

# Solids

7.11.2k

## Energy band formation in a solid

From Bohr's model of hydrogen like atoms we have seen that the spectra of this type of atoms consists of several sharp lines which ~~consist~~ <sup>characterise</sup> the atom. From quantum mechanics it has been proved that each such energy level will split into two such lines when two identical atoms are brought closer. This is also experimentally found to be correct. When 3 such atoms are brought closer, each line will split into 3 such lines. This type of splitting continues as more and more atoms are brought closer.



Energy levels due to single ~~electron~~ <sup>atom</sup>

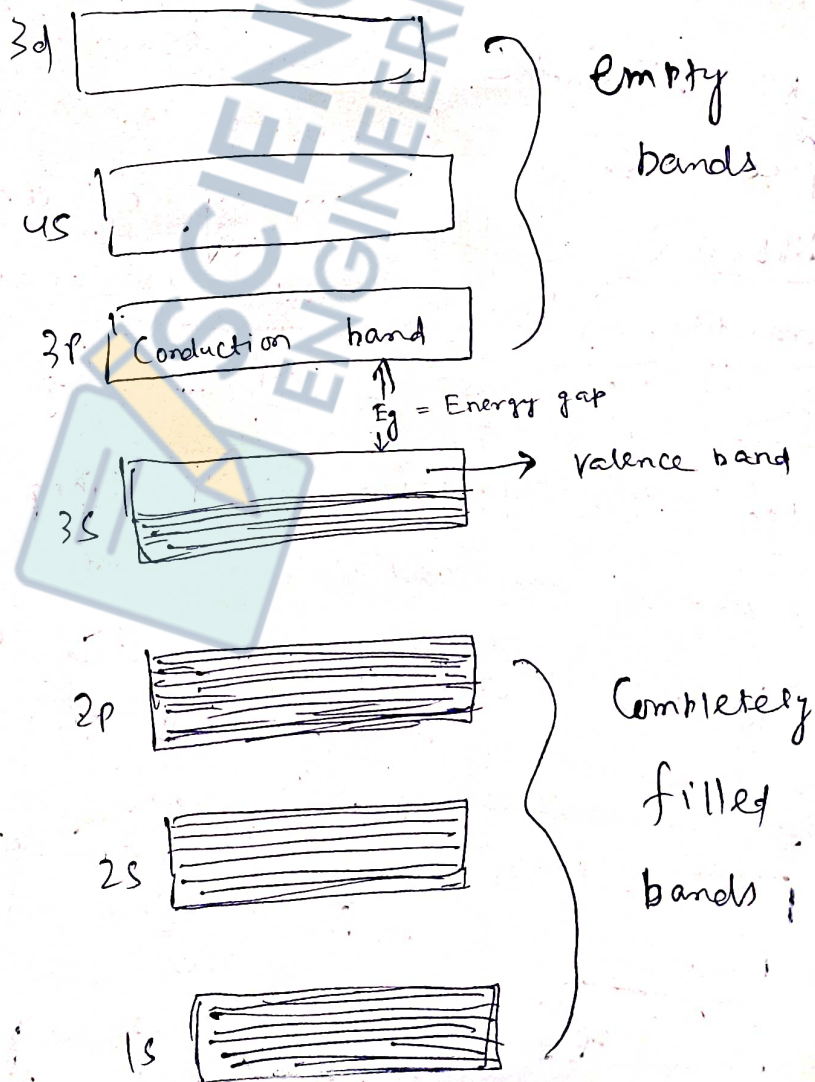
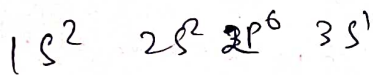
Energy level due to two atoms brought nearer.

Energy band formed in a solid

In the case of a solid, the atoms are very close to one another. Hence each one will split into large number of lines, being very close to one another. These group of lines will appear as a band. Therefore the solids are characterized by energy bands, which are separated from one another. Depending on the number of outer most electron & temperature the bands are occupied.

Ex → Solid He.

Electronic Configuration of a He atom is

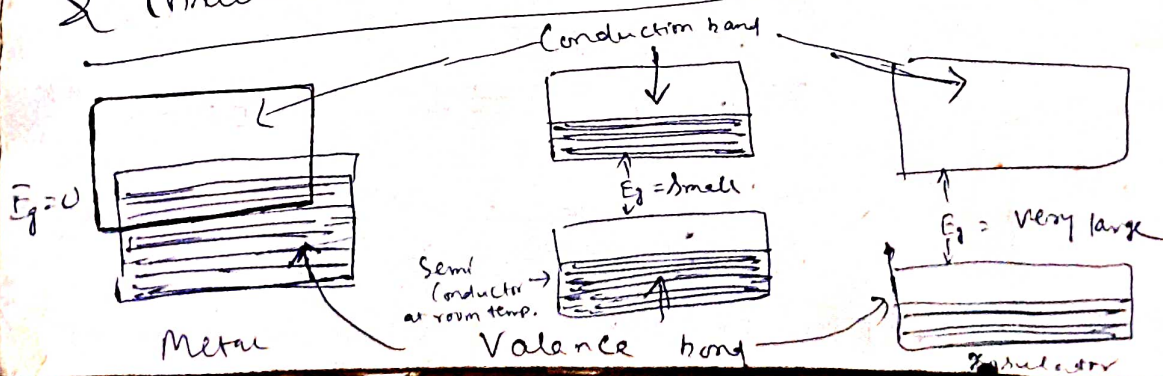


Energy Bands of solid sodium at very low temp.

At very low temperature the lower bands like 1s, 2s, 2p are completely filled, where as the 3s band is half filled. The other higher bands like 3p, 4s & 3d are empty. The highest filled band (may be partly or completely filled) is called the valency band.

The empty band that lies just above the valency band is called conduction band. When the temperature is increased, it is probable that some of the electrons of the valency band may jump over into the conduction band. due to thermal energy ( $\frac{1}{2}KT$ ). Then the electrons practically become free and conductivity increases. The difference of energy between the conduction & valency band is called energy gap or forbidden gap ( $E_g$ ). No electron can possess energy lying in between conduction band and valency band.

Classification of solids into metals, semiconductors & insulator on the basis of band theory



## Types of solids

### Metals (Good Conductor)

In the case of metals the outermost electrons of each atom gets detached from the atom form a sea of electrons. This is due to ~~the~~ overlapping of the valance band on the conduction band. The energy gap is zero, and the electrons of the valance band also belong to the conduction band. i.e why metals are good conductors even ~~at~~ <sup>at</sup> low temperatures. The energy gap is zero which ~~is~~ differentiates metals from the semiconductors & insulators. The highest energy of an electron at low ~~high~~ temp is called fermi energy ( $E_f$ ).

### Semi-conductor →

Germanium & Si are good semiconductors. The energy gap is found to be small, of the order of a few electron volts. Even with thermal energy ( $\frac{1}{2}kT$ ) per degree of freedom it is possible on the part of electrons to go over from the valance to the conduction band even at room temp. That is why semiconductors behave as conductor at high temp. With ~~the~~ the rise of temp, the conductivity of semiconductor increases. This character of the semiconductors distinguishes them from conductors.

Where conductivity decreases with the increase of temperature. In the case of insulators, the energy gap is found to be very large as a result of which, it is not possible for electrons of the valence band to move into the conduction band so that conductivity can increase. Therefore, insulators remain as non-conductors even at high temperature.

For Si  $E_g = 1.1 \text{ eV}$ , For Ge  $E_g = 0.7 \text{ eV}$

### Types of semiconductors →

Intrinsic  
(Pure)

Extrinsic  
(Impure)

n-type

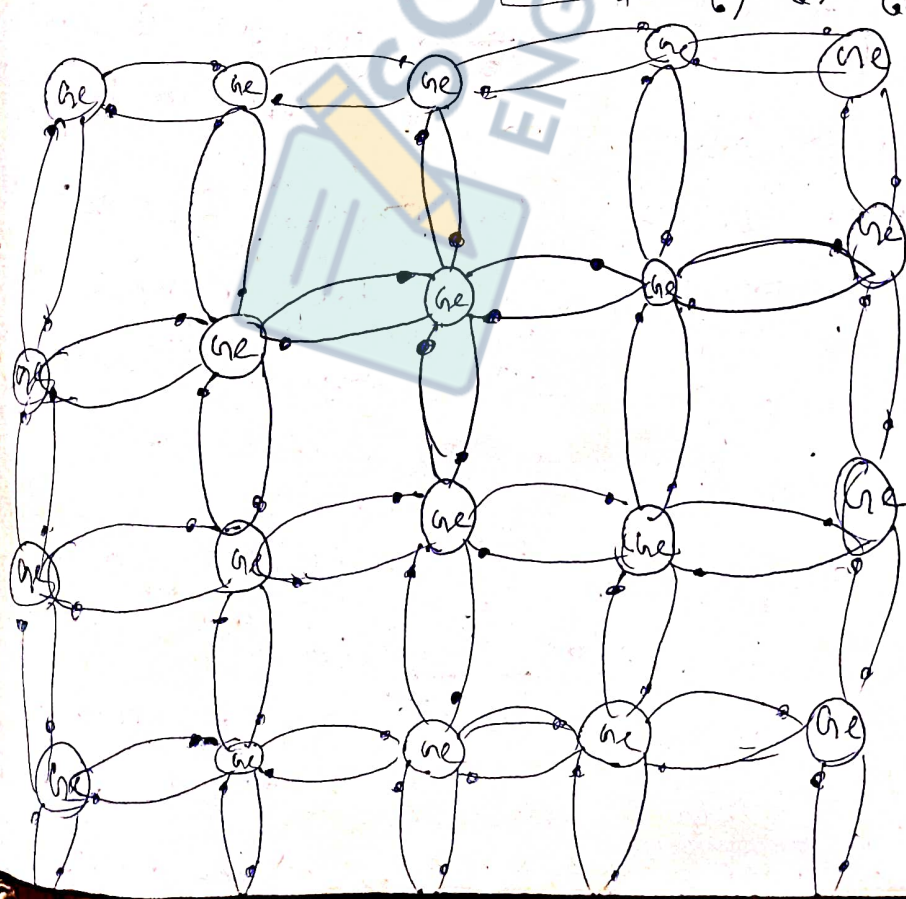
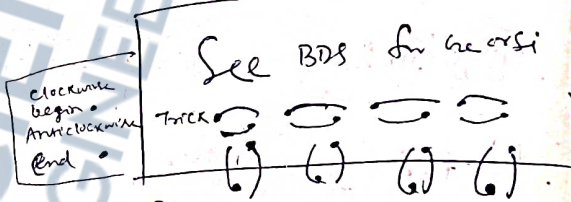
p-type

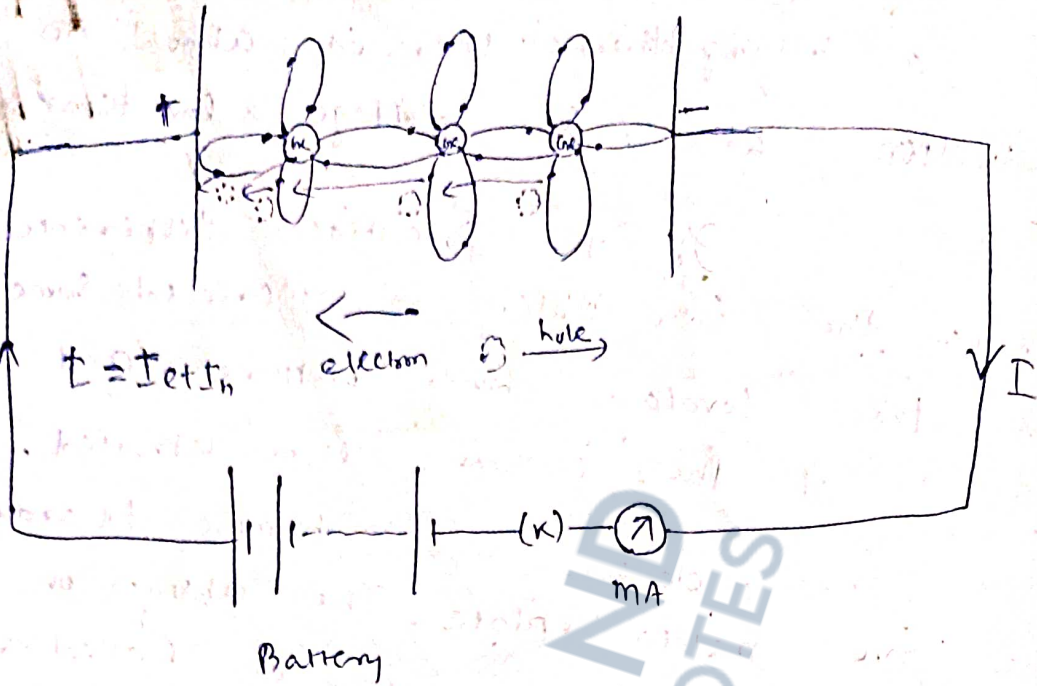
Pure form of Germanium & Silicon are called intrinsic semiconductors. The electronic configuration of Ge & Si are similar to that of C-atom, i.e. there are 4 outermost electrons and there is a deficit of 4 electrons to satisfy the octet (nearest inert gas structure).

Hence each atom of Ge or Si forms 4 covalent bonds with the nearest 4 similar atoms. When such a material is kept in between

2 plates, connected to the opposite terminals of a battery, then there is almost no conduction at low voltage & low temp.

If the potential difference between the two plates is increased, some of the covalent bonds rupture and some of the electrons will be uprooted. These free electrons start to move towards the +ve metallic plate. The absence of an electron in a covalent bond is called a hole - which appears to move towards the -ve plate. Therefore, a hole is regarded as a freely charged particle. Equal number of electrons & holes take part in the conduction process.





$I_e = I_h$  because no. of free electrons = no. of holes

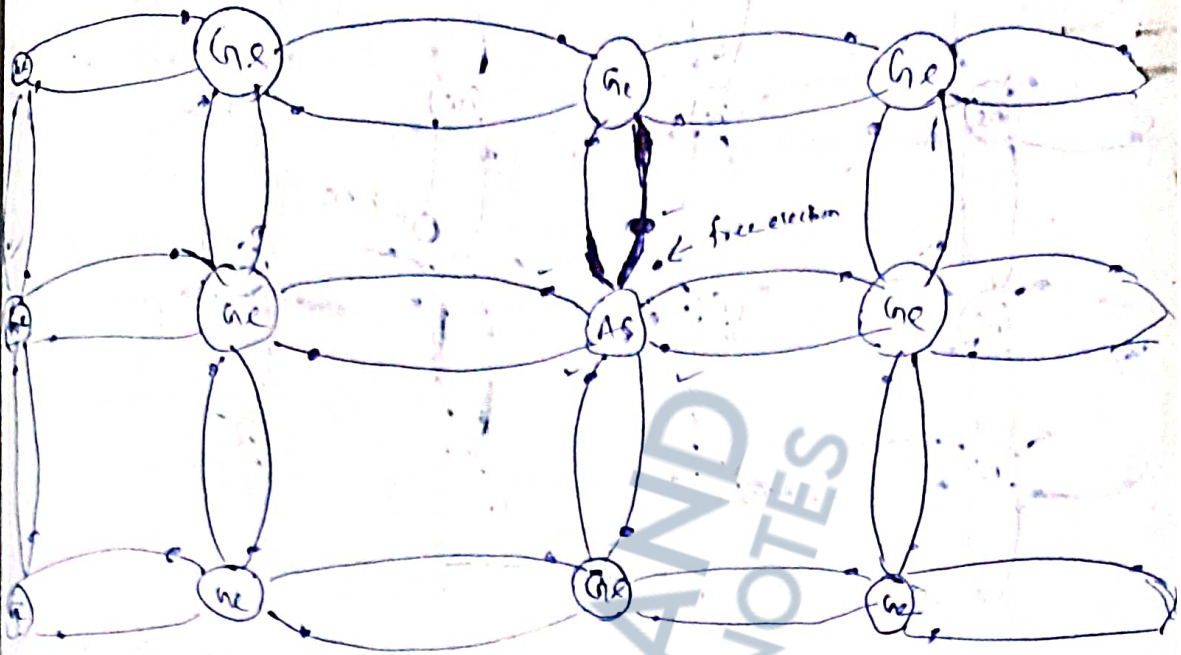
$I_e =$  electron current

$I_h =$  hole current

### Extrinsic Semiconductor →

Naturally occurring semiconductors like Ge, Si are found to be ~~given~~ giving out few electrons & holes which take part in the conduction process (when external field is applied). To increase the number of charge carriers, a procedure is followed called doping.

The metal is heated to a high temp when it just starts to melt. Then impurity atoms in powdered form is sprinkled on it. Adsorption takes place & some of the impurities atoms enter into the semiconductor. The

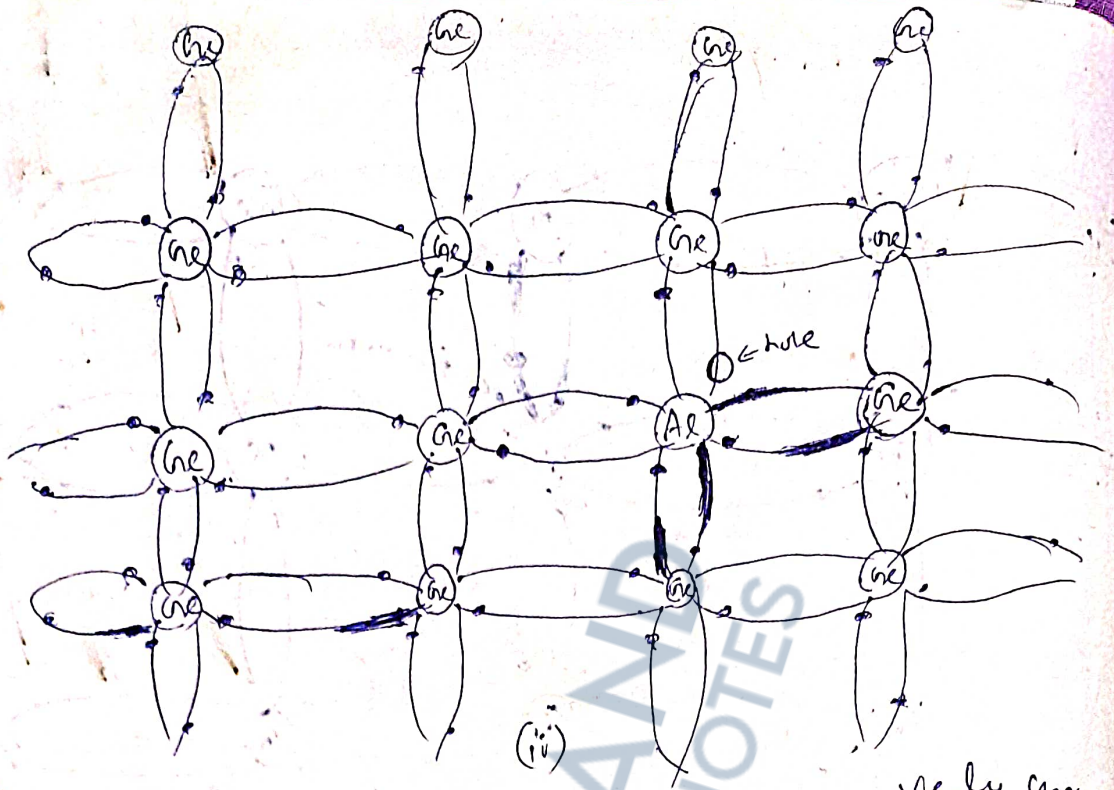


doping is done in such a manner that  
 the number of impurity atoms will be found  
 One in every 10<sup>6</sup> atom of Ge or Si.  
 Depending on the type of impurity atoms  
 extrinsic semiconductors are further classified  
 into (i) n-type (ii) p-type.

### n type extrinsic semiconductor

If the impurity atom be pentavalent in nature  
 like As, Sb, P, then there are 5 electrons  
 in the outermost orbit of an impurity atom. Out  
 of these 5 electrons 4 will be utilized to  
 satisfy the covalent bonds with the 4  
 neighbouring semiconducting atoms. One electron is  
 left unattached which behaves as a free  
 electron. Thus, when covalent bonds between  
 equal number of electron and holes are obtained as  
 before. When the free electron add to this,  
 the number of electrons become greater than the





(ii) Number of holes. Thus the  $-ve$  by energy electrons dominate. Hence this type of impure semiconductor are called  $n$ -type ( $-ve$  axis). The impure atom is called donor.

(ii) P-type extrinsic semiconductor

Trivalent impure atoms be introduced into pure crystals of Ge & Si on the proportion of one impure atom in  $10^6$  pure atoms, then all the 4 covalent bonds with the four neighbouring semiconductor atoms can not be satisfied. There is deficit of one electron in one of the bonds. This absence of electron in a bond behaves as a hole. Thus, no of holes as charge carrier increases than the number of electrons. Hence such an impure semiconductor is called p-type. The impure atom is called acceptor.

The acceptor atom may be Gallium, Indium, Boron, Aluminium etc

## Source of electrons

Free electrons can be obtained by any of the following 4 methods.

### (i) Thermionic emission

When a metal is heated to a high temp, it is found to emit electrons. These electrons are called thermions & the current arising due to these thermions is called thermionic current.

Richardson & Dushman gave a formula for the thermionic current obtained from the heating of some metal.

$$I = AT^2 e^{-\frac{b}{kT}}$$

where  $A$  &  $b$  are constants which differ from metal to metal.

$T$  = Absolute temp. of the metal.

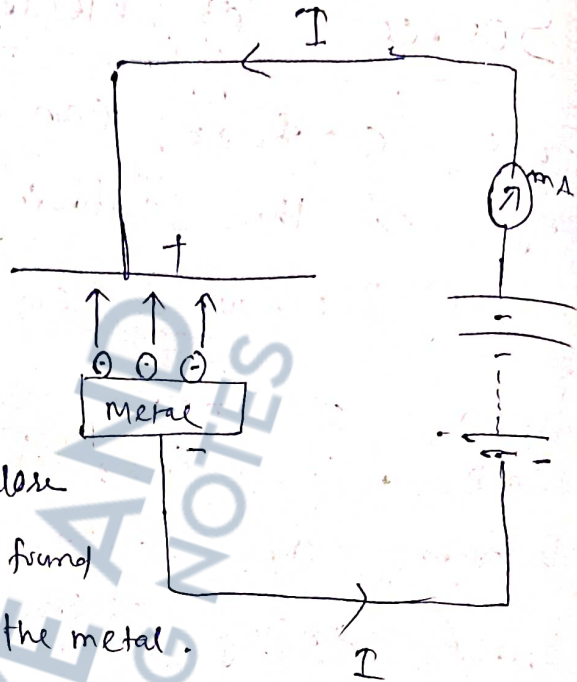
$b$  is found to be equal to the work function ( $W_0$ ) of the metal.

