Electronic Devices

Aakash Jog

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1 Lecturer Information

Tammy Ben-Yaacov

E-mail: tammybenyaacov@gmail.com

2 Instructor Information

Asia Shapira

E-mail: asiasapi@gmail.com

3 Required Reading

1. Streetman, B. Solid State Electronic Devices

4 Additional Reading

- 1. Bar-Lev, A. Semiconductor and Electronic Devices
- 2. S. M. Sze, Physics of Semiconductor Devices
- 3. Kittel, C. (2005). Introduction to Solid State Physics

Part I

1 Energy Bands in Semiconductors

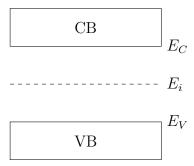


Figure 1: Energy Bands and Intrinsic Energy Level

2 Determining Factors for Number of Charge Carriers in Energy Bands

2.1 Density of Allowed States in Bands

The number of allowed states in the conduction and valence bands are denoted as $g_C(E)$ and $g_V(E)$, respectively.

There are no allowed states in the gap between the conduction and valence bands.

Theorem 1.

$$g_C(E) \approx \sqrt{E - E_C}$$

2.2 Probability of Occupancy of Allowed States

Definition 1 (Fermi function). The probability that an available energy state E will be occupied is

$$f(E) = f_{FD}(E)$$
$$= \frac{1}{1 + e^{\frac{E - E_f}{kT}}}$$

where E_f is the Fermi level or the Fermi energy, and T is the temperature.

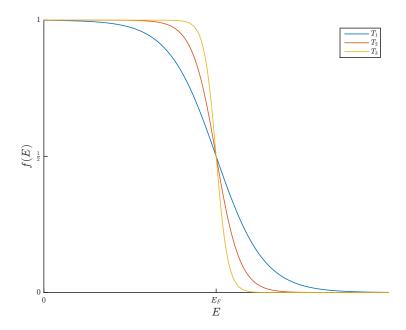


Figure 2: Graph of Fermi function for $T_1 > T_2 > T_3$

Therefore, the graph of f(E) with respect to E is as in Figure 2. If $E < E_f$,

$$\lim_{T \to 0} f(E) = 1$$

If $E = E_f$,

$$f(E) = \frac{1}{2}$$

If $E_f < E$,

$$\lim_{T \to 0} f(E) = 0$$

Therefore, at $0 \,\mathrm{K}$, f(E) is the unit step function. Therefore, at $0 \,\mathrm{K}$, all electrons are at available energy levels less than E_f , i.e. in the valence band. Hence, the conduction band is empty.

3 Number of Electrons in Conduction Band

Theorem 2. Electrons in semiconductors obey Fermi-Dirac statistics.

Consider an energy band of thickness dE, at energy E, in the conduction band. Therefore, the number of electrons in the band is

$$dn(E) = g_C(E)f(E) dE$$

Therefore, the total number of electrons in the conduction band are

$$n = \int_{E_C}^{E_C + \Delta} g_C(E) f(E) dE$$

where Δ is the thickness of the conduction band. As there are no allowed states above the conduction band,

$$n = \int_{E_C}^{\infty} g_C(E) f(E) \, \mathrm{d}E$$

4 Position of E_F

The graphs of f(E) and 1-f(E) are as in Figure 3 and ???. The area bounded by

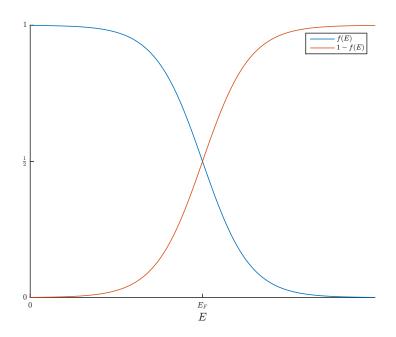


Figure 3: Graph of f(E) and 1 - f(E)

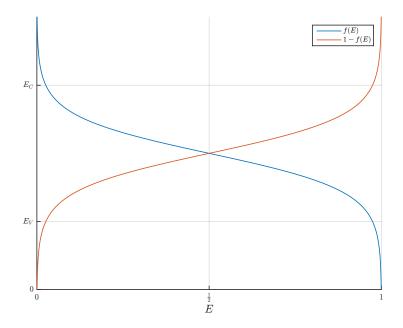


Figure 4: Graph of f(E) and 1 - f(E) with E_C and E_V

f(E) above E_C , in Figure 4, represents the number of electrons in the conduction band. Similarly, the area bounded by 1 - f(E) below E_V , in Figure 4, represents the number of holes in the valence band.

The densities of states for electrons and holes are given by

$$g_C(E) = \frac{4\pi}{h^3} (2m_e^*)^{\frac{3}{2}} (E - E_C)^{\frac{1}{2}}$$
$$g_V(E) = \frac{4\pi}{h^3} (2m_h^*)^{\frac{3}{2}} (E_V - E)^{\frac{1}{2}}$$

Therefore, substituting and solving,

$$n = \frac{2}{h^3} (2\pi m_e^* kT)^{\frac{3}{2}} e^{-\frac{E_C - E_F}{kT}}$$
$$p = \frac{2}{h^3} (2\pi m_h^* kT)^{\frac{3}{2}} e^{-\frac{E_F - E_V}{kT}}$$

The constants

$$N_C = \frac{2}{h^3} (2\pi m_e^* kT)^{\frac{3}{2}}$$

$$N_V = \frac{2}{h^3} (2\pi m_h^* kT)^{\frac{3}{2}}$$

are called the effective density of states in the conduction band and the valence band respectively.

4.1 Intrinsic Semiconductor

At 0 K, all electrons are in the valence band. Therefore, for all electrons to be below E_F , E_F must be between E_C and E_V . As the material is an intrinsic semiconductor,

$$n = p$$

$$\therefore (m_e^*)^{\frac{3}{2}} e^{-\frac{E_C - E_F}{kT}} = (m_h^*)^{\frac{3}{2}} e^{-\frac{E_F - E_V}{kT}}$$

$$\therefore \left(\frac{m_e^*}{m_h^*}\right)^{\frac{3}{2}} = e^{-\frac{E_F - E_V}{kT}} e^{\frac{E_C - E_F}{kT}}$$

$$= e^{\frac{E_C + E_V - 2E_F}{kT}}$$

Assuming $m_e^* = m_h^*$,

$$1 = e^{\frac{E_C + E_V - 2E_F}{kT}}$$

$$\therefore E_F = \frac{E_C + E_V}{2}$$

$$\therefore E_F = E_i$$

Realistically, $m_e^* > m_h^*$. Therefore, E_F is below E_i . However, as the difference between the effective masses, and hence E_F and E_i is small,

$$E_F \approx E_i$$

This can be used as a practical approximation.

4.2 N-type Semiconductor

In a N-type semiconductor,

As 0 K, all electrons must be either at in the valence band, or at E_d . Therefore, for all electrons to be below E_F , E_F must be between E_d and E_C . Therefore, as E_d is closer to E_C than to E_V , E_F is also closer to E_C than to E_V .

The number of electrons in the conduction band is equal to the area under the curve of f(E), to the right of E_C .

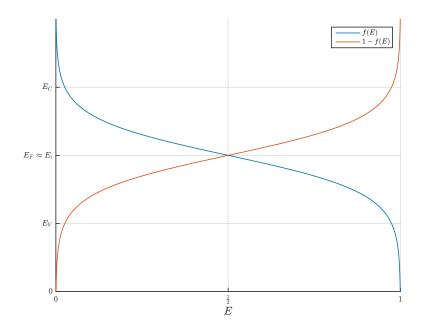


Figure 5: Position of E_F for an Intrinsic semiconductor

4.3 P-type Semiconductor

In a P-type semiconductor,

As 0 K, all electrons must be either at in the valence band, or at E_a . Therefore, for all electrons to be below E_F , E_F must be between E_a and E_C . Therefore, as E_a is closer to E_V than to E_C , E_F is also closer to E_V than to E_C .

