

① Prove that $\left(\nabla - \frac{h^2}{8\pi^2 m v^2} \right) \Psi = E \Psi$

→ The equation of a stationary wave is designated by.

$$\Psi = A \sin \frac{2\pi x}{\lambda}$$

Double Diff.

$$\begin{aligned} \frac{d\Psi}{dx} &= A \cos \frac{2\pi x}{\lambda} \cdot \frac{d}{dx} \left(\frac{2\pi x}{\lambda} \right) \\ &= A \cos \frac{2\pi x}{\lambda} \cdot \frac{2\pi}{\lambda} \end{aligned}$$

Now diff. again;

$$\frac{d^2\Psi}{dx^2} = -A \sin \frac{2\pi x}{\lambda} \cdot \frac{2\pi}{\lambda} \cdot \frac{2\pi}{\lambda}$$

$$\frac{d^2\Psi}{dx^2} = -\Psi \cdot \frac{4\pi^2}{\lambda^2}$$

$$\frac{d^2\Psi}{dx^2} + \frac{4\pi^2}{\lambda^2} \Psi(x) = 0 \quad \text{--- (1)}$$

Now,

$$\lambda = \frac{h}{mv}, \quad \lambda = \frac{h}{p} \Rightarrow p = \frac{h}{\lambda}$$

$$T.E = K.E + P.E$$

$$= \frac{1}{2}mv^2 + v$$

$$\Rightarrow \frac{1}{2}mv^2 = \frac{1}{2} \frac{m^2 v^2}{m}$$

$$= \frac{1}{2} \frac{(p)^2}{m}$$

$$k = \frac{p^2}{2m} = \frac{h^2}{2m\lambda^2}$$

$$\boxed{\lambda^2 = \frac{h^2}{2mk}}$$

Now, put λ^2 in eq- (1)

$$\frac{d^2\Psi}{dx^2} + \frac{4\pi^2 \times 2m (E - v)}{h^2} \Psi = 0$$

$$\frac{d^2\Psi}{dx^2} + \frac{8\pi^2 m}{h^2} (E - v) \Psi = 0$$

Now,
$$\left(\frac{-h^2}{8\pi^2 m^2} \frac{d^2}{dn^2} + V \right) \Psi = E \Psi$$

$$\Rightarrow \left(V - \frac{h^2}{8\pi^2 m^2} \frac{d^2}{dn^2} \right) \Psi = E \Psi.$$

Now,

$$\frac{d^2 \Psi}{dn^2} + \frac{d^2 \Psi}{dy^2} + \frac{d^2 \Psi}{dz^2} + \frac{8\pi^2 m^2}{h^2} (E - V) \Psi = 0$$

$$\frac{d^2 \Psi}{dn^2} + \frac{d^2 \Psi}{dy^2} + \frac{d^2 \Psi}{dz^2} = \nabla^2$$

$$\nabla^2 + \frac{8\pi^2 m^2}{h^2} (E - V) \Psi = 0 \rightarrow (2)$$

Now, multiply

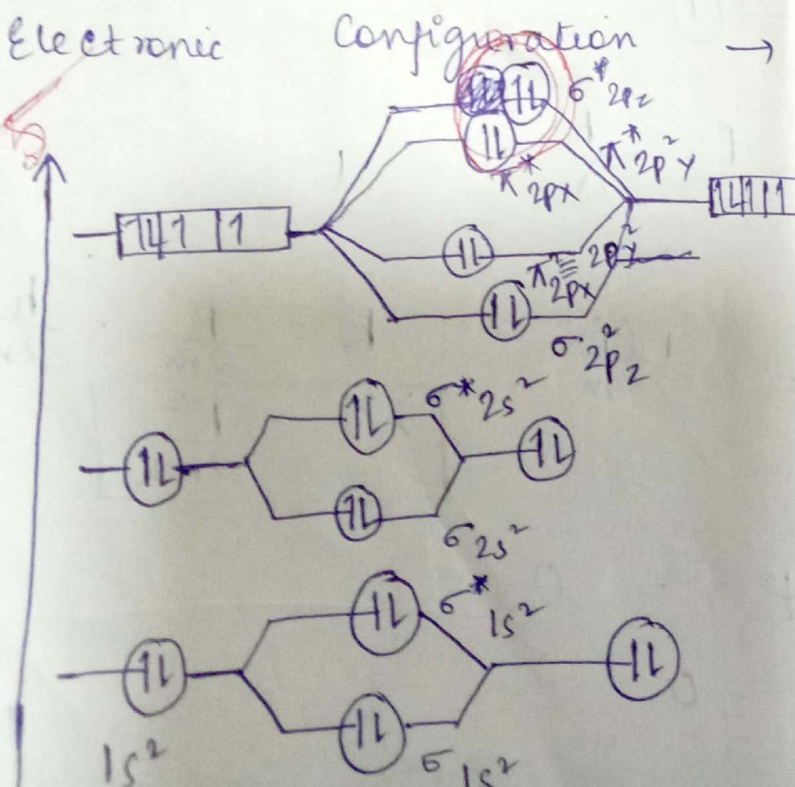
$$\frac{-h^2}{8\pi^2 m^2} \text{ in eq (2)}$$

