

Lecture-1

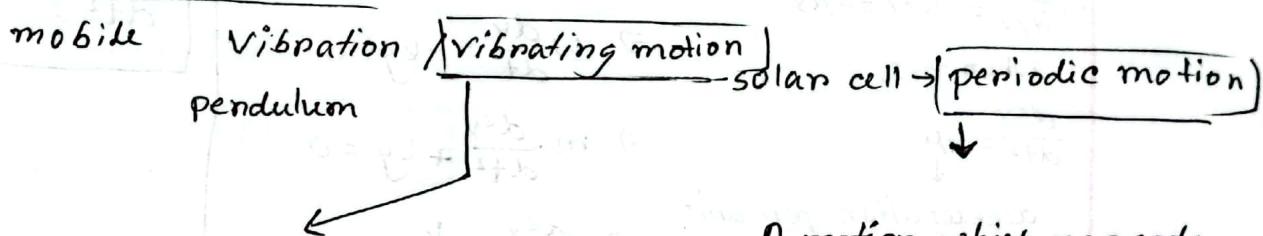
Kinetic theory of gases	- 35	Ornstein
Thermodynamics	- 35	$\frac{A}{4\pi} 2223 \text{ or } 37^2$ ans
Waves & Oscillations	- 35	105 marks
Lasers Physics	- 35	

Lecture-1

Date: 04.01.2023

Wednesday

Simple Harmonic motion:



When the particle, undergoing periodic motion, covers the same path back and forth about a mean position, it is said to be executing an oscillatory or vibrating motion.

A motion which repeats itself over and over again after a regular interval of time is referred to a periodic motion. The time required for its repetition is called time period.

Simple Harmonic motion:

Whenever a force acting on a particle, and hence the acceleration of the particle is proportional to its displacement from its equilibrium position but always directed in a direction opposite to the direction of the displacement and if the maximum displacement of the particle is same in either side of the main position, the particle is said to executing a simple Harmonic motion.

eqⁿ

Differential equation of SHM (Simple Harmonic Motion)

From the definition of SHM,

$$f \propto -y$$

$$\Rightarrow F = -ky \quad \text{---(1)}$$

[where k is
~~F~~ restoring force constant]

we know that, $F = ma \quad \text{---(2)}$

$$\therefore \frac{d^2y}{dt^2} = -\omega^2y = -ky$$

when, $y=1$,

$$\frac{d^2y}{dt^2} = -\mu$$

acceleration per unit
displacement of the
particle

$$ma = -ky$$

$$\Rightarrow m \frac{dv}{dt} = -ky$$

$$\Rightarrow m \cdot \frac{d^2y}{dt^2} + ky = 0$$

$$\Rightarrow \frac{d^2y}{dt^2} + \frac{k}{m}y = 0$$

$$\Rightarrow \frac{d^2y}{dt^2} + \omega^2y = 0$$

\downarrow
angular
velocity

Solution Of Differential eqⁿ of SHM:

$$\frac{d^2y}{dt^2} + \omega^2y = 0 \quad \text{---(1)}$$

$$\Rightarrow \frac{d^2y}{dt^2} = -\omega^2y$$

Multiplying both side by $\frac{2dy}{dt}$

$$\frac{2dy}{dt} \times \frac{d^2y}{dt^2} = -\omega^2y \times \frac{2dy}{dt}$$

$$\Rightarrow \frac{d}{dt} \left(\frac{dy}{dt} \right)^2 = -\omega^2 \frac{d}{dt} (y^2) \quad \text{---(2)} \quad \text{Integrating eqⁿ no. 2.}$$

$$\Rightarrow \int \frac{d}{dt} \left(\frac{dy}{dt} \right)^2 dt = \int -\omega^2 \frac{d}{dt} (y^2) dt$$

$$\Rightarrow \left(\frac{dy}{dt} \right)^2 = -\omega^2 y^2 + c \quad \text{---(3)}$$

When

if $y=a$, $\frac{dy}{dt}=0=v$
maximum displacement

From (3), we get, putting

$$c = \omega^2 a^2$$

putting this in eqn (ii),

$$\left(\frac{dy}{dt}\right)^2 = -\omega^2 y^2 + \omega^2 a^2$$

$$\Rightarrow \left(\frac{dy}{dt}\right)^2 = \omega^2 (a^2 - y^2)$$

$$\Rightarrow \frac{dy}{dt} = \pm \omega \sqrt{a^2 - y^2}$$

$\Rightarrow \frac{dy}{dt} = \omega \sqrt{a^2 - y^2}$ neglecting negative term

$$\Rightarrow \frac{dy}{dt} = \omega \sqrt{a^2 - y^2}$$

$$\Rightarrow \int \frac{dy}{dt} \left(\frac{y}{\sqrt{a^2 - y^2}} \right) dt = \int \omega dt$$

because velocity cannot be negative, and

$$\Rightarrow \int \frac{1}{\sqrt{a^2 - y^2}} dy = \omega dt$$

$$\Rightarrow \sin^{-1} \frac{y}{a} = \omega t + \varphi$$

$$\Rightarrow \frac{y}{a} = \sin(\omega t + \varphi)$$

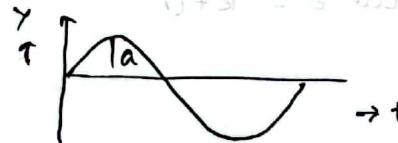
$$\Rightarrow y = a \sin(\omega t + \varphi)$$

Independent variable (time)

Graphical presentation of displacement:

$$y = a \sin(\omega t + \varphi)$$

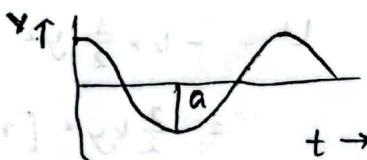
, when $\varphi = 0$:



Velocity:

$$v = \omega a \cos(\omega t + \varphi)$$

when $\varphi = 0$



$$v = \omega a \cos(\omega t + \varphi)$$

$$= \omega a \sin \sqrt{1 - \sin^2(\omega t + \varphi)}$$

$$= \omega a \sqrt{1 - \frac{y^2}{a^2}}$$

$$v = \omega \sqrt{a^2 - y^2}$$

$$a_v = \frac{dv}{dt}$$

When, $y=0$, $v_{\max} = \omega a \rightarrow \text{max}$

$$= \frac{d}{dt} \omega a \cos(\omega t + \delta)$$

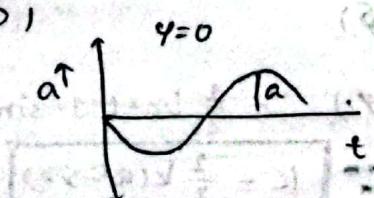
When, $y=a$, $v_{\min} = 0 \rightarrow \text{min}$

$$= -\omega^2 a \sin(\omega t + \delta)$$

When, $y=0$, $a_{v\min} = 0$

$$\Rightarrow a_v = -\omega^2 y$$

$y=a$, $a_{v\max} = \omega^2 a$



Time period of body executing SHM:

$$y = a \sin(\omega t + \varphi)$$

If the time is increased by $\frac{2\pi}{\omega}$

$$\begin{aligned} y &= a \sin\{\omega(t + \frac{2\pi}{\omega}) + \varphi\} \\ &= a \sin(\omega t + 2\pi + \varphi) \\ &= a \sin(2\pi + \omega t + \varphi) = a \sin(\omega t + \delta\varphi) \end{aligned}$$

Time period,

$$T = \frac{2\pi}{\omega} = \frac{2\pi}{\sqrt{\frac{k}{m}}} = 2\pi \sqrt{\frac{m}{k}}$$

$$\text{Frequency, } f = \frac{1}{T} = \frac{1}{2\pi} \sqrt{\frac{k}{m}}$$

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Tuesday

Energy of a particle executing SHM:

* Kinetic energy - $\frac{1}{2}mv^2$ (neglecting mass) $-ky^2 = F$

* Potential energy - U

$$\text{Total Energy } E = k + U$$

$$U = \frac{1}{2}ky^2$$

$$y = a \sin(\omega t + \varphi)$$

$$U = \frac{1}{2}ka^2 \sin^2(\omega t + \varphi) \quad (v)$$

$$K = \frac{1}{2}mv^2 =$$

$$U = \frac{dy}{dt}$$

$$= \frac{1}{2}m\left(\frac{dy}{dt}\right)^2$$

$$= \frac{d}{dt} a \sin(\omega t + \varphi)$$

$$= \frac{1}{2}m\omega^2 \cos^2(\omega t + \varphi) \approx \omega a \cos^2(\omega t + \varphi)$$

$$\approx \frac{1}{2}m\omega^2 a^2 \cos^2(\omega t + \varphi)$$

$$K = \frac{1}{2}ka^2 \cos^2(\omega t + \varphi) = \frac{1}{2}ka^2(1 - \sin^2(\omega t + \varphi))$$

$$\therefore K = \frac{1}{2}k(a^2 - y^2)$$

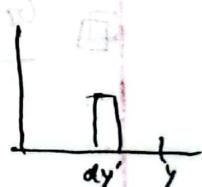
$$U = \int_0^y F dy'$$

$$= \int_0^y -ky^2 dy'$$

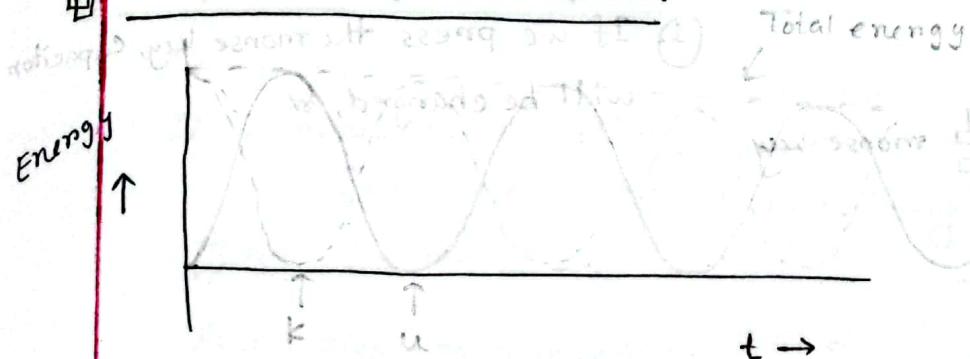
$$= -k \times \left[\frac{y^3}{3} \right]_0^y$$

$$U = -k \times \frac{1}{3}y^3$$

$$= \frac{1}{2}ky^2 \quad [\text{neglecting negative sign}]$$

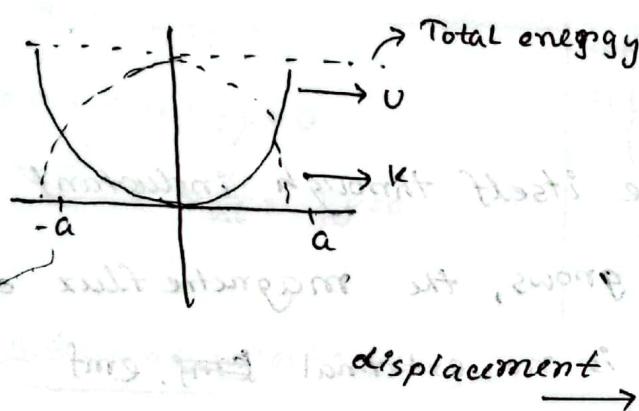


Graphical representation:



$$\text{Total energy, } E = U + K$$

$$\begin{aligned} &= \frac{1}{2}ka^2\sin^2(\omega t + \varphi) + \\ &\quad \frac{1}{2}ka^2\cos^2(\omega t + \varphi) \\ &= \frac{1}{2}ka^2 \end{aligned}$$



Average Potential energy,

$$U = \frac{1}{2}ka^2\sin^2(\omega t + \varphi)$$

$$U_{\text{avg}} = \frac{1}{T} \int_0^T \frac{1}{2}ka^2\sin^2(\omega t + \delta) \quad [\text{with respect to } T]$$

$$= \frac{ka^2}{4T} \int_0^T 2\sin^2(\omega t + \varphi)$$

$$= \frac{ka^2}{4T} \int_0^T \{1 - \cos(2\omega t + 2\varphi)\} \rightarrow \int_0^T / \int_0^T / \int_0^T 2\sin^2(\omega t + \varphi)$$

$$= \frac{Ka^2 T}{4T}$$

meaning

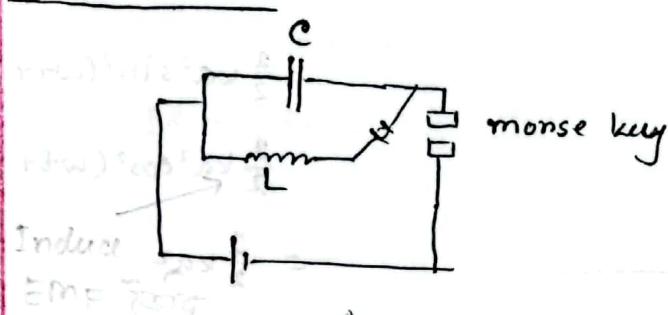
average zero + $\frac{1}{2}$ mean \Rightarrow displacement

$$\begin{aligned} &= \frac{ka^2}{2} \\ &\text{noticing initial position } \Rightarrow \frac{1}{2}(\text{total energy}) \end{aligned}$$

$$K_{\text{avg}} = \frac{1}{T} \int_0^T \frac{1}{2}ka^2\cos^2(\omega t + \varphi)$$

• SHM \rightarrow 2nd order \rightarrow sinusoidal wave

L.C. circuit :



If we add resistance system is not loss (damped)

- Capacitor discharge itself through inductant coil and current in the latter grows, the magnetic flux due to the current increase. There is no external emf.

$$\frac{Q}{C} = -L \frac{di}{dt}$$

$$\Rightarrow L \frac{di}{dt} + \frac{Q}{C} = 0$$

$$I = \frac{dQ}{dt}$$

$$\Rightarrow \frac{d}{dt} \left(\frac{dQ}{dt} \right) + \frac{Q}{LC} = 0$$

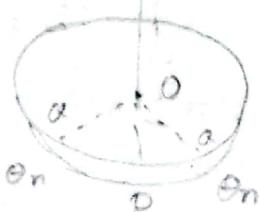
$$\frac{d^2Q}{dt^2} + \frac{Q}{LC} = 0$$

$$\omega^2 = \frac{1}{LC}$$

$$\Rightarrow \frac{d^2Q}{dt^2} + \omega^2 Q = 0$$

$$Q = Q_0 \sin(\omega t + \varphi)$$

Force F (tension)



