(37) SARDAR PATEL UNIVERSITY

External Examination

Class-T.Y. B. Sc. VI Semester

Date: - 04-04-2018, Day: - Wednesday

Time: - 10:00 am to 01:00 pm

Course: - US06CBNF05

Subject: Bioinformatics

Title: - Bioinformatics Applications II

Total Marks: 70

Q1.	Multip	Multiple choice questions (All are compulsory).							
•	(1)	Which route of drug administration is most likely to lead to the first-pass effect?							
	• •	a) Sublingual h) Oral c) Intravenous d) intramuscular							
	(2)	What is the term used for the automated in vitro testing of large numbers of compounds using							
		genetically modified cells?							
		a) robotic testing b) high throughput screening							
		c) multiscreening d) nanotechnology							
	(3)	How many codons are needed to specify five amino acids?							
		a) 3 b) 6 c) 15 d) 21							
r	(4)	What secondary structures are formed when single-stranded DNA or RNA is inverted and							
		complementary? A Adouble boliv b) B-DNA c) Z-DNA d) hairpin							
		A Juddie Helix							
	(5)	A bifurcating branch point in the phylogenetic tree is known as a) pade b) clade c) branch d) taxon							
		a) flode							
	(6)	Expand UPGMA.							
		a) Unweighted Pair Group Method with All Moan							
		b) Unweighted Pair Group Method with All Mean. c) Upregulated Gene Method with Arithmetic Mean.							
		d) Unregulated Genome Method with All Mean.							
	171	In standard microarrays, the probes are attached via surface engineering to a solid surface by							
	(7)	a to a chemical matrix	•						
		a) Metallic bond b) Chemical bond c) Covalent bond d) Aromaticity							
	(8)	Microarray data processing using							
	(0)	a) Generative topographic map b) Artificial intelligence							
		c) Artificial neural network d) Self-organizing map							
	(9)	Microarray analysis can be used to:							
	1-7	a) Determine the intron-exon organization of a gene.							
		b) Determine the concentration of a protein in a cell.							
		c) Determine the stage-specific expression of a gene							
		d)Determine the presence of a DNA sequence in a cell							
•	(10)	Species evolve by							
		a) diversification b) progression c) linear advancement d) magic	[20						
Q2.	Ansv	wer the following questions in short. (Attempt any 10) (each carry 2 marks)	[20						
	(1)	Differentiate average linkage and complete linkage.							
	(2)	How image processing occur in microarray.							
	(3)	and the state of t							
	(4)	Condition 1 to 1							
	, ,	and the state of t							
	(5)	s. A first the leader-constinue and CADD	(P.T						
	(6)	Ethist and two software a for pont budioBeneries arranges and a real	~						

	(7) V	(7) What is a basic structure of phylogenetic tree?											
(8) Explain preclinical and clinical trial in CADD.(9) Define Combinatorial Chemistry and docking.													
	, ,												
	(11) Differentiate between DNA and RNA structure.												
	(12) G	live the	e appl	icatio	ns and	d limit	ations	s of I	Afold method.				
Q3.	What is	micro	array î	Discu	uss its	meth	ods a	nd a	oplications in detail.	[10]			
	OR									[10]			
Q3.	Discuss the clustering method and its types.												
<u>Q</u> 4	What is a phylogenetic tree? Discuss its important methods and applications.												
									OR				
Q4.	Construct the tree using UPGMA method with following matrix table												
		А	В	С	D	Е	F	G					
	A												
	В	19.00	24.00	-				 					
	D	27.00 8.00	31.00 18.00	26.00			 		•				
	E	33.00	36.00	41.00	31.00								
	F	18.00	1.00	32,00	17.00	35.00		<u> </u>					
	G	13.00	13.00	29.00	14.00	28.00	12.00						
Q5.	Define (drug aı	nd its	basic '	featui	es. E	cplain	the:	steps for CADD	[10]			
									OR				
Q5.	Write a	short	note (on foll	owing	ξ;				[10]			
	i) ADMI	E ii) k	nocko	ut Ge	ne iii)	Lipin	ski's r	ule.					
										[5]			
Q6.	i) Explain M-fold method and its utility.												
	ii) Expla	in tert	iary s	tructu	re of	RNA.				[5]			
									OR				
Q6.	Write a									[10]			
	i) A, B and Z type of DNA ii) tRNA												
		·											