



Name :

Roll No. :

Invigilator's Signature :

CS/M.TECH(BT)/SEM-2/MBT-215D/2013

2013

GENOMICS AND PROTEOMICS

Time Allotted : 3 Hours

Full Marks : 70

The figures in the margin indicate full marks.

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

GROUP – A

(Multiple Choice Type Questions)

1. Choose the correct alternatives for the following : $10 \times 1 = 10$
 - i) Which is not a ion source in mass spectrometry ?
 - a) ESI
 - b) FAB
 - c) TOF
 - d) MALDI.
 - ii) Trypsin cleave the peptide bond containing
 - a) Arg or Lys
 - b) Glu or Asp
 - c) Met or Trp
 - d) none of these.
 - iii) The 2D Gel electrophoresis provides information about the proteins are
 - a) MW, pI and quantity
 - b) MW and pI
 - c) pI and quantity
 - d) none of these.
 - iv) Structural proteomics deals with
 - a) Cellular localization
 - b) PTMs
 - c) Edman degradation
 - d) 3D structure.



- v) In 2D Gel electrophoresis
 - a) Isoelectric focusing (IF) with SDS-PAGE
 - b) IF with another IF
 - c) SDS-PAGE with another SDS-PAGE
 - d) SDS-PAGE with ion exchange chromatography.
- vi) The most widely used program for multiple sequence alignment is
 - a) BLAST
 - b) FASTA
 - c) CLUSTAL
 - d) CHIME.
- vii) Protein-protein interactions are studied by
 - a) DNA foot printing
 - b) yeast two hybrid system
 - c) ligase chain reaction
 - d) mass spectrometry.
- viii) What is PEST sequence ?
 - a) Proline, Glutamic acid, Serine and Threonine
 - b) Proline, Ethylamine, Serine and Threonine
 - c) Proline, Glutamin, Serine and Threonine
 - d) None of these.
- ix) Sick cell anemia is resulted from the
 - a) mutation prion protein
 - b) mutation in transthyretin protein
 - c) mutation in haemoglobin, HbA
 - d) mutation in α -synuclein protein.
- x) Trypsin cleave the peptide bond containing
 - a) Arg or Lys
 - b) Glu or Asp
 - c) Met or Trp
 - d) none of these.



GROUP – B

(Short Answer Type Questions)

Answer any *three* of the following.

3 × 5 = 15

2. How do you characterize a protein ?
3. What is ubiquitin ? Write down its function in cellular process.
4. What is proteomics ? How many faces of proteomics are there ? State their significance.
5. What is cDNA library ? How will you make cDNA library ?
6. What is Transcriptomics ? What is its use ?

GROUP – C

(Long Answer Type Questions)

Answer any *three* of the following.

3 × 15 = 45

7.
 - a) "Every organism has one genome but many proteomes." Explain.
 - b) Discuss the principle of any one protein estimation method.
 - c) How will you prepare sample of 2D gel electrophoresis ?
 - d) Discuss any one procedure of protein fractionation for 2D gel electrophoresis.
8.
 - a) What is post transitional modification (PTM) ? Give few examples of PTM.
 - b) What is the purpose of PTM ?
 - c) Site an example where PTM changes character of the protein.
9.
 - a) Discuss the working principle of MALDI-TOF MS.
 - b) Write the advantages of MALDI-TOF over ESI-TANDEM.

4 + 4 + 4 + 3

5 + 5 + 5

7 + 8

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10. Write short notes on any *three* of the following : 3×5

- a) Proteasome
- b) Ribosome
- c) Molecular phylogenetics
- d) SUMO protein.

11. Write short notes on any *three* of the following : 3×5

- a) 2D-NMR
- b) Protein splicing
- c) DNA microarray
- d) SNP.

12. a) Write the applications of RFLP and AFLP.

b) Illustrate the procedure of AFLP. $(5 + 5) + 5$

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