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## **Enrolment No:**



## **UPES**

## **End Semester Examination, May 2025**

Course: Microbial Genetics Semester : VI

Program: Int. BSc.MSc. Microbiology Duration : 3 hours

Course Code: HSMB 3017 Max. Marks: 100

## **Instructions:**

S. No.	Section A	Marks	Cos	
	Short answer questions/ MCQ/T&F			
	(20Qx1.5M = 30 Marks)			
Q 1	Generalized transduction only transfers:	1.5	CO2	
	a) Random DNA			
	b) Phage DNA			
	c) Adjacent host genes			
	d) Entire chromosome			
Q 2	Identify which gene controls competence in <i>B. subtilis</i> .	1.5	CO2	
	a) com			
	b) lexA			
	c) comp			
	d) araC			
Q 3	Identify filamentous phage from the following.	1.5	CO2	
	a) T4			
	b) T7			
	c) M13			
	d) Lambda			
Q 4	Recall, which of these is a chemical mutagen.	1.5	CO1	
	a) UV radiation			
	b) X-rays			
	c) EMS			
	d) Heat			
Q 5	Transduction requires:	1.5	CO1	
	a) Pilus			
	b) Bacteriophage			
	c) Plasmid			
	d) Sigma factor			
Q 6	Two-factor crosses in phages are used to:	1.5	CO1	
	a) Determine operons			
	b) Create lysogens			
	c) Map genes			
	d) Observe transformation			
Q 7	The main function of antitermination in T4 phage is:	1.5	CO2	

	a) Prevent replication		
	b) Promote recombination		
	c) Continue transcription		
	d) Halt translation		
Q 8	Pick the correct answer. Which of the following follows a lytic life	1.5	CO2
	cycle?		
	a) Lambda phage		
	b) T4 phage		
	c) P2 phage		
	d) P4 phage		
Q 9	OriT refers to:	1.5	CO2
	a) Operator site		
	b) Origin of transcription		
	c) Origin of transfer		
	d) Outer region		
Q 10	Recombination tests help identify:	1.5	CO1
-	a) Gene expression levels		
	b) Regulatory proteins		
	c) Genetic linkage		
	d) DNA replication errors		
Q11	Spot, which mutation type is caused by tautomeric shifts.	1.5	CO2
	a) Frameshift		
	b) Transition		
	c) Deletion		
	d) Insertion		
Q12	Identify, which organism is naturally competent.	1.5	CO2
	a) E. coli		
	b) B. subtilis		
	c) Salmonella typhi		
	d) Mycobacterium leprae		
Q13	The lac operon is regulated by:	1.5	CO2
	a) Only lactose		
	b) Only glucose		
	c) Both positive and negative mechanisms		
	d) Phage integration		
Q14	Plasmid integration into the chromosome forms:	1.5	CO2
	a) Hfr strain		
	b) F- cell		
	c) Prime factor		
	d) Transposon		
Q15	The 'jumping genes' discovered by Barbara McClintock are known	1.5	CO1
	as:		
	a) Operons		
	b) Episomes		
	c) Transposons d) Plasmids		
Q16	Marker rescue is a technique used for:	1.5	CO1
<b>410</b>	a) Mutant enrichment	1.0	

	b) Gene mapping						
	c) Cloning based on functional restoration						
	d) Complementation testing						
Q17							
	a) Reversions		CO1				
	b) Intragenic or intergenic						
	c) Dominant						
	d) Recessive only	only					
Q18							
Q10	a) Generalized transduction	1.0	CO1				
	b) Specialized transduction						
	c) Transformation						
	d) Lysis only						
Q19	Identify, which method allows mapping of bacterial genes based on	1.5	CO1				
Q19	time of entry during conjugation.	1.3	COI				
	a) Transformation						
	b) Generalized transduction						
	c) Hfr conjugation						
	d) Site-specific recombination						
020		1.5	CO2				
Q20	Spot which system is an example of site-specific recombination	1.5	COZ				
	commonly used in genetic engineering.						
	a) lac operon						
	b) T4 phage lytic system						
	c) loxP-Cre system						
	d) SOS repair system  Section B						
	(4Qx5M=20 Marks)						
Q 1	Explain the role of base analogues in mutagenesis with example of	5	CO2				
Q I	one agent.	J	002				
Q 2	Distinguish between base excision repair and mismatch repair.	5	CO1				
	Differentiate between positive and negative selection of mutants in		CO2				
Q3	microbial genetics.	5	CO2				
Q 4	In case of lac operon the levels of structural proteins differs in ratio	5	CO1				
Q4	of 1:0.5:2 as you move from beginning to end of operon. Reason	3	COI				
	why.  Section C						
	Section C (2Qx15M=30 Marks)						
Q 1	Experiments using rII mutants of T4 phage were instrumental in	15	CO2				
<b>V</b> I	deciphering the genetic code. In one such experiment, a double	10	002				
	mutant was used to study intragenic recombination.						
	industr was used to study intrageme recombination.						
	a) Name the scientist who worked on rII locus? (1)						
	b) Spot the significance of rII mutants in phage genetics? (3)						
	c) Describe how complementation and recombination tests were						
	used in this context. (6)						
	d) Describe how complementation groups are created using the example below where zero means no complementation while _						

	Strain	1	2	3	4	5	6		
	1	0	0	+	0	0	0		
	2		0	+	+	0	0		
	3			0	+	+	0		
	4				0	+	+		
	5					0	+		
	6						0		
Q 2	A cancer arose i	in a heal	thy pers	son with r	no genet	ic or envi	ronmental	15	CO2
	A cancer arose in a healthy person with no genetic or environmental history. The doctor diagnosed a jumping gene which integrated into a tumor suppressor gene to be responsible for it. With your knowledge of microbial genetics and molecular biology, answer the following:  a. Define jumping genes. (1)  b. Name the scientist who discovered jumping genes and in which organism? (2)  c. Elaborate if there are different kinds/types of these jumping genes. Classify them. (5)  d. Explain, how do these genes cause mutations? (3)  e. Discuss the role of jumping genes in mapping. (4)								
				(2Qx10N					
Q1	<ul> <li>A student performs an interrupted mating experiment using an Hfr strain and a recipient F− strain. The order of gene transfer observed is: leu → thr → his → lac → gal.</li> <li>a) Explain the principle of interrupted mating and how it is used to map bacterial genes. (5)</li> <li>b) Name the scientists who used it first? (1)</li> <li>c) Based on the gene transfer order, construct a partial map of the</li> </ul>						10	CO1	
Q 2	a. Elaborate transcription highlighted b. Illustrate the	the onal reg d. (7)	life cy	and cho	oice of l		lysogeny	10	CO1