Exam.Code:0038

Sub. Code: 0987

P.T.O.

2071

B.Sc. (Hons.) Biotechnology Sixth Semester VI. I.T. Genetic Engineering

| | BIOT- Sem-VI-I-T: Genetic Engineering | |
|--------------------------------------|--|--------------------|
| Time allowed: 3 Hours Max. Marks: 67 | | |
| NOTE: | Attempt <u>five</u> questions in all, including Question No. I which is and selecting one question from each Unit. $x-x-x$ | compulsory |
| I. | Attempt the following:- | |
| | a) What are Neoschizomers? Give examples. | (3) |
| | b) What are Shuttle Vectors? | (2) |
| | c) What are Reverse transcriptases? | (2) |
| | d) Why is PCR called Exponential amplification? | (2) |
| | e) What are Concatamers? | (2) |
| | f) What are double digests and partial digests? | (2) |
| | g) What is Stuffer Fragment? | (2) |
| | <u>UNIT - I</u> | |
| II. | a) What are Type II Restriction Endonucleases? Discuss their Characteria | stic features. |
| | b) Draw comparison between DNA Pol I and Klenow Fragment. | (8,5) |
| III | . a) What are Hot start PCR and Inverse PCR? Explain'significance. | |
| | b) Deliberate on Applications of PCR. | (7,6) |
| IV | • | |
| | b) What is Insertional Inactivation? Explain application with example. | (7,6) |
| V | Write about various Cloning vectors in yeast. | $(2x6\frac{1}{2})$ |
| | <u>UNIT - III</u> | |
| V | | |
| | a) Colony Hybridization | $(2x6\frac{1}{2})$ |
| | b) Functional complementation based screening. | (25072) |

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(2)

VII. a) Discuss strategy for full length cDNA synthesis.

b) Write methods for mRNA enrichment.

 $(2x6\frac{1}{2})$

<u>UNIT - IV</u>

VIII. Write notes on:

- a) Pyro sequencing.
- b) Strand Selection mutagenesis.

(2x6½)

IX. a) Write about promoter designs for Recombinant protein production in Yeast.

b) Discuss problems with recombinant protein production in E.coli.

 $(2x6\frac{1}{2})$