Chapter 1: Electrons and Holes in Semiconductors

- □ Course slides are prepared with the aid of the following materials:
	- \Box The lecture slides accompanying the main textbook "Hu, Chenming. *Modern semiconductor devices for integrated circuits*. Prentice Hall, 2010."
	- [http://www.eecs.berkeley.edu/~hu/Book-Chapters-and-](http://www.eecs.berkeley.edu/~hu/Book-Chapters-and-Lecture-Slides-download.html)Lecture-Slides-download.html
	- **The lecture slides accompanying of the Semiconductor** Device Physics course offered by Dr.-Ing. Erwin Sitompul, President University, Indonesia.
	- [http://zitompul.wordpress.com/1-ee-lectures/2](http://zitompul.wordpress.com/1-ee-lectures/2-semiconductor-device-physics/) semiconductor-device-physics/

Chapter Objectives

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- \Box Provides the basic concepts and terminology for understanding semiconductors.
- □ Understand conduction and valence energy band, and how bandgap is formed.
- □ Understand carriers (electrons and holes), and doping in semiconductor
- □ Use the density of states and Fermi-Dirac statistics to calculate the carrier concentration

What is a Semiconductor?

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 \Box The conductivity (and at the same time the resistivity) of semiconductors lie between that of conductors and insulators.

What is a Semiconductor?

□ Semiconductors are some of the purest solid materials in existence, because any trace of impurity atoms called "dopants" can change the electrical properties of semiconductors drastically.

Completely ordered in segments

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No recognizable long-range order

Entire solid is made up of atoms in an orderly three- dimensional array

polycrystalline amorphous crystalline

□ Most devices fabricated today employ crystalline semiconductors.

Crystal Growth Until Device Fabrication

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Semiconductor Materials

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The Silicon Atom

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- ■14 electrons occupying the first 3 energy levels:
	- 1s, 2s, 2p orbitals are filled by 10 electrons.
	- 3s, 3p orbitals filled by 4 electrons.
- To minimize the overall energy, the 3s and 3p orbitals hybridize to form four tetrahedral 3sp orbital.
- **Each has one electron and is capable of** forming a bond with a neighboring atom.

1.1 Silicon Crystal Structure

- *Unit cell* of silicon crystal is cubic.
- *Each Si atom has 4 nearest neighbors***.**
- **Each cell contains:** 8 corner atoms 6 face atoms 4 interior atoms

5.43 Å

Exercise

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How Many Silicon Atoms per cm–³ ?

Silicon Wafers and Crystal Planes (Miller Indices)

 The standard notation for crystal planes is based on the cubic unit cell.

Silicon wafers are usually cut along the (100) plane with a flat or notch to help orient the wafer during IC fabrication.

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Crystallographic Planes of Si

Unit cell:

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View in <111> direction

View in <100> direction

View in <110> direction

1.2 Bond Model of Electrons and Holes

• When an electron breaks loose and becomes a *conduction electron*, a *hole* is also created.

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Doping - Manipulation of Carrier Numbers

By substituting an Si atom with a special impurity atom (elements from **Group III** or **Group V**), a hole or conduction electron can be created.

Acceptors: B, Ga, In, Al

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Donors: P, As, Sb

Boron, Gallium Indium, Aluminum

Phosphorus, Arsenic, Antimony

Doping Silicon with Acceptors

Example: Aluminum atom is doped into the Si crystal.

The Al atom accepts an electron from a neighboring **Si** atom, resulting in a missing bonding electron, or "hole".

The hole is free to roam around the Si lattice, and as a moving positive charge, the hole carries current.

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Doping Silicon with Donors

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Example: Phosphorus atom is doped into the Si crystal.

The loosely bounded fifth valence electron of the **P** atom can "break free" easily and becomes a mobile conducting electron.

This electron contributes in current conduction.

Dopants in Silicon

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- As, a Group V element, introduces conduction electrons and creates *N-type silicon,* and is called a *donor.*
- B, a Group III element, introduces holes and creates *P-type silicon*, and is called an *acceptor*.
- Donors and acceptors are known as dopants. Dopant ionization energy ~50meV (very low).

