Shiv Chhatrapati Shikshan Sanstha's

## Rajarshi Shahu Mahavidyalaya, Latur

(Autonomous)



Structure and Curriculum of Two Year Post Graduate Programme with Exit option

Postgraduate Programme of Science and Technology M.Sc.in Biotechnology

**Board of Studies** 

in शिव छत्रपती

Biotechnology

Rajarshi Shahu Mahavidyalaya, Latur

(Autonomous)

(PG-II) Rajarshi Shahu Mahavidyalaya,

w.e.f. June, 2024

(In Accordance with NEP-2020)

#### **Review Statement**

The NEP Cell reviewed the Curriculum of **M.Sc. in Biotechnology** Programme to be effective from the **Academic Year 2023-24.** It was found that, the structure is as per the NEP-2020 guidelines of Govt. of Maharashtra.

**Date:** 09/08/2023

Place: Latur

#### **NEP Cell**

Rajarshi Shahu Mahavidyalaya, Latur
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#### **CERTIFICATE**

I hereby certify that the documents attached are the Bonafide copies of the Curriculum of **M.Sc. in Biotechnology** Programme to be effective from the **Academic Year 2023-24.** 

Date: 14/07/2023

Place: Latur

(Dr. Sachin Kulkarni)

Chairperson
Board of Studies in Biotechnology
Rajarshi Shahu Mahavidyalaya, Latur
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शिव छत्रपती शिक्षण संस्था लातुर

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### Members of Board of Studies in the Subject Biotechnology Under the Faculty of Science and Technology

Sr. No.	Name	Designation	In position
1	Dr. Sachin S. Kulkarni	Chairperson	HoD
	Head, Department of Biotechnology,	-	
	Rajarshi Shahu Mahavidyalaya, <mark>Latu</mark> r		
	(Autonomous)		
2	Prof. Tukaram. A. Kadam	Member	V.C. Nominee
	Professor, School of Life Sciences SRTMU,		
	Nanded.		
3	Dr. Rahul. P. Bhagat	Member	Academic Council Nominee
	Asst. Professor, Department		
	ofBiotechnology, Govt. Institute of Science,		
	Aurangabad (Autonomous)	2.6	A 1 : 0 : 1N :
4	Dr. Rajesh M. Jorgewad	Member	Academic Council Nominee
	Asst. Professor, Department of		
	Biotechnology and Bioengineering, KIT		
5	college, Kolhapur (Autonomous)  Dr. Gunderao. H. Kathwate	Member	Expert from outside for
3	Asst. Professor, Dept. of Biotech.	Member	Special Course
	S. P. P. U. Pune		Special Course
6	Mr. Abhay. M. Desai	Member	Expert from Industry
	Wockhardt, Aurangabad		3
7	Dr. SantoshNar <mark>wade</mark>	Member	P.G. Alumni
	Serum Institute Pv <mark>t.Ltd. Pu</mark> ne	।शव छः	।पता
8	Dr. Manisha. A. Dh <mark>otre</mark>	Member	Faculty Member
9	Mr. Udaybhanu. P. S <mark>i</mark> rdeshm <mark>ukh</mark>	Member	Faculty Member
10	Dr. Ravindra. B. Ade	Member	Faculty Member
11	Dr. Sanghapal. S. Kshirsagar	Member	Faculty Member
12	Mr. Suraj. <mark>D. Kadam</mark>	Member	Faculty Member
13	Mr. Akash <mark>. J. Waghmare</mark>	Member	Faculty Member
14	Miss. Swati G. Swami	Member	Faculty Member
15	Mr. Sanket M. Bansode	Member	Faculty Member
16	Miss. Karun. S. Komatwar	Member	Faculty Member
17	Dr. Kakasaheb. S. Raut	Member	Member from same Faculty

#### From the Desk of the Chairperson...

Biotechnology as a subject is a highly interdisciplinary that combines biological sciences with engineering technologies to manipulate living organisms and biological systems to produce products that advances healthcare, medicine, agriculture, food, pharmaceuticals and environment. At its simplest, biotechnology is technology based on biology - which harnesses cellular and bimolecular processes to develop technologies and products that help to improve our lives and health of our planet.

Taking into consideration of the importance of Biotechnology, Rajarshi Shahu Mahavidyalaya, Latur (Autonomous), have taken an initiative to introduce a new emerging field as an undergraduate Programme in biotechnology under the faculty of science. B. Sc. Biotechnology is a Three-year graduate degree program which is started in the academic year 2004-05 followed by the postgraduate program started in academic year 2006-07.

National Education Policy (NEP) 2020 recognizes the relevance of biotechnology in the education system due to its interdisciplinary nature, potential for research and innovation, and its alignment with the development of 21st-century skills. By integrating

and opportunities of a rapidly advancing biotechnology driven world.

NEP-2020 has conceptualized the idea to develop well rounded competent individuals for making the nation a self-reliant and global leader. In the same spirit, we at Department of Biotechnology, have developed a curriculum framework to encompass the goals of NEP 2020. In the overall curriculum we have incorporated choice of courses of study, creating academic pathways having constructive combinations with exit point as well as focus on experiential learning for students by providing multidisciplinary and holistic approach to the courses taught as major courses along with electives of choice for equipping the students with adequate knowledge leading to the choice of better career paths.

With reference to global changes occurring in higher education in various national and foreign universities, the newly designed syllabi of M.Sc. Biotechnology as per NEP 2020 guidelines are effectively implemented from June, 2023. The committee members of Board of Studies in Biotechnology also took the local need and employability of graduate students into consideration while framing the given curriculum, keeping in view of the guidelines given in the University Grants Commission, New Delhi.

By aligning curriculum development, pedagogy, interdisciplinary connections, research opportunities, industry collaborations, teacher training, and available infrastructure with the institute, the department of biotechnology plans to integrate students with a comprehensive understanding of biotechnology, foster critical thinking and research skills, and prepare them for future careers in the field.

(Dr. Sachin Kulkarni)

Chairperson

Board of Studies in Biotechnology



## (Autonomous)

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# (Autonomous) Faculty of Science and Technology

## Structure for Two Year Postgraduate Degree Programme in Biotechnology in accordance with NEP-2020

Year Level	Sem	Major		Major		RM	OJT/FP	RP	Cum. Cr	Marks	Degree
		Mandatory	Elective	RMC	NA	NA	20Cr				
	I	Major I 4Cr	MEC-I(A)	4Cr				Theory:			
		Major II 4Cr	OR					1Cr=25M			
		Major III 4Cr	MEC-I(B)						PG		
			4Cr						Diploma		
I	II	Major IV 4Cr	MEC-II(A)	NA	OJT-I 4Cr	NA	20Cr		(After 03		
6.0		Major V 4Cr	OR		/FP-I 4Cr				Year B.A.		
		Major VI 4Cr	MEC-II(B)					OJT/FP:	Degree)		
			4Cr					1Cr=25M			
	Total	Major 24Cr	MEC 08Cr	RMC 04Cr	OJT/FP 04Cr	NA	40Cr				
	III	Major VII 4Cr	MEC-III(A)	NA	NA	RP-I	20Cr				
		Major VIII 4Cr	OR			4 Cr					
		Major IX 4Cr	M <mark>EC-III(</mark> B)					RPI &	PG		
			4Cr					RPII:	Degree		
II	IV	Major X 4Cr	MEC-IV(A)	NA	NA	RP-II	22Cr	1Cr=25M	(After 03		
6.5		Major XI 4Cr	OR			6Cr		101-25141	Year UG		
		Major XII 4Cr	MEC-IV(B)						Degree)		
		1/1	4Cr						Degreej		
	Total	Major 24 Cr	MEC	NA	NA	RP	42Cr	190			
			08 Cr		_	10 Cr					
	otal of I	Major	MEC	RMC	OJT/FP	RP	40+42		82 Credits		
& II Year		48 Cr	16Cr	04Cr	04Cr	10Cr	=82 Cr				

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#### **Department of Biotechnology**

#### PG Skeleton in Accordance with NEP-2020

#### Illustrative Credit Distribution Structure for Two Year M.Sc. Degree

Year	Sem	Major	•	Lab	RM	OJT/FP	RP	Cum.	Marks	Degree
Level				Course	_			Cr		
		Mandatory	Elective		RMC	NA	NA	20Cr	Theory:	
	I	Major I 3Cr	MEC I	LC-I 1Cr	4Cr				1Cr=25M	
		Major II 3Cr	3Cr	LC-II 1Cr	1				Lab	PG
		Major III 3Cr		LC-III 1 <mark>Cr</mark>					Course:	Diploma
				LC-IV <mark>1C</mark> r					1Cr=50M	(After
I	II	Major IV 3Cr	MEC II	LC-V 1 <mark>C</mark> r	NA	OJT-I 4Cr	NA	20Cr		03 Year
6.0		Major V 3Cr	3Cr	LC-VI <mark>1C</mark> r		/FPI 4Cr				B.Sc.
		Major VI 3Cr		LC-VII <mark>1Cr</mark>					OJT/FP:	Degree)
				LC-VIII <mark>1C</mark> r					1Cr=25M	Degree
	Total	Major	MEC	LC-8Cr	RMC	OJT/FP	NA	40Cr		
	Total	18Cr	06Cr		04Cr	04Cr				
		Exit Opti	ion: PG Dip	loma wit <mark>h 40</mark>	) Credit	s After 03 Y	ear B.Sc	c. Degree		
	III	Major VII 3Cr	MEC III	LC-IX 1 <mark>C</mark> r	NA	NA	RP-I	20Cr		1
		Major VIII 3Cr	3Cr	LC-X 1Cr			4Cr			
		Major IX 3Cr	VA.	LC-XI 1Cr					RPI &	PG
				LC-XII 1Cr					RPII:	Degree
II	IV	Major X 3Cr	MEC IV	LC-XIII 1Cr	NA	NA	RP-II	22Cr	1Cr=25M	(After
6.5		Major XI 3Cr	3Cr	LC-XIV 1Cr			6Cr		101-25W	03 Year
0.5		Major XII 3Cr		LC-XV 1Cr						UG
			1	LC-XVI 1Cr						Degree)
	Total	Major 18Cr	MEC	LC-8Cr	NA	NA	RP	42Cr		Degreej
			06Cr	0			10	-		
							Cr			
Cum. 7		Major	MEC	LC-16 <mark>C</mark> r	RMC	OJT/FP	RP	40+42		82
of I & I	II Year	36Cr	12Cr		04Cr	04Cr	10Cr	=82 Cr		Credits
						11.1		37.7		
	Exit Option: Two Years 04 Sem. PG Degree with 82 Credits After 03 Year UG Degree									

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#### **Abbreviations:**

1. MMC : Major Mandatory Course

2. MEC : Major Elective Course

3. RMC : Research Methodology Course

4. OJT : On Job Training (Internship/Apprenticeship)

5. FP : Field Project

6. RP : Research Project

7. Cum. Cr : Cumulative Credit



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# Rajarshi Shahu Mahavidyalaya, Latur (Autonomous)

# (Autonomous) Department of Biotechnology M.Sc. in Biotechnology

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Level	Semester	Course Code	Course Title	Credits	No. of Hrs.				
		602BIO3101	Genetic Engineering	03	45				
		(MMC VII)							
		602BIO3104	Lab Course-IX	01	30				
		602BIO3102	Microbial Biotechnology	03	45				
		(MMC VIII)							
		602BIO3105	Lab Course-X	01	30				
		602BIO3103	Plant and Agricultural	03	45				
		(MMC IX)	Biotechnology						
	III	602BIO3106	Lab Course-XI	01	30				
		602BIO3201	Advance Pharmaceutical	03	45				
		MEC <mark>-III</mark> (A)	Biotechnology						
		Or	Or						
		602 <mark>BIO3202</mark>	Enzyme and Protein						
		MEC-III (B)	Engineering						
		602BIO3203	Lab Course-XII	01	30				
II		601BIO1301	Research Project-I	04	120				
6.5		(RP-I)							
		Total (	Credits	20					
		602BIO4101	Animal and Livestock	03	45				
		(MMC X)	Biotechnology	200					
		602BIO4104	Lab Course-XIII	01	30				
		602BIO41 <mark>02</mark>	Clinical Research, IPR,	03	45				
		(MMC XI)	Bio entrepreneurship and						
	5.5		Start up						
	IV	602BIO4105	Lab Course-XIV	01	30				
	Daia	602BIO4103	Food and Nano	03	45				
	Raja	(MMC XII)	Biotechnology	ıya,					
		602BIO4106	Lab Course-XV	01	30				
		602BIO4201	Environmental	03	45				
		MEC-IV (A)	Biotechnology						
		Or	Or						
		602BIO4202	Omics Technology						

		MEC-IV(B)			
		602BIO4203	Lab Course-XIV	01	30
		(RP-II)	Research Project-II	06	180
		Total (	Credits	22	
Total Credits (Semester III & IV)					42



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#### **Faculty of Science & Technology**

I	Programme Outcomes (POs) for M.Sc. Programme					
PO 1						
PO 2						
PO 3						
PO 4						
PO 5						
PO 6						
PO 7						



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# Rajarshi Shahu Mahavidyalaya, Latur (Autonomous)

Progr	amme Specific Outcomes (PSOs) for M.Sc. Biotechnology
PSO No.	Upon completion of this programme the students will be able to -
PSO 1	Integrate basic principles of common analytical techniques of protein
	molecular structures to engage in hands-on practices for implementation
	of such techniques to facilitate the development of biopharmaceutical
	manufacturing.
PSO 2	Induce the understandings of basic principles of process units' operations
	of industrial products with hands-on practices for implementation of such
	techniques to facilitate the development of biopharmaceutical
	manufacturing.
PSO 3	Gain fundamental knowledge of molecular biotechnology, protein
	expression, and structural biology for the development of new products
	having clinical application.
PSO 4	Plan, condu <mark>ct, execute and write-up a pr</mark> op <mark>osal o</mark> f original research
	Practical skills.
PSO 5	Promote the entrepreneurship for self-growth and sustainability with the
	aim of promoting lab to land practices in, clinical, agriculture, food, nano,
	plant and animal biotechnology.
PSO 6	Integrate fundamental concepts of leadership, entrepreneurship and
	innovation, financial decision making and marketing to business
	enterprises.
PSO 7	Equip the students with the skills required for carrying out research in
	cutting edge areas of life sciences.
PSO 8	Make the students competent for dealing with the future problems and
	challenges of regional and global interest in overall development of society.

Latur (Autonomous)

## Semester - Third



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# (Autonomous) Department of Biotechnology

**Course Type: MMC-VII** 

**Course Title: Genetic Engineering** 

Course Code: 602BIO3101

Credits: 03 Max. Marks: 75 Lectures: 45Hrs.

#### **Learning Objectives**

- LO 1 To comprehend principles and techniques for DNA and RNA isolation, quantification
- LO 2 To master various DNA sequencing methods (Maxam-Gilbert, Sanger-Nicolson, Pyrosequencing) including principles, applications, advantages, and limitations.
- LO 3 To master the utilization of restriction endonucleases and DNA modifying enzymes for genetic engineering
- LO 4 To develop expertise in employing prokaryotic hosts and vectors, including plasmid vectors and bacteriophages, for genetic manipulation
- LO 5 To develop skills in the selection, screening, and analysis of recombinants using various techniques
- LO 6 To master the application of molecular biology techniques Polymerase Chain Reaction (PCR), (RFLP), RAPD, AFLP used for genetic analysis and identification of recombinants.
- LO 7 To develop proficiency in vector engineering and codon optimization techniques, along with host engineering strategies, to enhance gene expression in higher organisms.
- LO 8 To master the diverse strategies of gene delivery and expression in various organisms

#### **Course Outcomes**

- CO 1 Proficiently isolate and quantify nucleic acids
- CO 2 Apply understanding of DNA sequencing methods to analyze DNA sequences effectively
- CO 3 Apply advanced knowledge of genetic engineering tools
- CO 4 Demonstrate proficiency in constructing genomic and cDNA libraries
- CO 5 Demonstrate proficiency in employing a variety of techniques like Northern blotting, Southern blotting, Western blotting, PCR, RFLP, RAPD, AFLP screening of recombinants.

