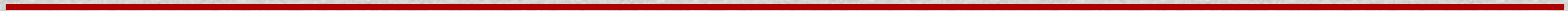
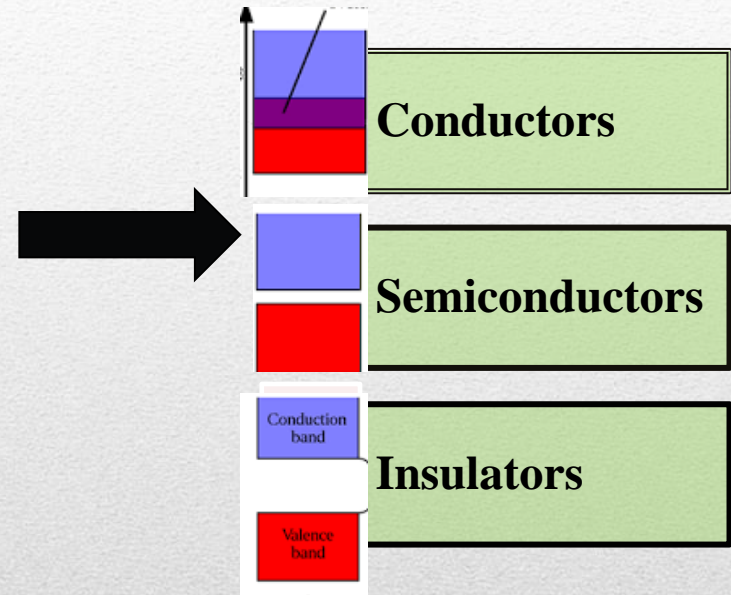
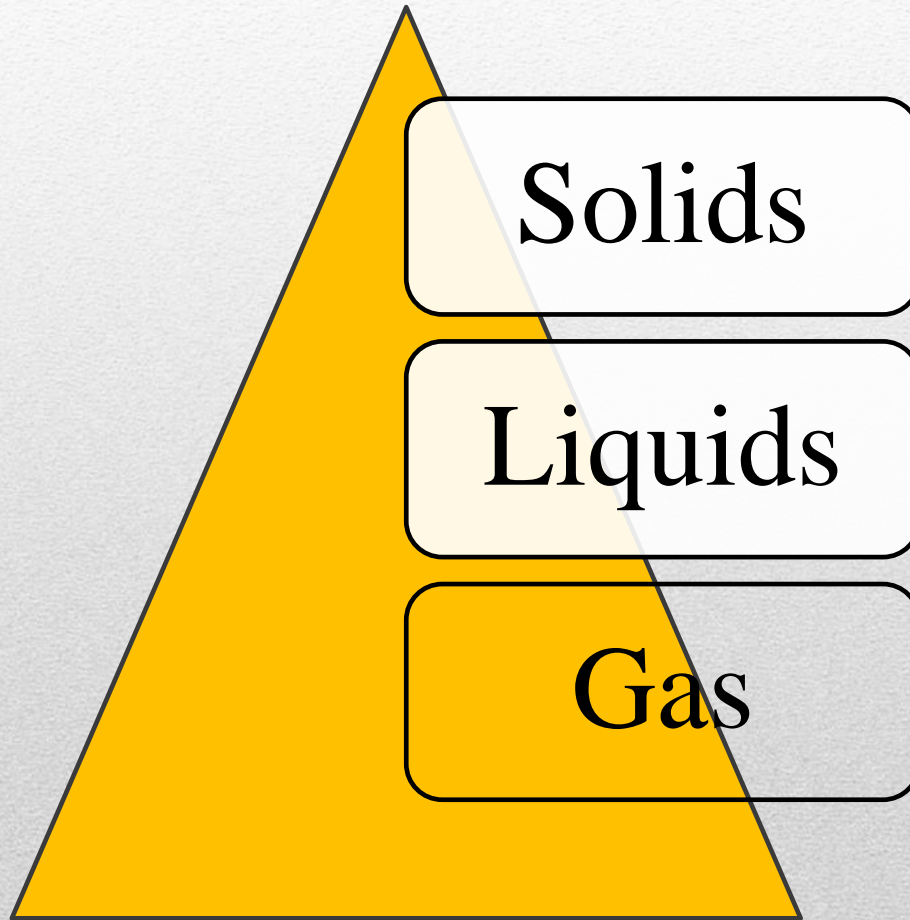


# SEMICONDUCTORS



# STATES OF MATTER



**Band Gap values**

**Conductors: VB and CB overlaps**

**Semiconductors:  $\sim 1-3$  eV**

**Insulators:  $> 5$  eV**

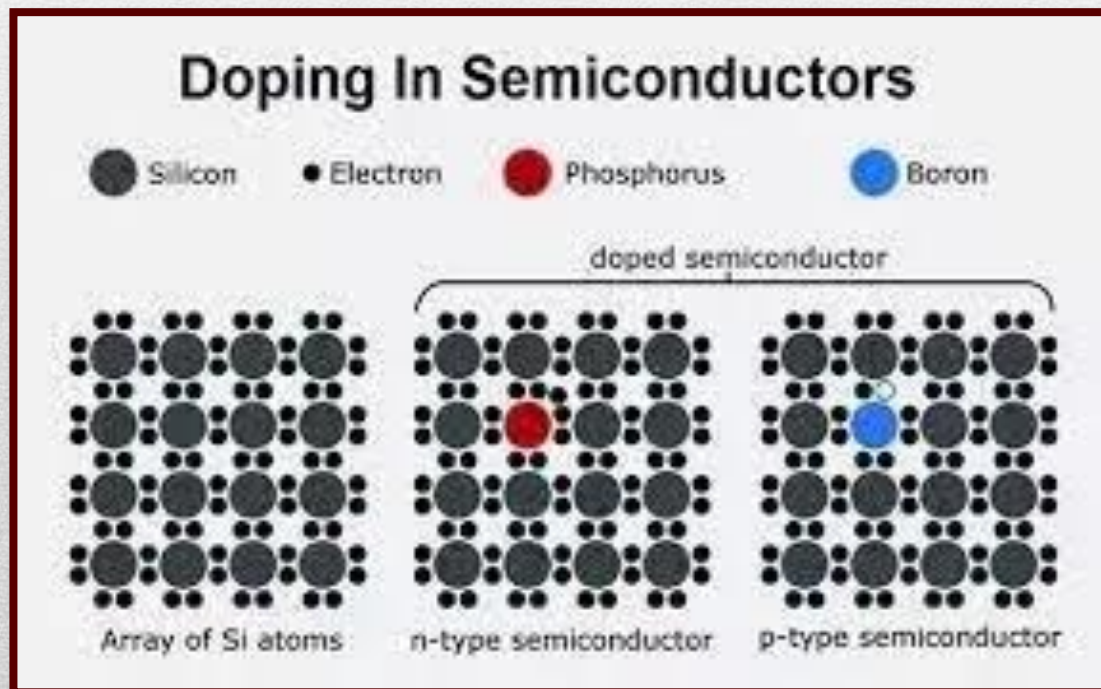


# DOPING IN SEMICONDUCTORS

- Semiconducting materials are very sensitive to impurities in the crystal lattice.
  - The controlled addition of these impurities is known as doping.
  - Allows the tuning of the electronic properties: technological applications.
  - Pure semiconductor are called 'intrinsic'.
  - Introduction of dopants → 'extrinsic semiconductors'.
  - Introduction of dopants → (i) new intra-band, energy levels  
(ii) Generation of positive or negative charge carriers.
-

- Extrinsic Semiconductors-

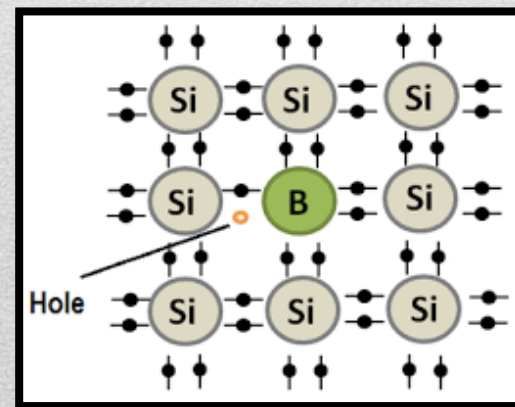
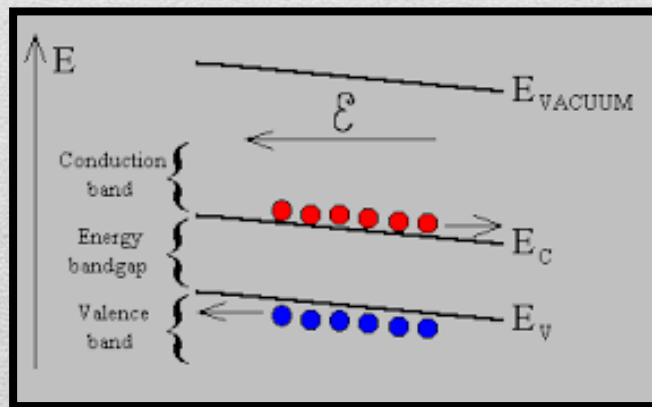
- (i) n-type: Pentavalent impurity- electrons as majority charge carriers (Donor Atoms)
- (ii) p-type: Trivalent impurity- holes as majority charge carriers (Acceptor atoms)



