"Complementi di Fisica" Lectures 25-26

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Trieste, 14/15-12-2015

in these lectures

Introduction

- Non or quasi-equilibrium: "excess" carriers injection
- Processes for "generation" and "recombination" of carriers
- Continuity equations for carriers
- Continuity equations: three important special cases
 - Steady-state injection from one side
 - "diffusion length" L_p
 - Minority carriers recombination at the surface
 - diffusion length and "surface recombination velocity" S_{lr}
 - The Haynes-Shockley experiment
 - Evidence for simultaneous diffusion, drift and recombination
- Are we describing the behaviour of *minority* carriers alone? What about *majority* carriers?
 - Why are "minorities" important? Some examples...
 - Built-in electric field (Gauss!) and "ambipolar" transport equations

In these lectures

Reference textbooks

- D.A. Neamen, Semiconductor Physics and Devices, McGraw-Hill, 3rd ed., 2003, p.189-230 ("6 Nonequilibrium excess carriers in semiconductors")
- R.Pierret, Advanced Semiconductor Fundamental, Prentice Hall, 2nd ed., p.134-174 ("5 Recombination-Generation Processes")
- J.Nelson, The Physics of Solar Cells, Imperial College Press, p. 79-117 (Ch.4, "Generation and Recombination")

Injection of "excess" carriers

Non-equilibrium!

(in some cases: quasi-equilibrium)

Carrier injection - introduction

- "carrier injection" = process of introducing "excess" carriers in a semiconductor, so that: $np > n_i^2$
 - Optical excitation:
 - shine a light on a semiconductor crystal;
 - if the energy of the photons is $h_V > E_g$, then
 - Photons absorbed
 - "excess" electron-hole *pairs* are created: $\Delta n = \Delta p$
 - Other methods:
 - Forward-bias a pn junction
 - ...
 - In an extrinsic semiconductor, the relative effect of ∆n = ∆p is very different for "majority" and "minority" carriers, since n ≠ p
 - Let us work out an example (n-type Si, $n_0 > p_0$ at equilibrium)

Carrier injection



Carrier injection



Carrier injection



Carrier injection - summary

- "carrier injection" = process of introducing "excess" carriers in a semiconductor
- Several methods (optical, etc.)
- Low-level injection: relative effect on concentration
 - Negligible on majority carriers
 - Important for minority carriers (also called "minority carriers injection")
- High-level injection
 - If very high, both concentrations become comparable
 - Sometimes encountered in device operation

Quasi-equilibrium



Example: "Injection" or "Generation" of an excess of electrons and holes by absorption of photons in a very short time, *about 10*⁻¹⁴ s

Excess electrons and holes relax separately to thermal equilibrium in about 10⁻¹² s and remain for a much longer time in this quasi-equilibrium state

Recombination of electrons and holes via several possible mechanisms takes typically *about* 10⁻⁶ s; plenty of time to do something useful with "stable" electrons and holes!

Quasi-equilibrium and quasi-Fermi levels



- In quasi-equilibrium conditions:
 - Two different "Quasi-Fermi Levels" F_N and F_P, describe the separate quasi-equilibrium concentrations of electrons and holes,
 - each population corresponding to a separate Fermi-Dirac pdf (one for electrons, another for holes)

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Quasi-Fermi levels: definition

- In quasi-equilibrium conditions:
 - Two different "Quasi-Fermi Levels" F_N and F_P, describe the separate quasi-equilibrium concentrations of electrons and holes:

• Electrons:

$$n \equiv n_i e^{(F_N - E_i)/kT} = N_C e^{-(E_C - F_N)/kT} \iff$$
$$F_N \equiv E_i + kT \ln(n/n_i) = E_C - kT \ln(N_C/n)$$

• Holes:

$$p \equiv n_i e^{(E_i - F_p)/kT} = N_V e^{-(F_N - E_V)/kT} \Leftrightarrow$$
$$F_P \equiv E_i - kT \ln(p/n_i) = E_V + kT \ln(N_V/p)$$

Modified mass action law

• Fermi level E_F at equilibrium:

$$n_0 = n_i e^{(E_F - E_i)/kT}$$
 $p_0 = n_i e^{(E_i - E_F)/kT}$
 $n_0 p_0 = n_i^2$