$k$  = Boltzmann's Constant.

From this formula we see that 2-factors  $T^2$  &  $e^{-\frac{b}{kT}}$  determine the current.

## (2) Cold emission or field emission

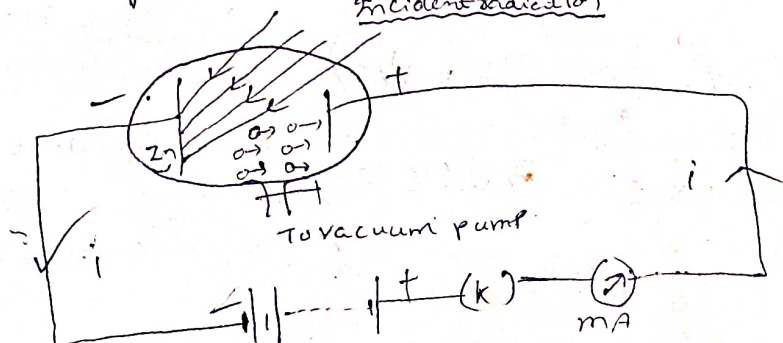
When the metal is connected to the -ve terminal of the battery & another plate is connected to the +ve terminal of the same battery and held over it, quite close to it, electrons are found to be emitted by the metal.



In this case no heating has been done, yet the strong attraction of the +ve plate for the electrons produces free electrons.

## (3) Photo electric emission

When ~~an~~ light (visible, U.V, X-rays) etc be incident on certain metals like Na, K, Zn, Cs etc electrons are found to be emitted by these metals. These electrons are called photoelectrons & the current arising out of them is called photo current.



## (4) Secondary emission

When high speed electrons are made to strike a metallic plate, more electrons are found to be ejected from the surface of the metal. This happens because of the fact that the incident energy imparted to the electrons such that the energy exceeds the work function. Then secondary electrons are found to affect the current & the characteristic curve of a triode. In case of pentode an extra grid is provided to push back the electrons so that the characteristic curve will become ~~more~~ smooth.

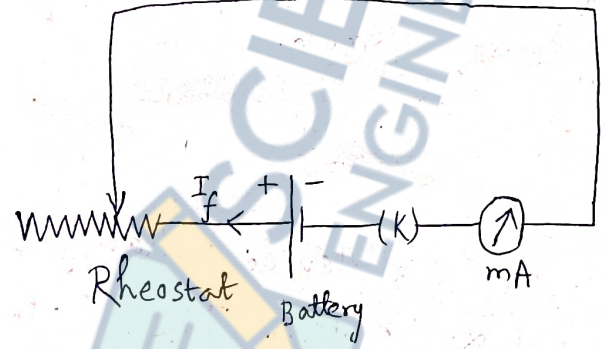
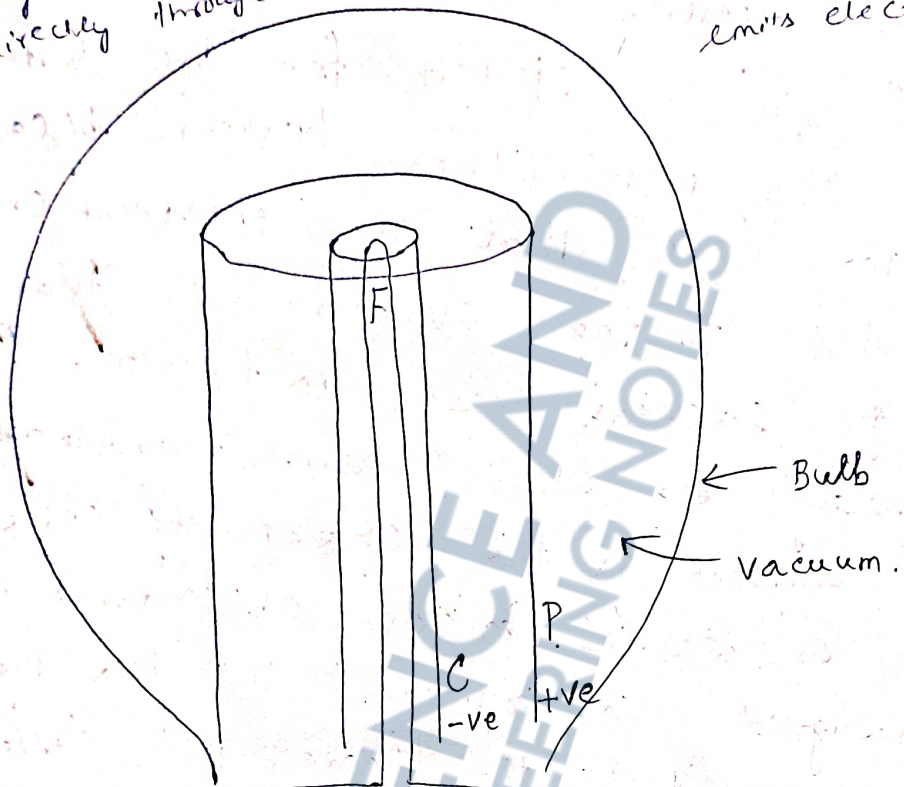
## Diode

A diode is a device having 2 electrodes called Cathode & plate. The Cathode emits the electrons & the plate receives them. Diode is used as a detector in radio circuits & as a rectifier to convert Alternating Voltage into direct voltage (A.C to D.C).

### Construction

There is a metallic wire heated by direct current or alternating current. It is called filament. It is surrounded by a thin metal sleeve and it emits electrons due to

With barium & strontium oxides. called the Cathode, ~~thermionic emission~~. The filament current can be varied by changing the resistance of the rheostat as shown in the figure. There is no electrical connection between heater & Cathode. The heating current is passed through filament & Cathode is heated indirectly through heat transfer from filament & the Cathode emits electron.



$I_f$  = Filament current

Symbol of diode



P = Plate  
C = Cathode  
F = Filament.

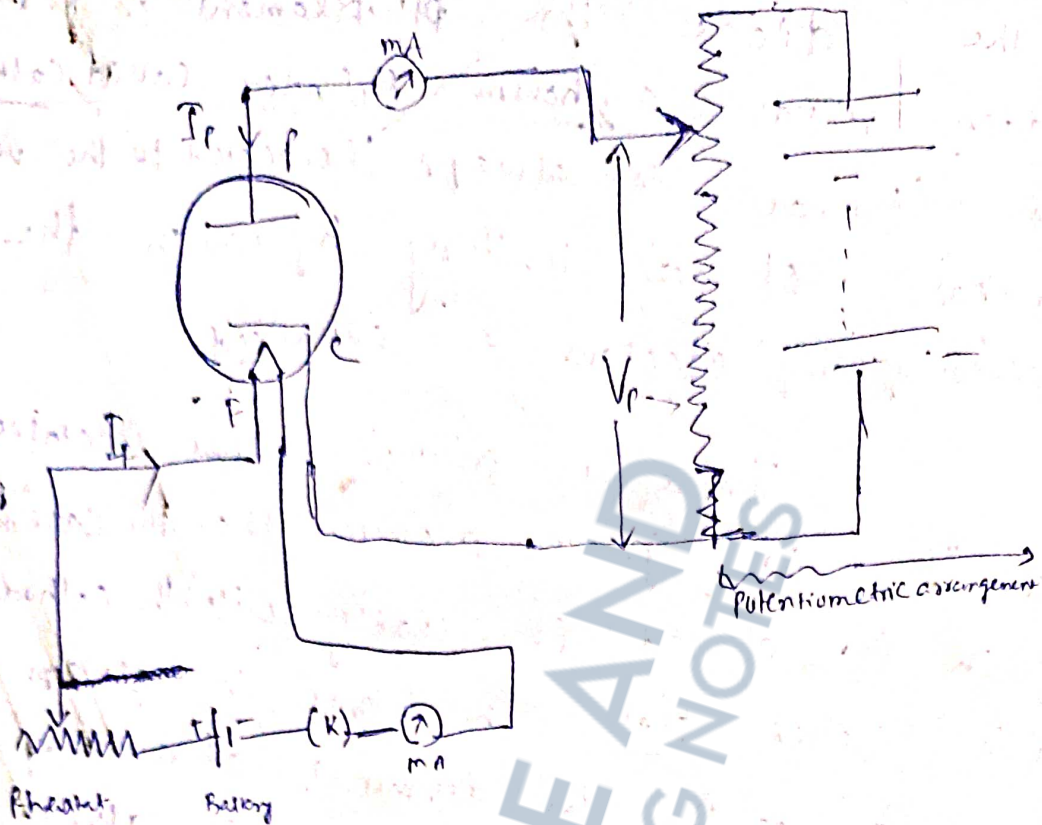
This Cathode is always connected to the -ve terminal of a battery by which the emission of electrons is enhanced.

There is another hollow metallic cylinder of greater diameter than the cathode which is placed co-axially with the cathode. It is called plate. The plate is always connected to the +ve terminal of a battery so that it will be able to attract the electrons. Due to these electrons, there is a small current flowing in the plate circuit. called plate current ( $I_p$ )

To avoid the collision of electrons with the atoms & molecules of air, the air inside the bulb is completely pumped out & vacuum is created. i.e. vacuum diode is also called vacuum diode

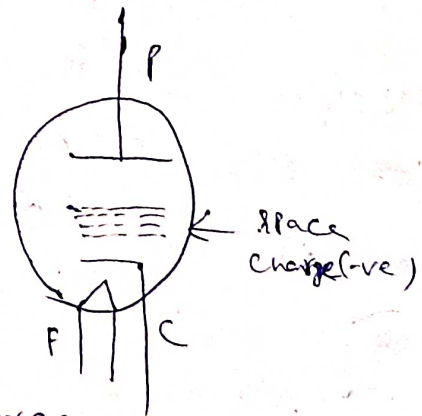
## Action

To study the functioning of the diode; the circuit diagram in the ~~previous~~ next page is used where potentiometric arrangement has been shown by which the plate voltage ( $V_p$ ) or  $E_b$  can be gradually changed. The corresponding plate currents ( $I_p$ ) or  $I_b$



are noted from the milliammeter. For each filament current, a set of observations can be taken.

When the plate voltage is low, all the electrons emitted by the cathode cannot be taken away by the plate. As a result, many electrons form a cloud of -ve charge near the cathode. 2 cm between the cathode and the plate. The electron cloud is called space charge. If space charge will be present, further emission of electrons become difficult. i.e. the diode will not function smoothly.



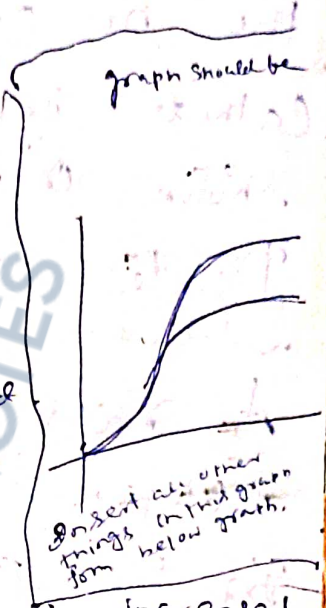
Under this space charge limited condition, the square of the plate current is found to be directly proportional to the cube of the plate voltage. This is called Child's Law

$$\therefore I_p^2 \propto V_p^3$$

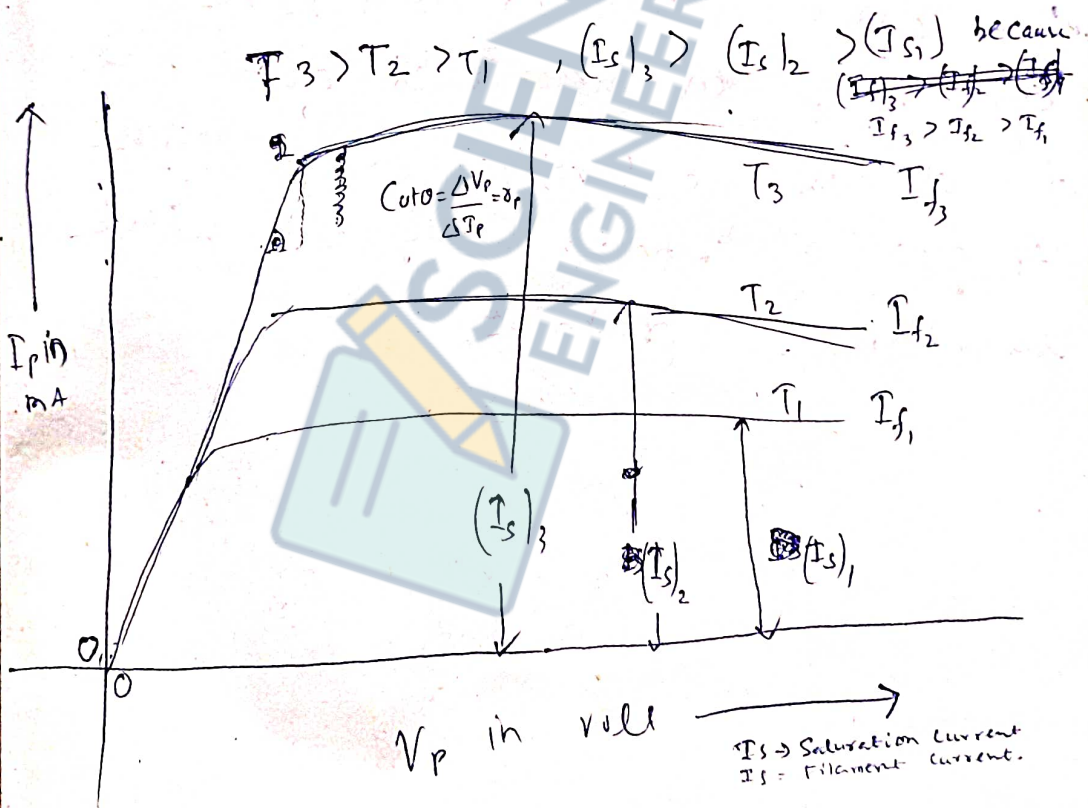
$$\text{or } I_p^2 = k V_p^3$$

where  $k$  is a constant

Also  $I_p \propto V_p^{3/2}$



When applied voltage is increased



Very much, the space charge vanishes because all the electrons emitted by the cathode are taken away by the plate. This is called saturation stage. The

endlessly. The plate current changes from anode to anode to anode.

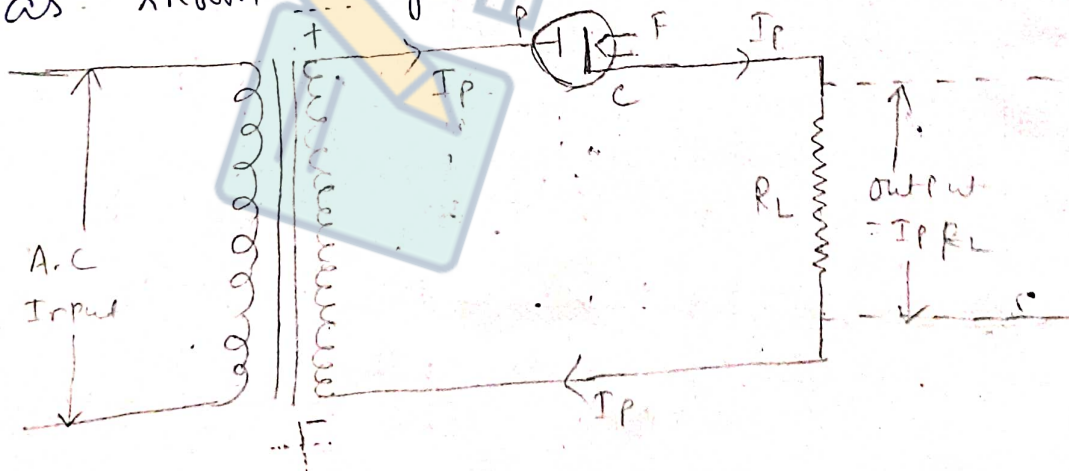


Current under this condition is called saturation current ( $I_s$ )

For different filament currents, the no of electrons emitted by the cathode become different. This has been shown on the graph drawn on the previous page.

## Diode as a rectifier

A rectifier is a device in which alternating voltage is converted to direct voltage. In this respect, its function resembles that of an eliminator. If one diode be used, then output voltage is found to be discontinuous as shown in the graph no-1. If two diodes be employed, it is possible to get a continuous supply of voltage as shown in graph-2.

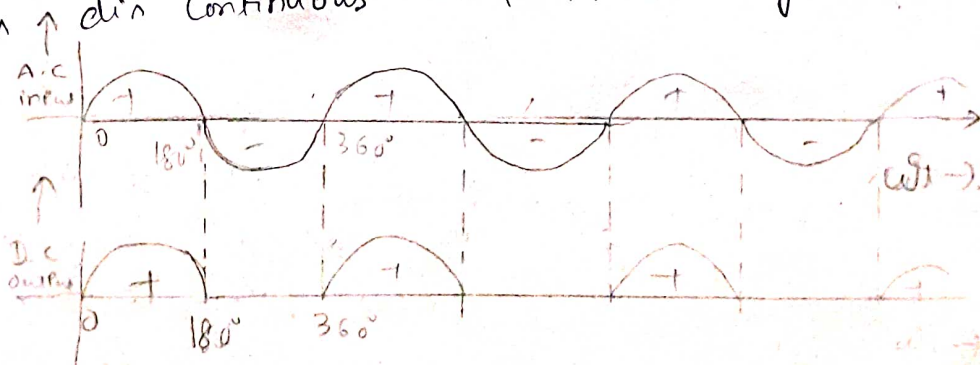


### 1. Diode as a half wave rectifier

Fig - 1

We know that alternating voltage changes sign twice during a

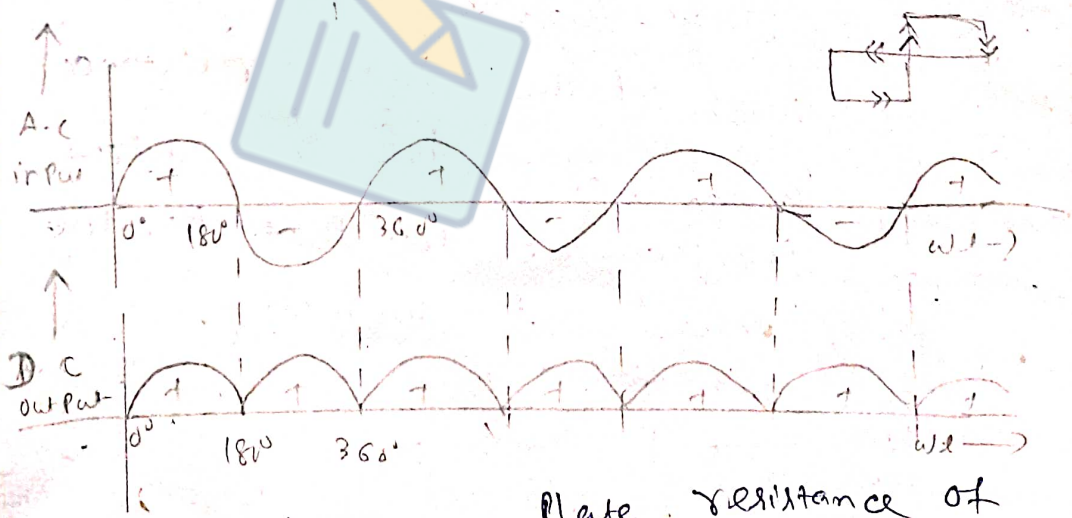
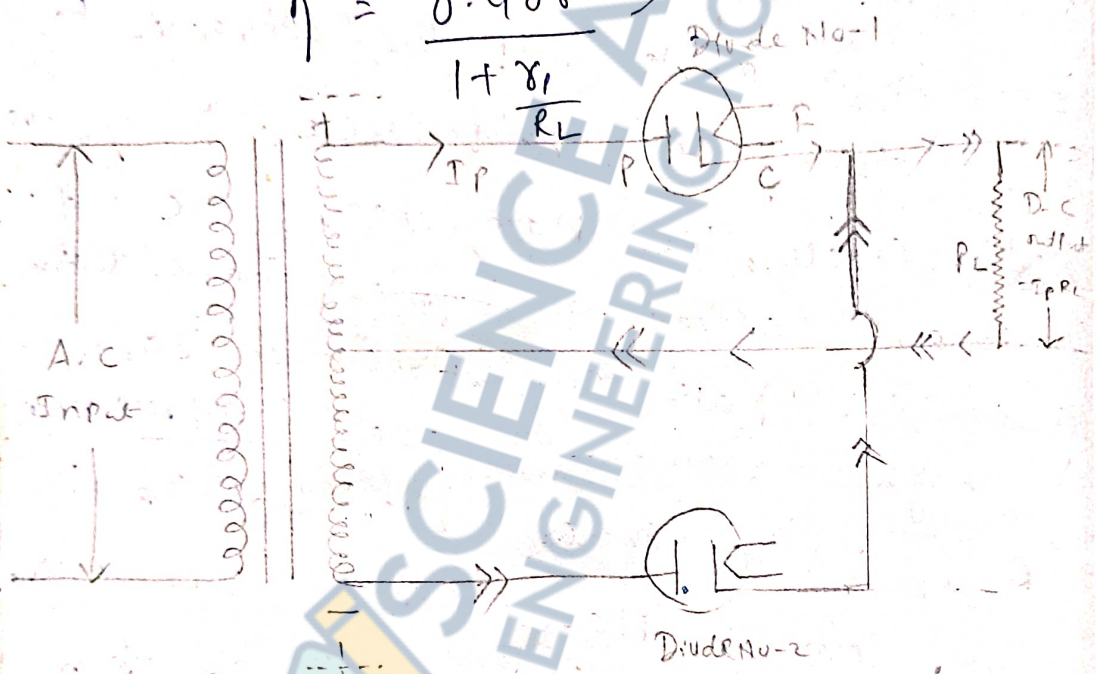
Complete cycle - when the voltage supplied to the diode is +ve, the plate becomes +ve and it can attract the electrons emitted by the cathode (which has been heated by the filament circuit, not shown in the diagram). As a result, there is a small current of the order of milli amperes in the plate circuit. This current increases from zero to a max<sup>m</sup> and then decreases to zero during +ve half cycle. During -ve half cycle, the plate becomes -ve and does not attract a ~~single~~ single electron and the plate current becomes zero. A high resistance called load resistance ( $R_L$ ) of the order kilo ohm up to Mega ohm is placed in the circuit. The output voltage can be derived from the two ends of the load resistance. In the graph no-1 the D.C output has been shown and we see that the output is discontinuous and fluctuating.



Since there is a unidirectional flow of electrons from the cathode towards the plate, it is compared to a valve and called diode valve.

