Paper / Subject Code: 30001 / GENETIC ENGINEERING

1T00415 - T.E.(BIOTECHNOLOGY)(Sem V) (CBSGS) / 30001 - GENETIC ENGINEERING

(3 Hours)	(Total marks: 80)
N.B: (1) Question no. 1 is compulsory.	
(2) Attempt any 3 out of remaining questions.	
(3) Draw diagrams wherever necessary.	
1. WIDITE GLIODE NOTE ON (ANN A)	2
1. WRITE SHORT NOTE ON: (ANY 4)	(20)
a) Cosmids b) RFLP c) Super helical & Relaxed molecule	s d) PCR
e) RNA interference technology	
2. a) What are plasmids? Discuss their size, conjugation & con	patibility. (10)
b) What are lambda insertional and replacement vectors? C	ive examples (10)
3. a) Explain the structure of Ti plasmid and its use as a clonic	ng vector? (10)
b) Explain the steps of making cDNA library? How will yo	screen for your clone of interest
from cDNA library.	(10)
4. a) Describe in detail the enzyme Restriction Endonucleases .	(10)
b) State and explain any two methods of transfer of DNA in	nto cells . (10)
5. a) Explain the role of following enzymes in genetic engineeri	ng (20)
(i) Terminal nucleotidyl transferase	
(ii) DNA ligase	
(iii)Polynucleotide kinase	
(iv)Reverse transcriptase	
(v) Alkaline phosphatase	
6. a) Compare the following	(10)
(i) Southern and Northern blotting	
(ii) Automated sequencing and conventional Sanger's method	od of sequencing
b) What is Insulin? How recombinant insulin is produced in ba	cteria? Why is recombinant human
insulin better for diabetic patients than pig or cow insulin?	(10)
 (iii)Polynucleotide kinase (iv)Reverse transcriptase (v) Alkaline phosphatase 6. a) Compare the following (i) Southern and Northern blotting (ii) Automated sequencing and conventional Sanger's method b) What is Insulin? How recombinant insulin is produced in band 	od of sequencing cteria? Why is recombinant hum