

**MASTER OF COMPUTER SC. & ENGG. 2<sup>ND</sup> SEM. EXAMINATION – 2024****BIOINFORMATICS****Time: Three Hours****Full Marks: 100****Answer any FOUR questions**

1. a) Briefly Describe the BLAST algorithm. Explain with respect to the following example (consider default word length as 4):  
 Input sequence: AILVPTV  
 Database sequence: MVQGWALYDFLKCRAILVGTVIAML
- b) Briefly discuss n-star consensus strategy in AMS algorithms for prediction of PTM sites.
- c) What is molecular docking? How different surface representation methods identify the regions of interest. Describe any two surface representation methods in brief. What is the basic working principle of the DOCK algorithm?

10+5+10 = 25

2. a) Use UPGMA to construct a phylogenetic tree using the following distance matrix:

|   | A     | B     | C     | D     | E     | F     | G   |
|---|-------|-------|-------|-------|-------|-------|-----|
| A | 0.0   |       |       |       |       |       |     |
| B | 19.00 | 0.0   |       |       |       |       |     |
| C | 27.00 | 31.00 | 0.0   |       |       |       |     |
| D | 8.00  | 18.00 | 26.00 | 0.0   |       |       |     |
| E | 33.00 | 36.00 | 41.00 | 31.00 | 0.0   |       |     |
| F | 18.00 | 1.00  | 32.00 | 17.00 | 35.00 | 0.0   |     |
| G | 13.00 | 13.00 | 29.00 | 14.00 | 28.00 | 12.00 | 0.0 |

- b) Consider the following DNA sequences. Construct the corresponding Position Frequency Matrix (PFM) and the resulting Position Probability Matrix (PPM).

GAGGTAAAC  
 TCCGTAAGT  
 CAGGTTGGA  
 ACAGTCAGT  
 TAGGTCATT  
 TAGGTACTG  
 ATGGTAACT  
 CAGGTATAC  
 TGTGTGAGT  
 AAGGTAAGT

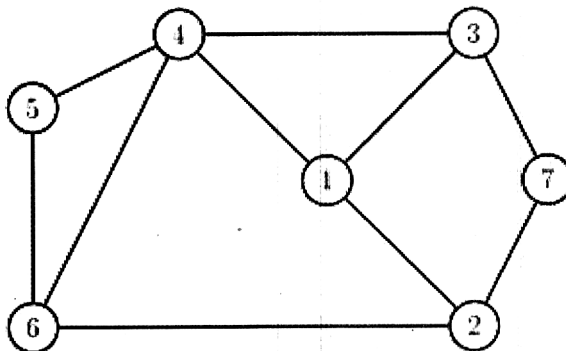
15+10=25

[ Turn over

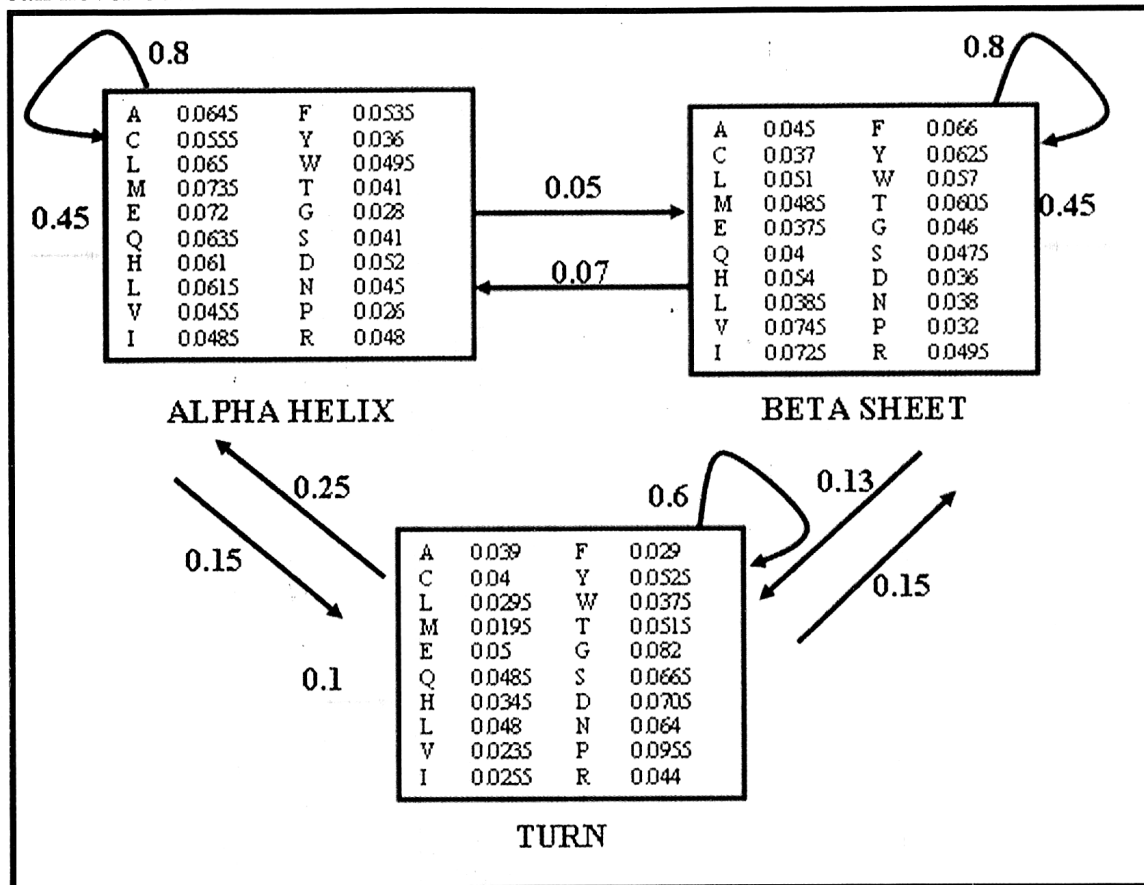
Sequence 1 = THISLINE  
Sequence 2 = ISALIGNED

[illegible]

- What is Protein Secondary Structure Prediction (PSSP)? Briefly discuss the PSIPRED method for PSSP.
- Consider the following graph and estimate the *degree distribution* and *clustering coefficient* for all the nodes in the graph.



- ;) Given the following probability distribution, use Hidden Markov Model to estimate the optimum secondary structure annotation for a primary sequence: **TAS**. T is the starting Amino Acid in the sequence, and assume that the first two Amino Acids in the sequence can never be a "TURN".



5+10+10 = 25

Write short notes on the following (any 5):

- DOT Plot
- Central Dogma of Molecular Biology
- BLOSUM
- Scale Free Network
- Ubiquitination
- KEGG

5x5 = 25