Efficiency of rectification is defined as the ratio of the d.c. output power to the A.C. input power. And it can be shown that

$$\eta = \frac{0.406}{1 + \frac{\gamma_p}{R_L}}$$



where  $\gamma_p$  = plate resistance of the diode, obtained as the reciprocal of the slope of the  $\phi$

Linear portion of the characteristic curve of the diode.

Since  $r_p \ll R_L$ , we can <sup>neglect</sup> the ratio  $\frac{r_p}{R_L}$  compared to one.

Then  $\eta = 0.406 = 40.6\%$ .

## 2. Diode as full wave rectifier

When the input voltage reaches diode No-1 as +ve, it receives electrons from the Cathode. At that time diode No-2 remains inactive as its plate receives -ve voltage.

Thus, during the first half cycle, diode No-1 only functions and produces a small fluctuating current of the order of few milli amperes. When this current flows over the load resistance, the voltage produced between its two ends can be tapped.

During the 2nd half cycle, diode No-2 becomes active where as diode No-1 becomes inactive. As shown in fig-2,

the circuit has been prepared in such a manner that the direction of flow of current through the load resistance remains the same. And the output voltage is unidirectional, yet fluctuating but continuous.

## Rectification efficiency

$$\eta = \frac{\text{D-c power output}}{\text{A-c power input}}$$

$$= \frac{0.812}{1 + \frac{\delta_p}{R_L}}$$

neglected and we

Since  $\delta_p \ll R_L$ ,  
the ratio of  
 ~~$\delta_p$~~   $\frac{\delta_p}{R_L}$  can be

$$\text{get } \eta = 0.812 = 81.2\%$$

## Triode →

Introduction → It is a device having three electrodes namely cathode, plate and grid. The grid is a piece of wire wound round the cathode rather loosely and it is connected to -ve terminal of a battery so as to control the flow of electrons. In this regard it is more efficient than a Diode. Triode is used to amplify a smaller voltage and it is also used as an oscillator by which electromagnetic waves can be generated.

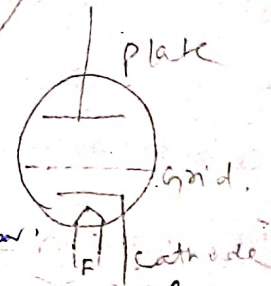
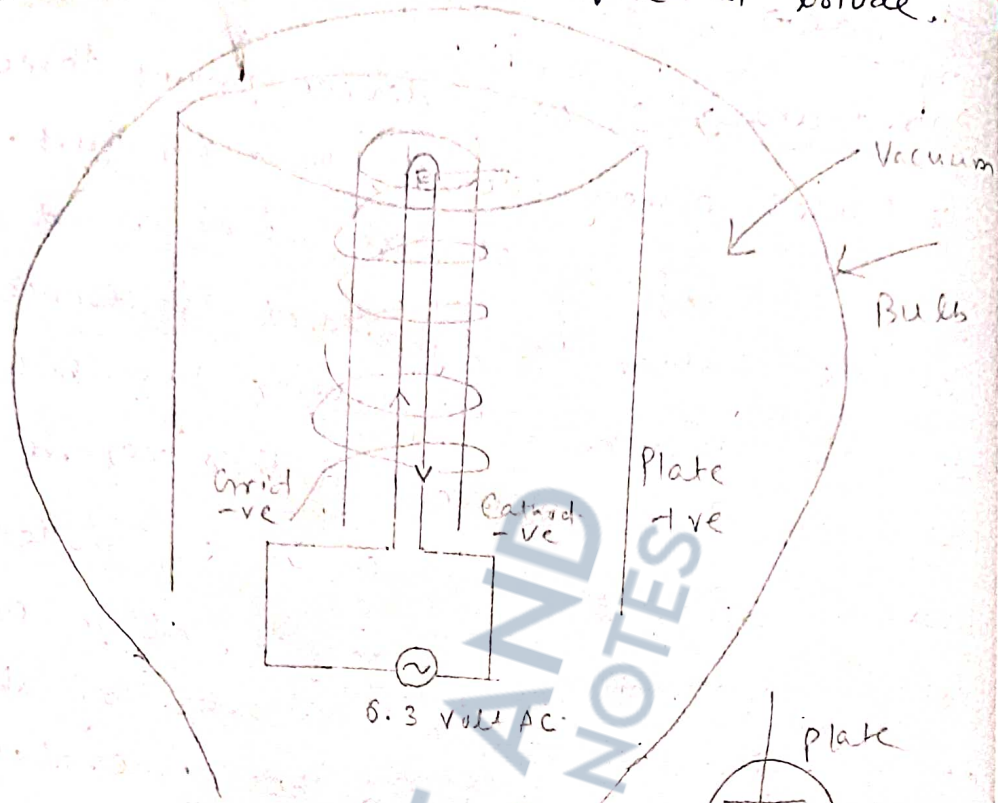
## Construction →

- 1) Filament
  - 2) Cathode
  - 3) Plate
- } Same as diode  
(write @ in exam, as of diode)

4) Grid → It is also called 'Controlled grid' because it is nearer to the cathode and slight change in its -ve voltage makes a greater change in the plate current. The grid is more effective than the plate. It will be connected to the +ve terminal of a battery then it will accept electrons and ~~it~~ <sup>EP decreases.</sup> grid current developed. Grid current is undesirable because it causes power loss in ~~the circuit.~~ Therefore, the grid is always connected to the -ve terminal of a battery.

All these are kept inside a hard glass bulb with the inside air pumped out.

Hence it is also called Vacuum triode.



### Action (working)

→ See 204 page Simulstap

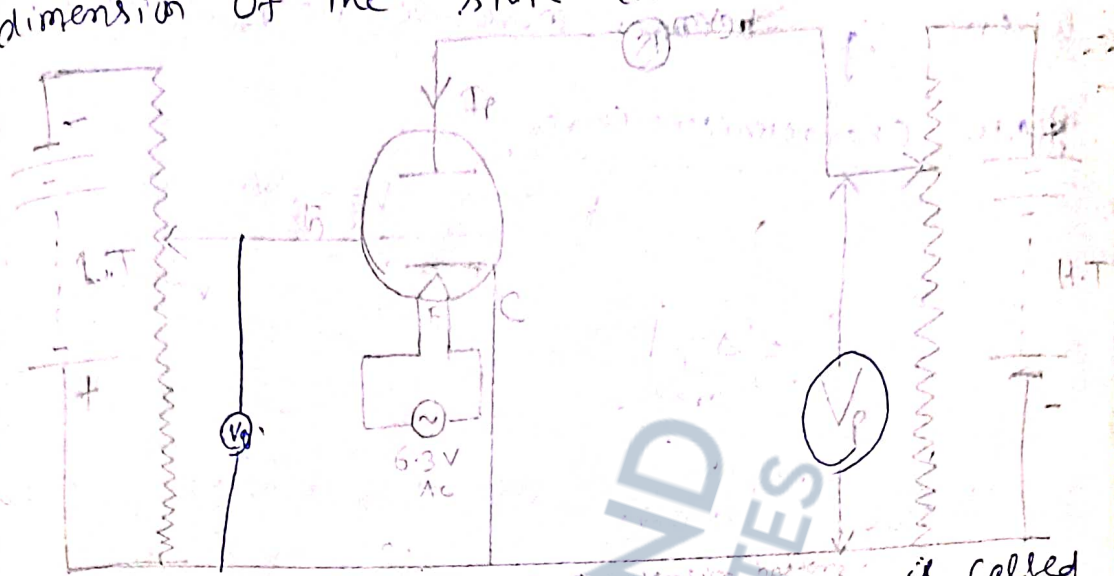
By using potentiometric arrangement for the plate voltage ( $V_p$ ) and grid voltage ( $V_g$ ) one can get different values of  $I_p$ . Two sets of characteristics curves can be drawn.

- ① ~~the~~ Plate current versus Plate voltage when the grid voltage is kept constant.
- ② Plate current Vs grid voltage when Plate voltage is kept constant.

### Plate Characteristic Curve →

with a particular value of grid voltage, the plate voltage is gradually increased. The plate current is found to increase also. Unlike the diode there is no saturation stage. The slope on any linear portion

of any curve can be found out. The dimension of the slope is that of conductance.



H.T. → High-tension battery, L.T. → Low-tension battery  
 The reciprocal of the slope is called plate resistance ( $r_p$ ).

$$i.e. \quad r_p = \left( \frac{\Delta V_p}{\Delta I_p} \right) V_g$$

This shows that the plate resistance of the same triode changes with the operating voltage.

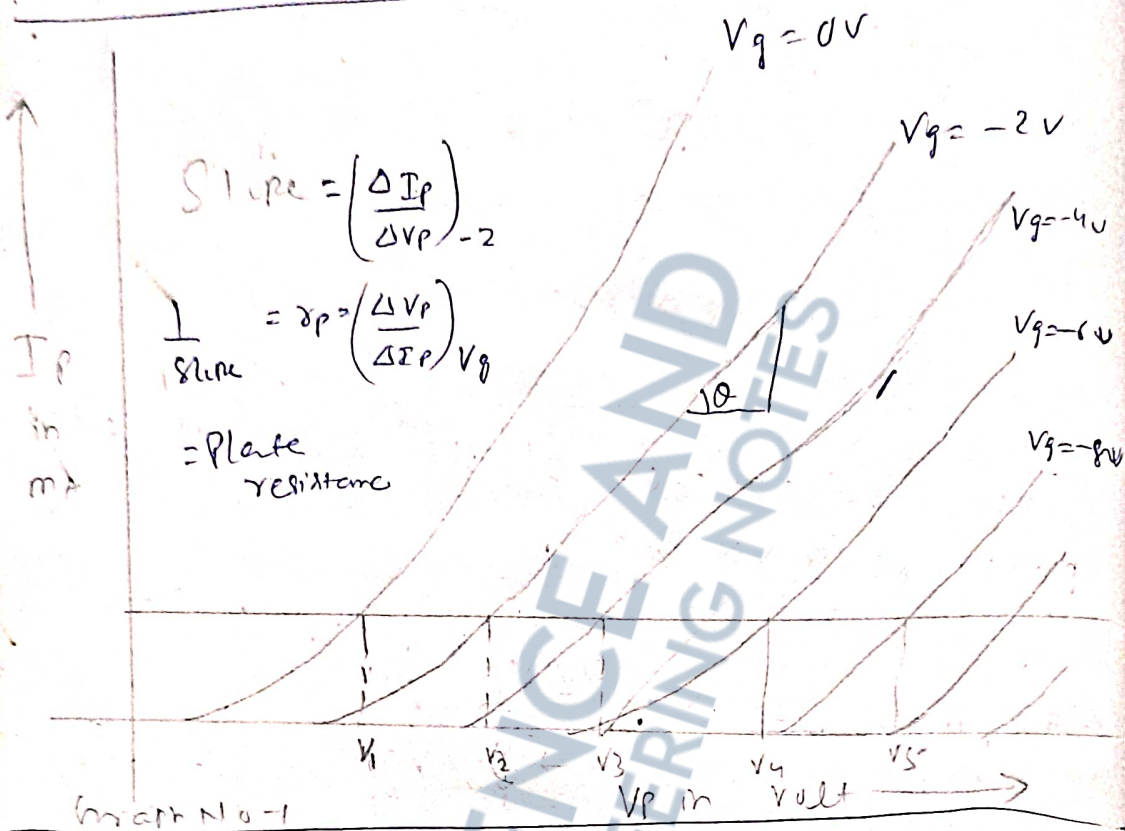
If the grid be made more -ve, the max. value of the plate current reduces very much and a certain grid voltage called cut-off grid voltage or critical grid voltage, the plate current will be zero.

However, if a great plate voltage is applied, if we draw a line // to x-axis for  $I_p = I_0$  (say), then it will cut different curves at different points. Dropping  $\perp$  lvs, we see that more plate voltage is necessary.



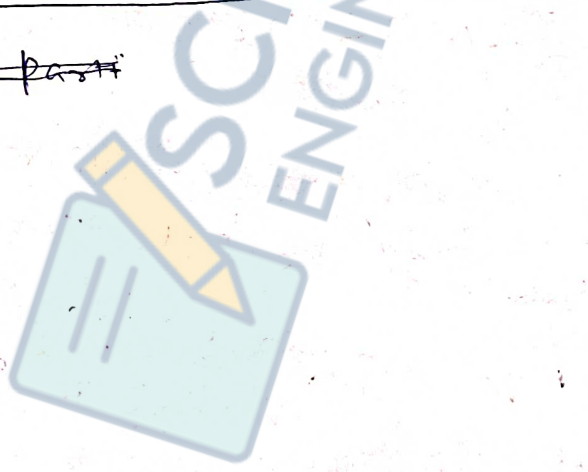
to get the same amount of plate current,  
 when  $V_g$  becomes more -ve (i.e.  $V_4 > V_3 > V_2 > V_1$ )

Plate characteristic curves



Mutual Characteristic Curves

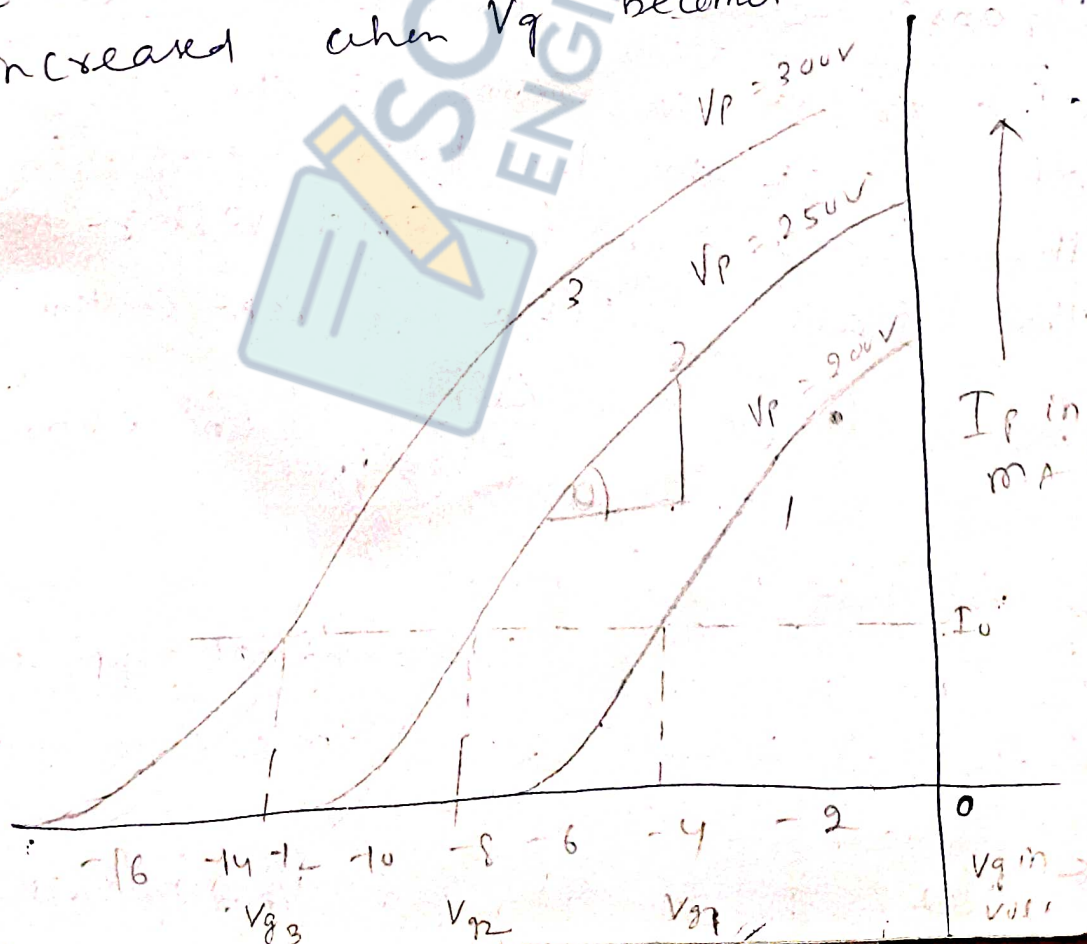
~~For a part~~



For a particular plate voltage  $V_p$  the grid voltage be gradually made more -ve. Then the plate current  $I_p$  will decrease. In graph No-2, ~~the~~ three such curves have been shown. Slope on any curve can be found out from the linear portion its dimension is the same as that of, conductance which we call as transconductance or mutual conductance ( $g_m$ ).

$$\therefore g_m = \left( \frac{\Delta I_p}{\Delta V_g} \right) V_p$$

If we draw a horizontal line parallel to x-axis, at a particular current  $I_0$ , we see  $V_p$  has to be increased when  $V_g$  becomes more -ve.



## Value Constants of a triode

A triode is characterized by 3 constants.

1. Plate resistance ( $r_p$ )
2. Transconductance ( $g_m$ )
3. Amplification factor ( $\mu$ )

Plate resistance of the ~~triode~~ triode is defined as the ratio of the change in the plate voltage to the change in the plate current when grid voltage is kept constant.

$$r_p = \left( \frac{\Delta V_p}{\Delta I_p} \right)_{V_g}, \text{ Unit Ohm.}$$

Transconductance or mutual conductance of a triode is defined as the ratio of change in the plate current to the change in the grid voltage when the plate voltage is kept constant.

$$g_m = \left( \frac{\Delta I_p}{\Delta V_g} \right)_{V_p}, \text{ Unit ohm}^{-1}.$$

Amplification factor  $\rightarrow$

To understand the meaning of amplification factor of triode, let us take a numerical example.

Let  $I_p = 10 \text{ mA}$ , when  $V_p = 200 \text{ volt}$

$$\& V_g = -6V$$

gr.  $V_p$  be increased to 250 volt  
 with  $V_g$  kept constant at  $-6V$ , then  
 $I_p$  will increase from 10 mA to 125 mA.  
 (say)

i.e.  $\Delta V_p = 50V$  has caused  $\Delta I_p = 25mA$

keeping  $V_p$  at 250 volt, let  $V_g$  be made  
 $-8V$  volt, the plate current will decrease  
 from 125 mA to 10 mA (say)

i.e.  $\Delta V_g = 2V$  has caused  $\Delta I_p = 25mA$ .

This shows that ~~the~~ a change  
 of 50V of the plate voltage is  
 equally effective as a change of  
 2V of the grid voltage

Defn of  $\mu$  → The ratio of change in  
 the plate voltage to the change in the  
 grid voltage to keep the plate current  
 constant is called amplification factor.

i.e.  $\mu = \left( \frac{\Delta V_p}{\Delta V_g} \right) I_p = 25$  in our example -

Relation among the valve constants

$$\mu = \frac{\Delta V_p}{\Delta V_g} = \frac{\Delta V_p}{\Delta I_p} \cdot \frac{\Delta I_p}{\Delta V_g} = \gamma_p \cdot g_m$$

$$\therefore \mu = \gamma_p \cdot g_m$$

## Triode as an amplifier →

Before deriving an expression for the voltage amplification by a triode, let us study fig-1. In this circuit, there is a fixed grid voltage & a fixed plate voltage. The grid voltage be  $-6V$  and the plate voltage be  $+200V$  (say). A high resistance called load resistance ( $R_L$ ) is also included in the plate circuit. From the two ends of  $R_L$ , output voltage can be derived, which is

$$I_p \cdot R_L = \text{Fixed.}$$

If an alternating small voltage, called signal voltage, say  $e = 6 \sin \omega t$  be included in the grid circuit as shown in fig(2), then the effective grid voltage will vary between  $0$  to  $-12$  volt.

because

$$(V_g)_{\max} = +6 + (-6) = 0V$$

$$(V_g)_{\min} = -6 + (-6) = -12 \text{ volt}$$

Accordingly, the plate current will change from  $I_p$  to  $(I_p \pm \Delta I_p)$ .