(ii) Torsional Pendulum:

$$\bullet T = -k\theta, \tau = I\alpha$$

$$\Rightarrow -k\theta = I\alpha$$

$$\Rightarrow y = \frac{1}{k}\theta$$

$$\Rightarrow I\alpha = k\theta = 0$$

$$\Rightarrow I \times \frac{d^2\theta}{dt^2} + k\theta = 0$$

$$\Rightarrow \frac{d^2\theta}{dt^2} + \frac{k}{I}\theta = 0$$

- Prob - Compute the kinetic & potential energies of a body when it has moved in halfway from its initial position towards the center of motion.

$$\frac{d^2\theta}{dt^2} + \omega^2\theta = 0$$

2nd Order

$$\omega^2 = \frac{k}{I}$$

Damped Harmonic Oscillations:

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Wednesday

$$F = -ky \quad (i)$$

If the medium is resistive,

$$F_r = -bv \quad (ii) \quad b \text{ is resisting force constant/damping force}$$

$$\text{Again, } F = ma \quad (iii)$$

From these three equations,

$$ma = -ky - bv$$

$$\Rightarrow ma + bv + ky = 0$$

$$\Rightarrow m \frac{dy}{dt^2} + b \frac{dy}{dt} + ky = 0$$

$$\left. \begin{aligned} & \frac{d^2y}{dt^2} + \frac{b}{m} \frac{dy}{dt} + \frac{k}{m} y = 0 \\ & \Rightarrow \frac{d^2y}{dt^2} + 2\gamma \frac{dy}{dt} + \omega^2 y = 0 \end{aligned} \right\} \quad (iv)$$

when, $2\gamma = \frac{b}{m}$

differential equation of damped harmonic oscillation

Solution of eqn (iv) :

Let the solution is, $y = Ae^{kt}$

$$\Rightarrow \frac{dy}{dt} = Ake^{kt}$$

$$\Rightarrow \frac{d^2y}{dt^2} = Ak^2e^{kt}$$

* From eqn (iv),

$$Ak^2e^{kt} + 2\gamma \times Ake^{kt} + \omega^2 Ae^{kt} = 0$$

$$\Rightarrow Ae^{kt}(k^2 + 2\gamma k + \omega^2) = 0$$

$$\Rightarrow k^2 + 2\gamma k + \omega^2 = 0 \quad (i)$$

$$\Rightarrow k = \frac{-2\gamma \pm \sqrt{4\gamma^2 - 4\omega^2}}{2}$$

$$\Rightarrow k = -\gamma \pm \sqrt{\gamma^2 - \omega^2}$$

amplitude increasing

$$from \ eqn(ii),$$

$$y = Ae^{(-\gamma + \sqrt{\gamma^2 - \omega^2})t}$$

$$y = Ae^{(-\gamma - \sqrt{\gamma^2 - \omega^2})t}$$

* In general case,

$$Y = A_1 e^{(-\gamma + \sqrt{\gamma^2 - \omega^2})t} + A_2 e^{(-\gamma - \sqrt{\gamma^2 - \omega^2})t} \quad (ii)$$

$$\frac{dy}{dt} = A_1 (-\gamma + \sqrt{\gamma^2 - \omega^2}) e^{(-\gamma + \sqrt{\gamma^2 - \omega^2})t} + A_2 (-\gamma - \sqrt{\gamma^2 - \omega^2}) e^{(-\gamma - \sqrt{\gamma^2 - \omega^2})t}$$

Let,

When $t=0$, $y=a_0 \rightarrow$ maximum displacement

From eqⁿ(ii),

$$y = A_1 + A_2 = a_0$$

when, $y = a_0 \rightarrow$ maximum displacement, $\frac{dy}{dt} = 0, t=0$.

$$(-\lambda + \sqrt{\lambda^2 - \omega^2})A_1 + (-\lambda - \sqrt{\lambda^2 - \omega^2})A_2 = 0$$

$$\Rightarrow -\lambda(A_1 + A_2) + \sqrt{\lambda^2 - \omega^2}(A_1 - A_2) = 0$$

$$\Rightarrow -\lambda a_0 + \sqrt{\lambda^2 - \omega^2}(A_1 - A_2) = 0$$

$$\Rightarrow A_1 - A_2 = \frac{\lambda a_0}{\sqrt{\lambda^2 - \omega^2}} \quad \text{(v)}$$

From eqⁿ(iv) and (v),

$$A_1 + A_2 = a_0$$

$$A_1 = \frac{1}{2} \left(a_0 + \frac{a_0 \lambda}{\sqrt{\lambda^2 - \omega^2}} \right) \quad \text{(vi)}$$

$$\Rightarrow A_1 - A_2 = \frac{a_0}{\sqrt{\lambda^2 - \omega^2}}$$

$$\Rightarrow \text{Again, } A_2 = a_0 - A_1 = a_0 - \frac{1}{2} a_0 - \frac{1}{2} \times \frac{a_0 \lambda}{\sqrt{\lambda^2 - \omega^2}} \\ = \frac{1}{2} \left(a_0 - \frac{a_0 \lambda}{\sqrt{\lambda^2 - \omega^2}} \right) \quad \text{(vii)}$$

The general solution is,

$$y = \frac{1}{2} a_0 \left(1 + \frac{\lambda}{\sqrt{\lambda^2 - \omega^2}} \right) e^{(-\lambda + \sqrt{\lambda^2 - \omega^2})t} + \frac{1}{2} a_0 \left(1 - \frac{\lambda}{\sqrt{\lambda^2 - \omega^2}} \right) e^{(-\lambda - \sqrt{\lambda^2 - \omega^2})t}$$

$$+ (\dots) \quad \text{or} \quad y = (a_0 + R_1 e^{\lambda t} + R_2 e^{-\lambda t}) \cos(\omega t) + R_3 e^{\lambda t} \sin(\omega t) \quad \text{(viii)}$$

④ When ($\lambda^2 > \omega^2$),

equation \rightarrow over damped Harmonic Oscillation

$e^{-\lambda t} \rightarrow$ negative exponential

Three important cases:

- i) $\lambda^2 > \omega^2$: Damping is large $\sqrt{\lambda^2 - \omega^2}$ is clearly a real quantity with positive value, less than λ . Then each of the

$e^{-\gamma t} \rightarrow$ damping factor

two terms on the right hand side of eqn (viii) and has an exponential term with a negative power and hence each decreases exponentially with time. In this case particle doesn't vibrate. There is no oscillations and the motion is therefore called overdamped.

(i) When $\gamma^2 = \omega^2$, In this case $\sqrt{\gamma^2 - \omega^2}$ is obviously equal to zero. In this case motion break down.

consider, $\sqrt{\gamma^2 - \omega^2} = h$; very small value

$$\begin{aligned} y &= A_1 e^{(-\gamma + ht)t} + A_2 e^{(-\gamma - ht)t} \\ \Rightarrow y &= e^{-\gamma t} \left[A_1 e^{ht} + A_2 e^{-ht} \right] \quad \text{A}_1, A_2 \xrightarrow{n} \begin{array}{l} \text{overdamped} \\ \text{critical} \\ \text{damping} \\ \text{under} \\ \text{damped} \end{array} \\ &= e^{-\gamma t} \left[(A_1 + A_2) - \left[A_1 \left[1 + ht + \frac{h^2 t^2}{2!} + \dots \right] + A_2 \left[1 - ht + \frac{h^2 t^2}{2!} - \dots \right] \right] \right] \\ &= e^{-\gamma t} \left[-(A_1 + A_2) + ht(A_1 - A_2) \right] \quad (\text{neglecting higher sign}) \end{aligned}$$

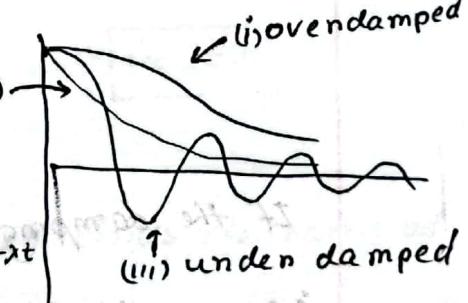
$$= e^{-\gamma t} \left[a_0 + h(A_1 - A_2)t \right] \quad \text{where,}$$

$$= e^{-\gamma t} \left[a_0 + \gamma a_0 t \right] \quad (\text{IX})$$

$$y = a_0 e^{-\gamma t} [1 + \gamma t]$$

$$\begin{aligned} A_1 + A_2 &= a_0 \\ A_1 - A_2 &= \frac{\gamma a_0}{\sqrt{\gamma^2 - \omega^2}} = \frac{\gamma a_0}{h} \end{aligned}$$

critical damping \rightarrow (ii)



(ii) The second form ($a_0 \gamma a_0 e^{-\gamma t}$) in eqn (IX) decays less rapidly than the first term $a_0 e^{-\gamma t}$ and the displacement of the oscillation first increases but as time increases exponential factor $e^{-\gamma t}$ becomes more important and the displacement decreases rapidly reaching the value of zero.

at the finite value of t . This is called critical damping.

(ii) $\lambda^2 < \omega^2$, The quantity is $\sqrt{\omega^2 - \lambda^2}$ is clearly imaginary, say equal to $i\gamma$, γ is imaginary.

$$y = A_1 e^{(-\lambda + i\gamma)t} + A_2 e^{(-\lambda - i\gamma)t}$$

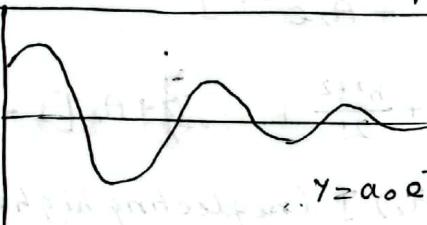
$$= e^{-\lambda t} [A_1 e^{i\gamma t} + A_2 e^{-i\gamma t}]$$

$$= e^{-\lambda t} [A_1 (\cos \gamma t + i \sin \gamma t) + A_2 (\cos \gamma t - i \sin \gamma t)]$$

$$= e^{-\lambda t} [(A_1 + A_2) \cos \gamma t + i(A_1 - A_2) \sin \gamma t]$$

$$= e^{-\lambda t} [A \cos \gamma t + B \sin \gamma t]$$

Power Dissipation of damped Harmonic Oscillations



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Tuesday

$$y = a_0 e^{-\lambda t} \sin(\gamma t + \phi)$$

$$\frac{dy}{dt} = a_0 - \lambda a_0 e^{-\lambda t} \cos(\gamma t + \phi) + a_0 e^{-\lambda t} \gamma \cos(\gamma t + \omega)$$

Kinetic energy, $E_k = \frac{1}{2} m v^2 = \frac{1}{2} m \left(\frac{dy}{dt} \right)^2$

$$= \frac{1}{2} m \times a_0^2 e^{-2\lambda t} [-\lambda \sin(\gamma t + \phi) + \gamma \cos(\gamma t + \phi)]^2$$

$$= \frac{1}{2} m a_0^2 e^{-2\lambda t} [\lambda^2 \sin^2(\gamma t + \phi) + \gamma^2 \cos^2(\gamma t + \phi) - 2\lambda \gamma \sin(\gamma t + \phi) \cos(\gamma t + \phi)]$$

If the damping is less than the term $a_0^2 e^{-2\lambda t}$ doesn't apparently, i.e., amplitude doesn't change apparently.

Average Kinetic energy, $E_{kav} = \frac{1}{2} m a_0^2 e^{-2\lambda t} \left[\frac{1}{2} \lambda^2 + \frac{1}{2} \gamma^2 t_0 \right]$

$$= \frac{1}{2} m a_0^2 e^{-2\lambda t} \left[\frac{1}{2} \lambda^2 + \frac{1}{2} \gamma^2 t_0 \right] \quad g = \sqrt{\omega^2 - \lambda^2}$$

For average value, $\sin^2(gt+\varphi)$ and $\cos^2(gt+\varphi)$ are $\frac{1}{2}$

Average value of $2\sin(gt+\varphi)\cos(gt+\varphi) = \sin 2(gt+\varphi)$ is zero

$$= \frac{1}{4} m a_0^2 g^2 e^{-2\alpha t}$$

Potential energy, $E_p = \int_0^y mg^2 y dy = mg^2 \left[\frac{y^2}{2} \right]_0^y$

$$= \frac{1}{2} mg^2 y^2$$

$$\therefore E_p = \frac{1}{2} mg^2 a_0^2 e^{-2\alpha t} \sin^2(gt+\varphi)$$

average value of $\sin^2(gt+\varphi)$ is $\frac{1}{2}$

So, average Potential energy, $E_{pav} = \frac{1}{2} mg^2 a_0^2 e^{-2\alpha t}$

Total Energy, $E = E_{pav} + E_{kav}$

$$= \frac{1}{2} m a_0^2 g^2 e^{-2\alpha t}$$

$$= \boxed{\text{Undamped}} e^{-2\alpha t}$$

Average power dissipation, = rate of loss energy

$$\Rightarrow - \frac{dE}{dt} = - \frac{d}{dt} \frac{1}{2} m a_0^2 g^2 e^{-2\alpha t}$$

$$= 2\alpha \frac{1}{2} m a_0^2 g^2 e^{-2\alpha t}$$

$$= 2E\alpha = 2\alpha E$$

$$\therefore P = 2E\alpha$$

Quality factor: $(\frac{g}{2\alpha})$

The quality factor, Q is defined as 2π times the ratio of between the energy stored and the energy lost per period,

$$Q = 2\pi \frac{\text{energy stored}}{\text{Energy lost per period}}$$

$$= 2\pi \frac{E}{P}$$

$$g = \frac{2\pi}{T}$$

$g \gg g$

$$= \frac{Eg}{P}$$

$$= \frac{Eg}{2\pi E} = \boxed{\frac{g}{2\pi}}$$

$$\boxed{Q} Q = \frac{\sqrt{\omega^2 - \alpha^2}}{2\pi} = \frac{\omega}{2\pi} = \frac{\sqrt{\frac{k}{m}}}{\frac{b}{m}} = \frac{\sqrt{km}}{b}$$

$$\therefore Q = \frac{\sqrt{km}}{b}$$

* * part $\frac{1}{2} = \sqrt{3}$



$$\boxed{Frequency} n = \frac{g}{2\pi}$$

$$= \frac{\sqrt{\omega^2 - \alpha^2}}{2\pi}$$

$$= \frac{1}{2\pi} \sqrt{\omega^2 - \alpha^2}$$

$$= \frac{1}{2\pi} \sqrt{\left(\frac{1}{Lc}\right)^2 + \left(\frac{R}{2L}\right)^2}$$

$$\therefore n = \frac{1}{2\pi} \sqrt{\frac{1}{Lc} - \frac{R^2}{4L^2}}$$

$$Q = \frac{g}{2\pi} = \frac{\sqrt{\frac{1}{Lc} - \frac{R^2}{4L^2}}}{\frac{R}{L}}$$

$$\Rightarrow L \frac{d^2Q}{dt^2} + R \frac{dQ}{dt} + \frac{1}{Lc} Q = 0$$

$$\Rightarrow \frac{d^2Q}{dt^2} + \frac{R}{L} \frac{dQ}{dt} + \frac{1}{Lc} Q = 0$$

$$\Rightarrow \frac{d^2Q}{dt^2} + 2\alpha \frac{dQ}{dt} + \omega^2 Q = 0$$

$$2\alpha = \frac{R}{L}, \quad \omega^2 = \frac{1}{Lc}$$

$$Q = Q_0 e^{-\alpha t} \sin(\omega t + \phi)$$

(K) instant value

Forced Vibration:

