## CHAUDHARY RANBIR SINGH UNIVERSITY, JIND

Syllabus of the Examination for Post Graduate Programme

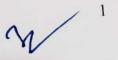
M.Sc. Biotechnology

(As Per NEP-2020) Curriculum and Credit Framework for Post Graduate Programme

With Multiple Entry-Exit, Internship and CBCS-LOCF With effect from the Session 2025-2026 (In Phased Manner)



Faculty of Life Sciences
Department of Biotechnology
Chaudhary Ranbir Singh University, Jind



Scheme of the Examination for Post Graduate Programme in M.Sc. Biotechnology

(As Per NEP-2020 Curriculum and Credit Framework for Post Graduate Programme)
With Multiple Entry-Exit, Internship and CBCS-LOCF with effect from the Session 2025-2026 (In Phased Manner)

	Course	Course Code	Nomenclature of course	Theory (T)/ Practica	C	redits	week	act hou (L:Le tical; T	cture	; P:	Internal Assessment Marks	End Term Examination Marks	Total Marks	Examinatio Hours
Sem.				l (P)		Total	L	T	P	Total		70	100	2
	CC 1	M24-BTY-101	Biomolecules	T	4		4	0	0	4	30	70	100	3
-	CC-1 CC-2	M24-BTY-102		T	4		4	0	0	4	30	70	100	3
- 1	CC-3	M24-BTY-103		T	4	1	4	0	0	4	30	70	100	3
L	CC-4	M24-BTY-104	Enzyme Technology	T	4	26	4	0	0	4	30	70	100	4
L	PC-1	M24-BTY-105	Bio-molecules & Enzyme	P	4		0	0	8	8	30			,
P	C-2	M24-BTY-106	Technology (Lab)  Molecular Cell Biology;  Microbiology & Biotechniques (Lab)	P	4		0	0	8	8	30	50	100	4
6	eminar	M24-BTY-107	Seminar	S	2		0	0	0	2	0	70	100	3
	C-5	M24-BTY-201	Genetic Engineering	T	4	26	4	0	0	4	30	70	100	_
	C-6	M24-BTY-202	Animal Cell & Tissue Culture	T	4		4	0	0	4	30	70	100	
	C-7	M24-BTY-203	Plant Cell &Tissue Culture	T	4		4	0	0	4	30	70	100	_
100000			Bioinformatics	T	4		4	0	0	4	30	70	10	
PC	C-8 C-3		Animal and Plant Cell & Tissue	P	4		0	0	8	8	30			
20	-4	M24-BTY-206	Culture Technology (Lab) Genetic Engineering &	P	4		0	0	8	8	30	70	10	0 4
			Bioinformatics (Lab)	T	2	-	2	0	1	) 2	15	35	5	3
Н	M	M24-CHM-201	Constitutional, Human and Moral values, and IPR	1			2	-	-		50	50	1	00
nte	rnship	M24-INT-200	Internship*										- ship C	m be either for

<sup>\*</sup>An internship course of 4 Credits of 4-6 weeks duration during summer vacation after 2<sup>nd</sup> semester is to be completed by every student. Internship can be eith enhancing the employability or for developing the research aptitude.



3 CC-9	)	M24-RTY-301	Plant Biotechnology	T	4	26	4	0	0	4	30	70	100	3
CC-1			Microbial Biotechnology	T	4	1	4	0	0	4	30	70	100	3
DEC-	1000		Molecular Genetics	T	4	1	4	0	0	4	30	70	100	3
DEC-		M24-BTY-305	-	T .	4		4	0	0	4	30	70	100	3
Any			Molecular Medicine & Diagnostic Techniques	T	4		4	0	0	4	30	70	100	3
PC-5		M24-BTY-307	Plant & Microbial Biotechnology (Lab)	P	4		0	0	8	8	30	70	100	4
PC-6	1		Molecular Genetics, Immunology/Molecular Medicine & Diagnostic Techniques (Lab)	P	4		0	0	8	8	30	70	100	4
OEC	?		Biotechnology & Human Welfare	T	2		2	0	0	2	15	35	50	3
ICC-11	M	124-BTY-401	Animal & Medical Biotechnology	Т	4	26	4	0	0	4	30	70	100	3
CC-12	-		Environmental Biotechnology	T	4	1	4	0	0	4	30	70	100	3
DEC-3			Food Biotechnology	T	4	1.	4	0	0	4	30	70	100	3
DEC-4	М	24-BTY-405	Genomics, Proteomics & Metabolomics	T	4	1	4	0	0	4	30	70	100	3
	34	24-BTV-406	Biosafety, Bioethics & IPR Issues of Biotechnology	Т	4		4	0	0	4	30	70	100	3
PC-7	M2	24-BTY-407	Food & Environmental Biotechnology (Lab)	P	4		0	0	8	8	30	70	100	4
PC-8	M2	24-BTY-408 A	Animal & Medical Biotechnology; Biosafety, Bioethics & IPR Isuues of Biotechnology/Genomics, roteomics & Metabolomics (Lab)	P	. 4		0	0	8	8	30	70	100	4
EEC	M2-	4-BTY-409	Entrepreneurship & Diagnostic Fechniques	T	2		2	0	0	2	15	35	50	3
	(N)	4 . If a Cam 1: 1-	te is Offered Dissertation Course, Th			SERTAT		II DE	C-3 D	FC-4 8	FFC from Al-	OVA COURS	es of Semest	er-4)
D:				ien ne/S	12	26	h	0	0	L	0	300	300	1
Dissertat Project W	Account to	STATE OF THE PARTY	Dissertation/Project Work	,	12	20	U	U	U		V	500		1
			TOTAL CREDITS			104				TOT	AL MARKS		2700	