At any instant of time, let the grid voltage be  $-2V$  when plate current =  $I_p + \Delta I_p$  as shown in fig 3,

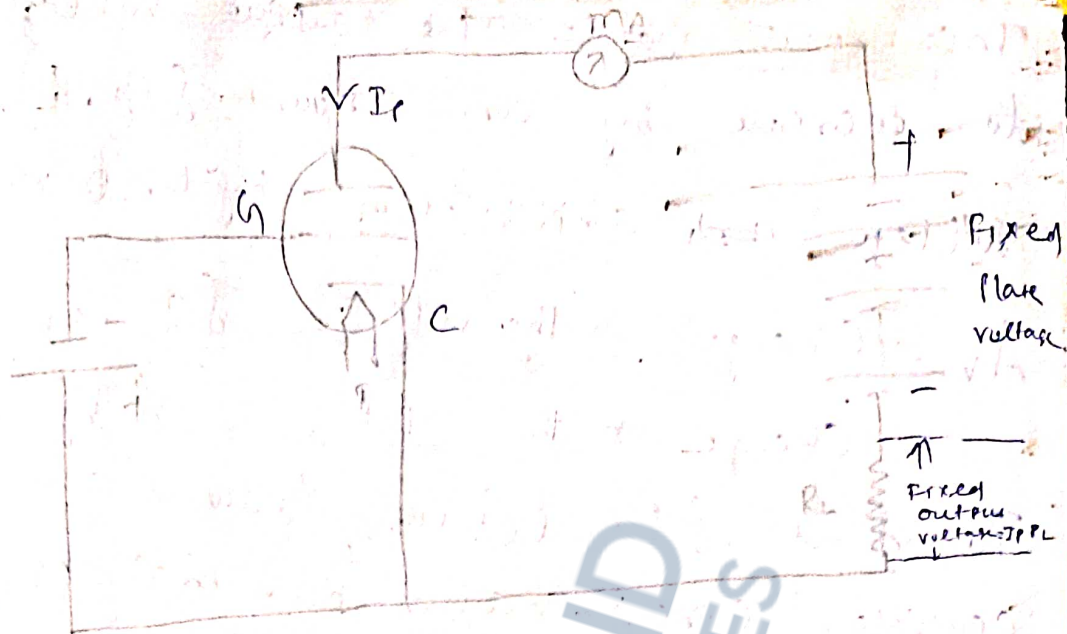
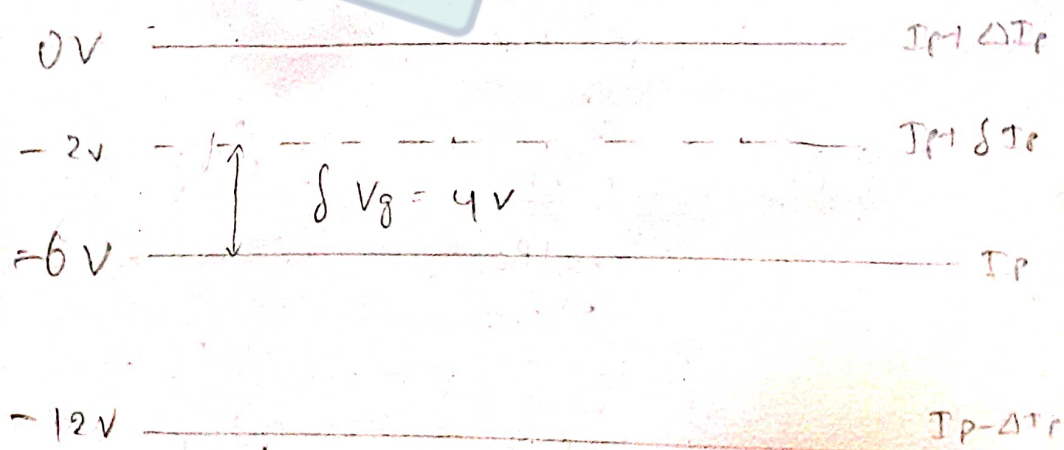
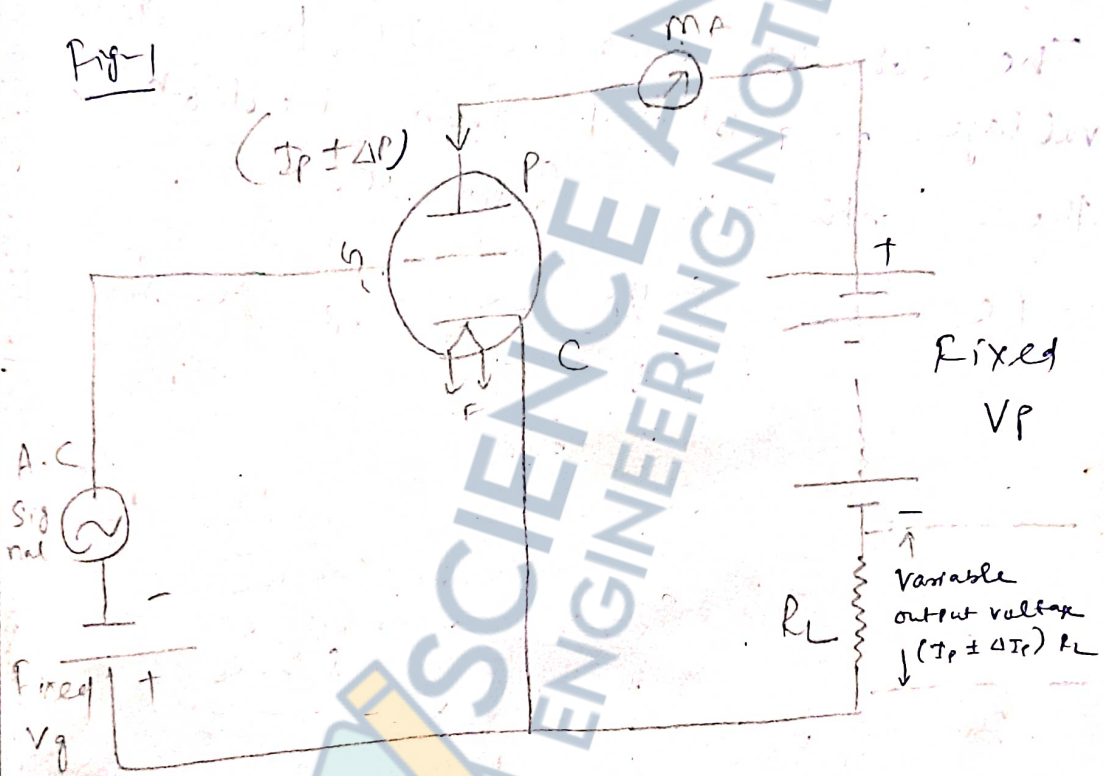


Fig-1

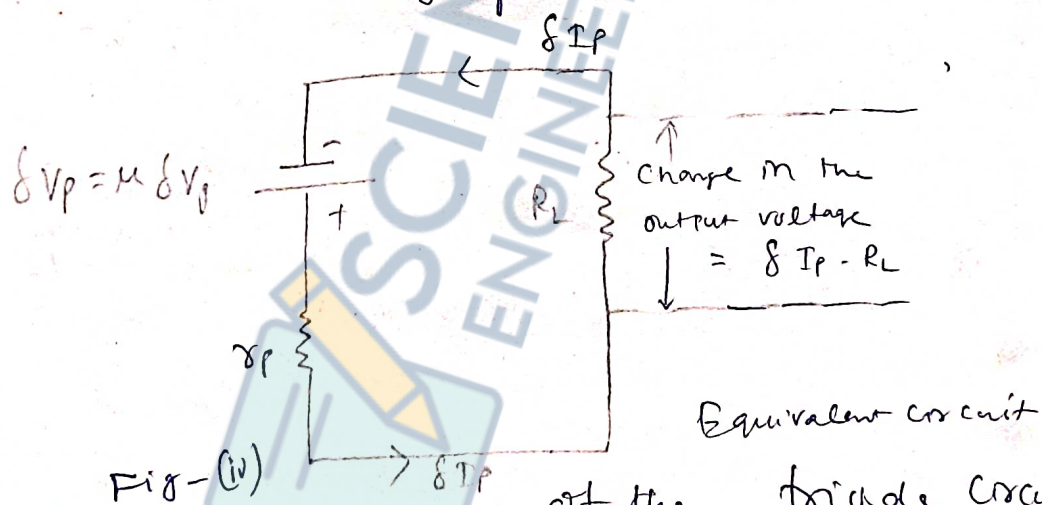


This will cause the output voltage to increase by an amount  $\delta I_p \cdot R_L$ .  
 Voltage ~~and~~ amplification by the triode

$$A_v = \frac{\text{Change in the output}}{\text{Change in the input}} = \frac{\delta I_p \cdot R_L}{\delta V_g}$$

To find  $\delta I_p$ , let us draw an equivalent circuit as shown in fig (4).  
 The effective increase of the plate voltage is obtained from the deflection or the amplification factor  $\mu$ .

$$\text{i.e. } \mu = \frac{\delta V_p}{\delta V_g} \Rightarrow \delta V_p = \mu \cdot \delta V_g$$



Total resistance of the triode circuit

$$= r_p + R_L$$

$$\therefore \delta I_p = \frac{\delta V_p}{(r_p + R_L)} = \frac{\mu \delta V_g}{r_p + R_L}$$

$$\therefore A_v = \frac{\mu \cdot \left( \frac{\delta V_g}{(r_p + R_L)} \cdot R_L \right)}{\delta V_g} = \frac{\mu R_L}{r_p + R_L}$$

$$A_v = \frac{\mu \cdot R_L}{r_p + R_L} \quad \text{No unit}$$

→ Ans for problem on the next page

① We know that in space charge limited region

$$I_p^2 = k \cdot (V_p)^3$$

$$(10 \text{ mA})^2 = k \cdot (120\text{V})^3$$

$$x^2 = k \cdot (240\text{V})^3$$

$$\Rightarrow \frac{10^4 \times \text{mA}^2}{x^2} = \left( \frac{120\text{V}}{240\text{V}} \right)^3 = \frac{1}{8}$$

$$\Rightarrow x^2 = 8 \times 10^4 \text{ mA}^2$$

$$\Rightarrow x = 2.83 \times 10^2 \text{ mA}$$

$$\Rightarrow x = 283 \text{ mA}$$

$$(10 \text{ mA})^2 = k \cdot (120\text{V})^3$$

$$x^2 = k \cdot (480\text{V})^3$$

$$\Rightarrow \frac{10^4 \text{ mA}^2}{x^2} = \left( \frac{1}{4} \right)^3$$

$$\Rightarrow x^2 = \frac{64}{27} \times 10^4 \text{ mA}^2$$

$$\Rightarrow x = 8 \times 10^2 \text{ mA}$$

$$\Rightarrow x = 800 \text{ mA}$$

②  $I_p = 10 \text{ mA}$ ,  $V_p = 100 \text{ volts}$ ,  $(2I_p)$

$$\Rightarrow \frac{(10 \text{ mA})^2}{(20 \text{ mA})^2} = \frac{k \cdot (100 \text{ V})^3}{k \cdot x^3} \Rightarrow x^3 = \frac{2}{20 \text{ mA}} \times 100 \text{ V}^3$$

$$x = \sqrt[3]{200} \text{ V} = 5.848 \text{ V}$$



$$\frac{1}{4} \times 2 = \frac{100^3 \text{ volt}^3}{2^3}$$

$$\Rightarrow m^3 = 4 \times 100^3 \text{ volt}^3$$

$$\Rightarrow x = 4^{\frac{1}{3}} \times 100 \text{ volt}$$

4.4  $y = 4^{\frac{1}{3}}$   
 $\log y = \frac{1}{3} \log 4 = \frac{1}{3} \times 2 \log 2 = \frac{1}{3} \times 2 \times 0.3010$   
 $\log y = \frac{1}{3} \times 0.602 = 0.2007$   
 $\frac{y}{10} = \frac{1}{3} = -2$

$$\Rightarrow y = \text{antilog}(0.2007) = 1.585$$

$$\therefore x = 1.585 \times 100 = 158.5 \text{ volt}$$

3. Efficiency =  $\frac{\text{D.C. output}}{\text{A.C. input}} = \frac{0.406}{1 + \frac{20}{800}} = \frac{0.406}{1 + \frac{1}{40}} = \frac{0.406}{1.025} = 39.6\%$

$R_L = 800 \Omega$   
 $r_p = 20 \Omega$   
 $I_p \cdot R_L = 10 \text{ watt}$

$$\frac{406}{80} = 5.075$$

$$32.48\% = 32.5\%$$

4.7

5.

$M = 30, R_L = 50k\Omega, r_p - gm = 30$

$M = 34, R_L = 85k\Omega, r_p - gm = 34$

Problem

4) The following readings were obtained from the linear portion of a vacuum triode. From that find:

- (a) Plate resistance (b) Mutual Conductance
- (c) Amplification factor

Ans: 11.1 Kilo ohm,  $4.666 \times 10^{-3}$  mho, 52

$V_p \text{ or } E_b$	150 V	150 V	100 V
$I_p \text{ or } I_b$	12 mA	5 mA	7.5 mA
$V_g \text{ or } E_c$	-1.5 V	-3 V	-1.5 V

$r_p = \text{Plate resistance} = \left( \frac{\Delta V_p}{\Delta I_p} \right) V_g$

$= \frac{(150 - 100) \text{ Volt}}{(12 - 7.5) \text{ mA}} = \frac{50 \times 10^3}{4.5} \text{ ohm}$

$= \frac{500}{45} = \frac{100}{9} = 11.11 \dots \text{ K ohm.}$

$g_m = \text{Mutual Conductance}$

$= \left( \frac{\Delta I_p}{\Delta V_g} \right) V_p = \frac{(12 - 5) \text{ mA}}{(-1.5 + 3) \text{ Volt}}$

$= \frac{7 \times 10^{-3}}{1.5} = \frac{70}{15} \times 10^{-3} = 4.666 \times 10^{-3} \text{ mho}$

$= 4.666 \text{ mill. ohm}$

$\mu =$  Amplification factor

$$= \mu_p \mu_m$$

$$= \frac{100}{9} \times 10^3 \text{ ohm} \times \frac{14}{3} \times 10^{-3} \text{ mho}$$

$$= \frac{1400}{27} = 51.851 \approx 52 \text{ Ans}$$

5. The voltage amplification factor is 30 with load resistance 50 K $\Omega$  and 34 with 85 Kilo ohm load resistance. Determine the value constants.

Ans: 20 Kilo ohm, 2.1 milliohm, 42

3. A half wave rectifier is used to supply 100 volts D.C from a load of 800 Ohms. The diode has a plate resistance of 200 Ohms. Find the rectification efficiency.

Ans: 32.5%

2. Plate current of a diode is 10 mA, when a plate voltage is 100 volt operating in space charge limited region. What is the plate voltage necessary to double the plate current?  
Ans: 158.7 volts

1. Under space charge limited condition, the plate current of a diode is 10 mA with a plate voltage of 120 volt. Find the plate currents with the

plate voltages of 240 and 480 volts.

Ans: 283 mA, 800 mA.

## P-N Junction

A P-N Junction can be prepared out of a crystal of Ge or Si by doping with pentavalent impurity atoms at one end and trivalent impurity atoms at the other end.

The action of such a device is understood by connecting it into 2 different ways

- (i) Forward biasing.
- (ii) Reverse biasing.

### Forward biasing

A P-N Junction is said to be forward biased when the +ve terminal of the battery is connected to the P end of the P-N Junction and -ve terminal of the battery is connected to the N end of P-N Junction.

The +ve plate repels the holes towards the junction layer where as the -ve plate repels the electrons towards the same layer. As a result, the electrons occupy the holes and leave behind holes in the N-region.

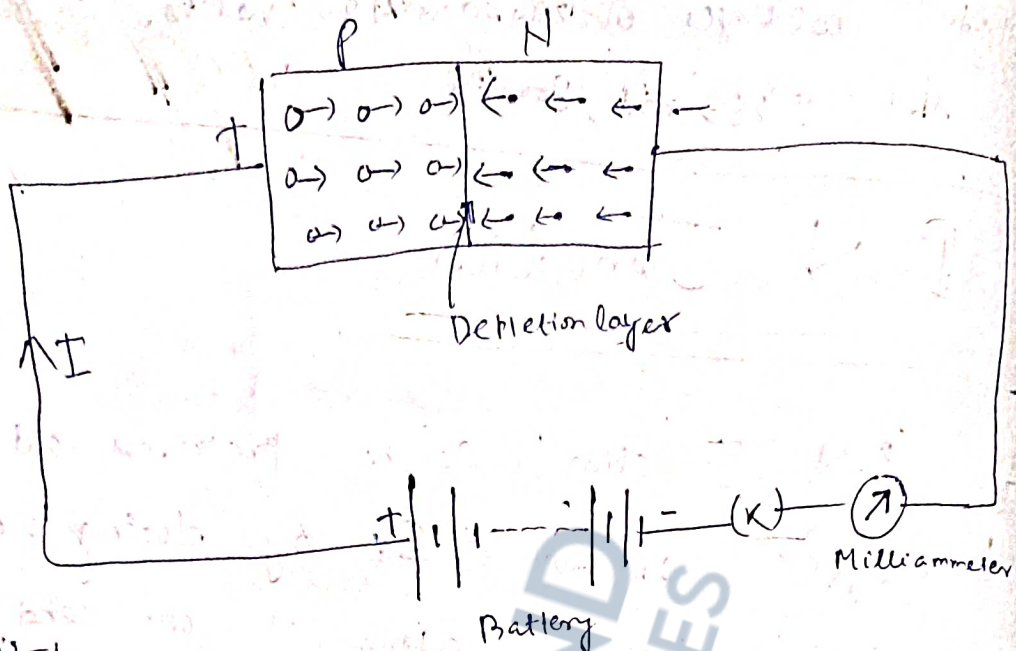


Fig-1

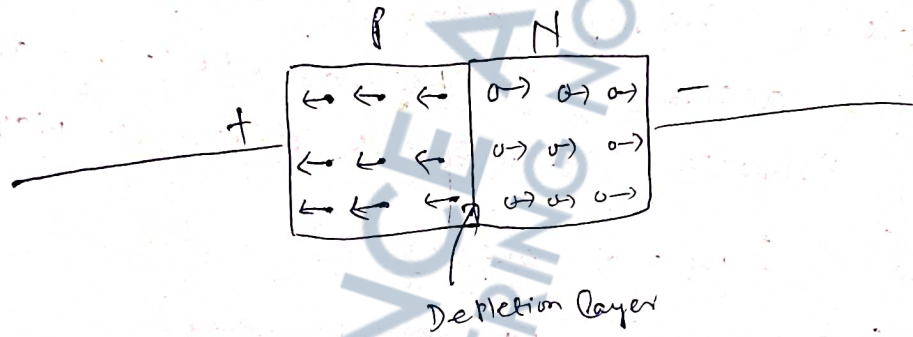


Fig-II

Intermediate stage

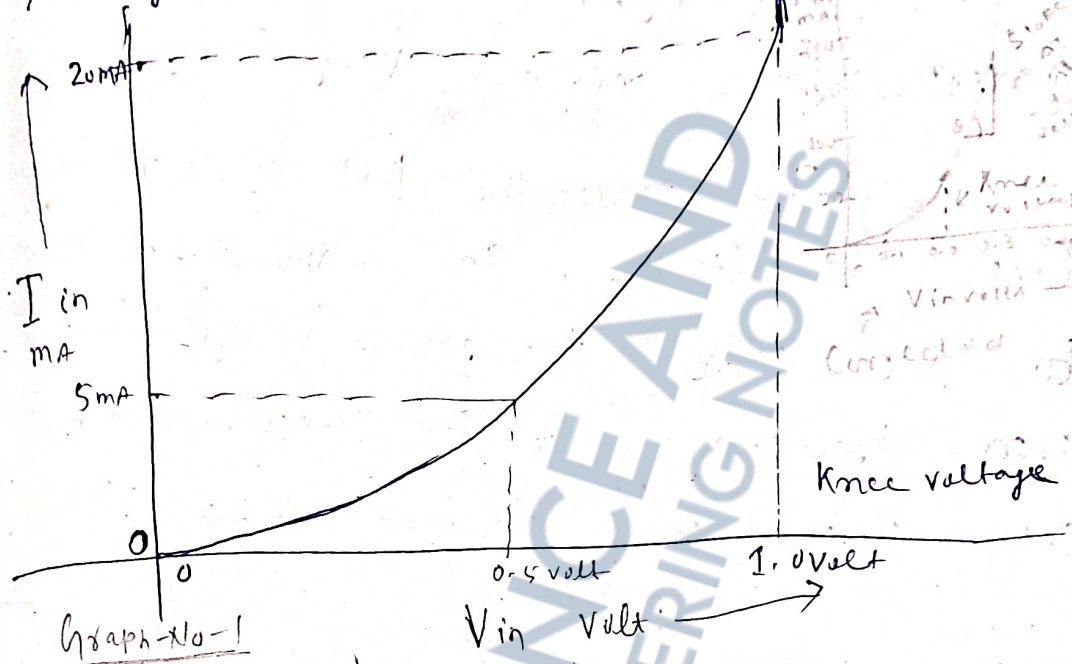
This has been shown in fig(ii).

Ultimately the electrons are absorbed by the +ve plate & holes are left behind in the P-region. This is just the situation shown in fig (i). These ~~abs~~ absorbed electrons move to the battery and then towards the N-region where they occupy the places of the holes.

Again the process is repeated.

A small current of the order of mA is produced. This current is small.

but very sensitive towards the small change of ~~the~~ applied voltage. This is indicate in graph no 1. The voltage at which the current increases very sharply is called Knee Voltage.



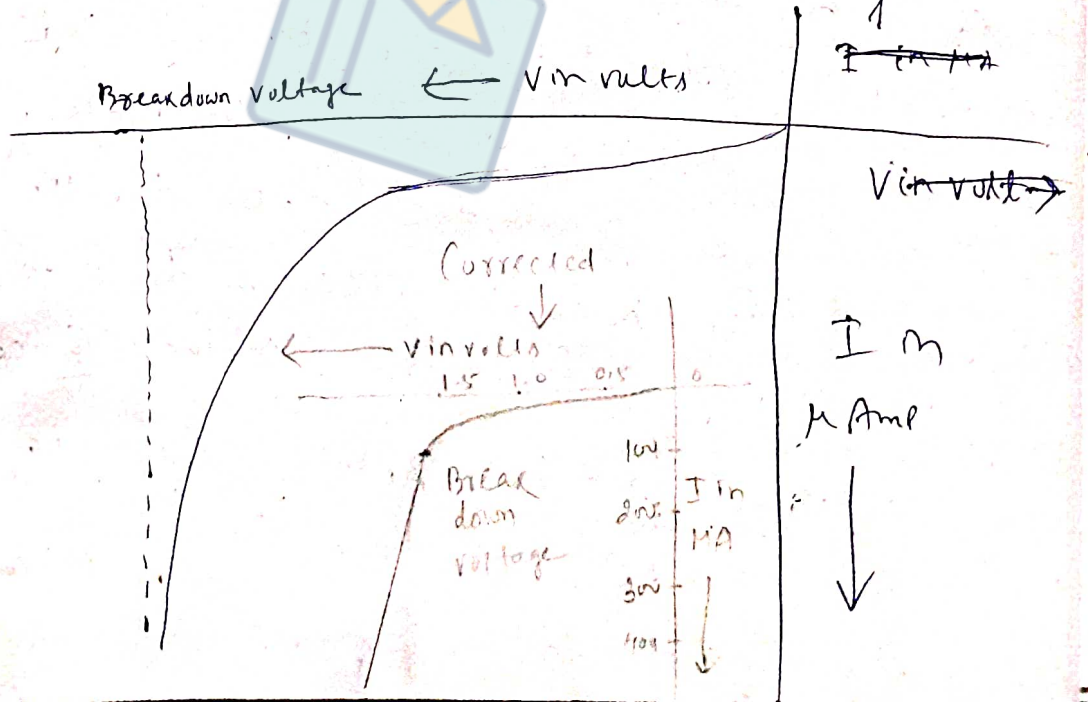
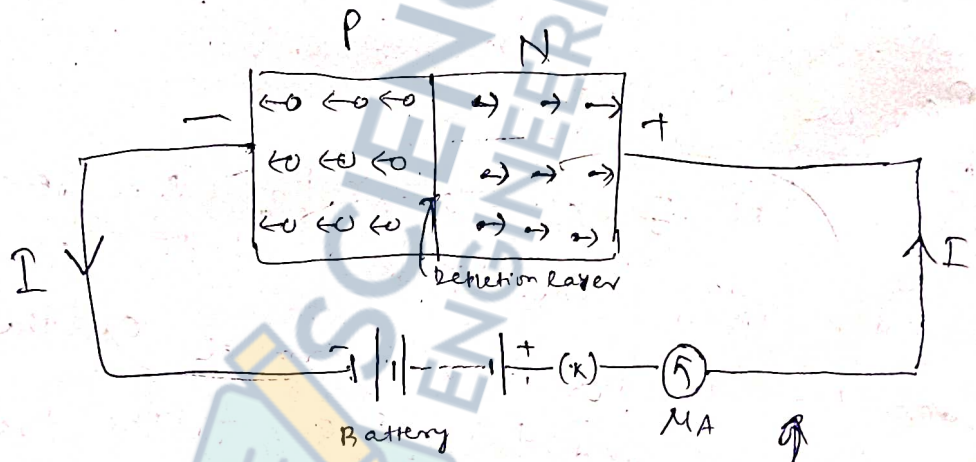
Reverse Biasing

The P-N Junction is said to be reverse biased when the P end of it is connected to the -ve terminal of the battery and the N end to the +ve terminal of the same battery. Due to the attraction of the -ve plate for the holes and the +ve plate for the electrons, the holes and electrons <sup>are</sup> drawn away from one another. Since the electrons and holes do not <sup>have not</sup> combined, the flow is almost stopped. Only a very small current of the order of microamperes

flows due to some stray electrons  
 and holes (minority charge carriers) formed on the specimen. A

Current of a few  $\mu A$  flows  
 till a voltage called breakdown voltage  
 is reached. Beyond <sup>(further)</sup> this voltage, the  
 current again increases very fast.