Let the applied sinusoidal force is $F = F_0 \sin \omega t$

$$\boxed{m \frac{d^2y}{dt^2}}$$

According to the definition of forced vibration

$$m \frac{d^2y}{dt^2} = -b \frac{dy}{dt} - ky + F_0 \sin \omega t$$

$$\Rightarrow m \frac{d^2y}{dt^2} + b \frac{dy}{dt} + ky = F_0 \sin \omega t$$

$$2\eta = \frac{b}{m}, \omega^2 = \frac{k}{m}, f_0 = \frac{\omega}{2\pi}$$

$$\Rightarrow \frac{d^2y}{dt^2} + \frac{b}{m} \cdot \frac{dy}{dt} + \frac{k}{m}y = F_0 \sin \omega t$$

$$\Rightarrow \frac{d^2y}{dt^2} + 2\eta \frac{dy}{dt} + \omega^2 y = F_0 \sin \omega t \quad \text{--- (i)}$$

Let the solution of equation (i)

$$y = A \sin(\omega t - \theta)$$

A is the amplitude and θ is the possible phase difference. frequency $= \frac{\omega}{2\pi}$

$$T_0 [3 \cos(\omega t) + (3 - \eta^2) \sin(\omega t)] \frac{1}{\eta^2} =$$

$$\textcircled{*} \quad \frac{d^2y}{dt^2} + 2\lambda \frac{dy}{dt} + \omega^2 y = f_0 \sin pt - \textcircled{1}$$

$$\Rightarrow y = A \sin(pt - \theta) - \textcircled{2}$$

$$\frac{dy}{dt} = A p \cos(pt - \theta)$$

$$\frac{d^2y}{dt^2} = -A p^2 \sin(pt - \theta)$$

Putting the values in eqⁿ(i),

$$-A p^2 \sin(pt - \theta) + 2\lambda A p \cos(pt - \theta) + \omega^2 A \sin(pt - \theta)$$

$$= f_0 \sin \underbrace{\{pt - \theta + \theta\}}_{= f_0 \sin(pt - \theta)}$$

$$= f_0 \sin(pt - \theta) \cos \theta + f_0 \cos(pt - \theta) \sin \theta - \textcircled{3}$$

- If this solution is to be hold good for all values of it, the respective low-efficiency of $\sin(pt - \theta)$ and $\cos(pt - \theta)$ on either side of eqⁿ(iii) must be equal.

$$A(\omega^2 - p^2) = f_0 \cos \theta - \textcircled{4}$$

$$\textcircled{5} \Rightarrow 2\lambda A p = f_0 \sin \theta - \textcircled{5}$$

Squaring and adding eqⁿ(iv) and (v) →

$$A^2 (\omega^2 - p^2)^2 + 4\lambda^2 A^2 p^2 = f_0^2$$

$$\Rightarrow A^2 [(\omega^2 - p^2) + 4\lambda^2 p^2] = f_0^2$$

$$\Rightarrow A^2 = \frac{f_0^2}{(\omega^2 - p^2)^2 + 4\lambda^2 p^2}$$

$$\Rightarrow A = \frac{f_0}{\sqrt{(\omega^2 - p^2)^2 + 4\lambda^2 p^2}}$$

Dividing eqⁿ(v) to (iv) ⇒

$$\frac{f_0 \sin \theta}{f_0 \cos \theta} = \frac{2\lambda A p}{A (\omega^2 - p^2)}$$

$$\Rightarrow \tan \theta = \frac{2\lambda p}{1(\omega^2 - p^2)}$$

$$\Rightarrow \theta = \tan^{-1} \left(\frac{2\lambda p}{\omega^2 - p^2} \right)$$

* Solution of forced vibration,

$$y = \frac{f_0}{\sqrt{(\omega^2 - p^2)^2 + 4\lambda^2 p^2}} \cdot \sin \left(pt - \tan^{-1} \frac{2\lambda p}{(\omega^2 - p^2)} \right)$$

Resonance

Maximum displacement of a driven oscillator,

$$A^2 = \frac{f_0^2}{(\omega^2 - \rho^2)^2 + 4\lambda^2\rho^2}$$

The amplitude is maximum when the denominator is minimum.

$$\frac{d}{dp} [(\omega^2 - \rho^2)^2 + 4\lambda^2\rho^2] = 0$$

$$\Rightarrow 2(-2\rho)(\omega^2 - \rho^2) + 4\lambda^2 \cdot 2\rho = 0$$

$$\Rightarrow -4\rho(\omega^2 - \rho^2) + 8\lambda^2\rho = 0$$

$$\Rightarrow \rho(\rho^2 - \omega^2) + 2\lambda^2\rho = 0$$

$$\Rightarrow \rho(\rho^2 - \omega^2 + 2\lambda^2) = 0 \quad \because \rho \neq 0$$

$$\therefore \rho^2 - \omega^2 - 2\lambda^2 = 0$$

$$\Rightarrow \rho^2 = \omega^2 - 2\lambda^2$$

$$\therefore \rho = \sqrt{\omega^2 - 2\lambda^2}$$

Then the amplitude will be maximum when the driven frequency

$$\frac{P_n}{2\pi} = \sqrt{\omega^2 - 2\lambda^2}$$

resonant frequency

The state of vibration when the amplitude of the driven oscillator is maximum is called resonance.

In the absence of damping, resonance take place when natural frequency of oscillator is equal to the frequency of driving force,

The maximum Amplitude,

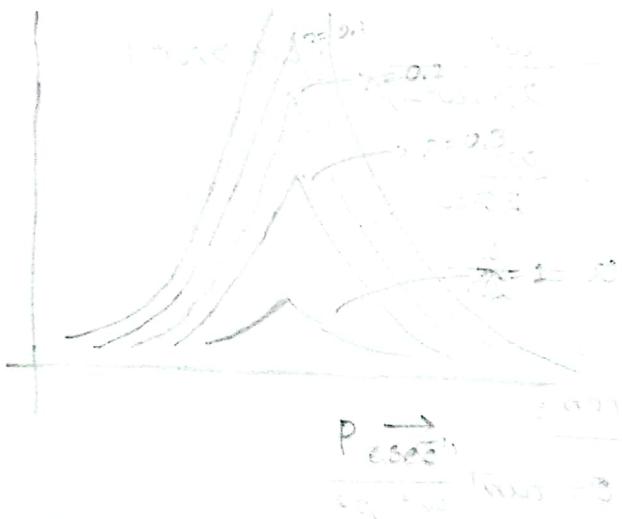
$$A_{max} = \frac{f_0}{\sqrt{(\omega^2 - \rho^2)^2 + 4\lambda^2\rho^2}}$$

$$\left(\frac{f_0}{\sqrt{(\omega^2 - \rho^2)^2 + 4\lambda^2\rho^2}} \right)_{max} = 3$$

$$\therefore A_{max} = \frac{f_0}{2\sqrt{\omega^2 - \gamma^2}}$$

$$= \frac{f_0}{\sqrt{4\gamma^4 + 4\gamma^2\omega^2 - 8\gamma^4}}$$

$$= \frac{f_0}{\sqrt{4\gamma^2\omega^2 - 4\gamma^4}}$$



$\gamma = 0$, No damping, $A_{max} \rightarrow \infty$

✓ Sharpness of Resonance :

It is a measure of the rate of fall in amplitude from its maximum value at resonant frequency on either side of it. The sharper the fall in amplitude sharper the resonance.

• Forced vibration of quality ~~factor~~ factor.

✓ Quality factor of forced vibration:

The ratio of the response of the oscillation when the driven frequency is equal to the resonant frequency to the response when the driven frequency is zero is called the quality factor of the forced vibration.

$$A = \frac{f_0}{\sqrt{(\omega^2 - p^2) + 4\gamma^2 p^2}} \quad \text{when } \begin{cases} p \text{ driving frequency} \\ p=0 \end{cases}$$

$$A_{p=0} = \frac{f_0}{\omega^2}$$

when, $\rho = \rho_n$ $A_{\max} = \frac{f_0}{2\pi\sqrt{\omega^2 - \eta^2}}$

$$\text{Quality factor} = \frac{A_{\max}}{P_{\rho=0}} = \frac{\frac{f_0}{2\pi\sqrt{\omega^2 - \eta^2}}}{\frac{f_0}{\omega^2}} = \frac{\omega^2}{2\pi\sqrt{\omega^2 - \eta^2}}$$

$$= \frac{\omega^2}{2\pi\sqrt{\omega^2 - \eta^2}} \quad [\omega^2 \gg \eta^2]$$

$$= \frac{\omega^2}{2\pi\omega}$$

$$\alpha = \frac{\omega}{2\pi}$$

Phase of the driven Oscillator:

$$y = A \sin(pt - \theta) \quad \theta = \tan^{-1} \frac{2\pi\rho}{\omega^2 - \rho^2}$$

The phase angle θ depends upon damping and relative values of ω and ρ , following cases are,

(i) when $\rho < \omega$; i.e

$\tan\theta$ will positive and values of θ lies between 0 to $\frac{\pi}{2}$ for all values of η .

(ii) when $\rho > \omega$; i.e, the $\tan\theta$ will be a negative quantity and the values of θ lies between $\frac{\pi}{2}$ to π for all values of η .

(iii) when, $\rho = \omega$

$$\tan\theta = 0, \quad \theta = \frac{\pi}{2}$$

absorption

Date: 24.01.2023

Tuesday

Power of driven Oscillator:

$$F = F_0 \sin pt$$

$\Rightarrow dt \rightarrow dy \rightarrow$ displacement

$$dE = F dy$$

$$\Rightarrow \frac{dE}{dt} = F \frac{dy}{dt} \quad \text{--- (i)}$$

$$y = A \sin(pt - \theta)$$

$$= \frac{f_0}{\sqrt{(\omega^2 - p^2)^2 + 4\lambda^2 p^2}} \times \sin(pt - \theta)$$

$$\frac{dy}{dt} = \frac{p f_0}{\sqrt{(\omega^2 - p^2)^2 + 4\lambda^2 p^2}} \cos(pt - \theta)$$

$$v = v_0 \cos(pt - \theta)$$

$$v_0 = \frac{p f_0}{\sqrt{(\omega^2 - p^2)^2 + 4\lambda^2 p^2}}$$

$$F = F_0 \sin pt$$

$$\Rightarrow F = m f_0 \sin pt$$

$$\Rightarrow P \Rightarrow \frac{dE}{dt} = F \frac{dy}{dt}$$

$$P_{ab} = m f_0 \sin pt = \frac{p f_0}{\sqrt{(\omega^2 - p^2)^2 + 4\lambda^2 p^2}} \cos(pt - \theta)$$

$$\Rightarrow P_{ab} = \frac{m f_0^2 p}{\sqrt{(\omega^2 - p^2)^2 + 4\lambda^2 p^2}} \sin pt \cos(pt - \theta)$$

Average power absorption:

$$P_{average} = \frac{1}{T} \int_0^T \frac{m f_0^2 p}{\sqrt{(\omega^2 - p^2)^2 + 4\lambda^2 p^2}} \times \sin pt \cos(pt - \theta)$$

$$= \frac{m f_0^2 p}{\sqrt{(\omega^2 - p^2)^2 + 4\lambda^2 p^2}} \times \frac{1}{T} \int_0^T \sin pt \cos(pt - \theta)$$

$$\text{Now, } \frac{1}{T} \times \frac{1}{2} \int_0^T 2 \sin pt \cos(pt - \theta)$$

$$= \frac{1}{2T} \int_0^T [\sin(2pt - \theta) + \sin \theta] dt$$

$$= \frac{1}{2T} \sin \theta T = \frac{1}{2} \sin \theta$$

$$P_{average} = \frac{m f_0^2 p}{\sqrt{(\omega^2 - p^2)^2 + 4\lambda^2 p^2}} \times \frac{1}{2} \sin \theta$$

$$f_0 \sin \theta = 2 \Delta AP$$

We know that,

$$f_0 \sin \theta = 2 \Delta AP$$

$$\Rightarrow \sin \theta = \frac{2 \Delta AP}{f_0}$$

$$\therefore P_{average} = \frac{m f_0^2 p}{\sqrt{(\omega^2 - p^2)^2 + 4\lambda^2 p^2}} \cdot \frac{1}{2} \cdot \frac{2 \Delta AP}{f_0}$$

$$= \frac{m f_0^2 p^2 \lambda}{\sqrt{(\omega^2 - p^2)^2 + 4\lambda^2 p^2}} \cdot \frac{f_0}{\sqrt{(\omega^2 - p^2)^2 + 4\lambda^2 p^2}}$$

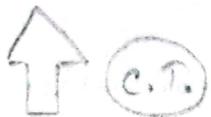
$$\therefore P_{average} = \frac{m f_0^2 p^2 \lambda}{(\omega^2 - p^2)^2 + 4\lambda^2 p^2}$$

$$\frac{F_0}{m} = f_0$$

The average power absorbed is maximum when, $P = \omega$

$$P_{\max} = \frac{m f_0^2 \rho^2 A}{4 \omega^2 P^2}$$

$$\boxed{P_{\max} = \frac{1}{4} \cdot \frac{m f_0^2}{A}}$$



Superposition :

Composition of two simple harmonic vibration at right angles to each other having equal frequencies but different in phase and amplitude.

$$x = a \sin(\omega t + \varphi) \quad \text{--- (i)}$$

$$y = b \sin \omega t \quad \text{--- (ii)}$$

$$\frac{x}{a} = \sin(\omega t + \varphi)$$

$$= \sin(\omega t) \cos \varphi + \cos \omega t \sin \varphi$$

$$= \sin \omega t \cos \varphi + \sqrt{1 - \sin^2 \omega t} \sin \varphi \quad \text{--- (iii)}$$

From eqn(ii),

$$\frac{y}{b} = \sin \omega t$$

From, eqn(iii),

$$\frac{x}{a} = \frac{y}{b} \cos \varphi + \sqrt{1 - \frac{y^2}{b^2}} \sin \varphi$$

$$\Rightarrow \frac{x^2}{a^2} + \frac{y^2}{b^2} \cos^2 \varphi - 2 \frac{x}{a} \cdot \frac{y}{b} \cos \varphi = \left(1 - \frac{y^2}{b^2}\right) \sin^2 \varphi$$

$$\Rightarrow \frac{x^2}{a^2} + \frac{y^2}{b^2} - \frac{2xy}{ab} \cos \varphi = \sin^2 \varphi \quad \text{--- (iv)}$$

This is the general equation of resultant vibrations.

