	(3 Hours)	(Total Marks: 100)
	Instructions to the candidates:-	
	1) All the questions are compulsory. Choice is internal.	
	2) Figures to the right indicate full marks .	
	3) All questions carry equal marks.	
	4) Draw flowcharts /diagrams wherever necessary .	
01.4)	Fill in the blanks:	
Q1 A)		
i)	The site for DNA replication is	
ii)	Thymine dimers occur due to light	
iii)	The enzyme used to join bits of DNA is	
iv)	Okazaki fragments are present on strand.	
Q1 B)	Write a note on (any one):	4
i)	DNA Polymerases	
ii)	SOS repair	
ŕ		
Q1 C)	Answer the following: (any two)	12
i)	Discuss the various proteins required to be synthesised in a cell to en	nable it to
	multiply.	\$\frac{1}{2}\tag{2}
ii)	Elaborate on the Excision repair mechanism.	
iii)	Discuss the structural mutations that can occur during or after replic	ation.
Q2 A)	Fill in the blanks:	4
i)	Post-transcriptional modification occurs in	
ii)	is a stop codon.	
iii)	The synthesis of polynucleotide chain of mRNA is catalysed by the	enzyme
1V)	Protein synthesis in bacteria takes place on/in	
88		
J. 00 1 1/2	Write a note on (any one):	4
(1)	Genetic code	
ii)		40
Q2 C)		12
(i)	Write a note on (i) inhibitors of mechanisms of central dogma of n	noiecular biology
	(ii) post-translational modifications.	lon
(ii)	Schematically ONLY represent the process of elongation in translati	IOII
ni)	Explain the process of initiation of protein synthesis	

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Q3 A)	Fill in the blanks:	4	
i)	EcoR1 cleaves DNA at		
ii)	In pUC vectors, if the gene of interest is inserted in gene		
iii)	Restriction endonucleases isolated from unidentified micro-organisms is named as		
iv)	chemical is used to increase the copy number of a plasmid.		
Q3 B)	Attempt the following: (any one)		
i)	Diagrammatically represent and explain the process of cloning a foreign gene into a plasmid cloning vector.		
ii)	Create a note on genetically modified food.		
Q3 C)	Answer the following: (any two)		
i)	Write a short note on probe. Briefly explain about labelling of probe and its applications.		
ii)	Elaborate on 'Molecular Scissors' and 'Molecular Stitchers.'		
iii)			
Q4 A)	Fill in the blanks:		
i)	At temperature denaturation of DNA double helix takes place in PCR.		
ii)	library only involves expressible genes.		
iii)			
iv)	chemical enhances transformation efficiency developed the polymerase chain reaction.		
Q4 B)	Attempt the following: (any one)		
i)	Highlight on the contribution of E.M. Southern to the field of recombinant DNA technology.	4	
ii)	Discuss the (a) experimental set-up and (b) advantages of a chemical method of gene transfer.		
Q4 C)	Answer the following: (any two)		
i)	Explain selection of recombinant cells by the use of antibiotic-resistance gene.		
ii)	Elaborate on a technique of DNA amplification.		
iii)	In a stepwise manner, explain the formation of a DNA library. Also, state the		
89 67 K	difference between a cDNA library and genomic library.		
Q5 A)	Define and explain:	8	
a)	Okazaki Fragments		
	OR		
b)	Aneuploidy		
c)	Shine-Dalgarno sequence		
	OR		
d)	TATA box		
e)	1 unit of restriction endonuclease		
	OR		
(f)	R plasmid		

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g) Colony hybridization

OR

h) Transformation

Q5 B) State True or False with justification:

- i) Hind III is a blunt-end cutter.
- ii) Reverse transcriptase has a DNA dependent DNA polymerase activity.
- iii) Lac selection method is also known as blue-white colony method.
- iv) All repair mechanisms are error free
- v) Meselson and Stahl proved that translation is by semi-conservative process
- vi) SSB are a requirement in repair

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