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

| CLASS:<br>BRANCH  | BTECH<br>BIOTECHNOLOGY   | SEMESTER : III<br>SESSION : MO/2022 |  |  |  |  |  |
|---|--|-------------------------------------|--|--|--|--|--|
| TIME:   | SUBJECT: BE205R1 BASICS OF BIOINFORMATICS<br>3:00 Hours  | FULL MARKS: 50                      |  |  |  |  |  |
| <ul> <li>INSTRUCTIONS:</li> <li>1. The question paper contains 5 questions each of 10 marks and total 50 marks.</li> <li>2. Attempt all questions.</li> <li>3. The missing data, if any, may be assumed suitably.</li> <li>4. Before attempting the question paper, be sure that you have got the correct question paper.</li> <li>5. Tables/Data hand book/Graph paper etc. to be supplied to the candidates in the examination hall.</li> </ul> |  |                                     |  |  |  |  |  |
| Q.1(a)<br>Q.1(b)  | Differentiate primary and secondary databases with example in bioinformati<br>Designate in terms of data content and application of followings inf<br>databases: NCBI, RCSB and CATH |                                     |  |  |  |  |  |
| Q.1(c)  | Write following file formats with example *.fasta, *.pdb, *.genbank, and *.dr  | nd. [5]                             |  |  |  |  |  |

- Q.2(a) Differentiate the substitution matrix PAM and BLOSUM with proper application.
- [2] Q.2(b) Compose the sequence searching algorithm BLAST stepwise. What is tBLAST? [3]
- Q.2(c) What is Sum of pairs method in multidimensional dynamic programming? [5] Using N-W algorithm for Global Alignment used in Dynamic Programming, complete the following alignment matrix (array) and predicts the possible alignment, (mismatch :-2, match: +2; Gap penalty: -2)

| 0      | T (-2) | C (-4) | G (-6) | A(-10) |  |  |  |
|--------|--------|--------|--------|--------|--|--|--|
| T (-2) |        |        |        |        |  |  |  |
|        |        |        |        |        |  |  |  |
| C (-4) |        |        |        |        |  |  |  |
|        |        |        |        |        |  |  |  |
| C(-6)  |        |        |        |        |  |  |  |
|        |        |        |        |        |  |  |  |
| A(-8)  |        |        |        |        |  |  |  |
|        |        |        |        |        |  |  |  |

- Q.3(a) Theorize with example: Dendogram and Kimura 2P model of DNA substitution with transversion and [2] transition.
- Q.3(b) For the following individual pairwise distances, construct the phylogenetic tree using UPGMA [3] method, (ab:16; ac:32; ad:48; bc28; bd:42; cd:12).
- Q.3(c) What is molecular clock? Synthesize the stepwise methodology for building phylogenetic tree, Fitch-[5] Margolish method OR maximum parsimony method.
- Q.4(a) Originate and illustrate RMSD value and Ramachandran plot.
- Develop and state the algorithms for Protein secondary structure prediction methods: Chou-Fasman Q.4(b) [3] method.

[2]

[2]

[3]

- Q.4(c) Write algorithms/flow-chart for Homology modelling for protein structure prediction. What is SCOP? [5]
- Q.5(a) Briefly discuss Chemoinformatics with different applicable softwares.
- Q.5(b) Briefly introduce Systems Biology.
- Q.5(c) Build the position specific scoring matrix (PSSM) for following motif, ctata atagg; cagcc; cggtt; tgcat. [5]

## :::::25/11/2022::::E