• Quasi-Fermi levels at quasi-equilibrium:

$$n \equiv n_i e^{(F_N - E_i)/kT} \qquad p \equiv n_i e^{(E_i - F_p)/kT}$$
$$np = n_i^2 e^{(F_N - F_p)/kT} > n_i^2$$

Difference in total chemical potentials or quasi-Fermi levels

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Quasi-Fermi levels: an example



Quasi-Fermi levels and currents

- At (quasi-)equilibrium, (quasi-)Fermi levels are constant
 - Why? Because from the thermodynamic point of view the Fermi level is the "total" chemical potential, including the "internal" chemical potential and "external" potential energy contributions, like for instance the electrostatic potential energy.
- Off-equilibrium, the net movement of carriers is related to the changing total chemical potential or (quasi-)Fermi level by:

$$J_{n,x} \equiv \mu_n n \frac{\partial F_N}{\partial x} \qquad \qquad J_{p,x} \equiv \mu_p p \frac{\partial F_P}{\partial x}$$

• From the definitions of F_N , F_P by substitution one obtains:

$$J_{n,x} = q\mu_n nE_x + q\mu_n \frac{k_B T}{q} \frac{\partial n}{\partial x} \qquad J_{p,x} = q\mu_p pE_x - q\mu_p \frac{k_B T}{q} \frac{\partial p}{\partial x}$$

drift diffusion D_n drift diffusion D_p

NB: A quasi-Fermi level that varies with position in a band diagram immediately indicates that current is flowing in the semiconductor! (see exercise 1 for an application)

Generation and Recombination

Charge carriers: electrons and holes

Electrons and holes

Generation rate G :

- G = number of free carriers generated (separating electrons from holes) per second and per unit volume
- G is usually a function of the available energy (temperature, etc.)

Recombination rate R:

- R = number of free carriers "disappearing" due to recombination per second and per unit volume
- *R* is usually proportional to the product of concentrations of "carriers" and "recombination centers" and to a "capture coefficient" defined as $c = v_{th}\sigma$, where v_{th} is the thermal velocity and σ is the recombination process "cross-section"

Net recombination rate: U = R - G

Generation processes

Generation processes





Photon absorption: ingredients



Figure 2.32 Simplified E:k plot for the calculation of dN/dE_p for direct transitions.

direct transitions



Figure 2.33 Simplified E:k plot for the calculation of α versus E_p for indirect transitions.

indirect transitions

Photon absorption coefficient α

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Figure 2.30 Attenuation factor (in nepers/cm) versus photon manium at two temperatures. [After W. C. Dash and R. Net 1151 (1955).]



gure 2.31 Attenuation factor (nepers/cm) versus photon energy for Ge (on linear iles). Note that for the indirect transition, α is proportional to $(E_p - E_g - E_{phon})^2$, ile for the direct transition α is proportional to $(E_p - E_{g0})^{1/2}$. Dash et al., loc. cit. previous figure.)

$$\xrightarrow{I(0)} x x + dx$$

$$\frac{dI}{dx} = -\alpha I(x) \implies I(x) = I(0)e^{-\alpha x}$$

Recombination processes

Recombination processes



Recombination via traps

- Recombination: often dominated by indirect processes through "recombination centers" or "traps" (direct recombination is negligible for Si)
- Example: in an n-type semicond., under low-injection conditions:
 - for the minority-carriers (holes !) excess-recombination, the bottleneck is "hole capture", that determines the hole "lifetime" τ_p
 - Once captured, the hole recombines quickly, since there are many electrons available



Recombination via traps: terminology

- "Capture", "emission":
 - From the point of view of the trap!
- In particular (figure, next slide):
 - (a) electron capture = (3)
 - (b) electron emission = (2)
 - (c) hole capture = (4)
 - (d) hole emission = (1)
- Detailed treatment: beyond our scope!
 - Shockley-Read-Hall model
 - See back-up slides and reference texts



Recombination via traps: "lifetime" approximation



Continuity equations

Overall conservation of charge!