Case - 1:

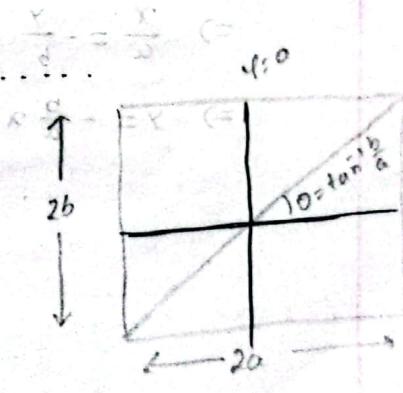
$$\varphi = 0, 2n, 4n, \dots \dots \quad n = 0, 1, 2, \dots \dots$$

$$\sin \varphi = 0, \quad \cos \varphi = 1$$

$$(iv) \rightarrow \frac{x^2}{a^2} + \frac{y^2}{b^2} - \frac{2xy}{ab} = 0$$

$$\Rightarrow \left(\frac{x}{a} - \frac{y}{b}\right)^2 = 0$$

$$\Rightarrow \frac{x}{a} = \frac{y}{b} \quad \Rightarrow y = \frac{b}{a}x$$



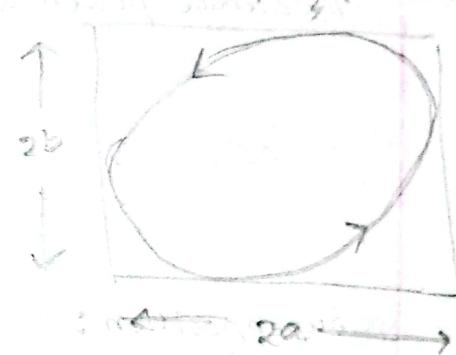
Case - 2:

$$\varphi = \frac{\pi}{4}$$

$$\cos \varphi = \sin \varphi = \frac{1}{\sqrt{2}}$$

$$\boxed{\frac{x^2}{a^2} + \frac{y^2}{b^2} - \frac{\sqrt{2}xy}{ab} = \frac{1}{2}}$$

anticlockwise



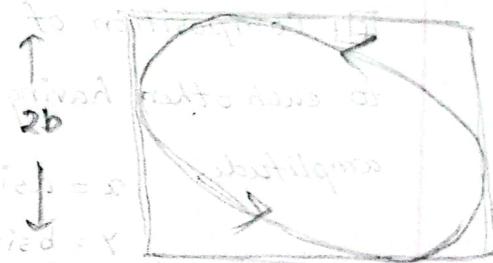
case - 4:

$$\varphi = \frac{3\pi}{4}$$

$$\frac{x^2}{a^2} + \frac{y^2}{b^2} + \frac{\sqrt{2}xy}{ab} = \frac{1}{2}$$

↑

$$(\varphi + \text{initial}) = \frac{\pi}{2} \rightarrow (\varphi + 3\omega)_{\text{final}} = \omega$$



case - 3:

Case - 3:

$$\varphi = \frac{\pi}{2}$$

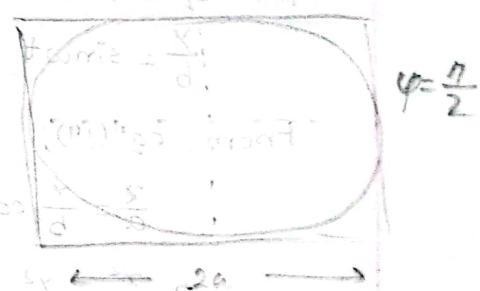
$$\sin \varphi = 1, \cos \varphi = 0.$$

$$\frac{x^2}{a^2} + \frac{y^2}{b^2} = 1$$

when, $a = b$,

$$x^2 + y^2 = a$$

$$x^2 \cos^2 \left(\frac{\pi}{2} - \theta \right) = y^2 \cos^2 \theta \cdot \frac{x^2}{a^2} + y^2 \cos^2 \theta \cdot \frac{b^2}{a^2}$$



Case - 5:

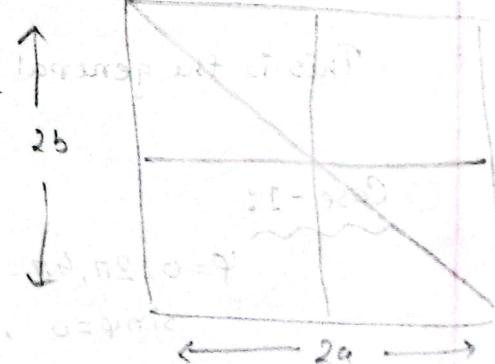
$$\varphi = \pi$$

$$\sin \varphi = 0, \cos \varphi = -1$$

$$\frac{x^2}{a^2} + \frac{y^2}{b^2} + \frac{2xy}{ab} = 0$$

$$\Rightarrow \frac{x}{a} = -\frac{y}{b}$$

$$\Rightarrow y = -\frac{b}{a}x$$



$$y = -\frac{bx}{a}, \frac{dy}{dx} = -\frac{b}{a}$$

$$0 = -\frac{b}{a} \left(\frac{x}{a} + \frac{y}{b} \right)$$

$$0 = -\frac{b}{a} \left(\frac{x}{a} + -\frac{b}{a}x \right)$$

$$0 = -\frac{b}{a} \left(\frac{x}{a} - \frac{b}{a}x \right)$$

Question
Hand

Sun/Monday
A.T. probability 50%

Lissasous figure:

The composition of two SHB in mutually perpendicular directions gives rise to an elliptical path. The actual shape of the wave will depend upon the phase difference ϕ between the two vibrations, and also on the ratio of the frequencies of the component of vibrations. These figures are called lissasous figure.

Composition of two SHB of same frequency but different phase and amplitude,

$$y_1 = a_1 \sin(\omega t + \alpha_1)$$

$$y_2 = a_2 \sin(\omega t + \alpha_2)$$

Resultant displacement,

$$Y = Y_1 + Y_2$$

$$\Rightarrow a_1 \sin(\omega t + \alpha_1) + a_2 \sin(\omega t + \alpha_2)$$

wave equation, amplitude, other values.

Thermodynamics

Date: 25.01.2023

Wednesday

Thermodynamic System

A system may be defined as a definite quantity of matter (solid, liquid or gases) bounded by some closed surface.

→ Homogenous System → phase uniform

When a system is completely uniform throughout, as such a gas or mixture of a gas.

4) Heterogenous System:

When a system consists of two or more phases.

A system maybe separated from its surrounding by a real or imaginary boundary throughout which heat or mechanical energy may pass.

Thermodynamic Variables:

The thermodynamic state or macroscopic state of a system is determined by four observable properties. These properties are the composition, pressure, volume & temperature, which are called the variable of state.

- For system, in general, there is a equation of state,

$$f(P, V, T) = 0$$

For, ideal gas,

$$PV = RT$$

For, Real gas,

$$(P + \frac{a}{V^2})(V - b) = RT, \text{ when } a \text{ and } b \text{ vanderwal's constant}$$

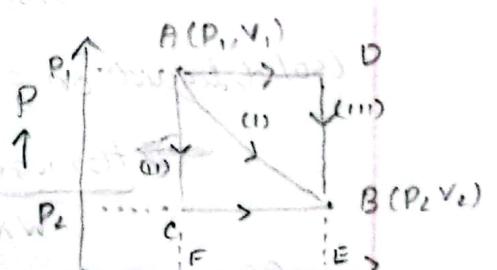
~~Work: Examples of variables of state~~

Must

A path dependant function!

A system can be taken from state A to state B by many ways in figure.

Let the co-ordinates of state A and B (P_1, V_1) and (P_2, V_2)



respectively,

(i) starting from A, the pressure is continuously decreased from P_1 to P_2 along the wave AB so that volume increased from V_1 to V_2 .

- The work done by the system,

$$W = \int_{V_1}^{V_2} P dV = \text{area } ABEF$$

(ii) Starting from point A, the volume V_1 is kept constant in going from A to C, the pressure decreases from P_1 to P_2 , and then P_2 kept constant from C to B.

- The work done in this process $W_L = P_2(V_2 - V_1) = \text{area } CB EF$

(iii) Again, starting from point A, the pressure P_1 is kept constant going from A to D and volume V_2 is kept constant from D to B.

The work done in this process,

$$W_3 = P_1(V_2 - V_1) = \text{area } ADEF$$

$$\therefore W_1 \neq W_2 \neq W_3$$

Work is a path dependant system

Thermodynamic Equilibrium:

(i) Mechanical equilibrium:

For a system to be mechanical equilibrium, there should be no macroscopic movement within the system or of the system with respect to its surrounding.

Bridget

- Explain zeroth law of thermodynamics

(ii) Thermal Equilibrium:

For a system to be thermal equilibrium, there should be no temp. between the parts of the system or between the system and surrounding.

(iii) Chemical Equilibrium:

Boxed Zeroth Law of thermodynamics:

If two bodies A and B are each separately in thermal equilibrium with a third body C, then A and B are also in their thermal equilibrium with each other.

Boxed Internal Energy:

The internal energy of a system is called interval energy. It is the sum of following form of energy of the system,

- Kinetic energy due to translational, rotational and vibrational motion of the molecular, all which depends on the temp.
- Potential energ due to inten-molecular force
- the energy of electron and nuclei.

First law of thermodynamics:

When a certain amount of heat (Q) is supplied to a system which does external work w in passing from state 1 to state 2, the amount of heat is equal to sum of the increase of the internal energy of the system and the external work done by the system.

$$Q = (U_2 - U_1) + w$$

For small change, $dQ = dU + dw$

Specific heat:

Heat capacity per unit mass is called specific heat.

$$\text{Specific heat} = \frac{Q}{m\Delta T} = c$$

The specific heat of materials is defined as the quantity of heat required to raise the temp. of unit mass of the material, through 1 degree.

$C_p \rightarrow$ at constant pressure

$$C_p = \frac{dQ}{dT}$$

 $C_p > C_v$

$C_v \rightarrow$ at constant volume

$$C_v = \frac{dQ}{dT}$$

Reason ?

* Adiabatic / Isothermal / Isotopic curve

Question

* Slope graph (?)

Application of first law of thermodynamics:

Date : 31.01.2023

Tuesday

Specific heat of a gas:

v, T , are independent variable. $U = f(v, T)$ —①

Differentiating this equation

$$dU = \left(\frac{\partial U}{\partial T}\right)_v dT + \left(\frac{\partial U}{\partial v}\right)_T dv \quad \text{—②}$$

From 1st law of thermodynamics,

$$dQ = dU + dw = dU + pdv \quad \text{—③}$$

Putting the value of dU in eqⁿ(③),

$$dQ = \left(\frac{\partial U}{\partial T}\right)_v dT + \left(\frac{\partial U}{\partial v}\right)_T dv + pdv$$

$$\Rightarrow \frac{dQ}{dT} = \left(\frac{\partial U}{\partial T}\right)_v + \left(\frac{\partial U}{\partial v}\right)_T \cdot \frac{dv}{dT} + p \frac{dv}{dT}$$

$$= \left(\frac{\partial U}{\partial T}\right)_v + \left[p + \left(\frac{\partial U}{\partial v}\right)_T\right] \frac{dv}{dT} \quad \text{—④}$$

At constant volume, ($\frac{dv}{dT} = 0$)

$$\frac{dQ}{dT} = \left(\frac{\partial U}{\partial T}\right)_v = C_v$$

At constant pressure,

$$\frac{dQ}{dT} = C_p$$

Putting this value in eqⁿ(④),

$$C_p = C_v + \left[p + \left(\frac{\partial U}{\partial v}\right)_T\right] \frac{dv}{dT}$$

$$\Rightarrow C_p - C_v = p \frac{dv}{dT} \quad \because \left(\frac{\partial U}{\partial v}\right)_T = 0$$

We know that,

$$PV = RT$$

$$\Rightarrow P \frac{dv}{dT} = R$$

$$\therefore C_p - C_v = R$$

For reversible adiabatic process, $dQ = 0$

From equation,

$$0 = \left(\frac{\partial U}{\partial T}\right)_V dT + \left(\frac{\partial U}{\partial V}\right)_T dV + pdV$$

$$= cvdT + \left[\left(\frac{\partial U}{\partial V}\right)_T + p\right] dV$$

$$= \frac{cvdT}{dV} + \left[p + \left(\frac{\partial U}{\partial V}\right)_T\right]$$

$$\therefore \frac{cvdT}{dV} = -p \quad \left[\left(\frac{\partial U}{\partial V}\right)_T = 0\right]$$

$$\Rightarrow \frac{cvdT}{dV} = \frac{cv - cp}{\alpha V}$$

$$\Rightarrow \frac{dT}{dV} = \frac{cv - cp}{\alpha V cv}$$

We know, Isochoric volume co-efficient expansion

$$\text{Memorise } \alpha = \frac{1}{V} \left(\frac{\partial V}{\partial T}\right)_P$$

$$\Rightarrow \alpha V = \left(\frac{\partial V}{\partial T}\right)_P$$

$$\Rightarrow \alpha V = \frac{cp - cv}{p}$$

$$\therefore \gamma_p = \frac{cp - cv}{\alpha V}$$

- This expansion holds for adiabatic reversible process.

