



Part I Fundamentals

Electron Theory : Matter Waves

Chap. 1 Introduction

Chap. 2 The Wave-Particle Duality

Chap. 3 The Schödinger Equation

**Chap. 4 Solution of the Schödinger Equation for
Four Specific Problems**

Chap. 5 Energy Bands in Crystals

Chap. 6 Electrons in a Crystal

Electromagnetic Theory : Maxwell Equations

Chap. 4 Light Waves

(Electrons in Solids, 3rd Ed., R. H. Bube)





3. The Schrödinger Equation



3.1 The Time-Independent Schrödinger Equation

- Time-independent Schrödinger equation: *a vibration equation*

$$\nabla^2 \psi + \frac{2m}{\hbar^2} (E - V) \psi = 0$$

$$\nabla^2 \psi = \frac{\partial^2 \psi}{\partial x^2} + \frac{\partial^2 \psi}{\partial y^2} + \frac{\partial^2 \psi}{\partial z^2}$$

where, m = the (rest) mass of the electron,

E = the total energy of the system,

E_{kin} = kinetic energy,

V = the potential energy (or potential barrier)

$$E = E_{\text{kin}} + V$$

- Applicable to the calculation of the properties of atomic systems in *stationary* conditions



3. The Schrödinger Equation

3.2 The Time-Dependent Schrödinger Equation

Time-dependent Schrödinger equation: *a wave equation*

$$\nabla^2 \Psi - \frac{2mV}{\hbar^2} \Psi - \frac{2mi}{\hbar} \frac{\partial \Psi}{\partial t} = 0$$

Since $\Psi(x, y, z, t) = \psi(x, y, z) \cdot e^{i\omega t}$

$$\frac{\partial \Psi}{\partial t} = \psi i \omega e^{i\omega t} = \Psi i \omega \quad \longrightarrow \quad \omega = -\frac{i}{\Psi} \cdot \frac{\partial \Psi}{\partial t}$$

and $E = \nu h = \omega \hbar \quad \longrightarrow \quad E = -\frac{\hbar i}{\Psi} \cdot \frac{\partial \Psi}{\partial t}$

Then $\nabla^2 \psi + \frac{2m}{\hbar^2} (E - V) \psi = 0 \quad \longrightarrow \quad \nabla^2 \Psi - \frac{2mV}{\hbar^2} \Psi - \frac{2mi}{\hbar} \frac{\partial \Psi}{\partial t} = 0$

Applying differential operators to the wave function

$$E = -\hbar i \frac{\partial}{\partial t} \quad \mathbf{p} = -\hbar i \nabla$$

(**Hamiltonian operators**)

$$E_{total} = E_{kin} + E_{pot} = \frac{p^2}{2m} + V \quad \longrightarrow \quad -\hbar i \frac{\partial \Psi}{\partial t} = \frac{\hbar^2 i^2}{2m} \nabla^2 \Psi + V \Psi$$



3. The Schrödinger Equation



3.3 Special Properties of Vibrational Problems

- When boundary conditions are imposed, only certain vibrational forms are possible. ex) a vibrating string
- Vibration problems determined by boundary conditions:
boundary (or eigenvalue) problems
A peculiarity of these problems : not all frequency values are possible and therefore, not all values for the energy are allowed because of $E = \nu h$

*The allowed values : **eigenvalues***

*The function belonging to the eigenvalues as a solution of the vibration equation : **eigenfunctions***

The **normalized eigenfunction**:
$$\int \psi \psi^* d\tau = \int |\psi|^2 d\tau = 1$$



4. Solution of Schrödinger Equation

4.1 Free Electrons

Suppose electrons propagating freely (i.e., in a potential-free space) to the positive x -direction.

Then $V = 0$ and thus

$$\nabla^2 \psi + \frac{2m}{\hbar^2} (E - V) \psi = 0 \quad \longrightarrow \quad \frac{d^2 \psi}{dx^2} + \frac{2m}{\hbar^2} E \psi = 0$$

The solution for the above differential equation for an undamped vibration with spatial periodicity, (see Appendix 1)

$$\psi(x) = A e^{i\alpha x}$$

where $\alpha = \sqrt{\frac{2m}{\hbar^2} E} \quad \longrightarrow \quad \alpha = \sqrt{\frac{2m}{\hbar^2} E} = \frac{p}{\hbar} = \frac{2\pi}{\lambda} = k \quad |\mathbf{k}| = \frac{2\pi}{\lambda}$

Thus $\Psi(x) = A e^{i\alpha x} \cdot e^{i\omega t}$

$$E = \frac{\hbar^2}{2m} \alpha^2 \quad \longrightarrow \quad E = \frac{\hbar^2}{2m} k^2$$

“energy continuum”

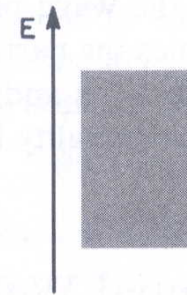


Figure 4.1. Energy continuum of a free electron (compare with Fig. 4.3).

