigenfunction, the eigenvalue is non-degenerate.

1.3 The Schrodinger Equation and a Particle in a Box

Let us consider a particle of mass m confined to a box along the x axis between $x = 0$ and $x = a$. This case is called the problem of a **particle in a one-dimensional box**. This simple model has at least a crude application to the re electrons in a linear conjugated hydrocarbon.

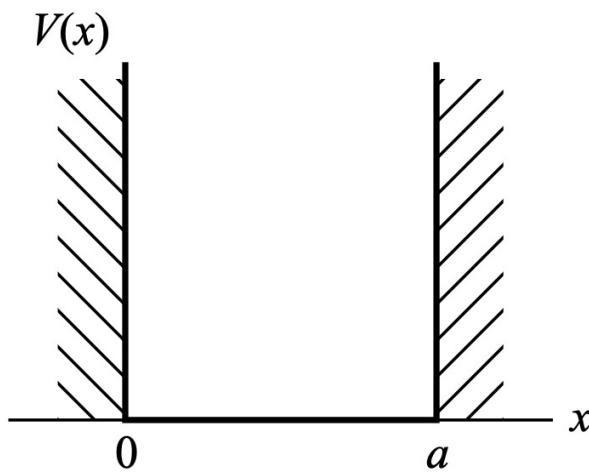


Figure 1.3.1: The potential energy $V(x)$ for a particle in a one-dimensional box of length a .

The potential energy for such a particle is given by $V(x) = \begin{cases} 0, & \text{if } 0 \leq x \leq a \\ \infty, & \text{if } x < 0, x > a \end{cases}$

The momentum operator is

$$\begin{aligned}\hat{p}_x &= -i\hbar \frac{d}{dx} \\ \hat{p}_x^2 &= \left(-i\hbar \frac{d}{dx}\right) \cdot \left(-i\hbar \frac{d}{dx}\right) \\ &= -\hbar^2 \frac{d^2}{dx^2}\end{aligned}$$

and the Hamiltonian operator is

$$\hat{H} = \frac{\hat{p}_x^2}{2m} + \hat{V}(\hat{x})$$

$$= -\frac{\hbar^2}{2m} \frac{d^2}{dx^2} + V(x)$$

Our aim is finding the acceptable **wavefunction** and **energy** of the confined particle solving the **Time-Independent Schrödinger Equation (TISE)**

$$\hat{H}\psi(x) = E\psi(x)$$

Outside the potential well, the **TISE** is given by

$$-\frac{\hbar^2}{2m} \frac{d^2\psi(x)}{dx^2} + \infty\psi(x) = E\psi(x)$$

for which the solution is simply $\psi(x) = 0$; the probability is zero for finding the particle outside the box where the potential is infinite.

Inside the box, the **TISE** is

$$\begin{aligned} -\frac{\hbar^2}{2m} \frac{d^2\psi(x)}{dx^2} &= E\psi(x) \\ \frac{d^2\psi(x)}{dx^2} &= -\frac{2mE}{\hbar^2}\psi(x) \\ \frac{d^2\psi(x)}{dx^2} + \frac{2mE}{\hbar^2}\psi(x) &= 0 \\ \frac{d^2\psi(x)}{dx^2} + k^2\psi(x) &= 0 \end{aligned} \tag{1.3.1}$$

$$k = \sqrt{\frac{2mE}{\hbar^2}} \tag{1.3.2}$$

The general solution of the homogeneous second order linear differential equation with constant coefficients (eq. 1.3.1) is

$$\psi(x) = A \cos(kx) + B \sin(kx) \tag{1.3.3}$$

where A and B are arbitrary constants (real or complex) of integration.

The constants A and B are determined by the *boundary conditions* placed on the solution $\psi(x)$. Since $\psi(x)$ must be continuous, the boundary conditions require that $\psi(x)$ vanish at each end

of the box so as to match the value of $\psi(x)$ outside the box, i.e., $\psi(0) = \psi(a) = 0$. At $x = 0$, the function $\psi(0)$ from eq. (1.3.3) is

$$\psi(0) = A \cos(0) + B \sin(0) = A$$

so that $A = 0$ and $\psi(x)$ is now

$$\psi(x) = B \sin(kx)$$

At $x = a$, $\psi(a)$ is

$$\psi(a) = B \sin(ka) = 0$$

The constant B cannot be zero, for then $\psi(x)$ would vanish everywhere and there would be no particle. Consequently, we have $\sin(ka) = 0$ or

$$ka = n\pi \quad n = 0, \pm 1, \pm 2, \pm 3, \dots \quad (1.3.4)$$

Important to note that the value of n is restricted to any positive integer greater than zero. The solution $n = 0$ would cause to vanish everywhere and is therefore not acceptable. Negative values of n give redundant solutions because $\sin(-\theta)$ equals $-\sin(\theta)$.

By using eq. (1.3.2) and (1.3.4), we find that

$$\begin{aligned} E_n &= \frac{k^2 \hbar^2}{2m} \\ &= \frac{\hbar^2}{2m} \frac{n^2 \pi^2}{a^2} \\ &= \frac{n^2 \pi^2 \hbar^2}{2ma^2} \\ &= \frac{n^2 \pi^2 h^2}{2ma^2 4\pi^2} \\ &= \frac{n^2 h^2}{8ma^2} \end{aligned} \quad (1.3.5)$$

The energy of the particle is said to be quantized and the integer n is called a quantum number. Note that the quantization arises naturally from the boundary conditions. The lowest allowed energy level is called the zero-point energy and is given by $E_1 = \frac{\hbar^2}{8ma^2}$.