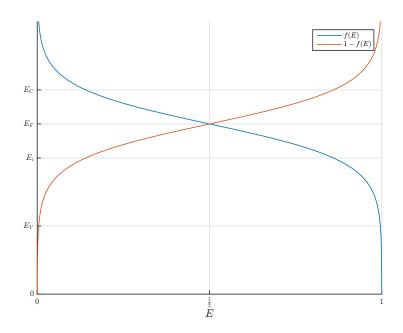


Figure 6: Position of E_F for an N-type semiconductor

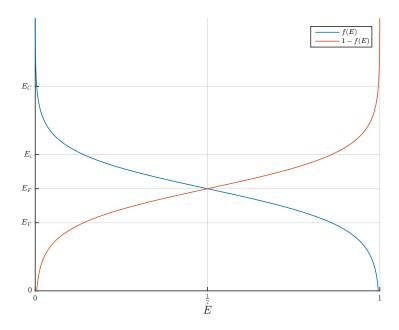


Figure 7: Position of E_F for an P-type semiconductor

5 Graphical Representation of Number of Electrons in the Conduction Band

The number of allowed states in the conduction band is given by

$$g_C(E) = \frac{4\pi}{h^3} (2m_e^*)^{\frac{3}{2}} (E - E_C)^{\frac{1}{2}}$$

$$\approx \sqrt{E - E_C}$$

The probability of an energy state E being occupied is given by

$$f(E) = \frac{1}{1 + e^{\frac{E - E_F}{kT}}}$$

Therefore, the electron distribution in the conduction band is

$$g_C(E)f(E) = \sqrt{E - E_C} \frac{1}{1 + e^{\frac{E - E_F}{kT}}}$$

Therefore, as $g_C(E)$ is zero for $E < E_C$, the graph of the electron distribution function is as in Figure 8.

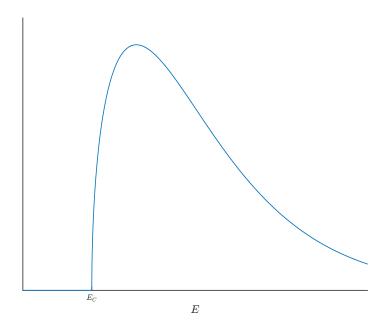


Figure 8: Electron distribution function in the conduction band

6 Maxwell-Boltzmann Approximation

The number of allowed states in the gap between the valence and conduction bands is zero. Therefore, although the Fermi function is non zero in the energy gap, the number of electrons in the conduction band can be approximated as

$$\frac{1}{1 + e^{\frac{E - E_F}{kT}}} \approx e^{-\frac{E - E_F}{kT}}$$

Therefore, integrating and solving,

$$n = N_C e^{\frac{E_C - E_F}{kT}}$$

where N_C is a constant which represents the effective density of states, and not a function of E.

Therefore, at equilibrium,

$$n_0 = N_C e^{-\frac{E_C - E_F}{kT}}$$
$$p_0 = N_V e^{-\frac{E_F - E_V}{kT}}$$

Therefore,

$$n_i^2 = n_0 p_0$$

$$= N_C N_V e^{-\frac{E_C - E_V}{kT}}$$

$$\therefore n_i = \sqrt{N_C N_V} e^{-\frac{E_{\text{gap}}}{2kT}}$$

Therefore,

$$n = N_C e^{-\frac{E_C - E_F}{kT}}$$

$$= N_C e^{-\frac{E_C - E_i}{kT}} e^{-\frac{E_i - E_F}{kT}}$$

For an intrinsic material, i.e. when $E_F = E_i$,

$$n = n_i$$

Therefore,

$$n = n_i e^{\frac{E_F - E_i}{kT}}$$

Similarly,

$$p = n_i e^{\frac{E_i - E_F}{kT}}$$

Therefore,

$$E_F - E_i = kT \ln \left(\frac{n}{n_i}\right)$$
$$E_i - E_F = kT \ln \left(\frac{p}{n_i}\right)$$

Exercise 1.

Consider a sample of Si, doped with

$$N_D = 10^{16} \frac{1}{\text{cm}^3}$$

such that

$$E_C - E_D = 0.05 \,\text{eV}$$

- 1. Draw E_F as a function of T.
- 2. How many dopants are ionized when $E_F = E_D$?
- 3. Given

$$N_C = 5 \times 10^{18} \frac{1}{\text{cm}^3}$$

calculate the temperature when

$$E_F = E_D$$

Solution 1.

1. Let the boundaries between the ionization and extrinsic regions and the extrinsic and intrinsic regions be T_1 and T_2 respectively. Therefore, for the ionization region, i.e. for $T < T_1$,

$$f(E < E_D) = 1$$

Therefore, E_F is between E_D and E_C .

As the temperature increases, the some of the electrons which were previously at E_D move to E_C . Therefore, the probability of electrons occupying the states at E_D reduces, and that of E_C increases. Therefore, the graph of f(E) spreads out. Therefore, E_F moves lower, and approaches E_D .

Similarly, in the extrinsic region, E_F is below E_D , and continues to move lower with increase in temperature. As the intrinsic carriers start ionizing, the rate of lowering of E_F increases.

In the intrinsic region,

$$E_F = E_i$$

2.

$$f(E_F) = \frac{1}{2}$$
$$\therefore f(E_D) = \frac{1}{2}$$

Therefore, exactly half of the dopant states are occupied. Therefore, exactly half of the dopants are ionized.

3.

$$n = N_C e^{-\frac{E_C - E_f}{kT}}$$

$$\therefore \frac{N_D}{2} = N_C e^{-\frac{E_C - E_D}{kT}}$$

$$\therefore \ln\left(\frac{N_D}{2N_C}\right) = -\frac{E_C - E_D}{kT}$$

$$\therefore \ln\left(\frac{2N_C}{N_D}\right) = \frac{E_C - E_D}{kT}$$

Therefore,

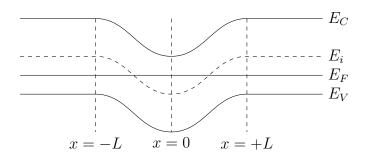
$$T = \frac{E_C - E_D}{k \ln\left(\frac{2N_C}{N_D}\right)}$$

$$= \frac{0.05 \text{eV}}{\left(8.6 \times 10^{-5} \frac{\text{eV}}{\text{K}}\right) \ln\left(\frac{2.5 \times 10^{18}}{10^{16}}\right)}$$

$$= 85 \text{K}$$

Exercise 2.

Consider a sample of Si at 300 K. The band diagram is as given.



Assume that at x = 0,

$$E_F - E_i = \frac{E_{\rm gap}}{4}$$

and at $x = \pm L$,

$$E_i - E_F = \frac{E_{\text{gap}}}{4}$$

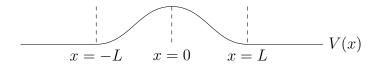
- 1. Sketch V(x), the electrostatic potential in the sample.
- 2. Sketch the electric field $\mathcal{E}(x)$.
- 3. Identify the different regions.
- 4. Sketch the concentrations of the electrons and the holes.
- 5. Find the directions of the electron currents.

Solution 2.

1.

$$E = -qV$$

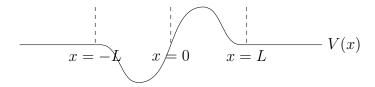
Therefore, the graph of V(x) is



2. By Poisson's Equation,

$$\frac{\mathrm{d}V}{\mathrm{d}x} = -\mathcal{E}$$

Therefore, the graph of the field is of the form



3. For x < -L, E_F is below E_i . Therefore, the material is P-type. Therefore,

$$p = n_i e^{\frac{E_i - E_f}{kT}}$$

$$= 10^{10} e^{\frac{E_{\text{gap}}}{4kT}}$$

$$= 3.92 \times 10^{14} \frac{1}{\text{cm}^3}$$

Therefore,

$$n = \frac{n_i^2}{p}$$

$$= \frac{10^{20}}{3.92 \times 10^{14}}$$

$$= 2.55 \times 10^5 \frac{1}{\text{cm}^3}$$

For x = 0, E_F is above E_i . Therefore, the material is N-type. Therefore,

$$n = n_i e^{\frac{E_i - E_f}{kT}}$$

$$= 10^{10} e^{\frac{E_{\text{gap}}}{4kT}}$$

$$= 3.92 \times 10^{14} \frac{1}{\text{cm}^3}$$

Therefore,

$$p = \frac{n_i^2}{n}$$

$$= \frac{10^{20}}{3.92 \times 10^{14}}$$

$$= 2.55 \times 10^5 \frac{1}{\text{cm}^3}$$

For x < -L, E_F is below E_i . Therefore, the material is P-type. Therefore,

$$p = n_i e^{\frac{E_i - E_f}{kT}}$$

$$= 10^{10} e^{\frac{E_{\text{gap}}}{4kT}}$$

$$= 3.92 \times 10^{14} \frac{1}{\text{cm}^3}$$

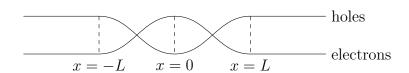
Therefore,

$$n = \frac{n_i^2}{p}$$

$$= \frac{10^{20}}{3.92 \times 10^{14}}$$

$$= 2.55 \times 10^5 \frac{1}{\text{cm}^3}$$

4.