4. Solution of Schrödinger Equation

4.2 Electron in a Potential Well (Bound Electron)

Consider an electron bound to its atomic nucleus.

Suppose the electron can move freely between two infinitely high potential barriers

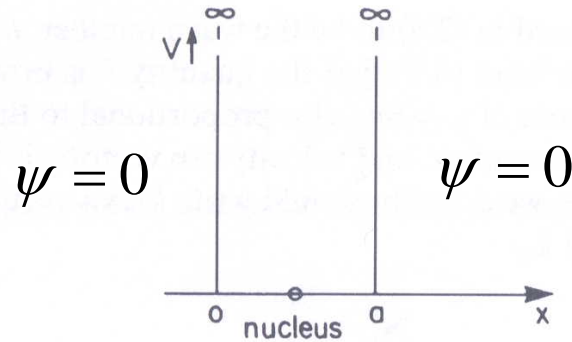


Figure 4.2. One-dimensional potential well. The walls consist of infinitely high potential barriers.

At first, treat 1-dim propagation along the x -axis inside the potential well

$$\frac{d^2\psi}{dx^2} + \frac{2m}{\hbar^2} E \psi = 0$$

The solution $\psi = Ae^{i\alpha x} + Be^{-i\alpha x}$ where $\alpha = \sqrt{\frac{2m}{\hbar^2} E}$

4. Solution of Schrödinger Equation

4.2 Electron in a Potential Well (Bound Electron)

Applying boundary conditions,

$$x = 0, \quad \psi = 0 \quad \longrightarrow \quad B = -A$$

$$x = a \quad \psi = 0 \quad \longrightarrow \quad 0 = Ae^{i\alpha a} \cdot Be^{-i\alpha a} = A(e^{i\alpha a} - e^{-i\alpha a})$$

With Euler equation, $\sin \rho = \frac{1}{2i}(e^{i\rho} - e^{-i\rho})$

$$A[e^{i\alpha a} - e^{-i\alpha a}] = 2Ai \cdot \sin \alpha a = 0$$

$$\alpha a = n\pi, \quad n = 0, 1, 2, 3, \dots \quad \text{“energy levels”}$$

Finally, $E_n = \frac{\hbar^2}{2m} \alpha^2 = \frac{\hbar^2 \pi^2}{2ma^2} n^2$

$$n = 1, 2, 3, \dots$$

“energy quantization”

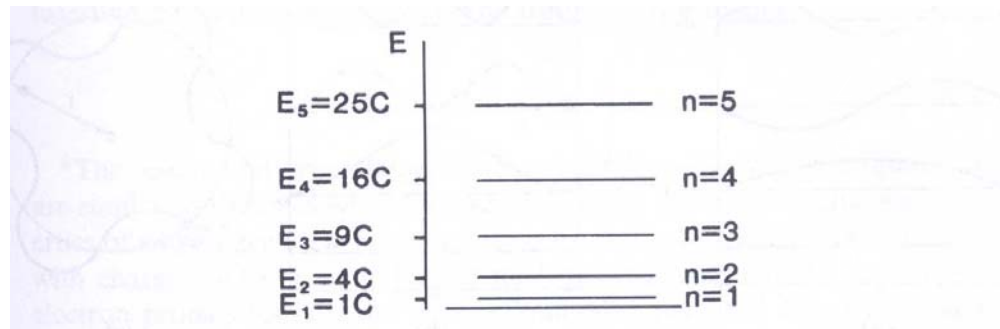


Figure 4.3. Allowed energy values of an electron that is bound to its atomic nucleus. E is the excitation energy in the present case. $C = \hbar^2 \pi^2 / 2ma^2$, see (4.18). (E_1 is the zero-point energy.)