# CHAUDHARY RANBIR SINGH UNIVERSITY, JIND

Faculty of Life Sciences Department of Biotechnology

Syllabus of the Examination for Post Graduate Programme In M.Sc. Biotechnology

(As Per NEP 2020)
Curriculum and Credit Framework for Post Graduate Programme

With Multiple Entry-Exit, Internship and CBCS-LOCF With effect from the Session 2024-2025 (In Phased Manner)



Faculty of Life Sciences

Department of Biotechnology

Chaudhary Ranbir Singh University, Jind

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## Chaudhary Ranbir Singh University, Jind

Syllabus for M.Sc. Biotechnology (CBCS) (III-Semester)

	Session: 202	25-26	
	Part A-Introd	luction	
Name of Programme	N	A.Sc. Biotechnology	
Semester		II	
Name of the Course		Plant Biotechnology	
Course Code		M24-BTY-301	
Course Type		CC-9	
Level of course (Anx-1)		500-599	
Pre-requisite for the course (if any)		NA	
completing this course, the learner will be able to:	CLO-1: Acquire Recent Kno of Plant Genome, Vectors, Methat are important for raising the Gene Silencing & Stacking CLO-2: Understand Genetic other value added Transger become entrepreneurs for biotechnology.  CLO-3: Attain knowledge bioactive/therapeutic Biomolof different cell culture systems and isolation of section and isolation of section and isolation of section and isolation.	and Molecular analysis of and Molecular analysis of and Molecular analysis of ang.  Engineering strategies for anic. They would be able various products and products and products of industrial important stems, bioreactors for their condary metabolites.	rmation and other aspect Transgenics. Understan quality improvement and to launch start-ups and cesses related to plan gh yielding of plan ance. Have an overviewen and technologies for the to GM crops.
Credits	Theory	Practical	Total
	4	0	4
Teaching Hours per week	4	0	4
Internal Assessment Marks	30	0	30
End Term Exam Marks	70	0	70
	100	0	
Max. Marks Examination Time	100 3 Hours	. 0	100

Instructions for Paper- Setter: The examiner will set 9 questions asking two questions from each unit and one compulsory question by taking CLOs into consideration. The compulsory question (Question No. 1) will consist at least 4 parts covering entire syllabus. The examinee will be required to attempt 5 questions, selecting one question from each unit and the compulsory question. All questions will carry equal marks.

Units	Topics	Contact Hours
ı	Plant Genetic Transformation: Organization of Plant, Nuclear, Chloroplast and Mitochondrial Genomes, Gene Tagging.  Chloroplast Transformation: Vector Designing, Method and Advantages. Agrobacterium mediated transformation-Ti and Ri Plasmids, Role of Virulence Genes, Mechanism of T-DNA transfer, Vectors based on Ti and Ri plasmids-Cointegrate and Binary vectors, technique and factors affecting Agrobacterium mediated transformation of plants.  Direct Gene Transfer: Particle Bombardment, ArF excimer laser, Electroporation, Microinjection and alternative methods. Screenable and Selectable markers.	20

