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|---|---|
| <p>a) Both A and R are Individually true and R is the correct explanation of A</p> <p>c) A is true but R is false</p> | <p>b) Both A and R are Individually true but R is not the correct explanation of A</p> <p>d) A is false but R is true</p> |
|---|---|
10. Which of the following algorithm is used in flexible docking CO6 [K<sub>1</sub>]
- |                             |                         |
|-----------------------------|-------------------------|
| a) Smith watermen algorithm | b) DOCK algorithm       |
| c) Conolly's algorithm      | d) Needle man algorithm |

**PART B (10 x 2 = 20 Marks)**  
**(Answer not more than 40 words)**

- |   |                       |
|---|-----------------------|
| 11. Calculate Levenstein and Hamming's distance for the sequences<br>A: TTCGATCCATTG<br>B: ATCAATCGATCG         | CO1 [K <sub>2</sub> ] |
| 12. Mention the application of dot plot analysis  | CO1 [K <sub>1</sub> ] |
| 13. What is PSI BLAST? Give its application   | CO2 [K <sub>1</sub> ] |
| 14. Classify different types of gaps with example.  | CO2 [K <sub>2</sub> ] |
| 15. Mention the drawbacks of iterative methods for MSA  | CO3 [K <sub>2</sub> ] |
| 16. Tell whether PILE UP is a method based on progressive or iterative alignment? Illustrate the steps involved | CO3 [K <sub>3</sub> ] |
| 17. Illustrate Newick format with an example  | CO4 [K <sub>1</sub> ] |
| 18. Mention the significance of bootstrapping in phylogenetic analysis.   | CO4 [K <sub>2</sub> ] |
| 19. List different methods used for protein secondary structure prediction. Give its significance               | CO5 [K <sub>1</sub> ] |
| 20. Discuss how virtual screening is useful in Computer Aided Drug Design?                                      | CO6 [K <sub>2</sub> ] |

**Answer any FIVE Questions:-**  
**PART C (5 x 14 = 70 Marks)**  
**(Answer not more than 350 words)**

- |  |   |     |                   |
|--|---|-----|-------------------|
| 21. a) Explain the various components of the biological sequence formats –FASTA, FASTQ, Genbank with examples. | 7 | CO1 | [K <sub>2</sub> ] |
| b) Describe the file format used in EMBL and PDB database.   | 7 | CO1 | [K <sub>2</sub> ] |
| 22. a) Explain how the scoring matrices PAM and BLOSUM are constructed? Give its significance in alignment.    | 7 | CO2 | [K <sub>2</sub> ] |
| b) Align the given sequences using Needleman Wunsch algorithm<br>A: GCATGCG<br>B:GATTACA                       | 7 | CO2 | [K <sub>3</sub> ] |

match =1, mismatch = -1,gap = -1

23. a) Illustrate the steps involved in progressive alignment with CLUSTALW as an example. 7 CO3 [K<sub>2</sub>]
- b) What is PSSM? Explain how the values in position specific scoring matrix(PSSM) are calculated and state its application 7 CO3 [K<sub>2</sub>]
24. a) Illustrate the steps involved in Maximum Likelihood method. Give its significance 7 CO4 [K<sub>2</sub>]
- b) Construct phylogenetic tree for the given distance matrix using UPGMA method. 7 CO4 [K<sub>2</sub>]

	B	C	D
A	14	11	7
B		11	13
C	-	-	8

25. a) What is homology modeling? Explain the steps involved in it for tertiary structure prediction. 10 CO5 [K<sub>2</sub>]
- b) Describe the tools by which the protein tertiary structures can be compared. 4 CO5 [K<sub>2</sub>]
26. a) Brief the applications of computer aided case study with a case study. 7 CO6 [K<sub>2</sub>]
- b) Elaborate the principle of rigid and flexible docking. Compare and contrast rigid and flexible docking. 7 CO6 [K<sub>2</sub>]

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