5. The concentration of the electrons is higher at the centre that at the ends. Therefore, the electrons flow outwards. Therefore, the electron diffusion current is directed outwards.

The electric field is directed leftwards for x < 0, and rightwards for x > 0. Therefore, the electron drift current is directed inwards.

7 Space Charge Neutrality

For a material to be neutral, the total negative charges must be equal to the total positive charges. Therefore,

$$n_0 + N_A^- = p_0 + N_D^-$$

Also,

$$n_0 = N_C e^{-\frac{E_C - E_F}{kT}}$$

$$p_0 = N_V e^{-\frac{E_F - E_V}{kT}}$$

$$N_A^- = N_A f(E_A)$$

$$N_D^+ = N_D \left(1 - f(E_A)\right)$$

$$N_C e^{-\frac{E_C - E_F}{kT}} + N_A f(E_A) = N_V e^{-\frac{E_F - E_V}{kT}} + N_D (1 - f(E_D))$$

Part II

PN Junctions

1 Step Junction

Definition 2 (Electron affinity). The electron affinity for a semiconductor is defined to be

$$q\chi = E_{\rm vac} - E_C$$

where E_{vac} is the energy of a free particle. The electron affinity is material dependent.

Definition 3 (Semiconductor work function). The semiconductor work function is defined to be

$$q\Phi_S = E_{\rm vac} - E_F$$

The semiconductor work function is doping dependent.

Consider a P-type semiconductor with $p = N_A$ and a N-type semiconductor with $n = N_D$.

Let the two be placed separately.

Let E_{F_p} and E_{F_n} be the Fermi levels for the P-type and the N-type semiconductors respectively.

Therefore,

$$q\chi = E_{\text{vac}} - E_C$$
$$q\Phi_{S_p} = E_{\text{vac}} - E_{F_p}$$
$$q\Phi_{S_n} = E_{\text{vac}} - E_{F_n}$$

Let these semiconductors be brought into contact as in Figure 9 and allowed to reach equilibrium. At equilibrium, the Fermi levels must match up, and E_F must

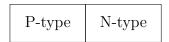


Figure 9: Structure of Step PN Junction

be constant.

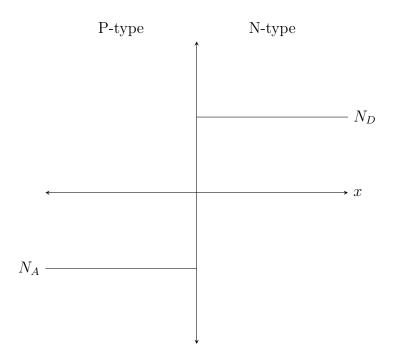


Figure 10: Carrier Concentrations in Step PN Junction

The carrier concentrations in the junction are as in Figure 10. Therefore, the carriers diffuse from regions of higher concentration to regions of lower concentrations, i.e., the electrons diffuse from the N-type region to the P-type region, and the holes diffuse from the P-type region to the N-type region.

This diffusion occurs in a region closer to the junction, and not in the regions near the ends. This region is called the depletion region.

As these mobile carriers diffuse, they leave behind the ionized dopants, which are fixed charges. This causes an internal electric field \mathcal{E} in the depletion region, directed from the N-type region to the P-type region. This generated electric field repels the diffusing carriers. Therefore, the diffusion is stopped. The width of the depletion region when the diffusion stops is denoted by W or W_d .

The width of the depletion region on the P-type side is denoted by x_p , and that on the N-type side is denoted by x_n .

The currents generated due to the drifting charge carriers are opposite in direction to the currents generated due to the diffusing charge carriers. Therefore, the net current in the PN junction is zero.

1.1 Charge Density

Let the cross-sectional area of the junction be A. Therefore, the total positive charge in the P-type region is

$$Q_{+} = qN_{D}x_{n}A$$

where q is the charge on an electron.

Therefore, the charge density in the N-type region is

$$\rho = \frac{Q_+}{x_n A}$$
$$= q N_D$$

Similarly, the charge density in the P-type region is

$$\rho = qN_A$$

Therefore, as the junction is electrically neutral,

$$Q_{+} = Q_{-}$$

$$\therefore qN_{D}x_{n}A = qN_{A}x_{p}A$$

$$\therefore N_{D}x_{n} = N_{A}x_{p}$$

1.2 Electric Field

By Gauss' Law,

$$\frac{\mathrm{d}\mathcal{E}}{\mathrm{d}x} = \frac{\rho}{\varepsilon\varepsilon_0}$$

where ε is the relative permittivity of the material, and ε_0 is the permittivity of vacuum.

The charge density in the junction is

$$\rho(x) = \begin{cases} -qN_A & ; & -x_p < x < 0 \\ qN_D & ; & 0 < x < x_n \end{cases}$$

Therefore, integrating,

$$\mathcal{E}(x) = \begin{cases} -\frac{qN_A}{\varepsilon\varepsilon_0}x + c_1 & ; & -x_p < x < 0\\ \frac{qN_D}{\varepsilon\varepsilon_0}x + c_2 & ; & 0 < x < x_n \end{cases}$$

At the boundaries of the depletion region,

$$\mathcal{E}(-x_p) = 0$$

$$\mathcal{E}(x_n) = 0$$

Therefore, substituting,

$$c_1 = -\frac{qN_A}{\varepsilon \varepsilon_0} x_p$$
$$c_2 = -\frac{qN_D}{\varepsilon \varepsilon_0} x_n$$

Therefore,

$$\mathcal{E}(x) = \begin{cases} -\frac{qN_A}{\varepsilon\varepsilon_0} x - \frac{qN_A}{\varepsilon\varepsilon_0} & ; & -x_p < x < 0\\ \frac{qN_D}{\varepsilon\varepsilon_0} x - \frac{qN_D}{\varepsilon\varepsilon_0} & ; & 0 < x < x_n \end{cases}$$

1.3 Electrostatic Potential

$$\frac{\mathrm{d}V}{\mathrm{d}x} = -\mathcal{E}$$

The electric field in the junction is

$$\mathcal{E}(x) = \begin{cases} -\frac{qN_A}{\varepsilon\varepsilon_0} x - \frac{qN_A}{\varepsilon\varepsilon_0} & ; & -x_p < x < 0\\ \frac{qN_D}{\varepsilon\varepsilon_0} x - \frac{qN_D}{\varepsilon\varepsilon_0} & ; & 0 < x < x_n \end{cases}$$

Therefore, integrating,

$$V(x) = \begin{cases} \frac{qN_A}{\varepsilon\varepsilon_0} \left(\frac{x^2}{2} + x_p x + c_1\right) & ; \quad -x_p < x < 0 \\ -\frac{qN_D}{\varepsilon\varepsilon_0} \left(\frac{x^2}{2} - x_n x + c_2\right) & ; \quad 0 < x < x_n \end{cases}$$

At the boundaries of the depletion region,

$$V(-x_p) = 0$$
$$V(x_n) = V_{\rm BI}$$

Therefore, substituting,

$$c_1 = \frac{x_p^2}{2}$$

$$c_2 = -\frac{\varepsilon \varepsilon_0 V_{\text{BI}}}{q N_D} + \frac{x_n^2}{2}$$

Therefore,

$$V(x) = \begin{cases} \frac{qN_A}{\varepsilon\varepsilon_0} \left(\frac{x^2}{2} + x_p x + \frac{x_p^2}{2}\right) & ; \quad -x_p < x < 0 \\ \frac{qN_D}{\varepsilon\varepsilon_0} \left(\frac{x^2}{2} - x_n x + \frac{x_n^2}{2}\right) + V_{\text{BI}} \\ = \begin{cases} \frac{qN_A}{2\varepsilon\varepsilon_0} (x + x_p)^2 & ; \quad -x_p < x < 0 \\ -\frac{qN_D}{2\varepsilon\varepsilon_0} (x - x_n)^2 + V_{\text{BI}} & ; \quad 0 < x < x_n \end{cases}$$

1.4 Energy Bands

$$\frac{\mathrm{d}E}{\mathrm{d}x} = -qV_{\mathrm{BI}}$$

Therefore, the energy band diagram for the junction is as in Figure 11.

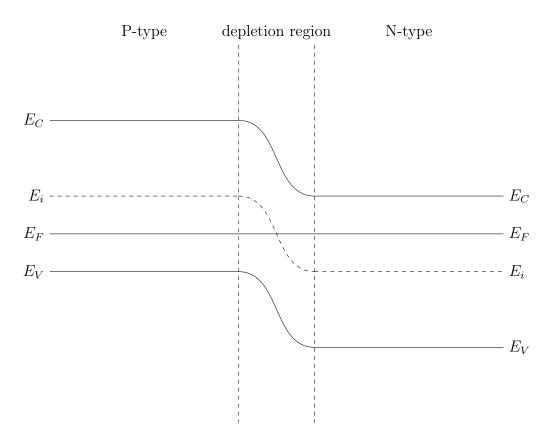


Figure 11: Energy Band Diagram for PN Junction

By definition, the builtin voltage is

$$qV_{\text{BI}} = (E_C)_{\text{P-type side}} - (E_C)_{\text{N-type side}}$$

$$= (E_i - E_F)_{\text{P-type side}} + (E_F - E_i)_{\text{N-type side}}$$

$$= kT \ln \left(\frac{p}{n_i}\right) + kT \ln \left(\frac{n}{n_i}\right)$$

$$= kT \ln \left(\frac{N_A}{n_i}\right) + kT \ln \left(\frac{N_D}{n_i}\right)$$

$$= kT \ln \left(\frac{N_A N_D}{n_i^2}\right)$$

$$\therefore V_{\text{BI}} = \frac{kT}{q} \ln \left(\frac{N_A N_D}{n_i^2}\right)$$

1.5 Width of Depletion Region

For step junctions,

$$W = \sqrt{\frac{2\varepsilon\varepsilon_0 V_{\rm BI}}{q} \left(\frac{N_A + N_D}{N_A N_D}\right)}$$
$$x_p = \frac{W N_D}{N_A + N_D}$$
$$x_n = \frac{W N_A}{N_A + N_D}$$

Exercise 3.

A silicon PN step junction at equilibrium at 300 K has

$$N_A = 10^{16} \frac{1}{\text{cm}^3}$$
$$N_D = 10^{16} \frac{1}{\text{cm}^3}$$

- 1. Sketch $\rho(x)$.
- 2. Calculate \mathcal{E}_{max} .

Solution 3.

1.

$$V_{\rm BI} = \frac{kT}{q} \ln \left(\frac{N_A N_D}{n_i^2} \right)$$
$$= (0.026 \, \text{V}) \ln \left(\frac{10^{32}}{10^{20}} \right)$$
$$= 0.718 \, \text{V}$$

Therefore,

$$W = \sqrt{\frac{2\varepsilon\varepsilon_0 V_{\rm BI}}{q} \left(\frac{N_A + N_D}{N_A N_D}\right)}$$

$$= \sqrt{\frac{(2)(11.8) \left(8.85 \times 10^{-14} \frac{\rm F}{\rm cm}\right) (0.718 \, \rm V)}{1.6 \times 10^{-19} \, \rm C}} \left(\frac{2 \times 10^{16}}{10^{32}} \rm cm^3\right)$$

$$= 4.32 \times 10^{-5} \rm cm$$

$$= 0.432 \, \mu m$$

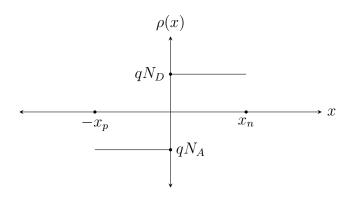
Therefore,

$$N_D x_n = N_A x_p$$

$$\therefore x_n = x_p$$

$$= \frac{W}{2}$$

$$= 0.216 \mu m$$



2. The field is maximum at the interface of the two regions. Therefore,

$$\begin{split} \mathcal{E}_{\text{max}} &= \frac{q N_D x_p}{\varepsilon \varepsilon_0} \\ &= \frac{\left(1.6 \times 10^{-19} \, \text{C}\right) \left(10^{16} \frac{1}{\text{cm}^3}\right) \left(2.16 \times 10^{-5} \text{cm}\right)}{(11.8) \left(8.85 \times 10^{-14} \frac{\text{F}}{\text{cm}}\right)} \\ &= 3.31 \times 10^4 \frac{\text{V}}{\text{cm}} \end{split}$$

Exercise 4.

A silicon PN step junction at equilibrium at $300\,\mathrm{K}$ has

$$N_A = 10^{16} \frac{1}{\text{cm}^3}$$

$$N_D = 10^{17} \frac{1}{\text{cm}^3}$$

- 1. Sketch $\rho(x)$.
- 2. Sketch $\mathcal{E}(x)$.
- 3. Sketch V(x).

Solution 4.

1.

$$N_A x_p = N_D x_n$$

$$\therefore 10^{16} x_p = 10^{17} x_n$$

$$\therefore x_p = 10 x_n$$

Therefore,

$$V_{\rm BI} = \frac{kT}{q} \ln \left(\frac{N_A N_D}{n_i^2} \right)$$
$$= 0.778 \,\mathrm{V}$$

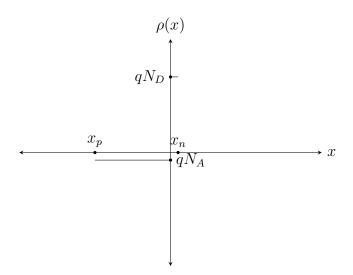
$$W = \sqrt{\frac{2\varepsilon\varepsilon_0 V_{\rm BI}}{q} \frac{N_A + N_D}{N_A N_D}}$$
$$= 0.334 \mu \rm m$$

Therefore,

$$x_n = 0.031 \mu \text{m}$$

 $x_p = 0.303 \mu \text{m}$

Therefore,



2. The field is maximum at the interface of the two regions.

$$\mathcal{E}_{max} = \frac{qN_D x_n}{\varepsilon \varepsilon_0}$$

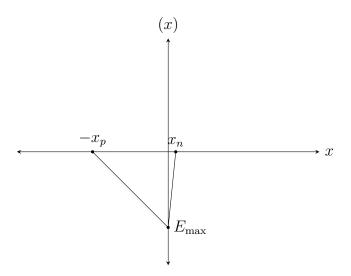
$$= \frac{\left(1.6 \times 10^{-19}\right) \left(10^{17}\right) \left(0.031 \times 10^{-4}\right)}{\left(11.8\right) \left(8.85 \times 10^{-14}\right)}$$

$$= 4.75 \times 10^4 \frac{V}{cm}$$

The field at the ends of the depletion region is

$$\mathcal{E}(-x_p) = 0$$

$$\mathcal{E}(x_n) = 0$$

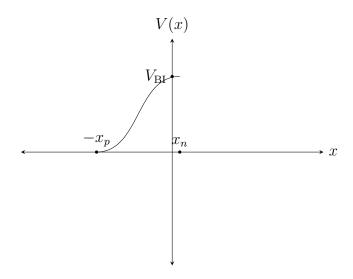


3. Let the potential at $x = -x_p$ be

$$V(-x_p) = 0$$

Therefore, by definition,

$$V(x_n) = V_{\rm BI}$$



Exercise 5.

A PiN junction is constructed as by inserting a thin intrinsic semiconductor of thickness x_i between the regions of a PN junction diode. It is kept at 300 K at equilibrium.

- 1. Find $\rho(x)$.
- 2. Find $\mathcal{E}(x)$.
- 3. Find V(x).

Solution 5.

