

(i) Printed Pages : 3

Roll No.

(ii) Questions : 9

Sub. Code :

2	9	9	4
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Exam. Code :

4	3	7
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M.Sc. 3rd Semester

1125

BIOTECHNOLOGY

Paper-MBIO-305 : Advances in Genomics and Proteomics

Time Allowed : Three Hours]

[Maximum Marks : 80

Note : Attempt five questions in all. Question No. 1 is compulsory. Select one question each from Unit I to Unit IV.

1. Write short answers on :

- (a) Oligonucleotide Fingerprinting
- (b) e-PCR
- (c) E-Value
- (d) Levenshtein distance
- (e) Log Odd Ratio
- (f) Electrospray Ionization (ESI)
- (g) Rooted vs. Un-Rooted Trees
- (h) ChIP-seq.

8×2=16

UNIT-I

- 2. (a) Discuss methods of eukaryotic gene prediction.
- (b) How bioinformatics is useful for discovering information related to protein structure, function and evolution ?

2×8=16

3. (a) How Phase Display technique is used to identify and extract polypeptides with desired properties from a large collection of variants ?
- (b) Explain role of proteomics technologies in crop improvement.
- $2 \times 8 = 16$

UNIT-II

4. (a) What are Expression Libraries ? Explain the process of generation of cDNA expression libraries.
- (b) What are DNA Chips ? How is it different from protein chips ? Give applications of DNA chips.
- $2 \times 8 = 16$
5. (a) Write about protein array technology. How the detection of protein-protein interactions is done on Protein microarrays ?
- (b) Give applications of protein chips in proteomics.
- $2 \times 8 = 16$

UNIT-III

6. (a) Explain Illumina sequence-by-synthesis approach of DNA sequencing.
- (b) Define BLAST. Describe its various forms used to analyze DNA and Protein sequences.
- (c) Differentiate between local and global alignment.
- $8 + 4 + 4 = 16$
7. (a) Explain DNA sequencing with Nanopores and its advantages over other next generation sequencing techniques.
- (b) Why there is a need for 3D structure prediction of macromolecules ? Explain Homology Modeling method of protein structure prediction.
- $2 \times 8 = 16$

UNIT-IV

8. (a) Explain the Mass spectrometry based protein identification method.
- (b) How DNA microarray can be used to identify disease gene expression profile ? $2 \times 8 = 16$
9. (a) Explain in detail ChIP method for Genome-wide identification of transcription factor binding sites.
- (b) Proteomics studies involve high-resolving Gel based or Liquid-Phase separation techniques. Explain Liquid-Phase proteomics technique for protein separation. $2 \times 8 = 16$