Isochoric Process:

$$dV = 0,$$

$$dQ = dU$$

Isobaric Process:

Pressure constant

$$dQ = dU + pdV$$

Isothermal process:

Temp constant

Adiabatic Process:

No heat transfer

$$dQ = 0$$

$$dU = -pdV$$

$$dU = 0$$

$$dQ = pdV$$

Isentropic Process:

No change of entropy

Work done during Isothermal Process

From the definition of work

$$dW = PdV$$

$$\Rightarrow \int dW = \int_{V_1}^{V_2} PdV$$

$$\Rightarrow W = \int_{V_1}^{V_2} PdV$$

$$W = RT \ln\left(\frac{V_2}{V_1}\right)$$

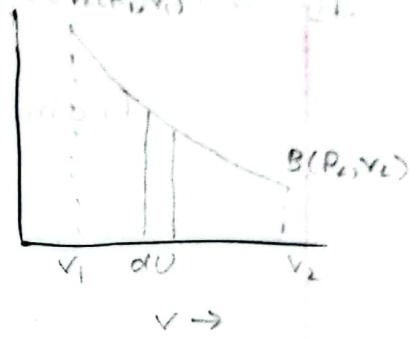
$$W = RT \ln\left(\frac{P_1}{P_2}\right)$$

$$PV = RT$$

$$\Rightarrow P = \frac{RT}{V}$$

$$P_1 V_1 = P_2 V_2$$

$$\Rightarrow \frac{P_1}{P_2} = \frac{V_1}{V_2}$$



Work done during Adiabatic process

$$\int dW = \int_{V_1}^{V_2} PdV$$

$$PV^r = K$$

$$\Rightarrow W = \int_{V_1}^{V_2} PdV$$

$$= \int_{V_1}^{V_2} K \frac{dV}{V^r}$$

$$= K \left[\frac{V^{-r+1}}{-r+1} \right]_{V_1}^{V_2}$$

$$= \frac{K}{1-r} [V_2^{1-r} - V_1^{1-r}]$$

$$= \frac{1}{1-r} \left[\frac{P_2 V_2^r}{V_2^{r-1}} - \frac{P_1 V_1^r}{V_1^{r-1}} \right]$$

$$= \frac{1}{1-r} [P_2 V_2 - P_1 V_1]$$

$$= \frac{1}{1-r} [RT_2 - RT_1]$$

$$= \frac{R}{1-r} [T_2 - T_1]$$

Reversible Process :

A process which can be retraced in the opposite direction so that the working substance passes through exactly the same states in all respects as in the direct process is called reversible process.

Inreversible Process:

A process which can not retracked in the opposite direction is called an irreversible process.

④ Slope of adiabatic and isothermal process:

Adiabatic slope is more steeper than isothermal slope.

$$PV = \cancel{RT} \propto$$

$$\Rightarrow \frac{d}{dV}(PV) = k$$

$$\Rightarrow P \frac{dV}{dV} + V \frac{dP}{dV} = \frac{d}{dV}(k)$$

$$PV = R T$$

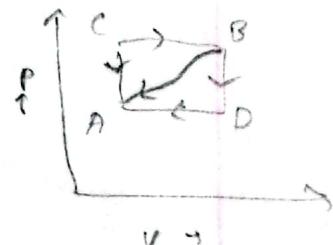
$$PV = R T$$

$$\Rightarrow P$$

④ When a system is taken from the state A to the state B, along the Path ACB, 80 joules of the heat flows into the system, and system does 30 joules of work:

(a) How much heat flows into the system

along the path ADB, if the work done is 10 Joules.



(b) The system is ~~retraced~~^{retarded} from B, the state B to the state A along curved path. The workdone on the system is 20 joules

Does the system absorb liberate heat and how much

(c) If $V_A = 0$, $V_D = 40$ joules, ~~final~~^{find} the heat absorbed in the process AD and DB

Date: 01.02.2023

Wednesday

Heat Engine:

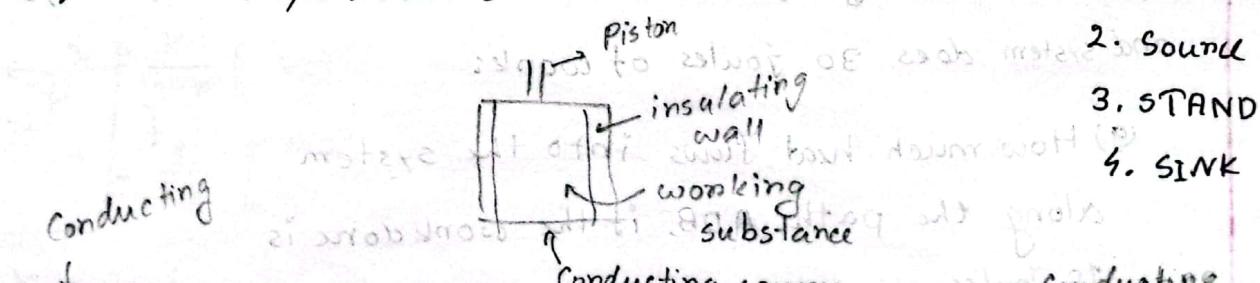
A practical machine which converts heat into mechanical work is called a heat engine. Heat engines in their operation absorb heat at a higher temperature, converts parts of it into mechanical work, reject remained heat at a low temperature.

Definition of Efficiency:

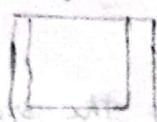
The efficiency η of a heat engine is defined as the ratio of the mechanical work done by the engine in one cycle to heat absorbed from the high temp. source, $\eta = \frac{Q_1 - Q_2}{Q_1}$

Carnot's Ideal Heat Engine:

Sadi Carnot conceived a theoretical engine which is free from all practical imperfections due to existing materials.



Conducting
↓



T₁

T₂



STAND

Stand

Insulating

wall

Conducting

The Cycle

Carnot's Ideal Heat Engine:

A cycle in which the working substance starting from a given condition of temperature, pressure, volume is made to undergo two successive expansion and then two successive compression at two end of which the working substance is brought back to its initial condition is called Carnot cycle.

Isothermal Expansion:

The cylinder is first placed on the source, so that the gas acquires the temperature T_1 of the source. Then the gas undergoes isothermal expansion at T_1 .

Let the working substance during isothermal expansion goes from its initial state $A(P_1, V_1, T_1)$ to state $B(P_2, V_2, T_1)$ along AB.

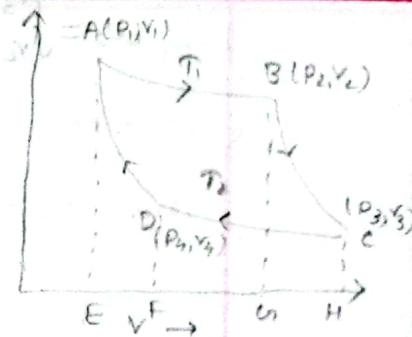
In this process substance absorbs heat Q_1 and does work, w

$$Q_1 = w = \int_{V_1}^{V_2} P dV = RT_1 \ln \frac{V_2}{V_1}$$

Adiabatic Expansion:

The cylinder is now removed from the source and is placed on the insulating stand. The gas is allowed to undergo slow adiabatic expansion.

This operation is represented by adiabatic BC. The temperature fall to T_2 . The work done in this process is,



$$W_2 = \int_{V_2}^{V_3} P dV = k \int_{V_2}^{V_3} \frac{dV}{V^r} = \frac{k V_3^{1-r} - k V_2^{1-r}}{1-r}$$

$$= \frac{P_3 V_3 - P_2 V_2}{1-r} = \frac{RT_2 - RT_1}{1-r}$$

$$W_2 = \frac{R}{1-r} [T_2 - T_1]$$

3. Isothermal Expansion:

The cylinder is now removed from the insulating stand and is placed on the sink which is at T_2 temperature. Thus gas undergoes isothermal compression at a constant temperature T_2 .

This operation is represented by CD.

In this process substance rejects heat Q_2 and does work w_3

$$Q_2 = w_3 = \int_{V_3}^{V_4} P dV = RT_2 \ln\left(\frac{V_4}{V_3}\right)$$

$\therefore w_3 = -RT_2 \ln\left(\frac{V_3}{V_4}\right)$ -ve sign indicates that work is done on the working substance.

4. Adiabatic Compression:

The cylinder is now removed from sink and again placed on the insulating stand. The adiabatic compression is continued till the gas is back to its original condition, i.e., state A (P_1, V_1) thus completing one full cycle, $w_4 = - \int_{V_1}^{V_2} P dV$

$$= \frac{-R(T_1 - T_2)}{r-1}$$

Work done by the engine per cycle:

During the above cycle, the working substance absorbs an amount of heat Q_1 from source and rejects Q_2 to the sink.

The net amount of heat absorbed = $Q_1 - Q_2$

$$\text{Net amount of work} = W_1 + W_2 + W_3 + W_4 = W_1 + W_3 \quad [W_2 = -W_4]$$

We know that,

Net heat absorbed = Net work done

$$Q_1 - Q_2 = W_1 + W_3 = RT_1 \ln \frac{V_2}{V_1} - RT_2 \ln \frac{V_3}{V_4} \quad (1)$$

Since, A and D lie on same adiabatic line.

$$T_1 V_1^{r-1} = T_2 V_4^{r-1}$$

$$\Rightarrow \frac{T_2}{T_1} = \left(\frac{V_1}{V_4}\right)^{r-1} \quad (ii)$$

Again,

B and C also lie on the same adiabatic line.

$$T_1 V_2^{r-1} = T_2 V_3^{r-1}$$

$$\Rightarrow T_2 = \left(\frac{V_2}{V_3}\right)^{r-1} \quad (iii)$$

$$\text{From (1) and (ii)} \Rightarrow \left(\frac{V_1}{V_4}\right)^{r-1} = \left(\frac{V_2}{V_3}\right)^{r-1} \Rightarrow \frac{V_1}{V_2} = \frac{V_3}{V_4} \quad \Rightarrow \frac{V_4}{V_1} = \frac{V_3}{V_2}$$

$$Q_1 - Q_2 = RT_1 \ln \frac{V_2}{V_1} - RT_2 \ln \frac{V_2}{V_1}$$

$$W = R(T_1 - T_2) \ln \frac{V_2}{V_1}$$

$$\text{Efficiency, } \eta = \frac{\text{net work done}}{\text{absorbed heat}} = \frac{Q_1 - Q_2}{Q_1} = \frac{R(T_1 - T_2) \ln \frac{V_2}{V_1}}{RT_1 \ln \frac{V_2}{V_1}}$$

$$\boxed{\eta = \frac{T_1 - T_2}{T_1} \times 100\%}$$

Second Law of Thermodynamics

Date : 07.02.2023

Tuesday

It is impossible for a self-acting machine working in a cyclic process, unaided by external agency, to transfer heat from a body at a lower temperature to a body at higher temperature.

Kelvin's statement:

It is impossible to engine which, working in complete cycle will produce no effort other than the raising of weight and the cooling of the heat reservoir.

Concent Entropy:

The quantity of entropy found to remain constant in adiabatic process just as temperature remain constant in an isothermal process. The entropy can be defined as the thermal property of a working substance which is constant in adiabatic process.

Change in Entropy :

Let us consider ~~non~~ reversible Carnot cycles bounded by the same two adiabatic L and isothermal T_1, T_2, T_3 as shown in Figure.

During ABCD cycle, an amount of heat Q_1 is absorbed in T_1 temperature and Q_2 heat reject at T_2 temperature.



Then the effectively,

$$\eta = \frac{Q_1 - Q_2}{Q_1} = \frac{T_1 - T_2}{T_1}$$

$$\Rightarrow \eta = 1 - \frac{Q_2}{Q_1} = 1 - \frac{T_2}{T_1}$$

$$\Rightarrow \frac{Q_1}{T_1} = \frac{Q_2}{T_2} \quad \text{---(i)}$$

Similarly for DCEF

$$\frac{Q_2}{T_2} = \frac{Q_3}{T_3} \quad \text{---(ii)}$$

From eq (i) and (ii)

$$\frac{Q_1}{T_1} = \frac{Q_2}{T_2} = \frac{Q_3}{T_3} = \text{constant}$$

$$\frac{Q}{T} = \text{constant}$$

If δQ a small amount of heat is rejected or absorbed

$$\frac{\delta Q}{T} = \text{constant}$$

Constant ratio is called the change of entropy.

$$\delta S = \frac{\delta Q}{T}$$

$$\Rightarrow \int_{S_A}^{S_B} dS = \int_A^B \frac{dQ}{T}$$

$$\Rightarrow [S_B - S_A = \int_A^B \frac{dQ}{T}]$$

Change in entropy in reversible process?

In reversible cycle, there are two isothermal and two adiabatic process, there is neither absorption nor rejection of heat and there no change in entropy.

During expansion an amount of heat Q_1 is absorbed at temperature T_1 and in compression there is . . . heat reject at T_2 temperature.

Change in entropy,

$$ds = \frac{Q_1}{T_1} - \frac{Q_2}{T_2} \quad \text{---(i)}$$

We know that for reversible cycle,

$$\frac{Q_1}{T_1} = \frac{Q_2}{T_2}$$

$$\Rightarrow ds = 0$$

i.e. s is constant

The entropy of a body in all reversible cycle remain constant.

Change in entropy in irreversible process:

Let us consider an engine performing reversible cycle in which working substance ... absorbs heat Q_1 at T_1 temperature and rejects heat Q_2 at T_2 temperature.

$$\text{Efficiency, } \eta' = \frac{Q_1 - Q_2}{Q_1} = \boxed{1 - \frac{Q_2}{Q_1}} \rightarrow \text{Inversible}$$

But according to carnot's theorem, this efficiency is less than that of ~~rever~~ reversible engine working in the same temperature

Then $\boxed{\eta = 1 - \frac{T_2}{T_1}}$

Reversible

Thus $\boxed{\eta' < \eta}$

$\boxed{1 - \frac{Q_2}{Q_1} < 1 - \frac{T_2}{T_1}}$

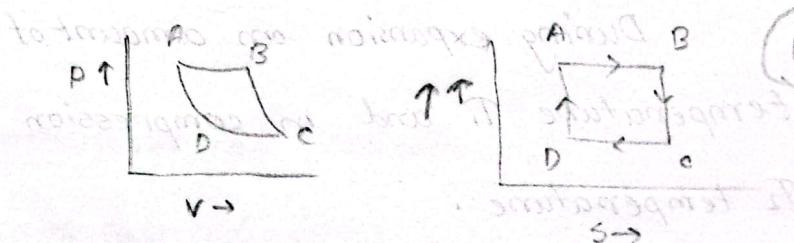
$$\Rightarrow \frac{Q_2}{Q_1} > \frac{T_2}{T_1}$$

$$\Rightarrow \frac{Q_2}{T_2} > \frac{Q_1}{T_1}$$

$$\Rightarrow \left(\frac{Q_2}{T_2} - \frac{Q_1}{T_1} \right) > 0$$

There is an increase in entropy of the system during irreversible process.