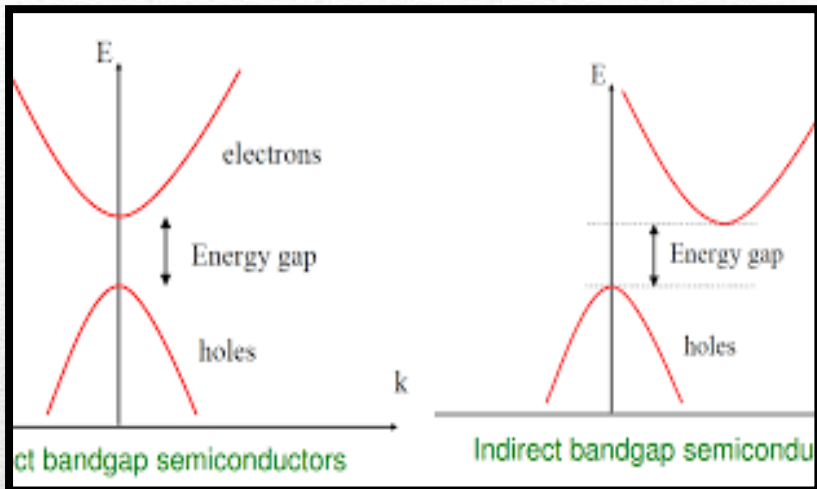


# CONCEPT OF HOLES

- A hole can be seen as the "opposite" of an electron.
- Holes have a positive charge.
- They are the *absence* of an electron in an atom (not physical particles).
- They are formed when electrons in atoms move out of the valence band to the conduction band.
- Holes can move from atom to atom in semiconducting materials as electrons leave their positions.

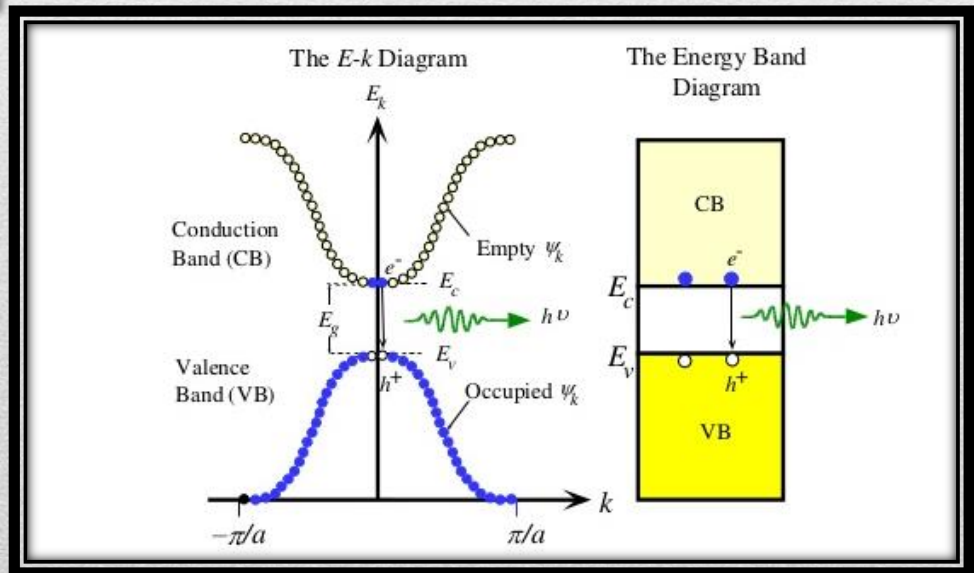


# BAND GAP IN SEMICONDUCTORS



Direct gap is important in optoelectronics as direct band gap materials have efficient radiative absorption and emission, which is what makes LEDs and laser diodes work

- An E-k diagram shows characteristics of a particular semiconductor material.
- It shows the relationship between the energy and momentum of available quantum mechanical states for electrons in the material.



<http://edetec106.blogspot.com/2016/01/differentiate-between-direct-and.html>

<https://www.slideshare.net/chinkitkit/chapter-4a-36657201>

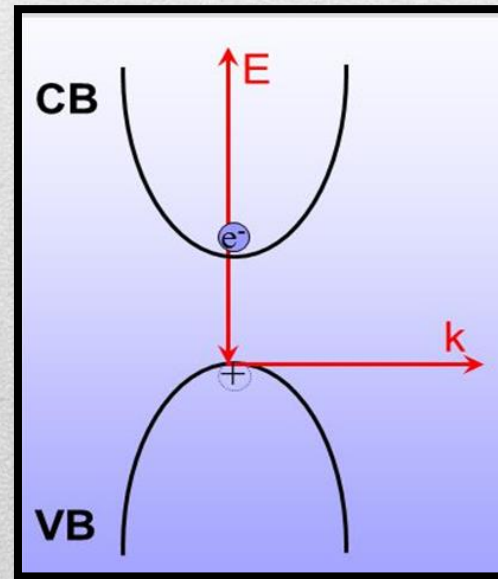


# EFFECTIVE MASS

- The effective mass represents the effect of all the internal forces on the motion of the electron in the conduction band.
  - Assumption: Mass of electron in solid is same as the mass of a free electron.
  - Experimentally: In some solids, electron mass is more while for some it is less than the free electron mass.
  - Effective mass: Experimentally determined electron mass.
  - Reason: Interaction between the drifting electrons and the atoms in the solids.
  - Sign of effective mass: determined from the sign of curvature of the E-k curve.
-

- The curvature of a graph at a minimum point is a positive quantity and while the curvature at a maximum point is a negative quantity.
- Particles (electrons) : near the minimum has a positive effective mass.
- Particles (holes) : near the valence band maximum has a negative effective mass.

- $$m^* = \frac{\hbar^2}{d^2 E / dk^2}$$



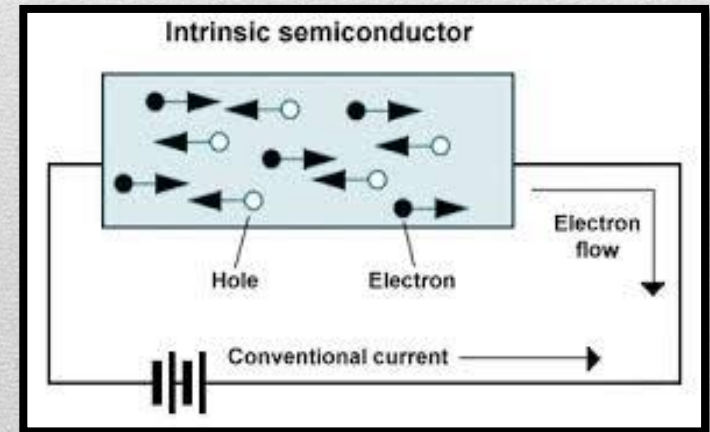


*For Derivation refer class notes*

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# MOBILITY

- When an electric field is applied across a solid, it accelerates the electrons in the direction of applied field.
- The moving electrons undergo repeated collisions with the atoms and hence moves with a steady velocity known as Drift velocity represented as  $v_d$ .
- $v_d \propto E \rightarrow v_d = \mu E$  where  $\mu$  is the mobility of electrons.
- Mobility: measure of how quickly an electron can move through a metal or semiconductor in presence of electrical field.
- Semiconductor mobility depends  
(i) defect concentration (ii) temperature.
- $\mu_e$  (metals) =  $10^{-3} \text{ m}^2/\text{Vs}$
- $\mu_e$  (semiconductors) =  $10^{-1} \text{ m}^2/\text{Vs}$ .





# CARRIER CONCENTRATION

- The density of electrons in a semiconductor  $\rightarrow$  the density of available states and the probability that each of these states is occupied.
- The density of occupied states per unit volume *is given as*

$$n(E) = g_c(E)f(E)$$

- Holes correspond to empty states in the valence band  $\rightarrow$  the probability of having a hole equals the probability that a particular state is not filled.
- Hole density per unit energy,  $p(E)$  is given as

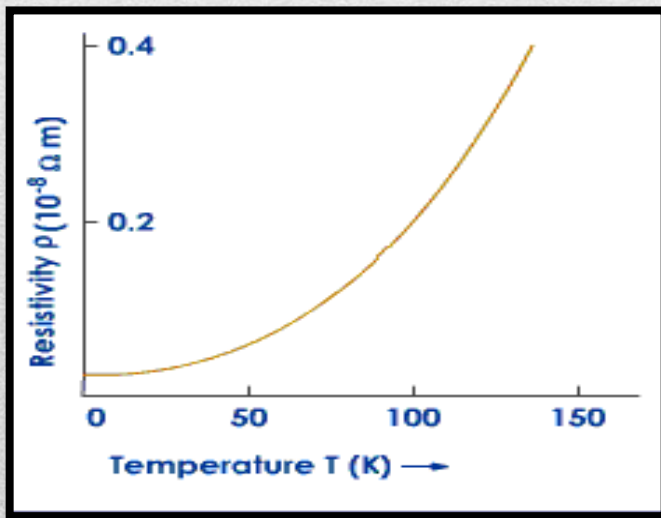
$$p(E) = g_v(E) [1 - f(E)]$$

where  $g_v(E)$  is the density of states in the valence band.

