



VI SEMESTER B.TECH (BIOTECHNOLOGY)

END SEMESTER EXAMINATIONS, APRIL 2016

SUBJECT: BIOINFORMATICS [BIO 3201]

Time: 3 Hours

MAX. MARKS: 50

Instructions to Candidates:

- ❖ Answer **ALL FULL** questions.
- ❖ Missing data may be suitably assumed.

- 1A. How many reading frames exist for a double-stranded DNA sequence? Is it possible to correlate the total number of ORFs and the actual number of genes? If yes, why? If no, why not? 3
- 1B. What do you mean by Low Complexity Regions (LCRs)? How would you process the sequence with LCRs? 3
- 1C. Consider the following piece of DNA
- | | | | |
|----|-----|---------------------------------------|----|
| 5' | ... | CCTCACTAGCGACCAGACTATAAATTACGTGACCTGT | 3' |
| 3' | ... | GGAGTGATCGCTGGTCTGATATTTAATGCACTGGACA | 5' |
- 1C. Map the fragments, a=GGTCA, b=AATTAC, c=AGTCTGG, and d=CACTAG occur in the DNA piece 4
- 2A. It is well known that hemoglobin and myoglobin are evolutionarily related proteins. Correlate their similarities at sequence, structure and functional level of hemoglobin and myoglobin 2
- 2B. Do a global alignment of the sequences *m* (CCTCACTA) and *n* (TAGTGAGG) using Needleman-Wunsch algorithm and also explain the trace-back step in detail. 5
- 2C. Explain the alignment strategy to pick a conserved pattern in the small disulfide-rich proteins of different lengths 3
- 3A. Differentiate conservation, semi- conservation, and conserved substitution with examples each 3
- 3B. To amplify the DNA between the two stretch of sequence shown below. Choose the listed primers pairs that will allow to amplify the DNA by PCR
- 5'-GACCTGTGGAGC ----- CATACGGGATTGA-3'
- Primers:
- | | |
|-------------------------|--------------------------|
| (1) 5'-GACCTGTCCAAGC-3' | (5) 5'-CTGGACACCTTCG-3' |
| (2) 5'-CGAAGGTGTCCAG-3' | (6) 5'-CATACGGGATTGA-3' |
| (3) 5'-GCTTCCACAGGTC-3' | (7) 5'-TCAATCCCGTATG-3' |
| (4) 5'-GTATGCCCTAACT-3' | (8) 5'-TGTTAGGGCATACT-3' |
- 4

3C. Illustrate Ramachandran's Plot based on dihedral angles. Also comment on the sterically favourable, partially favourable and disallowed regions. 3

Which graphical representation would you use in order to emphasize the following features of protein structure?

4A. (1) all-atoms (2) bonds and angles (3) atomic volumes (4) backbone (5) secondary structure (6) solvent accessibility (7) van der Waals radii (8) electrostatic potential (9) unit cell (10) ligand 5

4B. List out the steps involved in homology modeling process 3

4C. How do you measure the sensitivity and specificity of a software program? 2

5A. Assume you have eight taxa (A, B, C, D ... H). How many rooted and unrooted trees are possible with the same eight taxa? 2

Construct a cladogram based on six taxa (A to F) using the morphological features given below

Character	A	B	C	D	E	F
Eyes	1	1	1	1	1	1
Tail	1	1	1	0	0	1
Legs	1	1	0	0	0	0
Wings	0	1	0	0	1	1
Fur	0	0	0	0	1	1

It is common knowledge among biologists that mammals and reptiles all shared a common ancestor. The debate is whether (a) birds are part of the mammal clade or (b) whether they are part of the reptilia clade? Justify your argument.

5C.

