STELLA MARIS COLLEGE (AUTONOMOUS) CHENNAI 600 086 (For candidates admitted from the academic year 2004-2005 & thereafter)

SUBJECT CODE: BT/MO/IB64

B.Sc. DEGREE EXAMINATION, APRIL 2010 BRANCH V(a) – PLANT BIOLOGY AND PLANT BIOTECHNOLOGY SIXTH SEMESTER

COURSE	:	MAJOR – OPTIONAL	
PAPER	:	INTRODUCTION TO BIOINFORMATICS	
TIME	:	3 HOURS	MAX. MARKS: 100

Section-A						
I. Cho	oose the correct answ	er		(4×1=4)		
1.	The term gene was co	ined by				
	a) Johansson	b) Mendel	c) Morgan	d) Watson		
2. The actual location of genes on chromosome is determined by mapping						
	a) Chemical	b) genetic	c) physical	d) loop		
3.	Which of the followi	ng displays the result	of BLAST researches t	hat have been done		
	for every protein seq	uence?				
	a) SMART	b) BLINK	c) BEAUTY	d) PROW		
4.	4. In alpha helix, how many amino acid residues are present per turn?					
	a) 4 residues	b) 3.9 residues	c) 2 residues	d) 3.6 residues		
II. State true or false (4						
5. Computer scanning of database can be done by BLAST search.						

- 6. When the backbone of a polypeptide chain is extended, it forms alpha helix.
- 7. DNA sequence of the whole genome of Arabidopsis thaliana is yet to be completed.
- 8. In Drug design, the set of features common to the series of active molecule is known as proteome.

III. Fill in the blanks

- 9. The region of chromatin which is densely packed and stained diffusely during interphase is called ______.
- 10. Fredrick Sangers method is also called chain _____ method.
- 11. The collection of the biological data on a computer appearing as varying arrangements and subsets is regarded as a biological _____.
- 12. A signal molecule is also known as a _____.
- 13. The 3-D structure of pharmacore database is _____.

 $(5 \times 1 = 5)$

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IV. Match the following		(5 × 1 = 5)
14. AUG	a) protein	
15. Macrorestriction map	b) terminator	
16. Tertiary structure	c) gene expression	
17. Markers	d) initiation	
18. DNA microarray	e) high resolution	
	f) SNP	
V. Answer any six the follow	ing, each within 50 words	(6× 3=18)
19. Triplet		
20. cDNA map		
21. FASTA		
22. The primary structure of pr	oteins	
23. Linear space optimization		
24. Chip technology		
25. Gene tagging		
26.Heterochromatin		
	Section –B	
Answer any four questions, e	(4×6=24)	
27. Distinguish between top do		
28. What are the salient feature		
29. Write about protein -motif		
30. Discuss about local alignm	ent and end space alignments.	
31. Write about RNA secondar	y prediction.	
32. Write about applications of	bioinformatics.	
	Section – C	

Section – C

Answer any two questions, each answer within 1200 words (2×20=40)

- 33. Give an account on Gilbert and Maxam method of DNA sequencing.
- 34. Write about various sequence databases. Add a note on mapping databases.
- 35. Discuss the various models for protein folding.
- 36. Give a detailed account of Multiple sequence alignment.

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