4. Solution of Schrödinger Equation

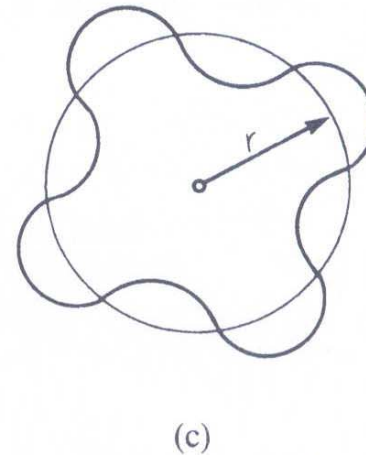
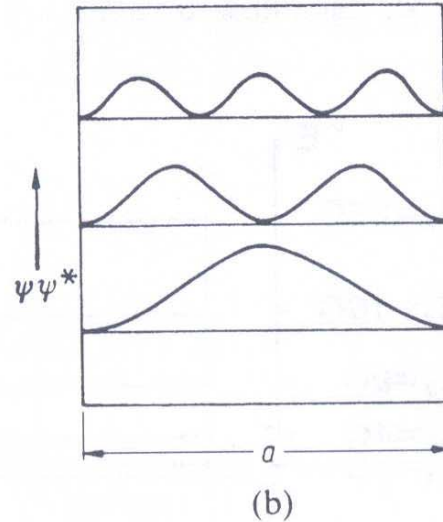
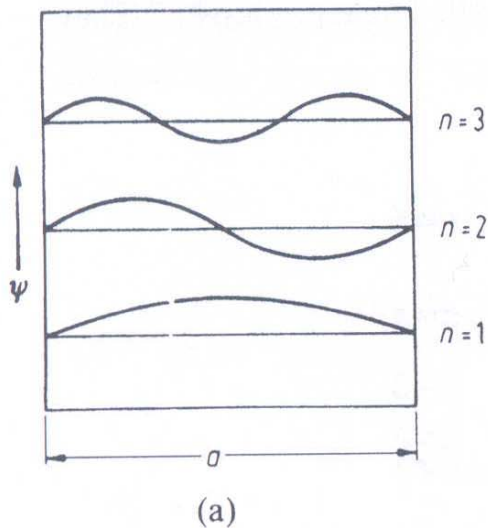
4.2 Electron in a Potential Well (Bound Electron)

Now discuss the wave function

$$\psi = 2Ai \cdot \sin \alpha x \quad \psi^* = 2Ai \cdot \sin \alpha x$$

$$\psi\psi^* = 4A^2 \sin^2 \alpha x \quad \int_0^a \psi\psi^* d\tau = 4A^2 \int_0^a \sin^2(\alpha x) dx = \frac{4A^2}{\alpha} \left[-\frac{1}{2} \sin \alpha x \cos \alpha x + \frac{\alpha x}{x} \right]_0^a = 1$$

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



$$2\pi r = n\lambda$$

$$r = \frac{\lambda}{2\pi} n$$

Figure 4.4. (a) ψ function and (b) probability function $\psi\psi^*$ for an electron in a potential well for different n -values. (c) Allowed electron orbit of an atom.

4. Solution of Schrödinger Equation

4.2 Electron in a Potential Well (Bound Electron)

For a hydrogen atom,
Coulombic potential

$$V = -\frac{e^2}{4\pi\epsilon_0 r}$$

$$E = \frac{me^4}{2(4\pi\epsilon_0\hbar)^2} \frac{1}{n^2} = -13.6 \cdot \frac{1}{n^2} \text{ (eV)}$$

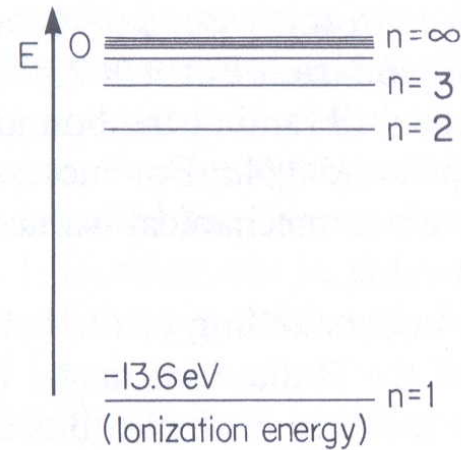


Figure 4.5. Energy levels of atomic hydrogen. E is the binding energy.