- CO 6 Apply advanced knowledge of molecular biology methods to analyze and interpret data obtained from screening processes
- CO 7 Demonstrate proficiency in utilizing vector engineering, codon optimization, and host engineering.
- CO 8 Apply advanced knowledge of gene expression mechanisms in bacteria, yeast, insects, insect cells, mammalian cells, and plants.

Unit No.	Title of Unit & Contents	Hrs.
I	DNA isolation and Sequencing methods.	9
	<ol> <li>Isolation of DNA and RNA.</li> <li>Quantification of nucleic acids.</li> <li>Radiolabeling of nucleic acids: End labeling, nick translation, labeling by primer extension.</li> <li>DNA sequencing: Maxam- Gilbert (Chemical) and Sanger-Nicolson (dideoxy/ enzymatic) sequencing method, Pyrosequencing, NGS.</li> </ol>	
	Unit Outcomes:	
	UO 1 Proficiently isolate and quantify nucleic acids UO 2 Apply understanding of DNA sequencing methods to analyze DNA sequences effectively	
II	Tools of Genetic Engineering	11
	<ol> <li>Nucleases, Restriction endonucleases: Types of restriction endonucleases, classification, and uses.</li> <li>Restriction mapping. DNA modifying enzymes: Polymerases, Phosphatases, and DNA ligases.</li> <li>Prokaryotic host.</li> <li>Plasmid vectors, Bacteriophage, T7 expression vectors, pET vector system.</li> <li>Construction of genomic and cDNA libraries</li> <li>Joining of DNA Fragments to vectors, Homopolymer tailing, cohesive and blunt end ligation, adaptors, and linkers.</li> <li>Unit Outcomes:</li> <li>U0 1 Apply knowledge of restriction endonucleases in molecular biology techniques such as DNA cloning, PCR, and DNA fingerprinting.</li> <li>U0 2 Proficiency in the design and utilization of plasmid vectors, bacteriophages, and T7 expression vectors for gene cloning and expression.</li> </ol>	

Unit No.	Title of Unit & Contents	Hrs.
III	Screening and Selection of Recombinants.	12
	1. Selection, screening, and analysis of recombinants.	
	2. Principle of hybridization. Northern blotting, Southern	
	blotting, Western blotting.	
	3. Polymerase chain reaction	
	4. Restriction fragment length polymorphism, RAPD,	
	AFLP, Map Construction.	
	Unit Outcomes:	
	UO 1 Able to discrimination between recombinant and non-recombinant clones.	
	UO 2 Explain the concept genetic analysis, gene expression studies, and disease diagnosis through precise molecular detection methods	
IV	Expression of gene in higher organism	13
	1. Vector Engineering and codon optimization, host engineering.	
	2. Strategie <mark>s of g</mark> ene delivery, <i>in vitro</i> translation,	
	3. Expressi <mark>on in bacteria, yeast, insects, an</mark> d insect cells.	
	Expres <mark>sion in mammalian cells and plan</mark> t	
	4. Chromosome engineering,	
	5. Targeted gene replacement, gene editing, gene	
	regulation & silencing.	
	Unit Outcomes:	
	UO 1 Apply specificity of gene expression through optimized	
	ve <mark>ctor de</mark> sign an <mark>d cod</mark> on usage.	
	UO 2 App <mark>ly advanced kn</mark> owledge of targeted delivery of	
	genetic materi <mark>al int</mark> o host cells for various applications	
	such as gene therapy and genetic engineering.	

### **Learning Resources:**

- 1. Principles of Gene Manipulation, R.N. Old & S.B. Primrose, Blackwell Publishing, 1994
- 2. From Genes to Clones, E.L. Winnaeker, Wiley VCH, 1987
- 3. Recombinant DNA, J.D. Watson, J. Witkowski, M. Gilman, & M. Zoller, W.H. Freeman & Co. Ltd., 1992

- 4. An Introduction to Genetic Engineering, D.S.T. Nicholl, Cambridge University Press, 2008
- 5. Molecular Biotechnology, J.J. Pasternak, American Society for Microbiology, 1996
- 6. The Biochemistry of Nucleic Acid, Adam et al., Springer, 1992
- 7. Genetic Engineering, J.K. Setlow, Springer, 2005
- 8. Molecular Cloning: A Laboratory Manual, J. Sambrook & D.W. Russell, Cold Spring Harbor Laboratory Press, 2012
- 9. Principles of Gene Therapy, T. Wirth & N. Parker, Humana Press, 2014
- 10. Genetic Engineering: Principles and Methods, J.K. Setlow & V.P. Setlow, Springer, 2016





#### (Autonomous)

#### **Department of Biotechnology**

**Course Type: Lab Course** 

Course Title: Lab Course -IX (Based on MMC-VII)

Course Code: 602BIO3104

Credits: 01 Max. Marks: 50 Hours: 30

#### **Learning Objectives**

- LO 1 To demonstrate proficiency in isolating nucleic acid
- LO 2 To master the technique of endonuclease digestion of nucleic acid
- LO 3 To understand the principles and procedures involved in thermal melting of DNA.
- LO 4 To acquire skills in isolating plasmid DNA through both mini preparation and large-scale isolation methods.
- LO 5 To develop expertise in performing in vitro DNA ligation and transformation of E. coli bacteria.
- LO 6 To gain proficiency in DNA blotting techniques and DNA hybridization.
- LO 7 To demonstrate competence in isolating cytoplasmic RNA and performing electrophoresis on denaturing gels.
- LO 8 To master techniques such as blotting, PCR/RFLP, along with their computational analysis.

#### **Course Outcomes**

- CO 1 Analyze chromatin structure through determination of mono-nucleosome size and chromatin gel electrophoresis.
- CO 2 Gain competence in analyzing DNA fragments following endonuclease digestion through agarose gel electrophoresis.
- CO 3 Understand the principles and procedures involved in thermal melting of dna.
- CO 4 Gain mastery in isolating plasmid DNA via both mini preparation and large-scale isolation methods.
- CO 5 Gain ability to perform in vitro DNA ligation and successfully transform e. Coli bacteria.
- CO 6 Adopt skill in employing DNA blotting and hybridization techniques effectively.
- CO 7 Become proficient in isolating cytoplasmic RNA and conducting electrophoresis on denaturing gels.
- CO 8 Receive competence in various advanced techniques such as northern blotting, nucleic acid sequencing, and PCR/RFLP analysis, including computational analysis and use current biochemical techniques to plan and carry out experiments.

Practical No.	Unit
1.	Isolation of nuclei and analysis of chromatin- i) determination of mononucleosome size ii) chromatin gel electrophoresis
2.	Endonuclease digestion of nuclei and analysis of DNA fragments by Agarose gel electrophoresis
3.	Thermal melting of DNA
4.	Isolation of plasmid DNA-i) mini preparation ii) large-scale isolation
5.	In vitro DNA ligation, t <mark>rans</mark> formation of E.coli.
6.	Techniques: a) DNA blotting technique b) DNA hybridization.
7.	Isolation of cytopla <mark>smic RNA</mark> .
8.	Electrophoresis of R <mark>NA on dena</mark> turing gels.
9.	Northern blotting tec <mark>hnique.</mark>
10.	Separation of poly A+RNA on oligo-dT column
11.	cDNA synth <mark>esis</mark> and cloning.
12.	RNA hybridization-dot and northern blots.
13.	In situ detection of RNA in embryos/tissue.
14.	In vitro translation.
15.	Nucleic acid Sequencing and its computational analysis. 16. PCR/RFLP technique.
16.	Isolation of nuclei and analysis of chromatin- i) determination of mononucleosome size ii) chromatin gel electrophoresis
17.	Endonuclease dig <mark>estio</mark> n of nuclei and analysis of DNA fragments by Agarose gel electrophoresis
18.	Thermal melting of DNA
19.	Isolation of plasmid DNA-i) mini preparation ii) large-scale isolation
20	In vitro DNA ligation, transformation of E.coli.
	tajarshi Shahu Mahavidyalaya,

N.B.: Any Ten Practicals from above.



# (Autonomous) Department of Biotechnology

**Course Type: MMC-VIII** 

**Course Title: Microbial Biotechnology** 

Course Code: 602BIO3102

Credits: 03 Max. Marks: 75 Lectures: 45 Hrs.

#### **Learning Objectives**

- LO 1 To understand the principles of microbial production and its significance in generating biochemicals, pharmaceuticals, and industrial enzymes.
- LO 2 To explore the various methods involved in the microbial production of organic acids, solvents, biofuels, and amino acids.
- LO 3 To examine the processes and techniques used in the microbial synthesis of vitamins, antibiotics, and recombinant products.
- LO 4 To analyze the environmental applications of microbes.
- LO 5 To investigate the production, immobilization techniques, and commercial applications of industrial enzymes.
- LO 6 To explore biotransformation reactions and their applications in transforming steroids, sterois, and nonsteroid compounds.
- LO 7 To evaluate the role of microbial biotechnology in sustainable production processes and its impact on various industries.
- LO 8 To critically assess the challenges and advancements in microbial production techniques, considering factors such as yield optimization, substrate utilization, and downstream processing.

#### Course outcomes

- CO 1 Demonstrate a comprehensive understanding of microbial production processes and their applications in generating biochemicals, pharmaceuticals, and industrial enzymes.
- CO 2 Acquire practical skills in microbial cultivation, fermentation techniques, and downstream processing necessary for the production of target compounds.
- CO 3 Be proficient in analyzing and optimizing microbial production systems to enhance yield, efficiency, and product purity.
- CO 4 Gain insights into the environmental implications of microbial biotechnology and its role in waste management, pollution control, and sustainable resource utilization.
- CO 5 Develop expertise in biocatalysis and its potential applications in various industries.
- CO 6 Equipped with the knowledge and skills required to work in biotechnology, pharmaceuticals, environmental science, and related fields, contributing to research, development, and innovation.

- CO 7 Capable of critically evaluating scientific literature, patents, and technological advancements in microbial production, fostering a culture of continuous learning and innovation.
- CO 8 Prepared to address contemporary challenges in microbial biotechnology, such as bioprocess optimization, strain engineering, and bioreactor design, contributing to the advancement of the field.

Unit No.	Title of Unit & Contents	Hrs.		
I	Microbial Production of Biochemicals	11		
	1. Organic acids Produc <mark>tion:</mark> Citric acid, Lactic acid, Acetic			
	acid			
	2. Solvents and Biofu <mark>els: Bu</mark> tanol, Ethanol, Brewing Industry			
	overview			
	3. Amino acids Prod <mark>uction:</mark>			
	Methods of Production			
	• Production of L-Glutamic acid, L Lysine, L-			
	Tryptophan			
	Unit Outcomes:			
	UO 1 Demonstrate a comprehensive understanding of organic			
	acid and amino acid production			
	UO 2 Gain insigh <mark>t i</mark> nto th <mark>e brewing industry a</mark> nd Proficiency in Solvents and Biofuels			
II	Microbial Production of Pharmaceuticals	10		
11	1. Vitamins Production: Vitamin B12, Riboflavin	10		
	2. Antibiotics Production: Beta-Lactam Antibiotics (e.g.			
	Penicillin), Carbohydrate Antibiotics (e.g. Streptomycin),			
	Tetracycline, Nucleoside Antibiotics, Aromatic Antibiotics			
	3. Recombinant Products:			
	Production of Hepatitis B Vaccine			
	Insulin Production			
	Erythropoietin Production			
	Unit Outcomes:			
	UO 1. Get acquainted with production process of vitamins and			
	antibiotics			
	UO 2. Develop expertise in the production of recombinant			
	pharmaceuticals.			
III	Environmental Applications of Microbes	10		
	1. Biomethanation			
	2. Bioleaching: Mechanism and Examples			
	3. Biosorption and Microbial Recovery of Petroleum			
	(MEOR)			
	4. Microbial Polysaccharides: Xanthan and Dextran			
	5. Production of Biopolymers and Bio-pesticides			
	Unit Outcomes:			
	UO 1. Understand the principles of Biomethanation, Bioleaching			
	and Biosorpion process			
	UO 2. Analyze the applications and environmental benefits of			

Unit No.	Title of Unit & Contents	Hrs.
	biopolymers and bio-pesticides	
IV	Microbial Enzyme Production and Biotransformation	13
	1. Enzyme Immobilization Techniques	
	2. Commercial Applications of Microbial Enzymes	
	3. Production of Key Microbial Enzymes: Amylases, Glucose	
	Isomerase, L-Asparaginase, Proteases, Pectinases and	
	Lipases	
	4. Biotransformation Reactions	
	5. Types of Bioconversio <mark>n Re</mark> actions	
	6. Microbial Transform <mark>ation</mark> of Steroids and Sterols	
	7. Microbial Transfor <mark>mation</mark> of Nonsteroid Compounds:	
	L-Ascorbic Acid	
	Prostaglandins	
	Antibiotics	
	Unit Outcomes:	
	UO 1. Understand the pr <mark>inciples and tech</mark> niques involved in	
	enzyme immobiliza <mark>tion and its applic</mark> ations in various	
	industries.	
	UO 2. Analyze the commercial significance of microbial enzymes	
	and their applications in different sectors, such as food,	
	pharma <mark>ceuticals, and biotechnology.</mark>	

#### **Learning Resources:**

- 1. Industrial Microbiology: An Introduction, Michael J. Waites, Neil L. Morgan, and John S. Rockey, Wiley-Blackwell, 2001.
- 2. Microbial Production of Biopolymers and Polymer Precursors: Applications and Perspectives" edited by Alane Beatriz Vermelho and Luciana Porto de Souza Vandenberghe, CRC Press, 2020.
- 3. Biotechnology: A Textbook of Industrial Microbiology, W. Crueger and A. Crueger, Sinauer Associates, 1989.
- 4. Bioprocess Engineering: Basic Concepts, Shuler, M.L., Kargi, F., Prentice Hall, 2001.
- 5. Biochemical Engineering Fundamentals, James E. Bailey and David F. Ollis McGraw-Hill Education, 1986.
- 6. Industrial Biotechnology: Principles and Applications, Christoph Wittmann and James C. Liao, Wiley-VCH, 2016.
- 7. Industrial Microbiology, 1st Edition, Casida, J.R., L.E., Willey Eastern Ltd, New Delhi, 2006.
- 8. Industrial Microbiology, 4<sup>th</sup> Edition, Prescott and Dunn, CBS Publishers, New Delhi, 1987.
- 9. Principles of Fermentation Technology, Stanbury, P.F., and Whitaker, A., 2<sup>nd</sup> edition Pergamon Press, Oxford,2005

- 10. Biotechnology. U Satyanarayana. Uppala Author Publisher Interlinks, Vijaywada, India, 2005.
- 11. Microbial Technology, Peppler & Perlman. Vol- I, II Academic Press
- 12. Basic Biotechnology, Bu'Lock J. and Kristansen B. (Eds), Academic Press Inc Ltd, London,1987.
- 13. Manual of Industrial Microbiology and Biotechnology, Demain A.L., Davies J.E. (Ed in Chief) ASM, 2nd Edition Washington, USA.,1999.
- 14. Industrial Microbiology, G. Reed, 4<sup>th</sup> edition, CBS Publishers (AVI Publishing Co.), 2020.
- 15. Comprehensive Biotechnology, Cooney & Humphery. Vol-3. Pergamon Press,
- 16. Text Book of Biotechnology, , H.K Das, 3rd edition, Willey India, 2008
- 17. Industrial Microbiology, A.H Patel, 2nd edition, Macmillan Publication, 2011.





#### (Autonomous)

#### **Department of Biotechnology**

**Course Type: Lab Course** 

Course Title: Lab Course -X (Based on MMC-VIII)

Course Code: 602BIO3105

Credits: 01 Max. Marks: 50 Hours: 30

#### **Learning Objectives**

- LO 1. To develop practical skills in the production processes of industrial products using microbial biotechnology techniques, including fermentation and cultivation methods.
- LO 2. To gain proficiency in quantitative analysis methods for assessing the yield, purity, and activity of microbial biotechnology products.
- LO 3. To acquire knowledge and proficiency in isolation techniques for microbial strains and biotechnological products.
- LO 4. To understand the principles and applications of purification techniques specific to microbial biotechnology, including chromatography and filtration methods.
- LO 5. To learn to optimize upstream processes such as media formulation, inoculum preparation, and bioreactor operation for efficient product synthesis.
- LO 6. To enhance understanding of downstream processing techniques for product recovery, purification, and characterization in microbial biotechnology.
- LO 7. To develop problem-solving skills by troubleshooting issues encountered during practical exercises and experiments in microbial biotechnology.
- LO 8. To foster teamwork and collaboration through group-based laboratory activities, promoting effective communication and cooperation among peers.

#### **Course outcomes**

- CO 1. Demonstrate competency in executing the production of industrial products using microbial biotechnology techniques, adhering to safety protocols and good laboratory practices.
- CO 2. Apply optimization strategies to both upstream and downstream processes, maximizing the yield, quality, and efficiency of microbial biotechnology products.
- CO 3. Perform quantitative analysis of microbial biotechnology products accurately, interpreting experimental data and drawing meaningful conclusions.
- CO 4. Execute isolation techniques effectively, obtaining pure cultures and biotechnological products suitable for further analysis and application.
- CO 5. Implement purification techniques to obtain highly purified microbial biotechnology products, meeting industry standards and regulatory requirements.

- CO 6. Successfully troubleshoot challenges encountered during production, isolation, and purification processes, demonstrating problem-solving abilities.
- CO 7. Collaborate effectively with peers in laboratory settings, contributing to a positive and productive learning environment.
- CO 8. Demonstrate proficiency in documenting experimental procedures, results, and observations accurately and systematically, adhering to scientific principles and protocols in microbial biotechnology research and practice.

Practical	Unit
No.	
1	Production of sauerkraut by Microorganisms
2	Production and quant <mark>itation</mark> of Antibiotics
3	Production and estim <mark>ation of lact</mark> ic acid by Lactobacillus Sp.
4	Production of ferment <mark>ed milk by Lact</mark> obacillus acidophilus.
5	Comparison of ethano <mark>l production usi</mark> ng various Organic wastes /raw
	Material
6	Laboratory sc <mark>ale</mark> prod <mark>uction of biofertilizers</mark>
7	Production a <mark>nd estimation of Bacterial Lipas</mark> e
8	Production and Estimation of Bacterial and Fungal Amylase
9	Production and estimation of alkaline protease from bacterial source
10	Produce bioplastics (e.g., polyhydroxyalkanoates) using microbial
	fermentation of renewable feedstocks and characterize the properties of
	the bioplastics.
11	Evaluate the ability of microorganisms to degrade environmental
	pollutants (e.g., hydrocarbons, heavy metals) using batch or continuous
	bioreactor systems
12	Production and isolation of bacterial exo-polysaccharides
13	Perform quality control tests on microbial products (e.g., antibiotics,
	enzymes) including microbial enumeration, purity testing, and potency
	assays Shahu Mahavidyalaya
14	Visit to Fermentation Industry

N.B.: Any Ten Practicals from above.



#### (Autonomous)

#### **Department of Biotechnology**

**Course Type: MMC IX** 

**Course Title: Plant and Agricultural Biotechnology** 

**Course Code: 602BIO3103** 

Credits: 03 Max. Marks: 75 Lectures: 45 Hrs.

#### **Learning Objectives**

- LO 1. To understand the advancements in the field of biotechnology with respect to plants.
- LO 2. To follow modern techniques and their applications in crop improvements, such as tissue culture, plant breeding, and transgenics.
- LO 3. To understand the modern trends in plant & agricultural biotechnology.
- LO 4. To understand pathological aspects of plant disease.
- LO 5. Analyze the applications of plant biotechnology in crop improvement, such as disease resistance, herbicide tolerance, and improved nutritional content.
- LO 6. Apply molecular biology tools and techniques for plant genetic analysis and manipulation.
- LO 7. Analyze the applications of plant biotechnology in crop improvement, such as disease resistance, herbicide tolerance, and improved nutritional content.
- LO 8. Evaluate the ethical, social, and environmental implications of genetically modified crops and biotechnological interventions in agriculture.

#### **Course outcomes**

- CO 1 Analyze the origins of cultivated plants and the concept of vavilov's centres of origin in plant diversity.
- CO 2 Explore the functions and importance of botanical gardens and herbaria in plant research, conservation, and education.
- CO 3 Understand the molecular basis of infection and disease resistance/defense mechanisms in plants.
- CO 4 Evaluate the physiological aspects of parasitism in plants and examine control measures, including the use of fungicides and other management strategies.
- CO 5 Understand the processes involved in the development of male and female gametophytes in plants and their significance in sexual reproduction
- CO 6 Explore the formation of somatic hybrids and cybrids and their potential applications in crop improvement and genetic manipulation.

- CO 7 Understand the principles and techniques involved in plant breeding, including introduction, selection, and hybridization methods such as pedigree, backcross, mass selection, and bulk method
- CO 8 Discuss the development of transgenic crops, their benefits, and the biosafety aspects associated with their cultivation and consumption.

Unit No.	Title of Unit & Contents	Hrs.
I	Plant Resource Development	10
	•	
	<ol> <li>Domestication and introduction of plants.</li> <li>Origin of cultivated plants, Vavilov's centres of</li> </ol>	
	origin.	
	3. Plants as sources for food, fodder, fibres, spices, beverages,	
	edible oils, drugs,	
	narcotics, insecticides, timber, gums, resins, and dyes;	
	4. latex, cellulose, starch, and its	
	products; 5. Energy planta <mark>tio</mark> ns;	
	6. Botanical Gar <mark>dens</mark> an <mark>d Herbaria</mark>	
	Unit Outcomes:	
	UO 1. Analyze the origins of cultivated plants and the concept	
	of Vavilov's centres of origin in plant diversity.	
	UO 2. Explore the functions and importance of botanical	
	gardens and herbaria in plant research, conservation,	
	and education.	
II	Plant Pathology	10
	1. crop diseases cau <mark>sed b</mark> y viruses, bacteria, mycoplasma,	
	fungi, and nematodes;	
	2. Modes of infection and dissemination; Molecular basis of	
	infection and disease resistance/	
	defense;	
	3. Physiology of parasitism and control measures. Fungal	
	toxins. Latur (Autonomous)	
	4. Modelling and disease forecasting;	
	5. Plant quarantine.	

Unit No.	Title of Unit & Contents	Hrs.
	Unit Outcomes:	
	UO 1. Understand the molecular basis of infection and	
	disease resistance/defense mechanisms in plants.	
	UO 2. Evaluate the physiological aspects of parasitism in	
	plants and examine control measures, including the use	
	of fungicides and othe <mark>r ma</mark> nagement strategies.	
III	Plant Biotechnology	12
	1. Development of male and female gametophytes.	
	2. Endosperm—its deve <mark>lopment</mark> and function.	
	3. Patterns of embryo de <mark>velopment; P</mark> olyembryony, apomixis;	
	4. Pollen haploids,	
	5. embryo rescue method <mark>s and their applica</mark> tions.	
	6. Protoplast cu <mark>ltu</mark> re.	
	7. Somatic hybrids and Cybrids;	
	8. Micropropag <mark>ation; Somaclonal variation and</mark> its	
	application <mark>s</mark>	
	Unit Outcomes:	
	UO 1 Understand the processes involved in the development of male and female gametophytes in plants and their significance in sexual reproduction  UO 2 Explore the formation of somatic hybrids and cybrids and their potential applications in crop improvement and genetic manipulation.	
IV	Agricultural Biotechnology	13
	1. Methods of plant breeding—introduction, selection and	
	hybridization (pedigree, backcross, mass selection, bulk	
	method);	
	2. Mutation, polyploidy, male sterility, and	
	heterosis breeding.	
	3. Use of apomixis in plant breeding	
	4. Genetic engineering—methods of transfer of genes;	
	5. Transgenic crops and biosafety aspects;	

Unit No.	Title of Unit & Contents	Hrs.
	6. Development and use of molecular markers in plant	
	breeding;	
	Unit Outcomes:	
	UO 1. Understand the principles and techniques involved in	
	plant breeding, including introduction, selection, and	
	hybridization methods such as pedigree, backcross,	
	mass selection, an <mark>d bulk</mark> method	
	UO 2. Discuss the devel <mark>opmen</mark> t of transgenic crops, their	
	benefits, and the <mark>biosafety</mark> aspects associated with their	
	cultivation and co <mark>nsumption.</mark>	

#### **Learning Resources:**

- 1. Biotechnology and Genomics, P.K. Gupta, Rastogi Publications, Meerut, India, 2004
- 2. Transgenic Plants: A Production System for Industrial and Pharmaceutical Proteins, M.R.L. Owen & J. Pen (Eds), John Wiley & Sons, England, 1996
- 3. Agricultural Biotechnology, S.S. Purohit, Agro Botanica, India, 1999
- 4. Plant Cell Biotechnology, R. Endress, Springer Verlag, Germany, 1994
- 5. Textbook of Modern Plant Pathology, K.S. Bilgrami & H.G. Dube, Vikas Publications, New Delhi, 1994
- 6. Genetics and Biotechnology in Crop Improvement, P.K. Gupta, Rastogi Publications, Meerut, 1998
- 7. Fundamentals of Plant Pathology, V.N. Pathak, N.K. Khatri, & M. Pathak, Agrobotanical Publications, Bikaner, 1996
- 8. General Microbiology, Vol. II, C.B. Powar & H.F. Daginawala, Himalaya Publishing House, Mumbai, 1990
- 9. Agricultural Biotechnology, S.S. Purohit, Agrobios India, Jodhpur, 2002
- 10. Biotechnology, U. Satyanarayana, Books and Allied Pvt. Ltd., Kolkata, 2007
- 11. Biofertilizer and Organic Farming, S.C. Vyas, S. Vyas, S. Vyas, & H.A. Modi, Akta Prakashan, Nadiad, G.S., Meerut, 1998
- 12. Experiments in Microbiology, Plant Pathology, Tissue Culture and Mushroom Cultivation, S.C. Vyas, S. Vyas, S. Vyas, & H.A. Modi, Vishwa Prakashan, New Age International (p) Ltd., New Delhi, 1998
- 13. Microbiology and Biotechnology: A Laboratory Manual, P.T. Kalaichelvan & P.C. Dandiya, MJP Publishers, Chennai, 2004
- 14. Laboratory Manual of Plant Biotechnology, S.S. Purohit & K.R. Aneja, Agrobotonical Pub. India, 1995

- 15. Methods in Biotechnology, Hans Peter Schmauder, Taylor and Francis, London, 1997
- 16. Methods in Plant Molecular Biology, 1st edition, M.A. Schuler & R.E. Zielinski, Academic Press, 1989
- 17. Methods in Biotechnology and Bioengineering, S.P. Vyas & D.V. Kohli, CBS Publishers and Distributors, New Delhi, 2002





#### (Autonomous)

#### **Department of Biotechnology**

**Course Type: Lab Course** 

Course Title: Lab Course -XI (Based on MMC IX)

Course Code: 602BIO3106

Credits: 01 Max. Marks: 50 Hours: 3

#### **Learning Objectives**

- LO 1 Acquire proficiency in aseptic techniques essential for maintaining sterile conditions in plant tissue culture laboratories, ensuring contamination-free cultures and reliable experimental outcomes.
- LO 2 Develop competence in preparing culture media tailored to specific plant tissue culture requirements, understanding the importance of nutrient composition and pH in supporting plant growth and development.
- LO 3 Gain insight into the principles and techniques of micropropagation, mastering the skills necessary for mass production of genetically uniform plants from a small explant, facilitating rapid multiplication of elite plant varieties.
- LO 4 Acquire skills in isolating bacterial and fungal plant pathogens from diseased plant tissues, enhancing knowledge of plant pathology and disease management strategies.
- LO 5 Understand the impact of different culture media formulations on plant growth and development, exploring the role of nutrients, hormones, and growth regulators in influencing plant physiology.
- LO 6 Engage in plant breeding experiments to study genetic variability, heritability, and trait inheritance patterns, fostering an understanding of plant breeding principles and techniques for crop improvement.
- LO 7 Develop proficiency in identifying bacterial, fungal, and viral plant diseases through morphological, physiological, and molecular techniques, enabling accurate disease diagnosis and management.
- LO 8 Enhance observational and analytical skills through visits to botanical gardens, gaining firsthand exposure to diverse plant species, habitats, and conservation efforts.

#### **Course outcomes**

After completion of the course, the student will be able to-

CO 1 Demonstrate proficiency in managing plant tissue culture laboratories, adhering to strict aseptic protocols and safety measures to ensure successful culture establishment and maintenance.

- CO 2 Formulate culture media suitable for various stages of plant tissue culture, optimizing nutrient concentrations and hormonal balances to support desired growth and development patterns.
- CO 3 Master micropropagation techniques, capable of efficiently propagating plants on a large scale for commercial and research purposes, contributing to crop production and genetic conservation efforts.
- CO 4 Develop skills in isolating and identifying bacterial, fungal, and viral plant pathogens, enabling accurate disease diagnosis and the development of effective disease management strategies.
- CO 5 Critically analyze experimental data from plant breeding experiments and disease identification assays, drawing conclusions and making recommendations based on scientific evidence.
- CO 6 Integrate theoretical knowledge with practical skills acquired through hands-on experiments and field visits, enhancing their understanding of agricultural and plant biotechnology concepts.
- CO 7 Recognize the importance of ethical considerations and sustainability in agricultural biotechnology, applying principles of responsible research conduct and environmental stewardship in their work.
- CO 8 Develop communication skills to effectively communicate scientific findings and collaborate with peers and professionals in the field of agricultural and plant biotechnology, fostering interdisciplinary teamwork and knowledge exchange.

Practical No.	Unit
1	Plant tissue culture laboratory design
2	Aseptic techniques
3	Media preparation
4	Micro propagation
5	Anther culture
6	Isolation of bacterial plant pathogens
7	Isolation of fungal plant pathogens
8	Effect of media on plant growth
9	Plant breeding experiments.
10	Visit botanical gardens.
11	Plant DNA isolation
12	Protoplast isolation
13	Synthetic seed preparation
14	Identification of bacterial diseases.
15	Identification of fungal diseases.
16	Identification of viral diseases.
17	Seed culture

18 Embryo Culture

N.B.: Any Ten Practicals from above.





# (Autonomous) Department of Biotechnology

**Course Type: MEC-III (A)** 

**Course Title: Advanced Pharmaceutical Biotechnology** 

Course Code: 602BIO3201

Credits: 03 Max. Marks: 75 Lectures: 45 Hrs.

#### **Learning Objectives**

- LO 1. Understand the principles of chemotherapy and the mechanisms of action of antimicrobial agents.
- LO 2. Identify and analyze the various mechanisms of microbial resistance to antibiotics and antimicrobial agents.
- LO 3. Classify antibiotics based on their chemical structures and mechanisms of action, with examples illustrating their therapeutic applications.
- LO 4. Describe the general characteristics of secondary metabolites, categorizing them based on their types and highlighting their medicinal applications.
- LO 5. Explain the structure, mechanism of action, and therapeutic applications of antibacterial, antifungal, antiviral, and anticancer drugs.
- LO 6. Evaluate drug discovery and development processes, including historical perspectives, molecular biology techniques, and combinatorial drug discovery approaches.
- LO 7. Discuss the principles of rational drug design and drug targeting, exploring strategies to optimize drug efficacy and minimize adverse effects.
- LO 8. Analyze the stability of drugs, pharmacokinetic and pharmacodynamic properties.

#### **Course outcomes**

- CO 1. Develop proficiency in identifying and selecting appropriate chemotherapeutic agents for the treatment of infectious diseases.
- CO 2. Apply knowledge of antibiotic classification and mechanisms of action in clinical decision-making, ensuring optimal patient care and antimicrobial stewardship.
- CO 3. Demonstrate an understanding of drug discovery and development processes.
- CO 4. Acquire skills in evaluating the efficacy, safety, and stability of drugs through preclinical and clinical trials.
- CO 5. Interpret the results of preclinical toxicity testing, including acute, sub-acute, and chronic toxicity studies.
- CO 6. Demonstrate knowledge of biosimilar technology and its implications in pharmaceutical development and regulatory approval processes.

- CO 7. Familiarize with Indian, international pharmacopoeias, and global regulatory guidelines, ensuring compliance with quality standards and regulatory requirements in drug manufacturing and marketing.
- CO 8. Critically analyze clinical trial phases and their significance in drug development, recognizing the ethical and regulatory considerations involved in human experimentation and patient safety.

Unit No.	Title of Unit & Contents	Hrs.
I	Chemotherapeutic Agents and Antibiotics	10
•	<ol> <li>Chemotherapy Antimicrobial Drug. Mechanism of action of antimicrobial agents.</li> <li>Microbial Resistance to antibiotics and antimicrobial agents (Types and Mechanism).</li> <li>Types of Antibiotics: Classification of antibiotics with example.</li> <li>General characteristics of an Secondary Metabolites: Types and Medicinal Applications</li> <li>Unit Outcomes:         <ul> <li>U0 1.Demonstrate a comprehensive understanding of chemotherapeutic agents and antibiotics in the management of infectious diseases.</li> <li>U0 2. Identifying different types of resistance and understanding the implications of resistance in clinical practice and public</li> </ul> </li> </ol>	10
	health	
II	<ol> <li>Antibacterial, Antifungal and Antiviral Drugs</li> <li>Chemotherapeutics Agents Structure, Mechanism of Action and Applications of Antibacterial drug: Sulfonamides, Quinolones.</li> <li>Antiviral drug: Amantadine, Azidothymidine.</li> <li>Antifungal drug: Nystatin, Griseofulvin.Mechanism of action of</li> <li>Anticancer drugs, Drugs acting on CNS, Insulin, Blood factor VIII.</li> <li>Unit Outcomes:         <ul> <li>U0 1 Understanding the structure, mechanism of action, and therapeutic applications of specific antibacterial, antifungal, and antiviral drugs.</li> <li>U0 2 Develop critical thinking skills to assess the potential benefits and limitations of different antibacterial, antifungal, and antiviral drugs.</li> </ul> </li> </ol>	10
III	Drug Discovery, Development and Targetting	13

Unit No.	Title of Unit & Contents	Hrs.
	1. Discovery and Development of drug, History, drug	
	targeting, Molecular Biology and Combinatorial drug	
	discovery, Rational Drug designing.	
	2. Stability of Drug,	
	<ol><li>Pharmacokinetics, Pharmacodynamics.</li></ol>	
	4. Drug delivery systems, Liposomes.	
	Unit Outcomes:	
	UO 1. Grasp the principles of drug discovery, development, and	
	targeting to the design and optimization of novel	
	therapeutic agents.  UO 2. Adept at applying molecular biology techniques and	
	rational drug design approaches to expedite the	
	development of targeted drugs.	
IV	Clinical Trials, Regulatory Guidelines and Pharmacopoeia	12
	1. Clinical Trials Phas <mark>es of Clinical tri</mark> als of drugs, Preclinical	
	drug evaluation of its biological activity, potency and	
	toxicity-	
	2. Toxicity te <mark>st</mark> in ani <mark>mals including acut</mark> e, sub-acute and	
	chronic to <mark>xici</mark> ty, E <mark>D50 and LD50 determ</mark> ination, special	
	toxicity te <mark>st like teratogenecity and mutag</mark> enecity.	
	3. Biosimilar Technology, Introduction to Indian,	
	4. International Pharmacopoeia and global regulatory	
	guide <mark>lines</mark>	
	Unit Outcomes:	
	UO 1. Gain a comprehensive understanding of the various phases of clinical trials, enabling them to design and	
	implement clinical research studies.	
	UO 2. Develop proficiency in navigating regulatory frameworks	
	and pharmacopoeia standards, ensuring adherence to	
	quality control measures and ethical considerations in	
	pharmaceutical development and marketing.	

शारोह तमसो ज्योतिः॥ Rajarshi Shahu Mahavidyalaya, Latur (Autonomous)

- 1. Pharmaceutical Microbiology, W. B. Hugo & A. D. Russell, Wiley India, 2021
- 2. Pharmacology and Pharmacobiotechnology, Ashutosh Kar, New Age, 2017
- 3. Essentials of Pharmaceuticals, FSK Barar, S.Chand, 2000
- 4. Molecular Biotechnology, B. Glick & J. Pasernak, ASM Press, 2020
- 5. Drug Designing, Doble, McGraw Hill, 2011
- 6. Pharmaceutical Biotechnology, S.P. Vyas & Dixit, CBS, 2007
- 7. Medicinal Chemistry, B. Razdan, CBS, 2018
- 8. Pharmacology and Pharmacotherapeutics, Satoskar & Bhandarkar, Popular, 2020
- 9. Pharmaceutical Biotechnology, Purohit & Saluja, Student Edition, 2003
- 10. Biotechnology: Secondary Metabolites, K.G. Ramawat & J.M. Merillon, Oxford, 2011
- 11. Chemistry of Natural Products, R.H. Thomson (Ed.), Springer, 1993
- 12. Biopharmaceuticals, S.N. Jogdand, Himalaya Publishing, 2006





# (Autonomous)

#### **Department of Biotechnology**

**Course Type: Lab Course** 

Course Title: Lab Course XII (Based on MEC -III (A))

Course Code: 602BIO3203

Credits: 01 Max. Marks: 50 Hours: 30

#### **Learning Objectives**

- LO 1. Understand the principles and techniques involved in biological assay for estimating penicillin/streptomycin.
- LO 2. Gain proficiency in chemi<mark>cal ass</mark>ay methods including titration and spectrophotometric techniques, and interpretation of analytical data.
- LO 3. Learn the procedures for assessing the antimicrobial activity of antibiotics through agar diffusion or broth dilution methods.
- LO 4. Develop skills in determining the Minimum Inhibitory Concentration (MIC) of antibiotics against specific microbial strains using broth dilution or agar dilution methods.
- LO 5. Acquire knowledge and techniques for determining the shelf life of antibiotics.
- LO 6. Gain practical experience in conducting sterility testing of commercial pharmaceutical products.

#### **Course outcomes**

After completion of the course, the student will be able to-

- CO 1. Demonstrate proficiency in performing biological and chemical assays for the estimation antibiotics.
- CO 2. Interpret MIC data accurately to guide clinical treatment decisions and optimize antibiotic dosing regimens for effective microbial control.
- CO 3. Evaluate the shelf life of antibiotics based on stability studies, ensuring product integrity and compliance with regulatory standards for pharmaceutical products.
- CO 4. Demonstrate competence in conducting sterility testing of pharmaceuticals, adhering to regulatory guidelines and ensuring product safety and efficacy.
- CO 5. Gain insights into pharmacopeial standards and global regulatory guidelines in the pharmaceutical industry, understanding their importance in ensuring product quality, safety, and efficacy.

Rajarshi Shahu Mahavidyalaya, Latur (Autonomous)

Practical	Unit
No.	
1	Estimation of penicillin/streptomycin by biological assay.
2	Estimation of penicillin/streptomycin by chemical assay.
3	Determination of Minimum Inhibitory Concentration (MIC) of Antibiotic
4	Determination of shelf life of antibiotics (Expired drugs)
5	Sterility testing of commercial pharmaceuticals.
6	Study of microbial spoilage of pharmaceuticals.
7	Sterility testing of injectable as per IP.
8	Effect of chemical disi <mark>nfectant</mark> on growth of bacteria
9	Study of Pharmacop <mark>eia and glob</mark> al regulatory guidelines in pharma
	industry
10	Study of dug action by using Zebra fish (Danio rerio) as model organism
11	Visit to Pharmaceutical industry

N.B.: Any Ten Practicals from above.



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# Rajarshi Shahu Mahavidyalaya, Latur

# (Autonomous)

#### **Department of Biotechnology**

Course Type: MEC I (B)

**Course Title: Enzyme Technology and Protein Engineering** 

Course Code: 602BIO3202

Credits: 03 Max. Marks: 75 Lectures: 45 Hrs.

# **Learning Objectives**

LO 1 To study the concept of classification of enzymes.

LO 2 To understand the role of enzymes in various sectors.

LO 3 To acquaint the knowledge about role of entropy in catalysis.

LO 4 To study the concept of allosteric enzymes with their applications.

LO 5 To study Enzyme Kinetics.

LO 6 To know the stabilization of biphasic aqueous-organic systems.

LO 7 To understand the kinetics of Immobilization & Protein Engineering.

LO 8 To learn the concept of Recombinant proteins.

#### **Course Outcomes**

After completion of the course, the student will be able to-

- CO 1. Describe structure, functions and the mechanisms of action of enzymes.
- CO 2. Learn kinetics of enzyme catalyzed reactions and enzyme inhibitory.
- CO 3. Perform immobilization of enzymes.
- CO 4. Get exposure of wide applications of enzymes and their future potential.
- CO 5. Acquire knowledge of significance of vmax and km.
- CO 6. Acquaint the knowledge of turnover number and end point kinetic assay.
- CO 7. Gain the knowledge about enzyme kinetics in biphasic liquid systems.
- CO 8. Understand the concept of biosensors.

Rajarshi Shahu Mahavidyalaya, Latur (Autonomous)

Unit No.	Title of Unit & Contents	Hrs.
I	Introduction To Enzymes & Enzyme Kinetics	12
	<ol> <li>The Enzyme-Introduction,</li> <li>Nomenclature of enzymes,</li> <li>Classification of enzymes,</li> <li>Applications in Industrial, Medical, Analytical, Chemical, Pharmaceutical and Food Sectors.</li> </ol>	
II	Unit Outcomes:  U0 1 Describe the nomenclature of enzymes.  U0 2 Discuss the Applications of enzymes in Industrial and Medical field.  Enzyme Kinetics	10
	1. Michaelis - Menten equation, Brigg's- Haldane equation,	
	2. Graphical procedures in enzymology	
	3. Advantag <mark>es</mark> and d <mark>isadvantages of altern</mark> ate plotting,	
	4. Estimatio <mark>n o</mark> f constants using graph <mark>ical technique,</mark>	
	Kinetics for reversible reactions, basics of enzymatic reaction,	
	5. Collision theory and transition state theory	
	6. Role of entropy in catalysis, presteady state kinetics,	
	7. Significance of Vmax and Km,	
	8. Kinetics of multi- substrate reactions,	
	9. Allosteric enzymes – The Monad – Changeux – Wyman model (MCW) and The Koshland – Nemethy – Filmer (KNF) model,	
	10. Enzyme inhibition - types of inhibitors- Mode of action and experimental determination.	
	11. Enzyme activity, international units, specific activity, turnover number, end point kinetic assay.	
	Unit Outcomes:	
	UO 1 Explain the advantages and disadvantages of alternate plotting.	

Unit No.	Title of Unit & Contents	Hrs.
	UO 2 Discuss the concept of enzyme inhibition and their	
	classification.	
III	Effect of Physical Factors & Enzyme Kinetics in Biphatic	10
	Reaction	
	1. Temperature dependence of rate constants of enzymatic	
	reaction,	
	2. Thermal deactivation, pH	
	3. Effect on rate con <mark>stants</mark> and protein structure.	
	4. pH dependence: i <mark>onizatio</mark> n of Acids and Bases.	
	5. Enzyme kinetics i <mark>n biphasic</mark> liquid systems,	
	6. Stabilization of bip <mark>hasic aqueou</mark> s-organic systems,	
,	7. Equilibria in bipha <mark>sic aqueous- orga</mark> nic systems.	
	Unit Outcomes:	
	UO 1. Understand the concept of enzyme kinetics in biphasic liquid systems.	
	UO 2. Explain the concept of temperature dependence of rate constants of enzymatic reaction.	
IV	Enzyme Immobilization, Kinetics of Immobilization & Protein Engineering	13
	1. Immobilization of Biocatalysts an Introduction,	
	2. Electrostatic Effect, effect of charged and uncharged	
	support, Kinetics of immobilized enzymes	
	3. Effect of external and internal mass transfer,	
	4. Damkohler number, effectiveness factor, Intraparticle	
	diffusion kinetics, Biot number.	
	5. Biosensors - glucose oxidase, cholesterol oxidase, urease	
	and antibodies as biosensors,	
	6. Introduction to protein engineering,	
	7. Structure prediction sequence structure relationship.	
	8. Recombinant proteins using fusion protein strategies for	
	enhanced recovery,	

Unit No.	Title of Unit & Contents	Hrs.
	7. Engineering protein for the affinity purification,	
	(engineering of streptavidin)	
	8. Stabilization of enzymes by protein engineering (e.g.	
	pseudomonas isoamylase)	
	Unit Outcomes:	
	UO 1 Explain the concept of protein engineering.	
	UO 2 Understand the concept of stabilization of enzymes by	
	protein engineeri <mark>ng.</mark>	

- 1. Biochemical Engineering Fundamentals, Bailey JE, Ollis, DF, McGraw Hill Education, 2nd edition, 2017.
- 2. Biochemical Engineering, Marcel Decker Blanch HW and Clark DS, University of California, 1995.
- 3. Bioreaction Engineering, modeling and control, Schugerl K., Bellgart KH (Eds): Springer-Verlag, Berlin, 2000.
- 4. Enzymes, palmer, East west publication, 2008.
- 5. Handbook of Enzyme Biotechnology, 3rd Edition, Wiseman, A: Ellis Horwood Publication, 1995.
- 6. Bioprocess technology, kinetics and reactors, Moser, A: Springer Verlag, 2011.
- 7. Biochemical Engineering Principles and functions, Syed Trnveer Ahmed Inamdar, PHI Learning Private limited, 2012.
- 8. Protein and enzyme engineering, Saurabh Bhatia, IOP Publishing Ltd, 2018.
- 9. Protein Engineering: Tools and Applications, Huimin Zhao, WILEY-VCH GmbH, 2021.
- 10. Enzyme Technology, Ashok Pandey, Springer Science & Business Media, 2006.



# (Autonomous)

#### **Department of Biotechnology**

**Course Type: Lab Course** 

Course Title: Lab Course -XII (Based on MEC-III (B)

Course Code: 602BIO3203

Credits: 01 Max. Marks: 50 Hours: 30

#### **Learning Objectives**

LO 1 To provide hands on isolation of high yielding microbial strains for the commercially important enzyme production.

- LO 2 To train students for standardizing medium composition for enzyme production.
- LO 3 To introduce students for the development of enzyme assay methods.
- LO 4 To provide hands on gel filtration method for determination of molecular weight.
- LO 5 To determine enzyme activity and specific activity.
- LO 6 To make understand to analyse the method of checking the purity of the enzymes by using SDS-PAGE.

#### **Course Outcomes**

- CO 1 Learn the kinetics of enzyme catalyzed reactions and enzyme inhibitory and regulatory process.
- CO 2 Perform immobilization of enzymes.
- CO 3 Get exposure of wide applications of enzymes and their future potential
- CO 4 Carry out enzyme isolation and purification protocols.

Practical No.	Unit
1.	Isolation of high yielding microbial strains for the production of commercially important enzymes.
2.	Production of commercially important enzymes from microbial sources.
3.	Standardization of medium composition for the optimum production of enzymes.
4.	Determination of enzyme activity and specific activity.

5.	Partial purification of isolated enzymes.
6.	Characterization of enzymes-Effect of pH, temperature, and inhibitors on enzyme activity etc.
7.	Molecular weight determination of enzyme by Gel filtration method.
8.	Method of checking the purity of the enzyme -SDS-PAGE
9.	Immobilization of enzymes –Different Techniques such as adsorption, entrapment, encapsulation and cross- linking.
10.	Strain improvement techniques- physical, chemical and genetic manipulation methods.
11.	Development of enz <mark>yme ass</mark> ay methods
12.	Formulation of enzyme stability.

