BOARD OF SCHOOL EDUCATION HARYANA

Syllabus and Chapter wise division of Marks (2025-26)Class: XIISubject: BiotechnologyCode: 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



Class-XII Subject-Biotechnology Code:871 Chapter Sr. No. Unit Marks Recombinant DNA Technology 1 Protein and Gene Manipulation Protein Structure and Engineering 30 Genomics and Bioinformatics Microbial Cell Culture and its Cell Structure and Genetic Application Manipulation 2 Plant Cell Culture and Application 30 Animal Culture Cell and Application Total 60 **Practical Examination** 20 Internal Assessment 20 **Grand Total** 100





Unit V: PROTEIN AND GENE MANIPULATION

Chapter 1: Recombinant DNA Technology

Introduction; Tools of rDNA Technology: Restriction enzymes, Restriction Fragment Length Polymorphism (RFLP), Other enzymes used in cloning, Vectors: Plasmids, vector based on bacteriophages, cosmids, YAC vectors, BAC vectors, Animal plant viral vectors, Host cells; Marking rDNA; and Introduction of rDNA to Host Cells: Transformation, Transfection, Electroporation, Microinjection, **Biolistics**: **Identification of Recombinants; Polymerase Chain Reaction** (PCR); Hybridisation Techniques: Southern Hybridisation Technique; DNA Library: DNA Sequencing: 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, Vander Wals forces, hydrophobic interactions; Structure-Function relationship in Proteins: Chymotrypsin _ a proteolytic enzyme, Molecular Disease- Sickle cell anaemia, printing-Peptide Protein Finger Mapping, 2-D Gel Electrophoresis; Purification of Proteins: Calculation of amount of bacterial ferment required, Downstream Processing, Aqueous two-phase partition, Industrial scale production of proteins, Special techniques for therapeutic /diagnostic proteins; Characterization of Proteins: Mass spectrometry; Protein Based Products: Blood products and vaccines, Therapeutic antibodies and enzymes, Therapeutic hormones and growth factors, Regulatory factors, Analytical applications, Industrial Functional non-catalytic proteins, enzymes, Nutraceutical (Protein Proteins; Designing **Proteins Engineering**): 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, Structural genomics, Functional genomics; Genome Sequencing Projects: Directed sequencing of Bacterial Artificial Chromosome (BAC) contigs, Random shotgun sequencing; Gene prediction and Genome Similarity, **SNPs** and counting: Comparative Genomics; **Functional Genomics:** Fluorescence in situ hybridization, Microarray Technology: principle, procedure, interpretation; **Proteomics:** Types of Proteomics: Expression proteomics, Structural proteomics, Functional proteomics, Genes and Proteins: Number of genes vs Number of proteins; History of 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 equipment's 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

9. Project work. (Any topic from theory syllabus)

Month wise Syllabus Teaching Plan (2025-26)

Class: XII

Subject: Biotechnology

Code: 871

Month	Subject-content	Teaching	Revision	Practical
	Co.	Periods	Periods	Work
April	Unit V Chapter 1: Recombinant DNA Technology Practical: Use of special equipment in biotechnology experiments	22	2	6
	Practical: Isolation of bacterial plasmid DNA	P	- A	2
	Practical: Detection of DNA by gel electrophoresis	ज्यो	9	2
	Practical: Estimation of DNA by UV spectroscopy			2
May	Unit V Chapter 2: Protein	22	2	

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	Engineering			
	Practical: Isolation			
	of bacteria from			2
	curd & staining of			
	bacteria			
June			1 States and a state of the sta	
	Summe	er Vacation (Pr	oject Work)
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July	Unit V		520	
	Chapter 3:		199	
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	Proteomics and			a)
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	Sequencing			
	Projects, Gene			
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	counting)		2	di.
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	Practical: Cell	2		<u> </u>
	viability assay			
	using Evan's blue	N	- XX	2
	dye exclusion			1
	method			
August	Unit V	20	4	
	Chapter 3:			
	Genomics,			
	Proteomics and			
	Bioinformatics			
	(Genome			
	Similarity, SNPs			
	and Comparative			
	Genomics,			



	Functional			
	Genomics,			
	Proteomics,			
	History of			
	Bioinformatics;			
	Sequences and			
	nomenclature,			
	Information			
	Sources)			
	Practical: Data	लिय		
	retrieval and			
	database search	0		>
	using internet site			6
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September	Revision for Half-		16	2
	Yearly Exam	Jean		
	Half-Yearly Exam			
October	Unit VI	20	4	
	Chapter 1:			
	Microbial Cell			
	Culture and its			
	Applications.			
November	Unit VI	22	2	



	Chapter 2: Plant			
	Cell Culture and			
	Applications.			
December	Unit VI			
	Chapter 3: Animal	22	2	
	Cell Culture and			
	Applications.		and the second se	
January	ONT	TAN		
	Revision	iviy	78	
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February	Revision		100	
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11	Exam	6		
March	Annual Exam			

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 (2025-26)

Class: XII

Subject: Biotechnology

Code: 871

Time 2¹/₂ Hours

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

T <mark>ypes</mark> Of	Marks	Number	Description	Tota l
Questions		\sim		<mark>Mark</mark> s
Objective	1	15	06 Multiple Choice	15
Questions	~	2	Questions,	
1			03 Fill in the Blanks	
20		18	Questions,	
No.			03 One Word Answer Type	
	1	Ç	Questions,	
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	- 22		Questions	
Very Short	2	6	Internal choice will be	12
Answer Type		1.00	given in any 2 questions	
Question			0411	
Short Answer	3	6	Internal choice will be	18
Туре			given in any 2 questions	
Question				
Essay Answer	5	3	Internal choice will be	15
Туре			given in all the questions	
Question				
Tota	ıl	30		60

