Paper / Subject Code: 66107 / Biochemistry III

Time: 2 hrs

N.B.: All Questions are compulsory.	Total Marks: 40
Q1. (a) Name two pyrimidine bases present in RNA.	(01)
(b) Name two drugs which inhibit topoisomerase.	\$\tag{01}
(c) Give role of spliceosomes in RNA splicing.	£ 6 6 7 7 7 8 7 (01)
(d) Name the initiation codon and its respective amino	acid. (01)
(e) Explain how Chloramphenicol and Streptomycin dr	ugs inhibit protein synthesis.(02
(f) Enlist all types of eukaryotic DNA polymerases.	(02)
Q2. (a) Discuss transcription in prokaryotes.	(03)
OR	
(a) Give details of promoter regions for initiation of eul	caryotic transcription. (03)
(b) Describe lac-operon.	(03)
(c) Explain Base excision process for DNA repair.	(02)
Q3. (a) Explain Holliday model for recombination.	(03)
(b) Describe Elongation stage of protein synthesis in ba OR	cterial cells. (03)
(b) Differentiate Initiation of translation between proka	ryotic and eukaryotic cells with
description of the steps for Initiation.	(03)
(c) With a schematic diagram depict Watson-Crick mod	del for DNA structure. (02)
Q4. (a) Write note on telomere and telomerase.	(03)
(b) Explain post-translational modifications.	(03)
# # # P P P P P P P P P P P P P P P P P	
(b) Compare solid phase peptide synthesis with biosynt	hesis. (03)
(c) What are mutagens? Give an example of chemical a	nd biological mutagens. (02)
Q5. (a) Explain the terms: Unidirectional and bidirection OR	onal mode of replication. (02)
(a) Give justification for the statement: 'DNA polymera	ase III is the principal replication
enzyme in E. coli while DNA polymerase II is involved	l in DNA repair'. (02)
(b) Discuss SOS DNA repair	(02)
(c) Give the post-transcriptional modifications of mRN	A. (02)
(d) Write a note on Single Nucleotide Polymorphisms.	(02)
37 V5 V5 V5 O V5 32 V5 V5 V7 V7 V6 V5 V6 V5 V6	

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[2 Hours]

Please check whether you have got the right question paper.

1. All questions are compulsory

N.B:

[Total Marks: 40]

	2. Figures to the right indicate full marks.				
Q.1	Answer the following: a) Give the regulatory requirements for manufacture of eye makeup products. b) What are the factors governing formulation of baby products? c) Enlist Nail speciality products d) What are pearlescent pigments? Give examples. e) What is the mechanism of working of hair conditioners? f) Distinguish between antiperspirants and deodorants.	(10)			
Q.2	Answer any two a) Vanishing creams are pearlescent and have a slightly acidic pH . Justify . b) Elaborate on colours used in cosmetics c) Describe any two quality control tests on nail lacquers. 	(6)			
Q.3	Answer any two a) Describe the packaging of lipsticks. b) Write a note on insect repellants. c) Classify surfactants used in shampoos with examples	(6)			
Q.4	Answer any two a) Write on large scale manufacturing of loose face powder. b) Enlist various types of shaving products and elaborate on any one. c) Give an account of raw materials used in toothpastes. 	(6)			
Q.5	a) Elaborate on methods used for sensorial evaluation of cosmeticsb) Write 2 in-vitro tests for skin irritation. Discuss sensitivity testing of cosmetic products done on animals.	(3)			
Q.6	a) Enlists various methods of processing of plant materials used in cosmetics. Ex any one process .b) Illustrate the significance of sensorial evaluation for cosmetics.c) Give the classification of different depilatory agents.	plain (2) (2) (2)			

Paper / Subject Code: 66106 / Microbial Genetics

(2 Hours) Marks: 40 NB: (1) All questions are Compulsory. (2) Draw neat labelled diagrams wherever necessary Q.1 2 How transposition can cause mutations in cells? 2 b Explain the two types of mobile genetic elements present in bacteria. 2 What is the importance of horizontal transfer of genetic material in bacteria? 2 What is specialized transduction? d Outline the steps for natural transformation in Bacillus subtilis. 2 е Q.2 Distinguish between Prototrophs and auxotrophs. 2 2 b How are F' factors generated? How does a geneticist doing interrupted mating experiments know that 2 the locus for the drug sensitive allele, used to eliminate the Hfr bacteria after conjugation, has crossed into the F- strain? Q.3 In a transduction experiment, the donor was g⁺ e⁺ f⁺ and the recipient g⁻ 3 e f. selection was for g. The four classes of transductants from this experiment are given below: **Number of Genetic composition** Class individuals g+ e+ f+ 1 57 g+ e+ f 2 76 g⁺ e⁻ f⁻ 3 365 4 2 g+ e-f+ Total 500 a) Determine the cotransduction frequency for g⁺ and e⁺. b) Determine the cotransduction frequency for g⁺ and f⁺. c) Which of the cotransduction frequencies calculated in (a) and (b) represents the greater actual distance between the genes? Why? b Explain the role of RuvABC complexin resolution of Holliday junction. 3 Explain the Holliday model for homologous recombination Q.4 What are plasmids? Explain briefly any 2 types 3 3 Write a short note on plasmid incompatibility. Or