	Analysis of Tonners in			
	Analysis of Transgenic Plants: For the prese transgenes and by biological assays. Gene Siler Gene Stacking in Plants: Motherland	ence, in	ntegration and expression of	
11				
	Introducing Strategies of Biotic and Abio Viral, Fungal, Insect, Herbicide resistance, Var Salinity, Ttemperature).  Genetic Engineering of Plants for Molecular of Antibodies, Vaccines and other Medically re Nutritional Enhancement of Plants (Carbol Vitamins), Manipulation of Flower color and Pulmportance.	ious Ab  r Farm lated Pr	ress Resistance/Tolerance: piotic stresses (like Drought, ning/Pharming: Production roteins in Plants. s. Seed storage Proteins,	16
III	Plant Cells as Biofactories for Production	on of	Secondary Metabolites:	14
	Secondary Metabolites, Types of Cell Culture	System	ns used for Production of	
	Secondary Metabolites and Advantages of their	in-vitro	production.	
	Strategies used for High Yield of Product: De			
	Yielding Cell-line Cultures, Optimization of Fac			
	(Physical Culture Conditions, Media and other I			
	Plant Cells, Bioreactors for Plant Cell, Organ &			
	Bio-transformation, Permeabilization of Cells an	d Remo	oval of Secreted Products.	10
IV	Intellectual Property Rights, Bio-safety & Eth	hical Is	sues: Intellectual Property	10
	Right (IPR): Patents, Trade Secrets, Copyr	right, C	Motorial Plant Breeders	
	Trademarks; GATT & TRIPPS; Patenting of Bi Rights (PBRs) & Farmers Rights; Clean General	Techr	pology: Current Status of	
	Transgenic Crops; Bane & Boon of GM Cro	ps: Coi	ncerns about GM Crops-	
	Environmental, Bio-safety & Ethical Issues.	p.,		
			<b>Total Contact Hours</b>	60
	Suggested Evaluation	Method	ds	
	Internal Assessment: 30		End Term Examination:	70
	Theory	30	Theory 70	
	❖ Class Participation	05	Written examination	
	Seminar/Presentation/Assignment/Quiz	10		
	or Class test, etc.	15		

Part C-Learning Resources

- 1. Elements of Biotechnology. PK Gupta, 4th Reprint (2nd Edition): 2019-2020, Rastogi Pub. 2. Plant Genetic Engineering. Vol: 1-6 (2003). Singh & Jaiwal (Eds.), Sci Tech. Publishing LLC, USA.
- 3. Introduction to Biotechnology. HS Chawla (2009), 3rd Ed., Science Pub. 4. Plant Tissue Culture: Theory & Practice. Bhojwani SS & Razdan M K (2005). Elsevier Publication.
- 5. Plant Biotechnology. Hammond J, Mc Garvey P & Yusibov V (2000) (Eds.) Springer Verlag, Germany.
- 6. Practical Application of Plant Molecular Biology. Henry RJC & Hall (1997).
- 7. Plants, Genes & Agriculture. Chrispeels MJ & Sadava DE (2002). 2nd Ed. Jones & Bartlett Pub. UK.
- 8. Plants, Genes & Agriculture. Chrispecis In Conduction of Plants (2<sup>nd</sup> Edition). Slater A, Scott N & Fowler M, Oxford Pub. (2008).
- 9. Crop Biotechnology: Genetic Modification and Genome Editing. Nigel G Halford (2018). World
- Scientific Pub. Europe Ltd. UK. 10. Plant Biotechnology: Principles & and Applications. Malik ZA, Usha K, Kamaluddin & Athar A (2017). Springer Nature, Singapore.

	Session: 2025		- 3
	Part A-Introdu	iction	
Si CN	N	A.Sc. Biotechnology	
Name of Programme		Ш	
Semester		Microbial Biotechnology	
Name of the Course		M24-BTY-302	
Course Code			
Course Type		CC-10	
Level of the course		500-599	
Pre-requisite for the course (if any)		NA	
Course Learning Outcome (CLO). After completing this course, the learner will be able to:  Credits	processes. Have insight a Developments in Fermentatio CLO-2: Attain knowledge various Media Optimizatio Industrial important Metale CLO-3: Get introduced to a Fermentation Broth. CLO-4. Understand the fermentations, knowledge Theory 4	of Micro-organisms in Specific Biot bout Industrially important Micro on Processes & various types of Ferminge about Designing of Industrial on strategies, Strategies for overproblites structure and functioning of various strategies of product recommon basic principles of microbial to solve critical problems.  Practical	bes, Recententations.  Strains & coduction of fermenter covery from
Teaching Hours per weel	k 4	0	
Internal Assessment Mark	30.00	0	30
End Term Exam Marks	70	0	70
Max. Marks	100	0	100
Examination Time	3 Hours		100
	Part B-Contents of	f the Course	
Question No. 1) will consi	question by taking CLOs in	9 questions asking two questions nto consideration. The compulsor	from each
o attempt 5 questions, sequestions will carry equal n	electing one question from onarks.  Topics	each unit and the compulsory que	y question
Jnits  Microbial Biotec Biology of Indu Metabolism Regu Preservation of Ir Fermentation System. Solid Sta  II Overproduction	Topics  Chnology: Scopes, Application strial Microorganisms: Indulation, Substrate Assimilation industrially Important Microorgystem: Batch, Fed Batch and the Fermentation & its Applicate Fermentation & its Application & its Applicati	n & Challenges.  ustrial Microorganisms, Growth n/Product Formation. Isolation & rganisms.	y question be required estion. All