1.

$$Q_{+} = qN_{D}A\left(x_{n} - \frac{x_{i}}{2}\right)$$
$$Q_{-} = qN_{A}A\left(x_{p} - \frac{x_{i}}{2}\right)$$

Therefore, as the sample is electrically neutral,

$$Q_{+} = Q_{-}$$

$$\therefore N_{D} \left(x_{n} - \frac{x_{i}}{2} \right) = N_{A} \left(x_{p} - \frac{x_{i}}{2} \right)$$

Therefore,

$$\rho(x) = \begin{cases} -qN_A & ; & -x_p < x < -\frac{x_i}{2} \\ 0 & ; & -\frac{x_i}{2} < x < \frac{x_i}{2} \\ qN_D & ; & \frac{x_i}{2} < x < x_n \end{cases}$$

2.

$$\frac{\mathrm{d}\mathcal{E}}{\mathrm{d}x} = \frac{\rho}{\varepsilon\varepsilon_0}$$

$$\mathcal{E}(x) = \begin{cases} -\frac{qN_A}{\varepsilon\varepsilon_0}x + c_1 & ; & -x_p < x < -\frac{x_i}{2} \\ c_2 & ; & -\frac{x_i}{2} < x < \frac{x_i}{2} \\ \frac{qN_D}{\varepsilon\varepsilon_0}x + c_2 & ; & \frac{x_i}{2} < x < x_n \end{cases}$$

The electric field at the ends of the depletion region is

$$\mathcal{E}(-x_p) = 0$$

$$\mathcal{E}(x_n) = 0$$

The electric field must be continuous at the ends of the intrinsic region. Therefore, solving,

$$c_{1} = -\frac{qN_{A}x_{p}}{\varepsilon\varepsilon_{0}}$$

$$c_{2} = -\frac{qN_{A}}{\varepsilon\varepsilon_{0}} \left(x_{p} - \frac{x_{i}}{2}\right)$$

$$= \frac{qN_{D}}{\varepsilon\varepsilon_{0}} \left(\frac{x_{i}}{2} - x_{n}\right)$$

$$c_{3} = -\frac{qN_{D}x_{n}}{\varepsilon\varepsilon_{0}}$$

Therefore,

$$\mathcal{E}(x) = \begin{cases} -\frac{qN_A}{\varepsilon\varepsilon_0} (x + x_p) & ; \quad -x_p < x < -\frac{x_i}{2} \\ -\frac{qN_A}{\varepsilon\varepsilon_0} (x_p - \frac{x_i}{2}) & ; \quad -\frac{x_i}{2} < x < \frac{x_i}{2} \\ \frac{qN_D}{\varepsilon\varepsilon_0} (x - x_n) & ; \quad \frac{x_i}{2} < x < x_p \end{cases}$$

3.

$$\frac{\mathrm{d}V}{\mathrm{d}x} = -\mathcal{E}$$

Therefore,

$$V(x) = \begin{cases} \frac{qN_A}{\varepsilon\varepsilon_0} \left(\frac{x^2}{2} + x_p x + c_1\right) & ; \quad -x_p < x < -\frac{x_i}{2} \\ \frac{qN_A}{\varepsilon\varepsilon_0} \left(x_p - \frac{x_i}{2}\right) x + c_2 & ; \quad -\frac{x_i}{2} < x < \frac{x_i}{2} \\ -\frac{qN_D}{\varepsilon\varepsilon_0} \left(\frac{x^2}{2} - x_n x + c_3\right) & ; \quad \frac{x_1}{2} < x < x_n \end{cases}$$

Let the potential at $x = -x_p$ be

$$V(-x_p) = 0$$

Therefore, by definition,

$$V(x_n) = V_{\rm BI}$$

Therefore, solving and sustituting,

$$V(x) = \begin{cases} \frac{qN_A}{2\varepsilon\varepsilon_0} (x + x_p)^2 & ; & -x_p < x < -\frac{x_i}{2} \\ \frac{qN_A}{\varepsilon\varepsilon_0} \left(x_p - \frac{x_i}{2} \right) \left(x_p + \frac{x_i}{2} + 2x \right) & ; & -\frac{x_i}{2} < x < \frac{x_i}{2} \\ V_{\text{BI}} - \frac{qN_D}{2\varepsilon\varepsilon_0} (x_n - x)^2 & ; & \frac{x_1}{2} < x < x_n \end{cases}$$

2 PN Junction with Linearly Graded Doping Profile

Consider a PN junction such that

$$N_D - N_A = ax$$

where a > 0.

Therefore, the doping profile is as in Figure 12.

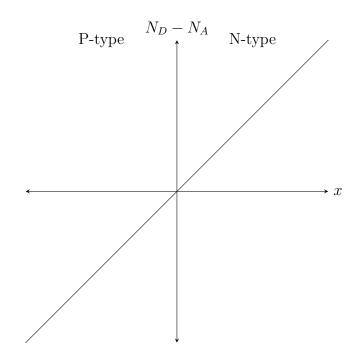


Figure 12: Doping Profile in Linearly Doped PN Junction

2.1 Charge Density

$$\rho(x) = \begin{cases} qax & ; & -x_p \le x \le 0 \\ qax & ; & 0 \le x \le x_p \end{cases}$$
$$= qax$$

for $-x_p \le x \le x_n$. Therefore, the charge density is as in Figure 13.

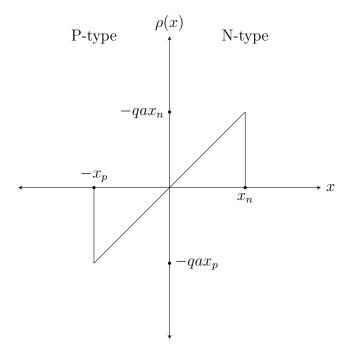


Figure 13: Charge Distribution in Linearly Doped PN Junction

2.2 Electric Field

By Gauss' Law,

$$\mathcal{E}(x) = \frac{qa}{\varepsilon\varepsilon_0} \left(\frac{x^2}{2} + c \right)$$

for $-x_p \le x \le x_n$.

As the electric field must be continuous,

$$\mathcal{E}(-x_p) = 0$$

$$\mathcal{E}(x_n) = 0$$

Therefore, solving and substituting,

$$\mathcal{E}(x) = \frac{q\alpha}{2\varepsilon\varepsilon_0} \left(x^2 - \frac{W^2}{4} \right)$$

2.3 Width of Depletion Region

As the junction is electrically neutral,

$$\frac{qax_n^2}{2} = \frac{qax_p^2}{2}$$

$$\therefore x_n = x_p$$

$$= \frac{W}{2}$$

Also,

$$\frac{\mathrm{d}V}{\mathrm{d}x} = -\mathcal{E}(x)$$

$$\therefore V(x) = -\frac{qa}{2\varepsilon\varepsilon_0} \left(\frac{x^3}{3} - \frac{W^2}{4}x + c\right)$$

Let the potential at $-x_p$ be zero.

Therefore, solving and substituting,

$$V(x) = -\frac{qa}{6\varepsilon\varepsilon_0} \left(x^3 - 3\frac{W^2}{4}x - \frac{2}{8}W^3 \right)$$

Therefore, by definition,

$$V_{\text{BI}} = V(x_n)$$

$$= -\frac{qa}{6\varepsilon\varepsilon_0} \left(x_n^3 - 3\frac{W^2}{4}x_n - \frac{2}{8}W^3 \right) = -\frac{qa}{6\varepsilon\varepsilon_0} \left(\frac{W^3}{8} - 4\frac{W^2}{4}\frac{W}{2} - \frac{2}{8}W^3 \right)$$

$$= \frac{qa}{12\varepsilon\varepsilon_0}W^3$$

$$\therefore W = \left(\frac{12V_{\text{BI}}\varepsilon\varepsilon_0}{q\alpha} \right)^{\frac{1}{3}}$$

2.4 Electrostatic Potential

$$\mathcal{E}(x) = -\frac{kT}{q} \frac{\frac{\mathrm{d}n}{\mathrm{d}x}}{n}$$
$$= -\frac{kT}{q} \frac{\frac{\mathrm{d}\alpha x}{\mathrm{d}x}}{\frac{\mathrm{d}x}{\alpha x}}$$
$$= -\frac{kT}{q} \frac{1}{x}$$

Therefore,

$$V(x) \propto \ln(x)$$

Therefore,

$$V_{\text{BI}} = \frac{kT}{q} \ln \left(\frac{N_D(x_n) N_A(-x_p)}{n_i^2} \right)$$
$$= \frac{kT}{q} \ln \left(\frac{\alpha x_n \alpha x_p}{n_i^2} \right)$$
$$= \frac{kT}{q} \ln \left(\frac{\alpha^2 \left(\frac{W}{2} \right)^2}{n_i^2} \right)$$

Exercise 6.