- The density of carriers is then obtained by integrating the density of carriers per unit energy over all possible energies within a band.
-

# CONDUCTIVITY IN METALS

- Conductivity is attributed to *free-charge carriers in metals*.
- Increase in temperature → increases the vibrations of the metal ions.
- Increased vibrations → causes frequent collisions between the electrons → drains out energy of the free electrons → restricts the movement of the delocalized electrons → drift velocity decreases → resistivity of the metal increases → current decreases → conductivity of the material decreases.



- $J = \sigma E$  (Ohm's law)  
 $\rho = 1/\sigma$

The temperature dependence is given as

$$\rho_t = \rho_0 [1 + \alpha (T - T_0)]$$



# Electrical Conductivity of Semiconductors:

$$I = I_e + I_h$$

$$I_e = neAv_e$$

$$I_h = peAv_h$$

So,  $I = neAv_e + peAv_h$

If the applied electric field is small,  
then semiconductor obeys Ohm's law.

$$\therefore \frac{V}{R} = neAv_e + peAv_h$$

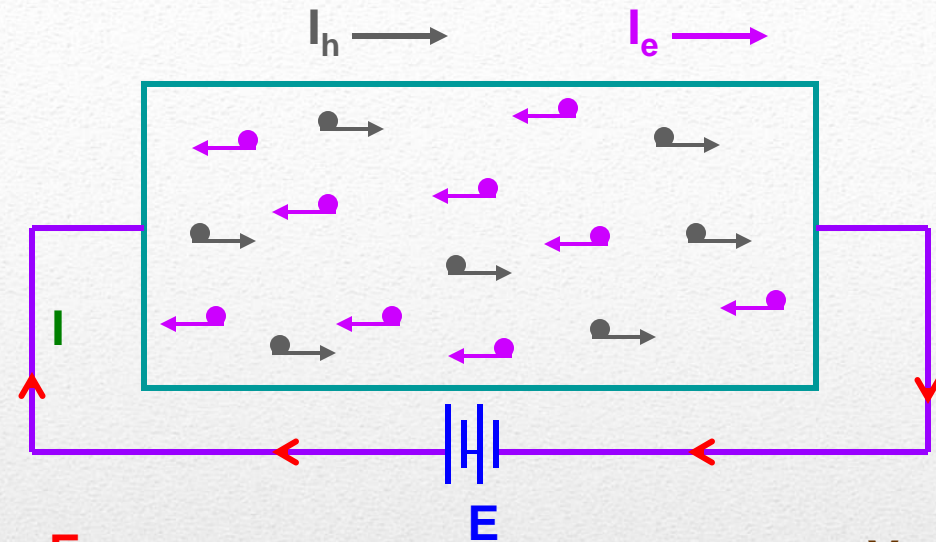
$$= eA (nv_e + pv_h)$$

Or  $\frac{VA}{\rho l} = eA (nv_e + pv_h)$

since  $R = \frac{\rho l}{A}$

**Note:**

1. The electron mobility is higher than the hole mobility.
2. The resistivity / conductivity depends not only on the electron and hole densities but also on their mobilities.
3. The mobility depends relatively weakly on temperature.



$$\frac{E}{\rho} = e (nv_e + pv_h) \quad \text{since } E = \frac{V}{l}$$

Mobility ( $\mu$ ) is defined as the drift velocity per unit electric field.

$$\therefore \frac{1}{\rho} = e (n\mu_e + p\mu_h)$$

Or

$$\sigma = e (n\mu_e + p\mu_h)$$

# CONDUCTIVITY IN INTRINSIC SC

- For Intrinsic semiconductors, the conductivity is given by

$$\sigma = e(n\mu_e + p\mu_h) = n_i e(\mu_e + \mu_h)$$

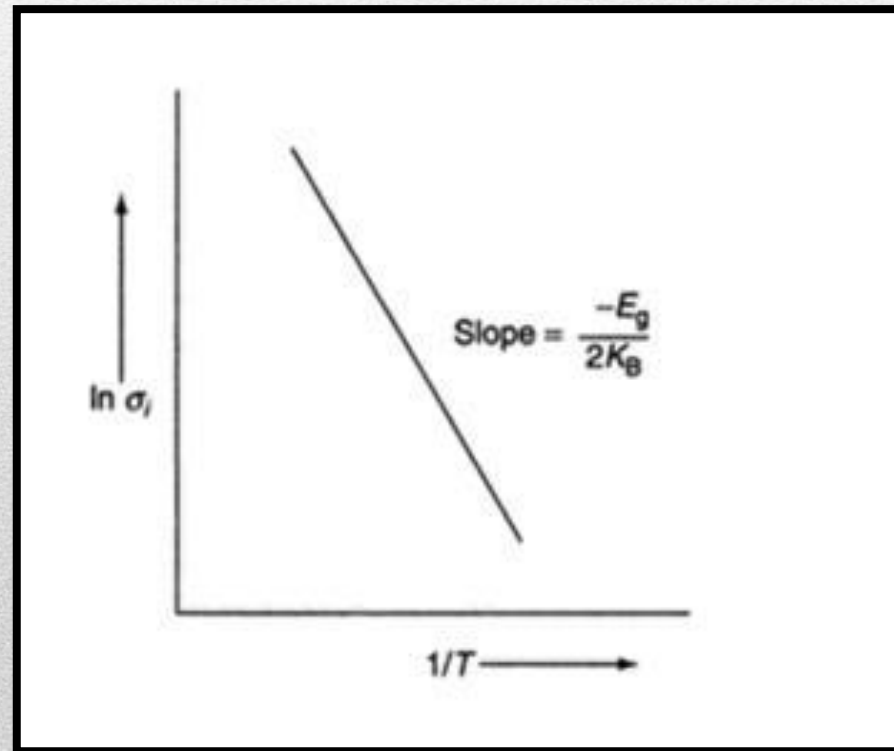
$\mu_n$  and  $\mu_p$  refer to the mobilities of the electrons and holes

$n$  and  $p$  refer to the density (concentration) of electrons and holes  
 $n_i$  is the intrinsic charge carrier density

- The electrons in the valence band gains energy  $\rightarrow$  moves to higher energy levels in the conduction band  $\rightarrow$  becomes charge carriers
  - Carrier concentration depends exponentially on the band gap and is given as  $n_i = \sqrt{N_c N_v} \exp\left(-\frac{E_g}{2k_B T}\right)$
-

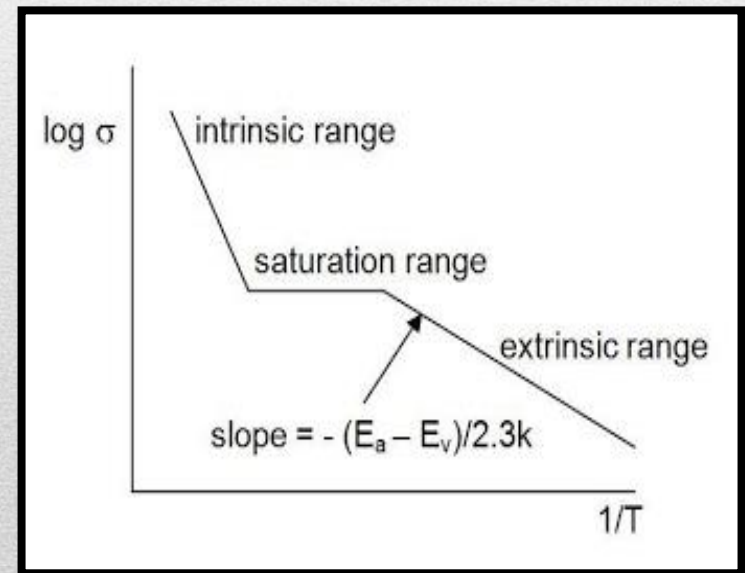


- In Intrinsic semiconductors → Conductivity increases with increasing temperature.
- $\log_e \sigma = \frac{-E_g}{2kT} + \log_e B$



# CONDUCTIVITY IN EXTRINSIC SC

- Low temperature  $\rightarrow$  frozen charge carriers  $\rightarrow$  resistivity is extremely high.
- Moderate increase in temperature  $\rightarrow$  Rapid decrease in resistivity with the increase of ionized charges.
- At sufficiently high temperature  $\rightarrow$  dopants are completely ionized  $\rightarrow$  conductivity decreases and the resistivity increases again.
- At still higher temperature  $\rightarrow$  resistivity decreases sharply due to appreciable excitation of all carriers and crossing the energy gap.





- In doped semiconductor (Extrinsic), majority carriers greatly outnumber the minority carriers, so that the equation can be reduced to a single term involving the majority carrier.