111	Industrial Production of Alcoholic Beverages Improvement by Genetic Engineering.	(Whisk)	v. Wine & Beer) and	15			
	Microbial Production of Food Additives: Amino-acids, Nucleosides & Vitamins						
	Microbial Production of Industrial Chemics Ethanol, Citric-acid, Acetic-acid, Gluconic-acid,	Glycero	d. Acetone & Butanol.				
	Microbial Production of Healthcare Production of Healthcare Productions), Vaccines (Bacterial cells & Toxin	ts: Ant	tibiotics (Penicillin &	15			
IV	Microbial Inoculants: Food Starter Cultures; Baker's Yeast, Starter Cultures for Dairy Industry, Meat Starter Cultures.  Biomass Production: Single Cell Protein (SCP) Production; Microbia Inoculants; Microbial Transformation of Steroids & Sterols.  Down-stream Processing: Separation processes for Microbial Cells and other						
	Inoculants: Microbial Transformation of Steroids  Down street Processing: Separation processes	& Stere	orobial Cells and other				
	Down-stream Processing: Separation processes Solids, Cell disruption, Centrifugation, Solids, Crystallization, Recovery schemes for Non-vo	& Stere for Mic	crobial Cells and other ecovery. Drying and				
	Inoculants: Microbial Transformation of Steroids  Down street Processing: Separation processes	& Stere for Mic	crobial Cells and other ecovery, Drying and Metabolites, Biomass,	60			
	Down-stream Processing: Separation processes Solids, Cell disruption, Centrifugation, Solv Crystallization. Recovery schemes for Non-vo-	for Mic for Mic vent re platile	crobial Cells and other ecovery, Drying and Metabolites, Biomass,  Total Contact Hours				
	Down-stream Processing: Separation processes Solids, Cell disruption, Centrifugation, Solv Crystallization. Recovery schemes for Non-vo- Extra-cellulars Polysaccharides & Enzymes.  Suggested Evaluation	for Mic for Mic vent re platile	crobial Cells and other ecovery, Drying and Metabolites, Biomass,  Total Contact Hours	tion: 70			
	Down-stream Processing: Separation processes Solids, Cell disruption, Centrifugation, Solv Crystallization. Recovery schemes for Non-vo- Extra-cellulars Polysaccharides & Enzymes.  Suggested Evaluation Internal Assessment: 30	for Mic for Mic vent re platile	crobial Cells and other ecovery, Drying and Metabolites, Biomass,  Total Contact Hours ds  End Term Examina Theory	tion: 70			
	Inoculants: Microbial Transformation of Steroids  Down-stream Processing: Separation processes Solids, Cell disruption, Centrifugation, Solve Crystallization. Recovery schemes for Non-volume Extra-cellulars Polysaccharides & Enzymes.  Suggested Evaluation  Internal Assessment: 30  Theory	for Micovent replatile	crobial Cells and other ecovery, Drying and Metabolites, Biomass,  Total Contact Hours  ds  End Term Examina	tion: 70			
	Down-stream Processing: Separation processes Solids, Cell disruption, Centrifugation, Solv Crystallization. Recovery schemes for Non-vo- Extra-cellulars Polysaccharides & Enzymes.  Suggested Evaluation Internal Assessment: 30	for Michael Method	crobial Cells and other ecovery, Drying and Metabolites, Biomass,  Total Contact Hours ds  End Term Examina Theory	tion: 70			

- 1. Principles of Fermentation Technology. Stansbury PF et al. (1997). Pergmon Press, Oxford.
- Fermentation Biotechnology: Principles, Process & Products. Ward OP (1998). Prentice Hall Publishing, New Jersey.
- 3. Microbial Biotechnology: Basic Research & Applications. Singh et al. (2020). Pub. Springer. (Ed.).
- 4. Modern Industrial Microbiology and Biotechnology. Nduka Okafor (2007). Science Publishers
- 5. Manual of Industrial Microbiology & Biotechnology. Arnold I Demain & Julian E Davies (1999). 2nd
- Edition, ASM Press, Washington DC.
- 6. Microbial Biotechnology. Glazer & Nikaido (1998). WH Freeman & Company, New York.
- 7. Biotechnology: A Textbook of Industrial Microbiology. Cruger & Cruger (2002), 2nd Edition,
- Panima Pub. Corporation, New Delhi.



	Session: 20	25-26	
	Part A-Intro	duction	
Name of Programme		M.Sc. