

P. P. SAVANI UNIVERSITY

Third Semester of M.Sc. Examination

December-2021

SSBT8090- Genetic Engineering: Theory and Application

09.12.2021, Thursday Time: 09:00 a.m. to 11:30 a.m. Maximum Marks: 60

Instructions:

1. The question paper comprises of two sections.
2. Make suitable assumptions and draw neat figures wherever required.

Section-I

Q.1 Very Short Questions (Attempt any five) [10]

- 1.1 What is a promiscuous plasmid?
- 1.2 Primer extension method for Site Directed Mutagenesis was developed in which year and by whom?
- 1.3 What is the cloning capacity of PAC and BAC?
- 1.4 What is a Cosmid?
- 1.5 Name two species of *Agrobacterium* which are commonly used for creating transgenic plants.
- 1.6 What are the four major steps in genomic library preparation?

Q.2 Write Short Notes (Attempt any two) [06]

- 2.1 Draw a structure of typical yeast promoter.
- 2.2 Chloroplast transformation
- 2.3 Name three commonly used selectable marker genes in animals along with principle of selection.

Q.3 Detail questions (Attempt any two) [14]

- 3.1 How you can construct a c-DNA library?
- 3.2 Explain four major strategies for gene transfer to plant cells?
- 3.3 Draw structure of YAC? Explain working, advantages and disadvantages of YAC vector?

Section-II

Q.1 Very Short Questions (Attempt any five) [10]

- 1.1 Name two promoters that can induce glucocorticoid hormone in mammals.
- 1.2 Name steroid hormone associated with molting in insects.
- 1.3 What is Exon shuffling?
- 1.4 What are positional orthologs?
- 1.5 Give genome size of *Pseudomonas aeruginosa* and *E. coli* in Mb
- 1.6 Give two examples of Site specific recombination systems

Q.2 Write Short Notes (Attempt any two) [06]

- 2.1 What are the three limitations associated with the use of endogenous inducible promoters?
- 2.2 Differentiate between orthologs and paralogs.
- 2.3 Explain about *Deinococcus radiodurans* remarkable ability.

Q.3 Detail questions (Attempt any two)

[14]

- 3.1** Explain in detail lac and tet repressor systems in bacteria.
- 3.2** How site specific recombination facilitates precise transgene integration?
- 3.3** Explain in detail comparative genomics of bacteria.