- For n-type semiconductor

$$\sigma_n = ne\mu_e$$

- For p-type semiconductor

$$\sigma_p = pe\mu_h$$

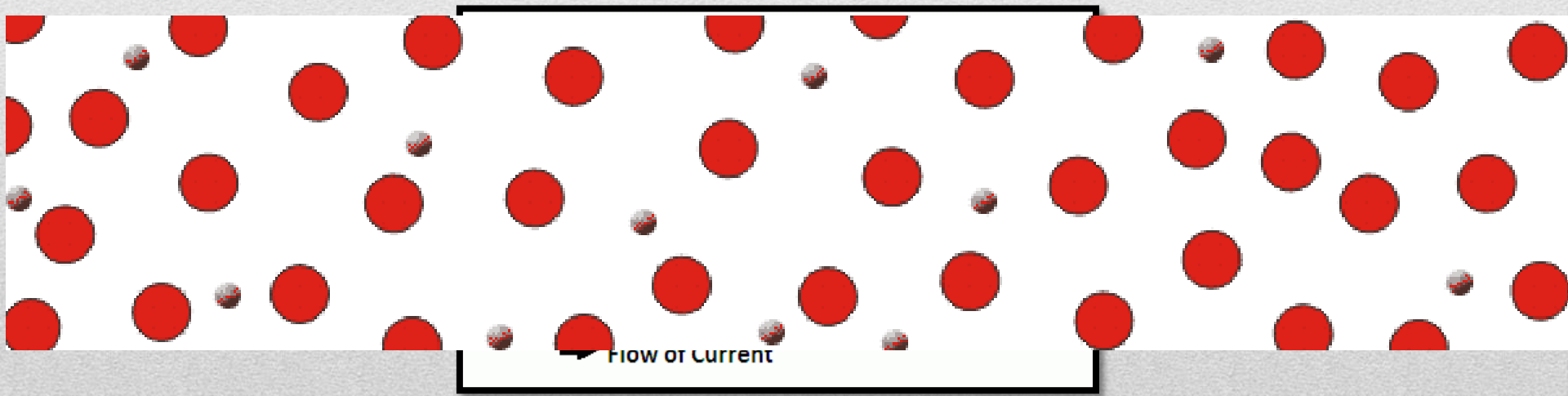
- Conductivity of a material is determined by two factors:
  - (i) concentration of free carriers available to conduct current
  - (ii) their mobility (or freedom to move).

**In a semiconductor, both mobility and carrier concentration are temperature dependent.**

---

# DRIFT CURRENT

- **Absence of field:** free electrons move in a conductor with random velocities and random directions.
- **Presence of field:** the randomly moving electrons experience an electrical force in the direction of the field.
- **Electrons shift towards higher potential with their random motion.**
- **The electrons will drift towards higher potential along with their random motions.**





- Electrons have a net velocity towards the higher potential end of the conductor known as the drift velocity of electrons.
- The drift movement of electrons inside an electrically stressed conductor, is known as drift current.
- The drift current density for hole and electrons are given by

$$J_{(drift)el} = -nev_e$$

$$J_{(drift)hole} = pev_h$$

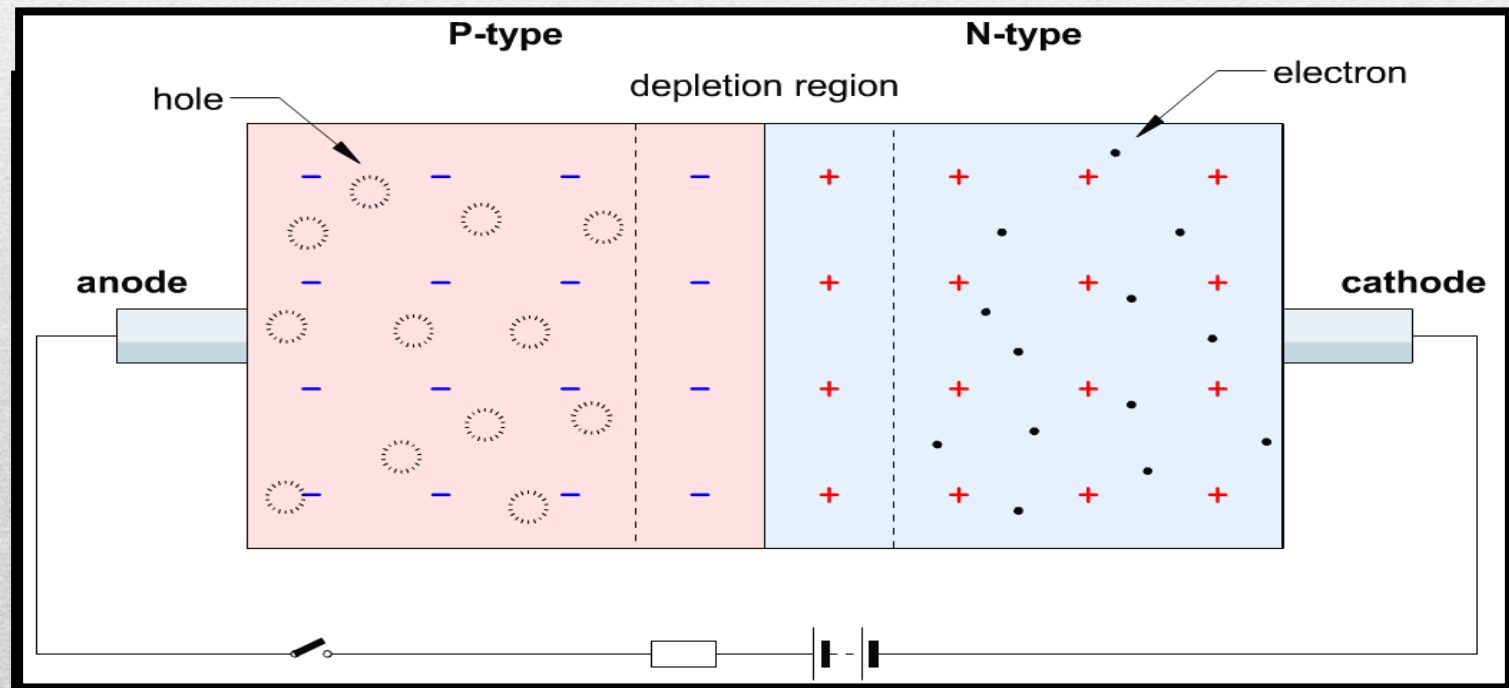
- where  $n$ ,  $p$  are the electron and hole densities,  $v_h$  and  $v_e$  are the drift velocities of holes and electrons respectively.
- Negative sign indicates that the electrons having -ve charge move in direction opposite to the applied field.
- Total drift current density

$$J_{(drift)Total} = J_{(drift)hole} + J_{(drift)el} = pev_h + nev_e$$

---

# DIFFUSION CURRENT

- In semiconducting material → Dopants are introduced to some region → even distribution of carriers takes place to maintain the uniformity → known as diffusion process
- Movement of the mobile charge carriers are responsible for the flow of diffusion current from one region to the other.
- No source of energy is required diffusion current.





# DIFFUSION CURRENT

- Non-uniformity of charge carriers (electrons/holes) → gives the diffusion current (is independent of the electric field) → depends on the concentration gradient.
  - Concentration of electrons ( $n$ ) and holes( $p$ ) varies with the distance  $x$ .
  - Diffusion current density for electrons is  $J_{(diff)el} = eD_n \frac{dn}{dx}$ , where  $D_n$  is the diffusion coefficient for electrons and  $dn/dx$  is the concentration gradient of electrons.
  - Diffusion current density for holes is  $J_{(diff)ho} = -pD_p \frac{dp}{dx}$ , where  $D_p$  is the diffusion coefficient for holes and  $dp/dx$  is the concentration gradient of holes.
  - Resultant diffusion current density for both holes and electrons is given as  $J_{(diff)Total} = eD_n \frac{dn}{dx} - pD_p \frac{dp}{dx}$
-

# TOTAL CURRENT DENSITY

Total current density in semiconductor is the sum of drift current and diffusion current is given by

$$J_{Total} = J_{drift} + J_{diffusion}$$

$$J_{Total} = J_{(drift)hole} + J_{(drift)el} + J_{(diffusion)el} + J_{(diffusion)hole}$$

$$J_{Total} = p e v_h + n e v_e + e D_n \frac{dn}{dx} - p D_p \frac{dp}{dx}$$

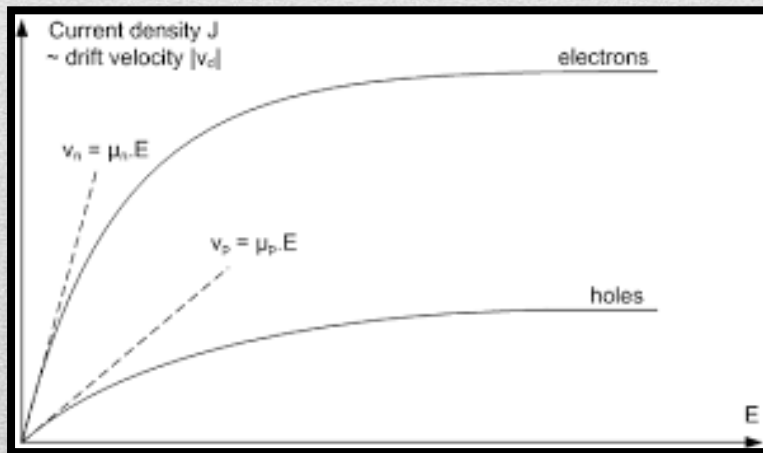


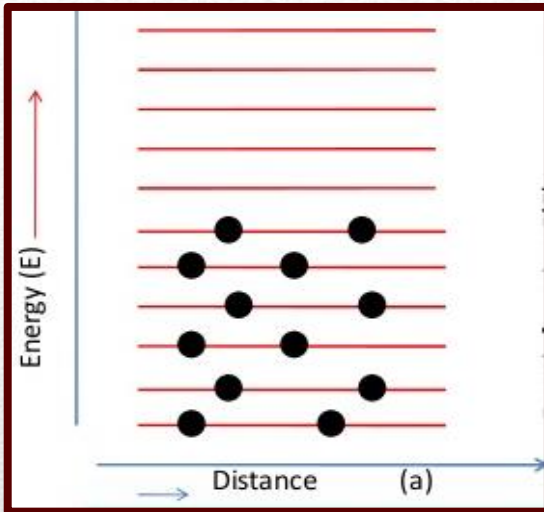
Figure shows the plot for the current density  $J_{drift}$  and the absolute value of the drift velocity, over the electric field  $E$ . The mobility of holes and electrons can be evaluated using the tangential of the drift velocity.



