V(5th Sm.)-Microbiology-G/DSE-A-1/CBCS

2021

MICROBIOLOGY — GENERAL

Paper : DSE-A-1

(Genetic Engineering and Biotechnology)

Full Marks : 50

The figures in the margin indicate full marks. Candidates are required to give their answers in their own words as far as practicable.

Group - A

1. Answer any five questions :

- (a) Write down any two features of pBR cloning vector.
- (b) Write down the differences between Type-I and Type-II restriction enzymes.
- (c) What are the differences between kinases and phosphatases?
- (d) Give two examples of vectors which can transfer eukaryotic gene.
- (e) Write down two importances of taking copyright.
- (f) What are the two properties of a good vector?
- 2. Write short notes on (any three) :
 - (a) SDS-PAGE
 - (b) Role of RDT in insulin production
 - (c) Bt-transgenic cotton
 - (d) Polymerase chain reaction
 - (e) Cosmid vectors.

Group - B

Answer any five from the following.

- 3. (a) What do you mean by Real-time PCR?
 - (b) What is the purpose of Northern blotting?
- 4. (a) Briefly describe any approach for identification and isolation of recombinant clone carrying the desired DNA insert.
 - (b) Name one type-II restriction endonuclease.

Please Turn Over

 2×5

5×3

 $2^{1/2}+2^{1/2}$

4 + 1

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- 5. Write notes on :
 - (a) Transformation of DNA as genetic material
 - (b) Significance of Patents.
- 6. (a) What is YC_p ?
 - (b) Write down two characteristic features of E. coli lac promoter and T7 promoter-based vectors.
- 7. Schematically describe the steps of cDNA library construction.
- 8. (a) Why do you think that DNA-sequencing of a gene is important?
 - (b) Write down Sanger's method of DNA sequencing. Describe with a suitable diagram.

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

- 9. (a) Write down two applications of genetic engineering in the field of human therapeutic interest.
 - (b) How much gene length can we transfer by using the bacteriophage lambda as cloning vector?
 - (c) What do you understand by shuttle vector? 2+1+2
- 10. Explain the role of the following in gene cloning :
 - (a) terminal deoxynucleotidyl transferase
 - (b) reverse transcriptase.

21/2+21/2

 $2^{1/2} \times 2$

1+(2+2)

5