### rDNA TECHNOLOGY

# Class: I, M. Sc., Biotechnology Sub Code: P16BT15

## Unit I

# 2 Mark ( Define the following)

- 1. Nuclein
- 2. Genetic material
- 3. Nucleosides
- 4. Nucleotides
- 5. Okazaki fragments
- 6. Polynucleotide
- **7.** Phosphodiester bond
- 8. Anti parallel
- 9. Palindromic DNA
- 10. Repetative DNA
- 11. Satellite DNA
- 12. Polyploid
- 13. Centromere
- 14. Heterochromatin
- **15.** Telomeres
- **16.** Chromosomin
- 17. Mono Cistronic mRNA
- 18. Polycistronic mRNA
- 19. Soluble RNA
- 20. Supernatant RNA
- 21. Anticodon loop
- 22. tRNA al
- 23. Polymerization
- **24.** RNA Primer
- 25. Ligase
- **26.** Restriction enzymes
- 27. Klenow enzyme
- 28. T4 dna polymerase
- **29.** Polynucletide kinase
- **30.** Alkaline phosphatise
- 31. Cohesive end ligation
- 32. Blent end ligation
- 33. Linkers
- 34. Adaptors
- **35.** Homopolmer tailing
- **36.** Nick translation
- **37.** Probe
- **38.** Radio active probe
- **39.** Non radio active probe
- 40. Dna foot printing

- 1. Given account on restriction enzymes.
- 2. Explain in detail about labelling of dna
- **3.** Explain in detail about colony hybridization.
- **4.** Explain about electromobility shift assay
- **5.** Explain about methyl interference assay.

- 1. Explain about the hybridization techniques.
- 2. Explain about northern blotting
- **3.** Explain about southern blotting
- **4.** An account on Dna protein interactions.
- 5. Explain in detail about dna foot printing

#### Unit II

# 2 Mark ( Define the following)

- **41.** Vector
- 42. Plasmid
- **43.** Col Plasmid
- 44. F Plasmid
- **45.** Ri Plasmid
- **46.** Ti Plasmid
- **47.** Copy Number
- **48.** PBR322
- **49.** Transposons
- **50.** Phagemids
- **51.** Blue script vectors
- **52.** Bacteriophage.
- **53.** Lambda phage vectors
- **54.** EMBL
- 55. Cosmid
- **56.** Fosmid
- **57.** YAC
- **58.** BAC
- **59.** Bacculo virus vectors
- **60.** Retroviral vectors
- **61.** Expression Vectors
- **62.** His-tag
- **63.** GST-tag
- **64.** Yeast vectors
- **65.** Shuttle vectors
- **66.** Pichia vectors
- **67.** MBP tag
- **68.** Protein based vectors

- **6.** Given account on structure of Plasmids and their Character
- 7. Discuss about the Difference between the Natural and Artificial Plasmid
- 8. Give an Brief account on Construction of Plasmid
- 9. Explain in detail about YAC

- 10. Explain in detail about BAC
- 11. Discuss about Protein based vectors
- 12. Explain about animal based vectors
- **13.** Explain about plant based vectors
- **14.** Explain in detail about expression based vectors.

- **6.** Explain about the types of Plasmids
- 7. Give an detailed note on Plasmid Transfer and its applications
- **8.** Describe about the Transposable genetic elements in Prokaroyotes and Eukaryotes and their uses
- 9. Explain about cosmid and its types.
- 10. Explain about artificial chromosomes.

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- 23. Polymerization
- 24. RNA Primer
- 25. Ligase

- 1. Describe about the Nucleosides
- 2. Give an account on Nucleotides
- 3. Write a note on Properties and function of DNA
- 4. Brief account on Types of DNA

- 5. Describe about the Chromosome structure and function
- 6. Give an account on types of Chromosome
- 7. Discuss about the differentiation between DNA and RNA
- 8. Describe about the Rolling Circle mechanism of DNA replication
- 9. Give an account on inhibition of DNA repair

- 1. Explain about the Specialized chromosomes
- 2. What is DNA replication and its types
- 3. Explain about the types of RNA and its function
- 4. Explain about the DNA replication in Eukaryotes
- 5. Describe about the DNA repair mechanism