In 3-dim potential

$$E_n = \frac{\hbar^2 \pi^2}{2ma^2} (n_x^2 + n_y^2 + n_z^2)$$

The same energy but different quantum numbers: “degenerate” states

4. Solution of Schrödinger Equation

4.3 Finite Potential Barrier (Tunnel Effect)

Suppose electrons propagating in the positive x -direction encounter a potential barrier $V_0 (> \text{total energy of electron, } E)$

- **Region (I) $x < 0$**

$$\frac{d^2\psi}{dx^2} + \frac{2m}{\hbar^2} E\psi = 0$$

- **Region (II) $x > 0$**

$$\frac{d^2\psi}{dx^2} + \frac{2m}{\hbar^2} (E - V_0)\psi = 0$$

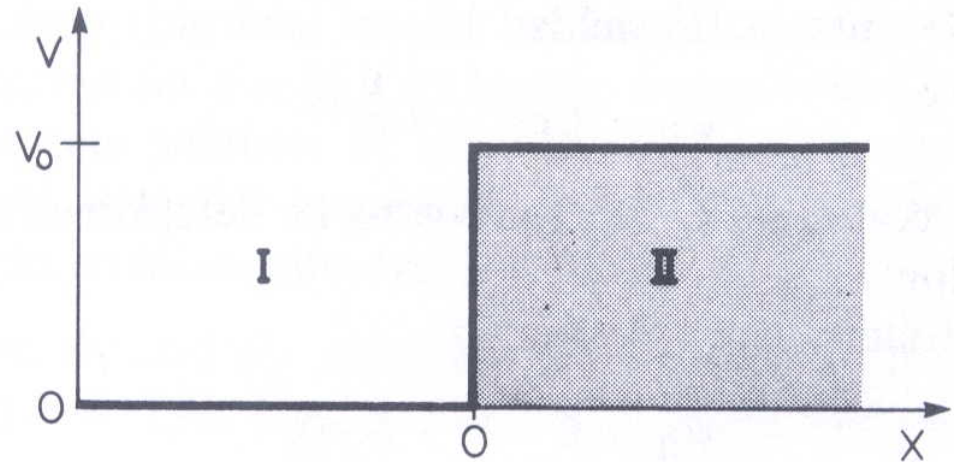
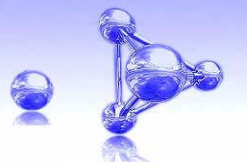


Figure 4.6. Finite potential barrier.

The solutions (see Appendix 1)

$$\psi_I = Ae^{i\alpha x} + Be^{-i\alpha x} \quad \alpha = \sqrt{\frac{2m}{\hbar^2} E}$$

$$\psi_{II} = Ce^{i\beta x} + De^{-i\beta x} \quad \beta = \sqrt{\frac{2m}{\hbar^2} (E - V_0)}$$



4. Solution of Schrödinger Equation



4.3 Finite Potential Barrier (Tunnel Effect)

Since $E - V_0$ is negative, $\beta = \sqrt{\frac{2m}{\hbar^2}(E - V_0)}$ becomes imaginary.

To prevent this, define a new parameter, $\gamma = i\beta$

Thus, $\gamma = \sqrt{\frac{2m}{\hbar^2}(V_0 - E)}$, and $\psi_{II} = Ce^{i\beta x} + De^{-i\beta x} \rightarrow \psi_{II} = Ce^{\gamma x} + De^{-\gamma x}$

Determination of C or D by B.C. For $x \rightarrow \infty$ $\psi_{II} = C \cdot \infty + D \cdot 0$

Since $\Psi \Psi^*$ can never be larger than 1, $\psi_{II} \rightarrow \infty$ is no solution, and thus $C \rightarrow 0$, which reveals Ψ -function decreases in Region II

$$\psi_{II} = De^{-\gamma x}$$

Using (A.27) + (4.39) in textbook, the damped wave becomes

$$\Psi = De^{-\gamma x} \cdot e^{i(\omega t - kx)}$$



4. Solution of Schrödinger Equation

4.3 Finite Potential Barrier (Tunnel Effect)

As shown by the dashed curve in Fig 4.7, a potential barrier is penetrated by electron wave : **Tunneling**

* For the complete solution,

(1) At $x = 0$ $\psi_I = \psi_{II}$: continuity of the function

$$Ae^{i\alpha x} + Be^{-i\alpha x} = De^{i\gamma x} \longrightarrow A + B = D$$

(2) At $x = 0$ $\frac{d\psi_I}{dx} \equiv \frac{d\psi_{II}}{dx}$: continuity of the slope of the function