N.B.: Any Ten Practicals from above.



# Semester - Fourth



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Rajarshi Shahu Mahavidyalaya, Latur (Autonomous)



# (Autonomous) Department of Biotechnology

**Course Type: MMC-X** 

**Course Title: Animal and Livestock Biotechnology** 

Course Code: 602BIO4101

Credits: 03 Max. Marks: 75 Lectures: 45 Hrs.

# **Learning Objectives:**

- LO 1 To Understand the basic principles of cell culture, including cell types, growth requirements, and culture techniques.
- LO 2 To Learn and practice sterile techniques necessary for cell culture to prevent contamination and maintain cell health.
- LO 3 To Understand the composition of cell culture media and learn to prepare different types of media suitable for specific cell types.
- LO 4 To Learn to operate and maintain cell culture equipment such as incubators, biosafety cabinets, and microscopes.
- LO 5 To Development of affordable new generation vaccines and diagnostics against major diseases of livestock, dogs and poultry
- LO 6 To Study assisted reproductive technologies (ARTs) like artificial insemination, in vitro fertilization (IVF), and cloning, and their applications in animal breeding and conservation.
- LO 7 To Learn about techniques such as gene cloning, PCR, DNA sequencing, and gene editing (e.g., CRISPR/Cas9) and their applications in animals.
- LO 8 To Genome analysis and genetic characterization of indigenous livestock breeds

#### **Course Outcomes:**

- CO 1 Acquire a solid understanding of the principles and techniques of animal cell culture, including cell biology, sterile technique, and culture maintenance.
- CO 2 Develop proficiency in a variety of cell culture techniques, including cell line establishment, maintenance, and manipulation, as well as cell counting and viability assessment.
- CO 3 Learn to design and execute cell culture experiments, including the selection of appropriate cell lines, culture conditions, and assays.
- CO 4 Understand and practice safe laboratory techniques, including proper handling and disposal of hazardous materials.
- CO 5 Gain knowledge about the genetics of livestock species, including breeding strategies, genetic diversity, and the role of genomics in livestock improvement.

- CO 6 Learn and practice biotechnological techniques used in livestock research and production, such as gene editing, cloning, and transgenic animal production.
- CO 7 Understand assisted reproductive technologies (arts) used in livestock breeding, including artificial insemination, embryo transfer, and in vitro fertilization.
- CO 8 Learn how biotechnology can improve livestock production efficiency, including traits related to growth rate, feed efficiency, and environmental adaptation.

Unit No.		Hrs.
I	Basics of Animal Cell Culture	12
	<ol> <li>Introduction to Animal Cell Culture, Planning and Layout of animal tissue culture laboratory.</li> <li>Cell Culture Media and Reagents: Types, Growth supplements, serum free media, balanced salt solution, other cell culture reagents.</li> <li>Tissue Culture Techniques: Primary and secondary culture, continuous cell lines, suspension culture, Tissue Specific Culture and organ culture.</li> <li>Behaviour of cells in culture conditions, their growth pattern, cell metabolism, estimation of cell number.</li> <li>Cell Line Development and Maintenance.: Establishment, Characterization, Maintenance and Contamination Control</li> </ol>	
	Unit Outcomes:	
	<ul><li>UO 1. To understand basics of animal cell culture including laboratory setup.</li><li>UO 2. To understand the process of media preparation, cell</li></ul>	
	handing, cell characterization and contaminants in animal cell culture.	
II	Scale-Up and Commercial Applications of Animal Biotechnology	10

Rajarshi Shahu Mahavidyalaya, Latur (Autonomous)

Unit No.		Hrs.
	<ol> <li>Cell culture reactors; Scale-Cell Culture Reactors and Scale-Up Techniques.</li> <li>Advanced Bioreactor Systems: Rotating chambers, Perfused suspension cultures, Fluidized bed reactors and their role in suspension culture, Scale-up in monolayers.</li> <li>Multi Surface propagators Multiarray disks, spirals and tubes, Roller culture; Microcarriers, Perfused monolayer cultures; Membrane perfusion.</li> <li>Hollow fiber perfusion; Matrix perfusion; Microencapsulation, Growth monitoring</li> <li>Transfection and transformation of cells; Commercial scale production of animal cells, Application of animal cell culture for in vitro testing of drugs.</li> <li>Testing of toxicity of environmental pollutants in cell culture; Application of cell culture technology in production of human and animal viral vaccines and pharmaceutical proteins.</li> <li>Unit Outcomes:</li> <li>U0 1. To understand industrial level applications of cell culture and related processes in culturing cells.</li> </ol>	
111	UO 2. To understand applications of animal tissue culture in industry and human welfare.  Molecular Techniques and Applications in Livestock Biotechnology	10
III		10
	<ol> <li>Genome organization of any suitable Livestock with example</li> <li>Physical and genetic map, current status of genome maps of livestock.</li> <li>Marker Assisted Selection (MAS), Polymerase Chain Reaction (PCR), its types and applications.</li> <li>Molecular markers and its applications - RFLP, RAPD, AFLP, Microsatellite/ Minisatellite markers, SNP markers.</li> <li>DNA fingerprinting. DNA sequencing, Genome sequencing, Genomic Library, Genomics database of Livestock. gene editing (e.g., CRISPR/Cas9)</li> <li>Unit Outcomes:</li> </ol>	
	<ul><li>UO 1 To understand the concept of marker assisted selection in Livestock biotechnology.</li><li>UO 2 To understand molecular marker study and its applications in research.</li></ul>	

Unit No.		Hrs.
IV	Breeding Techniques and Transgenic Technology in Livestock	13
	1. History of development of important breeds of dairy	
	cattle.	
	2. Methods of cross breeding and its types, assisted	
	reproductive technologies (ARTs) like artificial	
	insemination, in vitro fertilization (IVF), and cloning,	
	and their applications in animal breeding and	
	conservation	
	3. milk quality and p <mark>roductio</mark> n efficiency, Transgenesis and	
	methods of gene t <mark>ransfer in a</mark> nimals.	
	4. Statistical techniq <mark>ues for analyzi</mark> ng molecular genetic	
	data, Quantitative Trait Loci (QTL) mapping and its	
	applicati <mark>on</mark> in anim <mark>al breeding</mark>	
	5. Genome scan, Candidate gene approach, Genomic	
	selection in livestock, Applications of transgenic	
	technology in livestock improvement and molecular	
	biopharming.	
	Unit Outcomes:	
	UO 1. To understand livestock breeding and its quality	
	improvement by the application of biotechnology.  UO 2. To understand QTL mapping and its applications in livestock biotechnology.	

- 1. Culture of Animal Cells: A Manual of Basic Technique and Specialized Applications, Freshney, R. Ian, Wiley-Blackwell, Seventh Edition, 2016.
- 2. Animal Cell Culture and Technology, Butler, M., & Griffiths, B, Taylor & Francis, Second Edition, 2004.
- 3. Basic Cell Culture Protocols, Helgason, C., & Miller, C., Springer, Fourth Edition, 2013.
- 4. Principles of Tissue Engineering, Lanza, R., Langer, R., & Vacanti, J., Academic Press, Fourth Edition, 2013.

- 5. Animal Cell Biotechnology: Methods and Protocols, Pörtner, R., & Zeng, A. P., Humana Press, Third Edition, 2007.
- 6. Livestock Biotechnology: Breeding, Genetics, and Genomics, Abdullah, M., & Salim, H. M., Springer, Second Edition, 2020.
- 7. Biotechnology in Animal Husbandry, Pandey, A. K., & Singh, R. K., Daya Publishing House, First Edition, 2016.
- 8. Livestock Production and Biotechnology, Cai, Y., Nova Science Publishers, First Edition, 2017.
- 9. Applications of Biotechnology in Animal Health and Production, Maqbool, A., & Singh, S. P., Springer, First Edition, 2019.
- 10. Animal Biotechnology: Models in Discovery and Translation, Bhan, S., & Singh, S., CRC Press, First Edition, 2018.





# (Autonomous) Department of Biotechnology

**Course Type: Lab Course** 

Course Title: Lab Course-XIII (Based on MMC-X)

Course Code: 602BIO4104

Credits: 01 Max. Marks: 50 Hours: 30

# **Learning Objectives**

- LO 1 To master sterile techniques for handling cells and media to prevent contamination and maintain cell viability.
- LO 2 To Learn to prepare cell culture media with the appropriate nutrients and supplements for different cell types.
- LO 3 To Practice techniques for passaging and sub-culturing cells to maintain healthy cultures and promote cell growth.
- LO 4 To Understand the principles and theory behind MAS, including the role of genetic markers in livestock breeding.
- LO 5 To Become proficient in genotyping techniques used for molecular marker analysis, such as PCR, DNA sequencing, and genotyping arrays.
- LO 6 To Learn to analyze molecular marker data, including marker-trait association analysis and genomic selection.

#### **Course Outcomes**

After completion of the course, the student will be able to-

CO 1 demonstrate proficiency in maintaining sterility during cell culture procedures to prevent contamination.

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- CO 2 Prepare cell culture media with the appropriate components and concentrations for specific cell types.
- CO 3 Successfully maintain animal cell cultures, including regular passaging and subculturing, to promote cell growth and viability.
- CO 4 ability to select appropriate molecular markers (e.g., SNPs, microsatellites) for MAS based on their linkage to traits of interest in specific livestock species.

- CO 5 Proficiency in genotyping techniques used for molecular marker analysis, such as PCR, DNA sequencing, and genotyping arrays.
- CO 6 ability to analyze molecular marker data, including marker-trait association analysis and genomic selection, using statistical software.

Practical No.	Unit
1	Packing and sterilization of glass and plastic wares for cell culture.
2	Preparation of reagents and media for cell culture.
3	Primary culture technique for chicken embryo fibroblast.
4	Secondary culture of chicken embryo fibroblast.
5	Cultivation of continuous cell lines.
6	Quantification of cells by trepan blue exclusion dye.
7	Isolation of lymphocytes and cultivation of lymphocytes
8	Study of effect of toxic chemicals on cultured mammalian cells
9	Study of effect of virus on mammalian cells.
10	Suspension culture technique
11	Cryopreservation of cell primary cultures and cell lines
12	Practicals based on genomic databases of livestock and its analysis.
13	A study of molecular markers by using RFLP and AFLP.

N.B.: Any Ten Practicals from above.





#### (Autonomous)

#### **Department of Biotechnology**

**Course Type: MMC XI** 

Course Title: Clinical Research, IPR, Bio-entrepreneurship and Start Up

Course Code: 602BIO4102

Credits: 03 Max. Marks: 75 Lectures: 45 Hrs.

#### **Learning Objectives:**

- LO 1. To develop experts or skilled professionals to handle large clinical data procedure with correct guidelines
- LO 2. To acquire a basic understanding of the concepts and practices of clinical trials in pharmaceutical industry.
- LO 3. To enable students to understand issues surrounding the risks and benefits of drugs.
- LO 4. To encourage the students for start up development.
- LO 5. To understand the basic concepts of pharmacology.
- LO 6. To get the knowledge about toxicology studies.
- LO 7. To learn about the basic concepts of IPR.
- LO 8. To understand the process of data management and safety monitoring in clinical trials.

#### **Course Outcomes:**

- CO 1. Extend understanding good manufacturing practices and good clinical practices.
- Co 2. Understand the principles of good laboratory practices.
- Co 3. Know the new drug development process.
- Co 4. Conduct the clinical trials of nascent drugs.
- Co 5. Know safety monitoring and reporting in clinical trials.
- Co 6. Understand the regulatory and ethical requirements in clinical research.
- Co 7. Conduct the clinical trials of nascent drugs.
- Co 8. Acquire the knowledge about stages of pharmacokinetics and pharmacodynamics.

Unit No.		Title of Unit & Contents	Hrs.
I	Int	roduction to Clinical Research	10 Hrs.
	1.	Introduction to Drug Discovery and Drug development	
	2.	Introduction to Clinical Research Industry	
	3.	Types of Clinical research	
	4.	Phases of clinical research	
	5.	Drug development process	
	6.	Manufacturing of drugs and Good Manufacturing Practices (GMP)	

Unit No.	Title of Unit & Contents	Hrs.
	7. Toxicology: Mutagenicity, teratogenicity and carcinogenicity, Systemic toxicology (Single dose and	
	repeat dose toxicity studies)	
	Unit Outcomes:	
	UO 1. Extend understanding of drug ddiscovery and drug	
	development	
	UO 2. Describe the types & phases of clinical research.	
II	<ul><li>IPR</li><li>1. Basic concepts of Intellectual Property rights</li></ul>	12 Hrs.
	1 7 6	
	2. Evolution of ethics in clinical research	
	3. Ethics and Ethical Guidelines for Clinical Trials and Good	
	Clinical Practice (GCP)	
	4. Human rights in clinical research	
	5. Principles of Good Laboratory Practices	
	6. Good Manufacturing Practices & Good Clinical Practices.	
	7. Types of clinical trials, single blinding, double blinding	
	8. Open access	
	9. Randomized trials and their examples preclinical studies	
	10. Concepts and Application in clinical trials	
	11. Quality Assurance and Quality Control in Clinical Trials	
	Unit Outcomes: UO 1. Understand the ethics and ethical guidelines for clinical trials	
	and good clinical practice.	
	UO 2. Understand the quality assurance and quality control in	
	clinical trials.	
III	General Pharmacology	13 Hrs.
	1. Introduction, definitions and scope of pharmacology	
	2. Routes of administration of drugs	
	3. New drug discovery process	
	4. New Drug Application and Approval.	
	5. Pharmacokinetics (absorption, distribution, metabolism and	
	excretion)	
	6. Pharmacodynamics, stages of pharmacodynamics.	
	Unit Outcomes:	

Unit No.	Title of Unit & Contents	Hrs.
	UO 1. Understand the different routes of drug administration.	
	UO 2. Describes the concept of pharmacokinetics and	
	pharmacodynamics.	
IV	Bio-entrepreneurship	10 Hrs.
	1. Concept of Bio-entrepreneurship	
	2. Scope for biotechnology students	
	3. Bio- entrepreneurship Importance	
	4. Steps of Bio-entrepreneurship development	
	5. Data Management in cl <mark>inical R</mark> esearch	
	6. Safety monitoring in cl <mark>inical tri</mark> als.	
	7. Clinical Trial Start up activities: Site Feasibility Studies, Pre-	
	study visit, Site initiatio <mark>n visit.</mark>	
	Unit Outcomes: UO 1. Understand the steps of bio-entrepreneurship development.	
	UO 2. Gain the knowledge about various clinical trial start up	
	activities.	

- 1. Handbook of Clinical Research, Julia Lloyd and Ann Raven, Churchill Livingstone Publications, 1994
- 2. Principles of Clinical Research, Giovanna di Ignazio & Di Giovanna, CRC Press / BSP Books, 2018
- 3. Basic and Clinical Pharmacology, B.G. Katzung, Prentice Hall International, 2021
- 4. Textbook of Clinical Trials, David Machin, Simon Day, & Sylvan Green, John Wiley and Sons, 2005
- 5. Basic Principles of Clinical Research and Methodology, S.K. Gupta, JPB Publication, 1st edition, 2007
- 6. Handbook of Good Clinical Research Practice, World Health Organization, 2005
- 7. Fundamentals of Clinical Trials, Lawrence M. Friedman & Curt, 5th edition, 2015
- 8. Designing Clinical Research, Dr. Stephen B Hulley, MD, MPH, Steven R Cummings, MD, Warren S Browner, MD, 4th edition, 2013
- 9. Biotechnology Entrepreneurship: Starting, Managing, and Leading Biotech Companies, Craig Shimasaki, Academic Press, 1st Edition, 2008
- 10. Good Clinical Practice: A Question & Answer Reference Guide, Susanne Prokscha, SAGE Publications, Inc, 2nd Edition, 2017



# (Autonomous)

#### **Department of Biotechnology**

**Course Type: Lab Course** 

Course Title: Lab Course XIV (Based on MMC-XI)

Course Code: 602BIO4105

Credits: 01 Max. Marks: 50 Hours: 30

#### **Leaning Objectives:**

LO 1 To identify and describe various techniques for collecting biological specimens.

