## BIRLA INSTITUTE OF TECHNOLOGY, MESRA, RANCHI (END SEMESTER EXAMINATION)

CLASS: MSC/PRE-PHD SEMESTER: II/I BRANCH: BIOTECH SESSION: SP/2024

**SUBJECT: BT417 BIOINFORMATICS** 

TIME: 3 Hours FULL MARKS: 50

## **INSTRUCTIONS:**

- 1. The question paper contains 5 questions each of 10 marks and total 50 marks.
- 2. Attempt all questions.
- 3. The missing data, if any, may be assumed suitably.

4. Befor		the question	n paper, be	sure that y		the correct question paped in the examinates in the examinates in the examinates.		ll. 	
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Q.1(a)							[5]	1	1
Q.1(b)	i. PubMed ii. PDB iii. Swiss-Prot iv. Expasy v. SCOP What is the necessity of multiple sequence alignments (MSA)? Briefly explain the three main classifications of MSA methods.							1,2	2,3
Q.2(a)	Interpret the dot matrix plots shown below, and provide examples of hypothetica sequences that could generate these dot matrix plots.							2,3	4,5
Q.2(b)	Perform the alignment of following sequences using the Needleman-Wunsch algorithm taking the following criteria. Also write the final alignment(s).  Sequence1: AGT; Sequence2: AAGC; [Match: +1, Mismatch: -1, Gap: -2]						[5]	1,2	3
Q.3(a)	List the applications of phylogenetic trees and explain the following terms in the context of a phylogenetic tree by illustrating them on phylogenetic tree. i. Branches, ii. Taxa, iii. Node, iv. Root Node.						[5]	2,3	1,2
Q.3(b)	Construct the phylogenetic trees for the given distance matrix using the UPGMA algorithm, and report the final tree with all the distances.						[5]	2	3
		Α	В	С	D	Е			
	Α	-	22	39	39	41			
	В	-	-	41	41	43			
	С	-	-	-	18	20			
	D	-	-	-	-	10			
	E	-	-	-	-	-			
Q.4(a)	What is Ramachandran plot? Discuss its importance and explain why glycine and proline have distinct Ramachandran plots compared to other amino acids.						[5]	3,4	2,4
Q.4(b)	List all the steps of homology modeling in sequential order and explain all the factors that should be considered for template selection.						[5]	3,4	1,2
Q.5(a)	Explain one of the topics mentioned below, along with its applications and [5] limitations.							3	2
Q.5(b)	i) Docking ii) Pharmacophore iii) QSAR iv) Microarray v) NGS vi) Motif Analysis Imagine a situation like COVID-19 where we were dire need of drug/vaccine, in such scenario how you will utilize the bioinformatics techniques mentioned in question number 5a and design a personalized drug/vaccine. Provide a detailed technical explanation of the computational methodologies, databases involved, accompanied by a concise flowchart detailing each step.								

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