T.E- I Sem- Biotech

Genetic Engineering

QP Code: 31070

(3 Hours)

[Total Marks: 80]

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	Question No. 1 is compulsory.	
2)	Attempt any three questions from Q. Nos. 2 to 6	
3)	Draw neat labelled diagram wherever necessary.	
1.00		4-97
•	Write short notes on the following (any four) (5x4	= 20)
	a) Alkaline Phosphatase	
	b) Homopolymer tailing	1
	c) .Cosmids	1
*	d) RFLP	
	e) Antisense Technology	7
	f) Chromosome Walking	37
	· · ·	
2_	a) Discuss any two strategies for gene transfer in host?	(4.5)
27.10.	y - 10 - 10 of a constraint and of a constraint and a con	(10)
	birDescribe expression wastern with with a site bland of	9831_00
	b) Describe expression vectors with suitable example?	(10)
3.	a) Expisin bow the general library is an analysis of the same	
	a) Explain how the genomic library is created and now to identify a cione)
	from the library?	(10)
	b)What advantage does a thermostable DNA-polymerase offer in	
	PCR? Explain.	(10)
ł.	a)Describe Southern Blotting explaining the rational of each step involve	ed?
		(10)
	b)Comment on single stranded phages as vectors for gene cloning?	(10)
		And Annual of
5.	a) Describe Sanger and Maxim and Gilbert method of nucleic	\$• 3 .6
	acid sequencing?	(10)
		(10)
	b) Discuss types, nomenclature and use of restriction endonucleases in	•
	gene cioning?	NL 32-1088403
		(10)
5.	a) Discuss the echnique of transformation in E. Coli?	/465
•	as Discuss the scorningue of transformation in E. Colls	(10)

b) Write the applications of recombinant DNA technology in medicine?