(i)	Print	ed Pages: 2	Roll No	•••••••
(ii)	Ques	tions : 9	Sub. Co	
		*	Exam. Co	ode: 0 0 3 8
		B.Sc. (Hons.)	Biotechnology	6 th Semester
			(2054)	
		GENE'	TIC ENGINEER	RING
		Pa	per : BIOT-601-7	\mathbf{T}
Time	[Maximum Marks: 67			
Note		Attempt five que Attempt one que mpt the following	uestion from each	stion No. 1 is compulsory. Unit.
	(a)	What are Links	ers and Adaptors (?
	(b)		oolymeric Tailing?	
	(c)	What are Rever	se transcriptases?	?
	(d)	Define Insertiona	al inactivation.	2
	(e)	What is the r constructs?	ole of antibiotic	resistance in Vector
	(f)	What are doubl	e digests and partia	
	(g)	What is Stuffer	Fragment?	2
			UNIT—I	
2.	(a)	What are Type I characteristic fea	I Restriction Endon atures.	nucleases? Discuss their 7
	(b)	Fragment.	son between DNA	A Pol I and Klenow 6
09	987/P	C-946	1	[Turn over

3.	(a)	What are Real time PCR and Inverse PCR? Expl	ain
		significance.	1
	(b)	Deliberate on Applications of PCR.	6
		UNIT—II	
4.	(a)	What are Insertion and Replacement vectors? Discuss vexamples.	with 6
	(b)	List cloning vectors of E.coli. Elaborate on features of pBR and pUC8.	327 7
5.	Wh	at is Insertional Inactivation? How is Blue White select	tion
	app	lied for identification of recombinants?	13
		UNIT—III	
6.	Wr	ite notes on :	
	(a)	Colony Hybridization	
	(b)	DNA probe labelling strategies. 6.5	5×2
7.	(a)	Discuss strategy for full length cDNA synthesis.	
	(b)	Write methods for mRNA enrichment. 6.	5×2
		UNIT—IV	
8.	W	rite notes on:	
	(a)) Pyro sequencing Technique	
	(b)) Site directed mutagenesis. 6.	.5×2
9.	(a)) Write about promoter designs for Recombinant proproduction in E.coil.	otein 7
	(b	 Explain limitations and advantages of E, coli as a host ce recombinant protein production. 	ll for 6