Hydrogen:
$$
E_{ion} = \frac{m_o q^4}{8e_o^2 h^2} = 13.6 \text{ eV}
$$

Carrier-Related Terminology

- **Donor**: impurity atom that increases *n* (conducting electron). **Acceptor**: impurity atom that increases *p* (hole).
- *n***-type material**: contains more electrons than holes. *p***-type material**: contains more holes than electrons.
- **Majority carrier**: the most abundant carrier. **Minority carrier**: the least abundant carrier.
- **Intrinsic semiconductor**: undoped semiconductor *n* = $p = n_i$. **Extrinsic semiconductor**: doped semiconductor.

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1.3 Energy Band Model $2p$ (conduction band) **Empty upper bands**

- · Energy states of Si atom (a) expand into energy bands of Si crystal (b).
- The lower bands are filled and higher bands are empty in a semiconductor.
- · The highest filled band is the *valence band.*
- · The lowest empty band is the *conduction band* .

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1.3.1 Energy Band Diagram

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- · *Energy band diagram* shows the bottom edge of conduction band, *E^c* , and top edge of valence band, *E^v* .
- · *E^c* and *E^v* are separated by the *band gap energy***,** *E^g* .

Measuring the Band Gap Energy by Light Absorption

• *E^g* can be determined from the minimum energy (*hn*) of photons that are absorbed by the semiconductor.

Bandgap energies of selected semiconductors

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Donor / Acceptor Levels (Band Model)

Ionization energy of selected donors and acceptors in Silicon (E_G = 1.12 eV)

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1.4 Semiconductors, Insulators, and Conductors

- · Totally filled bands and totally empty bands do not allow current flow. (Just as there is no motion of liquid in a totally filled or totally empty bottle.)
- · Metal conduction band is half-filled.
- · Semiconductors have lower *E g* 's than insulators and can be doped.

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1.5 Electrons and Holes

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- · Both electrons and holes tend to seek their lowest energy positions.
- · Electrons tend to fall in the energy band diagram.
- · Holes float up like bubbles in water **.**

1.5.1 Effective Mass

The electron wave function is the solution of the three dimensional Schrodinger wave equation

$$
-\frac{\hbar^2}{2m_0}\nabla^2\psi + V(r)\,\psi = \psi
$$

The solution is of the form $exp(\pm {\bf k} \cdot {\bf r})$ k = wave vector = 2π /electron wavelength For each k, there is a corresponding E.

acceleration
$$
= -\frac{q\varepsilon}{\hbar^2} \frac{d^2E}{dk^2} = \frac{F}{m}
$$

effective mass $\equiv \frac{\hbar^2}{d^2E/dk^2}$

1.5.1 Effective Mass

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In an electric field, ε, an electron or a hole accelerates.

$$
a = \frac{-q \ell}{m_n}
$$
 electrons

$$
a = \frac{q \ell}{m_p}
$$
 holes

Electron and hole effective masses

1.5.2 How to Measure the Effective Mass

Centripetal force = Lorentzian force

$$
\frac{m_n v^2}{r} = qvB
$$

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B

- *•f_{cr}* is the Cyclotron resonance frequency.
- •It is independent of *v* and *r*.
- •Electrons strongly absorb microwaves of that frequency.
- •By measuring f_{cr} , m_n can be found.

1.6 Density of States

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1.7 Thermal Equilibrium and the Fermi Function

1.7.1 An Analogy for Thermal Equilibrium

• There is a certain probability for the electrons in the conduction band to occupy high-energy states under the agitation of thermal energy.

