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SARDAR PATEL UNIVERSITY

M.Sc. Biotechnology (SEMESTER-II)

Paper: PS02CBIT03- Genetic Engineering and Bioinformatics Saturday, 15th April, 2017. 10.00 a.m to 01.00 p.m

Marks: 7<u>0</u>

		(O 1)
 Q1. Choose the most appropriate answer: i) Which of the following is NOT true of type II rests a) Modification by methylation within the sequence b) They are symmetrical made of 4-8 bases c) They signal the attachment of RNA polymers d) They are often palindromic 	riction enzymes? ence prevents restriction	(8 marks)
 ii) A thionucleotide is used to improve site directed in a) allows the restriction enzymes to cleave the b) it protects the mutant DNA strand from become it enhances the binding of DNA polymera d) it reduces the non specific binding of primary 	ne mutant DNA strand bing cleaved by a restriction en se	ızyme
 iii) Eukaryotic genes may not be expressed properly a) Lack of intron splicing mechanisms in bacte b) Destruction of the cloned DNA by native en c) Promoters not being recognized by bacterial d) All of the above 	ria donucleases	cause of
iv) The modified nucleotide used in Sanger's method	d of sequencing known is as _	
a) Dideoxy ribonucleotide c) met	hyl Adenine xy ribonucleotide	
v) Which of the following is not a protein database a) PIR b) SWISS PROT	c) DDBJ d) '	TrEMBL
vi) Which of the following is not a gene prediction a) Chou-Fasman method b) Ab initio method	method c) Homology Method d) Comparative method	
vii) Which of the following is/are desirable characta) Event should occur in deep timeb) Gene that changes very slowly	eristic of phylogenetic markers c) Gene should be highly cor d) All of the above	s nserved
viii) What is meant by a lead compound in drug di a) A drug containing the element lead	scovery?	
b) A leading drug in a particular area of m	edicine	-
c) A compound that acts as the starting point d) A drug which is normally the first to be	nt for drug design and develop	ment ient

Q2. Ar	nswer any seven of the following questions in brief:	(14 marks)
2.	Homopolymer tailing Blue-white screening	
3.	Dideoxynucleotide	
4.	Taqman probe	
5.	Composite database	
6.	Dot Plot	
	ExPASy	
8.	CATH	
9.	3D Viewing software	
3.	a) Explain the principles and most common methods for the removal of protein from DNA samples.	is 06
	b) Discuss the principle and advantages of modified PCR based method for site directed mutagenesis	06
	OR	
	b) Write a note on the reaction parameters and significance of ligation in DNA cloning	06
4.	a) Discuss the principle and procedure for the Sanger's method for DNA seque Explain how this method is better than Maxam-Gilbert's method.	encing. 06
	b) Outline the principle of Taqman probe method for real time PCR. What	
	are the advantages and limitations of this method?	06
	OR	
	b) Write notes on i) SCAR markers ii) AFLP	06
5.	a) Enlist various databases and explain any two in details.	06
-,	b) Explain various BLAST and their working in detail. OR	06
	b) Write notes on multiple sequence alignment (MSA)? Discuss how it is help to find out evolutionary relationship among organisms.	ful 06
6	6. a) Discuss any two methods for the protein secondary structure prediction. 06 b) Discuss the mechanism of molecular phylogeny. Explain different types of	
	phylogenic tree.	06
	OR	1
	b) What are the different stages in drug discovery? Explain how bioinformatics significant role in drug discovery.	plays a
	XXXXXXXXXX	

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Sardar Patel University M Sc II Semester Microbiology/Biotechnology PS02CMIC01/BIT01 Bioprocess and Biochemical Engineering External Theory Examination