Detailed accounting of local carrier density as a function of time: Generation, recombination, drift, diffusion

Summary of Classical Physics



(A) Conservation of charge: continuity equations

• Any net flow of charge must come from some supply!

$$\oint_{S} \vec{J} \cdot \hat{n} \, dS = \oint_{V} \vec{\nabla} \cdot \vec{J} \, dV = -\frac{d}{dt} \oint_{V} \rho \, dV = -\frac{dQ}{dt}$$
$$\vec{\nabla} \cdot \vec{J} = \frac{\partial J_{x}}{\partial x} + \frac{\partial J_{y}}{\partial y} + \frac{\partial J_{z}}{\partial z} = -\frac{\partial \rho}{\partial t}$$

- The flux of a current from a closed surface is equal to the decrease of the charge inside the surface
- $-~\rho$ is the net charge density (negative and positive, algebraic sum)
- Let us consider electrons and holes, separately, in a semiconductor, in a simple one-dimensional case

Continuity for electrons



Continuity for electrons and holes

One-dimensional

Three-dimensional

$$\frac{\partial n}{\partial t} = \frac{1}{|q|} \frac{\partial J_n}{\partial x} + (G_n - R_n)$$
$$\frac{\partial p}{\partial t} = -\frac{1}{|q|} \frac{\partial J_p}{\partial x} + (G_p - R_p)$$

$$\frac{\partial n}{\partial t} = \frac{1}{|q|} \vec{\nabla} \cdot \vec{J}_n + (G_n - R_n)$$
$$\frac{\partial p}{\partial t} = -\frac{1}{|q|} \vec{\nabla} \cdot \vec{J}_p + (G_p - R_p)$$

Continuity for electrons and holes

Minority carriers:

Electrons in p-type

holes in n-type

$$J_{n,x} = n_p |q| \mu_n E_x + D_n \frac{\partial n_p}{\partial x} \qquad J_{p,x} = p_n |q| \mu_p E_x - D_p \frac{\partial p_n}{\partial x}$$

One-dimensional, under low-injection conditions, for minority carriers:



Continuity: generation, recombination, drift, diffusion

Summary and real-life applications

Generation, Recombination, Continuity - 1



Generation, Recombination, Continuity - 2

minority carriers diffusion lengths


Device simulations

 In real life, device designers use programs performing numerical integrations in discrete space and time steps, to obtain (*):



"Equilibrium" vs "steady state"



"Equilibrium": detailed balance, for *each* process

"steady state": overall balance Continuity equations: applications

Three examples

(1) Steady state injection from one side(2) Recombination at the surface(3) Haynes-Shockley experiment

System of differential equations

(A) Continuity (transport) equations for minority carriers, 1-d case (Sze notations):

$$\frac{\partial n_p}{\partial t} = n_p \mu_n \frac{\partial E_x}{\partial x} + \mu_n E_x \frac{\partial n_p}{\partial x} + D_n \frac{\partial^2 n_p}{\partial x^2} + G_n - \frac{n_p - n_{p0}}{\tau_n}$$
$$\frac{\partial p_n}{\partial t} = -p_n \mu_p \frac{\partial E_x}{\partial x} - \mu_p E_x \frac{\partial p_n}{\partial x} + D_p \frac{\partial^2 p_n}{\partial x^2} + G_p - \frac{p_n - p_{n0}}{\tau_p}$$

(B) Gauss' law, relating the divergence of the electric field with the local charge density, 1-d case:

$$\frac{\partial E_x}{\partial x} = \frac{\rho}{\varepsilon}$$
Globally neutral, locally can be unbalanced!
$$\rho = |q| (p - n + N_D^+ - N_A^-) \approx |q| (p - n + N_D - N_A)$$

$$N = N_D - N_A$$

To be solved with given boundary conditions!