The wave function corresponding to E_n is (using eq. (1.3.4))

$$\begin{aligned}\psi_n(x) &= B \sin(kx) \\ &= B \sin\left(\frac{n\pi x}{a}\right) \quad n = 1, 2, 3, \dots\end{aligned}\tag{1.3.6}$$

We will determine the constant B shortly.

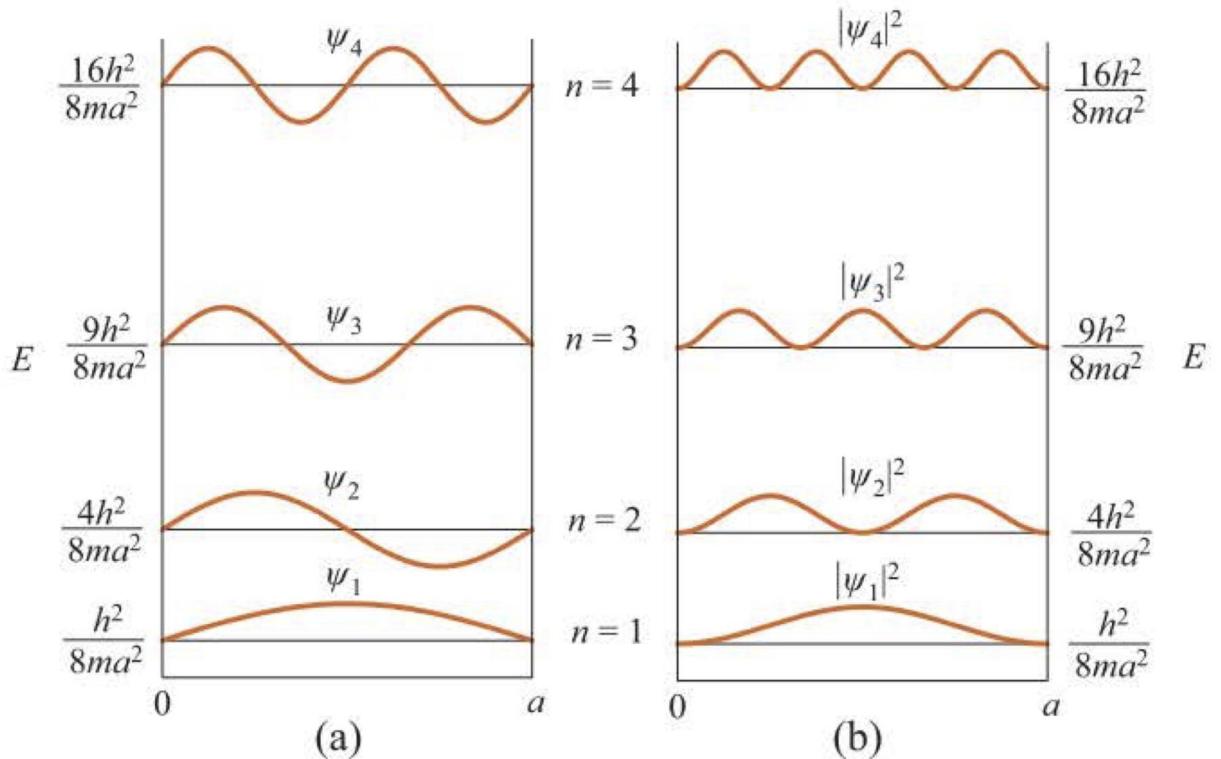


Figure 1.3.2: The energy levels, wave functions (a), and probability densities (b) for the particle in a box. Note that the energy increases with the number of nodes.

$$\int_0^a \psi_n^*(x) \psi_n(x) dx = 1 \tag{1.3.7}$$

$$|B|^2 \int_0^a \sin^2\left(\frac{n\pi x}{a}\right) dx = 1 \tag{1.3.8}$$

Let $\frac{n\pi x}{a} = z$, then $\frac{n\pi}{a} dx = dz$, $dx = \frac{a}{n\pi} dz$. The limits: when $x = 0, z = 0$ but when

$$x = a, z = n\pi$$

$$\begin{aligned}
\frac{a}{n\pi} \int_0^{n\pi} \sin^2 z \, dz &= \frac{a}{2n\pi} \int_0^{n\pi} 2 \sin^2(z) \, dz \\
&= \frac{a}{2n\pi} \int_0^{n\pi} (1 - \cos(2z)) \, dz \\
&= \frac{a}{2n\pi} \left[\int_0^{n\pi} dz + \int_0^{n\pi} \cos(2z) \, dz \right] \\
&= \frac{a}{2n\pi} \left[n\pi + \frac{\sin(2z)}{2} \Big|_0^{n\pi} \right] \\
&= \frac{a}{2n\pi} [n\pi + \sin(2n\pi) - \sin(0)] \\
&= \frac{a}{2n\pi} [n\pi + \cancel{\sin(2n\pi)}^0 - \sin(0)] \quad n = 1, 2, 3, \dots \\
&= \frac{a}{2}
\end{aligned} \tag{1.3.9}$$

From eqs. (1.3.8) and (1.3.9), we obtain

$$\begin{aligned}
|B|^2 \frac{a}{2} &= 1 \\
B &= \sqrt{\frac{2}{a}}
\end{aligned} \tag{1.3.10}$$

Inserting eq. (1.3.10) in eq.(1.3.6), we get

$$\psi_n(x) = \sqrt{\frac{2}{a}} \sin\left(\frac{n\pi x}{a}\right) \quad n = 1, 2, 3, \dots \tag{1.3.11}$$