### **Unit III**

## 2 Mark (Define the following)

- 1. Transcription
- 2. Translation
- 3. Central Dogma
- 4. Promoter
- 5. Initiation site
- 6. Amino Acyl Adenylate
- 7. Elongation
- 8. Translocase
- 9. Anti termination
- 10. Post Transcripitional modification
- 11. Post Trnaslational Modification
- 12. hnRNA
- 13. Poly Adenylation
- 14. Catalytic RNA
- 15. Codon
- 16. Anticodon
- 17. Protein Stability
- 18. Wobble Base Pairs
- 19. Splicing
- 20. Site directed mutagenesis.
- 21. Conjugation
- 22. Transfection
- 23. Transduction
- 24. Gene libraries
- 25. c dna library
- 26. Genomic cloning

- 1. Give a brief note on Plyadenylation
- 2. Describe about the RNA Splicing
- **3.** How the post Transcriptional modification occur?
- **4.** Describe about the Post translational modification
- **5.** Explain in detail about site directed mutagenesis.
- **6.** Isolation of foreign dna into host cell.

7. Isolation of mRna and t Rna procedure.

### 10 Mark

- 1. What is Genetic code? Explain the Characterization of Genetic code.
- 2. Explain about the Transcription
- 3. Give an detailed note on Translation
- 4. Explain about the Ribosome composition and its assembly.
- 5. Explain in detail about genomic cloning.
- 6. Explain in detail about c dna cloning and libraries.
- 7. Explain in detail about protein protein interactions cloning.
- 8. Discuss about yeast two hybrid system.
- 9. Explain in detail about gene expression.

### **Unit IV**

# 2 Mark ( Define the following)

- 1. Gene Transfer
- 2. Transformation
- 3. Gene Amplification
- 4. Inducer
- 5. Repressor
- 6. House keeping genes
- 7. Ara Operon
- **8.** Operon
- 9. Gene Dosage
- 10. Sigma Factor
- 11. Anti Sense RNA
- 12. DNA finger printing.
- **13.** DNA profiling
- 14. Multiplex pcr
- **15.** Real time pcr
- 16. Touch down per
- 17. Hot start per
- **18.** SSCP
- **19.** DGGE
- **20.** RFLP
- **21.** OLA
- **22.** MCC
- **23.** ASA
- **24.** PTT
- 25. T-vectors.

- 1. Discuss about the per working principle and its applications.
- 2. Short note on SSCP
- 3. Write the role and application of RFLP.
- 4. Explain in detail about DGGE and OLA.
- 5. Explain in detail about MCC, ASA and PTT detection.

- 1. Explain in detail about PCR and its types.
- **2.** Explain in detail about proof reading enzymes.
- **3.** Discuss about pcr in gene recombination.
- **4.** Explain in detail about per In molecular diagnosis.
- 5. Discuss about primer design.

#### Unit V

# 2 Mark ( Define the following)

- 1. Transgenic animals
- 2. Transgenic Plants
- 3. IVF
- 4. Herbicides
- 5. Pesticides
- 6. Organic products
- 7. DNA Vaccines.
- 8. Pharmaceutical products.
- 9. Insulin.
- 10. Transgenic genes
- 11. Gene theraphy
- 12. Gene knock outs
- 13. Dna sequencing
- 14. Rna sequencing
- 15. Gene silencing
- 16. Oligonucletides
- 17. Micro rna vectors
- 18. Gene replacement
- 19. Gene targeting
- 20. Protein array
- 21. Si RNA vectors.

## 5 Mark

- 1. Methods of transgenic animal production.
- 2. Explain in detail about Dna sequencing
- 3. Explain in detail about Rna sequencing
- 4. Discuss about gene silencing
- 5. Explain about chemical synthesis of oligonucletides.

- 1. Explain about the DNA vaccine production
- 2. Explain about the Designing of Vaccine
- 3. Explain the human genome project.
- 4. Explain about somatic and germ line theraphy
- 5. Discuss about protein array.
- 6. Explain in detail about gene knockout mice.
- 7. Explain in detail about principle, construction and applications of sirna vectors.