# 2021

### **BIOCHEMISTRY — HONOURS**

Paper: SEC-A-2

## (Protein Purification Techniques)

Full Marks: 80

The figures in the margin indicate full marks.

Candidates are required to give their answers in their own words as far as practicable.

#### 1. Answer any ten questions:

 $2\times10$ 

- (a) Write down the separating principle of SDS-PAGE.
- (b) What is mass spectrometry?
- (c) What do you mean by isoelectric pH of a protein?
- (d) Give example of an anion exchanger and a cation exchanger.
- (e) Write two differences between mobile phase and stationary phase.
- (f) What is 'salting in' of protein?
- (g) What is dialysis?
- (h) Why ammonium sulphate  $\lceil (NH_4)_2 SO_4 \rceil$  is most commonly used in salting out of proteins?
- (i) What are the common buffers used in HPLC?
- (j) Why is  $C_{18}$  column mostly used in HPLC?
- (k) What is Svedberg unit?
- (l) What is blank run on HPLC analysis?

#### 2. Answer any four questions :

- (a) (i) An enzyme of M.W. = 26,000 and pI = 5.5 is contaminated with a protein of similar M.W. but with pI = 7.2 and another protein of M.W. = 120,000 and pI = 5.4. Suggest a purification strategy.
  - (ii) What are the functions of APS and  $\beta$ -mercaptoethanol in SDS-PAGE?

3+2

- (b) (i) What is  $R_f$  value in TLC? On which factors does  $R_f$  value depend?
  - (ii) Define specific activity of proteins with unit. Why does it increase during protein purification? (1+2)+(1+1)
- (c) (i) What are the differences between adsorption chromatography and affinity chromatography?
  - (ii) Name two common matrix used in partition chromatography.

3+2

Please Turn Over

- (d) (i) Write down the principle of gel filtration chromatography and state one of its uses.
  - (ii) Tropomysin, a 70kd muscle protein, sediments more slowly than does haemoglobin (65kd). Their sedimentation coefficients are 2.6 S and 4.31 S. Which structural features of Tropomysin account for its slow sedimentation? (2+1)+2
- (e) (i) The octapeptide Ala-Val-Gly-Trp-Arg-Val-Lys-Ser was digested with the enzyme trypsin. Would ion-exchange or gel filtration chromatography be most appropriate for separating the products? Explain.
  - (ii) Suppose that the peptide was digested with chymotrypsin. What would be the optimal separation technique?

#### 3. Answer any four questions :

- (a) (i) How would you purify integral and peripheral membrane proteins?
  - (ii) What is solvent fractionation?
  - (iii) Name two membranes used in dialysis.
  - (iv) When would you employ dialysis during protein purification?
  - (v) Differentiate between ultrafiltrations and dialysis.

2+2+2+2+2

- (b) (i) What is isopycnic centrifugation?
  - (ii) What do you mean by gradient elution of ion exchange chromatography?
  - (iii) In what order would the following proteins emerge upon gel filtration of a mixture on sephadex G 200: Myoglobin (M = 16,000), Catalase (M = 500,000), Cytochrome C (M = 12,000), Chymotrypsinogen (M = 26,000) and Serum albumin (M = 65,000)? Explain.
  - (iv) What are the working principles of horizontal electrophoresis using agarose gel?
  - (v) Write down the application of ultracentrifugation.

2+2+2+2+2

- (c) (i) What is void volume?
  - (ii) In a chromatographic column, the stationary phase has an exclusion limit of 80,000 MW. If you tried to use this column material to separate alcohol dehydrogenase (M.W. = 150,000) from  $\alpha$ -amylase (M.W. = 55,000), what would happen and why?
  - (iii) What is katal?
  - (iv) What is the difference between SDS-PAGE and NATIVE PAGE?
  - (v) Why is extraction of proteins from a source done under ice-cold solution? 2+2+2+2+2
- (d) (i) A solution containing egg albumin (pI = 4.6),  $\beta$ -lactoglobulin (pI = 5.2) and chymotrypsinogen (pI = 9.5) was loaded on to a column of DEAE allulose at pH = 5.4. The proteins are then eluted with buffer with pH = 5.5, with an increasing salt concentration. Predict the elution pattern.
  - (ii) Define the following: Partition coefficient, retention time, resolution, reverse phase chromatography.  $2+(2\times4)$

## (3)

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- (e) (i) Is ion exchange a chemisorption process?
  - (ii) What are the disadvantages of HPLC?
  - (iii) In what direction will the following proteins move in an electric field?
    - (A)  $\beta$ -lactoglobulin (pI = 5.2) at pH = 5.0 and pH = 7.0
    - (B) Ribonuclease (pI = 9.5) at pH = 4.5 and pH = 9.5
  - (iv) From a mixture of Glu (pI = 3.08), His (pI = 7.64) and Arg (pI = 10.76), how can you retrieve individual amino acids?