Consider a silicon PN junction, at equilibrium at 300 K, with doping

$$N_D - N_A = \begin{cases} -N_A & ; \quad x < 0 \\ \alpha x & ; \quad x > 0 \end{cases}$$

Find

- 1. Charge density
- 2. Relation between x_n and x_p
- 3. Electric field
- 4. Electrostatic potential

Solution 6.

1.

$$\rho(x) = \begin{cases} -qN_A & ; & x < 0 \\ q\alpha & ; & x > 0 \end{cases}$$

2. As the junction is electrically neutral, the area under the curve of $\rho(x)$ on the P-type side must be equal to that on the N-type side. Therefore,

$$qN_A x_p = \frac{1}{2} (q\alpha x_n)(x_n)$$
$$\therefore N_A x_p = \frac{1}{2} \alpha x_n^2$$

3.

$$\mathcal{E}(x) = \begin{cases} -\frac{qN_A}{\varepsilon\varepsilon_0}(x+c_1) & ; & -x_p \le x \le 0\\ \frac{q\alpha}{\varepsilon\varepsilon_0}\left(\frac{x^2}{2}+c_2\right) & ; & 0 \le x \le x_n \end{cases}$$

At the boundaries of the depletion region, the electric field must be zero. Therefore,

$$\mathcal{E}(-x_p) = 0$$

$$\mathcal{E}(x_n) = 0$$

Therefore, solving,

$$c_1 = x_p$$

$$c_2 = -\frac{{x_n}^2}{2}$$

Therefore,

$$\mathcal{E}(x) = \begin{cases} -\frac{qN_A}{\varepsilon\varepsilon_0}(x + x_p) & ; \quad -x_p \le x \le 0\\ \frac{q\alpha}{2\varepsilon\varepsilon_0}(x_2 - x_n^2) & ; \quad 0 \le x \le x_n \end{cases}$$

4.

$$\frac{\mathrm{d}V}{\mathrm{d}x} = \mathcal{E}(x)$$

$$\therefore V(x) = \begin{cases} \frac{qN_A}{\varepsilon\varepsilon_0} \left(\frac{x^2}{2} + x_p x + c_1\right) & ; \quad -x_p \le x \le 0\\ -\frac{q\alpha}{\varepsilon\varepsilon_0} \left(\frac{x^3}{3} - x_n^2 x + c_2\right) & ; \quad 0 \le x \le x_n \end{cases}$$

Let the potential at $-x_p$ be zero. Therefore, by definition,

$$V_{\rm BI} = V(x_n)$$

Therefore, substituting and solving,

$$c_1 = \frac{x_p^2}{2}$$

$$c_2 = -\frac{2\varepsilon\varepsilon_0 V_{\text{BI}}}{q\alpha} + \frac{2}{3}x_n^3$$

$$V(x) = \begin{cases} \frac{qN_A}{\varepsilon\varepsilon_0} \left(\frac{x^2}{2} + x_p x + \frac{x_p^2}{2}\right) & ; \quad -x_p \le x \le 0\\ -\frac{q\alpha}{\varepsilon\varepsilon_0} \left(\frac{x^3}{3} - x_n^2 x - \frac{2\varepsilon\varepsilon_0 V_{\text{BI}}}{q\alpha} + \frac{2}{3}x_n^3\right) & ; \quad 0 \le x \le x_n \end{cases}$$

3 Biased PN Junctions

Consider a PN step junction at equilibrium.

Let the width of the depletion region on the P-side be x_{p_0} , and that on the N-type side be x_{n_0} .

Let the builtin voltage be $V_{\rm BI}$.

Let the internal electric field in the depletion region be \mathcal{E} . Therefore, \mathcal{E} is directed from the N-type region to the P-type region.

Let the diode be connected to a voltage source as in Figure 14. Modelling the diode

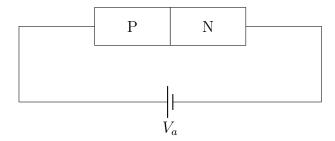


Figure 14: PN Junction in Bias

as resistors, let the resistance of the depletion region be R_D , that of the remaining P-type part be R_P , and that of the remaining N-type part be R_N .

The resistivity in a region is

$$\rho = \frac{1}{q(\mu_n n + \mu_p p)}$$

In the depletion region, there are very few free charge carriers, compared to the remaining P-type and N-type regions. Therefore,

$$\rho_D >> R_P$$
 $\rho_D >> R_N$

Therefore,

$$R_D >> R_P$$

 $R_D >> R_N$

Therefore, for practical purposes, it can be assumed that the entire voltage drop in on the depletion region.

Therefore,

$$E(x) = -qV(x)$$

Therefore, if V(x) is increased, the energy bands shift downwards, as compared to their equilibrium positions. If V(x) is decreased, the energy bands shift upwards, as compared to their equilibrium positions.

Therefore, if the applied external voltage V_a is positive, the energy bands shift upwards, and if it is negative, the energy bands shift downwards.

Therefore, the difference between the energy bands at the two terminals, increases by qV_a .

If V_a is positive, the PN junction is said to be in forward bias, and if it is negative, the PN junction is said to be in reverse bias.

Therefore, due to V_a , there will be an electric field generated, directed in the direction of V_a .

If $V_a > 0$, the generated electric field opposes the internal electric field. Therefore, the net electric field is smaller in magnitude than at equilibrium.

Therefore, the width of the depletion region is smaller than that at equilibrium.

By Gauss' Law, the slope of the curve of the electric field represents the charge density. Therefore, as the charge density must be the same as at equilibrium, the slope of the electric field also must be the same as at equilibrium.

Exercise 7.

Consider a PN step junction at 300 K. Given

$$N_A = 10^{16} \frac{1}{\text{cm}^3}$$
$$N_D = 10^{16} \frac{1}{\text{cm}^3}$$

find the width of the depletion region on each side of the junction, for

1.
$$V_a = 0$$

2.
$$V_a = 0.5 \,\mathrm{V}$$

3.
$$V_a = -0.5 \,\text{V}$$

Solution 7.

1. As the junction is a step junction,

$$W = \sqrt{\frac{2\varepsilon\varepsilon_0}{q}(V_{\rm BI} + V_a)\left(\frac{N_A + N_D}{N_A N_D}\right)} = \sqrt{\frac{2\varepsilon\varepsilon_0}{q}(V_{\rm BI} + 0)\left(\frac{N_A + N_D}{N_A N_D}\right)}$$

Therefore, substituting and solving,

$$W = 0.432 \mu m$$

Why?

Add breakdown of Fermi levels As the junction is electrically neutral,

$$N_A x_p = N_D x_n$$

$$\therefore x_p = x_n$$

$$= \frac{W}{2}$$

Therefore,

$$x_p = x_n$$
$$= 0.216 \mu m$$

2. As the junction is a step junction,

$$W = \sqrt{\frac{2\varepsilon\varepsilon_0}{q}(V_{\rm BI} + V_a)\left(\frac{N_A + N_D}{N_A N_D}\right)} = \sqrt{\frac{2\varepsilon\varepsilon_0}{q}(V_{\rm BI} + 0.5\,{\rm V})\left(\frac{N_A + N_D}{N_A N_D}\right)}$$

Therefore, substituting and solving,

$$W = 0.238 \mu m$$

As the junction is electrically neutral,

$$N_A x_p = N_D x_n$$

$$\therefore x_p = x_n$$

$$= \frac{W}{2}$$

Therefore,

$$x_p = x_n$$
$$= 0.119 \mu m$$

3. As the junction is a step junction,

$$W = \sqrt{\frac{2\varepsilon\varepsilon_0}{q}(V_{\rm BI} + V_a)\left(\frac{N_A + N_D}{N_A N_D}\right)} = \sqrt{\frac{2\varepsilon\varepsilon_0}{q}(V_{\rm BI} - 0.5\,{\rm V})\left(\frac{N_A + N_D}{N_A N_D}\right)}$$

Therefore, substituting and solving,

$$W = 0.563 \mu m$$

As the junction is electrically neutral,

$$N_A x_p = N_D x_n$$

$$\therefore x_p = x_n$$

$$= \frac{W}{2}$$

Therefore,

$$x_p = x_n$$
$$= 0.281.5 \mu m$$

4 Currents in PN Junctions

At equilibrium, the concentration of electrons is higher in the N-type region is higher than that in the P-type region. Therefore, they tend to diffuse from the N-type region to the P-type region. However, as there is an energy barrier in the depletion region, only the electrons with $E > E_C$ can diffuse and cross over. Therefore, an electron diffusion current is generated.