$$Ai\alpha e^{i\alpha x} - Bi\alpha e^{-i\alpha x} = -\gamma D e^{-\gamma x}$$

With $x = 0$ $Ai\alpha - Bi\alpha = -\gamma D$

Consequently, $A = \frac{D}{2} \left(a + i \frac{\gamma}{\alpha} \right)$

$$B = \frac{D}{2} \left(1 - i \frac{\gamma}{\alpha} \right)$$

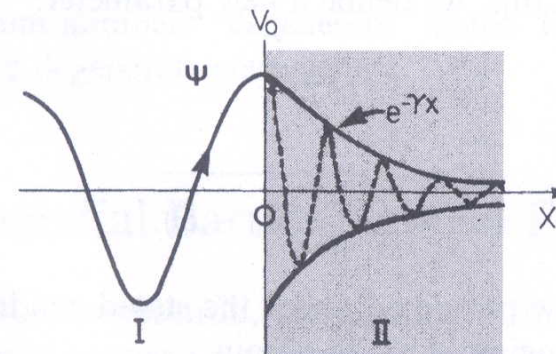


Figure 4.7. ψ -function (solid line) and electron wave (dashed line) meeting a finite potential barrier.

4. Solution of Schrödinger Equation

4.3 Finite Potential Barrier (Tunnel Effect)

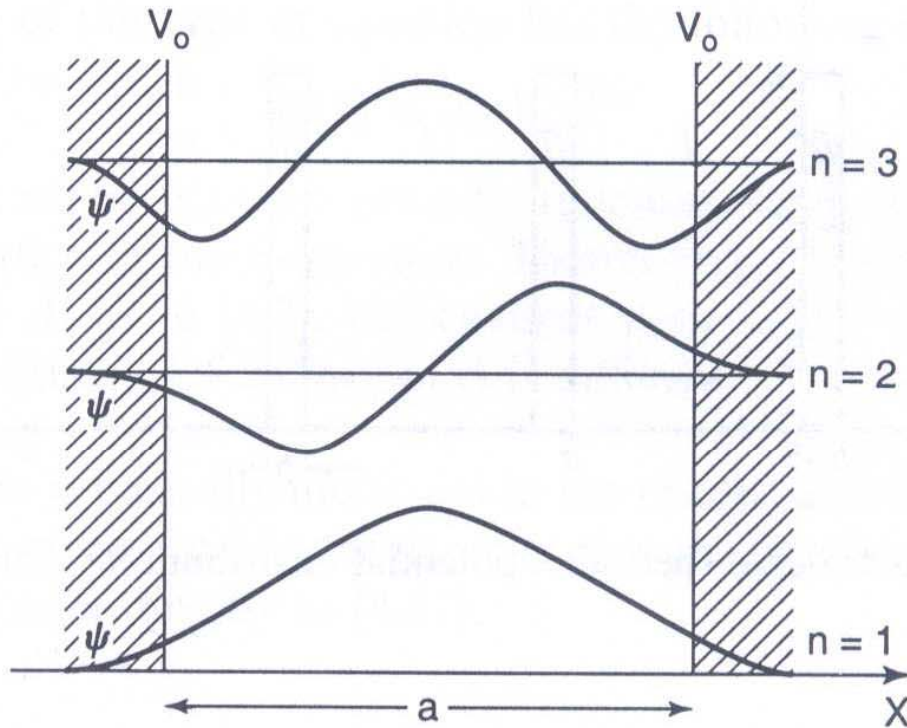


Figure 4.8. Square well with finite potential barriers. (The zero points on the vertical axis have been shifted for clarity.)

4. Solution of Schrödinger Equation

4.4 Electron in a Periodic Field of Crystal (the Solid State)

The behavior of an electron in a crystal → A motion through periodic repetition of potential well

well length : a

barrier height : V_0

barrier width : b

Region (I)

$$\frac{d^2\psi}{dx^2} + \frac{2m}{\hbar^2} E\psi = 0$$

Region (II)

$$\frac{d^2\psi}{dx^2} + \frac{2m}{\hbar^2} (E - V_0)\psi = 0$$

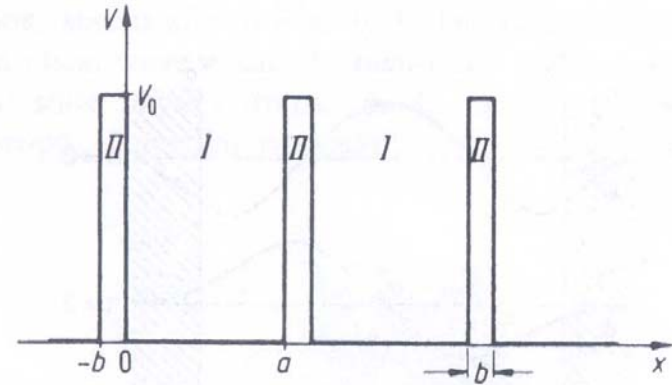


Figure 4.9. One-dimensional periodic potential distribution (simplified) (Kronig-Penney model).

