

4. Draw a schematic to illustrate (any two) :

(7.5×2=15)

(i) Renaturation of denatured ribonuclease.

(ii) A discontinuous SDS-PAGE.

(iii) Oxygen saturation curves of haemoglobin & myoglobin.

5. (a) Why do you think Sanger's protein sequencing isn't a preferred method over any other method you know? Nevertheless, justify his choice of using FDNB as a tag for N-terminal labelling.

(b) Are there chances/risks of a patient developing an immune reaction on receiving a therapeutic protein made in *E. coli*? Give reasons and solutions. (8,7)

6. (a) Justify the use of pectinase and xylanase in juice processing and proteases in detergents? To purify such enzymes, would you prefer high-pressure chromatography or low-pressure chromatography?

(b) Explain why i) Poor-quality data and ii) Absence of informed consent constitute ethical issues while considering proteomic data in biomedical applications? (8,7)

(200)

[This question paper contains 4 printed pages.]

Your Roll No.....

Sr. No. of Question Paper : 9836

K

Unique Paper Code : 3183010016

Name of the Paper : Protein Structure and
Function: Advanced Concepts
and Biomedical Applications

Name of the Course : B.Sc. (H) Biomedical
Sciences. Discipline
Specific Elective (NEP)

Semester : VII

Duration : 2 Hours

Maximum Marks : 60

Instructions for Candidates

1. Write your Roll No. on the top immediately on receipt of this question paper.
2. Attempt **four** questions in all. Question No. 1 is compulsory.
3. Subparts of the questions should be attempted together.
4. Draw illustrations or diagrams wherever necessary.

P.T.O.

1. (a). Define : (5×1=5)

- (i) Ampholytes
- (ii) HRP
- (iii) Enzyme replacement therapy
- (iv) Domain swapping
- (v) Helix-turn-helix motif

(b) Give one word for : (5×1=5)

- (i) A large, barrel-shaped molecular machine that assists in the proper folding of proteins.
- (ii) The manual sequencing method that uses Phenylisothiocyanate (PITC) to cleave the N-terminal residue.
- (iii) An amino acid constituent of an electrophoresis that plays a key role in protein separation.
- (iv) A technique created to provide protein-friendly alternatives to high-pressure liquid chromatography.
- (v) A chemical reagent that specifically cleaves peptide bonds at the C-terminal of methionine residues.

(c) Give two major differences between the following :
(2.5×2=5)

- (i) Myoglobin and haemoglobin.
- (ii) FPLC & PMF

2. Give the principle and applications/significance of (any two) : (7.5×2=15)

- (i) Post-translational modification.
- (ii) Alkaline Phosphatase (ALP) in protein detection assays. Give an example.
- (iii) ABC Transporter Proteins as a major class of membrane proteins.

3. Write a short note (any two) on the following :
(7.5×2=15)

- (i) DNA-binding regulatory proteins.
- (ii) Different methods for determining protein stability.
- (iii) Chaperonin's energy-intensive assistance in protein-folding.