T-S Diagram:



Explain

Third law of thermodynamics:

At absolute zero temperature the entropy tends to zero and molecular a substance or a system are in the perfect order.

Entropy of a perfect gas:

Consider 1 gm of perfect gas occupies a volume v at P and T ,
Let dQ be the heat supplied to the gas,

$$dQ = dU + dW \quad \text{(i)}$$

$$dU = dQ - dW \quad dU = C_v dT$$

$$dQ = C_v dT + P dV$$

Change of Entropy,

$$\Delta S = \int \frac{dQ}{T} = \int \frac{1}{T} (C_v dT + P dV)$$

$$\Delta S = \int C_v \frac{dT}{T} + \int \frac{P}{T} dV \quad \text{(ii)}$$

(i) S in terms of temperature and volume,

$$PV = RT \quad \left(T = \frac{PV}{R}, \frac{dV}{T} = \frac{V}{T^2} \right) \quad \left(\frac{P}{T} = \frac{R}{V} \right)$$

$$\begin{aligned} \Delta S &= \int_{T_1}^{T_2} C_v \frac{dT}{T} + \int_{V_1}^{V_2} \frac{R}{V} dV \\ &= C_v \left[\ln T \right]_{T_1}^{T_2} + R \left[\ln V \right]_{V_1}^{V_2} \end{aligned}$$

$$\boxed{\Delta S = C_v \ln \frac{T_2}{T_1} + (C_p - C_v) \ln \frac{V_2}{V_1}}$$

(ii) S in terms of temperature and - - - - - ,

$$PV = RT$$

$$\Rightarrow V = \frac{RT}{P}$$

$$PdV + VdP = RdT$$

$$\Rightarrow PdV = RdT - VdP$$

$$\begin{aligned} \Delta S &= \int C_v \frac{dT}{T} + \int \frac{RdT - VdP}{T} \\ &= \int C_v \frac{dT}{T} + R \int \frac{dT}{T} - \int \frac{VdP}{T} \\ &= \int C_v \frac{dT}{T} + (C_p - C_v) \int \frac{dT}{T} - P \int \frac{dP}{T} \\ &= C_p \int \frac{dT}{T} + (C_p - C_v) \int \frac{dP}{P} \end{aligned}$$

$$\therefore \Delta S = C_p \left[\ln T \right]_{T_1}^{T_2} - (C_p - C_v) \left[\ln P \right]_{P_1}^{P_2}$$

$$\therefore \Delta S = C_p \ln \frac{T_2}{T_1} - (C_p - C_v) \ln \frac{P_2}{P_1}$$

(iii) S in terms of pressure and volume,

$$PV = RT$$

$$\Rightarrow T = \frac{PV}{R}$$

$$\Rightarrow dT = \frac{Pdv + vdp}{R}$$

$$\Delta S = C_v \int \frac{dT}{T} + \int \frac{P}{T} dv$$

$$= C_v \int \frac{Pdv + vdp}{RT} + \int \frac{R}{V} dv$$

$$= C_v \int \frac{Pdv}{RT} + \int C_v \frac{vdp}{RT} + R \int \frac{dv}{V}$$

$$= C_v \int \frac{dv}{V} \times \frac{T}{R} + C_v \int \frac{dp}{P} \times \frac{1}{R} + (C_p - C_v) \int \frac{dv}{V}$$

$$\cancel{\Delta S} = \frac{C_v T}{R} \ln \frac{V_2}{V_1} + \frac{C_v}{R} \ln \frac{P_2}{P_1} + (C_p - C_v) \times \ln \frac{V_2}{V_1}$$

$$PV = RT$$

$$\Rightarrow \frac{P}{T} = \frac{R}{V}$$

NOT RIght

Diagrams - does not correspond to correct in 2 iii

$$T_{bog} = q_{bog} + V_{bog}$$

$$q_{bog} - T_{bog} = V_{bog}$$

$$T_b = Vq$$

$$\frac{T_b}{q} = V$$

$$\frac{q_{bog} - T_{bog}}{T} + \frac{T_{bog}}{T} V \} = 2A$$

$$\frac{q_{bog}}{T} \left(1 - \frac{T_{bog}}{T} \right) + \frac{T_{bog}}{T} V \} = 2A$$

$$\frac{q_{bog}}{T} \left(1 - \frac{T_{bog}}{T} \right) + \frac{T_{bog}}{T} V \} = 2A$$

LASER PHYSICS

Date: 08.02.2023

Wednesday

- Quantum Optics

- **LASER** is a photonic device

Photonics

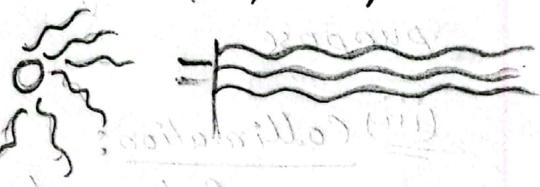
Light Amplification through
Stimulated emission of
Radiation

not amplify

- Light Oscillation through Stimulated

Emission of Radiation, **LASER**

Normal light \rightarrow tungsten light,
mercury lamps



- ④ The unique property of laser is

that its light waves travel very long distance with a very little divergence. In case of conventional

Light source, the light is emitted in jumble of separate waves that cancel each other at random and hence can travel very short distance.

In laser, the light waves are exactly in step with each other and have a fixed phase relationship.

■ Definition of Laser:

Laser is an optical electronic device that generate an intense beam of single color light by pumping photon with more energy through collision with other photon.

Properties of laser:

(i) Monochromativity:

The light emitted from laser is monochromatic.

It means that LASER light has single color.

(ii) Directionality:

LASER emit light that is a highly directed or undirected

Light, because of its directional nature, it can carry energy and data to very long distance for remote diagnosis and communication purpose.

(iii) Collimation:

A laser beam is collimated, meaning it consists of waves travelling parallel to each other in a single direction with very little divergence.

(iv) Coherent:

The light from a laser is said to be coherent which means the wavelength of the laser light, are in phase in space and time.

(v) Intensity:

A laser beam is extremely intense and brighter source compared to other conventional sources such as sun. Brightness is defined as the power emitted per unit area per solid angle.

$g_{f, n_2 > n_1}$

laser will emit

$E_2 \dots N_2$

$E_1 \dots N_1$

$N_1 >> N_2$

Thermal equation:

Boltzmann

A material medium is composed of identical atoms, which are characterized by a specific system of energy levels. These energy levels are common to all atoms in the medium. We can therefore say that certain number of atoms occupy a certain energy level.

- The number of atoms per unit volume that occupy a given energy level is called population of that energy level.

Let E_1 be the ground level and E_2 the excited level. Atoms are distributed ~~equally~~ differently in ~~these~~ two energy levels. Let the population at the levels E_1 and E_2 are N_1 and N_2 ,

$E_2 \dots N_2$

$E_1 \dots N_1$

At thermal equilibrium, the population at the energy levels can be found with the Boltzmann statistic,

$$N_1 = e^{-\frac{E_1}{kT}}$$

$$N_2 = e^{-\frac{E_2}{kT}}$$

- The relative population (N_2/N_1),

$$\left(\frac{N_2}{N_1}\right) = e^{-(E_2 - E_1)/kT}$$

- $\left(\frac{N_2}{N_1}\right)$ is dependent on two factors,

(i) The temp. (T) and

(ii) The energy difference ($E_2 - E_1$)

Effect of Temperature:

In case of hydrogen atom,

$$E_1 = -13.6 \text{ eV}, E_2 = -3.39 \text{ eV}, E_2 - E_1 = 10.21 \text{ eV}$$

① At room temperature, ($T = 300 \text{ K}$)

$$kT = 0.025 \text{ eV}$$

$$\frac{N_2}{N_1} = e^{-\frac{10.21}{0.025}} = e^{-408.4} \approx 0$$

$$\Rightarrow N_2 \approx 0$$

$$N_2 \approx 0$$

② At 6000 K ($T = 6000 \text{ K}$)

$$kT = 0.516 \text{ eV}$$

$$N_1 \gg N_2$$

$$\frac{N_2}{N_1} = e^{-\frac{10.21}{0.516}} = \frac{1}{10^{19}}$$

③ When $T = \infty$

$$\frac{N_2}{N_1} \approx e^0 = 1$$

$$N_2 \approx N_1$$

When, $E_2 - E_1 = 0$

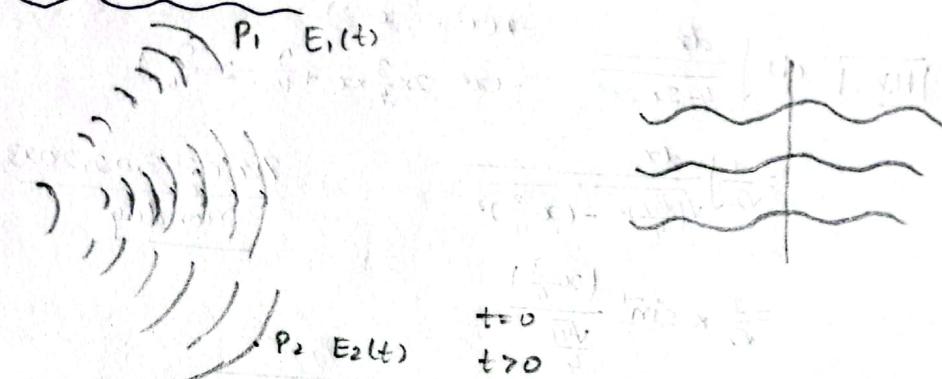
$$\frac{N_2}{N_1} \approx 0 \approx 1$$

$$\Rightarrow N_2 \approx N_1$$

Both this limiting cases indicate, as long as medium is in thermal equilibrium, the population in ~~higher~~ energy level can not exceed the lower energy level.

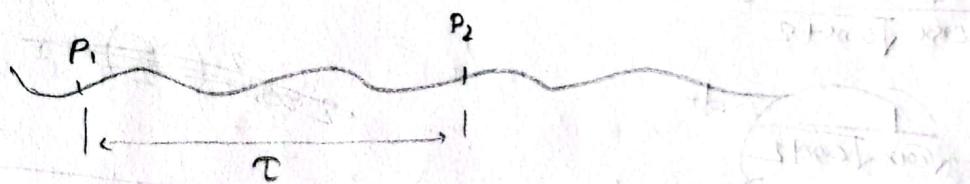
Physics (LASER)

Spatial coherence:



Let us consider two points P_1 and P_2 that, at time $t=0$, lie on the same wavefront of same given em.f. wave and let $E_1(t)$ and $E_2(t)$ be the corresponding electric field at $t=0$. If the phase difference of two electric field is remain zero after $t > 0$ then we will say that there is a perfect coherence between the two points.

If this occur for any two points of e.m. wavefront, we say that wave has perform spatial coherence.



Temporal coherence:

We now consider the electric field of the e.m. wave at a given point p , at times t and $(t + \tau)$. If for a given time delay τ , the phase difference between two field remain the same for any time. then we will say there is a temporal coherence.

Active medium:

Atoms in medium which can produce more than stimulated emissions than spontaneous emission and cause amplification of light is called active center. The rest of the medium acts as host and supports active center. The medium which resting the active center is called the active medium.

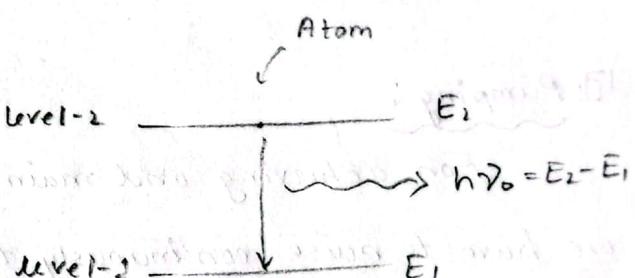
a) Spontaneous emission:

We assume that the atom is initially at level-2. Since $E_2 > E_1$, the atoms tends to be decay to level-1. The corresponding energy difference must be released by the atom. When the energy is delivered in the form of emt wave, the process is called spontaneous emission.

The frequency of the radioactive wave,

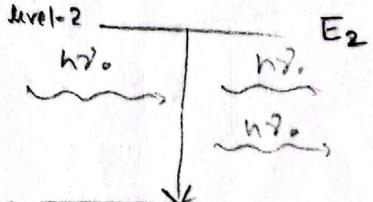
$$\boxed{h\nu_0 = E_2 - E_1}$$

$$\nu_0 = \frac{E_2 - E_1}{h}$$



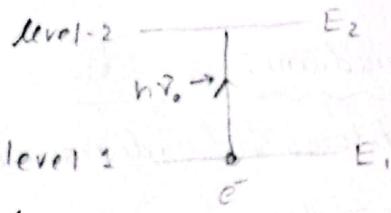
(b) Stimulated Emission:

Let the atom be initially at level-2. An em wave frequency ν_0 which is equal to that of spontaneous emission is incident on the material. The incident wave will force the atom to make a transition from level-2 to level-1. This is characterized by an emission of a photon with energy equal to $(E_2 - E_1)$. This phenomenon is known as Stimulated Emission.



(c)

Absorption:



Let the atom is now initially at level-1, level-1. The atom remains in this level unless some external energy is applied to the material when an incident wave with the energy equal to $E_2 - E_1$, is applied to the material then the atom will be raised to level-2. This process is absorption.

Pumping:

For achieving and maintaining the condition of population, we have to raise continuously the atoms in the lower energy level to upper energy level. It requires energy to be supplied to the system. Pumping is the process of supplying energy to the ~~less~~ lower medium with a view to transfer it into the state of population inversion.