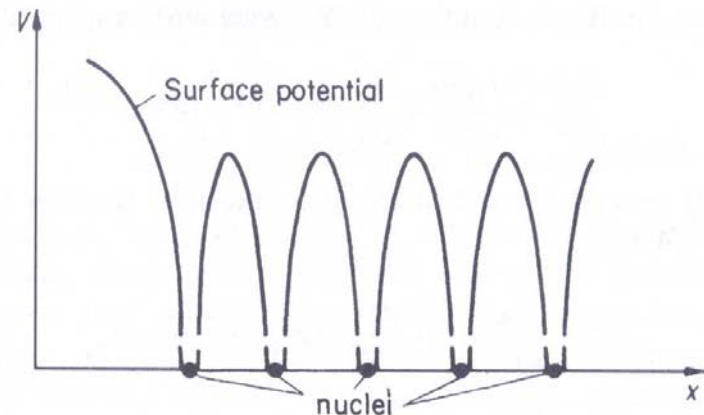


Figure 4.10. One-dimensional periodic potential distribution for a crystal (muffin tin potential).



4. Solution of Schrödinger Equation



4.4 Electron in a Periodic Field of Crystal (the Solid State)

(Continued) For abbreviation

$$\alpha^2 = \frac{2m}{\hbar^2} E \quad \gamma^2 = \frac{2m}{\hbar^2} (V_0 - E)$$

The solution of this type equation (not simple but complicate)

$$\psi(x) = u(x) \cdot e^{ikx} \quad (\text{Bloch function})$$

Where, $u(x)$ is a periodic function which possesses the periodicity of the lattice in the x -direction

The final solution of the Schrödinger equations;

$$P \frac{\sin \alpha a}{\alpha a} + \cos \alpha a = \cos ka \quad \text{where} \quad P = \frac{maV_0b}{\hbar^2}$$





4. Solution of Schrödinger Equation



4.4 Electron in a Periodic Field of Crystal (the Solid State)

Mathematical treatment for the solution : *Bloch function*

$$\psi(x) = u(x) \cdot e^{ikx}$$

Differentiating the Bloch function twice with respect to x

$$\frac{d^2\psi}{dx^2} = \left(\frac{d^2u}{dx^2} + \frac{du}{dx} 2ik - k^2u \right) e^{ikx}$$

Insert 4.49 into 4.44 and 4.45 and take into account the abbreviation

$$\frac{d^2u}{dx^2} + 2ik \frac{du}{dx} - (k^2 - \alpha^2)u = 0 \quad (\text{I}) \quad \frac{d^2u}{dx^2} + 2ik \frac{du}{dx} - (k^2 + \gamma^2)u = 0 \quad (\text{II})$$

The solutions of (I) and (II)

$$u = e^{-ikx} (Ae^{i\alpha x} + Be^{-i\alpha x}) \quad (\text{I}) \quad u = e^{-ikx} (Ce^{-\gamma x} + De^{\gamma x}) \quad (\text{II})$$





4. Solution of Schrödinger Equation



4.4 Electron in a Periodic Field of Crystal (the Solid State)

(Continued) From continuity of the function ψ and $\frac{d\psi}{dx}$

$$A + B = C + D$$

du/dx values for equations (I) & (II) are identical at $x = 0$

$$A(i\alpha - ik) + B(-i\alpha - ik) = C(\gamma - ik) + D(\gamma - ik)$$


Further, ψ and u is continuous at $x = a + b \rightarrow$ Eq. (I) at $x = 0$ must be equal to Eq. (II) at $x = a + b$, Similarly, Eq. (I) at $x = a$ is equal to Eq. (II) at $x = b$

$$Ae^{(i\alpha - ik)a} + Be^{(-i\alpha - ik)a} = Ce^{(ik + \gamma)b} + De^{(ik - \gamma)b}$$

Finally, du/dx is periodic in $a + b$

$$Ai(\alpha - k)e^{ia(\alpha - k)} - Bi(\alpha + k)e^{-ia(\alpha + k)} = -C(\gamma + ik)e^{(ik + \gamma)b} + D(\gamma - ik)e^{(ik - \gamma)b}$$

limiting conditions : using 4.57- 4.60 in text and eliminating the four constant A-D, and using some Euler eq.(see Appendix 2)

$$\frac{\gamma^2 - \alpha^2}{2\alpha\gamma} \sin(\gamma b) \cdot \sin(\alpha a) + \cos(\gamma b) \cdot \cos(\alpha a) = \cos k(a + b)$$




4. Solution of Schrödinger Equation



4.4 Electron in a Periodic Field of Crystal (the Solid State)

If V_0 is very large, then E in 4.47 is very small compared to V_0 so that

$$\gamma = \sqrt{\frac{2m}{\hbar^2}} \sqrt{V_0} \times b \quad \rightarrow \quad \gamma b = \sqrt{\frac{2m}{\hbar^2}} \sqrt{(V_0 b)b}$$

Since $V_0 b$ has to remain finite and $b \rightarrow 0$, γb becomes very small.

