

PSG COLLEGE OF ARTS & SCIENCE  
(AUTONOMOUS)

MSc DEGREE EXAMINATION MAY 2023  
(Second Semester)

Branch – BIOTECHNOLOGY

RECOMBINANT DNA TECHNOLOGY

Time: Three Hours

Maximum: 50 Marks

SECTION-A (5 Marks)

Answer ALL questions

ALL questions carry EQUAL marks

(5 x 1 = 5)

- 1 Choose the Restriction site for Bam HI.  
(i) CCCGGG (ii) GGCC  
(iii) GGATCC (iv) AAGCTT
- 2 Plasmids containing lambda DNA with cos site are referred as \_\_\_\_\_.  
(i) Phasmids (ii) BAC Vectors  
(iii) Cosmids (iv) YAC vectors
- 3 Identify the commonly used vector in genomic library construction.  
(i) λEMBL (ii) λZAP  
(iii) C2XB (iv) pET
- 4 Which of the following contributes to errors during PCR amplification due to misincorporation of nucleotide  
(i) Random Primers (ii) Temperature variation  
(iii) Taq Polymerase (iv) dNTPs
- 5 Name the vector used in site-directed mutagenesis.  
(i) pUC18 (ii) pBR322  
(iii) M13 (iv) pSC101

SECTION - B (15 Marks)

Answer ALL Questions

ALL Questions Carry EQUAL Marks

(5 x 3 = 15)

- 6 a Discuss the role of Methyl transferase in DNA methylation.  
OR  
b Analyze the role of kinases and phosphatases.
- 7 a Show how cloning in *E.coli* is performed?  
OR  
b Illustrate on phagemids and packing of DNA.
- 8 a Sketch *in vitro* packaging of λ vectors.  
OR  
b Discuss protein expression Vectors.

Cont...

- 9 a Explain autoradiography.  
OR  
b Show how capillary based gel electrophoresis is used in sequencing.
- 10 a Explain selection of mutants.  
OR  
b Discuss PCR based Tn mutagenesis.

**SECTION -C (30 Marks)**

Answer ALL questions

ALL questions carry EQUAL Marks

(5 x 6 = 30)

- 11 a Classify Restriction enzymes and explain the nomenclature with examples.  
OR  
b Construct a generalized cloning strategy.
- 12 a Construct pBR322 and discuss its structure and biology.  
OR  
b Design cloning strategy using any one yeast vector.
- 13 a Construct a cDNA library with any one commonly used method.  
OR  
b Enumerate the role of Tissue specific vectors and explain the protein purification techniques used after expression.
- 14 a Justify how Sangers Di-deoxy method is applied in DNA sequencing.  
OR  
b Categorize PCR and note on its applications.
- 15 a Plan a site-directed mutagenesis using M13 vector.  
OR  
b Create a mutation using CRISPR/cas9 gene editing.

Z-Z-Z

END