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1.7.2 Fermi Function–The Probability of an Energy State Being Occupied by an Electron

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

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Ef is called the *Fermi energy* or the *Fermi level.*

Boltzmann approximation:

$$
E_{f} + 3kT
$$
\n
$$
E_{f+2kT}
$$
\n
$$
E_{f} + kT
$$
\n
$$
E_{f} - kT
$$
\n
$$
E_{f} - 3kT
$$
\n
$$
0.5 \t 1
$$
\n
$$
f(E)
$$

$$
\left| f(E) \approx e^{-(E-E_f)/kT} \right| \quad E-E_f >> kT
$$

$$
f\left(E\right)\thickapprox1\!-\!e^{-\left(E_{_{f}}-E\right)/kT}\quad E\!-\!E_{_{f}}<<\!-kT
$$

Remember: there is only one Fermi-level in a system at equilibrium.

Effect of Temperature on *f(E)*

No electrons can be above the valence band at 0K, since none have energy above the Fermi level and there are no available energy states in the band gap.

At high temperatures, some electrons can reach the conduction band and contribute to electric current.

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Effect of Temperature on *f***(***E***)**

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Equilibrium Distribution of Carriers - Intrinsic

 \blacksquare *n*(*E*) is obtained by multiplying $g_c(E)$ and $f(E)$, $p(E)$ is obtained by multiplying $g_v(E)$ and $1-f(E)$.

Intrinsic semiconductor material

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Equilibrium Distribution of Carriers – n-type

*n***-type semiconductor material**

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Equilibrium Distribution of Carriers – p-type

*p***-type semiconductor material**

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Nondegenerately Doped Semiconductor

The expressions for n and p will now be derived in the range where the Boltzmann approximation can be applied:

The semiconductor is said to be **nondegenerately doped (lightly doped)** in this case.

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Degenerately Doped Semiconductor

 \blacksquare If a semiconductor is very heavily doped, the Boltzmann approximation is not valid.

■ For Si at
$$
T = 300
$$
 K,
\n $E_c - E_F < 3kT$ if N_D > 1.6 × 10¹⁸ cm⁻³
\n $E_F - E_v < 3kT$ if N_A > 9.1 × 10¹⁷ cm⁻³

The semiconductor is said to be degenerately doped (heavily doped) in this case.

• N_{D} = total number of donor atoms/cm³

• N_A = total number of acceptor atoms/cm³

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Important Constants

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Electronic charge, $q = 1.6 \times 10^{-19}$ C **P** Permittivity of free space, $\varepsilon_0 = 8.854 \times 10^{-12}$ F/m Boltzmann constant, $k = 8.62 \times 10^{-5}$ eV/K **Planck constant,** $h = 4.14 \times 10^{-15}$ eV·s, $\hbar = h/2\pi$ **Figure** Free electron mass, $m_0 = 9.1 \times 10^{-31}$ kg Thermal energy, $kT = 0.02586$ eV (at 300 K) Thermal voltage, $kT/q = 0.02586$ V (at 300 K) Silicon energy band gap, $E_G = 1.12eV$ ■ Intrinsic Si carrier concentration $n_i = 1 \times 10^{10}$ cm⁻³ (at 300 K)

1.8 Electron and Hole Concentrations

1.8.1 Derivation of *n* **and** *p* **from** $D(E)$ **and** $f(E)$

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Electron and Hole Concentrations

$$
n = N_c e^{-(E_c - E_f)/kT}
$$

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$$
N_c \equiv 2 \left[\frac{2\pi m_n kT}{h^2} \right]^{3/2}
$$

$$
p = N_{v}e^{-(E_f - E_v)/kT}
$$

$$
N_v \equiv 2 \left[\frac{2\pi m_p kT}{h^2} \right]^{3/2}
$$

Nc is called the *effective density of states (of the conduction band) .*

Nv is called the *effective density of states of the valence band.*

Remember: the closer E_f **moves up to** N_c **, the larger** *n* **is;** the closer E_f moves down to \mathbf{E}_{ν} , the larger p is. **For Si,** $N_c = 2.8$ $\hat{10}^{19}$ cm⁻³ and $N_v = 1.04$ $\hat{10}^{19}$ cm⁻³.