(ex.1) Steady-state injection from one side



(ex.1) Diffusion length - typical values

"Diffusion length"
$$L_p = \sqrt{D_p \tau_p}$$
 $L_n = \sqrt{D_n \tau_n}$

	μ _n [cm²/Vs]	D _n [cm ² /s]	μ_p [cm ² /Vs]	D _p [cm²/s]
Si	1350	35	480	12.4
GaAs	8500	220	400	10.4
Ge	3900	101	1900	49.2

example
$$L_p = \sqrt{D_p \tau_p} = \sqrt{(12.4)(5 \times 10^{-7})} = 25 \ \mu \text{m}$$

(ex.2) Recombination at the surface



(ex.2) Solution with boundary conditions

General solution:
$$\Delta p_n(x) = \begin{bmatrix} Ae^{x/L_p} + Be^{-x/L_p} \end{bmatrix} + \begin{bmatrix} G_L \tau_p \end{bmatrix}$$

"complementary" "particular"
(homegeneous)
 $L_p = \sqrt{D_p \tau_p}$

 \mathbf{v}

$$\Delta p_n(x) \xrightarrow[x \to +\infty]{} G_L \tau_p \implies A = 0$$

$$\Delta p_n(x) \xrightarrow[x \to 0]{} \Delta p_n(0) \implies \Delta p_n(0) = B + G_L \tau_p$$

$$B = \Delta p_n(0) - G_L \tau_p$$

after some algebra, substituting A and B, our solution:

$$p_{n}(x) = p_{n0} + G_{L}\tau_{p}\left(1 + \frac{\Delta p_{n}(0) - G_{L}\tau_{p}}{G_{L}\tau_{p}}e^{-x/L_{p}}\right)$$

$$S_{lr}$$

$$\Delta p_n(0) = ???$$



(ex.2) Solution with surface recomb. velocity

$$-J_{x}(0) = -(v_{th}\sigma_{p}N_{st})\Delta p_{n}(0) \implies D_{p}\left(\frac{d\Delta p_{n}}{dx}\right)_{x=0} = S_{lr}\Delta p_{n}(0)$$

cm² s⁻¹ cm⁻⁴ cm s⁻¹ cm⁻³

from the general solution and boundary conditions:

$$\left(\frac{d\Delta p_n}{dx}\right)_{x=0} = -\frac{B}{L_p} \qquad \Rightarrow \quad D_p \left(-\frac{B}{L_p}\right) = S_{lr} \left(G_L \tau_p + B\right)$$
$$\Delta p_n(0) = G_L \tau_p + B \qquad \Rightarrow \quad B = \frac{-S_{lr} G_L \tau_p}{D_p / L_p + S_{lr}}$$

Solution expressed in terms of the surface recombination velocity:

$$p_{n}(x) = p_{n0} + G_{L}\tau_{p} \left(1 - \frac{S_{lr}\tau_{p}}{L_{p} + S_{lr}\tau_{p}}e^{-x/L_{p}}\right)$$



(ex.2) Limiting cases

Neglecting surface recombination:

$$\begin{split} S_{lr} \tau_p &<< L_p \implies p_n(x) = p_{n0} + G_L \tau_p \\ p_n(0) &= p_{n0} + G_L \tau_p \\ \end{split} \quad \text{as expected!} \end{split}$$

Large ("immediate") surface recombination:

$$\begin{split} S_{lr}\tau_p >> L_p & \Rightarrow \quad p_n(x) = p_{n0} + G_L \tau_p \Big(1 - e^{-x/L_p} \Big) \\ p_n(0) = p_{n0} & \text{as expected!} \end{split}$$

(ex.3) The Haynes-Shockley experiment



Experimental set-up

excess carrier distributions at successive times t_1 and t_2 , no applied field

excess carrier distributions at successive times t_1 and t_2 , with a constant applied field

(ex.3) The Haynes-Shockley experiment

After a light pulse: $G_L = 0$ no bulk generation $\frac{\partial E_x}{\partial x} = 0$ constant applied field

Transport equation for excess minority carriers (n-type semiconductor):

$$\frac{\partial \Delta p_n}{\partial t} = \mu_p E_x \frac{\partial \Delta p_n}{\partial x} + D_p \frac{\partial^2 \Delta p_n}{\partial x^2} - \frac{\Delta p_n}{\tau_p} \qquad \Delta p_n = p_n - p_{n0}$$

Solution, no applied field:

$$\Delta p_n(x,t) = \frac{N}{\sqrt{4\pi D_p t}} \exp\left(-\frac{x^2}{4D_p t} - \frac{t}{\tau_p}\right) \qquad \qquad \text{diffusion,} \\ \text{recombination}$$