To make the P-N junction act  
 like a diode, care should be taken  
 to see that the applied voltage is low.  
 Then, practically the P-N Junction conducts  
 only when it is forward biased ( $\therefore I_{MAX} \approx 1mg$ )



From the above discussion, we see that the P-N Junction conducts only when it is forward biased, provided the applied voltage is low. This is similar to that of a vacuum diode valve. For this reason, the P-N Junctions have almost replaced the vacuum diodes.

The advantages are

- ① P-N Junction is very small in size
- ② It is much cheaper compared to vacuum diodes.
- ③ Operating voltage necessary for P-N junction is very small. Even, a torch battery can do the work. ( $V = 1.5 \text{ V}$ )
- ④ Unlike the vacuum diodes no extra heating device is necessary

P-N Junction as a half wave rectifier

We know that alternating voltage changes its sign twice during a complete cycle. When the voltage supplied to the P-N junction is +ve i.e. the P end is connected to the +ve terminal of the battery & N-end is connected to the -ve terminal of the battery then the

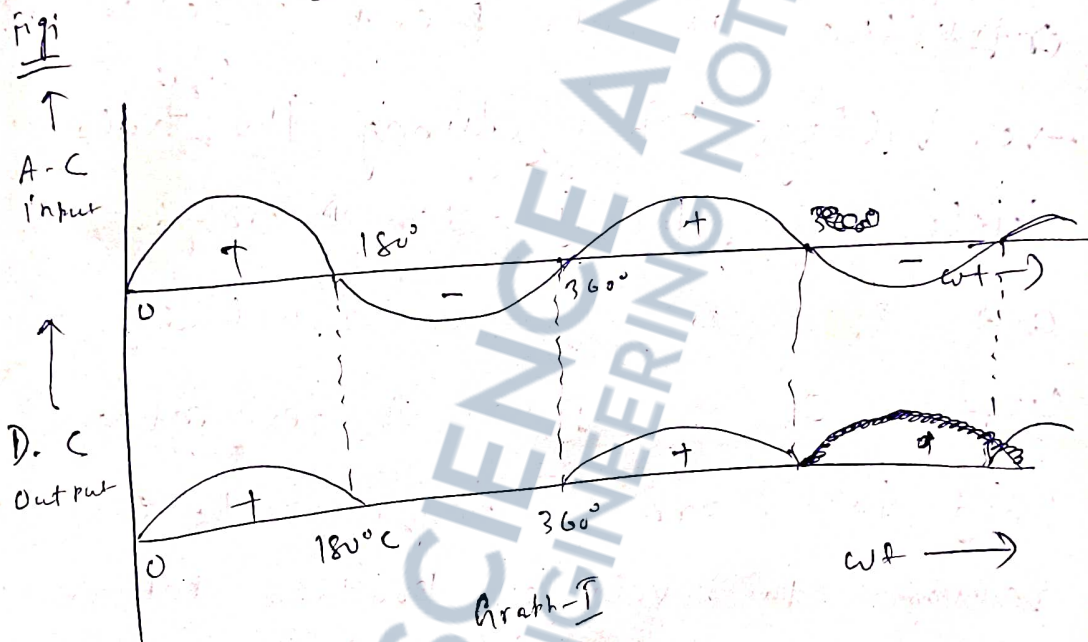
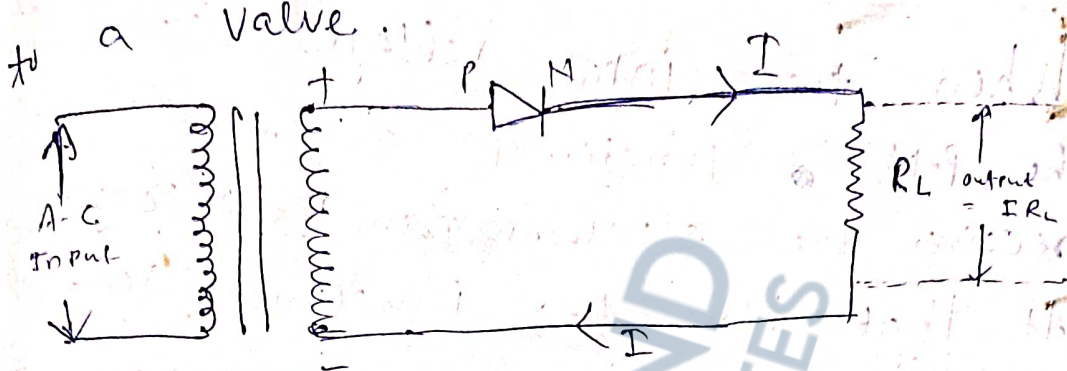


The P end repels the holes, and the  $-ve^N$  end repels the electrons, towards the junction layer. As a result, the electrons occupy the holes and they are absorbed by the +ve plate and a small current of the order of a few milliamperes is developed in the circuit. This current increases from 0 to a max and then decreases to zero during the +ve half cycle.

During the -ve half cycle the P end becomes -ve and attracts the holes and N end becomes +ve attracts the electrons. As a result holes & electrons do not combine and thus the current produced is of the order of a few milliamperes which can not be easily detected. A high resistance called load resistance ( $R_L$ )

of the order of kilo ohm up to mega ohm is placed in the circuit. The output voltage can be derived from the two ends of this load resistance. In the graph (1) the d.c output has been shown & we see that the output is discontinuous

and fluctuating. Since there is unidirectional flow of electrons from the P-end towards the N-end, it is compared to a valve.



Efficiency of rectification is defined as the ratio of d.c. output power to the a.c. input power and is shown as

$$\eta = \frac{0.406}{1 + \frac{\delta_p}{R_L}}$$

where  $\delta_p$  = Resistance of the p-n junction

obtained as the reciprocal of the slope of the graph of the action of the junction

Since  $\delta_p \ll R_L$ , we can neglect the ratio  $\frac{\delta_p}{R_L}$  compared to 1. Then

$$\eta = \text{Efficiency} = 0.406 = 40.6\%$$

P-N junction as full wave rectifier

When the input voltage reaches the P-N junction (1) as +ve, it receives the electrons from the N-region.

At that time P-N junction (2) remains ~~active~~ <sup>inactive</sup> as its P-end receives

-ve voltage. Thus, during the <sup>first</sup> half-cycle the P-N junction (1) only functions

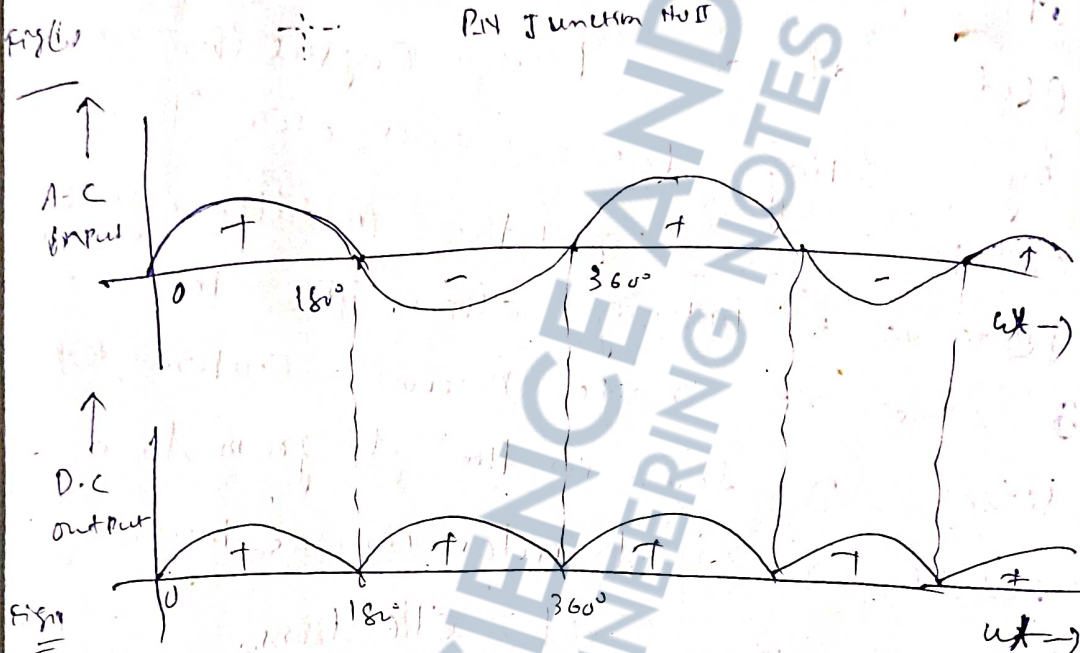
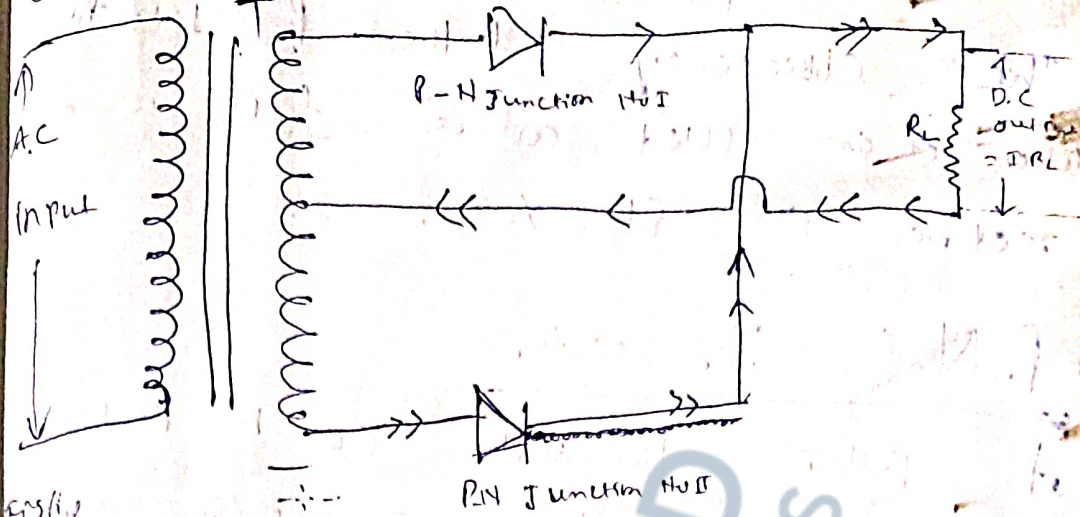
and produces a small fluctuating current of the order of few milliamperes.

When this current flows over the load resistance, the voltage produced between the two ends can be ~~can~~ tapped.

During the second half cycle, the P-N junction (2) becomes active where as the P-N junction (1) remains <sup>is</sup> inactive. As shown in fig (11), the circuit

has been prepared in such a manner that the direction of flow of current through the load resistance remains the same. And the output voltage is unidirectional ~~get~~

fluctuating : But it's continuous



Rectification efficiency is defined as the ratio of d.c. output power to the a.c. input power and is shown as  $\eta =$

$$\text{Efficiency} = \frac{0.812}{1 + \frac{r_p}{R_L}}$$

where  $r_p =$  Resistance of the P-N junction

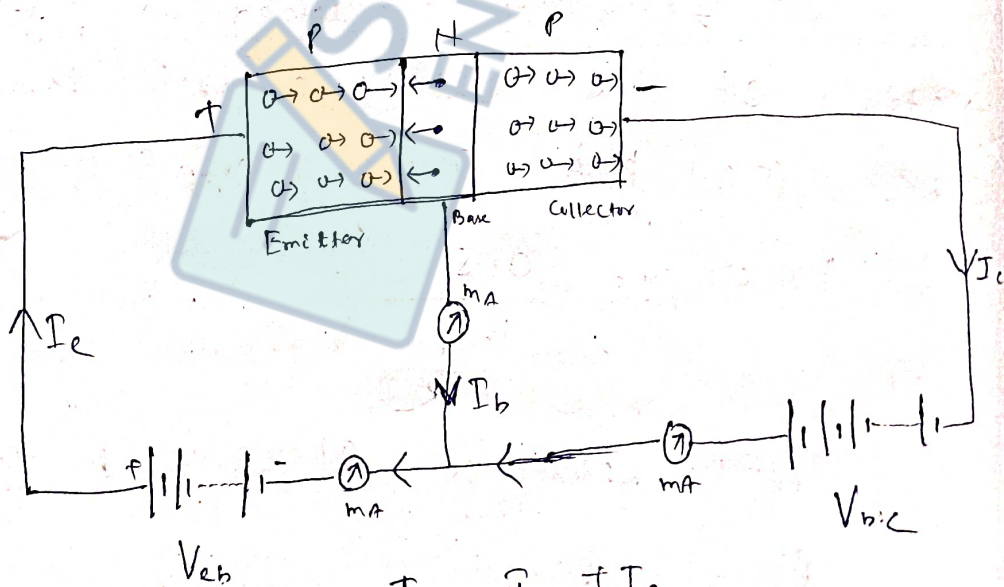
$R_L =$  Load resistance

Since  $r_p \ll R_L$ , then we can the ratio of  $\frac{r_p}{R_L}$  as compared to 1, then the

Efficiency  $\eta = 0.82$  i.e. 81.2%. Thus  
 The efficiency of the P-N junction  
 when it is used as a full wave  
 rectifier is 81.2%.

## P. N. P Transistor

A P-N-P transistor can be prepared  
 out of a pure crystal of Ge or Si  
 by doping the 2 ends with trivalent  
 impurity atoms and the middle region by  
 pentavalent impurity atoms. The middle  
 zone is made narrow compared to the  
 sides as shown in the diagram. As a  
 result, no. of holes become much higher  
 than the no. of electrons.



$$I_e = I_b + I_c$$

A P-N-P transistor thus formed  
 can be regarded as a P-N junction connected  
 in series with an N-P junction. To study

the action of the P-N-P transistor. The P-end of the P-N Junction is connected to the +ve terminal of a battery  $V_{eb}$ . i.e. this P-N junction is forward biased and this P end is called emitter. The other P end of the P-N-P transistor is connected to the -ve terminal of another battery  $V_{bc}$  and we call that P-end as the collector. The middle zone is called base. This base is connected to the -ve end of  $V_{eb}$  and to the +ve end of  $V_{bc}$ .

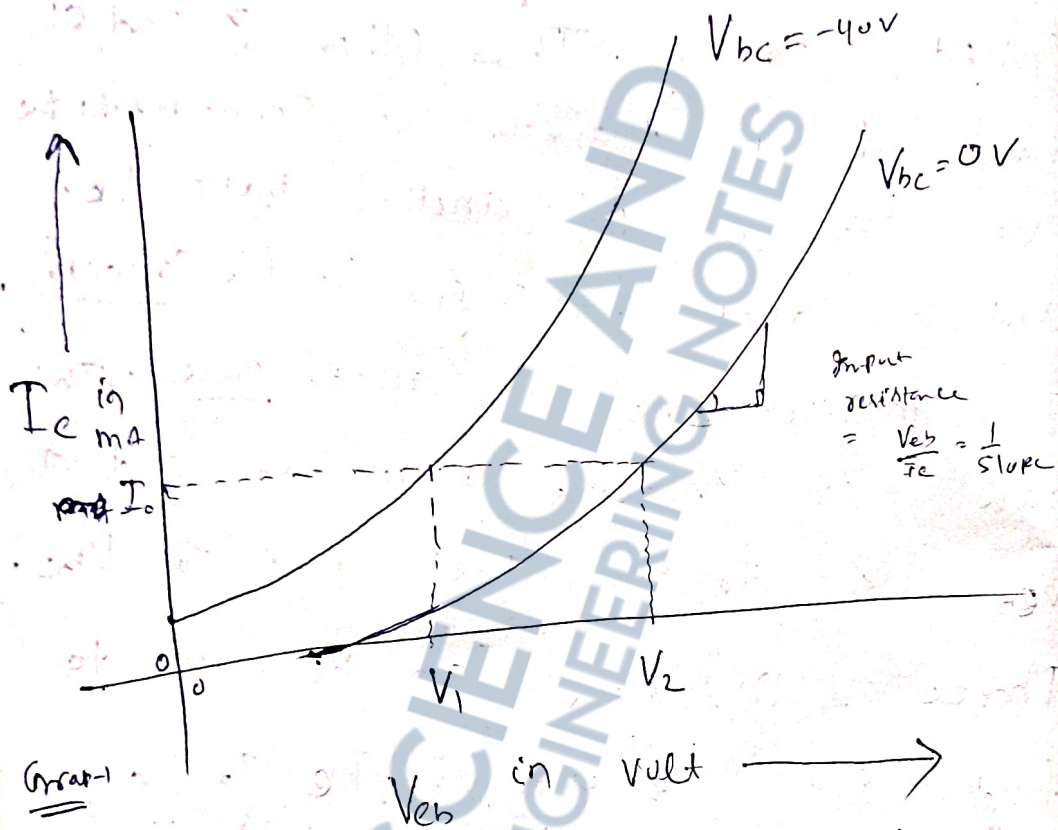
Three  $m_A$  Ammeters can be used to

find the 3 currents -  $I_e, I_b, I_c$

Emitter is heavily doped to provide large number of charge carriers, base is lightly doped & this it passes most of the emitted injected carriers to the collector. Collector is moderately doped.

Action → Since the emitter is connected to the +ve terminal of a battery, the ~~that~~ holes of that end are pushed towards the base. The electrons of the base region are attracted towards the emitter. Hence they combine, but no. of electrons is less compared to the no. of holes. Majority of the holes cross over into the collector zone. The ~~less~~ electrons ~~reach~~ reach the +ve end and get absorbed by the +ve plate. They move through

The conducting wire upto the junction point and some of them go towards the base while the others go towards the collector zone. These electrons will occupy the holes present in the base & collector zones.



Due to the flow of electrons through the conductors a current like  $I_b$  is produced in the base circuit and  $I_c$  along the collector circuit such that their sum equals to that of the emitter current  $I_e$ .

$$\therefore I_e = I_b + I_c \quad \text{Obviously } I_e < I_c$$

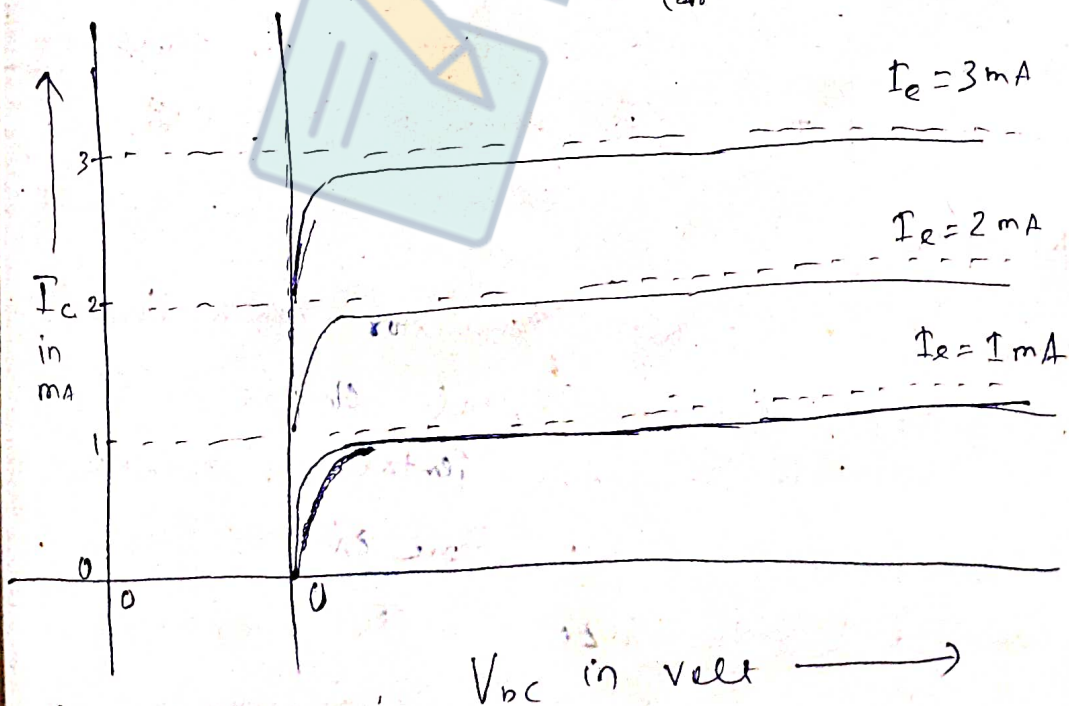
To sets of characteristic curves can be drawn for PNP and NPN transistors.

- (i)  $I_e$  versus  $V_{be}$  when  $V_{bc}$  or  $I_c$  is kept constant.

# (Input Characteristic Curves)

(ii)  $I_c$  versus  $V_{bc}$  when  $V_{eb}$  or  $I_e$  is kept constant. (output characteristic curves.)

As shown in graph no (i), the emitter current increases slowly with the increase of emitter base voltage ( $V_{eb}$ ) when  $V_{bc}$  is made 0. On the other hand, emitter current increases very quickly with the change of  $V_{eb}$  when  $V_{bc} = -40V$  (say). If we draw a horizontal line // to x-axis for a constant current  $I_o$ , then it cuts the 2 curves at 2 different points. It indicates that voltage necessary to have  $I_o$  for  $V_{bc} = -40$  volt is  $V_1$  which is less than  $V_2$ .





(Voltage necessary for same current  $I_c$  when  $V_{bc} = 0$ )

Input resistance can be found out as the reciprocal of the slope

from the linear portion of the curves.

With the emitter current kept fixed the base collector voltage is to be gradually changed so that different collector currents are obtained.