LO 2 To recognize commonly used instruments in experimental pharmacology.

LO 3 To gain knowledge of laboratory animals.

LO 4 To learn about types of preclinical experiments.

LO 5 To understand standard operating protocols in clinical research.

LO 6 To explore the use of anesthetics in laboratory animals.

LO 7 To learn how to apply for a patent.

LO 8 To gain insights into starting a startup.

#### **Course Outcomes:**

- UO 1 Describe and differentiate various techniques used for specimen collection.
- UO 2 Identify & describe commonly used instruments in experimental pharmacology.
- UO 3 Get the knowledge about knowledge of laboratory animal.
- UO 4 Apply knowledge of experimental design principles to plan and studies effectively.
- UO 5 Describe about various routes of drug administration.
- UO 6 Gain insights into the principles and practice of anaesthesia and analgesia in laboratory animals.
- UO 7 Acquire knowledge of the essential steps and considerations involved in starting a startup.
- UO 8 Develop skills in methods of collection of specimens.

Practical No.	Unit
1	To study different Techniques of specimen collection.
2	Commonly used instruments in experimental pharmacology.
3	Study of laboratory animals.
4	Types of preclinical experiments.
5	Techniques of blood collection from animals.
6	Standard operating protocols in clinical research/trails.
7	Study of different routes of drugs administration.
8	Study of use of anesthetics in laboratory animals.

9	How to apply for a patent.
10	How to start a Start Up.
11	Visit to Biotechnology industry/ Research Institute.

N.B.: Any Ten Practicals from above.





#### (Autonomous)

#### **Department of Biotechnology**

**Course Type: MMC-XII** 

Course Title: Food and Nano Biotechnology

**Course Code: 602BIO4103** 

Credits: 03 Max. Marks: 75 Lectures: 45 Hrs.

# **Learning Objectives:**

- LO 1. To understand the fundamental principles of biotechnology and its applications in food production.
- LO 2. To describe the various biotechnological processes involved in the synthesis of food ingredients, flavors, colors, and sweeteners.
- LO 3. To analyze the importance of food safety measures, including the Hazard Analysis and Critical Control Points (HACCP) system, in ensuring the quality and safety of food products.
- LO 4. To evaluate different methods of food preservation and their effectiveness in extending shelf life and maintaining food quality.
- LO 5. To apply genetic engineering techniques to modify microorganisms and plants for enhanced food production and nutritional quality.
- LO 6. To investigate the role of traditional fermentation processes in food production and their relevance in modern bioprocessing techniques.
- LO 7. To recognize the potential of nanobiotechnology in improving food quality, safety, and sustainability.
- LO 8. To develop critical thinking and problem-solving skills to address challenges and opportunities in the field of biotechnology for food ingredients.

#### Course Outcomes:

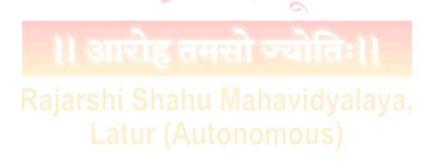
- CO 1 Demonstrate a comprehensive understanding of biotechnological principles and their applications in the food industry.
- CO 2 Equipped with the knowledge and skills necessary to synthesize a wide range of food ingredients, flavors, colors, and sweeteners using biotechnological approaches.
- CO 3 Implement food safety protocols, including haccp, to ensure the production of safe and high-quality food products.
- CO 4 Select and apply appropriate food preservation methods based on product characteristics and storage requirements.
- CO 5 Demonstrate proficiency in utilizing genetic engineering techniques to enhance the nutritional content and functional properties of food ingredients.
- CO 6 Gain insight into traditional fermentation processes and their integration with modern bioprocessing technologies for food production.
- CO 7 Aware of the potential applications of nanobiotechnology in addressing challenges related to food quality, safety, and sustainability.

CO 8 Contribute to the advancement of biotechnological innovations in the food industry through critical thinking, problem-solving, and research skills.

Unit No.	Title of Unit & Contents	Hrs.
I	Biotechnology in Food Production	10 Hrs.
	<ol> <li>Metabolic Engineering of Bacteria for Food Ingredients</li> <li>Biotechnology of Microbial Polysaccharides in Food</li> <li>Microbial Biotechnology for Food Flavor Production</li> <li>Food Safety: Introduction to HACCP System and Food Protection</li> <li>Responsibility for Ensuring Food Safety</li> <li>Food Additives: Definition, Types, and Functional Characteristics</li> <li>Natural Colors in Food: Types and Applications</li> <li>Sweeteners: Types and Applications</li> <li>Causes of Food Spoilage</li> <li>Food Preservation Methods</li> <li>Unit Outcomes:         <ul> <li>U0 1 Understand and Apply Biotechnology in Food Production.</li> </ul> </li> </ol>	
	UO 2 Ensure and Evaluate Food Safety and Quality	
II	Food Applications and Functional Foods  1. Solid State Fermentations for food applications	12 Hrs.
	<ol> <li>Genetic Engineering of baker's yeast</li> <li>Biotechnology of wine yeast</li> <li>Genetic Modification of Plant Oils for Food uses</li> <li>Biotechnology of -carotene from Dunaliella</li> <li>SCP: Spirulina and Chlorella</li> <li>Biotechnological approaches to improve nutritional quality and shelf life of fruits and vegetables</li> <li>Unit Outcomes:         <ul> <li>U0 1 Understand Biotechnological Processes in Food Production.</li> <li>U0 2 Apply Biotechnological Principles in Food Industry</li> </ul> </li> </ol>	
III	Food Safety, Traditional Fermentations and Novel	13 Hrs.
	Bioprocessing  1. Molecular Evolution and Diversity of food borne pathogens.	
	2. Application of Microbial Molecular Techniques for food	
	systems.	

Unit No.	Title of Unit & Contents	Hrs.
	3. Application of ELISA assays for detection and quantitation	
	of toxins in foods and E.coli in food	
	4. Biosensors for food quality assessment	
	5. Biotransformation applicable to food industries	
	6. Fermentation technology for traditional food of the Indian	
	subcontinent	
	7. Functional foods: Conc <mark>ept</mark> of Prebiotics, Probiotics and	
	Nutraceuticals.	
	Unit Outcomes: UO 1 Proficiency in Molecular Techniques and Assays for	
	Food Safety.	
	UO 2 Comprehensive understanding of functional foods and	
	their compone <mark>nts and Abil</mark> ity to apply	
	biotransf <mark>or</mark> mation processes in food industries.	
IV	Nano Biotechnology	10 Hrs.
	1. Introduction to Nano-Biotechnology	
	2. The Nanoscale Dimension and Paradigm	
	3. Types of Nanomaterials and Their Classifications, Structures	
	and Characteristics of Nanomaterials: D, 2D, and 3D,	
	Nanocrystal, Nanoparticle, Quantum dot, Quantum Wire and	
	Quantum Well etc.	
	4. Polymer, Carbon, Inorganic, Organic and Biomaterials –	
	Structures and characteristics	
	5. Green Synthesis of Nanoparticles Using Bacteria and Plants	
	6. Characterization of Nanoparticles	
	7. Applications of Nanobiotechnology	
	8. Relevance of nanotechnology in the food industry	
	Unit Outcomes: UO 1 Understand Nanobiotechnology Fundamentals and Applications.	
	UO 2 Competence in evaluating and proposing strategies for	
	green synthesis of nanoparticles	

- 1. Food Biotechnology, CRC Taylor & Francis Kalidas Shetty G. Paliyath, 2<sup>nd</sup> edition, A Pometto R,E. Levin, 2005.
- 2. Food Microbiology, Adam M.R and Moss M.O, 2<sup>nd</sup> edition, New Age International Pub., 2006.
- 3. Food Microbiology, Frazier W.C and Westhoff D.C 4<sup>th</sup> Edition., Tata McGraw Hill, 2013.
- 4. Food Processing and Preservation, Sivsankar B Prentice Hall of India, 2002.
- 5. Food Microbiology Protocols. Spencer J.F.T. and de Spencer A.L.R. ,Humana Press, 2008.
- 6. Modern Food Microbiology, Jay J.<mark>M., Cha</mark>pman and Hall,4<sup>th</sup> Ed., New York ,NY, USA,
- 7. Bio-Nano technology concept and applications, Madhuri Sheron, Sunil Pande- Ane Books New Delhi, 2013.
- 8. Nanotechnology, Pearson Mark Ratner, 1st edition, Daniel Ratner, 2002.
- 9. Nanotechnology an Introduction, Ramsden, 1st edition, Elsevier, 2012.
- 10. Advances in food Biotech<mark>nol</mark>ogy, S<mark>mith, M. A., & Dávila,</mark> G. (Eds.), John Wiley & Sons, 2018.
- 11. Food engineering: integrated approaches, Gómez-López, V. M. (Ed, CRC Press. 2017
- 12. Applications of biotechnology in the food industry. In Biotechnology Molecular Studies and Novel Applications for Improved Quality of Human Life, Silva, C., & others. Intech Open, 2019.
- 13. Emerging Applications of Nanotechnology in the Food Industry: A Critical Review. Nanomaterials in Food ,Pareek, S., & others. Elsevier,2020.
- 14. Nanotechnology in the Agri-Food Sector: Implications for the Future ,Sanz, T., Salvador, A., & Fiszman, S. M. ,Academic Press,2017.





# (Autonomous)

#### **Department of Biotechnology**

**Course Type: Lab Course** 

Course Title: Lab Course -XV (Based on MMC -XII)

Course Code: 602BIO4106

Credits: 01 Max. Marks: 50 Hours: 30

#### **Leaning Objectives**

- LO 1 Understand the process of fermentation in food.
- LO 2 Acquire skills in quantitative analysis.
- LO 3 Develop skills in microscopic examination of food and milk using the breed method.
- LO 4 Assess the quality of pasteurized milk
- LO 5 Acquire practical skills in performing these food processing techniques.
- LO 6 Learn techniques for isolating and characterizing probiotic bacteria.
- LO 7 Understand the principles of nanoparticle synthesis and characterization
- LO 8 Acquire skills in evaluating the efficacy of nanoparticles as antimicrobial agents

#### **Course outcomes**

- CO 1 Identify and characterize microorganisms involved in food fermentation processes, which can be applied to various food industries
- CO 2 Quantify biomolecules and nutritional content in food samples, which is crucial for assessing nutritional quality and stability of food products.
- CO 3 Analyze food samples for microbial contamination using microscopic techniques, aiding in food quality control
- CO 4 Determine the adequacy of milk pasteurization through mbrt and phosphatase testing, ensuring microbiological safety.
- CO 5 Gain knowledge and practical skills in isolating, characterizing, and evaluating the probiotic properties, facilitating their understanding of probiotic culture selection and application in food production
- CO 6 Design and conduct sensory evaluation tests to assess differences and similarities in food products, enabling them to understand consumer preferences and product quality attributes
- CO 7 Assess the antimicrobial potential of nanoparticles and understand their applications in food preservation and biomedical fields
- CO 8 Gain knowledge and practical experience in isolating and detecting nanoparticles from plant extracts, contributing to their understanding of nanotechnology applications in food and biomedical sciences

Practical No.	Unit
1	Isolation and Characterization of food fermenting organisms from idli batter
2	Estimation of ascorbic acid from given food sample by the titrimetric
	method.
3	Analysis of mycotoxin (Aflatoxin) in fungus-contaminated food material.
4	Microscopic examination of Food/Milk by breed method.
5	Estimation of lactose from milk.
6	Quality characterization of pasteurized milk by MBRT method.
7	To judge efficiency o <mark>f pasteurizati</mark> on of milk by Phosphatase test.
8	Detection of microbial count in Milk by SPC method.
9	Measurement of fat c <mark>ontent using Soxhl</mark> et extraction or solvent extraction methods.
10	Estimation of protein content using Kjeldahl method or Dumas combustion method.
11	Demonstration of various food processing techniques such as blanching, pasteurization, sterilization, and freeze-drying.
12	Preparation of food products like jams, jellies, sauces, and pickles.
13	Isolation and biochemical testing of probiotic cultures (Lactobacilli) from food samples (curd, intestine, sauerkraut, dosa, etc)
14	Check the potential of bacterial culture as probiotic culture by testing bile i) salt tolerance ii) acid tolerance iii) heat tolerance
15	Conducting discrimination tests (e.g., triangle test, duo-trio test) and descriptive analysis to assess sensory characteristics.
16	Isolation and detection of nano particles from plant extract (silver nano particles)
17	Spectrophotometric analysis (UV/IR) of nano particles
18	Antimicrobial activity of nano particles
N.B.: Any Ten Pra	acticals from above.



#### (Autonomous)

# **Department of Biotechnology**

Course Type: MEC-IV (A)

**Course Title: Environment Biotechnology** 

**Course Code: 602BIO4201** 

Credits: 03 Max. Marks: 75 Lectures: 45 Hrs.

# **Learning Objectives**

- LO 1 To understand the structure and functions of ecosystems, including the roles of abiotic and biotic components.
- LO 2 To describe the flow of energy through ecosystems and analyze food chains and food webs
- LO 3 To classify different types of pollution and pollutants, including their properties and effects on the environment.
- LO 4 To discuss sustainable management practices and conservation strategies for the environment.
- LO 5 To understand the processes involved in wastewater treatment plants, including physical, chemical, and biological unit operations.
- LO 6 To describe different types of bioremediation techniques, including microbial and phytoremediation.
- LO 7 To demonstrate knowledge of remote sensing principles, terminologies, and applications in various fields.
- LO 8 To understand the objectives and guidelines of Environmental Impact Assessment (EIA) and its classification.

#### **Course Outcomes**

- CO 1 Develop a comprehensive understanding of ecosystem dynamics and the factors influencing ecological balance.
- CO 2 Analyze the role of biotic and abiotic components in shaping ecosystems and their resilience to disturbances.
- CO 3 Apply knowledge of pollution classification and pollutants to assess environmental risks and propose mitigation measures

- CO 4 Understand the significance of renewable energy sources and their potential contribution to reducing environmental impacts.
- CO 5 Describe biotechnological solutions to address environmental issues of pollution.
- CO 6 Explain emerging technologies that are important in the area of environmental biotechnology.
- CO 7 Explain Remote sensing & GIS.
- CO 8 Demonstrate competency in conducting Environmental Impact Assessments and proposing recommendations for sustainable development.

Unit No.	Title of Unit & Contents	Hrs.
I	Ecology and Environment	12
	<ol> <li>Ecosystem structure and functions,</li> <li>abiotic and biotic component.</li> <li>Energy flow,</li> <li>Food chain, food web.</li> <li>Ecological Pyramids-types.</li> <li>Biogeochemical cycles.</li> <li>Ecological succession,</li> <li>Ecads and ecotypes.</li> <li>Sustainable management and conservation of environment</li> </ol>	
	Unit Outcomes:  U0 1 Develop a comprehensive understanding of ecosystem dynamics and the factors influencing ecological balance.  U0 2 Analyze the role of biotic and abiotic components in shaping ecosystems and their resilience to disturbances.	
II	Environmental Pollution	10
	<ol> <li>Classification of pollution</li> <li>Classification of pollutants.</li> <li>Air pollution and their properties,</li> <li>Water pollutants and their properties.</li> <li>Environmental pollution and associated hazards to crops, animals and humans.</li> <li>Greenhouse effect and global warming.</li> </ol>	

Unit No.	Title of Unit & Contents	Hrs.
	Unit Outcomes:	
	UO 1 Apply knowledge of pollution classification and	
	pollutants to assess environmental risks and propose	
	mitigation measures	
	UO 2 Understand the significance of renewable energy	
	sources and their potential contribution to reducing	
	environmental i <mark>mpact</mark> s.	
III	Biotechnological processes	15
	1. Waste water trea <mark>tment pl</mark> ant-	
	2. Physical, Chemica <mark>l and Biolog</mark> ical Unit	
	operations/proces <mark>ses-overview,</mark>	
	3. Activated Sludge P <mark>rocess,</mark>	
	4. Trickling <mark>Fil</mark> ters,	
	5. UASB rea <mark>ctor</mark>	
	6. Introdu <mark>ction to bioremediation,</mark>	
	7. Types of Bioremediations	
	8. Microbial bioremediation- Types	
	9. Phytoremediation- Types	
	10. Energy & Biofuels: Non-conventional or renewable	
	sources of energy,	
	11. En <mark>ergy fr</mark> om Bio <mark>mass</mark> .	
	Unit Outcomes:	
	UO 1 Describe biotechnological solutions to address	
	environmental issues of pollution.	
	UO 2 Explain emerging technologies that are important in	
	the area of environmental biotechnology.	
IV	Advancement in environmental technology	08
	1. Remote sensing and GIS- Principal, terminologies and	
	objectives.	
	2. Energy sources for remote sensing,	
	3. Types of remote sensing.	

