M.Sc. BIOTECHNOLOGY THIRD SEMESTER GENETIC ENGINEERING MBT-301

(Use separate answer scripts for Objective & Descriptive)

| Dı | ration: 3 hrs. | F | full Marks: 7 |
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| Γi | (<u>PART-A: O</u> me: 20 min. | <u>bjective</u>) | Marks: 2 |
| Choose the correct answer from the following: | | | 1x20=20 |
| 1. | The 300Kb DNA can be inserted into | b. YAC | |
| | c. Cosmid | d. BAC | |
| 2. | Sticky ends are preferable because | | 11 |
| 3. | Restriction modification system requires Me a. Polymerasesc. Kinases | thylases andb. Restriction enzymes d. Phosphatases | |
| 4. | RNase H digestsa. RNA-DNA hybrid c. dsDNA | b. ssDNA and RNA d. ssDNA and ds DNA | |
| 5. | 3' OH is connected to 5' P group of DNA by a. Ligase c. Isoschizomers | b. Type I nucleases d. Type II nucleases | |
| 6. | Usuallysequence is added to the a. 5'CCCCCCC3' c. 5'AAAAAAAA3' | primer used against 3' region b. 5'GGGGGGGG3' d. 5'TTTTTTTTTT3' | of RNA. |
| 7. | The function of Alkaline phosphatases is to. a. Add phosphate c. Synthesize P | b. Remove P d. None of the above | |
| 8. | Molecular tools are | b. RE and ligases d. Both b and c | |
| 9. | Shuttle vector's example is | b. BAC d. Cosmid | |
| 10 | . Reverse transcriptase is not used in | b. Retrovirusesd. All are correct | |

11. Dideoxy nucleotides are required in: a. In place of normal nucleotides at high concentration. b. Chain termination method. c. Chemical method. d. All are correct. 12. The marker used in DNA finger printing is: a. Mini satellites b. SNP d. ISSR c. Micro satellites 13. The most common vector for plants are: a. SV 40 Vectors and Bovine Papilloma virus vectors. b. Lambda phage and M13 phage vectors. c. Cauliflower mosaic virus and Gemini virus vectors. d. T4 phage vectors. 14. Solid matrix mostly used in Southern is: a. Glucose membrane b. Nitrocellulose membrane c. Nylon membrane d. All can be used 15. Which portion of Ti plasmids is transferred to plants to cause crown gall disease? b. Ori a. T-DNA region d. All of the above c. Vir region **16.** *Pfu* and *Vent* polymerase are more efficient than Taq polymerase because: a. Of more efficient polymerase activity b. Of proof reading activity d. None of these c. Both a and b 17. Which of the following would be eliminated by Hot Start PCR? a. Aerosol contamination from the barrel of pipetors. b. Addition of nucleotide to the terminal end of PCR products. c. Infidelity of DNA copying by Tag DNA polymerase. d. Formation of primer-dimers. 18. Which of the following bio-molecule has self repair mechanism?

a. DNA, RNA and protein

b. DNA and RNA

c. DNA and protein

d. DNA only

19. Which of the following methods for introducing DNA into cells is used only for plants?

a. A gene 'gun

b. Microinjection

c. Electroporation

d. Transformation of competent cells

20. The DNA chain acting as template for RNA synthesis has the following order of bases, AGCTACGA. What will be the order of bases in mRNA?

a. TCGATGCT

b. TCGAUGCT

c. UCGUAGCU

d. UCGAAGCU

PART-B: Descriptive

Time: 2 hrs. 40 min. Marks: 50

[Answer question no.1 & any four (4) from the rest]

5+5=10 1. How will you transform plants using Ti plasmids? Discuss any two physical mediated methods used for transferring gene of interest into the cell. 2. Explain restriction endonucleases with suitable restriction site. Explain 5+5=10 function of ligase with suitable diagram. 4+6=10 3. Discuss the restriction mapping mechanism in brief. Write basic differences between BAC and YAC. 6+4=10 4. Explain homopolymer tailing with its significance. Explain sticky end is preferable in genetic engineering. 6+4=10 5. Explain plasmid with suitable structure. Explain the characteristics of a good vector. 10 6. Using Sanger sequencing method how will you determine the sequence of the DNA template: ATCGATCGATCTTAGCCATA? Explain with the help of suitable diagram. 2+3+5=10 7. What is the function of MgCl₂ and dNTPs in polymerase chain reaction (PCR)? Calculate the annealing temperature of the primer: GACTCCTATAGTCTACAAATGCC. Briefly explain Hot start PCR? 8. Write short notes on: 2x5=10

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a. Blotting technique to identify the product of a gene of interest

b. PCR based molecular marker