Printed Pages: 3			688	NBT502
(F	ollow	ing Paper II	and Roll No Answer Bo	. to be filled in your ok)
Paper ID : 154502			Roll No.	
,			B.Tech.	
		(SEM. V) T	HEORY EXA	M. 2015-16
		BIO-	INFORMATI	CS-I
[Time:3 hours]				[Total Marks: 100]
			Section-A	
1.	Attempt all parts. All parts carry equal marks. Write			
	answer of each part in shor		art in short.	(2x10=20)
	(a)	Define hom	ology.	
	(b)	What is reg	ular expression	1?
	(c)	Define the p	orotein domain	 :
	(d)	Define pair	wise alignment	i .
	(e)	What is Clu	stalW?	
ŧ	(f)	What is PHI	-BLAST?	
	(g)	Define phyl	ogeny.	

(h)

Define alpha helix.

- (i) What is the template structure?
- (j) What is the use of Jmol program?

Section-B

Note: Ateempt any five questions from this section.

(10x5=50)

- 2. Perfrom an optimal global alignment for the DNA sequences GAATTC and GATTA using dynamic programming (scoring: +2 for a match, -I for a mismatch, amd 2 for a linear gap penalty).
- 3. How do PAM and BIOSUM can be used in database similarty serach? Explain.
- 4. What is the role of multiple sequence alignment? Explain the ClustalW program.
- 5. Define the tertiary structure of a protein? Discuss the method of homology modeling.
- 6. Explain the Chou-Fasman method for analysis of protein secondary structrue elements.
- 7. Define RMSD. Discuss the uses of RASMOL program.

(2)

- 8. Explain the rational drug design approach.
- 9. What are the various file formats?

Section-C

Note: Attempt any two questions from this section.

(2x15=30)

- Explain the concept of dendogram. Descuss the UPGMA method for phylogenetic tree construction.
- 11. Write short notes on:
 - i) IEDB
 - ii) EMBL
- 12. What do you mean by protein structure visualization?

 Describe recently developed tools available for visualization and analysis of protein structures.