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Exam. Code : 107406

Subject Code: 1876

B.Sc. (Biotechnology) 6th Semester rDNA TECHNOLOGY—B

Paper: BT-1

Time Allowed—Three Hours] [Maximum Marks—40

SECTION-A

Note: — Attempt ALL questions. 1 mark each.

- 1. What are the applications of TAC vector?
- 2. Explain features of pET280.
- 3. What are lambda vectors?
- 4. What is hot start PCR?
- 5. How the blunt end can be converted into sticky ends for cloning?
- 6. What is cassette mutagenesis?
- 7. What is biopanning?
- 8. What are oilgonucleotide arrays?

SECTION—B

Note: — Attempt FIVE questions by selecting ONE from each unit. 4 marks each.

UNIT-I

- 1. Explain Ti plasmid in details and its application.
- 2. What kind of promoters are used in expression vectors?

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UNIT—II

- 3. Explain construction of genomic libraries.
- 4. Explain self priming method of cDNA synthesis and cloning.

UNIT—III

- 5. What is multiplex and touch-down PCR?
- 6. Explain, how can we analyze gene expression by microarrays?

UNIT—IV

- Explain chemical degradation based method of DNA sequencing.
- 8. How the site directed mutagenesis improves the function of proteins ?

SECTION—C

Note: Do any TWO questions. 6 marks each.

- 1. Explain BAC vectors in details. How they are different from YAC and expression vectors?
- 2. What are expression libraries? Explain their construction and screening.
- 3. Explain the principle of PCR. How can you do qPCR and RT-qPCR?
- 4. Explain phage display method for selection of mutants.