

Exam #1 — EE 482
Winter 2010Mean: 26.4
Median: 28
Std. Dev: 10.4

The test is open book/open notes. Show all work. Be sure to state all assumptions made and **check** them when possible. The number of points per problem are indicated in parentheses (50 total).

1. The electron concentration in a region of silicon depends linearly on depth with concentration of $5 \times 10^{15} \text{ cm}^{-3}$ at surface ($x = 0$) and 10^{15} cm^{-3} at depth of $x = 500 \text{ nm}$. If the vertical electron current density in this region is constant at $J_n = 100 \text{ A/cm}^2$, calculate the electric field near $x = 500 \text{ nm}$. You may assume that the mobility is constant at $1250 \text{ cm}^2/\text{Vs}$. (12)

$$J_n = q(\mu_n n E + D_n \frac{dn}{dx})$$

$$\mu_n = 1250 \frac{\text{cm}^2}{\text{V}\cdot\text{s}} \quad D_n = \frac{kT}{q} \mu_n$$

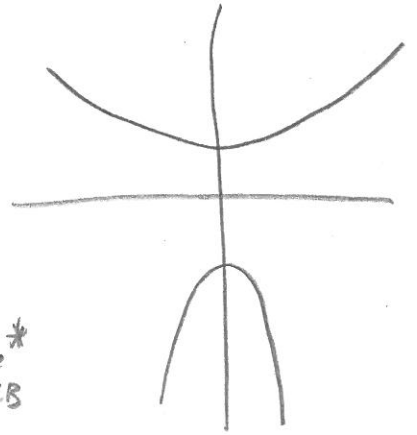
$$J_n = q \mu_n \left(n E + \frac{kT}{q} \frac{dn}{dx} \right)$$

$$E = \frac{1}{n} \left[\frac{J_n}{q \mu_n} - \frac{kT}{q} \frac{dn}{dx} \right] = \frac{1}{10^{15} \text{ cm}^{-3}} \left[\frac{100 \text{ A/cm}^2}{(1.6 \times 10^{-19} \text{ C})(1250 \frac{\text{cm}^2}{\text{V}\cdot\text{s}})} - (0.026 \text{ V})(8 \times 10^{19} \text{ cm}^{-4}) \right]$$

$$= 10^{-15} \text{ cm}^3 \left[\frac{100 \text{ V/cm}^4}{2 \times 10^{16}} - 2.08 \times 10^{18} \frac{\text{V}}{\text{cm}^4} \right]$$

$$= 2.58 \times 10^3 \frac{\text{V}}{\text{cm}} = 2580 \frac{\text{V}}{\text{cm}}$$

2. (a) Sketch the band diagram (E vs. k along main symmetry directions) for a semiconductor which has $\mu_p > \mu_n$ and $N_v < N_c$. The qualitative relations should be clear from your sketch. Assume scattering lifetimes are equal for holes and electrons. Briefly note reasoning. (10)



$$i) \mu_p = \frac{q\tau_p}{m_h^*}, \mu_n = \frac{q\tau_n}{m_e^*} \quad \mu_p > \mu_n \Rightarrow \frac{1}{m_h^*} > \frac{1}{m_e^*}$$

$$\tau_p = \tau_n \Rightarrow \left(\frac{d^2E}{dk^2}\right)^{VB} > \left(\frac{d^2E}{dk^2}\right)^{CB}$$

$$ii) N_v = 2N_{max} \left(\frac{2\pi m_h^* kT}{h^2} \right)^{3/2}$$

$$N_v < N_c \Rightarrow m_h^* < m_e^* \text{ and/or } N_{max}^{VB} < N_{min}^{CB}$$

- (b) Calculate the intrinsic carrier concentration at 300 K in a semiconductor which has a single valence band maxima given by $E = E_v - (\hbar^2 k^2 / 2m_0)$ and eight conduction band minima in the $\langle 111 \rangle$ directions with $E = E_c + 2\hbar^2(k - k_0)^2 / m_0$ where $k_0 = (\pm a, \pm a, \pm a)$, $E_c - E_v = 1$ eV and m_0 is the free electron mass. (10)

$$n_i = \sqrt{N_c N_v} \exp(-E_g / 2kT)$$

$$E_g = E_c - E_v = 1 \text{ eV}$$

$$N_c = 2N_{min} \left(\frac{2\pi m_e^* kT}{h^2} \right)^{3/2}$$

$$m_e^* = \frac{\hbar^2}{\left(\frac{d^2E}{dk^2}\right)_{CB}} = \frac{\hbar^2}{4\hbar^2/m_0} = \frac{m_0}{4}$$