# FERMI DIRAC DISTRIBUTION FUNCTION

- The probability density functions describes the probability that particles occupy the available energy levels in a given system.
  - Fermions : half-integer spin particles- Electrons are Fermions- obeys Pauli exclusion principle → one Fermion occupies a single quantum state → fills the available states in an energy band.
  - Fermi function : The probability that an energy level at energy,  $E$ , in thermal equilibrium with a large system, is occupied by an electron.
  - Fermi Dirac distribution function is given as:
  - $$f(E) = \frac{1}{1 + \exp^{(E - E_F)/kT}}$$
  - $E_F$  : Fermi energy;  $k$  : Boltzmann constant;  $T$ : Temperature
-

# FERMI LEVEL IN A CONDUCTOR



Fermi function is  $f(E) = \frac{1}{1 + \exp^{(E - E_F)/kT}}$

**Case 1: At T= 0 K;  $E < E_F \Rightarrow (E - E_F)$  is negative**

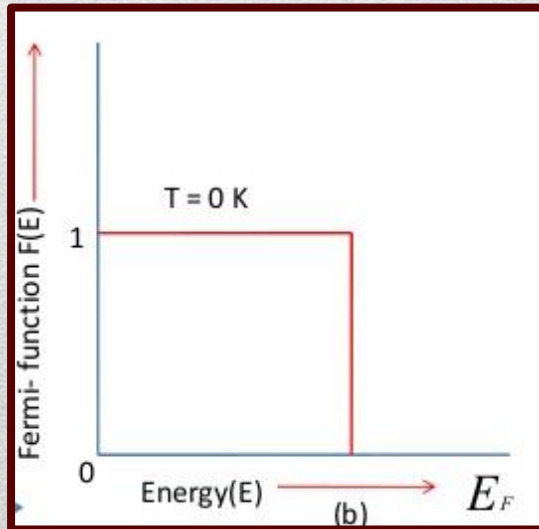
$$\therefore f(E) = \frac{1}{1 + e^{-(E - E_F)/0}} = \frac{1}{1 + e^{-\infty}} = \frac{1}{1 + 0} = 1$$

*All levels lying below  $E_F$  are occupied.*

**Case 2: At T= 0 K;  $E > E_F \Rightarrow (E - E_F)$  is positive**

$$\therefore f(E) = \frac{1}{1 + e^{(E - E_F)/0}} = \frac{1}{1 + e^{\infty}} = \frac{1}{1 + \infty} = \frac{1}{\infty} = 0$$

*All energy levels lying above  $E_F$  are vacant*

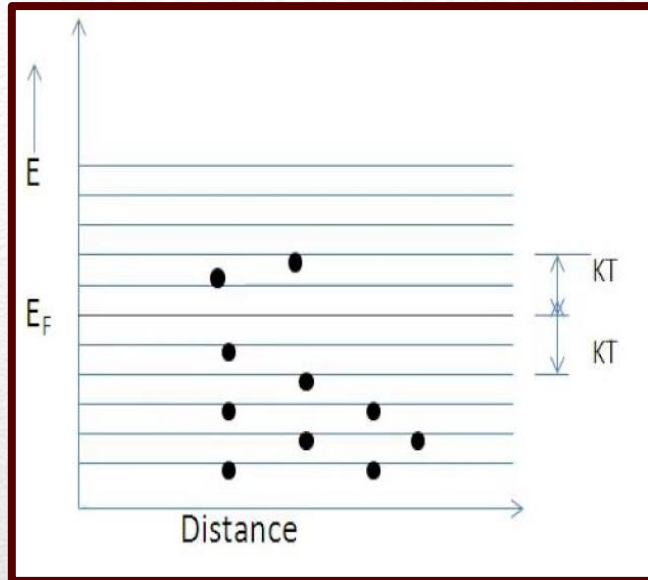






**What happens to the fermi level at high temperature ?**

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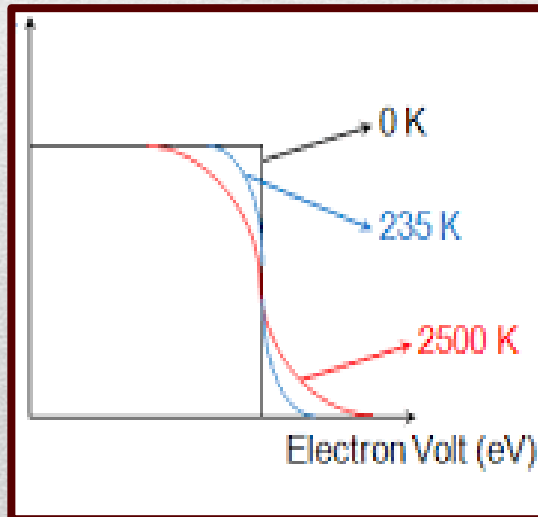
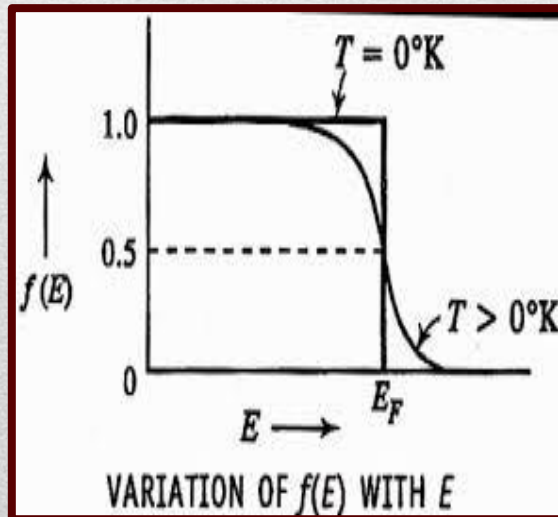


**Case 3: At  $T > 0$  K;  $E = E_F$**

$$\therefore f(E) = \frac{1}{1 + e^{\left(\frac{0}{kT}\right)}} = \frac{1}{1 + e^0} = \frac{1}{1 + 1} = \frac{1}{2}$$

The probability of occupancy at any temperature  $T > 0$  K is 50 %.

**Fermi energy:** Average energy possessed by electrons participating in conduction at temperature above 0K.



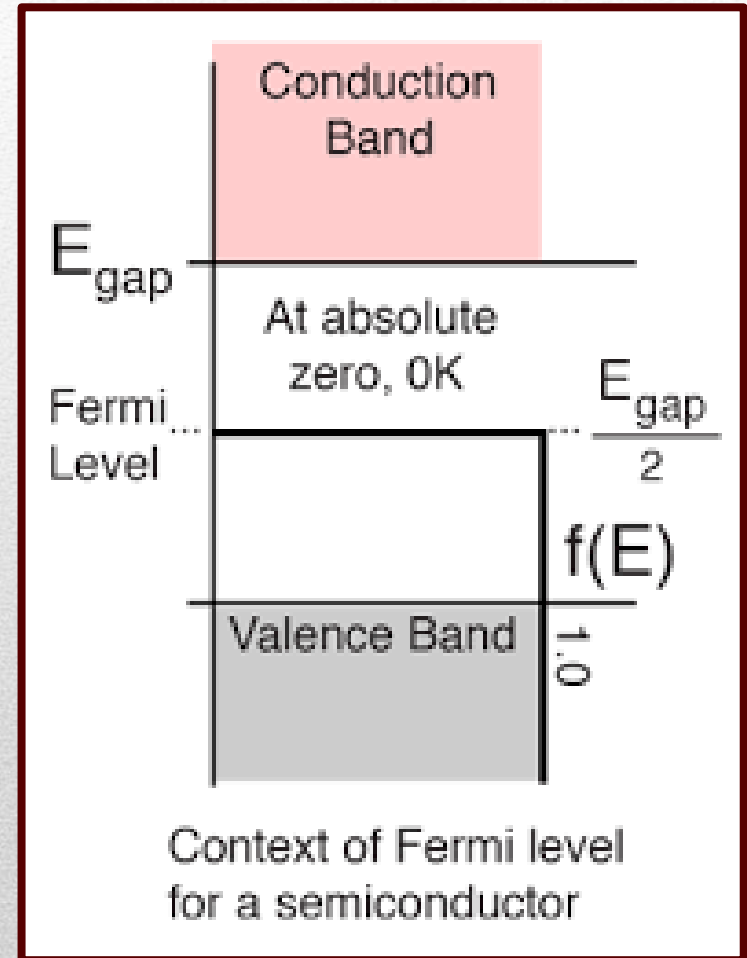
**Fermi velocity:  $v_F$ :** It is the velocity of the electrons in the highest occupied states in metals at zero temperature.