Solution, with applied field:

$$\Delta p_n(x,t) = \frac{N}{\sqrt{4\pi D_p t}} \exp \left(-\frac{\left(x - u_p E_x t\right)^2}{4D_p t} - \frac{t}{\tau_p}\right) \quad \begin{array}{c} \text{drift,} \\ \text{diffusion,} \\ \text{recombination} \end{array}\right)$$

The role of Gauss' law

Dielectric relaxation Ambipolar transport

The role of Gauss' law



Dielectric relaxation time constant



Dielectric relaxation: Debye length

"Debye length" L_D (~ 10⁻⁵ cm):

$$L_D \approx \sqrt{D_p \left(\frac{\varepsilon}{\sigma}\right)} = \sqrt{D_p \tau_d} \qquad \tau_d \equiv \varepsilon/\sigma \qquad \text{(~10^{-12} s)}$$

Expect no significant departures from electrical neutrality, over distances greater than about 4 L_D to 5 L_D in *uniformly doped extrinsic* material, at thermal equilibrium (also true off-equilibrium!); this process is much faster than the typical excess carrier lifetime (10⁻⁷ s)

Numerical example for n-type Si, doped with donor concentration $N_D = 10^{16}$ cm⁻³

$$\tau_{d} = \frac{\varepsilon}{\sigma} \approx \frac{\varepsilon_{r} \varepsilon_{0}}{q_{e} \mu_{n} N_{D}} = \frac{(11.7)(8.85 \times 10^{-14})}{(1.6 \times 10^{-19})(1200)(10^{16})} \frac{\text{F} \cdot \text{cm}}{(\Omega \cdot \text{cm})^{-1}}$$
$$= 5.4 \times 10^{-13} \text{ s} = 0.54 \text{ ps}$$

"ambipolar" transport

Two examples: (1) Bipolar diffusion (2) Shockley experiment

"Ambipolar transport" - equations

Combining the *transport equations* for electrons and holes with *Gauss' law*, under some simplifying assumptions, (see back-up slides and reference texts):

$$p' \equiv p - p_0$$
 $n' \equiv n - n_0$ $n' \approx p'$ $|E_{int}| << |E_{app}|$

 \Rightarrow equations of *coupled* continuity for excess concentrations

$$D'\frac{\partial^2 n'}{\partial x^2} + \mu' E_x \frac{\partial n'}{\partial x} + g - R = \frac{\partial n'}{\partial t}$$

"ambipolar transport equation" Non-linear!

With "ambipolar diffusion coefficient" and "ambipolar mobility":

$$D' = \frac{\mu_n n D_p + \mu_p p D_n}{\mu_n n + \mu_p p} \qquad \mu' = \frac{\mu_n \mu_p (p - n)}{\mu_n n + \mu_p p}$$

Example 1: "Ambipolar diffusion"

Excess electrons and holes produced by light close to the surface, in large concentrations compared to the equilibrium (dark) ones.

Electrons have larger mobility and move faster: electrons and holes partly separate (net charge positive close to the surface, negative inside)



The resulting electric field is directed so as to compensate for the different mobilities (electrons slowed down, holes accelerated)

The coupled motion is called ambipolar diffusion. Since electrons and holes move with the same velocity in the same direction, there is no net charge current associated with this motion !

Example 2: Heynes-Shockley experiment





Lecture 32 - exercises

- Exercise 1: An intrinsic Si sample is doped with donors from one side such that $N_D = N_0 \exp(-ax)$. (a) Find an expression for the built-in electric field E(x) at equilibrium over the range for which $N_D >> n_i$. (b) Evaluate E(x) when a = $1\mu m^{-1}$.
- Exercise 2: An n-type Si slice of thickness L is inhomogeneusly doped with phosphorous donor whose concentration profile is given by $N_D(x) = N_0 + (N_L N_0)(x/L)$. What is the formula for the electric potential difference between the front and the back surfaces when the sample is at thermal and electric equilibria regardless of how the mobility and diffusivity vary with position? What is the formula for the equilibrium electric field at a plane x from the front surface for a constant diffusivity and mobility?