Unit No.	Title of Unit & Contents	Hrs.
	4. Applications of Remote Sensing- Agricultural, Forestry,	
	Water Resource, Urban Planning, Wildlife Ecology,	
	Disaster Assessment.	
	5. Environmental Impact Assessment: Introduction,	
	Objectives, Classification, Guidelines.	
	Unit Outcomes:	
	UO 1 Explain Remote sensing & GIS.	
	UO 2 Demonstrate competency in conducting	
	Environmenta <mark>l Impact</mark> Assessments and proposing	
	recommendat <mark>ions for sus</mark> tainable development.	

- 1. Environmental Biotechnology, 2nd edition, Allan Scragg, OUP Oxford, 2005
- 2. Environmental Biotechnology, Prof. Jogdand, Himalayan Publication, 2010
- 3. Environmental Biotechnology, Foster C.F. & John Ware D.A., Ellis Horwood Ltd., 1987
- 4. Biotechnology and Biodegradation, Karrely D., Chakrabarty K., & Omen G.S., Portfolio Publishing Co Inc., U.S., 1990
- 5. Bioremediation Engineering: Design and Application, John T. Cookson Jr., McGraw Hill, Inc., 1994
- 6. Environmental Biotechnology, 3rd edition, A.K. Chatterjee, Prentice Hall India Learning Private Limited, 2011
- 7. Environmental Biotechnology, Bimal Bhattachraya & Ritu Banerjee, Oxford University Press, 2007
- 8. Environmental Pollution Control Engineering, 4th edition, C.S. Rao, New Age International Publishers, 2021
- 9. Environmental Biotechnology: Theory and Application, 1st edition, Gareth Evans & Judith Furlong, John Wiley and Sons Ltd., 2002
- 10. Environmental Biotechnology: Concept and Application, 1st edition, edited by Hans-Joachim Jördening & Josef Winter, Wiley VCH Verlag GmbH & Co. KGaA, 2004

Rajarshi Shahu Mahavidyalaya, Latur (Autonomous)



# (Autonomous)

#### **Department of Biotechnology**

**Course Type: Lab Course** 

Course Title: Lab Course -XVI (Based on MEC-IV (A))

Course Code: 602BIO4203

Credits: 01 Max. Marks: 50 Hours: 30

#### **Learning Objectives**

- LO 1. Gain proficiency in various analytical methods for the determination of water and soil quality parameters.
- LO 2. Acquire skills in isolating bacteria and fungi from different environmental sources.
- LO 3. Develop competence in composting organic waste materials to produce nutrient-rich compost.
- LO 4. Gain insight into the process of conducting environmental impact assessments and evaluating the potential environmental consequences of various projects, fostering environmental stewardship and responsibility.
- LO 5. Understand the principles and techniques involved in ecological restoration projects.
- LO 6. Learn techniques for the production of microbial-based biofertilizers.
- LO 7. Explore symbiotic relationships between plants and beneficial microorganisms.
- LO 8. Develop critical thinking skills by analyzing case studies related to environmental restoration projects and ecological assessments.

#### **Course Outcomes**

- CO 1 Proficiently employ various analytical techniques to determine water and soil quality parameters, ensuring accurate environmental assessments.
- CO 2 Master the techniques for isolating bacteria and fungi from polluted soil, water, and air samples, showcasing proficiency in microbial ecology and diversity studies.
- CO 3 Apply composting principles to effectively manage organic waste materials, producing nutrient-rich compost for soil amendment and waste reduction.

- CO 4 Capable of conducting comprehensive environmental impact assessments, evaluating the potential ecological consequences of proposed projects and activities.
- CO 5 Develop skills to design and implement ecological restoration plans, promoting biodiversity conservation and ecosystem resilience.
- CO 6 Produce microbial-based biofertilizers using nitrogen-fixing bacteria, phosphate-solubilizing bacteria, and plant growth-promoting rhizobacteria, contributing to sustainable agricultural practices.
- CO 7 Understand the significance of symbiotic relationships between plants and beneficial microorganisms, applying this knowledge to enhance plant growth and environmental sustainability.
- CO 8 Analyze environmental case studies to understand real-world applications of environmental science and propose informed solutions to environmental challenges.

Practical No.	Unit
1.	Determination of total solids.
2.	Determination of alkalinity
3.	Determination of COD
4.	Determination of DO
5.	Determination of BOD
6.	Determination of hardness of water
7.	Isolation of bacte <mark>ria f</mark> rom polluted soil.
8.	Isolation of bacteria from polluted water.
9.	Isolation of bacteria from polluted air.
10.	Isolation of fungi from polluted soil.
11.	Isolation of fungi from polluted water.
12.	Isolation of fungi from polluted air.
13.	Composting of organic waste materials to produce nutrient-rich compost for soil amendment.

14.	Investigation of composting parameters such as temperature, moisture content, carbon-to-nitrogen ratio, and microbial activity.
15.	Study of symbiotic relationships between plants and beneficial microorganisms (e.g., mycorrhizal fungi, nitrogen-fixing bacteria) for enhancing plant growth and environmental sustainability.
16.	Production of microbial-based biofertilizers using nitrogen-fixing bacteria, phosphate solubilizing bacteria, and plant growth-promoting rhizobacteria (PGPR).
17.	Case study on Ecological restorartion projects
18.	Case study on Envir <mark>onmen</mark> tal Impact Assesment.

N.B.: Any Ten Practicals from above.



### Rajarshi Shahu Mahavidyalaya, Latur



#### (Autonomous)

#### **Department of Biotechnology**

**Course Type: MEC-II (B)** 

**Course Title: Omics Technology** 

Course Code: 602BIO4202

Credits: 03 Max. Marks: 75 Lectures: 45 Hrs.

#### **Learning Objectives**

- LO 1 To explain the principles behind genomics, transcriptomics, proteomics, and metabolomics, including the technologies used and their applications in biological research.
- LO 2 To describe the process of generating omics data, including sample preparation, data acquisition, and quality control measures.
- LO 3 To apply bioinformatics tools and methods to analyze omics data, including data preprocessing, normalization, statistical analysis, and interpretation of results.
- LO 4 To interpret omics data to identify patterns, trends, and biological insights relevant to the research question.
- LO 5 To design omics experiments, including selecting appropriate technologies, experimental conditions, and controls.
- LO 6 To critically evaluate omics data and scientific literature to draw meaningful conclusions and propose hypotheses for further investigation.
- LO 7 To aware the students about ethical issues related to omics analysis, including data privacy, consent, and responsible conduct of research.
- LO 8 To apply omics analysis skills in research or professional settings, including academic research, biotechnology industry, healthcare, or regulatory agencies.

#### **Course outcomes**

After completion of the course, the student will be able to-

- CO 1 demonstrate an understanding of genomics, transcriptomics, proteomics, metabolomics, and other omics technologies, including their principles, methodologies, and applications in biological research.
- CO 2 generate omics data using relevant technologies and apply bioinformatics tools and methods to analyze and interpret omics data sets.
- CO 3 design omics experiments, including selecting appropriate technologies, experimental conditions, and controls, to address specific biological questions.
- CO 4 develop critical thinking skills and be able to evaluate omics data and scientific literature to identify research gaps, formulate hypotheses, and propose experimental approaches.

- CO 5 demonstrate an awareness of ethical issues related to omics technologies, including data privacy, consent, and responsible conduct of research, as well as the societal implications of omics research.
- CO 6 effectively communicate omics analysis results and conclusions through written reports, oral presentations, and visualizations to both scientific and non-scientific audiences.
- CO 7 think about integrative approach of omics technologies and its application in research.
- CO 8 Prepare the student for careers in academia, industry, healthcare, or regulatory agencies that require omics analysis skills, including the ability to adapt to new technologies and research paradigms in the rapidly evolving field of omics.

Unit No.	Title of Unit & Contents	Hrs.
I	Introduction to omics technologies	06
	<ol> <li>Overview of omics technologies,</li> <li>Historical perspective.</li> <li>Importance and applications in biology.</li> <li>Ethical issues in omics research.</li> <li>Data sharing and privacy concerns.</li> <li>Public perception and communication of omics research.</li> <li>Advances in omics technologies.</li> </ol>	
	Unit Outcomes: U0 1. To study historical perspective and fundamentals of omics technologies. U0 2. To study ethical issues and advances in omics technologies.	
II	Genomics and Transcriptomics	12
	<ol> <li>Basics of genomics</li> <li>Sequencing technologies- Next-generation sequencing technologies</li> <li>genome assembly, and annotation</li> <li>Comparative genomics</li> <li>Data analysis method databases</li> <li>tools for Genomics data analysis</li> <li>Basics of transcriptomics</li> <li>Microarrays</li> <li>RNA-Seq and other transcriptome profiling techniques</li> <li>Differential gene expression analysis</li> </ol>	

Unit No.	Title of Unit & Contents	Hrs.					
	<ul> <li>Unit Outcomes:</li> <li>UO 1. To study NGS technologies fundamentals and data analysis in genomics.</li> <li>UO 2. To study transcriptomics, microarrays, and RNA-seq for the study of differential gene expression analysis.</li> </ul>						
III	Proteomics	12					
	1. Basics of proteomics,						
	2. Protein separation techniques						
	3. Mass spectrometry-based proteomics						
	4. protein identification and quantification						
	5. Data analysis methods						
	6. databases, tools for proteomics data analysis						
	7. Protein-protein interactions						
	Unit Outcomes:						
	UO 1. To study fundamentals of protein purification and						
	mass spectrosco <mark>py-based proteom</mark> ics. UO 2. To learn use <mark>of proteomics d</mark> atabases and its						
	application in data analysis.						
IV	Metabolomics	15					
	1. Basics of metabolomics						
	2. Analytical techniques in metabolomics						
	3. Metabolite identification and quantification						
	4. Data analysis methods,						
	5. databases, tools for metabolomics data analysis Metabolic						
	pathway analysis						
	Unit Outcomes:						
	UO 1. To understand analytical techniques used in						
	m <mark>etabo</mark> lites an <mark>d met</mark> abolomics analysis.						
	UO 2. To learn about databases of metabolomics and its						
	application i <mark>n met</mark> abolomics data analysis.						

#### **Learning Resources:**

- 1. Bioinformatics for Omics Data: Methods and Protocols, Bernd Mayer, Humana Press, 1st edition, 2011.
- 2. Statistical Methods in Bioinformatics: An Introduction, Warren J. Ewens and Gregory R. Grant, Springer, 2nd edition, 2010.
- 3. Introduction to Bioinformatics, Arthur M. Lesk, Oxford University Press, 4th edition, 2019.
- 4. Computational Biology: A Practical Introduction to BioData Processing and Analysis with Linux, MySQL, and R, Röbbe Wünschiers, O'Reilly Media, 1st edition, 2013.

- 5. Bioinformatics: Sequence and Genome Analysis, David W. Mount, Cold Spring Harbor Laboratory Press, 2nd edition, 2002.
- 6. Bioinformatics and Functional Genomics, Jonathan Pevsner, Wiley-Blackwell, 3rd edition, 2015.
- 8. Data Mining Techniques in Bioinformatics: Theory and Models, Xue-Wen Chen, Mohammed J. Zaki, and Hagit Shatkay, Wiley-Blackwell, 1st edition, 2008.
- 9. Genomics and Proteomics: Principles, Technologies, and Applications, Stewart Sell and Prem Seth, Wiley, 1st edition, 2004.
- 10. Transcriptomics and Gene Regulation, Nikolai V. Dokholyan, Brian D. Sykes, and Jeffrey Skolnick, Springer, 1st edition, 2006.
- 11. Metabolomics: Methods and Protocols, Wolfram Weckwerth and Aimin Liu,, Humana Press, 1st edition, 2018.





## Rajarshi Shahu Mahavidyalaya, Latur

#### (Autonomous)

#### **Department of Biotechnology**

**Course Type: Lab Course-XIV** 

**Course Title: (based on omics technologies)** 

Course Code: 602BIO4203

Credits: 01 Max. Marks: 50 Hours: 30

#### **Learning Objectives**

- LO 1.To learn techniques for DNA and RNA extraction, quantification, and quality assessment.
- LO 2.To Understand principles of polymerase chain reaction (PCR) and agarose gel electrophoresis.
- LO 3.To gain proficiency in using bioinformatics tools and databases for sequence alignment, genome assembly, variant calling, and functional annotation.
- LO 4.To prepare samples for proteomic and metabolomic analysis, including protein extraction, purification, and metabolite extraction.
- LO 5.To analyze proteomic and metabolomic data using bioinformatics tools and databases to identify proteins and metabolites,

#### **Course outcomes**

After completion of the course, the student will be able to-

- CO 1 familiar with common experimental techniques used in genomics, such as DNA extraction, and PCR.
- CO 2 use bioinformatics tools and databases to analyze genomic data, including sequence alignment, variant calling, and functional annotation.
- CO 3 Students should be able to interpret genomic data and draw meaningful conclusions about genetic variation, gene function, and evolutionary relationships
- CO 4 prepare samples for proteomic and metabolomic analysis, including protein extraction, digestion, and metabolite extraction.
- CO 5 identify proteins and metabolites from mass spectrometry data using database search algorithms and other identification techniques.

Latur (Autonomous)

Practical	Unit				
No.					
1.	Isolation of genomic DNA from bacteria, plant and animal tissues.				
2.	To amplify specific DNA segments by using PCR and run the agarose gel				
	electrophoresis to study the amplified products of DNA.				
3.	Retrieve the genomic database of a particular bacteria/plant/animal for				
	comparative studies, and other characterization studies.				
4.	Take any suitable example to understand in silico process of gene				
	prediction, genome an <mark>notati</mark> on and data analysis.				
5.	Isolation of RNAs fro <mark>m bacter</mark> ia, plant and animal tissue by using Trizol				
	method.				
6.	Introductory practical on RNA-seq data analysis by using any online				
	suitable tool.				
7.	Isolation, pr <mark>ec</mark> ipitatio <mark>n, dialysis and pu</mark> rification of proteins from				
	animal/plant tissues/bacteria.				
8.	A study of proteomics databases, data retrieval for proteomics data				
	analysis.				
9.	To identify the Proteins from given proteomics database file by using any				
	available online tool.				
10.	To extract and purify the secondary metabolites from given plant material.				
11.	A introductory study of metabolomics databases and its application in				
	plant research and industry.				

N.B.: Any Ten Practicals from above.





## Shiv Chhatrapati Shikshan Sanstha's Rajarshi Shahu Mahavidyalaya, Latur

# (Autonomous) PG First Year

#### **Extra Credit Activities**

Sr.	Course Title	Credits	Hours
No.			T/P
1	MO0Cs	Min. of 02 credits	Min. of 30 Hrs.
2	Certificate Courses	Min. of 02 credits	Min. of 30 Hrs.
3	IIT Spoken	Min. of 02 credits	Min. of 30 Hrs.
	Tutorial Courses		

#### **Guidelines:**

#### Extra -academic activities

- 1. All extra credits claimed under this heading will require sufficient academic input/contribution from the students concerned.
- 2. Maximum 04 extra credits in each academic year will be allotted.
- 3. These extra academic activity credits will not be considered for calculation of SGPA/CGPA but will be indicated on the grade card.

