2020

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.

Day 2

Group - A

1. Answer any five questions:

 2×5

- (a) A laboratory strain of *E.coli* is resistant to ampicilin. Can this *E.coli* be used as host in recombinant DNA technology? Explain your answer.
- (b) What is the speciality of type II restriction enzyme?
- (c) What is primer dimer in PCR?
- (d) What is Klenow?
- (e) What is cosmid vector?
- (f) Write the name of an octacutter restriction enzyme.
- (g) What is blunt end ligation?
- 2. Write short notes on (any three):

5×3

- (a) Liposomes
- (b) Ti-plasmid
- (c) BACs
- (d) SDS-PAGE
- (e) T4 poly nucleotide kinase.

Group - B

Answer any five questions.

- 3. (a) Describe the advantages of λ -phage based rectors over plasmid vectors.
 - (b) Describe the function of IPTG for the over expression of a protein from lac-promoter based expression vector in *E.coli*. $2\frac{1}{2}+2\frac{1}{2}$

Please Turn Over

T(5th Sm.)-Microbiology-G/DSE-A-1/CBCS/Day-2 (2)

- 4. (a) Describe the difference between southern blotting and western blotting.
 - (b) "Blocking of the membrane with a non-specific protein like BSA is very important during western blot analysis"— justify the statement.

 2½+2½
- 5 (a) Why is the presence of unique restriction enzyme site(s) at the non-essential part of a cloning vector very important?
 - (b) What is the difference between an endonuclease and an exonuclease?
 - (c) What do you know about GM-crops?

2+1+2

- **6.** (a) How do you prepare human growth hormone by the help of RDT?
 - (b) Why is partial digestion of genomic DNA with restriction endonucleases done during preparation of genomic DNA library? 2½+2½
- 7. (a) Name one vector that is suitable for cloning in mammalian cell. Give two characteristic features of it.
 - (b) Write down two advantageous features of M13-based vectors.
 - (c) Give an example of a low copy cloning vector.

(1+1)+2+1

8. How will you prepare genetically engineered recombinant human insulin?

5

- 9. (a) Describe the different steps of RT-PCR.
 - (b) How is it different from real time PCR?

 $3\frac{1}{2}+1\frac{1}{2}$

- **10.** (a) Describe the difference between genomic DNA-library and cDNA library. Mention the importance of cDNA library in genetic engineering.
 - (b) What is 'intellectual property right'?

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