$$v_F = \sqrt{2E_F/m}$$



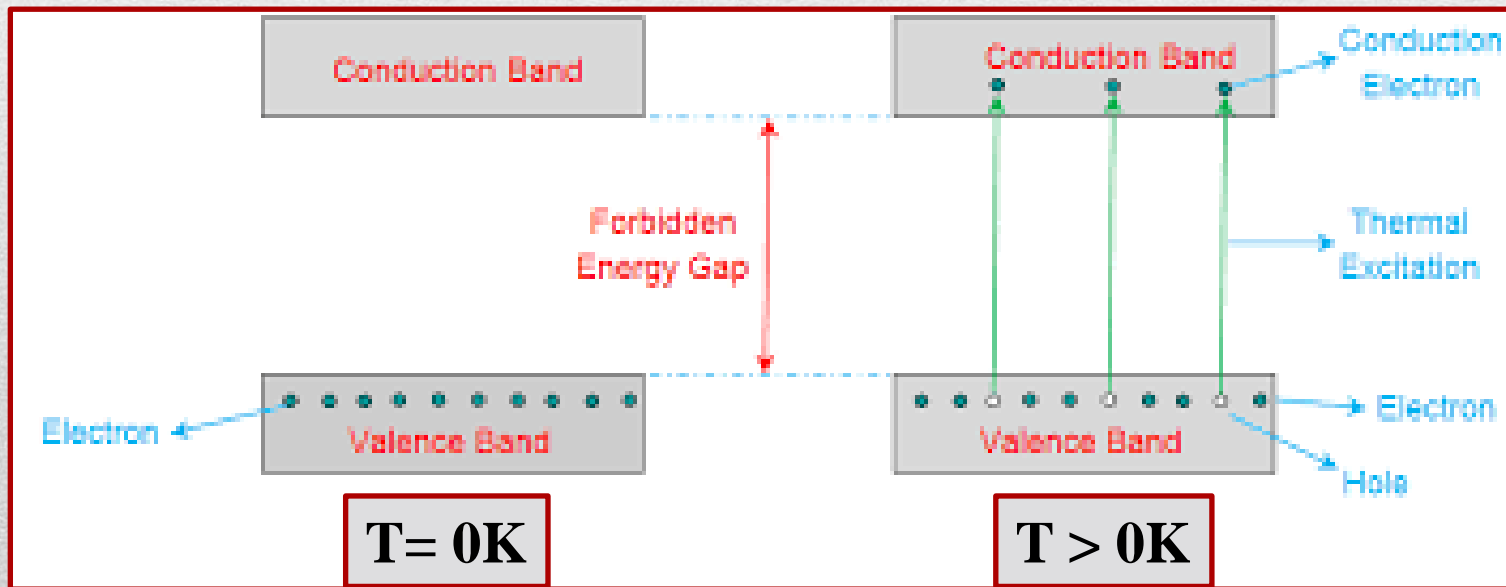
# WHAT IS FERMI LEVEL

- The highest energy level that an electron occupies at the absolute zero temperature is known as the Fermi Level.
- The Fermi level lies between the valence band and conduction band. At  $T=0$  K, the electrons are all in the lowest energy state  $\rightarrow$  Fermi level can be considered as the sea of fermions (or electrons) above which no electrons exist.
- The Fermi level changes as the temperature increases or electrons are added to or withdrawn from the solids.



# FERMI LEVEL IN INTRINSIC SC

- At  $T=0$  K, the valence band will be full of electrons  $\rightarrow$  impossible to cross the energy barrier  $\rightarrow$  acts as an insulator.
- At  $T > 0$  K  $\rightarrow$  the electron movement from the valence band to the conduction band increases  $\rightarrow$  create holes in the valence band in place of electrons.
- The electron concentration ' $n$ ' is equal to hole concentration ' $p$ '.





# POSITION OF FERMI LEVEL IN INTRINSIC SC

Let,  $n$  be the number of electrons in the semiconductor band.

$p$  be the number of holes in the valence band.

At temperature  $T > 0$  K

$$n = N_c e^{-(E_c - E_F)/kT} \quad p = N_v e^{-(E_F - E_V)/kT}$$

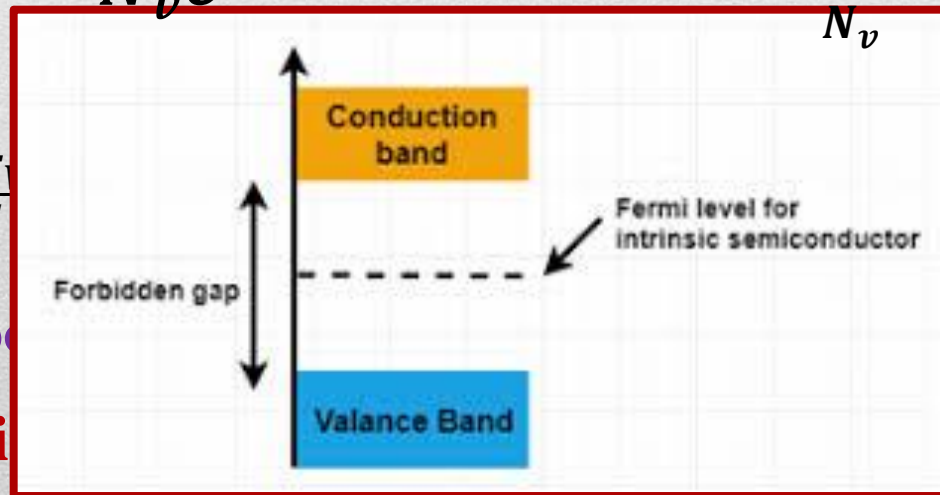
$N_c$  is the effective density of states in the conduction band and  $N_v$  is the effective density of states in the valence band. For an intrinsic semiconductor,  $n_e = n_v$

$$N_c e^{-(E_c - E_F)/kT} = N_v e^{-(E_F - E_V)/kT} \Rightarrow \frac{N_c}{N_v} = \frac{e^{-(E_F - E_V)/kT}}{e^{-(E_c - E_F)/kT}}$$

$$\Rightarrow e^{\frac{-(2E_F - E_c - E_V)}{kT}}$$

Taking log on both sides

$\therefore$  Fermi level is

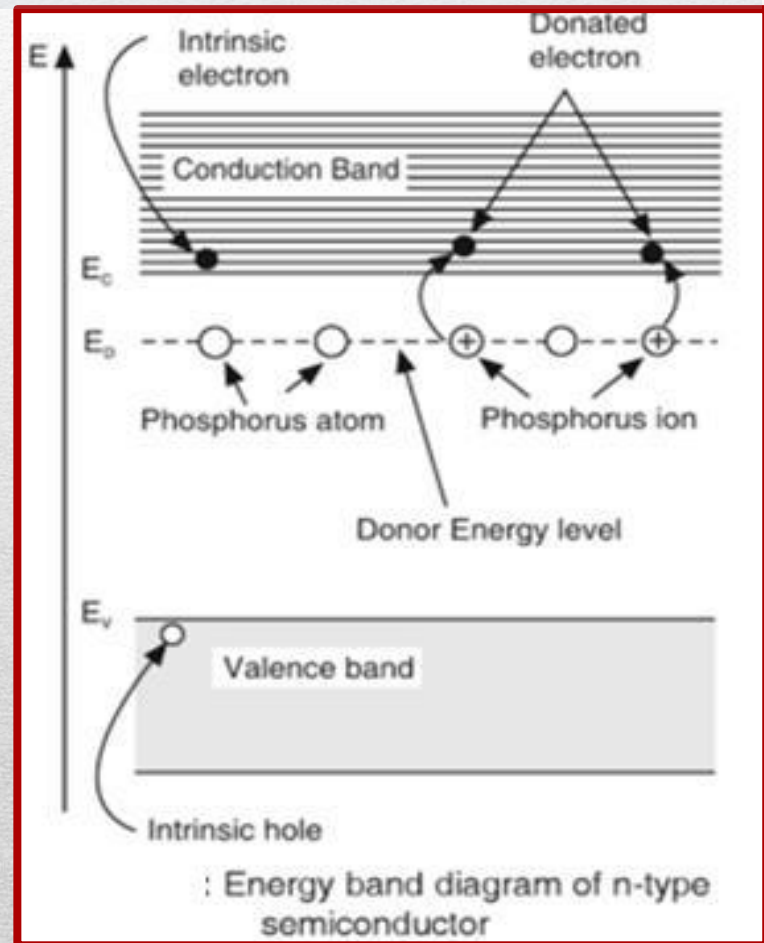
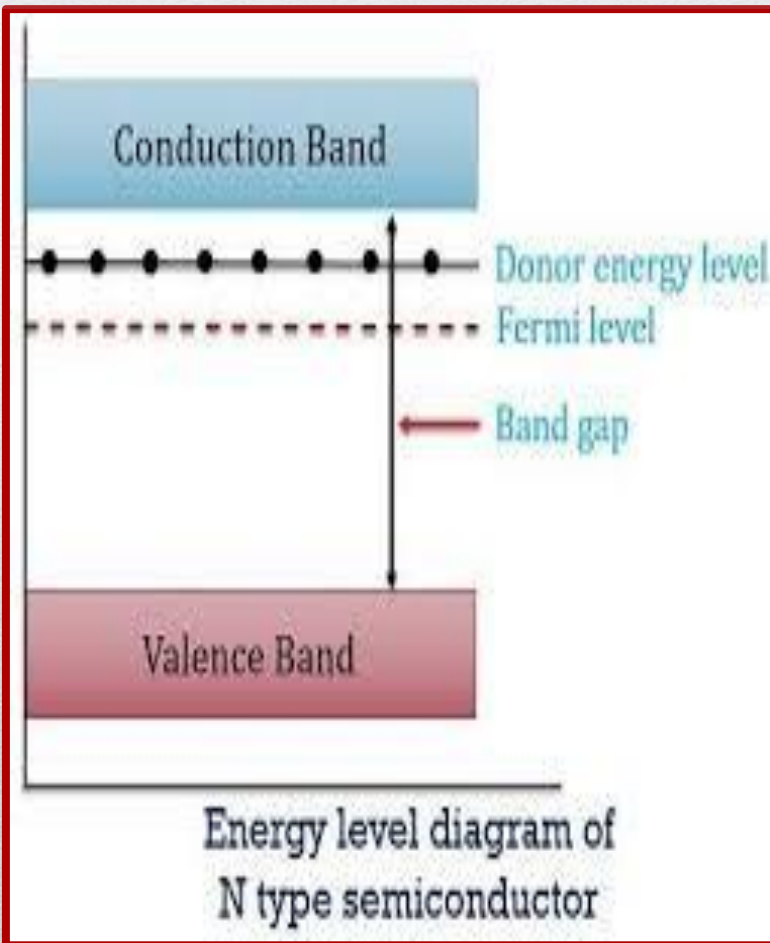