	: 10 th April, 2017 : 10:00 am to 1:00 pm		Max Marks : 70
Q. 1	Choose the correct answer:		(08)
1.	The cyclone column reactor a) hydrodynamic system c) pneumatic system	b) air lift reactor	
2.	The tube and shell are a typa) bioreactors c) filters	be of b) heat exchangers d) stirrer seals	S
3.	Lyophilizers are used in a) sterilization c) Cell disruption	b) culture preser d) Cell separation	
4.	Which of these is a nitroger a) Molasses c) Corn steep liquor	b) Corn starch d) vegetable oils	and the second s
5.	θm is associated with a) agitation aeration c) cell disruption	b) solvent extraction	on
6`.	PT-100 is a a) biosensor c) controller	b) filter d) none of these	
7.	The mixing time is symboliz a) Del factor c) μ	zed as b) K _L a d) θm	
8.	Dynamic gassing out technia) K_La c) air flow rate	ique is used for determi b) mixing time d) filtration rate	nation of

Q. 2	Explain the terms in brief: (any seven)	(14)
	 a) Feedback systems b) Supercritical fluid extraction c) Imperfectly mixed bioreactors d) Scale up and scale down e) Mass transfer f) Dcrit g) Pneumatic systems h) Feed forward loop i) Holding time 	
Q. 3	A) Explain secondary screeningB) Write a note (Any one)a) Carbon sourcesb) Industrially important organism	(06) (06)
Q. 4	A) Explain the kinetics of a batch sterilization process B) Discuss the design of a fixed laboratory scale fermentor OR	(06) (06)
	B) Discuss with a diagram the design pressure cycle reactor	(06)
Q. 5	A) Explain microprocessor based control systems B) Write a note on (any one) i) PID controllers ii) Fedbatch cultivation	(06) (06)
Q. 6	A) Discuss downstream processing strategies B) Write a note on (any one) i) Cross flow filtration ii) Cell dusruption	(06) (06)
	* * * * * * * * *	

(2)

SARDAR PATEL UNIVERSITY

M.Sc (II Semester) Biotechnology Examination (CBCS) Wednesday, 19thApril, 2017

Wednesday, 19thApril, 2017 Time - 10:00 am to 1:00 pm PS02EBIT02 –Toxicology

TOTAL MARKS: 70

•			
Q.1 Tick mark / select the correct a number needs to be written in provi	nswer for the following ided answer book)	. (Only correct option	against given question (08 Marks)
 Cycasin (methyl azoxy method) (a) Nasal route Which of the following toxi 	(b) Dermal route	(c) Oral route	it is exposed by (d) any route
(a) Cyanide3. If the toxic response of the effect is known as	(b) Botulinum toxin two toxicants is greater	(c) caffeine than the individual re	(d) Paraquat sponse than this
4. The toxic effect of a toxican	etentiation or Synergism t is affected by	-	(d) None
(a) Dose of toxicant (b) from	equency of exposure	(c) route of exposure	(d) all of the above
5. The larger the	, the better the	chemotherapeutic effec	et ·
(a) therapeutic index	(c) toxic dose		
(b) therapeutic dose	(d) selective toxicity		
6. Benzoic acid, a food additive,(a) Preservative(b) Emulsifier	is used as (c) Flavouring agent (d) antioxidant		
7. All are greenhouse gases exce	pt:		
(a) Methane	(c) Ozone		
(b) Nitrous oxide	(d) Nitrogen		
8. Herbicides 2, 4 D and 2, 4 T a	re		
(a) Photosynthetic inhibitors	(c) Respiratory inhibite	ors	
(b) Growth stimulators	(d) Growth inhibitors		
	MI		

Q.2 Answer any seven from the following:

(14 marks)

