





BOARD OF SCHOOL EDUCATION HARYANA

Syllabus and Chapter wise division of Marks (2024-25)
Class: XII Subject: Biotechnology Code: 871

General Instructions:

- 1. There will be an Annual Examination based on the entire syllabus.
- 2. The Annual Examination will be of 60 marks, Practical Examination will be of 20 marks and 20 marks weightage shall be for Internal Assessment.
- 3. For Practical Examination:
 - i) Two experiments of 6 marks each.
 - ii) One activity of 3 marks.
 - iii) Practical record of 2 marks.
 - iv) Viva-voce of 3 marks.
- 4. For Internal Assessment:

There will be Periodic Assessment that would include:

- i) For 4 marks- Two SAT exams will be conducted and will have a weightage of 04 marks towards the final Internal Assessment.
- ii) For 2 marks- One half-yearly exam will be conducted and will have a weightage of 02 marks towards the final Internal Assessment.
- iii) For 2 marks- One Pre-Board exam will be conducted and will have a weightage of 02 marks towards the final Internal Assessment.
- iv) For 2 marks- Subject teacher will assess and give maximum 02 marks for CRP (Classroom participation).
- v) For 5 marks- A project work to be done by students and will have a weightage of 05 marks towards the final Internal Assessment.
- vi) For 5 marks- Attendance of student will be awarded 05 marks as:

75% to 80% - 01 marks

80% to 85% - 02 marks

85% to 90% - 03 marks

90% to 95% - 04 marks

95% to 100% - 05 marks













Course Structure (2023-24)

Class-XII

Subject-Biotechnology

Code:871

| Sr. No. | Unit | Chapter | Marks | |
|---------|---|-------------------------------------|-------|--|
| 1 | | Recombinant DNA Technology | | |
| | Protein and Gene Manipulation | Protein Structure and Engineering | 30 | |
| | | Genomics and Bioinformatics | | |
| 2 | Cell Structure and Genetic Microbial Cell Culture and its Application | | 20 | |
| | | Plant Cell Culture and Application | 30 | |
| | | Animal Cell Culture and Application | | |
| | Total | | | |
| | Practica | 20 | | |
| | Interna | 20 | | |
| | Gr | and Total | 100 | |













Unit V: PROTEIN AND GENE MANIPULATION

Chapter 1: Recombinant DNA Technology

Introduction; Tools of rDNA Technology: enzymes, Restriction Fragment Length Polymorphism (RFLP), Other enzymes used in cloning, Vectors: Plasmids, vector based on bacteriophages, cosmids, YAC vectors, BAC vectors, Animal and plant viral vectors, Host cells; Marking rDNA; Introduction of rDNA to Host Cells: Transformation, Electroporation, Transfection, Microinjection, **Biolistics:** Identification of Recombinants; Polymerase Chain Reaction (PCR); Hybridisation Techniques: Southern Hybridisation DNA Library: DNA Sequencing: Technique; Dideoxynucleotide chain termination method; Site-directed **Mutagenesis**

Unit V: PROTEIN AND GENE MANIPULATION

Chapter 2: Protein Structure and Engineering

Introduction to the World of Proteins; 3-D Shape of **Proteins:** Non-covalent bonds: ionic bonds, hydrogen bonds, Van der Waals forces, hydrophobic interactions; Structure-Function relationship in Proteins: Chymotrypsin proteolytic enzyme, Molecular Disease- Sickle cell anaemia, printing-Peptide Mapping, Protein Finger Electrophoresis; Purification of Proteins: Calculation amount of bacterial ferment required, Downstream Processing, Aqueous two-phase partition, Industrial scale production of proteins, Special techniques for therapeutic /diagnostic proteins; Characterisation of Proteins: Mass spectrometry; Based Products: Blood products and vaccines, Therapeutic antibodies and enzymes, Therapeutic hormones and growth factors, Regulatory factors, Analytical applications, Industrial enzymes, Functional non-catalytic proteins, Nutraceutical **Designing Engineering**): Proteins; **Proteins** (Protein Improving laundry detergent Subtilisin, Creation of Novel Proteins, Improving nutritional value of cereals and legumes.

Unit V: PROTEIN AND GENE MANIPULATION

Chapter 3: Genomics and Bioinformatics













Introduction: Progress in stages, Evolving approaches, genomics, Functional Genome Structural genomics; **Sequencing Projects:** Directed sequencing of Bacterial Artificial Chromosome (BAC) contigs, Random shotgun Gene prediction and counting: sequencing: Similarity, SNPs and Comparative Genomics; Functional **Genomics:** Fluorescence in situ hybridization, Microarray Technology: principle, procedure, interpretation; **Proteomics:** Types of Proteomics: Expression proteomics, proteomics, Functional proteomics, Genes and Proteins: Number of genes vs Number of proteins; **History** Bioinformatics; Sequences and nomenclature: DNA and protein sequences, The concept of directionality, Different types of sequences; **Information Sources:** Major databases: NCBI, Database retrieval tools, BLAST family of search tools, Resources for gene level sequences, Analysis using Bioinformatics tools.

