

SEMICONDUCTOR OPTOELECTRONICS

Questions & Problems for Revision

PART-I: Semiconductor Physics for Optoelectronics

Note: The bold numbers in brackets indicate typical marks allocated to the question in a one-hour written test for 25 marks (or 2 hr written-test for 50 marks)

- 1) What subject does the course title *Semiconductor Optoelectronics* refer to, and what is its importance? (Explain briefly) (2)
- 2) Draw qualitatively the $E-k$ diagrams for an n -type and a p -type *degenerate semiconductor* at 0 K. (For both cases, indicate the positions of the Fermi levels; show the filled portions of the bands by shaded regions.) (2)

- 3) Given that the density of states in a semiconductor is

$$\rho_c(E) = \frac{1}{2\pi^2} \left(\frac{2m_c}{\hbar^2} \right)^{3/2} (E - E_c)^{1/2} \rightarrow \text{Conduction Band}$$

$$\rho_v(E) = \frac{1}{2\pi^2} \left(\frac{2m_v}{\hbar^2} \right)^{3/2} (E_v - E)^{1/2} \rightarrow \text{Valence Band}$$

- a) Obtain expressions for the carrier concentrations n and p in a semiconductor, under the 'Boltzmann approximation'. (3)
 - b) Consider a particular sample of InP ($E_g = 1.36$ eV at 300 K) for which E_f lies exactly in the middle of the bandgap. Calculate the concentration of electrons in the conduction band of this sample. (Given: $m_v = 0.5m_0$; $m_c = 0.07m_0$) (2)
 - c) Is the above sample an n -type or a p -type semiconductor? (Give reason for your answer) (1)
- 4)
 - a) What do you understand by the term 'density of states'? Why is it necessary to know the density of states in the study of semiconductor devices? (Explain briefly) (3)
 - b) Show that the *density of states* for a semiconductor *quantum well* at energies $E(q = 1)$, $E(q = 2)$, etc. in the conduction band, for example, are the same as that for the bulk semiconductor. (3)
 - 5) The carrier concentrations in the valence band and conduction band of a semiconductor in thermal equilibrium (under the Boltzmann approximation) are given by -

$$p = N_v e^{(E_v - E_f)/kT}; \quad n = N_c e^{(E_f - E_c)/kT}$$

Show that the *law of mass action* is not valid in the junction region of a forward biased $p-n$ diode. (3)

- 6) Indium Phosphide (InP) is a direct bandgap material with $m_v = 0.5m_0$. For this material, calculate the number of states between the valence band edge and an energy level that is 0.1eV below the valence band edge. (3)

- 7) Consider a certain type of semiconductor material in which the *density of states* (in the *k-space* is given by

$$\rho(k) = \frac{k}{\pi d}$$

where d is a constant. Obtain an expression for $\rho_c(E)$ near the band edge, and plot qualitatively its variation with E . (2)

- 8) If the effective mass of electrons in the conduction band for the above semiconductor (in the previous question) is $0.10 m_0$ and $d = 100$ nm, determine the number of available states in the conduction band up to the level 0.2 eV above E_c . (3)

- 9) Show that under Boltzmann approximation, the carrier concentration of holes in the valance band of a semiconductor is given by

$$p = N_v e^{-\frac{(E_v - E_f)}{kT}} \quad (3)$$

- 10) What is meant by *bandgap engineering*? State any two methods that could be employed to 'engineer' the bandgap of semiconductor devices. (Explain briefly how do these methods result in bandgap engineering). (4)

- 11) What are *ternary-* and *quaternary* (compound) semiconductors? State two important reasons for using ternary and quaternary semiconductors in Optoelectronics. (No explanation) (2)

- 12) Consider a double heterostructure made up of $\text{GaAs}_{0.6}\text{P}_{0.4}/\text{GaAs}/\text{GaAs}_{0.6}\text{P}_{0.4}$ layers, wherein the thickness of the GaAs layer is 10 nm; the dopant concentrations are such that both GaAs and $\text{GaAs}_{0.6}\text{P}_{0.4}$ are characterized by the same Fermi energy. The direct bandgap energy (in eV at 300 K) of the ternary compound $\text{GaAs}_{1-x}\text{P}_x$ is given by

$$E_g(x) = 1.424 + 1.150x + 0.176x^2; \quad 0 \leq x \leq 0.45$$

- a) If the conduction- and valence band-edge discontinuities are in the ratio of $60:40$, draw qualitatively the energy band diagram corresponding to the double heterostructure. (Indicate clearly the values of the energy discontinuities.) (3)
- b) Does the above structure form a quantum well? (Justify your answer by your own calculations/data). Given $m_c = 0.067m_0$ (2)
- 13) $\text{Ga}_{0.47}\text{In}_{0.53}\text{As}$ (with $E_g = 0.72$ eV) is a widely used photodetector material in the receivers of optical fiber communication systems. Give any two important reasons for the choice of this detector material. (2)