Tuesday

$$N_2 \xrightarrow{\text{Excitation}} \text{Emission} \xrightarrow{\text{Decay}} N_1$$

$$N_1 \xrightarrow{\text{Decay}} E_1$$

$$N_2 \xrightarrow{\text{Decay}} E_2$$

(a) Normal state

Thermal equilibrium

$$(N_1 > N_2)$$

(b) Inverted State

Non equilibrium condition

$$(N_2 > N_1)$$

Population inversion:

We know from thermal equilibrium condition,

$$\frac{N_2}{N_1} = e^{-\frac{(E_2 - E_1)}{kT}}$$

$$\frac{N_2}{N_1} = e^{\frac{R(E_2 - E_1)}{kT}}$$

We may somehow enhance the number of atoms in the excited level than the lower such that the population ratio ($\frac{N_2}{N_1}$) momentarily increase without change in temperature.

* Population inversion is the non-equilibrium condition of the material in which population of the upper energy level N_2 unmomentarily ~~exceeds~~ exceeds the population lower energy level N_1 , $\boxed{N_2 > N_1}$

Three level laser:

In a three level laser the atoms are in some way raised from ground level to level 3 if the material is such that after an atom has been raised to level 3, it decays rapidly to level 2, then the population inversion can be obtained between level 2 to 1, in this case $N_3 \approx 0$ due to fast decays

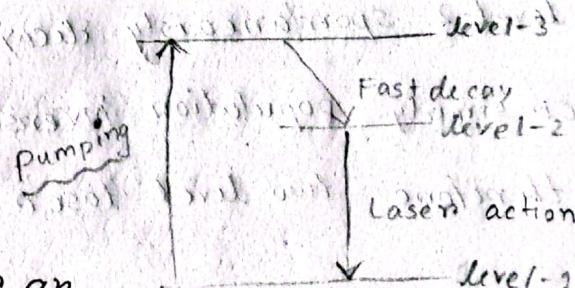


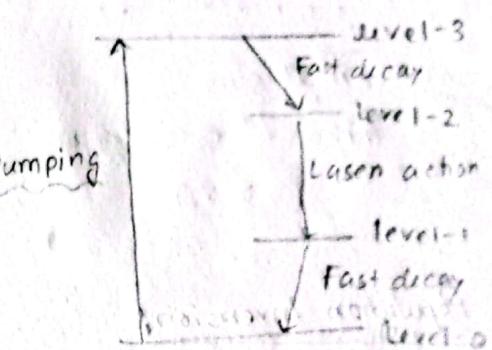
Fig: 3 level laser

For $N_2 > N_1$, $(N_2 - N_1)$ has a definite value, there is a minimum value of $(N_2 - N_1)$ for which we get laser action between level-1 and level-2.

This minimum value is known as critical inversion denoted by N_1 .

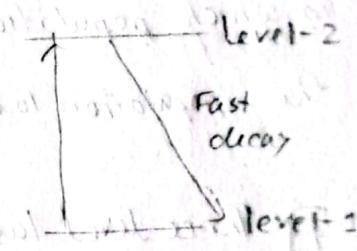
Four level laser:

In a four level laser atoms are raised from ground level-0 to level-3, if the atom then rapidly decay to level-2, a population inversion obtain from level-2 to level 1. Once oscillation starts in four level laser, the atom will then be transferred to level-1, though stimulated emission. For continuous wave operation it is therefore necessary that the transition 1 to 0 should be very fast.



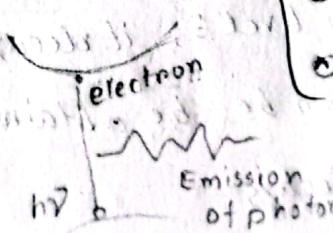
Two level laser: (Reason why it is not possible)

In two level laser population inversion is not possible after pumping process, the population from level-2 spontaneously decays to level-1. Critical inversion or simply population inversion is not possible, in two level system, therefore two level laser is possible.

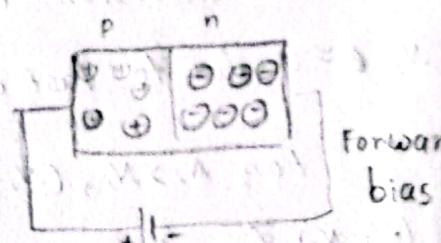


Semiconductor diode laser:

It is specifically fabricated p-n junction diode. This diode emits laser light when it is forward bias.



- ④ Semiconductor laser
- ④ Ruby laser (solid state laser)
- ④ He-Ne (gas laser) laser



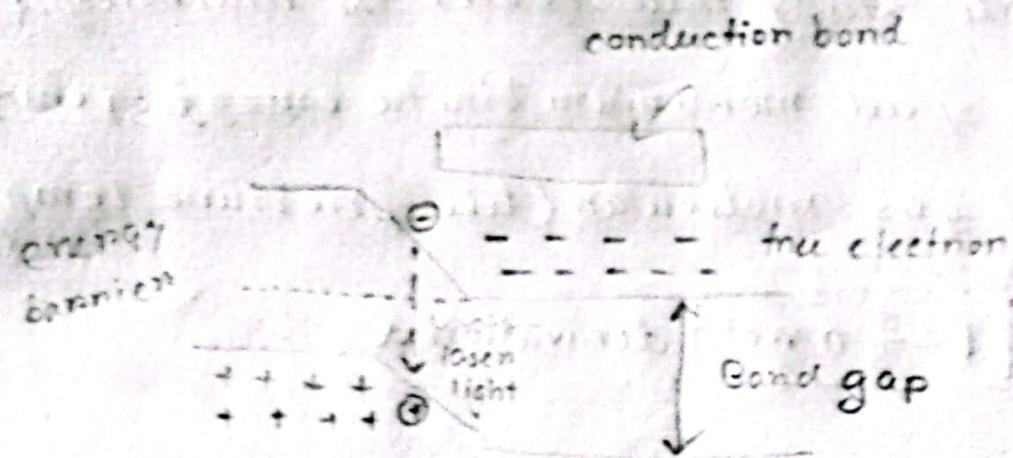


Fig: Solid state laser.

Characteristics:

1. It is a solid state laser.
2. A p-n junction made from single crystal of gallium arsenide is used in active medium.
3. Pumping method: The direct conversion by method is used in pumping action.
4. The power out of this level is ~~more than~~ 1 mW.

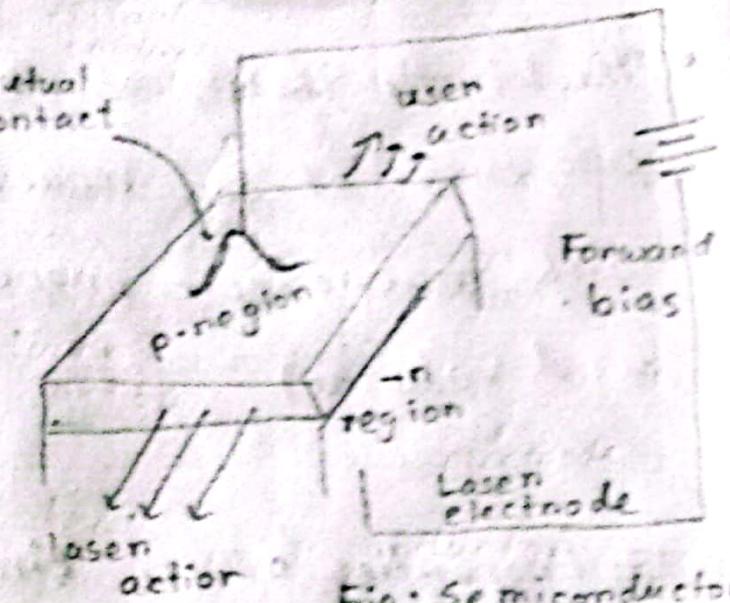


Fig: Semiconductor laser

$$E_g = h\nu = h\frac{c}{\lambda}$$

$$\lambda = \frac{hc}{E_g}$$

Date: 14.03.2023

Tuesday

Kinetic Theory of gases:

- volume, Pressure, Temperature

Volume → The volume is ~~the~~ result of the freedom ^a of the atoms have to spread throughout the container.

Pressure → The pressure is a result of the emission of the atom with the container wall.

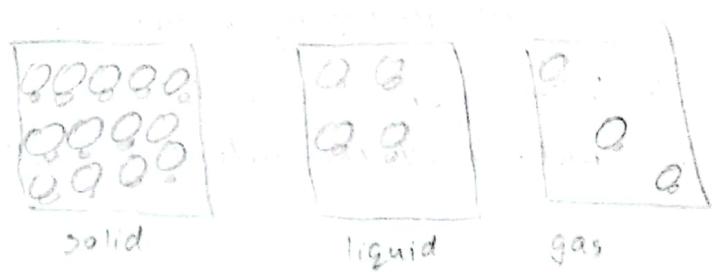
Temperature → The temperature has to do with the kinetic energy of the atoms.

- The kinetic theory of gases is the study that relates the macroscopic properties of gas molecule (like speed, momentum, kinetic ~~energy~~) energies with macroscopic properties of gas molecular (like pressure, temp and volume).

$$VP = \frac{1}{3} mn\bar{c}^2 \quad | \text{ derivation } \times$$

Properties of the kinetic theory of gases:

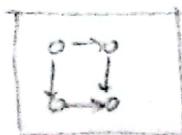
- The gas is composed of small indivisible particles called molecules. The properties of the individual molecules are the same as that of the gas as a whole.
- The distance between the molecules is large as compared to that of solid and liquid hence force of intermolecular attraction is negligible.



- Any infinitely small volume of a gas contains many molecules under standard conditions of temperature and pressure there are nearly 3×10^{25} molecules in a cubic meter.
- The molecules continuously move about in random direction. Every directions is equally probable.



- The size of the molecules is much less than the average distance between them.



⑥ The molecules of the a gas exert no forces on each other except when they collide i.e., there are no intermolecular forces. In the absence of external force the molecules will move in uniform rectilinear motion and their direction will change only by collision with walls of the container or with other molecules

⑦ Collision between molecules and with the walls are perfectly elastic i.e., there is no loss in their kinetic energy in a collision.



⑧ The molecules are distributed uniformly in the absence of an external field and there is no preferential direction of motion.



⑨ The molecules move with all speeds from 0 to infinity. This means that all molecules do not have the same speed.

⑩ The time of impact is negligible in comparison with to the time taken to traverse the free path.

Brownian motion:

Light suspended particles in a gas or in low viscous fluid are found to move continuously in a random manner in short zigzag path such a motion if suspended particle is known as Brownian Motion.

Characteristics:

1. Brownian motion is continuous spontaneous, perpetual and random
2. With the rise of temp, the motion becomes more vigorous.
3. The smaller particles are agitated & move.
4. The motion depends on the chemical nature of the particle.
5. At the same temp., particles of the same size are equally agitated.
6. The less the viscosity of the fluid, the more vigorous the motion.
7. The motion is independent of the movement of the containing vessel.
8. Brownian motion is not affected by electric and magnetic field.

Degree of freedom:

The degree of freedom for a dynamic system is the number direction in which a particle can move freely or the minimum number of independent coordinates needed to specify the position and configuration of a thermodynamic system in space. It is denoted f.

Total number of freedom,

$$f = f_t + f_r + f_v$$

suppose, we have N number of gas molecules, degree of freedom.

$$f = 3N$$

But if the system has q number of constraints, then degrees of freedom, $f = 3n - q$

Monatomic gas : (He, Ar, ...)

$$f_{trans} = 3$$

$$\downarrow \text{transition} \quad f_{rot/notation} = 0 \quad \begin{array}{l} \text{No bond,} \\ \text{no vibration} \end{array}$$

$$f_{vib} = 0$$

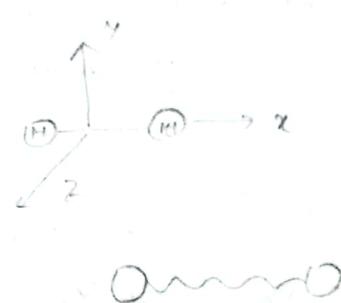
$$f_{total} = 3$$

Diatom gas (H_2, N_2)

$$f_{trans} = 3$$

$$f_{not} = 2 \quad \text{notation of freedom}$$

$$f_{vib} = 1 \quad \text{vibration of freedom}$$



Below 100K $\rightarrow f_r \rightarrow$ no notation

100K < T < 1000K $\rightarrow f_{trans} + f_{not}$

T > 1000K $\rightarrow f_{trans} + f_{not} + f_{vib}$

Triatomic gas

1. Linear triatomic (CO_2)

$$f_{trans} = 3$$

$$f_{not} = 2$$

$$f_{vib} = (3N-5)$$

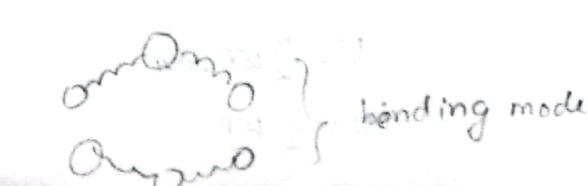
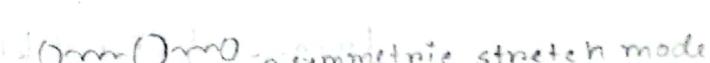
$$= 4$$

$$f_{total} = 4+2+3$$

$$= 9$$



symmetric stretch mode



2. Non-linear triatomic gas (H_2O)

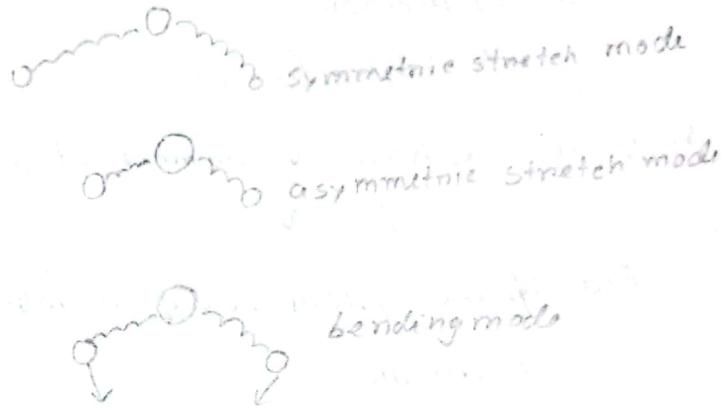
$$f_{\text{trans}} = 3$$

$$f_{\text{rot}} = 3$$

$$f_{\text{vib}} = 3N - 6 = 3$$

$$f_{\text{total}} = 3 + 3 + 3$$

$$= 9$$



Principle of Equipartition of Energy:

According to this law, the total energy of a dynamic system in thermal equilibrium is shared equally by all its degrees of freedom, the energy associated per molecules per degree of freedom being a constant equal to $\frac{1}{2}kT$.