Similarly, the holes with $E < E_V$ diffuse from the P-type region to the N-type region, and a hole diffusion current is generated.

When the electrons and holes diffuse and cross over into the other region, they leave behind fixed charges. This results in an internal electric field being generated in the depletion region.

Due to the internal electric field, electrons in the conduction band of the P-type region drift from the P-type region to the N-type region. Therefore, an electron drift current is generated.

Similarly, the holes in the valence band of the N-type region drift from the N-type region to the P-type region, and a hole drift current is generated.

As the junction is at equilibrium,

$$J_{\text{total}} = 0$$

Therefore,

$$J_{\text{diffusion}_p} + J_{\text{drift}_p} = 0$$
$$J_{\text{diffusion}_n} + J_{\text{drift}_n} = 0$$

The magnitude of the diffusion current is determined by the height of the energy barrier.

The magnitude of the drift current is determined by the magnitude of the internal electric field and the concentration of minority carriers only. Therefore, the height of the energy barrier does not affect the magnitude of the drift current.

Therefore, if the height of the energy barrier is lowered, the magnitude of the diffusion currents increases, but that of the drift currents does not.

The electron and hole diffusion currents are directed from the P-type region to the N-type region, and the electron and hole drift currents are directed from the N-type region to the P-type region. In equilibrium, the two are equal in magnitude and opposite in direction. Therefore, there is no net current.

If the junction is in forward bias, the height of the energy barrier is less than that at equilibrium. Therefore, the magnitude of the diffusion current is more than that of the drift current. Therefore, the net current is directed from the P-type side to the N-type side.

Similarly, if the junction is in reverse bias, the height of the energy barrier is greater than that in equilibrium, and hence, the net current is directed from the N-type side to the P-type side.

5 Carrier Profiles

Let the hole concentration in the P-type region be p_P .

Let the electron concentration in the N-type region be n_N .

Let the electron concentration in the P-type region be n_{P0} .

Let the electron concentration at the interface of the P-type region and the depletion region be n_N . Let the hole concentration in the P-type region be p_{N_0} .

Let the electron concentration at the interface of the N-type region and the depletion region be p_N .

$$V_{\text{BI}} = \frac{kT}{q} \ln \left(\frac{N_D(x_n) N_A(-x_p)}{n_i^2} \right)$$
$$= \frac{kT}{q} \ln \left(\frac{n_N p_P}{n_i^2} \right)$$

If the junction is in equilibrium,

$$n_P = \frac{n_i^2}{p_P}$$

$$= n_N e^{-\frac{qV_{\text{BI}}}{kT}}$$

$$= n_{P0}$$

$$p_N = \frac{n_i^2}{n_N}$$

$$= p_P e^{-\frac{qV_{\text{BI}}}{kT}}$$

$$= p_{N0}$$

If the junction is in bias,

$$\begin{split} n_P &= n_N e^{-\frac{q(V_{\rm BI} - V_a)}{kT}} \\ &= n_{P0} e^{\frac{qV_a}{kT}} \\ p_N &= p_P e^{-\frac{q(V_{\rm BI} - V_a)}{kT}} \\ &= p_{N0} e^{\frac{qV_a}{kT}} \end{split}$$

In general, if the junction is in forward bias, the curves of the minority carriers bend upwards in the form of an exponential, and in reverse bias, the curves bend downwards in the form of an exponential.

The P-type and N-type regions can be considered as semiconductors illuminated from one end only. Therefore, setting up the x-axes for each of the region such that origin is at the interface of the respective region and the depletion region, with the positive direction directed outwards,

$$n_{P} = n_{P0}e^{\frac{qV_{a}}{kT}}$$

$$= n_{P0} + \hat{n}(x = 0)$$

$$p_{N} = p_{N0}e^{\frac{qV_{a}}{kT}}$$

$$= p_{N0} + \hat{p}(x = 0)$$

Also,

$$\hat{n}(x=0) = n_{P0} \left(e^{\frac{qV_a}{kT}} - 1 \right)$$

$$\hat{p}(x=0) = p_{N_0} \left(e^{\frac{qV_a}{kT}} - 1 \right)$$

Therefore,

$$\hat{n}(x) = n_{P0} \left(e^{\frac{qV_a}{kT}} - 1 \right) e^{-\frac{x}{L_n}}$$

$$\hat{p}(x) = p_{N0} \left(e^{\frac{qV_a}{kT}} - 1 \right) e^{-\frac{x}{L_p}}$$

Therefore,

$$J_{\text{diffusion}_n} = qD_n \frac{\mathrm{d}n}{\mathrm{d}x}$$

$$= qD_n \frac{\mathrm{d}\hat{n}}{\mathrm{d}x}$$

$$= -q \frac{D_n}{L_n} n_{P0} \left(e^{\frac{qV_a}{kT}} - 1 \right) e^{-\frac{x}{L_n}}$$

$$J_{\text{diffusion}_p} = qD_p \frac{\mathrm{d}p}{\mathrm{d}x}$$

$$= qD_p \frac{\mathrm{d}\hat{p}}{\mathrm{d}x}$$

$$= -q \frac{D_p}{L_p} p_{N0} \left(e^{\frac{qV_a}{kT}} - 1 \right) e^{-\frac{x}{L_p}}$$

The continuity equation for electrons in the depletion region is

$$\frac{\partial \hat{n}}{\partial t} = \frac{1}{q} \frac{\partial J_n}{\partial x} - \frac{\hat{n}}{\tau_n} + G$$

Assuming that no generation or recombination of EHPs occurs in the depletion region,

$$G = 0$$
$$\tau_n = 0$$

As the junction is in steady state,

$$\frac{\partial \hat{n}}{\partial t} = 0$$

Therefore,

$$\frac{1}{q} \frac{\partial J_n}{\partial x} = 0$$

Similarly,

$$\frac{1}{q}\frac{\partial J_p}{\partial x} = 0$$

Therefore as J_n and J_p are constant in the depletion region,

$$J_n(-x_p < x < x_n) = J_n(x = -x_p)$$

 $J_p(-x_p < x < x_n) = J_p(x = x_n)$

Therefore,

$$J_{\text{total}} = J_{\text{diffusion}_n}(x=0) + J_{\text{diffusion}_p}(x=0)$$
$$= q \left(\frac{D_n}{L_n} n_{P0} + \frac{D_p}{L_p} p_{N0}\right) \left(e^{\frac{qV_a}{kT}} - 1\right)$$

where each x = 0 corresponds to the respective x-axis. Let

$$J_0 = q \left(\frac{D_n}{L_n} n_{P0} + \frac{D_p}{L_p} p_{N0} \right)$$

Therefore,

$$J_{\text{total}} = J_0 \left(e^{\frac{qV_a}{kT}} - 1 \right)$$

As the total current in the junction must be constant, the majority and minority currents in each of the regions must add up to J_{total} .

Exercise 8.

Consider a silicon sample with

$$N_D = 10^{16} \frac{1}{\text{cm}^3}$$

$$\tau_n = 1 \text{µs}$$

$$\tau_p = 1 \text{µs}$$

$$D_n = 40 \frac{\text{cm}^2}{\text{s}}$$

$$D_p = 25 \frac{\text{cm}^2}{\text{s}}$$

is maintained at steady state at 300 K. The hole concentration at the left end is $10^{14} \frac{1}{\text{cm}^3}$. What is p(x) at steady state?