- 1) Name the antidote used for toxicity caused by N-acetylbenzoquinoneimine, a phase I product of paracetamol. 2) Is cytochrome P450 enzyme polymporphic? Discuss in brief. 3) Where does phase II metabolsim take place?
- 4) Enumerate the list of enzymes in phase I reactions involving xenobiotic metabolism?
- 5) Define the term rancidity. Which broad category of food additives are used to prevent rancidity?
- 6) Differentiate between acute and chronic toxicity.
- 7) What are the advantages of using Drosophila as a test organism in toxicology studies?
- 8) What is 'risk assessment' with reference to toxicology?
- 9) What is the importance of structure-activity relationship in pharmacodynamics?
- Q.3 A:Draw a dose response curve and show NOEL, LD50 and maximum toxicity levels in the graph. (6 marks)
- Q.3 B: Explain the role of Cytochrome P450 enzymes in metabolism of toxicants. (6 marks)

OR

Q.3 B: Differentiate between

(6 marks)

- (i) Pharmacokinetics and Pharmacodynamics
- (ii) Synergism and Antagonism.
- Q.4 A: Explain the bacteria reversed mutation assay (Ames test) to detect mutagenic properties of test chemicals. (6 marks)
- Q.4 B: Give examples of any six toxicants with their toxic effects.

(6 marks)

Q.4 B:What is fluctuation test? What is it used for?

(6 marks)

- Q.5 A:Write short notes on: (6 marks)
 - 1. General measures for management of poisoned patients
 - 2. Methanol toxicity
- Q.5 B:Explain the toxicity of organophosphorous insectides with suitable examples. (6 marks)
- Q.5 B: Write a note on various types of insecticides.

(6 marks)

Q.6 A: Explain the metabolism and chronic toxicity of paracetamol.

(6 marks)

Q.6 8: What is Therapeutic Drug Monitoring (TDM)? What are the major criteria for valid therapeutic drug monitoring? OR

(6 marks)

Q.6 B: Explain any two manifestations of plumbism.

(6 marks)

- (B) What is reaction wood? What are its salient features? What is it significance?
- (C) Giving any two suitable examples to justify how ethnobotany differs from economic botany.
- (D) How does heart wood differs from sapwood? Which of these two more economically important? Why?
- (E) Name any two Indian Ethnobotanists? What is their major contribution?
- (F) What are bordered pits? What is their significance?
- (G) List any two plant conservation centres of national importance. Where are they located?
- (H) Give botanical names of any four gum yielding plants.
- (I) What are botanical pesticides? What are their advantages? Give botanical names of two sources of such pesticides.

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Q3.	(A) "Define Ethnobotany. Why is that it is said to be multidisciplinary subject? Justify your answer with reasons.	(6)
	(B) Give a comprehensive account on ethnomedicobotanical data collection.	(C)
	OR OR	(6)
	(B) (i) The concept of Sacred Groves is a proved lesson on phytorsource conservation. Justify the statement.	(3)
	(ii) What is the botanical source of Indian saffron? What are its uses?	(3)
Q4		(6)
	(B) "We obtain around 95% of our daily requirement of energy from a wide diversity of phytoresoures available to us. Do you agree with the statement? Justify your answer in either the case giving suitable examples.	(6)
	OR	
	(B) Name any four little known phytoresources, having scope for wider usage for their merits. Offer your innovative ideas for making them popular.	(6)
Q5	(A) "Describe the origin, cultivation, useful products and uses of any two oil yielding crops studied by you.	(6)
	(B) What are the important criteria used for determination of the botanical source and quality of wood?	(6)
	OR WHITE WAR WAS A STATE OF THE	
	(B) Write notes on:	
	(i) Fiber yielding plants	(3)
	(ii) Gene banks	(3)
26	(A) Write short notes on:	(0)
	(i) Role of Botanical gardens in conservation of threatened phytoresources.	(3)
	(ii) Major threats to agribiodiversity	(3)
	(B) Justify any two of the following statements with suitable examples:	(0)
	(i) "Many of the phytoresources can be potential alternatives for conservation of fossil fuels.	(3)
	(ii) "Palms and Fruit yielding trees are not good choice for plantations along highways"	(3)
	(iii) "Traditional knowledge on phytoresources is more threatened than the phytoresources."	(3)