Unit VI: CELL CULTURE AND GENETIC MANIPULATION

Chapter 1: Microbial Cell Culture and Its Applications

Introduction; Microbial culture techniques: Nutrients for microbial culture, Culture Procedures, Equipment for microbial culture, Types of microbial culture: Batch culture, Fed-batch culture, continuous culture; Measurement and kinetics of microbial growth: Measurement of microbial growth, Growth kinetics and specific growth rate; Scale-up of microbial processes; Isolation of microbial products; Strain isolation, improvement and preservation: Strain isolation, Strain improvement: Mutation Selection, Genetic Engineering Techniques, Metagenomics, Strain preservation, Culture Collections Centers; Applications of microbial culture technology; Biosafety issues in Microbial Technology.

Unit VI: CELL CULTURE AND GENETIC MANIPULATION

Chapter 2: Plant Cell Culture and Applications

Introduction; Cell and Tissue Culture Techniques: Basic Technique, Nutrient Media, Types of cultures: organ culture, explant culture, callus culture, cell suspension culture, mass cell culture, protoplast culture, protoplast fusion, Plant Regeneration pathways; Applications of Cell and Tissue Culture: Micropropagation, Virus-free plants, Artificial seeds, Embryo rescue, Haploids and triploids, Somatic hybrids and cybrids, Production of secondary metabolites, Somaclonal variation, *In*













vitro plant germplasm conservation; Gene transfer methods in plants: Vector-mediated or indirect gene transfer, Vectorless or direct gene transfer, Transgene analysis; Transgenic plants with beneficial traits: Stress tolerance, Biotic stress tolerance: Herbicide tolerance, Pest resistance, Disease resistance, Virus resistance, Fungi and bacteria, Abiotic stress tolerance: Delayed fruit ripening, Male sterility, Transgenic plants as bioreactors, Nutrient quality, Diagnostic and therapeutic proteins, Edible vaccines, Biodegradable plastics, Metabolic engineering and secondary products, Other applications; Biosafety in Plant Genetic Engineering.

Unit VI: CELL CULTURE AND GENETIC MANIPULATION

Chapter 3: Animal Cell Culture and Applications

Introduction; Animal Cell Culture Techniques: Features of animal cell growth in culture, Primary Cell Cultures, Secondary Cell Cultures and Cell Lines, Types of Cell Lines: Finite Cell Lines, Continuous Cell Lines, Physical environment for culturing Animal Cells: temperature, pH, osmolality, Medium, serum and antibiotics, vessels and equipments required for animal cell culture, tissue culture hood, CO₂ incubator, centrifuge, inverted microscope; Characterization of Cell Lines: storage and revival of cells; Methods of Gene Delivery into Cells; Scale-up of Animal Culture Process; Applications of Animal Cell culture: Erythropoietin, Factor VIII, Factor IX, Tissue Plasminogen Activator (tPA), Hybridoma Technology for Monoclonal Antibody Production, Therapeutic mAb - OKT3, Therapeutic mAb - Herceptin; Stem Cell Technology: ES Cell culture and its applications; Tissue engineering.

Practicals:

- 1. Use of special equipment in biotechnology experiments
- 2. Isolation of bacterial plasmid DNA
- 3. Detection of DNA by gel electrophoresis
- 4. Estimation of DNA by UV spectroscopy
- 5. Isolation of bacteria from curd & staining of bacteria
- 6. Cell viability assay using Evan's blue dye exclusion method
- 7. Data retrieval and database search using internet site NCBI and download a DNA and protein sequence from internet, analyse it and comment on it
- 8. Reading of a DNA sequencing gel to arrive at the sequence













Month wise Syllabus Teaching Plan (2024-25)

Class: XII Subject: Biotechnology Code: 871

| Month | Subject-content | Teaching | Revision | Practical |
|--------------------|---|----------|----------|-----------|
| | | Periods | Periods | Work |
| April | Unit V | 22 | 2 | |
| | Chapter 1: | | | |
| | Recombinant DNA | | | |
| | Technology | | | |
| | CRET | MIT | | |
| _ | Practical: Use of | 1619 | 1.6 | |
| file of the second | special equipment | | 1/20 | 6 |
| 1 | in biotechnology | 0 | 77 | × 1 |
| 1 1 | experiments | 7) | | = 4 1 |
| 103 | | N | | 92 |
| 17 | Practical: Isolation | - Lbu | | |
| no | of bacterial plasmid | | | 2 |
| 1" | DNA | 1 | - 1 | |
| | | | | |
| | Practical: Detection | | | 2 |
| | of DNA by gel | 1 | | |
| N. | electrophoresis | | - | |
| N. | | | | 1 |
| | Practical: | | 4 | 2 |
| | Estimation of DNA | 9 | 100 | |
| | by UV | N MAN | | |
| | | | | |
| May | Unit V | 22 | 2 | |
| | | 0.50 | | |
| | Structure and | | | |
| | | | | |
| | | | | |
| | Practical: Isolation | | | |
| | of bacteria from | | | 2 |
| | | | | |
| | bacteria | | | |
| | | | | |
| May | Practical: Estimation of DNA by UV spectroscopy Unit V Chapter 2: Protein Structure and Engineering Practical: Isolation of bacteria from curd & staining of | 22 | 2 | |













