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electrospray ionization mass spectrometry and the following data were obtained:

m/z	773.9	825.5	884.3	952.3	1031.3
Abundance (%)	59	88	100	66	37

Given that  $n_2 = (m_1-1) / (m_2-m_1)$  and  $M = n_2 (m_2 - 1)$ and assuming that the only ions in the mixture arise by protonation, deduce average molecular mass for the protein by this method.

5 + 3 + 4 = 12

- 7.(a) Write the names of different steps of 2 D PAGE and describe the basic principles of the two major steps of 2D PAGE with labeled diagram.
  - (b) Describe the protein-protein interaction study by Yeast two hybrid assay with labelled diagram.
  - (c) Describe the steps of biomarker discovery for cancer using proteomics with a labeled diagram.

4 + 4 + 4 = 12

## Group - E

- 8.(a) What are the pulse sequences that are typically used to characterize a protein molecule using NMR?
  - (b) What specific interactions do these pulse sequences measure?
  - (c) What is a typical number of ensemble models of a protein structure produced by NMR analysis?
  - (d) Name and briefly explain four methods by which NMR resonance signals can be improved.

3 + 3 + 2 + 4 = 12

- 9.(a) Explain why phospho-proteomics represents a prime example of protein modification. Draw the structures of three amino acids that are known to be phosphorylated in a biological context. Draw a flowchart of techniques for the analysis of phosphoproteins.
  - (b) Draw a labeled diagram of a SPR sensorgram. Explain, the industrial application of SPR to proteomics in the area of two component binding reactions.

(1+2+3)+(3+3)=12

# M.TECH/BT/2ND SEM/BIOT 5241/2017 GENOMICS AND PROTEOMICS (BIOT 5241)

Time Allotted: 3 hrs

Full Marks: 70

Figures out of the right margin indicate full marks.

Candidates are required to answer Group A and

<u>Any 5 (five)</u> from Group B to E, taking <u>at least one</u> from each group.

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

Group – A

### (Multiple Choice Type Questions)

1. Choose the correct alternative for the following:

10 ×1=10

- (i) Which type of genomics study describes the transcripts and proteins expressed by a genome?
  (a) Comparative genomics
  (b) Structural genomics
  (c) Functional genomics
  (d) Trancripto-genomics.
- (ii) X-ray diffraction on a protein requires
  (a) a well-ordered protein crystal
  (b) integration of an electron density map
  (c) surface entropy increase
  (d) all of the above.
- (iii) The first genome sequencing project was carried on
  (a) Haemophilus influenza
  (b) E. coli
  (c) ØX 174
  (d) Drosophila melanogaster.
- (iv) Which of the following information about proteins can be obtained by 2D-Gel Electrophoresis?
  (a) MW, pl and quantity
  (b) MW and pl
  (c) pl and quantity
  (d) none of these.
- (v) Individual genetic maps in a given species are
  (a) genetically similar
  (b) genetically identical
  (c) genetically dissimilar
  - (d) not useful in species analysis.

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- (vi) Which of the following statements is not true about sequencing peptides with mass spectroscopy?
  - (a) The entire protein can be sequenced all at once using mass spectroscopy.
  - (b) Two rounds of mass spectroscopy are used to determine sequence.
  - (c) some purified protein must be digested with proteases to eliminate undesirable characteriscts such as hydrophobicity and solubility.
  - (d) In order to determine the sequence, a pure sample of protein is obtained through 2D-PAGE or HPLC.
- (vii) Why co-immunoprecipitation is used?
  - (a) to determine if a protein-of-interest binds to a specific DNA sequence.
  - (b) to examine protein-protein interaction in the nucleus instead of in the cytoplasm.
  - (c) to examine protein-protein interactions in the cytoplasm instead of the nucleus.
  - (d) to allow protein to be expressed in mammalian cell culture.

## (viii) Structural proteomics is concerned with

- (a) coverage of fold space
- (b) low throughput determination of protein structures
- (c) high-throughput determination of protein structures(d) all of the above.
- (ix) Expression of genes can be analyzed by
  - (a) Northern analysis
  - (b) Southern analysis
  - (c) Comparative genomics
  - (d) RNA interference.
- (x) A lab-on-a-chip is
  - (a) a laboratory on a microchip
  - (b) a DNA microarray
  - (c) a protein chip
  - (d) a small device that can perform electrophoretic separations.

# Group - B

- 2. (a) Define STS. Analyze the usage of STS in genomics research.
  - (b) What do you mean by polymorphic STS?

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- (c) What is snip-SNP?
- (d) Define pseudogene.

(2+3)+3+2+2=12

- 3.(a) Discuss the applications of genomics in tracing human migration and evolution.
  - (b) Illustrate the process of TaqMan assay for identifying SNPs.
  - (c) Describe with a flow diagram the technique of AFLP.

4 + 4 + 4 = 12

### Group - C

- 4.(a) "In the transcriptome analysis by quantitative analysis of gene expression fragment lengths of more than 200 nucleotides are used" justify the statement with suitable experimental procedure.
  - (b) (i) To tackle the huge nonredundant data in transcriptomics, some special processing is needed for this purpose. Mention the processes citing suitable reasons.
    - (ii) Describe briefly any such database which is used for this purpose.

## 6 + (3+3) = 12

- 5.(a) "Microarray based method is followed to study the global gene expression profiling". Mention the detailed procedure using oligo nucleotide followed for this microarray based method.
  - (b) To design an optimal oligonucleotide probe certain criteria are needed to follow. Mention those criteria citing the reason behind it.

6 + 6 = 12

## Group - D

- 6.(a) Describe the basic principles of determination of mass of protein by ESI TOF MS with labled diagram.
  - (b) Write the advantages of MALDI TOF over ESI TOF.
  - (c) A protein was isolated from human tissue and subjected to a variety of investigations. Relative molecular mass determinations gave values of approximately 12000 by size exclusion chromatography and 13000 by gel electrophoresis. After purification, a sample was subjected to

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