

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  $\lambda$ -phage based vectors 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*. 2½+2½

**Please Turn Over**

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? 3½+1½
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+1½)+1½
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