Paper / Subject Code: 66106 / Microbial Genetics

		write the principle and method of isolation of plasmids.			
Q.5	а	What is artificial transformation in bacter same.	ia. Explain two methods for the		
	b	Write a short note on composite transpos	ons.		
		Or			
		Integrons are major mechanism for spread Explain why?	d of multidrug resistance.		
Q.6	а	What is catabolite repression?			
	b	on by attenuation			
		 i. Indicate whether for the following permease will be produced or not inducer: 			
		Inducer absent	Inducer present		
		β-galactosidase permease a) I ⁺ P ⁺ O ^c Z ⁺ Y ⁻	3-galactosidase permease		
		b) I-P+O+Z+Y-	2		
		nstitutive gene expression and			
			100 10 10 10 10 10 10 10 10 10 10 10 10		

[Total Marks: 40] (2 Hours) N.B: (1) All the questions are **compulsory**. (2) Figures to the right indicate full marks. (3) Draw neat labelled sketches wherever necessary Q1. a. State what is meant by – primary packaging and secondary packaging. Give two examples of each 1Mb. State any two defects observed in glass containers. 1M c. Explain the issues related to biocompatibility of plastics 2Md. Enlist various types of aerosol valves. 1**M** e. Mention/State basic components of Blister Packs. 1M f. Enlist types of labels 1M g. Give one example of child resistant cap. 1Mh. State any two Q.C. tests for paper and paper board. 2MQ2. a. Discuss the different types of glass w.r.t. chemical composition. 3M b. Discuss sterilization of plastic containers. 3M OR **b.** Elaborate on sterilization of rubber closures. 3M Q3. a. Explain in detail fabrication processes for plastics. 3M b. Describe in detail printing processes of labels 3M Q4. a. Elaborate on cans and collapsible tubes for pharmaceuticals. 3M b. Write a note on Packaging and the ICH guidelines 3M OR b) Elaborate on photostability testing 3M Q5. a. Elaborate on coatings and laminates 3M b. Describe Strip packaging process w.r.t. materials used and machinery. 3M b) Elaborate on shrink wrapping and stretch wrapping processes in packaging. 3M Q6. a. Give the functions of a package. 2Mb. Elaborate on ancillary packaging materials. 2Mc. Write on elastomeric closures for parenteral products. 2M

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		[Time: 2 Hours]	[Marks:40]
		Please check whether you have got the right question paper. N.B: 1. All questions are compulsory.	
Q.1		Answer the following:	12
	i)	What are functional foods? Give examples.	
	ii)	Give limitations of nutraceuticals (any 2)	
	iii)	Give the source & structure of Resveratrol.	
	iv)	Name one marketed preparation and the therapeutic uses of Curcumin.	
	v)	State the role of tocopherol as nutraceuticals.	
	vi)	Write the occurrence and use of Glutathione.	
	vii)	Give the structure and recommended does of Astaxanthin.	
	viii)	Name any 2 processing challenges for liquid oral dosage forms.	
	ix)	Give one examples of food-nutraceutical interaction.	5
	x)	Give any one adverse effect of nutraceutical with suitable examples.	
	xi)	What does FPO stand for?	
	xii)	Write a pharmacoepial specification for nutraceuticals.	
Q.2	i)	Write a note on nutrition and aging.	04
	ii)	Write a note on labeling of nutraceuticals.	03
Q.3	i)	Discuss adulteration of nutraceuticals.	04
	ii)	Discuss marine nutraceuticals.	03
Q.4	i) (Comment on AGMARK regulatory aspects for nutraceuticals.	04
a constant	ii)	Write a note on flax Lignans.	03
Q.5	i)	Discuss the challenges involved in extraction and concentration of nutraceutica constituents.	al 04
	ii)	Write a note on the occurrence and role of prebiotics and probiotics.	03

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Time: 2 Hours Marks: 40 N.B.: 1. All Questions are compulsory

2. Figures to right indicate full marks

Q.1. Answer the following in brief. Draw structures wherever required.
i. Define: a. Functional group interconversion b. Synthon
ii. Suggest suitable synthetic equivalents for the following synthons
2M

iii. Identify synthon A and B in the given retrosynthesis reaction

iv. Discuss any two disconnection rules by giving suitable example
v. Identify A and B in the given heterocyclic retrosynthesis

2M

$$\begin{array}{c}
O \\
HN^{\frac{1}{2}} \\
O \\
H
\end{array}$$
C-N disconnection
$$A + B$$

Q.2. i. Identify the following conversion as Functional Group Interconversion (FGI) or Functional Group Addition (FGA) or Functional Group Removal (FGR) and give suitable justification3M

ii. Suggest the economical and simple retrosynthetic pathway and synthetic scheme for

4-methoxyacetophenone **OR** Ethyl 1,4-diphenylpiperidin-2,6-dione-3-carboxylate

3M

iii. Complete the following retrosynthetic pathway

4M

4M

3M

2M

Q.3. i. Write the "Order of events" for retrosynthesis of m-nitrochlorobenzene 3M

ii. Write the retrosynthesis of furan **OR** N-methylpyrrole using "1+4" strategy. 3M

iii. Discuss the retrosynthetic pathway for Benzocaine **OR** Sulfadiazine

Q.4. i. Suggest a suitable retrosynthetic scheme for any one of the following compounds

ii. Discuss the term "Umpolung" using suitable example

iii. Discuss the retrosynthesis of Ibuprofen **OR** Propranolol **4M**