Solution 8.

Let the left end of the sample correspond to x = 0. Therefore,

$$\begin{split} \frac{\partial \hat{p}}{\partial t} &= -\frac{1}{q} \frac{\partial J_p}{\partial x} - \frac{\hat{p}}{\tau_p} + G_{\text{optical}} \\ J_p &= q \mu_p p \mathcal{E} - q D_p \frac{\mathrm{d}p}{\mathrm{d}x} \end{split}$$

As $\mathcal{E} = 0$,

$$J_p = -qD_p \frac{\mathrm{d}p}{\mathrm{d}x}$$
$$= -qD_p \frac{\mathrm{d}\hat{p}}{\mathrm{d}x}$$

As the sample is in steady state,

$$\frac{\partial \hat{p}}{\partial t} = 0$$

As the sample is illuminated at the end only,

$$G_{\text{optical}} = 0$$

Therefore, substituting,

$$0 = D_p \frac{\mathrm{d}^2 \hat{p}}{\mathrm{d}x^2} - \frac{\hat{p}}{\tau_p}$$

$$\therefore \frac{\mathrm{d}^2 \hat{p}}{\mathrm{d}x^2} = \frac{1}{D_p \tau_p}$$

$$\therefore \frac{\mathrm{d}^2 \hat{p}}{\mathrm{d}x^2} = \frac{1}{L_p^2}$$

Therefore,

$$\hat{p}(x) = c_1 e^{\frac{x}{L_p}} + c_2 e^{-\frac{x}{L_p}}$$

Substituting the initial conditions,

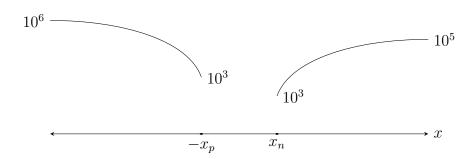
$$c_1 = 0$$
$$c_2 = 10^{14}$$

Therefore,

$$\hat{p}(x) = 10^{14} e^{-\frac{x}{L_p}}$$

Exercise 9.

Consider a silicon PN step junction at $300\,\mathrm{K}$. The steady state distribution is as shown.



- 1. Is the junction in forward bias or reverse bias?
- 2. Find V_a .

Solution 9.

1.

$$n_P = n_{P0}e^{\frac{qV_a}{kT}}$$
$$\therefore 10^3 = 10^6 e^{\frac{qV_a}{kT}}$$

Therefore,

$$e^{\frac{qV_a}{kT}} < 1$$
$$\therefore V_a < 0$$

Therefore, the junction is in reverse bias.

2.

$$e_{\frac{qV_a}{kT}} = \frac{n_P}{n_{P0}}$$
$$= \frac{p_N}{p_{N0}}$$
$$= 10^{-3}$$

Therefore,

$$V_a = \frac{kT}{q} \ln \left(10^{-3}\right)$$
$$= -0.18 \,\mathrm{V}$$

Exercise 10.

Consider a silicon PN step junction at 300 K such that

$$N_A = 3 \times 10^{16} \frac{1}{\text{cm}^3}$$

$$N_D = 2 \times 10^{15} \frac{1}{\text{cm}^3}$$

$$D_n = 50 \frac{\text{cm}^2}{\text{s}}$$

$$D_n = 10 \frac{\text{cm}^2}{\text{s}}$$

$$\tau_n = \tau_p$$

Given that

$$n_P = 10^{15}$$

 $n_{P0}(3\mu\text{m}) = 3.7 \times 10^{14}$

- 1. Find the minority current density at x > 0.
- 2. Find the applied voltage.
- 3. Find J_{total} for $V_a = 0.6 \,\text{V}$.

Solution 10.

1.

$$\hat{n}(x) = \hat{n}(x = 0)e^{-\frac{x}{L_n}}$$

$$\therefore L_n = 3\mu m$$

Therefore,

$$J_{\text{diffusion}_n}(x=0) = -q \frac{D_n}{L_n} n_{P_0} \left(e^{\frac{qV_a}{kT}} - 1 \right)$$
$$= -\left(1.6 \times 10^{-19} \right) \left(\frac{50}{3 \times 10^{-4}} \right) \left(10^{15} \right)$$
$$= -26.67 \frac{A}{\text{cm}^2}$$

2.

$$\hat{n}(x=0) = n_{P0} \left(e^{\frac{qV_a}{kT}} - 1 \right)$$

As the minority carrier concentration at the interface of the depletion region is greater than that inside the region, the function is in forward bias. Therefore,

$$\hat{n}(x=0) = n_{P0}e^{\frac{qV_a}{kT}}$$

Therefore,

$$V_a = \frac{kT}{q} \ln \left(\frac{\hat{n}(x=0)}{n_{P_0}} \right)$$

$$= 0.026 \ln \left(\frac{N_A \hat{n}(x=0)}{n_i^2} \right)$$

$$= 0.026 \ln \left(\frac{3 \times 10^{16} 10^{15}}{10^{20}} \right)$$

$$= 0.687 \text{ V}$$

3.

$$J_{\text{total}} = q \left(\frac{D_p}{L_p} p_{N_0} + \frac{D_n}{L_n} n_{P_0} \right) \left(e^{\frac{qV_a}{kT}} - 1 \right)$$

As the junction is in forward bias,

$$J_{\text{total}} = q \left(\frac{D_p}{L_p} p_{N_0} + \frac{D_n}{L_n} n_{P_0} \right) \left(e^{\frac{qV_a}{kT}} \right)$$

Also,

$$L_p = \sqrt{\tau_p D_p}$$

$$= \sqrt{\tau_p D_p} \frac{\sqrt{D_p}}{\sqrt{D_p}}$$

$$= \sqrt{\tau_p D_n} \sqrt{\frac{D_p}{D_n}}$$

$$= \sqrt{\tau_n D_n} \sqrt{\frac{D_p}{D_n}}$$

$$= L_n \sqrt{\frac{D_p}{D_n}}$$

Therefore, as $L_n = 3\mu m$,

$$L_p = 3\sqrt{\frac{10}{50}}$$

= 1.34 × 10⁻⁴cm

Therefore, solving,

$$J_{\text{total}} = \left(\left(\frac{10}{1.34 \times 10^{-4}} \right) \left(\frac{10^{20}}{2 \times 10^{15}} \right) + \left(\frac{50}{3 \times 10^{-4}} \right) \left(\frac{10^{20}}{3 \times 10^{16}} \right) \right) e^{\frac{0.6}{0.026}}$$
$$= 7.12 \frac{A}{\text{cm}^2}$$

Exercise 11.

Consider a silicon PN step junction such that

$$au_n = 0.1 \mu s$$
 $au_p = 0.1 \mu s$

$$au_p = 0.1 \mu s$$

$$au_n = 25 \frac{\text{cm}^2}{\text{s}}$$

$$au_p = 10 \frac{\text{cm}^2}{\text{s}}$$

Find the ratio of the doping such that 95% of the current in the depletion region is due to electrons.

Solution 11.

$$J_{\text{total}} = q \left(\frac{D_p}{L_p} p_{N_0} + \frac{D_n}{L_n} n_{P_0} \right) \left(e^{\frac{qV_a}{kT}} - 1 \right)$$

Therefore, the contribution of the electrons corresponds to $\frac{D_n}{L_n}n_{P0}$ and that of the holes corresponds to $\frac{D_p}{L_p}p_{N0}$. Therefore, for the required ratio,

$$0.95 = \frac{\frac{D_n}{L_n} n_{P0}}{\frac{D_n}{L_n} n_{P0} \frac{D_p}{L_p} p_{N0}}$$

$$\therefore \frac{D_n}{L_n} n_{P0} = 19 \frac{D_p}{L_p} p_{N0}$$

$$\therefore \frac{D_n}{L_n} n_i^2 N_A = 19 \frac{D_p}{L_p} \frac{n_i^2}{N_D}$$

$$\therefore \frac{N_A}{N_D} = \frac{D_n}{L_n} \frac{L_p}{D_p} \frac{1}{19}$$

$$= \frac{D_n}{\sqrt{D_n \tau_n}} \frac{\sqrt{D_p \tau_p}}{D_p} \frac{1}{19}$$

$$= 0.083$$