

PSG COLLEGE OF ARTS & SCIENCE
(AUTONOMOUS)

BSc DEGREE EXAMINATION MAY 2022
(Fourth Semester)

Branch – BIOTECHNOLOGY

RECOMBINANT DNA TECHNOLOGY

Time: Three Hours

Maximum: 75 Marks

SECTION-A (10 Marks)

Answer ALL questions

ALL questions carry EQUAL marks (10 x 1 = 10)

1. Single stranded unpaired extensions formed by restriction enzyme upon cleavage is called as
 - (i) Blunt ends
 - (ii) Flush ends
 - (iii) sticky ends
 - (iv) none of these
2. Which of the following ions are required for the activity of Type II restriction enzymes
 - (i) Ca²⁺
 - (ii) Mg²⁺
 - (iii) Cl⁻
 - (iv) Mn²⁺
3. The first engineered plasmid vector is
 - (i) pBR 322
 - (ii) pUC vectors
 - (iii) pUC101
 - (iv) pUC19
4. Vectors designed to replicate in cells of two different species are called
 - (i) Phasmids
 - (ii) transfer vectors
 - (iii) shuttle vectors
 - (iv) phagemids
5. Plasmids which are maintained as multiple copy number per cell are known as
 - (i) Stringent plasmids
 - (ii) relaxed plasmids
 - (iii) cryptic plasmids
 - (iv) none of these
6. Cosmid vectors are
 - (i) plasmids that contain fragment of lambda DNA including the cos site
 - (ii) Phages that lack cos site
 - (iii) Plasmids that have no selection marker
 - (iv) cryptic plasmids
7. PCR is used in
 - (i) site specific recombination
 - (ii) site directed mutagenesis
 - (iii) both (i) and (ii)
 - (iv) site specific translocation
8. Pfu and Vent polymerase are more efficient than Taq polymerase because
 - (i) of more efficient polymerase activity
 - (ii) of proof reading activity
 - (iii) both (i) and (ii)
 - (iv) none of these
9. Which of the following properties is improved by site directed mutagenesis?
 - (i) Physical property
 - (ii) Chemical property
 - (iii) kinetic property
 - (iv) integrity
10. Which phage is used in oligonucleotide directed mutagenesis?
 - (i) M13
 - (ii) Cosmid
 - (iii) Phagemid
 - (iv) Lambda Phage

Cont...

SECTION - B (25 Marks)

Answer ALL questions

ALL questions carry EQUAL Marks

(5 x 5 = 25)

- 11 a Write a short notes on DNA methyltransferase.
OR
b Describe about End modifications process
- 12 a Brief about lambda phage cloning vector.
OR
b What are the criteria of an ideal vector?
- 13 a Elaborate on bacterial expression system.
OR
b How the recombinants can be selected by blue white screening?
- 14 a Write about Maxam – Gilbert method of DNA sequencing.
OR
b Discuss about RT – PCR.
- 15 a Explain about site directed mutagenesis.
OR
b Elaborate on site specific mutagenesis.

SECTION - C (40 Marks)

Answer ALL questions

ALL questions carry EQUAL Marks

(5 x 8 = 40)

- 16 a Elucidate the origin and types of restriction enzymes.
OR
b How the genes can be cloned in recombinant DNA technology.
- 17 a With a neat diagram explain the cloning vectors.
OR
b Write in detail about the types of vectors.
- 18 a Distinguish between genomic library and cDNA library.
OR
b Explain in detail about invitropackaging.
- 19 a Illustrate the methods of nucleic acid sequencing.
OR
b Describe in detail about the applications of PCR.
- 20 a Describe about PCR based mutagenesis.
OR
b Explain the process of production of glucagon through recombinant DNA technology.

Z-Z-Z

END