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(iii) Protein disordered region

(iv) Genome Browsers

(4,4,4,3)

[This question paper contains 8 printed pages.]

Your Roll No.....

Sr. No. of Question Paper: 1171

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Unique Paper Code

: 2493010008

Name of the Paper

: In-silico Tools in Proteomics

and Genomics

Name of the Course

: B.Sc. (Hons.) Biochemistry

Semester

: V

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. There are 6 questions in all.
- 3. Attempt any 4 questions.
- 4. Question no. 1 is compulsory.
- 5. All questions carry equal marks.

- 1. (a) Expand the following terms -
 - (i) HTML
 - (ii) OMIM
 - (iii) DDBJ
 - (iv) CDS
 - (v) KEGG
 - (vi) PDB
 - (vii) PIR
 - (b) Comment on the following -
 - (i) Sequence similarity is different from sequence identity

- 5. (a) What is Yeast two hybrid (Y2H) technique? What are its advantages?
 - (b) Discuss the current approaches employed in the drug discovery process emphasizing target identification, target validation, lead identification and lead optimization.
 - (c) Discuss the difference between local and global alignment with suitable examples.

(5,5,5)

- 6. Write short notes on the following:
 - (i) Role of AI in Genomics
 - (ii) Human Genome Project

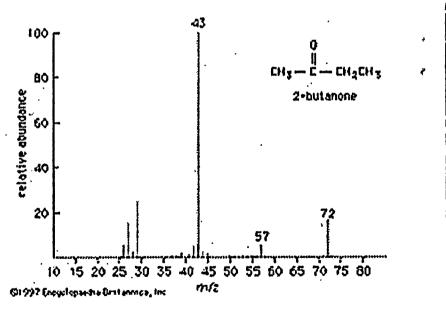
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(ii) NCBI is a composite database.

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- (iii) Lipinsky's rule of 5 distinguishes drug-like and non-drug-like molecules
- (iv) FASTA is a sequence file format
- (c) Fill in the following blanks -
 - (i) _____ is a protein visualization tool.
 - (ii) Demis Hassabis and John M. Jumper won the 2024 Nobel Prize in Chemistry for their work on ______, an AI tool that predicts protein structures.
 - (iii) _____ tool can be used for predicting secondary structures in proteins.

- (b) You have a polynucleotide sequence. Describe the6 reading frames that can be generated?
- (c) Describe the principle of mass spectrometry. The figure above has mass spectrometry data for 2-butanone. Explain each of the peaks obtain in the spectra.



(iv) GLIMMER is a tool for predicting
_____ in microbial DNA, especially the
genomes of bacteria, archaea, and viruses.

(7,4,4)

- 2. (a) Differentiate between the following:
 - (i) Primary and Secondary Databases
 - (ii) Prokaryotic and eukaryotic genome organization
 - (iii) Functional Genomics and Comparative

 Genomics
 - (iv) Pairwise Sequence Alignment & Multiple
 Sequence Alignment

- (b) Describe Ramachandran plot. Explain how it is useful in protein modelling. (12,3)
- 3. (a) Describe Next Generation Sequencing (NGS).

 What are its advantages and disadvantages?
 - (b) Expand and describe BLAST. What is the difference between BLASTN and TBLASTN?
 - (c) Describe any 2 applications of Bioinformatics.
 (6,6,3)
- 4. (a) You have a protein of unknown function from a bacterium. You have made a knock-out mutant, but the bacteria die immediately without the corresponding gene. You have sequenced the protein. Describe the methods to predict the structure of the protein using in silico tools.