$$\left(\frac{-h^2}{8\pi^2 m^2} \nabla^2 + V \right) \Psi = E \Psi$$

$$\hat{H} \Psi = E \Psi. \quad \text{Proved}$$

(2) Paramagnetic Behaviour of O₂

It has unpaired e⁻.



$1s^2 2s^2 2p^4$

$\uparrow \uparrow \quad \uparrow \uparrow \quad \uparrow \uparrow \uparrow \uparrow$

$\sigma 1s^2 < \sigma^* 1s^2 < \sigma 2s^2 < \sigma^* 2s^2$

$\sigma 2p_z^2 < \pi 2p_x^2 \equiv \pi 2p_y^2 < \pi^* 2p_x^2 \equiv \pi^* 2p_y^2 < \sigma^* 2p_z^2$

$$\Rightarrow \frac{N_b - N_A}{2} = (2)$$

$$\Rightarrow \frac{10 - 6}{2} = \frac{4}{2} = (2)$$

Because O_2 is having the M.O picture:

③ Vander Waals' forces :- ~~It~~ Vanderwaals forces are weak electrostatic forces that attract neutral forces in liquid or air vibrates constantly each other including ~~rotate~~ collide with each other including media particle such as water. This is called BROWNIAN MOTION.

④ NATURE :-

- The gas molecules are moving in a random, chaotic motion.
- The collision between the gas molecules and to the walls is perfectly elastic.
- The attraction between the gas molecules (repulsive / attraction) is negligible.

④ Equation of state for real gas -

$$\left(P + \frac{a}{V^2}\right)(V-b) = RT$$

P → Pressure of Real gas

V → volume of Real gas.

R → gas constant = $R = 8.314 \text{ J/mol}$

A gas obeys the ideal gas equation.

$PV = nRT$ under all the condition

of temperature and pressure called Ideal gas. There is no gas which ^{doesn't} obeys the Ideal gas

under all conditions of temp and pressure.
Significance of constant a and b →

a → Constant a provides the correction for the Intermolecular forces.

b → Constant b provides the volume occupied by a gas particle. It represents the correction for finite molecular size and its value is volume of one mole of the atom/molecule.

⑤ CHEMICAL POTENTIAL :- Chemical potential of a substance is the chemical energy per mole of the substance. In this Gibbs free energy is chemical energy, and substance can be a pure substance or a multicomponent of system.

DERIVATION :-

~~ΔG and ΔH~~ Temp and ΔG

~~$\Delta G = \Delta H - T\Delta S$~~

$$\Delta G = \Delta H - T\Delta S$$

$$W = nFE \text{ (cell)}$$

W = work

nFE → total energy

$$\Delta G = -W$$

$$\Delta G = -nFE$$

$$\Delta G^\circ = -nFE^\circ_{\text{cell}}$$

$$\Delta G^\circ = -RT \ln K$$

$$-nFE^\circ_{\text{cell}} = -RT \ln K$$

$$E^{\circ}_{\text{cell}} = \left(\frac{RT}{nF} \right) \ln K$$

Relation Between E_{mp} of cell and ΔH

$$\Delta H = \Delta A - U \Delta T$$

$$nFE = \text{total energy}$$

$$\Delta_r H = -nFE^{\circ}_{\text{cell}}$$

$$\Delta_r H = -RT \ln K$$

$$-nFE^{\circ}_{\text{cell}} = -RT \ln K$$

$$E^{\circ}_{\text{cell}} = \frac{RT}{nF} \ln K$$