According to kinetic theory,

$$\frac{1}{2}mv^2 = \frac{3}{2}kT$$

$$\bar{v}^2 = \bar{v_x}^2 + \bar{v_y}^2 + \bar{v_z}^2$$

$$\Rightarrow \bar{v_x}^2 = \bar{v_y}^2 = \bar{v_z}^2$$

$$\text{So, } \frac{1}{2}m\bar{v_x}^2 = \frac{1}{2}m\bar{v_y}^2 = \frac{1}{2}m\bar{v_z}^2$$

$$\Rightarrow \frac{1}{2}m(\bar{v_x}^2 + \bar{v_y}^2 + \bar{v_z}^2) = \frac{1}{2}m\bar{v}^2$$

$$\Rightarrow \frac{1}{2}m(3\bar{v_x}^2) = \frac{1}{2}m\bar{v}^2$$

$$\Rightarrow \frac{1}{2}m(3\bar{v_x}^2) = \frac{3}{2}kT$$

$$\Rightarrow \boxed{\frac{1}{2}m\bar{v_x}^2 = \frac{1}{2}kT}$$

$$\frac{1}{2}m\bar{v_y}^2 = \frac{1}{2}kT$$

$$\frac{1}{2}m\bar{v_z}^2 = \frac{1}{2}kT$$

Ratio of specific heat:

1. monoatomic gas-

$$E_{\text{trans}} = \frac{1}{2}m\bar{v_x}^2 + \frac{1}{2}m\bar{v_y}^2 + \frac{1}{2}m\bar{v_z}^2$$

$$E_{\text{trans}} = \frac{1}{2}kT + \frac{1}{2}kT + \frac{1}{2}kT = \frac{3}{2}kT$$

$$U = \frac{3}{2}kT \cdot N_A$$

$$U = \frac{3}{2}RT$$

$$C_V = \frac{dU}{dT} = \frac{3}{2}R$$

$$C_P = C_V + \frac{3}{2}R \\ = \frac{5}{2}R$$

$$\gamma = \frac{C_P}{C_V} = \frac{5R}{3R/2} = 1.67$$

2. Diatomic gas:

① At normal temp ,

$$E_T = \frac{5}{2} kT$$

$$V = \frac{5}{2} RT$$

$$C_V = \frac{\partial V}{\partial T} = \frac{5}{2} R$$

$$C_P = C_V + R = \frac{5}{2} R + R = \frac{7}{2} R$$

$$\gamma = \frac{C_P}{C_V} = \frac{7/2 R}{5/2 R} = 1.40$$

② High temp:

3. Triatomic gas:

non linear , $E_T = \frac{6}{2} kT$, $V = \frac{6}{2} RT$

$$C_V = \frac{\partial V}{\partial T} = \frac{6}{2} R = 3R \quad C_P = C_V + R = 4R \quad \gamma = \frac{4}{3} = 1.33$$

Date : 27.03.2023
Tuesday

Distribution of velocity:

Determination of num of molecules having a certain velocity but range of velocities infinite, num of molecules finite.

$$f(c) = 4\pi \left(\frac{m}{2\pi kT} \right)^{3/2} c^2 e^{-\frac{mc^2}{2kT}}$$

$$dN \rightarrow c \quad c+dc$$

Let dN = num of molecules having velocities within range c and $(c+dc)$

$$dN \propto dc$$

$$\Rightarrow dN = adc \quad (1)$$

a is proportionately constant and a is a function of c .

$$a = f(c) \quad (2)$$

$$\text{But, } dN \propto N$$

Hence $f(c)$ is a distribution function

We can write,

$$dN = N f(c) dc \quad (3)$$

Considering $dc = 2$, $f(c) = \frac{dN}{N}$,

Fraction of molecule whose velocities are within unit interval near c ,

$$f(c) = \frac{dN}{N} \quad (4)$$

$\frac{dN}{N}$ is the probability of any molecule in unit volume having velocity that is within unit interval near velocity c .

(u, v, w) describes a velocity point.

Probability of molecules having velocities u and $u+du$, $f(u)du$

" " " " " " " " v and $v+dv$, $f(v)dv$

" " " " " " " " w and $w+dw$, $f(w)dw$

$$dN = N f(u) f(v) f(w) du dv dw$$

$$\Rightarrow dN = N \varphi(c^2) du dv dw$$

$$\Rightarrow \frac{dN}{du dv dw} = N f(u) f(v) f(w) = N \varphi(c^2)$$

$\frac{dN}{du dv dw}$ is ρ which is the num density in velocity space.

$$\rho = N f(u) f(v) f(w) = N \varphi(c^2)$$

$$\Rightarrow d\rho = d[N f(u) f(v) f(w)] = d[N \varphi(c^2)] \quad (5)$$

For a fixed c , RHS = 0.

$$d[N f(u) f(v) f(w)] = 0$$

$$\Rightarrow f'(u) f(v) f(w) du + f(u) f'(v) f(w) dv + f(u) f(v) f'(w) dw = 0$$

Dividing by $[f(u) f(v) f(w)]$ we get,

$$\Rightarrow \frac{f'(u)}{f(u)} du + \frac{f'(v)}{f(v)} dv + \frac{f'(w)}{f(w)} dw = 0 \quad (6)$$

We know,

$$c^2 = u^2 + v^2 + w^2$$

$$\Rightarrow 2udu + 2vdv + 2wdw = 0$$

$$\Rightarrow udu + vdv + wdw = 0$$

$$\Rightarrow \beta udu + \beta vdv + \beta wdw = 0$$

—(7)

where, β is undetermined multiplier of language

• Adding equation (6) & (7)

$$\left(\frac{f'(u)}{f(u)} + \beta u \right) du + \left(\frac{f'(v)}{f(v)} + \beta v \right) dv + \left(\frac{f'(w)}{f(w)} + \beta w \right) dw = 0$$

$$f(x, y, z) + \lambda \varphi(x, y, z)$$

$$F(x, y, z) = f(x, y, z) + \varphi(x, y, z)$$

$$\frac{f'(u)}{f(u)} = -\beta u \quad \rightarrow \text{Integrating}, \quad \ln f(u) = -\frac{\beta u^2}{2} + \ln a$$

[a is constant]

$$\frac{f'(v)}{f(v)} = -\beta v$$

$$b = \frac{\beta}{2}$$

$$\frac{f'(w)}{f(w)} = -\beta w$$

$$\begin{aligned} \Rightarrow f'(u) &= a e^{-\frac{\beta u^2}{2}} \\ &= a e^{-bu^2} \end{aligned}$$

Similarly, $f(v) = a e^{-bv^2}$

$$f(w) = a e^{-bw^2}$$

$$f(u)f(v)f(w) = a^3 e^{-b(u^2+v^2+w^2)}$$

$$dN = N f(u) f(v) f(w) du dv dw$$

$$= N a^3 e^{-b(u^2+v^2+w^2)} du dv dw$$

$$dN = N a^3 e^{-b(c^2)} du dv dw$$

$$dN = N a^3 e^{-b(c^2)} du dv dw \quad (9)$$

Finding the volume in velocity space,

$$du dv dw = c^2 d\cos\theta d\phi \sin\phi \quad (10)$$

$$(x, y, z) \rightarrow (v, \theta, t)$$

$$dx dy dz \rightarrow n^2 dn \sin\theta d\theta d\phi$$

$$du dv dw \rightarrow c^2 d\cos\theta d\phi \sin\phi \quad (10)$$

$$(dN) = N a^3 e^{-b(c^2)} c^2 \sin\theta d\phi \sin\phi \quad (11)$$

$$\sum dN = N a^3 \int_0^\infty e^{-b(c^2)} c^2 dc \int_0^\pi \sin\theta d\theta \int_0^{2\pi} d\phi \quad (11)$$

$$\sum dN = 4\pi N a^3 \int_0^\infty e^{-b(c^2)} c^2 dc$$

$$\Rightarrow \frac{\sum dN}{N} = 4\pi a^3 \int_0^\infty e^{-b(c^2)} c^2 dc$$

$$\Rightarrow \int f(c) dc = 4\pi a^3 \int_0^\infty e^{-b(c^2)} c^2 dc = 1$$

$$\Rightarrow 4\pi a^3 \int_0^\infty e^{-b(c^2)} c^2 dc = 1$$

standard integral,

$$\int_0^\infty e^{-\frac{1}{2}c^2} c^2 dc = \frac{1}{4}\sqrt{\frac{\pi}{2}}$$

$$\Rightarrow \int_0^\infty c^{-b(c^2)} c^2 dc = \frac{1}{4}\sqrt{\frac{\pi}{b}}$$

$$\Rightarrow 4\pi a^3 \frac{1}{4}\sqrt{\frac{\pi}{b}} = 1$$

$$\Rightarrow \pi^{3/2} a^3 \sqrt{\left(\frac{2}{\beta}\right)^3} = 1$$

$$\Rightarrow \pi^{3/2} a^3 \cdot 2^{3/2} \beta^{-3/2} = 1$$

$$\Rightarrow a^3 = \frac{\beta^{3/2}}{(2\pi)^{3/2}}$$

$$\Rightarrow a = \sqrt{\frac{\beta}{2\pi}} \quad [b = \frac{\beta}{2}]$$

Expectation value of mean square velocity,

$$\langle c^2 \rangle = \frac{N_1 c_1^2 + N_2 c_2^2 + N_3 c_3^2 + \dots}{N_1 + N_2 + N_3 + \dots}$$

$$= \frac{1}{N} \int_0^\infty c^2 dN$$

$$= \int_0^\infty c^2 f(c) dc$$

$$= \int_0^\infty c^2 \cdot 4\pi a^3 e^{-\frac{\beta}{2}c^2} c^2 dc$$

$$= 4\pi a^3 \int_0^\infty e^{-\frac{\beta}{2}c^2} c^4 dc$$

$$\int_0^\infty e^{-5x^2} x^4 dx = \frac{3}{8} \sqrt{\frac{\pi}{5^5}}$$

$$\int_0^{\infty} e^{-sx^2} x^2 dx = \frac{1}{4} \sqrt{\frac{n}{s^3}}$$

$$= 4\pi a^3 \frac{3}{8} \sqrt{\frac{\pi}{(\beta_F)^5}}$$

$$\langle c^2 \rangle = 4\pi \left(\sqrt{\frac{\beta}{2\pi}} \right)^3 \frac{3}{8} \sqrt{\frac{\pi}{(\beta/2)^5}} = \frac{\beta}{3}$$

We know,

$$\frac{1}{2} m \langle c^2 \rangle = \frac{3}{2} kT$$

$$\Rightarrow \frac{1}{2} m \left(\frac{\beta}{3} \right) = \frac{3}{2} k T$$

$$\Rightarrow \beta = \frac{m}{kT}$$

$$\Rightarrow a = \sqrt{\frac{B}{2\pi}} = \sqrt{\frac{m}{2\pi kT}}$$

$$f(c)dc = 4\pi a^3 e^{-bc^2} c^2 dc$$

$$f(c) = 4\pi \left(\frac{mc^2}{2\pi kT} \right)^{3/2} c^2 e^{-\frac{mc^2}{2kT}}$$

$$dN = 4\pi N \left(\frac{m}{2\pi kT} \right)^{3/2} c^2 e^{-\frac{mc^2}{2kT}} dc$$

$$= \frac{1}{N} \int_0^{\infty} 4\pi N \left(\frac{m}{2\pi kT} \right)^3 c^3 e^{-\frac{mc^2}{2kT}}$$

$$= \frac{4\pi}{3} \left(\frac{m}{2\pi kT} \right)^3 \int_0^{\infty} e^{-\frac{mc^2}{2kT}} c^3 dc$$

$$= \frac{1}{2} \pi \left(\frac{m}{2\pi kT} \right)^{3/2} \frac{1}{2 \left(\frac{m}{2kT} \right)^2}$$

$$= \cancel{m} \cancel{\pi} 2x \frac{m^{3/2} \cdot \pi^{3/2} \cdot L^2 T^2}{2^{3/2} \cdot \pi^{3/2} \cdot L^{3/2} T^{3/2}}$$

$$= \frac{2 \cdot 6 \sqrt{2} \sqrt{3} \cdot y}{x_2 \sqrt{2}} = \sqrt{\frac{8 k T}{m x}}$$

Root mean square velocity

$$C_{rms}^2 = \frac{N_1 C_1^2 + N_2 C_2^2 + N_3 C_3^2 + \dots}{N_1 + N_2 + N_3 + \dots}$$

$$c^2_{rms} = \frac{1}{N} \int^{\infty} c^2 dN$$

$$= \frac{1}{N} \int_0^{\infty} 4\pi N \left(\frac{m}{2\pi kT} \right)^{3/2} c^2 e^{-\frac{mc^2}{2kT}} c^2 dc$$

Average velocity / mean velocity

$$\text{mean} = \frac{N_1 C_1 + N_2 C_2 + \dots}{N_1 + N_2 + \dots}$$

$$= \frac{1}{N} \int_0^\infty c dN$$

$$\int_0^{\infty} e^{-ax^2} x^2 dx = \frac{1}{2a^3}$$

$$= \int_{-\infty}^{\infty} e^{-\frac{m\omega^2}{2kT}} \cos \omega t$$

$$L = \frac{1}{2 \left(\frac{m}{2kT} \right)^2}$$

$$c_{rms} = \sqrt{\frac{3kT}{m}}$$

\therefore Most Probable velocity, $\frac{dN}{dc} = F$

$$\Rightarrow F = 4\pi N \left(\frac{m}{2\pi kT}\right)^{3/2} c^2 e^{-\frac{mc^2}{2kT}}$$

$$\Rightarrow \frac{dF}{dc} = 4\pi N \left(\frac{m}{2\pi kT}\right)^{3/2} \left\{ c^2 e^{-\frac{mc^2}{2kT}} \left(-\frac{2m}{2kT}\right) + e^{-\frac{mc^2}{2kT}} 2c \right\}$$

$$\Rightarrow 0 = c^3 e^{-\frac{mc^2}{2kT}} \left(-\frac{m}{kT}\right) + e^{-\frac{mc^2}{2kT}} 2c$$

$$\Rightarrow c \left\{ c^2 \left(-\frac{m}{kT}\right) + 2 \right\} = 0$$

$$c \neq 0, c_m = \sqrt{\frac{2kT}{m}}$$

constant