$$N_v = 2 \left(\frac{2\pi m_h^* kT}{h^2} \right)^{3/2} \quad m_h^* = m_0$$

$$n_i = 2 \left(\frac{2\pi m_0 kT}{h^2} \right)^{3/2} \left[8 \left(\frac{1}{4} \right)^{3/2} (1)^{3/2} \right]^{1/2} \exp\left(-\frac{1 \text{ eV}}{0.052 \text{ eV}}\right)$$

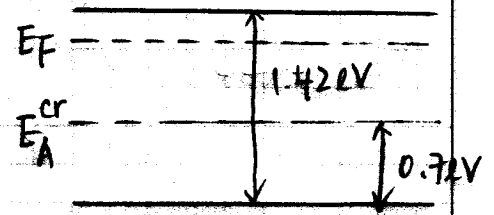
$$= 2 \left(\frac{2\pi (9.11 \times 10^{-31} \text{ kg}) (0.026 \text{ eV})}{(6.63 \times 10^{-34} \text{ J s})^2} \right)^{3/2} 4.45 \times 10^{-9} = 1.12 \times 10^{17} \text{ m}^{-3}$$

$$= 1.12 \times 10^{11} \text{ cm}^{-3}$$

Exam 1.

3(A) $N_D = 10^{16} \text{ cm}^{-3}$ $N_A = 2 \times 10^{15} \text{ cm}^{-3}$ → assume all shallow dopants → fully ionized.
 assume that this is an n-type material

Steps:
 - Proof of assumptions
 - Neutrality @ $n = 10^9 \text{ cm}^{-3}$ = $N_C \exp\left(\frac{E_C - E_F}{KT}\right)$



$$E_F - E_C = -KT \ln \frac{N_C}{n} = -25 \text{ meV} \ln \left(\frac{4.3 \times 10^{17} \text{ cm}^{-3}}{10^9 \text{ cm}^{-3}} \right) \quad (4)$$

$$= -0.497 \text{ eV} \quad (0.514 \text{ eV if } 25.9 \text{ meV was used})$$

$$E_F - E_A^{Cr} = E_g - (E_C - E_F) - (E_A^{Cr} - E_V) \quad (4)$$

$$= (1.42 - 0.497 - 0.7) \text{ eV} = 0.223 \text{ eV} \quad (>> 3KT)$$

Cr is fully ionized because E_A^{Cr} is below E_F .

Material neutrality gives: $n = N_D^+ - N_A^- - N_A^{Cr\bar{o}}$

Main issue: - used pre-doping level $N_A^{Cr\bar{o}} = (10^{16} - 2 \times 10^{15} - 10^9) \text{ cm}^{-3}$ to calculate E_F

- assume ionization without proving. $\approx 8.00 \times 10^{15} \text{ cm}^{-3}$ (3 s.f.) (2)

- flipped the e^- & hole relationship?

(b) $n = 10^9 \text{ cm}^{-3}$

- n is the majority carrier, not p .

$$p = \frac{n_i^2}{n} = \frac{(2.4 \times 10^6)^2}{10^9} \text{ cm}^{-3} = 5760 \text{ cm}^{-3} \quad (3) \leftarrow \text{1 for correct } \mu.$$

$$R = \frac{1}{\frac{q\mu_p p}{1} + \frac{q\mu_n n}{1}} = \frac{1}{(4.6 \times 10^{-19} \text{ C})(6 \times 10^3 \frac{\text{cm}^2}{\text{Vs}})(10^9 \text{ cm}^{-3})} \quad (1)$$

$$\text{negligible} = 1.04 \text{ M}\Omega \cdot \text{cm}$$

$$R_Q = \frac{1}{\int_0^{200 \mu\text{m}} q\mu_n n dx} = \frac{p}{200 \times 10^{-4} \text{ cm}} \quad (3)$$

$$= 52.1 \text{ M}\Omega \quad (1)$$

$$\text{Total doping} = (10^{16} + 2 \times 10^{15} + 8 \times 10^{15}) \text{ cm}^{-3} = 2 \times 10^{16} \text{ cm}^{-3}$$

Main issue: - used pre-doping p to calculate P .
 - forgot that μ s are related to total doping