Lecture 33 - exercises

- Exercise 1: Calculate the electron and hole concentration under steadystate illumination in an n-type silicon with $G_L = 10^{16} \text{cm}^{-3} \text{s}^{-1}$, $N_D = 10^{15} \text{cm}^{-3}$, and $\tau_n = \tau_p = 10 \ \mu \text{s}$.
- Exercise 2: An n-type silicon sample has $2x10^{16}$ arsenic atoms/cm³, $2x10^{15}$ bulk recombination centers/cm³, and 10^{10} surface recombination centers/ cm². (a) Find the bulk minority carrier lifetime, the diffusion length, and the surface recombination velocity under low-injection conditions. The values of σ_p and σ_s are $5x10^{-15}$ and $2x10^{-16}$ cm², respectively. (b) If the sample is illuminated with uniformly absorbed light that creates 10^{17} electron-hole pairs/(cm²s), what is the hole concentration at the surface?
- Exercise 3: The total current in a semiconductor is constant and is composed of electron drift current and hole diffusion current. The electron concentration is constant and equal to 10^{16} cm⁻³. The hole concentration is given by $p(x)=10^{15} \exp(-x/L) \text{ cm}^{-3}$ (x>0), where L = $12\mu\text{m}$. The hole diffusion coefficient is $D_p=12\text{cm}2/\text{s}$ and the electron mobility is $\mu_n=1000\text{cm}^2/(\text{Vs})$. The total current density is J = 4.8 A/cm². Calculate (a) the hole diffusion current density as a function of x, (b) the electron current density versus x, and (c) the electric field versus x.

Lecture 34 - exercises

- Exercise 1: Excess electrons have been generated in a semiconductor so that at t = 0 the excess concentration is $\Delta n(0) = 10^{15} \text{ cm}^{-3}$. Assuming an excess-carrier lifetime $\tau_n = 10^{-6}$ s, calculate the excess electron concentration and the recombination rate for t = 4µs.
- Exercise 2: Excess electrons and holes are generated at the end of a silicon bar (at x = 0); the silicon bar is doped with phosphorus atoms to a concentration N_D = 10^{17} cm⁻³. The minority lifetime is 10^{-6} s, the electron diffusion coefficient is D_n = 25 cm²/s, and the hole diffusion current is D_p = 10 cm²/s. Determine the steady-state electron and hole concentrations as a function of x (for x >0) and their diffusion currents at x = 10μ m.

Back-up slides

(topics *not* included in the standard program!)

Recombination via traps

Shockley-Read-Hall model

"Low-injection minority lifetime" approximation



"Shockley-Read-Hall" lifetimes - 1

- What happens if these approximations are not valid?
 - *n*, *p* may be comparable (no longer true that n >> p or p >> n)
 - carrier lifetime no longer dominated by availability of:

p-type: "empty" traps for "electron capture" $(N_t^0 = N_t(1-F) \approx N_t)$ n-type: "full" or "ionized" traps for "hole capture" $(N_t^- = N_tF \approx N_t)$



4=c "hole capture" (in a trap)

Net recombination rates for electrons and holes separately:

 $R_c = c_p N_t F$

 $U_n = R_a - R_b$

 $U_{p} = R_{c} - R_{d}$

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"Shockley-Read-Hall" lifetimes - 3

From equilibrium conditions

 $G_L = 0$; detailed balance: $R_a - R_b = R_c - R_d = 0$)

- emission coefficients (e_n, e_p) in terms of:
- capture coeff. ($c_n = v_{th}\sigma_n$, $c_p = v_{th}\sigma_p$)

$$e_n = c_n n_1$$
 $n_1 = n_i e^{(E_i - E_i)/kT}$
 $e_p = c_p p_1$ $p_1 = n_i e^{(E_i - E_i)/kT}$



"Shockley-Read-Hall" lifetimes - 4

non-equilibrium steady-state

 $(G_L = constant \neq 0, and: U_n = R_a - R_b = U_p = R_c - R_d \neq 0$): see SZE eq. (63)





- A special case: the previous "Low-injection minority lifetime" result
- For very high doping concentrations, direct transitions become likely: this can be modeled by making \(\tau_n\) and \(\tau_p\) concentration-dependent\)