For a small γb , we obtain (see tables of the hyperbolic function)

$$\cosh(\gamma b) \approx 1 \quad \text{and} \quad \sinh(\gamma b) \approx \gamma b$$

Finally, neglect α^2 compared to γ^2 and, b compared to a so that 4.61 reads as follow

$$\frac{m}{\alpha \hbar^2} V_0 b \sin \alpha a + \cos \alpha a = \cos ka$$

Let $P = \frac{maV_0b}{\hbar^2}$, then $P \frac{\sin \alpha a}{\alpha a} + \cos \alpha a = \cos ka$



4. Solution of Schrödinger Equation

4.4 Electron in a Periodic Field of Crystal (the Solid State)

“Electron that moves in a periodically varying potential field can only occupy certain allowed energy zone”

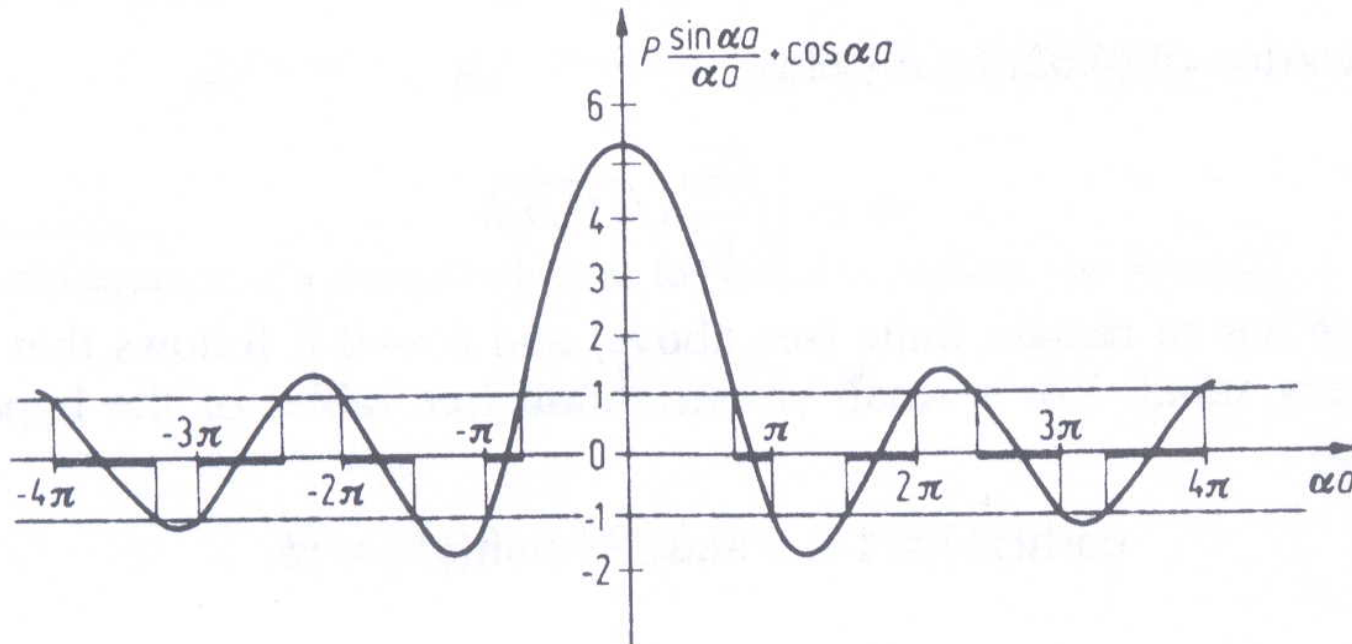


Figure 4.11. Function $P(\sin \alpha a / \alpha a) + \cos \alpha a$ versus αa . P was arbitrarily set to be $(3/2)\pi$.

4. Solution of Schrödinger Equation

4.4 Electron in a Periodic Field of Crystal (the Solid State)

The size of the allowed and forbidden energy bands varies with P .

For special cases

(a) If the potential barrier strength, V_0b is large, P is also large and the curve on Fig 4.11 steeper. The allowed bands are narrow.