| June | | | | |
|--------|--------------------------------|-------|------|-------------|
| | Summer Vacation (Project Work) | | | |
| | | , | | |
| July | Unit V | | | |
| | Chapter 3: | | | |
| | Genomics, | | | |
| | Proteomics and | | | |
| | Bioinformatics | | | |
| | (Introduction, | 18 | 2 | |
| | Genome | TOTOT | | |
| | Sequencing | 1614 | 78 | |
| 6 | Projects, Gene | | 1/20 | |
| / / | prediction and | 0 | 47 | > |
| 1 6 | counting) | | | -4 |
| 103 | | N | | 92/ |
| 17 | Practical: Cell | - Dh | / | |
| no | viability assay | | | Cu |
| | using Evan's blue | 1 | | 2 |
| | dye exclusion | | | |
| | method | | | |
| August | Unit V | 20 | 4 | 1 |
| N. | Chapter 3: | | ~ | 1 |
| 1 | Genomics, | -3 | | 1 |
| | Proteomics and | | | |
| | Bioinformatics | | 100 | |
| | (Genome | | XX | |
| | Similarity, SNPs | | | 1 |
| | and Comparative | 7.52 | 9 / | |
| | Genomics, | | | |
| | Functional | | | |
| | Genomics, | | | |
| | Proteomics, | | | |
| | History of | | | |
| | Bioinformatics; | | | |
| | Sequences and | | | |
| | nomenclature, | | | |
| | Information | | | |













| | () | | | |
|-----------|--|------|-----|-------|
| | Sources) | | | |
| | Practical: Data retrieval and database search using internet site NCBI and download a DNA and protein sequence from internet, analyse it and comment on it. Practical: Reading of a DNA sequencing gel to | लिय | | 6 |
| ho | arrive at the | - 10 | . 1 | (sh |
| | sequence. | A | | |
| September | Revision for Half- Yearly Exam | | 16 | |
| | Half-Yearly Exam | 20 | | |
| October | Unit VI | 20 | 4 | |
| | Chapter 1: Microbial Cell | V | 100 | J. J. |
| | Culture and its | M | | |
| | Applications. | | | |
| November | Unit VI | 22 | 2 | |
| | Chapter 2: Plant | | | |
| | Cell Culture and | | | |
| | Applications. | | | |
| December | Unit VI | | | |
| | Chapter 3: Animal Cell Culture and | 22 | 2 | |













| | Applications. | | | |
|----------|------------------|-------|-----------|--|
| January | | | | |
| | Revision | | | |
| | | | 16 | |
| | | | | |
| February | Revision | | | |
| | | | 12 | |
| | | | | |
| | Annual Practical | | Section 1 | |
| | Exam | TOTAL | 1 | |
| March | Annual Exam | 7 | 18 | |

Note:

• Subject teachers are advised to direct the students to prepare notebook of the Terminology/Definitional Words used in the chapters for enhancement of vocabulary or clarity of the concept.

Prescribed Books:

- 1. Biotechnology Text book for Class XII, CBSE Publication
- 2. Laboratory Manual-Biotechnology-Class XII, CBSE Publication













Question Paper Design (2024-25)

Class: XII Subject: Biotechnology Code: 871

Time 2½ Hours

| Competencies | Marks | Percentage |
|---------------|-------|------------|
| Knowledge | 24 | 40% |
| Understanding | 18 | 30% |
| Application | 12 | 20% |
| Skill | 6 | 10% |
| Total | 60 | 100% |

| Types Of Marks Questions | | Number | Description | Total Marks | |
|--------------------------|-------|--------|----------------------------|----------------|--|
| Objective | 1 | 15 | 06 Multiple Choice | 15 | |
| Questions | | 9 | Questions, | | |
| | | ~ | 03 Fill in the Blanks | | |
| 1 | | | Questions, | | |
| X. | | 1 | 03 One Word Answer Type | | |
| 1) | | 1 | Questions, | | |
| /\ | | | 0 3 Assertion Reason | | |
| | | | Questions | | |
| Very Short | 2 | 6 | Internal choice will be | 12 | |
| Answer Type | | | given in any 2 questions | | |
| Question | | 200 | -MC | | |
| Short Answer | 3 | 6 | Internal choice will be | 18 | |
| Type | | | given in any 2 questions | | |
| Question | | | | | |
| Essay Answer | 5 | 3 | Internal choice will be | 15 | |
| Type | | | given in all the questions | | |
| Question | | | | | |
| Tota | Total | | | 60 | |