$$E_F = \frac{E_c + E_V}{2}$$

at the center of the band gap.

# FERMI LEVEL IN EXTRINSIC SC

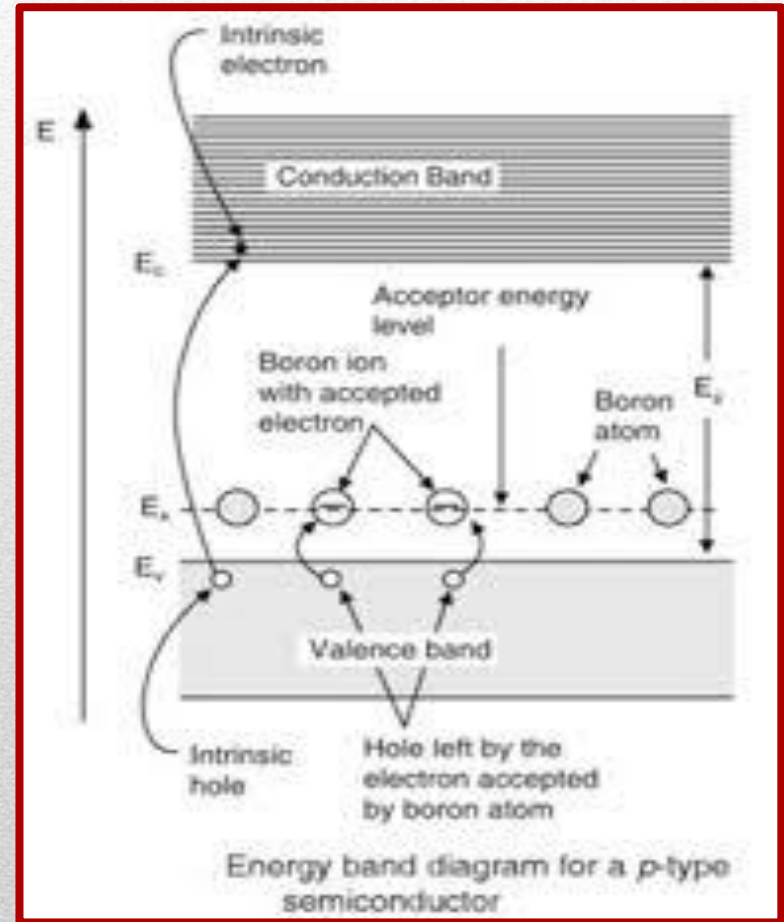
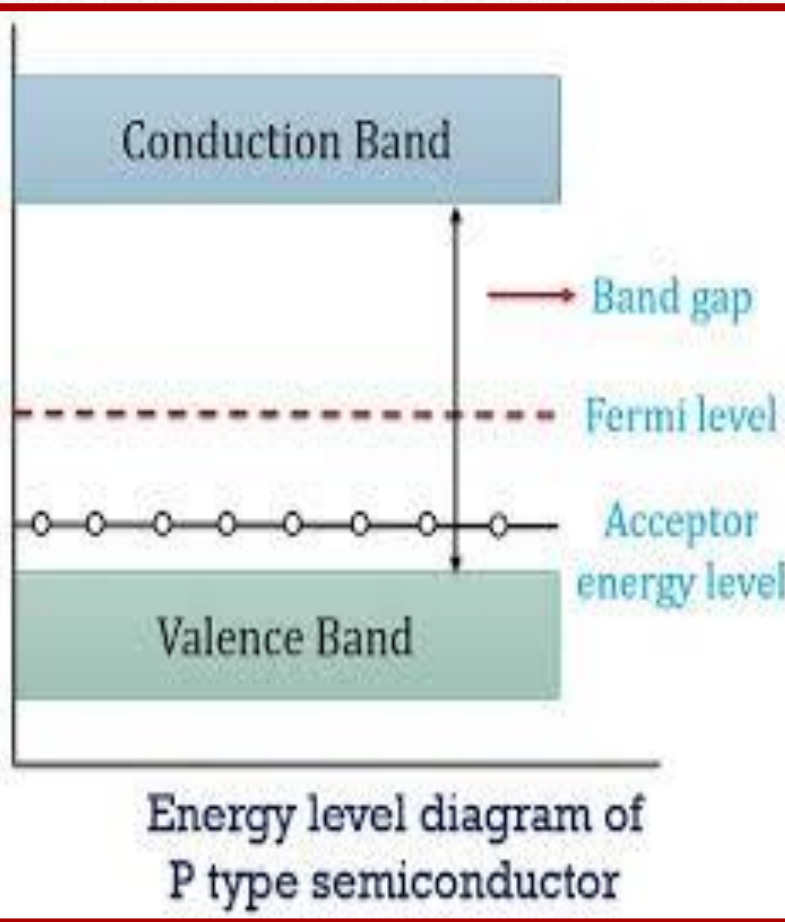
N-type SC- pentavalent impurity : electrons as majority charge carriers: donor impurities





# FERMI LEVEL IN EXTRINSIC SC

P-type SC- trivalent impurity : holes as majority charge carriers: acceptor impurities



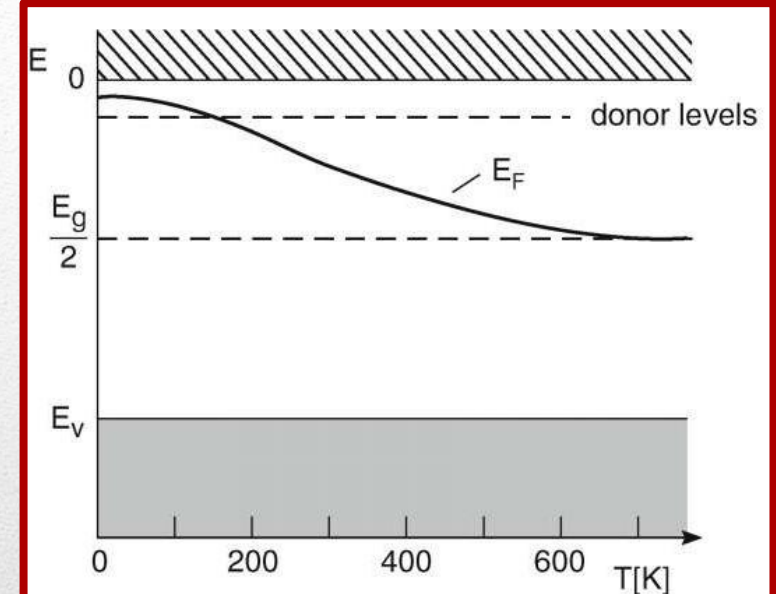
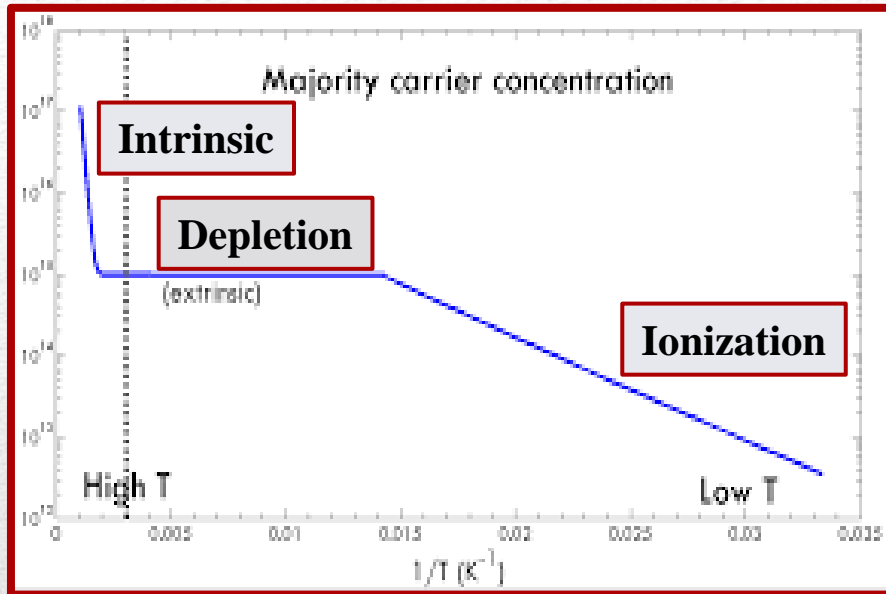


