

This question paper contains **3** printed pages]

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S. No. of Question Paper : **5562**

Unique Paper Code : **2492013603**

Name of the Paper : **Fundamentals of Recombinant DNA Technology**

Name of the Course : **B.Sc. (Hons.) Biochemistry (NEP)**

Semester : **VI**

Duration : **2 Hours**

Maximum Marks : **60**

(Write your Roll No. on the top immediately on receipt of this question paper.)

There are **6** questions.

Attempt any *four* questions.

All questions carry equal marks.

Question No. **1** is compulsory.

1. (A) Write the contributions of the following :

- (i) Kary Mullis
- (ii) Paul Berg
- (iii) Bolivar and Rodriguez.

(B) State True/False and justify (any *six*) :

- (i) Hybrid promoters are preferred for expression vectors.
- (ii) Blunt ends increase the efficiency of ligation.
- (iii) All three types of restriction endonucleases are useful in genetic engineering.

P.T.O.

- (iv) Factor VIII cannot be expressed in *E. coli*.
- (v) Bacterial cells are naturally competent.
- (vi) Lambda replacement vectors are used to make genomic libraries.
- (vii) Any *E. coli* strain can be used for recombinant DNA technology.

(3,12)

2. Differentiate between the following :

- (i) Real time PCR and Multiplex PCR
- (ii) Genomic and cDNA library
- (iii) Linker and Adapter
- (iv) pUC18 and Ti-Plasmid
- (v) YAC and BAC.

(15)

3. (A) 200 DNA molecules were used as a template in a PCR reaction. Calculate the number of amplicons produced after 30 cycles.

(B) What are the uses of the following in genetic engineering ? Explain with the help of a diagram :

- (i) BAL31
- (ii) T4 Polynucleotide Kinase
- (iii) Alkaline Phosphatase
- (iv) EcoRI
- (v) S1 Nuclease
- (vi) DNA ligase.

(3,12)

4. (A) What are the different classes of viral vectors ? Discuss their advantages and disadvantages in genetic engineering.
- (B) Describe the principle of Sanger's method of DNA sequencing. List *two* limitations of Sanger's method.
- (C) What are the essential features of a typical expression vector ? Give *two* examples. (6,4,5)
5. (A) Discuss the factors that affect the efficiency of a PCR reaction.
- (B) Briefly explain *two* methods of DNA uptake by cells.
- (C) Draw a labelled diagram of pGEM3Z vector. How is it different from pBR322 cloning vector ? (5,5,5)
6. Write short notes on the following (any *five*) : (15)
- (i) Pyrosequencing
 - (ii) Site directed mutagenesis
 - (iii) Homopolymer Tailing
 - (iv) Colony Hybridization
 - (v) Fusion tags and their role in purification of recombinant proteins
 - (vi) Blue-White Selection.