#### Additional Credits for Online Courses:

- 1. Courses only from SWAYAM and NPTEL platform are eligible for claiming credits.
- Students should get the consent from the concerned subject Teacher/Mentor/Vice Principal and Principal prior to starting of the course.
- 3. Students who complete such online courses for additional credits will be examined/verified by the concerned mentor/internal faculty member before awarding credits.
- 4. Credit allotted to the course by SWAYAM and NPTEL platform will be considered as it is.

#### **Additional Credits for Other Academic Activities:**

- 1. One credit for presentation and publication of paper in International/National/State level seminars/workshops.
- 2. One credit for measurable research work undertaken and field trips amounting to 30 hours of recorded work.
- 3. One credit for creating models in sponsored exhibitions/other exhibits, which are approved by the concerned department.
- 4. One credit for any voluntary social service/Nation building exercise which is in collaboration with the outreach center, equivalent to 30 hours
- 5. All these credits must be approved by the College Committee.

#### **Additional Credits for Certificate Courses:**

- 1. Students can get additional credits (number of credits will depend on the course duration) from certificate courses offered by the college.
- 2. The student must successfully complete the course. These credits must be approved by the Course Coordinators.
- 3. Students who undertake summer projects/internships/training in institutions of repute through a national selection process, will get 2 credits for each such activity. This must be done under the supervision of the concerned faculty/mentor.

#### Note:

- 1. The respective documents should be submitted within 10 days after completion of Semester End Examination.
- 2. No credits can be granted for organizing or for serving as office bearers/volunteers for Inter-Class / Associations / Sports / Social Service activities.
- 3. The office bearers and volunteers may be given a letter of appreciation by the respective staff coordinators. Besides, no credits can be claimed for any services/activities conducted or attended within the college.
- 4. All claims for the credits by the students should be made and approved by the mentor in the same academic year of completing the activity.
- 5. Any grievances of denial/rejection of credits should be addressed to Additional Credits Coordinator in the same academic year.
- 6. Students having a shortage of additional credits at the end of the third year can meet the Additional Credits Coordinator, who will provide the right advice on the activities that can help them earn credits required for graduation.



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### Shiv Chhatrapati Shikshan Sanstha's Rajarshi Shahu Mahavidyalaya, Latur

# (Autonomous) Examination Framework

#### Theory:

40% Continuous Assessment Tests (CATs) and 60% Semester End Examination (SEE)

#### **Practical:**

50% Continuous Assessment Tests (CATs) and 50% Semester End Examination (SEE)

Course	Marks	CAT & Mid Term Theory				CAT Practical		Best Scored CAT & Mid Term	SEE	Total
				3		4				
1	2	Att.	CAT I	Mid	CAT II	Att.	CAT	5	6	5 + 6
				Term						
Research	100	10	10	20	10	-	-	40	60	100
Methodology										
DSC/DSE	75	05	10	15	10	-	-	30	45	75
Lab Course	50	-	<b>  -</b>	/ -	-	05	20	-	25	50
Field Project	100	10	10	20	10	-	-	40	60	100

#### Note:

- 1. All Internal Exams are compulsory
- 2. Out of 02 CATs best score will be considered
- 3. Mid Term Exam will be conducted by the Exam Section
- 4. Mid Term Exam is of Objective nature (MCQ)
- 5. Semester End Exam is of descriptive in nature (Long & Short Answer)
- 6. CAT Practical (20 Marks): Lab Journal (Record Book) 10 Marks, Overall Performance 10 Marks.

शि आरोह तमसो ज्योतिः।। Rajarshi Shahu Mahavidyalaya, Latur (Autonomous)

#### Shiv Chhatrapati Shikshan Sanstha's



## Rajarshi Shahu Mahavidyalaya, Latur

(Autonomous)

## **Semester End Examination Paper Pattern**

#### Pattern - I

Course: Theory Max. Marks: 45 Time: 2 Hrs

Q.1 Answer the following questions (3 Marks each)

12 Marks

- a) Based on Unit I
- b) Based on Unit II
- c) Based on Unit III
- d) Based on Unit IV

Q.2 Answer any THREE of the following (5 Marks each)

15 Marks

- a) Based on Unit I
- b) Based on Unit II
- c) Based on Unit III
- d) Based on Unit IV

Q.3 Answer any ONE of the following

08 Marks

- a) Based on Unit I
- b) Based on Unit II

Q.4 Answer any ONE of the following

10 Marks

- a) Based on Unit III
- b) Based on Unit IV

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Latur (Autonomous)

# Shiv Chhatrapati Shikshan Sanstha's



# Rajarshi Shahu Mahavidyalaya, Latur

(Autonomous)

# **Semester End Examination Paper Pattern**

## Pattern - I

Cour	rse : 1	rneory	Max. Marks :	60	Time: 2.30 Hrs
Q.1	Ansv	wer the following quo	estio <mark>ns (4</mark> Marks	each)	16 Marks
	a)	Based on Unit - I			
	b)	Based on Unit - II			
	c)	Based on Unit - III			
	d)	Based on Unit - IV			
Q.2	Ansv	wer any THREE o <mark>f t</mark> he	e foll <mark>owing (6 Ma</mark>	<mark>rks eac</mark> h)	18 Marks
	a)	Based on Unit - <mark>I</mark>			
	b)	Based on Unit - <mark>II</mark>			
	c)	Based on Uni <mark>t - III</mark>			
	d)	Based on U <mark>nit - IV</mark>			
Q.3	Ansv	wer any T <mark>WO of the f</mark>	ollowing (8	3 Marks each)	16 Marks
	(Bas	ed on <mark>any two U</mark> nits)			
	a)				
	b)		121	व छत्रप	ती
	c)		far	थाग जं	ESIT
<b>Q.4</b>	Ansv	wer any ONE of the fo	llowing		10 Marks
	(Bas	ed on remaining two U	Inits)	الأح	
	a)				
	b)				

# Shiv Chhatrapati Shikshan Sanstha's



# Rajarshi Shahu Mahavidyalaya, Latur

(Autonomous)

# **Semester End Examination Paper Pattern**

## Pattern - I

Cou	rse : I	Numerical	Max. Mar	ks : 60	Time	: 2.30 Hrs
Q.1	Ans	wer the following qu	estio <mark>ns (4</mark> Ma	arks each)		16 Marks
	a)	Based on Unit - I				
	b)	Based on Unit - II				
	c)	Based on Unit - III				
	d)	Based on Unit - IV				
Q.2	Ans	wer any TWO of t <mark>h</mark> e f	follow <mark>ing (9 I</mark>	Marks each)		18 Marks
	(Bas	ed on any two uni <mark>ts)</mark>				
	a)					
	b)					
	c)					
Q.3	Ans	wer any ON <mark>E of the f</mark> o	ollowing			16 Marks
	(Bas	ed on <mark>remaining</mark> two i	ınits)			
	a)					
	b)			शिव छः	त्रपती	
	c)			शिक्षण	<b>जंज्</b> शा	
<b>Q.4</b>	Ans	wer any ONE of the fo	oll <mark>owin</mark> g	लान्य	(1)	10 Marks
	(On	any Unit)		ande		
	a)				7/11	
	b)					

### **Summary of cross cutting issues:**

Biotechnology encompasses a wide array of technologies that utilize biological processes to create innovative products and solutions. These technologies range from traditional practices like brewing and bread-making to modern techniques such as genetic engineering, hybridization, and gene manipulation. As a transformative technology for the new millennium, biotechnology plays a crucial role in various sectors, including agriculture, medicine, food processing, environmental conservation, and even nanoelectronics.

To ensure comprehensive student development, the biotechnology curriculum integrates several cross-cutting issues, focusing on professional ethics, gender equality, environmental sustainability, and human values. These critical topics are embedded in various courses to enhance employability, foster entrepreneurship, and equip students with ethical and practical skills for the future.

# Cross-cutting issues relevant to Professional Ethics, Gender, Environment and Sustainability, and Human Values into the curriculum:

Sr. No.	Course Name	Code	Relevant to Professional	Description
	pro-		<b>Ethics/Environment</b>	
1	Genetic	MMC-	Professional Ethics	Prepares students for
	Engineering	VII		job opportunities in
				research institutes and
	- A			biotech industries.
2	Microbial	MMC-	Professional Ethics	Equips students with
	Biotechnology	VIII		skills for employability
				in fermentation
			1914 705	industries.
3	Plant and	MMC-	Professional Ethics	Opens pathways for
	Agriculture	IX	्राणश्राद	careers in research
	Biotechnology	1		institutes, agro-
		2	लातर	industries, and biotech
4	A.J	MEC	D. C	industries.
4	Advanced	MEC-	Professional Ethics	Provides job
	Pharmaceutical	III (A)	तमसा ज्यात	opportunities in the
	Biotech <mark>nology</mark>			pharmaceutical and
	Dajarch	i Ch	ahu Mahayidy	biopharmaceutical sectors.
5	Enzyme and Protein	MEC-	Professional Ethics	Enhances expertise in
)	Engineering	III (B)	1 Tolessional Ethics	enzyme and protein
	Linginicering	ш(в)	, mionomono)	technology for various
				industrial applications.
6	Animal and	MMC-	Professional Ethics	Prepares students for
	Livestock	X	11010001011di Liliico	careers in animal cell
	Biotechnology			culture laboratories.
L		L		

7	Clinical Research, IPR, Bio- entrepreneurship, and Start-Up	MMC- XI	Professional Ethics	Provides job opportunities in clinical research and biotech industries.
8	Food and Nano Biotechnology	MMC- XII	Professional Ethics	Equips students with skills for employability in the food, dairy, and fermentation industries.
9	Environmental Biotechnology	MEC- III (A)	Environment and Sustainability	Trains students to understand and address environmental problems, with opportunities to work as consultants or environment officers.
10	Omics Technology	MEC- III (B)	Professional Ethics	Provides insights into advanced omics technologies and their applications.
11	Research Project-I	RP-I	Professional Ethics	Encourages independent research, fostering ethical scientific inquiry and analysis.
12	Research Project-II	RP-II	Professional Ethics	Builds on research skills, emphasizing professional ethics in scientific research.

This reframed table highlights the integration of professional ethics, environment, and sustainability into biotechnology courses, ensuring students are prepared to tackle global challenges while adhering to ethical standards.



# Curricula developed and implemented have relevance to the local, national, regional and global developmental needs

The Biotechnology curriculum has been designed to align with developmental needs at the local, national, regional, and global levels. Each course focuses on addressing key challenges and opportunities in Biotechnology across various sectors, providing students with knowledge and skills to contribute to sustainable development and innovation.

Sr. No.	Course code	Course Name	Linkage with Local/National/Regional/Global development	
1	MMC- VII	Genetic Engineering	Addresses global advancements in genetic manipulation for healthcare, agriculture, and industry.	
2	MMC- VIII	Microbial Biotechnology	Solutions to local and global challenges in fermentation, waste management, and bioremediation.	
3	MMC-IX	Plant and Agriculture Biotechnology	Supports national and regional agricultural development through crop improvement and sustainability.	
4	MEC-III (A)	Advanced Pharmaceutical Biotechnology	Contributes to the pharmaceutical industry, enhancing national and global healthcare.	
5	MEC-III (B)	Enzyme and Protein Engineering	Facilitates regional industrial applications of enzymes and proteins, enhancing global competitiveness.	
6	MMC-X	Animal and Livestock Biotechnology	Addresses local and global challenges in animal health, breeding, and livestock improvement.	
7	MMC-XI	Clinical Research, IPR, Bio-entrepreneurship, and Start-Up	Encourages bio-entrepreneurship, addressing national and global demands for innovative biotech solutions.	
8	MMC- XII	Food and Nano Biotechnology	Links to regional and global food security challenges, with applications in food safety and nanoengineering.	
9	MEC-III (A)	Environmental Biotechnology	Addresses pressing global environmental issues, offering local and regional solutions for sustainability.	
10	MEC-III (B)	Omics Technology	Advances regional and global research in genomics, proteomics, and bioinformatics.	
11	RP-I	Research Project-I	Promotes local and national research initiatives, preparing students for global challenges.	

12	RP-II	Research Project-II	Fosters independent research aligned with
			national and global scientific progress.

#### Courses having focus on employability/entrepreneurship/skill development

The following biotechnology courses are designed to enhance employability, foster entrepreneurship, and develop essential skills. Each course is aligned with industry and research needs, equipping students with the tools required to succeed in their professional careers.

Sr N o.	Name of the Course	Cour se Code	Activities/C Employabil developme	ect bearing on urship/ Skill	Year of introduct ion		
			Employabi <mark>l</mark> y	it	Entrepreneu rship	Skill development	
1	Genetic Engineering	MMC -VII		or in ch	Provides foundational knowledge for starting genetic modification or recombinant DNA technology ventures.	Develops practical skills in gene manipulation and recombinant DNA technology.	2018-19
2	Microbial Biotechnolog y	MMC -VIII	Opens care opportunities in fermentation industries and bioprocessing labs.	es n	Encourages the creation of bioprocessing startups using microbial fermentation.	Provides hands-on experience in microbial genetics, fermentation technologies, and industrial applications.	2018-19

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3	Plant and Agriculture Biotechnolog y	MMC -IX	Enables employability in agro- industries, plant breeding companies, and research institutions.	Promotes entrepreneurs hip in crop improvement and plant tissue culture startups.	Trains students in plant tissue culture, genetic engineering, and crop improvement techniques.	2023-24
4	Advanced Pharmaceutic al Biotechnolog y	MEC- III (A)	Prepares students for roles in pharmaceutic al and biopharmace utical companies.	Supports entrepreneuri al initiatives in drug development and biopharmaceu tical startups.	Focuses on advanced drug development techniques and biopharmace utical production.	2018-19
5	Enzyme and Protein Engineering	MEC- III (B)	Offers employability in enzyme production, biocatalysis, and protein engineering sectors.	Encourages entrepreneurs hip in enzyme technology and industrial biotech applications.	Develops expertise in enzyme technology, protein modification, and biocatalysis.	2024-25
6	Animal and Livestock Biotechnolog y	MMC -X	Provides job opportunities in animal biotechnolog y labs, veterinary industries, and research institutes.	Supports startups in animal health, livestock improvement, and veterinary biotechnology.	Trains students in animal cell culture, genetic modification, and livestock biotechnolog y.	2023-24
7	Clinical Research, IPR, Bio- entrepreneur ship, and Start-Up	MMC -XI	Creates employability in clinical research, IPR management, and biotech industries.	Encourages bio- entrepreneurs hip through clinical research ventures and biotech startups.	Develops knowledge of clinical trials, IPR management, and bio- entrepreneur ship skills.	2023-24
8	Food and Nano	MMC -XII	Opens job opportunities in the food	Promotes entrepreneurs hip in food	Trains students in food	2018-19

	Biotechnolog y		industry, dairy industry, and nano- biotechnolog y sectors.	safety, dairy processing, and nanotechnolo gy-based ventures.	biotechnolog y, fermentation technologies, and nanoengineer ing.	
9	Environment al Biotechnolog y	MEC- III (A)	Provides employability in environmenta l consultancy, waste management, and bioremediatio n industries.	Supports environmenta l solutions- based startups, such as bioremediatio n and sustainable development consultancies.	Develops skills in bioremediatio n, sustainable development, and environmenta l impact assessment.	2018-19
10	Omics Technology	MEC- III (B)	Offers career opportunities in genomics, proteomics, and bioinformatic s sectors.	Encourages innovation and entrepreneurs hip in omics technologies and bioinformatics services.	Trains students in genomics, proteomics, and bioinformatic s tools and techniques.	2024-25
11	Research Project-I	RP-I	Fosters research aptitude, leading to employability in research institutions.	Encourages independent innovation and entrepreneurs hip in biotechnology research.	Develops project management, research, and analytical skills through independent research.	2024-25
12	Research Project-II	RP-II	Enhances employability through advanced research experience in biotechnolog y fields.	Supports entrepreneuri al thinking by promoting innovative research ideas and prototypes.	Builds advanced research skills, data analysis, and scientific writing abilities.	2018-19