- 14) Consider the formation of a heterojunction between $p\text{-Ga}_{0.7}\text{Al}_{0.3}\text{As}$ and $n\text{-GaAs}$. Given that the *electron affinity* of GaAs is 4.1 eV, $(E_f - E_v)_{\text{GaAlAs}} = 0.3$ eV, $(E_c - E_f)_{\text{GaAs}} = 0.2$ eV. Draw the energy band diagram of the (unbiased) $p\text{-}n$ junction. Calculate the magnitude of *band-edge discontinuities*, and indicate the same in the energy band diagram. (You may assume that the conduction- and valence band-edge discontinuities are in the ratio of $65:35$) (3)

15) The direct bandgap energy (in eV) of the ternary compound $\text{GaAs}_{1-x}\text{P}_x$ is given by

$$E_g(x) = 1.424 + 1.150x + 0.176x^2; \quad 0 \leq x \leq 0.45$$

What alloy composition of $\text{GaAs}_{1-x}\text{P}_x$ would correspond to a bandgap wavelength of 680 nm? (3)

16) What is meant by *Strained layer epitaxy*? From a device engineer's point of view, explain briefly the advantage and constraints associated with it. (3)

17) What are strained-layer quantum wells? Explain briefly with the help of relevant diagrams, and mention their importance in *Semiconductor Optoelectronics*. Name two widely used processes to fabricate semiconductor quantum well structures. (3+1)

18) The *critical thickness* of layers, for epitaxial growth of defect-free strained layers on a semiconductor substrate, is given by

$$d_c = \frac{a_s}{2|\varepsilon|}$$

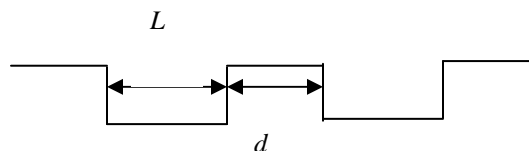
where ε is the *strain* and a_s is the lattice constant of the substrate. What does the above mean to a device-engineer? (Elucidate, with the help of some typical materials and numbers involved, how the critical thickness varies with strain). (3)

19) When do we talk of *quasi Fermi levels* in a semiconductor? What does the energy separation between these levels indicate? State one situation each when the separation between the quasi Fermi levels would be positive, and negative. (2+2)

20) Consider a thin sheet of GaAs of dimensions $1\text{mm} \times 1\text{mm} \times 5\text{nm}$ that forms a quantum well. If the effective mass of electrons in the conduction band is $0.07 m_0$, draw a schematic variation of the density of states with electron energy in the conduction band. Mark the axes with relevant numbers. (4)

21) Consider a semiconductor laser whose active (emission) region comprises of the double heterostructure $\text{Ga}_{0.7}\text{Al}_{0.3}\text{As}/\text{GaAs}/\text{Ga}_{0.7}\text{Al}_{0.3}\text{As}$. Which of the following wavelengths may the laser emit: 630 nm, 730 nm, 830 nm, 930 nm? Justify your answer. (3)

22) Consider a semiconductor heterostructure device comprising of alternate layers of high- and low bandgap materials, that result in a periodic potential variation for conduction band electrons as shown in the figure:



Assume that the width of the potential well (L) is small enough so as to support only one electron eigenstate E_0 in a single (isolated) well. Plot qualitatively the energy eigenvalues of electrons as a function of the number of wells in the device when i) $d = 0.2 L$, ii) $d = L$, and iii) $d = 10 L$. (Plot three different figures corresponding to the above three cases). (3)

23) A particular double heterostructure (DH) has 80nm thick layer of a semiconductor **A** ($E_g = 1.5$ eV) sandwiched between two thicker layers of another semiconductor **B** ($E_g = 2.0$ eV). Given that the effective mass of electrons at room temperature is $m_c = m_0$ in both semiconductors **A** and **B**. Determine whether the given DH forms a quantum well or not? (3)

24) Consider the following four binary semiconductors:

GaP: (5.45, 2.26) AlP: (5.46, 2.45) GaAs: (5.65, 1.42) AlAs: (5.66, 2.16)

The numbers in the brackets give the lattice constant (in Å) and bandgap energy (in eV), respectively.

- Making use of the given numbers, draw qualitatively the positions of these compounds on the 'lattice constant' vs. 'bandgap energy' diagram. (1)
- Which ternary compound is lattice matched to GaP? (1)
- Which ternary compounds would correspond to a bandgap wavelength of $0.62 \mu\text{m}$? Also, from your plot, approximately estimate their compositions. (2)
- Which compound (with different compositions) would represent the area enclosed by the polygon (obtained by joining the points corresponding to the binary compounds)? (1)

25) What is an *Ohmic contact*? Under what conditions does a Schottky junction behave like an Ohmic contact? (Explain briefly with the help of relevant diagrams.) (3)

26) Draw typical I-V characteristics of a Schottky junction diode and an ideal ohmic contact. How would one achieve a near ohmic contact using a Schottky junction diode, in spite of a Schottky barrier? (Explain briefly with the help of energy band diagram) (4)