1.8.2 The Fermi Level and Carrier Concentrations

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1.8.2 The Fermi Level and Carrier Concentrations

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1.8.3 The *np* **Product and the Intrinsic Carrier Concentration**

Multiply
$$
n = N_c e^{-(E_c - E_f)/kT}
$$
 and $p = N_v e^{-(E_f - E_v)/kT}$

$$
np = N_c N_v e^{-(E_c - E_v)/kT} = N_c N_v e^{-E_g/kT}
$$

$$
np = n_i^2
$$

$$
n_i = \sqrt{N_c N_v} e^{-E_g/2kT}
$$

- In an intrinsic (undoped) semiconductor, $n = p = n_i$.
- n_i is the *intrinsic carrier concentration*, $\sim 10^{10}$ cm⁻³ for Si.

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Alternative Expressions: $n(n_i, E_i)$ and $p(n_i, E_i)$

In an intrinsic semiconductor, $n = p = n_i$ and $E_F = E_i$, where E_i denotes the intrinsic Fermi level.

$$
n = N_{\rm c}e^{(E_{\rm F}-E_{\rm c})/kT}
$$
\n
$$
p = N_{\rm v}e^{(E_{\rm v}-E_{\rm c})/kT}
$$
\n
$$
\Rightarrow N_{\rm c} = n_{\rm i}e^{-(E_{\rm i}-E_{\rm c})/kT} \Rightarrow N_{\rm v} = n_{\rm i}e^{-(E_{\rm v}-E_{\rm c})/kT}
$$
\n
$$
n = n_{\rm i}e^{-(E_{\rm i}-E_{\rm c})/kT} \cdot e^{(E_{\rm F}-E_{\rm c})/kT}
$$
\n
$$
p = n_{\rm i}e^{-(E_{\rm v}-E_{\rm i})/kT}
$$
\n
$$
p = n_{\rm i}e^{-(E_{\rm v}-E_{\rm i})/kT}
$$
\n
$$
p = n_{\rm i}e^{(E_{\rm c}-E_{\rm i})/kT}
$$
\n
$$
E_{\rm F} = E_{\rm i} + kT \ln \left(\frac{n}{n_{\rm i}}\right)
$$
\n
$$
E_{\rm F} = E_{\rm i} - kT
$$

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$$
p = N_{\rm V} e^{(E_{\rm v}-E_{\rm F})/kT}
$$

$$
p_{i} = N_{v}e^{(E_{v}-E_{i})/kT}
$$

\n
$$
\Rightarrow N_{v} = n_{i}e^{-(E_{v}-E_{i})/kT}
$$

$$
p = n_{\mathrm{i}}e^{-(E_{\mathrm{v}}-E_{\mathrm{i}})/kT} \cdot e^{(E_{\mathrm{v}}-E_{\mathrm{F}})/kT}
$$

$$
p = n_{\mathrm{i}}e^{(E_{\mathrm{i}}-E_{\mathrm{F}})/kT}
$$

$$
E_{\rm F} = E_{\rm i} - kT \ln \left(\frac{p}{n_{\rm i}} \right)
$$

Intrinsic Fermi Level, E.

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EXAMPLE: Carrier Concentrations

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Question: What is the hole concentration in an N-type semiconductor with 10¹⁵cm-3 of donors?

Solution: n = 10¹⁵ cm-3 .

$$
p = \frac{n_i^2}{n} \approx \frac{10^{20} \text{cm}^{-3}}{10^{15} \text{cm}^{-3}} = 10^5 \text{cm}^{-3}
$$

After increasing T by 60C, n remains the same at 10¹⁵cm-3 while p increases by about a factor of 2300 because $n_i^{\,2} \propto e^{-E_g/kT}$ $n_i^{\;\;\prime} \varpropto e^{-\frac{1}{2} \varpropto \varrho}$

Question: What is n if p = 10¹⁷cm-3 in a P-type silicon wafer?