"ambipolar" transport

Two examples: (1) Bipolar diffusion (2) Shockley experiment

"Ambipolar transport" - equations

Special case: homogeneous semiconductor \Rightarrow Thermal equilibrium concentrations n₀, p₀ constant (time and space)

$$D_{p} \frac{\partial^{2} p'}{\partial x^{2}} - \mu_{p} \left(E_{x} \frac{\partial p'}{\partial x} + p \frac{\partial E_{x}}{\partial x} \right) + g_{p} - \frac{p}{\tau_{p}} = \frac{\partial n'}{\partial x}$$
$$D_{n} \frac{\partial^{2} n'}{\partial x^{2}} + \mu_{n} \left(E_{x} \frac{\partial n'}{\partial x} + n \frac{\partial E_{x}}{\partial x} \right) + g_{n} - \frac{n}{\tau_{n}} = \frac{\partial n'}{\partial x}$$
$$\vec{\nabla} \cdot \vec{E} = \frac{\partial E_{x}}{\partial x} = \frac{q}{\varepsilon} (p' - n') \qquad p' \equiv p - p_{0} \quad n' \equiv n - p'$$

Assume:

- Small internal electric field, $|E_{int}| << |E_{app}|$ with respect to the applied field - Almost complete balance $n' \approx p'$ of electron and hole concentrations - Generation, recombination $g_n = g_p \equiv g$ $\frac{n}{\tau_{nt}} = \frac{p}{\tau_{pt}} \equiv R$

 n_0

"Ambipolar transport" - equations

We get then :

$$D_{p} \frac{\partial^{2} n'}{\partial x^{2}} - \mu_{p} \left(E_{x} \frac{\partial n'}{\partial x} + p \frac{\partial E_{x}}{\partial x} \right) + g - R = \frac{\partial n'}{\partial t} \qquad \times \mu_{p} p$$
$$D_{n} \frac{\partial^{2} n'}{\partial x^{2}} + \mu_{n} \left(E_{x} \frac{\partial n'}{\partial x} + n \frac{\partial E_{x}}{\partial x} \right) + g - R = \frac{\partial n'}{\partial t} \qquad \times \mu_{n} n$$

Multiply (see above), add and divide by $\mu_n n + \mu_p p$

$$D'\frac{\partial^2 n'}{\partial x^2} + \mu' E_x \frac{\partial n'}{\partial x} + g - R = \frac{\partial n'}{\partial t}$$

"ambipolar transport equation" Non-linear!

With "ambipolar diffusion coefficient" and "ambipolar mobility":

$$D' = \frac{\mu_n n D_p + \mu_p p D_n}{\mu_n n + \mu_p p} \qquad \mu' = \frac{\mu_n \mu_p (p - n)}{\mu_n n + \mu_p p}$$
"Ambipolar transport"

In an extrinsic semiconductor under low injection, the ambipolar mobility coefficients reduce to the minority-carrier parameter values, that are constant

p-type
minority: electrons
$$D_n \frac{\partial^2 n'}{\partial x^2} + \mu_n E_x \frac{\partial n'}{\partial x} + g' - \frac{n'}{\tau_{n0}} = \frac{\partial n'}{\partial t}$$

n-type
minority: holes
$$D_p \frac{\partial^2 p'}{\partial x^2} - \mu_p E_x \frac{\partial p'}{\partial x} + g' - \frac{p'}{\tau_{p0}} = \frac{\partial p'}{\partial t}$$

The behaviour of excess majority carriers follows that of minority!!!

Example 1: "Ambipolar diffusion"

Excess electrons and holes produced by light close to the surface, in large concentrations compared to the equilibrium (dark) ones.

Electrons have larger mobility and move faster: electrons and holes partly separate (net charge positive close to the surface, negative inside)



The resulting electric field is directed so as to compensate for the different mobilities (electrons slowed down, holes accelerated)

The coupled motion is called ambipolar diffusion. Since electrons and holes move with the same velocity in the same direction, there is no net charge current associated with this motion !