# **EFFECT OF TEMPERATURE ON THE FERMI LEVEL OF N-TYPE SEMICONDUCTOR**

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# EFFECT OF TEMPERATURE ON $E_F$ OF N- TYPE SC



Region I: Ionization region:  $E_{F_n} = \frac{E_c + E_D}{2}$

Region II: Depletion Region:  $E_{F_n} = E_D$

Region III: Intrinsic Region:  $E_{F_n} = E_{F_i} = \frac{E_g}{2}$

Fermi level position in n type semiconductor with respect to intrinsic Fermi level is given as

$$E_{F_n} - E_{F_i} = k_B T \ln \frac{n}{n_i}$$

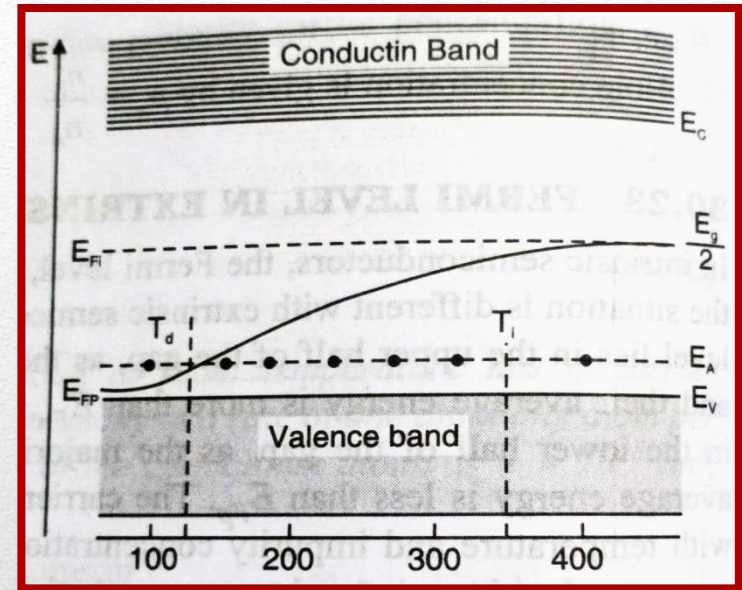
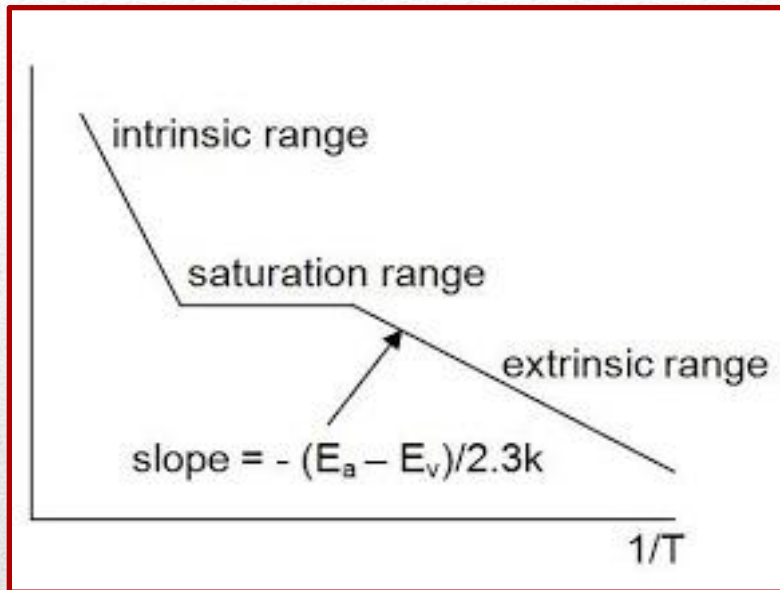


# **EFFECT OF TEMPERATURE ON THE FERMI LEVEL OF P-TYPE SEMICONDUCTOR**

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# EFFECT OF TEMPERATURE ON $E_F$ OF P- TYPE SC



Region 1: Ionization region:  $E_{F_p} =$

$$\frac{E_v + E_a}{2}$$

Region 2: Depletion Region:  $E_{F_p} = E_a$

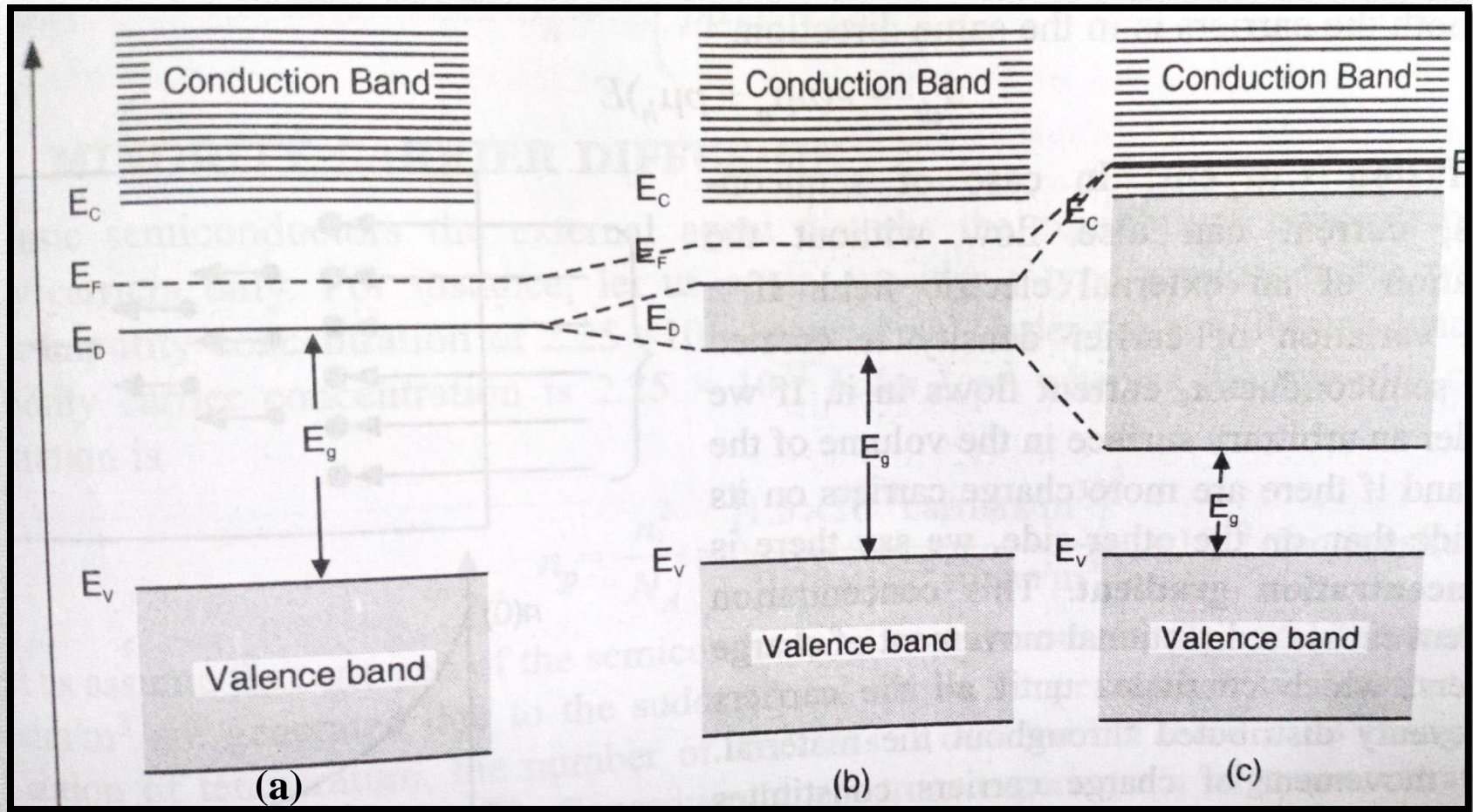
Region 3: Intrinsic Region :  $E_{F_p} =$

$$E_{F_i} = \frac{E_g}{2}$$

Fermi level position in p type semiconductor with respect to intrinsic Fermi level is given as

$$E_{F_p} - E_{F_i} = -k_B T \ln \frac{p}{n_i}$$

# EFFECT OF IMPURITY CONCENTRATION ON $E_F$ OF N- TYPE SC





# EFFECT OF IMPURITY CONCENTRATION ON $E_F$ OF P- TYPE SC

