

[This question paper contains 4 printed pages.]

990

Your Roll No. ....

**B.Sc. (Hons.) / II**

**C**

**BIOCHEMISTRY – Paper VI**

(Biochemical and Biophysical Techniques)

(Admissions of 2000 and onwards)

*Time : 3 Hours*

*Maximum Marks : 60*

*(Write your Roll No. on the top immediately  
on receipt of this question paper.)*

*Attempt Five questions in all, including  
Q. No. 1 which is compulsory.*

1. (a) Explain briefly :

- (i) Glass cuvettes are not used to measure absorbance at 260 nm
- (ii) Precipitation of proteins by organic solvent is carried at low temperature
- (iii) Gel filtration columns can be used for desalting

(b) Give one word for the following :

- (i) A molecule or part of molecule that is responsible for absorbance of light

P.T.O.

- (ii) A fluorescent molecule that is added to a macromolecule to study its property
  - (iii) Gel filtration columns can be calibrated for void volume using this dye
  - (iv) A strongly basic anion exchanger
  - (v) An ion electrostatically bound to the ion exchanger
  - (vi) The gas used in GLC
- (c) Define the following :
- (i) Exclusion limit of the gel
  - (ii) Sedimentation coefficient
  - (iii) Partition coefficient
  - (iv) Generation time (6.6.4)
2. (a) State Beer's Law and Lambert's Law. What are the limitations of Beer's Law ?
- (b) A solution containing ATP is found to have an absorbance of 0.17 at 260 nm in a 1.0 cm cuvette. If the molar extinction coefficient of ATP at 260 nm is  $1.54 \times 10^4 \text{ M}^{-1} \text{ cm}^{-1}$  calculate the concentration of ATP solution.
- (c) Discuss the working of a GM counter. (4,4,3)

3. (a) How do you detect the following :
- (i) Amino acids on paper
  - (ii) DNA in an agarose gel
  - (iii) Sugars on TLC plate
  - (iv) Proteins in a polyacrylamide gel
- (b) Distinguish between the following :
- (i) Intrinsic and extrinsic colours
  - (ii) Stacking gel and separating gel
- (c) Discuss why Ammonium sulfate is preferred over other salts for precipitating proteins. (4.4.3)
4. (a) Discuss the principle of ion-exchange chromatography.
- (b) Classify the following Ion Exchangers as Cation or Anion Exchanger
- (i) DEAE-Sephadex
  - (ii) P-Cellulose
  - (iii) SP-Sepharose
  - (iv) CM-Cellulose
- (c) Describe the process of dialysis. (4.4.3)
5. (a) Mention the role of the following :
- (i) Bromophenol blue in the sample buffer

- (ii) PMT in spectrophotometer
  - (iii) Ampholytes in isoelectric focusing
  - (iv) High temperature in GLC
- (b) How do you purify an enzyme by affinity chromatography ?
- (c) Give two methods for sterilization of growth media. (4,4,3)
6. (a) Explain briefly how nucleic acids can be purified using affinity chromatography.
- (b) Give two methods of coupling a ligand to the affinity matrix.
- (c) Write a short note on phase contrast microscopy. (4,4,3)
7. Give the applications of the following :
- (a) Gel-filtration chromatography
  - (b) Radioisotopes in Biology
  - (c) Isoelectric focusing (4,4,3)
8. Write short notes on the following :
- (a) Determination of molar extinction coefficient of a solute
  - (b) Precipitation of proteins by inorganic salt
  - (c) Density gradient centrifugation (4,4,3)
- (200)