(b) V_0b and P are small, the allowed band becomes wider.

(c) If V_0b goes 0, thus, $P \rightarrow 0$

From 4.67, $\cos \alpha a = \cos ka$

$$E = \frac{\hbar^2 k^2}{2m}$$

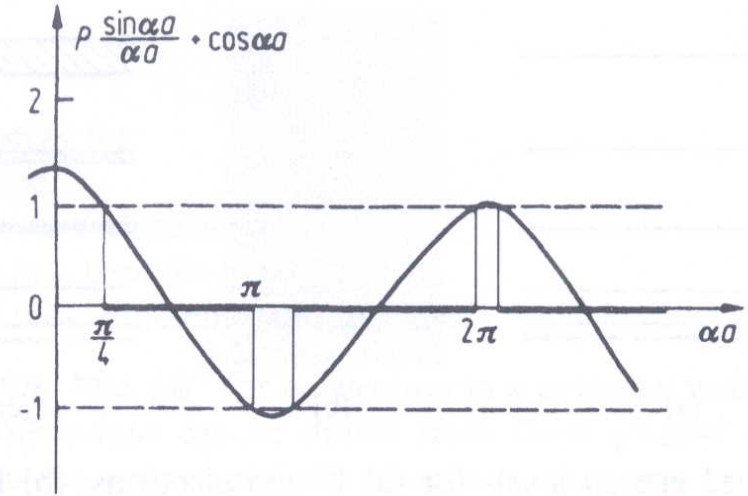


Figure 4.12. Function $P(\sin \alpha a / \alpha a) + \cos \alpha a$ with $P = \pi/10$.

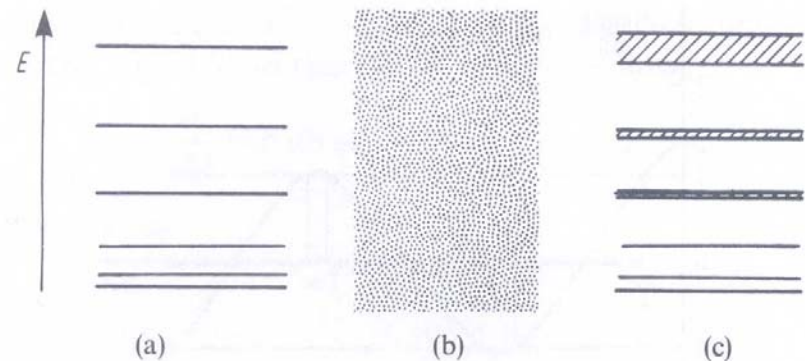


Figure 4.13. Allowed energy levels for (a) bound electrons, (b) free electrons, and (c) electrons in a solid.

4. Solution of Schrödinger Equation

4.4 Electron in a Periodic Field of Crystal (the Solid State)

(d) If the $V_0 b$ is very large, $P \rightarrow \infty$

$$\frac{\sin \alpha a}{\alpha a} \rightarrow 0$$

$$\sin \alpha a \rightarrow 0 \quad \alpha a = n \pi$$

$$\alpha^2 = \frac{n^2 \pi^2}{a^2} \quad \text{for } n = 1, 2, 3, \dots$$

Combining 4.46 and 4.69

$$E = \frac{\pi^2 \hbar^2}{2m a^2} \cdot n^2$$

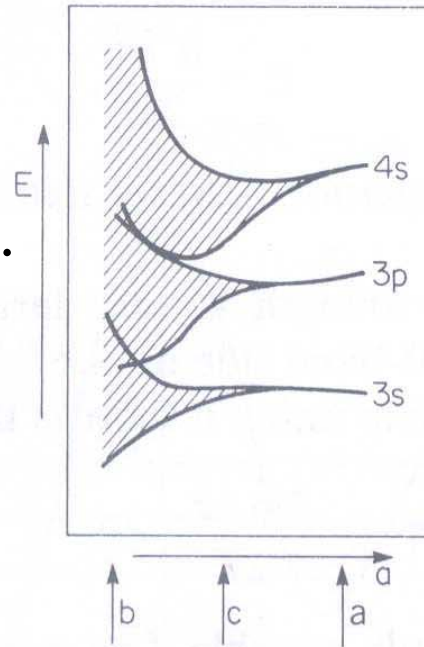


Figure 4.14. Widening of the sharp energy levels into bands and finally into a quasi-continuous energy region with decreasing interatomic distance, a , for a metal (after calculations of Slater). The quantum numbers are explained in Appendix 3.