1×10=10

## NRT/KS/19/2162

## Bachelor of Science (B.Sc.) Semester–V Examination MOLECULAR BIOLOGY AND rDNA TECHNOLOGY

## **Optional Paper-2**

## (Biotechnology)

Time: Three I	Hours] [Maximum Marks	: 50
<b>N.B.</b> :— (1)	All questions are compulsory and carry equal marks.	
(2)	Draw diagrams wherever necessary.	
1. Describe	e in detail the attachment of amino acids to tRNA.	10
	OR	
Describe	e in detail how the genetic code was deciphered.	10
2. Describe	e the initiation process of prokaryotic protein biosynthesis.	10
	OR	
(a) Des	scribe the role of release factors in prokaryotic translation.	5
(b) Dese	scribe the role of antibiotics affecting translation process.	5
3. Describe cells.	e the technique of transformation and transfection. Add a note on selection of transfor	med
	OR	
(a) Des	scribe briefly the PUC series of vectors.	5
	scribe briefly the restriction endonucleases.	5
	e in detail the applications of rDNA technology in medicine and agriculture.	10
	OR	
Write sho	nort notes on :	
(a) Exp	pression vectors	21/2
• • • • •	mer designing	21/2
(c) cDN	NA library	21/2
(d) Step	ps in PCR technique	21/2
5. Solve any	ny <b>ten</b> of the following:	
(I) To	which end of tRNA, the amino acid is attached?	
(II) Wl	Tho proposed Wobble hypothesis ?	
(III) Gi	ive any one role of Shine-Dalgarno sequence.	
(IV) Na	ame any one elongation factors used in protein biosynthesis.	
(V) Na	ame the factor which separates the large and small subunit of ribosomes	
(VI) WI	That is meant by autogenous control?	
(VII) WI	That is meant by PBR322 ?	
(VIII) WI	hat is meant by "EcoRI"?	
(IX) Na	ame the enzyme efficient in blunt-end ligation.	
(X) Gi	ive any one advantage of cDNA library over genomic library.	
(XI) Na	ame any one rDNA product used in the field of medicine.	
(XII) W	Thy two primers which are 90% complimentary to each other cannot be used as prime	rs in

the PCR technique? Give any one reason.