When this current plotted ~~vs~~ along y-axis &  $V_{bc}$  along x-axis, a curve is obtained which

is similar to the static characteristic curve of a diode value, saturation

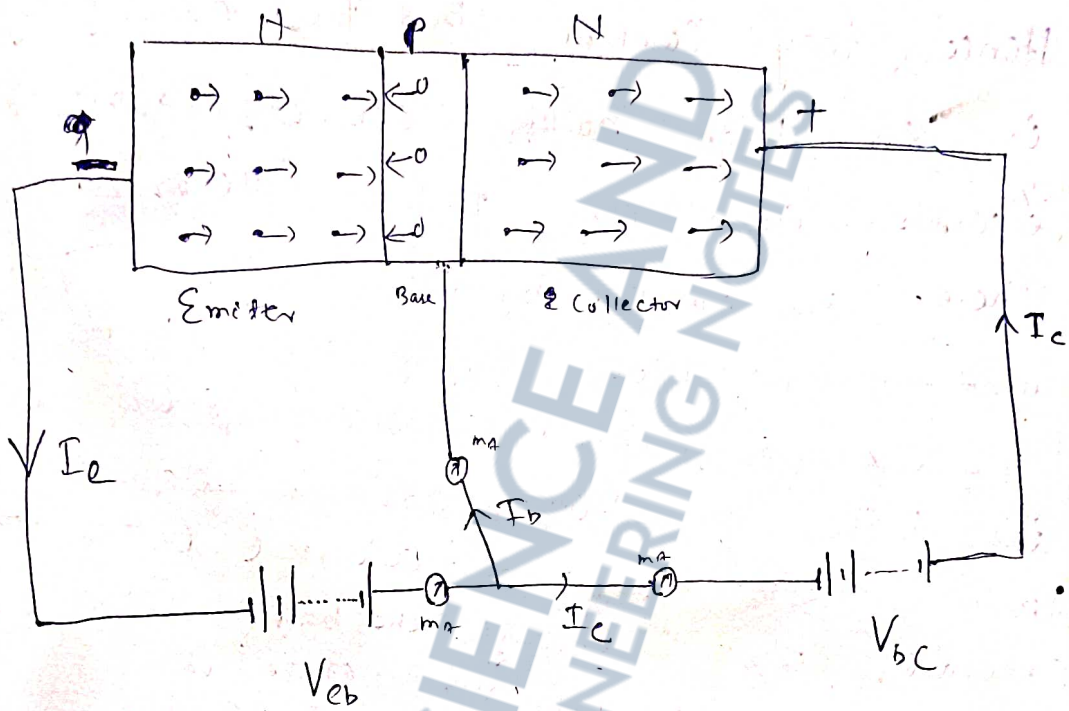
stage is quickly raised. A set of similar curves are obtained for different values of  $I_e$  as shown in graph next.

N-P-N transistor

A N-P-N transistor can be prepared out of a pure crystal of Ge or Si by doping the ends with pentavalent impurity atoms and the middle zone is made narrow compared to the size as shown in fig. As a result, no. of electrons becomes much higher

them no. of holes.

A N-P-N transistor thus formed can be regarded as a ~~N-P~~ N-P junction connected in series with P-N junction. To study the action of the N-P-N junction transistor the N end of the the N-P junction is



$$I_E = I_B + I_C$$

Connected to the -ve terminal of a battery  $V_{EB}$ . i.e. this N-P junction is forward biased. & this N end is called emitter.

The other N end of N-P-N transistor is connected to the +ve terminal of another battery  $V_{BC}$  and we called that N-end as collector. The middle zone is called base.

This base is connected to the -ve end of  $V_{EC}$  & +ve end of  $V_{EB}$ . Three milliammeters can be used to find the 3 currents  $I_E$ ,  $I_B$  &  $I_C$ .

Action  $\rightarrow$  Since the emitter is connected to the -ve terminal of the battery, the electrons of that end are pushed towards the base. The holes of the base are attracted towards the emitter. Hence they combine, but the no of holes is less than no of electrons. Majority (95%) electrons cross over to the Collector Zone. There are absorbed by the +ve plate. They

~~move through the conducting wire upto the junction point & some of them go towards the base while others go towards emitter.~~

About 5% electrons diverted to the external circuit through the base. The majority electrons come out of the -ve terminal of the emitter which moves towards emitter region. Then the process is repeated. Thus, current through the transistor is due to motion of electrons from collector to emitter. ~~current motion of electrons from collector to emitter.~~

From fig, we see that  
~~Sum of~~ emitter current = sum of base & collector current.  

$$I_e = I_b + I_c$$

The characteristic curves are similar to p-n-p transistor.

30.11.21

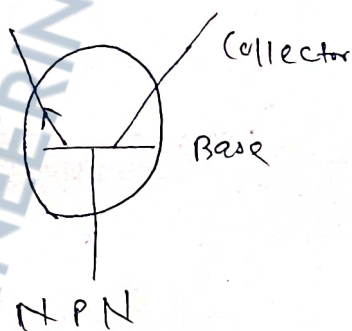
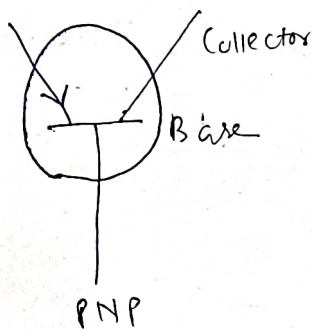
# Types of Transistor Connections

A transistor can be connected in 3 different ways.

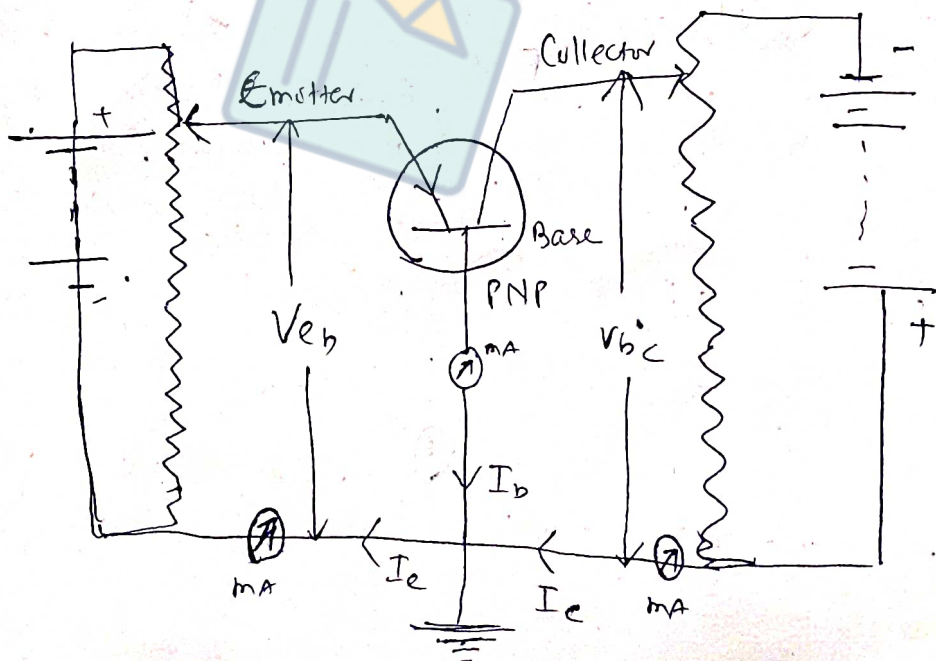
- (i) Common base
- (ii) Common emitter
- (iii) Common collector.

In all such connections, the emitter must be forward biased & collector must be reverse biased.

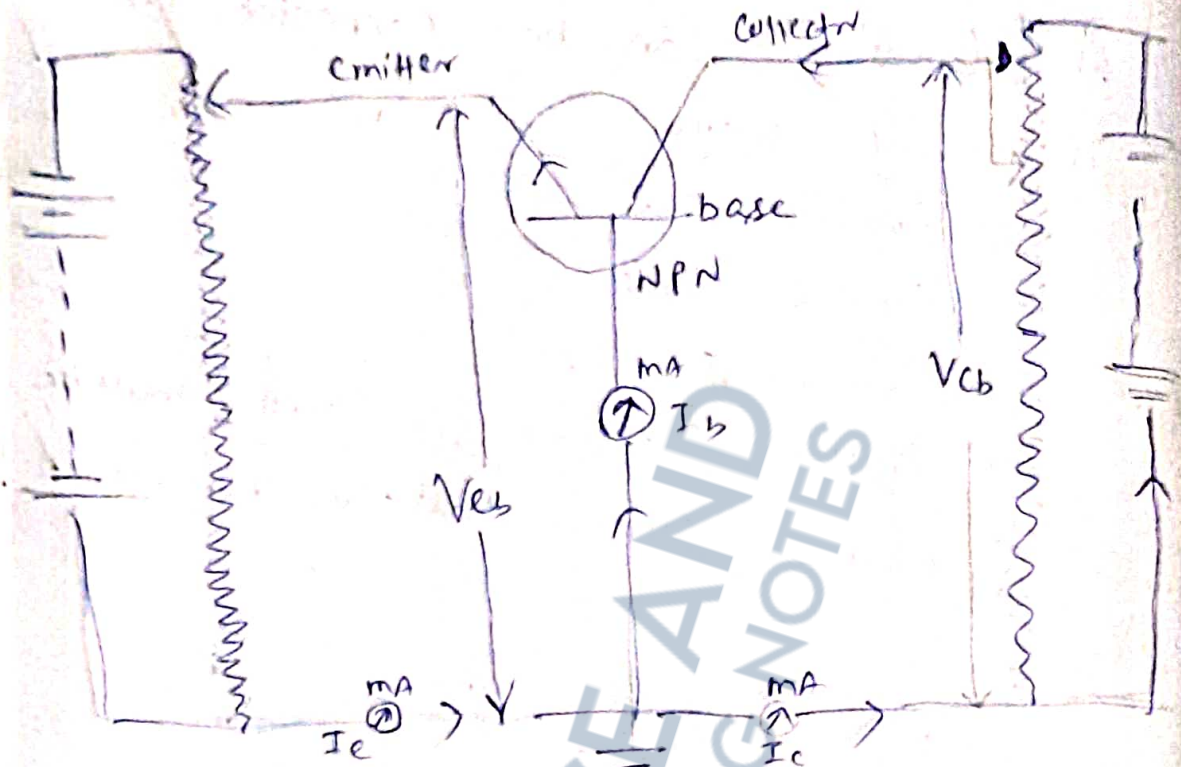
## Symbols of Transistors



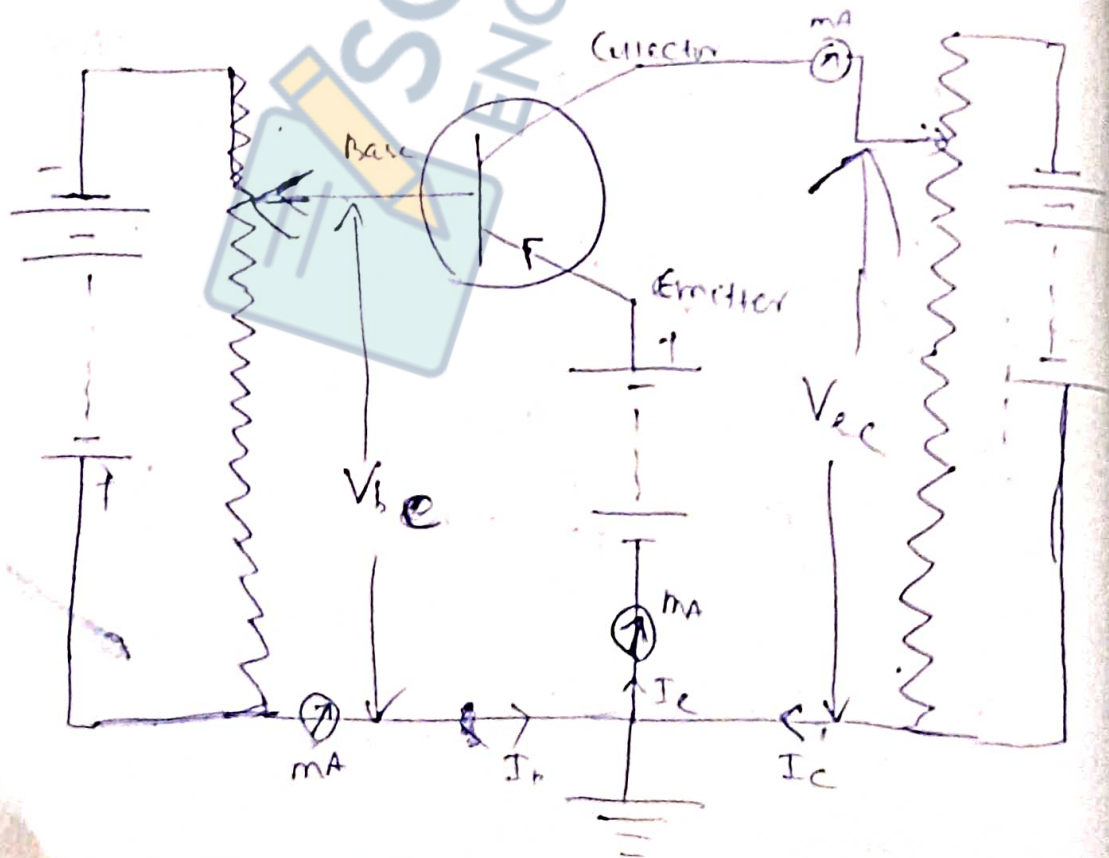
## 1(a) Common base PNP transistor connection



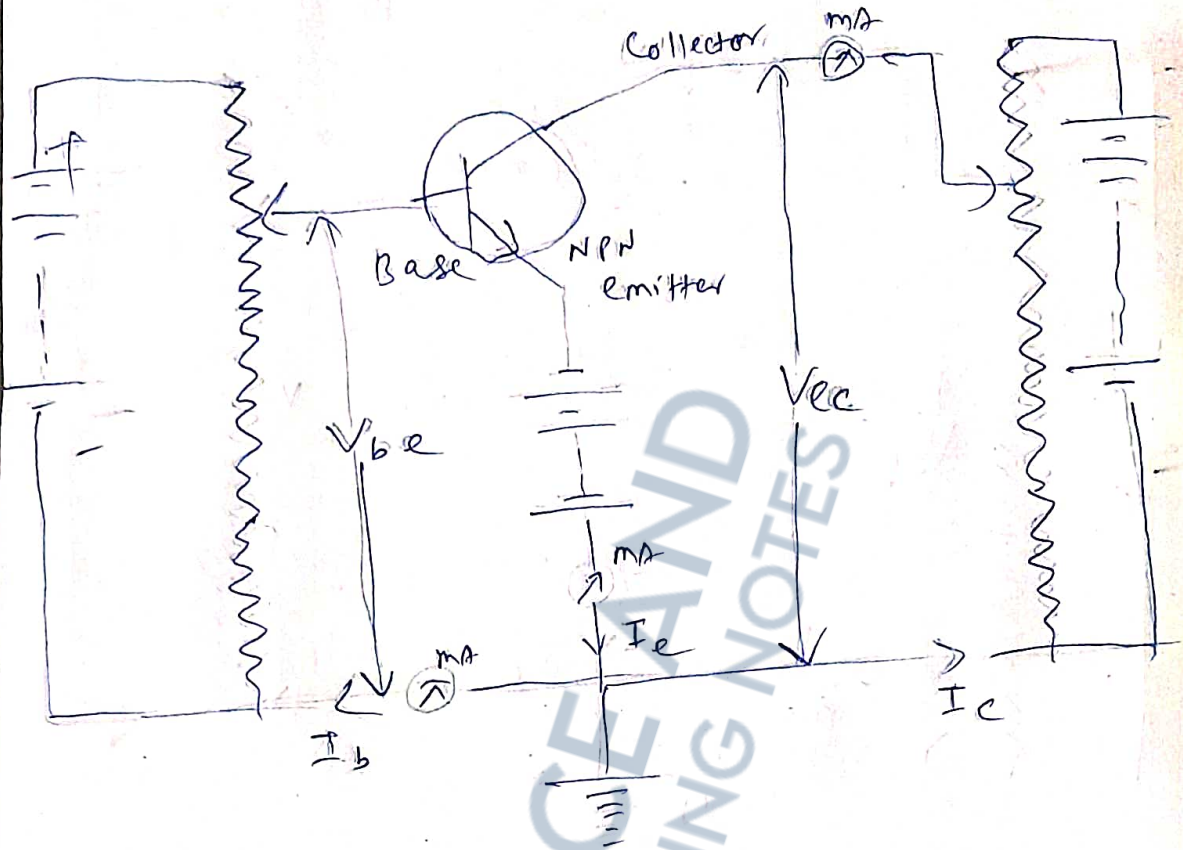
1 (b) Common base NPN transistor connection



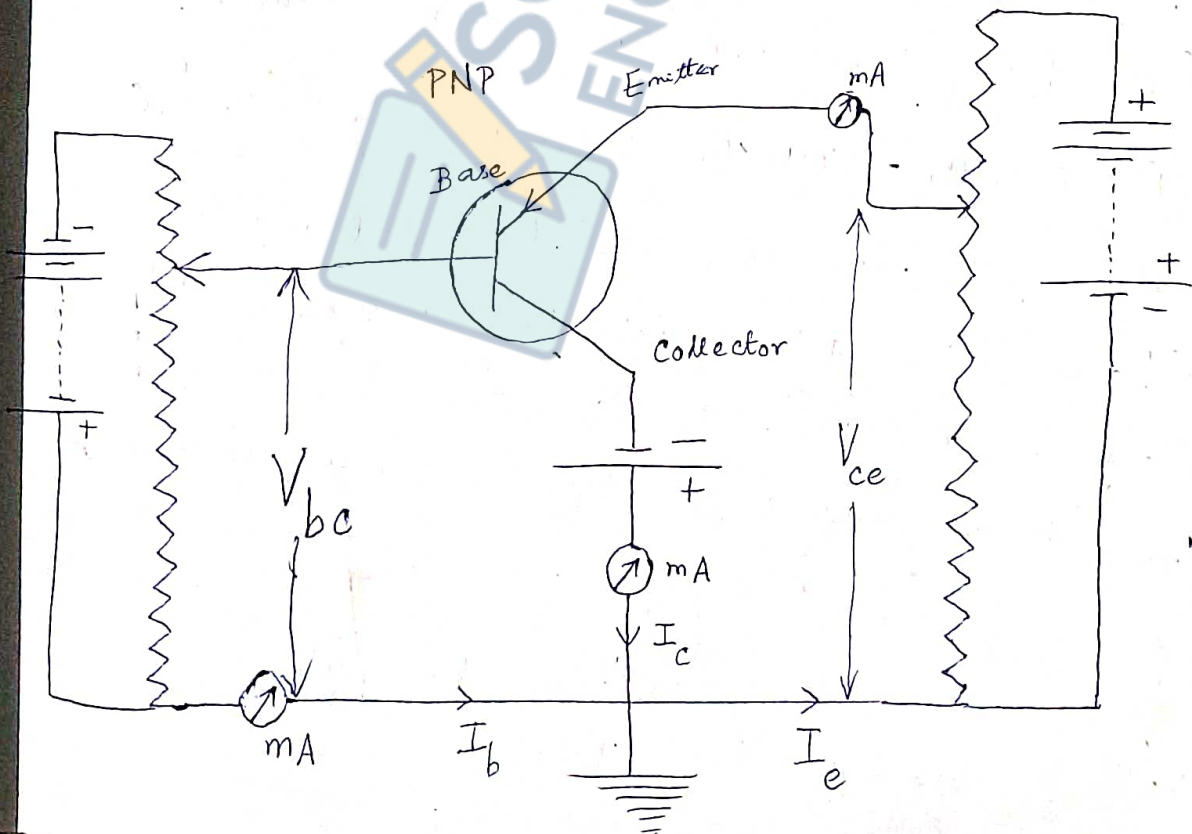
2 (a) Common emitter PNP transistor connection



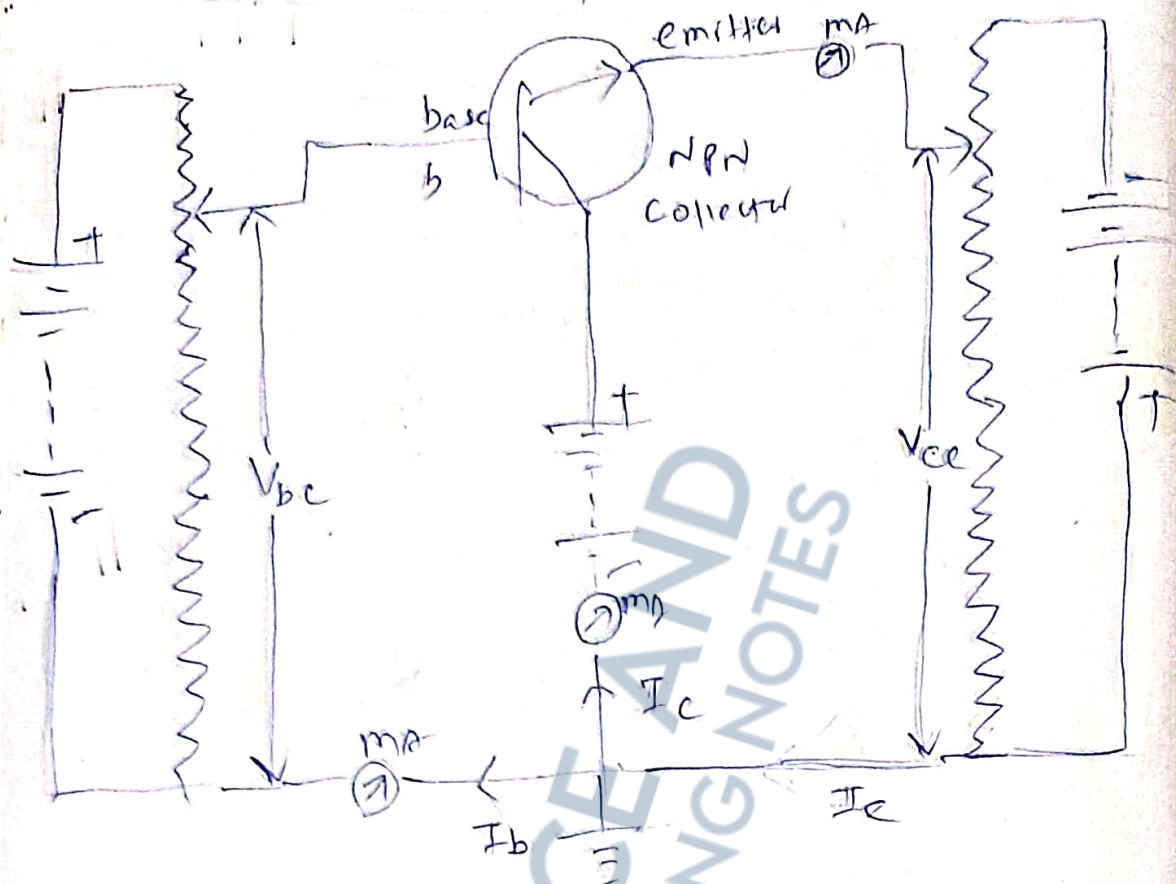
2(b) Common emitter NPN transistor connection



3(a) Common collector PNP transistor connection



### 3 (b) Common collector / NPN Connection.



### Transistor Parameters

There are 3 different ways of connecting the transistor and accordingly 3 constants for the same transistor can be defined.