Example 1: "Ambipolar diffusion"

Charge currents for electrons and holes:

$$j_{x,n} = |q|D_n \frac{\partial n}{\partial x} + |q|n\mu_n E_x$$

$$j_{x,p} = -|q|D_p \frac{\partial p}{\partial x} + |q|p\mu_p E_x$$

The net current density vanishes! Associated electric field:

$$j_x = j_{x,n} + j_{x,p} = 0 \implies E_x = \frac{D_n \partial n / \partial x - D_p \partial p / \partial x}{n\mu_n + p\mu_p}$$

The particle currents are therefore equal for electrons and holes:

$$j_{e} = j_{h} = \frac{D_{n}n\mu_{n} + D_{p}p\mu_{p}}{n\mu_{n} + p\mu_{p}} \frac{\partial n}{\partial x} = D_{amb} \frac{\partial n}{\partial x}$$
$$D_{amb} = \frac{D_{n}n\mu_{n} + D_{p}p\mu_{p}}{n\mu_{n} + p\mu_{p}}$$
is the "ambipolar diffusion coefficient"

A "steady-state" example: locally illuminated semiconductor bar

Ingredients and qualitative expectations



n-type; non-equilibrium; open-circuit; Local steady illumination

Diffusion of excess carriers (p', n')

$$p' \equiv p - p_0 = \Delta p$$
 $n' \equiv n - n_0 = \Delta n$

Diffusion currents, but also drift currents due to the electric field E_x $J_h = q\mu_h pE_x - qD_h \frac{dp'}{dx}$ $J_e = q\mu_e nE_x + qD_e \frac{dn'}{dx}$ $J = J_e + J_h = 0$ Electric field E_x (charge unbalance!) $\frac{dE_x}{dx} = \frac{q}{\varepsilon} (p' - n') \neq 0$ The local charge unbalance is small! $\left| \frac{p' - n'}{p'} \right| \approx \left| \frac{p' - n'}{n'} \right| <<1$

Ingredients and qualitative expectations

holes (h): minority electrons (e): majority

drift (e,h): n >> p $q\mu_e nE_x >> q\mu_h pE_{\star}$ $J_{h} = q\mu_{h}pE_{x} - qD_{h}\frac{dp'}{dx}$ $J_{e} = q\mu_{e}nE_{x} + qD_{e}\frac{dn'}{dx}$ $J = J_{e} + J_{h} = 0$ Diffusion currents, but also drift currents due to the elect Diffusion (e,h): opposite currents, comparable sizes Electric field E (charge units) $\Rightarrow \left| q\mu_{h} pE_{x} \right| << \left| qD_{h} \frac{dp'}{dx} \right|$ $\Rightarrow J_h \approx -qD_h \frac{dp'}{dx}$ in this case

n-type; non-equilibrium; open-circuit; Local steady illumination

Diffusion of excess carriers (p', n')

$$p' \equiv p - p_0 = \Delta p$$
 $n' \equiv n - n_0 = \Delta n$

drift currents due to the electric field E_{x}

Electric field E_x (charge unbalance!)

$$\frac{dE_x}{dx} = \frac{q}{\varepsilon} (p' - n') \neq 0$$

The local charge unbalance is small! $\left|\frac{p'-n'}{n'}\right| \approx \left|\frac{p'-n'}{n'}\right| << 1$

minority carriers flow mainly by diffusion

Under conditions of:

- comparable mobilities
- small injection in uniform extrinsic material

the minority-carrier current will be comparable to the majority-carrier current only if minority carriers flow mainly by diffusion



Under conditions of: - comparable mobilities - small injection in uniform extrinsic material the minority-carrier current will be comparable to the majority-carrier current only if minority carriers flow mainly by diffusion

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The large supply of majority carriers effectively "shields" the minority ones from producing any significant space charge.

The small fields that are generated by slight departures from neutrality serve to adjust the majority-carrier current to the general conditions of the problem, without producing significant effects on minority carriers.

An approximate calculation of J_h , J_e , E_x , p', n' in the "quasi-neutral" n' \approx p' approximation (*without* enforcing Gauss' law with p' - n' = 0) will be quite satisfactory; of course, the small p' - n' will not be very accurately determined from E_x found in this way

Approximate quantitative solution



Assuming "quasi-neutral" behaviour:

$$p' \approx n'$$
 $\frac{dp'}{dx} \approx \frac{dn'}{dx}$

(well justified in most cases)



Approximate charge unbalance (dE_x/dx)

Nearly exact solution



Nearly exact solution