Solution:
\n
$$
n = \frac{n_i^2}{p} \approx \frac{10^{20} \text{cm}^{-3}}{10^{17} \text{cm}^{-3}} = 10^3 \text{cm}^{-3}
$$

Example: Energy-Band Diagram

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For Silicon at 300 K, where is E_F **if** $n = 10^{17}$ **cm⁻³?**

Silicon at 300 K,
$$
n_i = 10^{10} \text{ cm}^{-3}
$$

\n
$$
E_F = E_i + kT \ln \left(\frac{n}{n_i} \right)
$$
\n
$$
= 0.56 + 8.62 \cdot 10^{-5} \cdot 300 \cdot \ln \left(\frac{10^{17}}{10^{10}} \right) \text{ eV}
$$
\n
$$
= 0.56 + 0.417 \text{ eV}
$$
\n
$$
= 0.977 \text{ eV}
$$

1.9 General Theory of n and p

EXAMPLE: Complete ionization of the dopant atoms

N^d = 10¹⁷ cm-3 . What fraction of the donors are not ionized?

Solution: First assume that all the donors are ionized.

$$
n = N_d = 10^{17} \text{cm}^{-3} \Rightarrow E_f = E_c - 146 \text{ meV}
$$

\n
$$
\frac{1}{10^{-7} \text{cm}^{-3}} = \frac{1}{1 + \frac{1}{2} e^{(E_d - E_f)/kT}} = \frac{1}{1 + \frac{1}{2} e^{((146 - 45) \text{meV})/26 \text{meV}}} = 0.04
$$

\n
$$
\frac{1}{1 + \frac{1}{2} e^{(E_d - E_f)/kT}} = \frac{1}{1 + \frac{1}{2} e^{((146 - 45) \text{meV})/26 \text{meV}}} = 0.04
$$

\nTherefore, it is reasonable to assume complete ionization, i.e., $n = N_d$.

1.9 General Theory of n and p (2)

$$
Change \text{ neutrality: } n + N_a = p + N_d
$$

$$
np = n_i^2
$$

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1.9 General Theory of on n and p (3)

I.
$$
N_d - N_a \gg n_i
$$
 (i.e., N-type) $n = N_d - N_a$
 $p = n_i^2/n$

$$
\text{If } N_d >> N_a \quad, \quad n = N_d \quad \text{and} \quad p = n_i^2 / N_d
$$

II.
$$
N_a - N_d
$$
 $> n_i$ (i.e., P-type) $p = N_a - N_d$
\n $n = n_i^2 / p$

If
$$
N_a \gg N_d
$$
, $p = N_a$ and $n = n_i^2/N_a$

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EXAMPLE: Dopant Compensation

What are n and p in Si with (a) $N_d = 6 \times 10^{16}$ *cm⁻³ and* $N_a = 1$ *210¹⁶ cm-3 and (b) additional 610¹⁶ cm-3 of N^a ?* $n - 4 \times 10^{16}$ cm⁻³

(a)
$$
n = N_d - N_a = 4 \times 10^{16} \text{ cm}^{-3}
$$

\n $p = n_i^2 / n = 10^{20} / 4 \times 10^{16} = 2.5 \times 10^3 \text{ cm}^{-3}$
\n(b) $N_d = 2 \times 10^{16} + 6 \times 10^{16} = 8 \times 10^{16} \text{ cm}^{-3} > N_d$
\n $p = N_a - N_d = 8 \times 10^{16} - 6 \times 10^{16} = 2 \times 10^{16} \text{ cm}^{-3}$
\n $n = n_i^2 / p = 10^{20} / 2 \times 10^{16} = 5 \times 10^3 \text{ cm}^{-3}$
\n $N_d = 6 \times 10^{16} \text{ cm}^{-3}$
\n $N_d = 6 \times 10^{16} \text{ cm}^{-3}$
\n $N_d = 6 \times 10^{16} \text{ cm}^{-3}$

 $p = 2 \times 10^{16}$ cm⁻³

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Carrier Concentration *vs.* **Temperature**

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1.11 Chapter Summary

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Energy band diagram. Acceptor. Donor. m_n , m_p . Fermi function. *E^f* .

$$
n = N_c e^{-(E_c - E_f)/kT}
$$

$$
p = N_v e^{-(E_f - E_v)/kT}
$$

$$
n = N_d - N_a
$$

$$
p = N_a - N_d
$$

 $np = n_i$