#### (1) Emitter - Collector Current gain factor ( $\alpha$ )

It is defined as the ratio between change in the collector current to the change in the emitter current when the base collector voltage is kept constant.

$$\therefore \alpha = \left( \frac{\Delta I_c}{\Delta I_e} \right)_{V_{bc}}$$

It is obtained from the common base type or transistor connection.

Its value ranges from 0.9 to 0.99.

## ② Base - Collector Current gain factor ( $\beta$ )

It is defined as the ratio between the change in collector current to the change in base current, ~~when~~

$$\therefore \beta = \left( \frac{\Delta I_c}{\Delta I_b} \right)_{V_{CE}}$$

It is obtained from the common emitter type of transistor connection. Its value ranges from ~~20~~ 20 to 500.

## ③ Base - emitter current gain factor ( $\gamma$ )

It is defined as the ratio between the change in emitter current to the change in base current

$$\therefore \gamma = \frac{\Delta I_e}{\Delta I_b}$$

It is obtained from the common collector type of transistor connection.

## Relation between $\alpha$ & $\beta$

For all types of transistor connections,

We know that  $I_e = I_b + I_c$

$$\Rightarrow \Delta I_e = \Delta I_b + \Delta I_c$$



Dividing both the sides of the above eqn by  $\Delta I_e$ , we get

$$\Rightarrow I = \frac{\Delta I_b}{\Delta I_e} + \frac{\Delta I_c}{\Delta I_e} \quad \text{--- (i)}$$

$$= \frac{\Delta I_b}{\Delta I_c} \cdot \frac{\Delta I_c}{\Delta I_e} + \alpha$$

$$\Rightarrow 1 - \alpha = \frac{1}{\beta} \cdot \alpha$$

$$\Rightarrow \boxed{\beta = \frac{\alpha}{1 - \alpha}}$$

Relation between  $\alpha$  &  $\gamma$

From eqn (i), we get

$$1 = \frac{1}{\gamma} + \alpha$$

$$\Rightarrow 1 - \alpha = \frac{1}{\gamma}$$

$$\Rightarrow \gamma = \frac{1}{1 - \alpha}$$

Relation Among  $\alpha$ ,  $\beta$ ,  $\gamma$

$$\text{Now } \frac{\beta}{\gamma} = \frac{\frac{\alpha}{1 - \alpha}}{\frac{1}{1 - \alpha}} = \alpha$$

$$\Rightarrow \boxed{\beta = \alpha \gamma}$$

## Problems

1. In a common base connection,  
 $I_e = 1.5 \text{ mA}$ . Find  $I_b$  &  $I_c$  when  
the current amplification factor is 0.94.

Sol<sup>n</sup> : Given  $\alpha = 0.94$

$$\Rightarrow \frac{\Delta I_c}{\Delta I_e} = 0.94$$

$$\Rightarrow \frac{I_c - 0}{I_e - 0} = 0.94$$

$$\Rightarrow \frac{I_c}{1.5} = 0.94$$

$$\Rightarrow I_c = (1.5 \times 0.94)$$

$$= 1.41 \text{ mA}$$

$$= 1.41 \text{ mA}$$

$$I_b = I_e - I_c = 1.5 - 1.41 = 0.09 \text{ mA}$$

2. Find  $I_e$  when  $\beta = 40$  &  $I_b = 10 \mu\text{A}$

$$\beta = \frac{\Delta I_c}{\Delta I_b}$$

$$\Rightarrow \beta = \frac{I_c - 0}{I_b - 0} = \frac{I_c}{10 \mu\text{A}}$$

$$\Rightarrow 40 = \frac{I_c}{10 \mu\text{A}}$$

$$\Rightarrow I_c = 400 \mu\text{A}$$

$$I_e = I_b + I_c = 10 + 400 = 410 \mu\text{A}$$

225  
2310  
204  
94  
15  
211 0  
138

(3) Find the value of  $\beta$  &  $\gamma$  when

(i)  $\alpha = 0.9$

(ii)  $\alpha = 0.98$

(iii)  $\alpha = 0.99$

Ans  $\therefore (9, 10), (49, 50), (99, 100)$

(i)  $\beta = \frac{\alpha}{1-\alpha} = \frac{0.9}{1-0.9} = \frac{0.9}{0.1} = 9$

$\gamma = \frac{\beta}{\alpha} = \frac{9}{0.9} = 10$

(ii)  $\beta = \frac{\alpha}{1-\alpha} = \frac{0.98}{1-0.98} = \frac{0.98}{0.02} = 49$

$\gamma = \frac{\beta}{\alpha} = \frac{49}{0.98} = 50$

(iii)  $\beta = \frac{\alpha}{1-\alpha} = \frac{0.99}{1-0.99} = \frac{0.99}{0.01} = 99$

$\gamma = \frac{\beta}{\alpha} = \frac{99}{0.99} = 100$

(4) For a transistor in C-Emitter Connection  
the voltage drop across a collector load  
of  $1\text{K}\Omega$  is 1.5 volt. The value  
of  $\alpha = 0.96$ . Find the value  $\beta, I_c, I_b$   
&  $I_e$ .

Ans: 24, 1.5 mA, 0.06 mA, 1.56 mA

Ans: In Common emitter Connection  
Voltage = 1.5 volt

$$R_L = 1 \text{ k}\Omega$$

$$I_C = \frac{V}{R} = \frac{1.5}{1 \times 10^3} = 1.5 \times 10^{-3} = 1.5 \text{ mA (Ans)}$$

$$\alpha = 0.96$$

$$\Rightarrow \frac{\Delta I_C}{\Delta I_E} = 0.96$$

$$\Rightarrow \frac{I_C - 0}{I_E - 0} = 0.96$$

$$\Rightarrow \frac{1.5}{I_E} = 0.96$$

$$\Rightarrow I_E = \frac{1.5}{0.96} = \frac{150}{96} = 1.56 \text{ mA (Ans)}$$

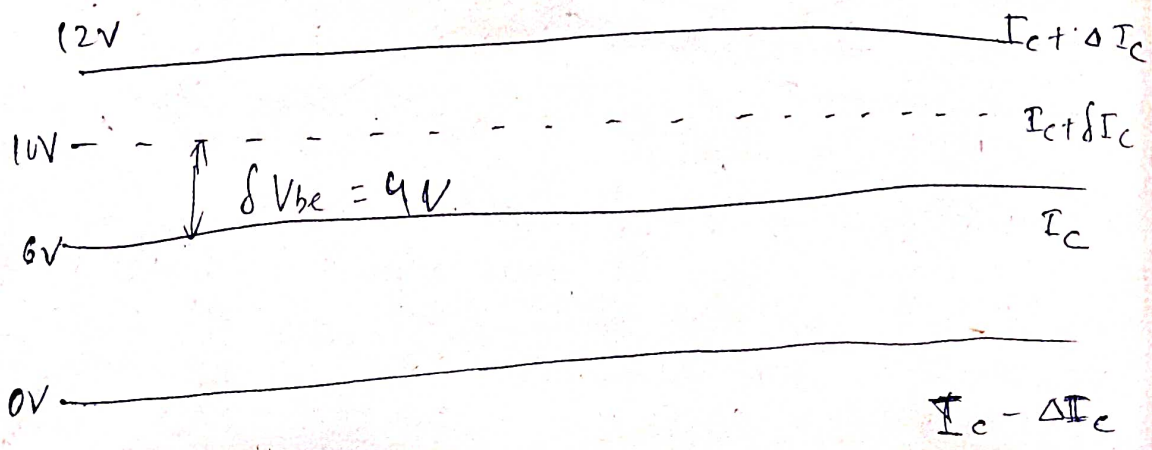
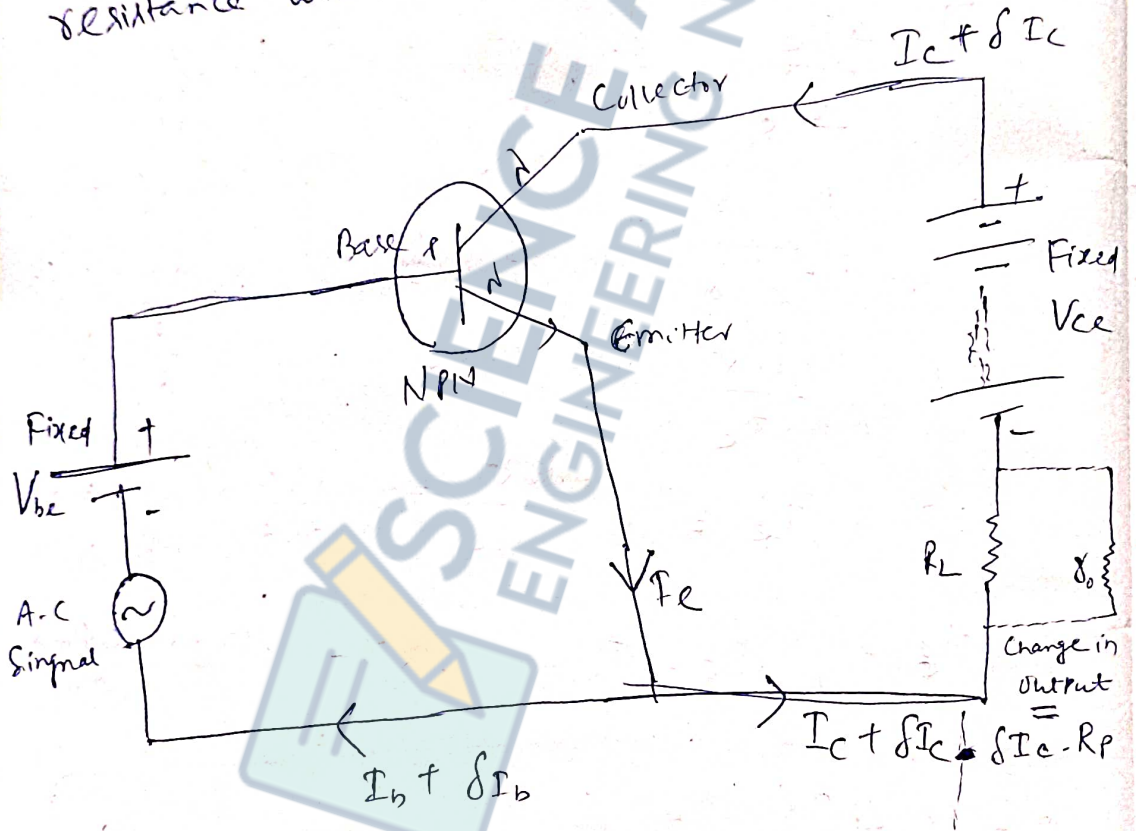
$$I_B = I_E - I_C = 1.56 - 1.5 = 0.06 \text{ mA (Ans)}$$

$$\beta = \frac{\Delta I_C}{\Delta I_B} = \frac{I_C - 0}{I_B - 0} = \frac{1.5}{0.06} = \frac{150}{6} = 24$$

$$\beta = \frac{\alpha}{1 - \alpha} = \frac{0.96}{1 - 0.96} = \frac{0.96}{0.04} = 24 \text{ (Ans)}$$

# Transistor as Amplifier

Let's take the case of an NPN transistor connected as common emitter in the circuit. Suppose, there is no A.C signal in the circuit. Due to fixed  $V_{be}$  & fixed  $V_{ce}$ , the currents in different branches will be fixed. Hence the output voltage derived from the 2 ends of the load resistance will be also fixed like  $I_c \cdot R_L$ .



where  $R_p \equiv$  Equivalent resistance between  $R_L$  &  $r_o$ .

$r_o =$  Resistance of the output circuit.

$$\frac{1}{R_p} = \frac{1}{R_L} + \frac{1}{r_o}$$

$$\Rightarrow R_p = \frac{R_L r_o}{R_L + r_o}$$

Due to the introduction of the alternating small voltage (Signal) represented by  $e = \sin \omega t$  (say), the net voltage between the base and emitter will change. To maintain the base at some +ve voltage, we must have a fixed  $V_{be} = 0.7$  Volt (min value)

At any instant of time, suppose the net voltage between the base and the emitter be 10 Volt.

$$\Rightarrow \Delta V_{be} = 9 \text{ Volt}$$

Due to  $\Delta V_{be}$ , there will be a change in the base current as well as the collector current.

If  $\Delta I_c$  be the change in the collector current, then the change in the output

$$= \delta I_c \cdot R_c$$

$A_v$  = Voltage Amplification by the transistor

$$= \frac{\text{Change in the output}}{\text{Change in the input}}$$

$$= \frac{\delta I_c \cdot R_c}{\delta I_b \cdot r_i}$$

where  $\delta V_{be}$  = Change in the input voltage  
 $= \delta I_b \cdot r_i$

$r_i$  = Total resistance between the base & emitter.

$$A_v = \left( \frac{\delta I_c}{\delta I_b} \right) \cdot \left( \frac{R_c \cdot r_o}{R_c + r_o} \right) \cdot \frac{1}{r_i}$$

$$= \beta \cdot \left( \frac{R_c \cdot r_o}{R_c + r_o} \right) \cdot \frac{1}{r_i}$$

where  $\beta$  = ~~Current~~ Base-Collector Current gain factor

$$= \frac{\Delta I_c}{\Delta I_b}$$

$$\therefore A_v = \frac{\beta \cdot r_o}{r_i} \cdot \left( \frac{R_c}{R_c + r_o} \right)$$

$$A_v = \beta' \cdot \frac{R_L}{r_o + R_L}$$

where  $\beta' = \frac{\beta \cdot r_o}{r_i}$  = Dimensionless quantity.

This expression for the voltage amplification is very much similar to the corresponding expression due to a triode as an amplifier.

$$\left( A_v = \frac{\mu R_L}{r_p + R_L} \right)$$

In multistage amplifiers the output voltage of one amplifier is used as the signal voltage for the other amplifier. As a result the amplification becomes very large - we can

write  $(A_v)_{net} = (A_v)_1 \times (A_v)_2 \times (A_v)_3 \dots$

### Problems

In a common-emitter amplifier circuit

$\beta = 60$  , Input resistance =  $500 \Omega$  &

Output resistance =  $500 \Omega$ .

What is the voltage gain  $(A_v)$  if  $R_L = 10 k\Omega$

(Ans = 40)



Ans:

$$A_V = \frac{B \cdot 10^3}{\gamma_1} \left( \frac{R_L}{R_L + R_C} \right)$$

$$= \frac{60 \times 10^3}{5000} \left( \frac{10 \times 10^3}{5000 + 10 \times 10^3} \right)$$

$$= 600 \left( \frac{10^3}{15 \times 10^3} \left( \frac{10}{5+10} \right) \right)$$

$$= 600 \times \frac{10}{15}$$

$$= 400$$

### Binding in solids

Compared to gaseous and liquid states, the atoms and molecules in a solid are more closely packed and they are held together by strong forces of attraction. These interatomic forces are basically electrostatic in nature. When atoms come closer and finally unite to form molecules, their electrons rearrange themselves in such a way as to achieve stable configuration. This <sup>re</sup>arrangement of electrons gives rise to different types of bonds which hold the atoms together in a solid state. The ~~principle~~ principal

types of bonds in solid are,

1. Ionic or Electrovalent bonds

Ex: NaCl, MgO, CaF<sub>2</sub>

2. Covalent ~~bonds~~ or Homopolar bonds

Ex: C, Si, Ge

### 3. Metallic bonds

Ex = Na, Al, Cu, Ag

### 4. Molecular bonds due to Vander-waals

force Ex → Solid Argon, Solid Kr and Solid Xe.

### ① Ionic bonds

This type of bonds are mainly formed in inorganic compounds like NaCl, KCl etc and never in pure elements. This type of bonds are very strong and develop between two atoms out of which one gives out an electron and other accepts it. i.e. There is complete transfer of electrons from one atom to the other.

For example, let us discuss the formation of NaCl molecule. The electronic configuration of Na atom is  $1s^2 2s^2 2p^6 3s^1$ . It can lose one electron to become  $Na^+$  ion. The electron configuration of Cl is  $1s^2 2s^2 2p^6 3s^2 3p^5$ . Hence Cl can accept the electron donated by Na and it will become  $Cl^-$  ion.

Now  $Na^+$  and  $Cl^-$  will attract each other by strong electrostatic force of attraction (due to Coulomb's law)

## Properties of Ionic Solids →

Solids having ionic bonds have the following characteristic properties.

1. Crystal Structure ∴ X-ray diffraction study has revealed that such solids have well defined crystalline structure. They are hard and brittle.

2. Melting & boiling points → Since, powerful electrostatic forces exist between the ions, considerable energy is required to overcome these forces. Hence ionic solids have high melting & boiling points.

3. Electrical Conductivity → Since free electrons are not available, ionic solids are insulators. However, in the fused state, some of the ions move and conductivity increases with the increase of temp.

4. Solubility →

Ionic solids are readily soluble in solvents like water & liquid ammonia.

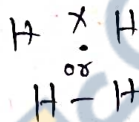
5. Ionic reaction → Ionic reactions are practically instantaneous. For ex, when  $\text{AgNO}_3$  soln is added to  $\text{NaCl}$  or  $\text{BaCl}_2$  etc, white ppt of  $\text{AgCl}$  is immediately formed.

# Covalent bond →

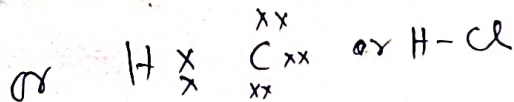
A Covalent bond is formed when two similar or dissimilar atoms achieve stability by sharing valence electrons between themselves.

The shared electrons become the common electrons of both the atoms. By this, the atoms achieve the noble gas configuration.

Ex-1  $H_2$  molecule formation.

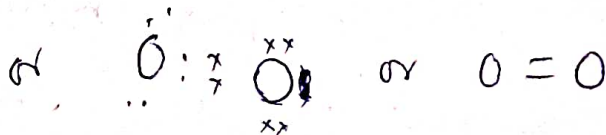
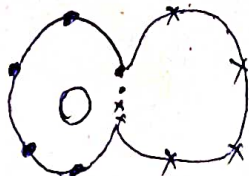


Ex-2 →

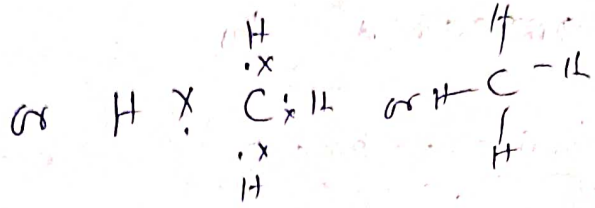
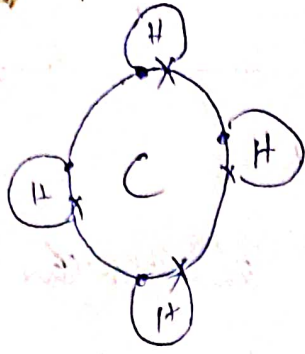


Ex-2

$O_2$  molecule formation



## EX-1 CH<sub>4</sub> molecule formation



The following additional points are to be noted.

1. Attainment of noble gas configuration is not essential.
2. The two electrons forming a pair must have opposite spin.
3. A covalent bond may be either polar or non polar depending on the fact whether the electron pair is shared unequally or equally between two atoms.

### Properties of covalent bond

The characteristic properties are discussed below.

(1) Physical state → Covalent compounds may be solids, or gases. For ex → Cl<sub>2</sub> is gas, Br<sub>2</sub> is liquid & I<sub>2</sub> is solid at normal temp & pressure.

(2) Crystal structures → The following three types of covalent compounds are found.

(a) Covalent solids like sulphur and

Iodine possess molecules or atoms that are held together by weak forces. Hence they are soft & easily fusible.

(b) In the case of Diamond, the atoms are found to form giant molecules. Each 'C' atom forms 4 covalent bonds with four neighbouring atoms and these bonds form the corners of a regular tetrahedron. ( $\theta = 109.5^\circ$  = angle between two bonds)

(c) In the case of graphite, the 'C' atoms forms regular hexagons & layers of atoms are found with weak forces ~~are found~~ between adjacent layers. This is the cause of softness and lubricating action of graphite.

### (3) Melting & boiling points →

Since covalent bonds are ~~not~~ not that strong as the ionic bonds, covalent compounds have comparatively low M.P & B.P.

### (4) Electrical conductivity →

All covalent ~~is~~ solids are basically insulators because free electrons are not available for conduction process. However, metals like Ge & Si are semiconductors & their conductivity increases with the increase of temp & introduction of impurity

Returns

### (5) Solubility →

Covalent Compounds are soluble in nonpolar solvents like Benzene & CCl<sub>4</sub>.

### (3) Metallic bond →

When atoms or molecules unite to form a metal, the outermost electron of each atom or molecule gets detached from the parent atom and these electrons form a cloud of -ve charge called sea of electrons. The ions are found at regular lattice points. The electron cloud is present in between two +ve ions & is ~~attached~~ attracted equally by the ions. The metallic bond is comparatively weaker than the ionic and covalent bonds, it is sometimes called unsaturated covalent bond.



### Properties of Metal →

(1) They have crystalline structure.

(2) Since free electrons are available in plenty, the electrical conductivity is extremely high even at low temp.

(3) Since the metallic bonds are not very strong, metals have moderate to high melting temperatures

(4) Metals have high thermal conductivity due to electrons taking part in the conduction process.

(5) Since free electrons in a metal absorb light energy, all metals are opaque to light.

#### (4) Molecular bonds →

This type of bonds occur in those elements or compounds which have noble gas configuration.

Due to van der Waals forces, the ~~molecules~~ molecules attract each other with a weak force.

Only at very low temp, such solids are formed. Two such molecules have many attractive & repulsive forces, yet calculation

shows that the net force is attractive (as shown by van der Waal.)

#### Properties of molecular solids →

(1) Molecular structures can be both crystalline and non crystalline.

(2) These solids have low densities.



③. Because of weak molecular bonds, such solids have low melting points.

4. Since free electrons are not available, they are good insulators.

⑤. They are usually transparent in nature.

⑥. They are soluble in both polar & non-polar ~~the~~ liquids.

## Crystal Structure 26. 12. 21

Solids are mainly of two types

1. Crystalline
2. Amorphous

X-ray diffraction by some solids clearly shows that ~~there~~ there is ~~fixed~~ periodic arrangement of atoms or molecules & these are called Crystals.

Crystal Structure = Lattice + Basis

The internal design or arrangement scheme is called lattice, whereas

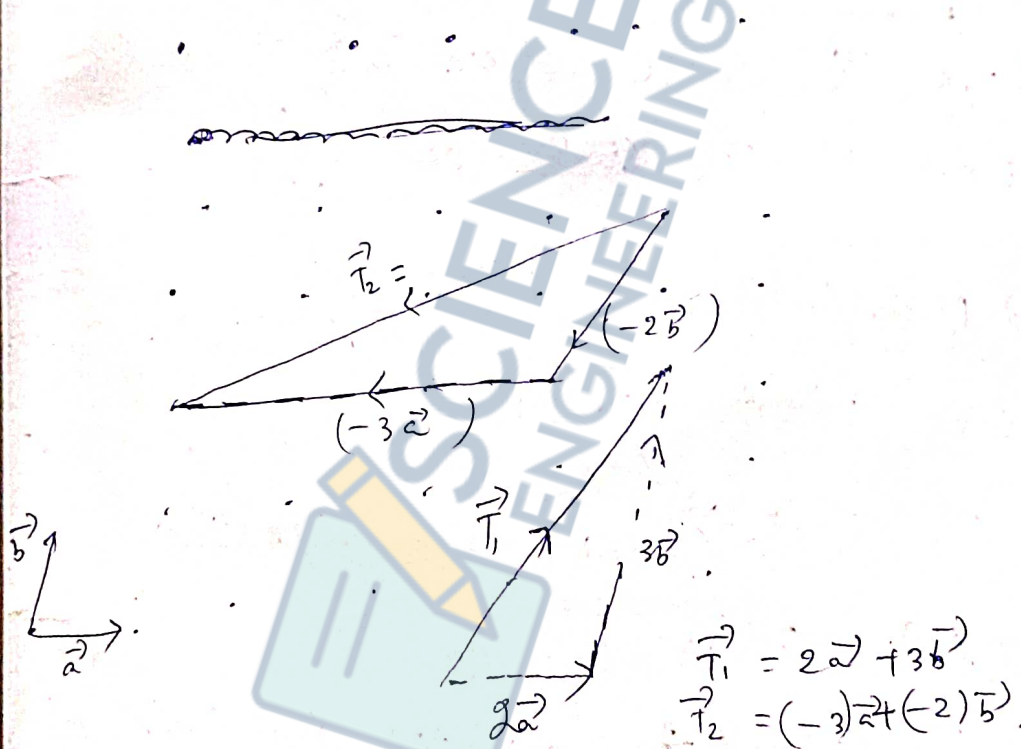
the atom or groups of atoms placed at regular lattice points is called basis.

EX → The NaCl molecule is found at the corner of a cube & also at the 6 face centres of NaCl crystal.

In the case of amorphous solids there is no such orderliness found as in the case of crystals.

EX-1) Glass.

In this regard liquids can be compared with amorphous solids. But they (liquids) differ in having a definite boiling point & freezing point whereas amorphous solids don't possess such temperatures.



### Primitive translation vectors

In three ~~dimensional~~ dimension, there are three primitive translation vectors  $\vec{a}, \vec{b}, \vec{c}$ . Crystal structure is confirmed if the vector joining any lattice point to another lattice point, called translation vector is of the form

$$\vec{T} = n_1 \vec{a} + n_2 \vec{b} + n_3 \vec{c}$$

where  $n_1, n_2, n_3$  are integers (both +ve & -ve). In the diagram two dimensional translation vector have been shown.

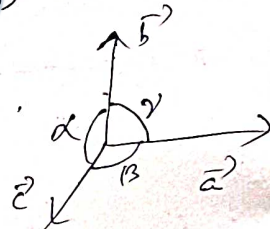
## Types of Crystal Structures

Mainly there are seven types of crystal structures.

① Cubic,  $a = b = c, \alpha = \beta, \gamma = 90^\circ$

where  $\alpha =$  angle between  $\vec{b}$  &  $\vec{c}$ ,

Ex  $\rightarrow$  NaCl, CaF<sub>2</sub> etc.



② Monoclinic  $\rightarrow a \neq b \neq c, \alpha = \beta = 90^\circ \neq \gamma$

Ex: FeSO<sub>4</sub>, Na<sub>2</sub>SO<sub>4</sub>

③ Triclinic  $\rightarrow a \neq b \neq c, \alpha \neq \beta \neq \gamma \neq 90^\circ$

Ex  $\rightarrow$  CuSO<sub>4</sub>, K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>

④ Tetragonal  $\rightarrow a = b \neq c, \alpha = \beta = \gamma = 90^\circ$

Ex  $\rightarrow$  ~~Na~~ NiSO<sub>4</sub>, SnO<sub>2</sub>

⑤ Orthorhombic  $\rightarrow a \neq b \neq c, \alpha = \beta = \gamma = 90^\circ$

Ex: BaSO<sub>4</sub>, MgSO<sub>4</sub>

⑥ Trigonal  $a = b = c, \alpha = \beta = \gamma \neq 90^\circ$

Ex: CaSO<sub>4</sub>

7. Hexagonal :  $a = b \neq c, \alpha = \beta = 90^\circ, \gamma = 120^\circ$

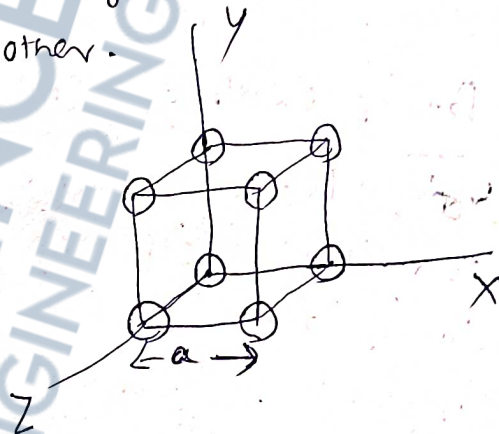
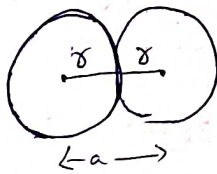
Ex  $\rightarrow$   $\text{SiO}_2, \text{AgI}$

Cubic Structure  $\rightarrow$

There are 8 atoms at the 8 corners points of a unit cube of side length 'a'. But each atom is shared by 8 such cubes. Hence effectively there is only one atom in the unit cell. It is called

Simple cubic structure.

For close packing of spheres, they should touch each other.



$\therefore$  atomic radius  $= r = \frac{a}{2}$

Packing fraction is defined as the ratio of the volume actually occupied by atom or atoms present in a unit cell to the volume of the unit cell.

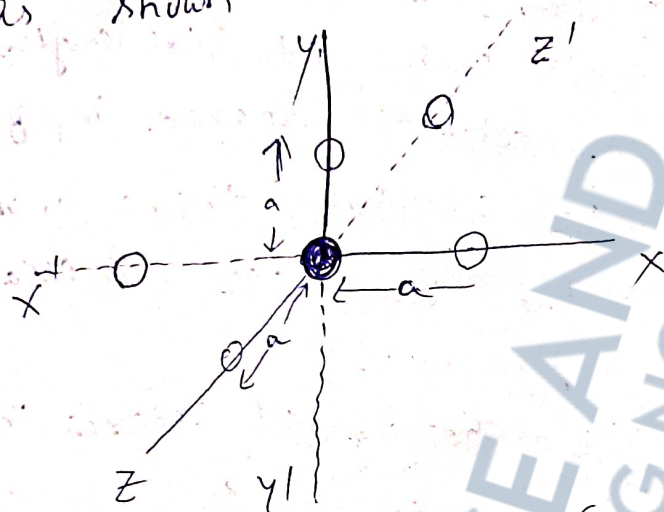
For S.C structure,  $f = \frac{\frac{4}{3} \pi r^3}{a^3}$

$$= \frac{\frac{4}{3} \cdot \frac{22}{7} \cdot \left(\frac{a}{2}\right)^3}{a^3} = \frac{88}{21} \times \frac{1}{8} = \frac{11}{21} = 0.523$$

$= 52.3\%$ . This shows that a lot

of empty space exist inside the solid.

Co-ordination Number is defined as the number of nearest neighbours of an atom. It is 6 for SC structure as shown below.



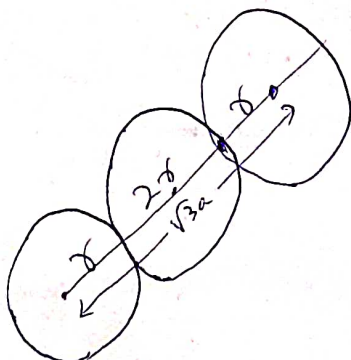
### Body Centred Cubic (BCC)

There is one atom at the centre of each unit cell in addition to the 8 atoms at the 8 corners points, a BCC crystal is formed.

Number of atoms in a unit cell

$$= \left(\frac{1}{8} \times 8\right) + 1 = 2$$

For closest packing of spheres, they must touch along the diagonal.



packing fraction

$$= f = \frac{\text{Volume of two atoms in the unit cell}}{\text{Volume of the unit cell}}$$

$$= \frac{2 \cdot \frac{4}{3} \pi r^3}{a^3} = \frac{2 \cdot \frac{4}{3} \cdot \pi \cdot \left(\frac{\sqrt{3}a}{4}\right)^3}{a^3}$$

$$= \frac{\frac{8}{3} \cdot \pi \cdot \frac{3\sqrt{3}a^3}{4 \times 16}}{a^3} = \frac{2 \times 22}{7} \times \frac{1.732}{16}$$

$$= 0.6804 = 68.04\%$$

Co-ordination number  $\rightarrow$  (8) Because the body centres are nearest to the atom at any corner point  $\left(\frac{\sqrt{3}a}{2} = 0.866a\right)$

Face Centred Cubic (FCC)  $\rightarrow$

In addition to the 8 atoms at the 8 corner points, there are 6 atoms present at the 6 face centres.

Hence no. of atoms on the unit cell

$$= \left(\frac{1}{8} \cdot 8\right) + \left(\frac{1}{2} \cdot 6\right) = 1 + 3 = 4$$

For closest packing of spheres, they must touch along the face diagonal.

$$\therefore 4r = \sqrt{2} a$$

$$\therefore \text{atomic radius} = r = \frac{\sqrt{2}}{4} a$$

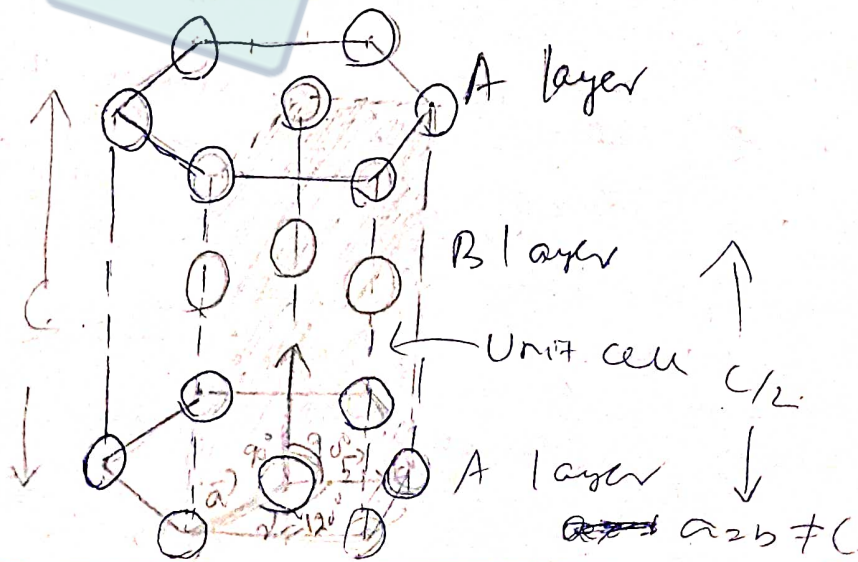
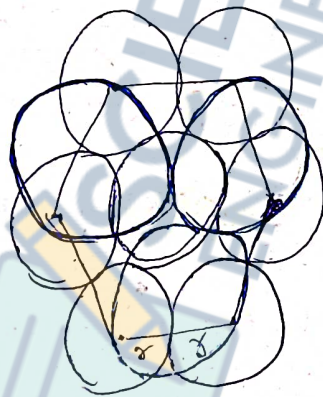
$$\text{Packing fraction} = \frac{\text{Volume occupied by 4 atoms on the unit cell}}{\text{Volume of unit cell}}$$

$$= \frac{4 \times \frac{4}{3} \pi r^3}{a^3} = \frac{4 \times \frac{4}{3} \pi \left(\frac{\sqrt{2}a}{4}\right)^3}{a^3}$$

$$= \frac{11 \times 1.414}{21} = 74.06 = 74.06 \%$$

Co-ordination no = 12 (Because the face centred atoms are the nearest neighbours)  
Hexagonal Close packed structure (HCP)

Here the arrangement of atoms is like AB, AB, AB --- The unit cell (where the repetition gives the crystal structure) is a rhombus as shown in the 2nd diagram.



$$\begin{aligned} \alpha &= 90^\circ \\ &= \beta \\ \gamma &= 120^\circ, \infty \end{aligned}$$

$$a \neq b \neq c$$

Atomic radius =  $r = \frac{a}{2}$

Volume of the unit cell = Base area  $\times$  Height

$$= \left( \frac{\sqrt{3}}{4} a^2 - a \right) \cdot c$$

$$= \frac{\sqrt{3}}{2} a^2 c$$

Number of atoms in the unit cell

$$= \left( \frac{1}{8} \times 8 \right) + 1 = 1 + 1 = 2$$

$\therefore$  Packing fraction =  $f = \frac{\text{Volume of two atoms in the unit cell}}{\text{Volume of the unit cell}}$

$$= \frac{\frac{4}{3} \pi r^3 \times 2}{\frac{\sqrt{3}}{2} a^2 c}$$

$$= \frac{\frac{4}{3} \pi \left( \frac{a}{2} \right)^3 \cdot 2}{\left( \frac{\sqrt{3}}{2} \right) a^2 c}$$

$$= \frac{\frac{8}{3} \cdot \frac{11}{7} \cdot \frac{a^3}{8}}{\frac{\sqrt{3}}{2} a^2 c}$$

$$= \frac{44a}{21} \cdot \frac{1}{\sqrt{3}c} = \frac{44}{21\sqrt{3}} \times \frac{a}{c}$$

But  $\frac{c}{a}$  ratio =  $\frac{2\sqrt{2}}{\sqrt{3}}$

$\therefore f =$  Packing fraction =  $\frac{44}{21\sqrt{3}} \times \frac{\sqrt{3}}{2\sqrt{2}}$

$$= \frac{11}{21\sqrt{2}} = 11 \times \frac{0.707}{21}$$

$$= 0.7406 = 74.06\%$$

Problems  $\rightarrow$

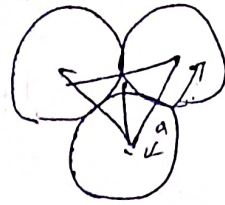
1. Prove that  $\frac{c}{a} = \frac{2\sqrt{2}}{\sqrt{3}}$  for hcp

Structure. Ans. Height of the ~~the~~ triangle h



$$\leq \frac{\sqrt{3}}{2} a$$

$$x = \frac{2}{3} h = \frac{2}{3} \cdot \frac{\sqrt{3}}{2} a = \frac{a}{\sqrt{3}}$$



Since B layer of atoms touch the A-layer on the sides, we can apply Pythagoras theorem.



$$a^2 = \left(\frac{c}{2}\right)^2 + x^2$$

$$\Rightarrow a^2 = \frac{c^2}{4} + \left(\frac{a}{\sqrt{3}}\right)^2$$

$$\Rightarrow a^2 = \frac{c^2}{4} + \frac{a^2}{3}$$

$$\Rightarrow a^2 - \frac{a^2}{3} = \frac{c^2}{4}$$

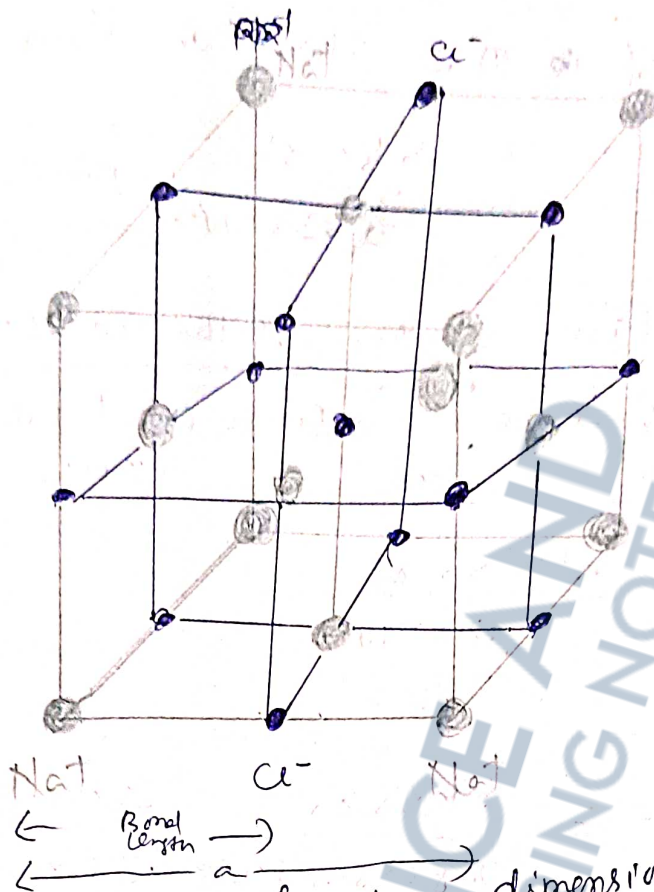
$$\Rightarrow \frac{2a^2}{3} = \frac{c^2}{4}$$

$$\Rightarrow \frac{\sqrt{2} a}{\sqrt{3}} = \frac{c}{2}$$

$$\Rightarrow \frac{c}{a} = \frac{2\sqrt{2}}{\sqrt{3}} = \frac{2 \times 1.414}{1.732} = 1.6328$$

$$\Rightarrow \frac{a}{c} = \frac{\sqrt{3}}{2\sqrt{2}}$$

# NaCl Structure →



Calculation of the dimensions of a unit

Cell →

The dimension of the unit cell or the interatomic distance in a crystal lattice can be calculated from the knowledge of molecular weight ( $M$ ) or atomic weight ( $A$ ), Avogadro Number ( $N_0$ ), density of the material ( $\rho$ ) and number of atoms or molecules in the unit cell ( $n$ )

Ex → 1 BCC Crystal

In the  $\alpha$ -iron crystal, there are two atoms of iron in the unit cell. Atomic weight =  $A = 55.85$ , density of iron =  $7.86 \text{ gm/cc}$   
 Number of atoms of iron present in

$$55.85 \text{ gm} = M_u = 6.023 \times 10^{23}$$

$$\therefore \text{Mass of one atom or ion} = \frac{55.85}{6.023 \times 10^{23}} \text{ gm}$$

$$\text{Density} = \frac{\text{Mass}}{\text{Volume}} = \frac{\text{Mass of the unit cell}}{\text{Volume of the unit cell}}$$

$$7.86 = \frac{2 \times 55.85}{a^3 \times 6.023 \times 10^{23}}$$

$$\Rightarrow a^3 = \frac{2 \times 55.85}{7.86 \times 6.023 \times 10^{23}} = \left( \frac{20 \times 55.85}{7.86 \times 6.023} \right) \times 10^{-24} \text{ cm}^3$$

$$\Rightarrow a = \left( \frac{20 \times 55.85}{7.86 \times 6.023} \right)^{\frac{1}{3}} \times 10^{-8} \text{ cm}$$

$$\therefore a = 2.87 \times 10^{-8} \text{ cm} = 2.87 \text{ \AA}$$

Formula

$$f = \frac{n \cdot A}{\frac{N_0}{a^3}}$$

$$\Rightarrow \boxed{a^3 f = \frac{nA}{N_0}}$$

Example  $\rightarrow 2$

Molecular weight of  $\text{NaCl} = 23 + 35.5 = 58.5$

Number of molecules in the unit cell

$$n = 4$$

$$\text{Density} = f = 2.18 \text{ gm/cc}$$

Number of molecules of Hacc present  
in 58.5 gm =  $6.023 \times 10^{23}$

$$\text{Mass of one molecule of Hacc} = \frac{58.5}{6.023 \times 10^{23}} \text{ gm.}$$

$$\text{Density} = \frac{\text{Mass}}{\text{Volume}} = \frac{\text{Mass of the unit cell}}{\text{Volume of the unit cell.}}$$

$$2.18 = \frac{4 \times \frac{58.5}{6.023 \times 10^{23}}}{a^3}$$

$$\Rightarrow a^3 = \frac{4 \times 58.5}{2.18 \times 6.023 \times 10^{23}} = \left( \frac{40 \times 58.5}{2.18 \times 6.023} \right) \times 10^{-27} \text{ cc}$$

$$\Rightarrow a = \left( \frac{40 \times 58.5}{2.18 \times 6.023} \right)^{\frac{1}{3}} \times 10^{-8}$$

$$= 5.63 \times 10^{-8} \text{ cm} \\ = 5.63 \text{ \AA}$$

$$\therefore \text{Bond length} = \frac{a}{2} = \frac{5.63}{2} = 2.815 \text{ \AA.}$$

Formula

$$f = \frac{nM}{N_0} / a^3 \Rightarrow a^3 f = \frac{nM}{N_0}$$

## Seebeck effect

A thermo-couple is constituted by a pair of metals (Copper-iron, bismuth-antimony, etc) with their ends fused into one another. The soldered ends are called junctions of the thermo-couple.

Consider two rods, one of iron and another of copper, having common junction A & B.

A galvanometer G is also connected in the circuit (as shown in figure). Since there is no source of e.m.f. in

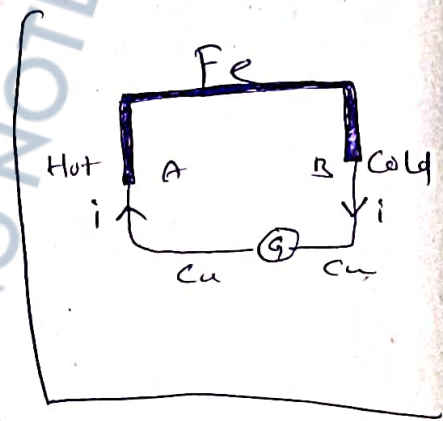
the circuit, therefore the galvanometer shows no deflection.

Now if one of the junctions, A is heated keeping the other cold, the galvanometer gives deflection, indicating an electric current starts flowing through the circuit. This current is called

Thermoelectric current.

The phenomenon by virtue of which a thermoelectric current is generated in the circuit comprising of two different metals, when the junctions are maintained at a difference of temp. is known as Seebeck effect or thermoelectric effect.

The e.m.f. generated is known as - thermo e.m.f.



hot copper  $\rightarrow$  At the hot junction, current flows from Copper to Fe

In cold junction current flows from Fe to Cu.

In Bi and Sb,

In hot junction  $\rightarrow$  Bismuth to Sb

In cold junction  $\rightarrow$  Antimony to Bismuth,

Bi-Sb thermo ~~cell~~  $\rightarrow$  Cu-Fe thermo emb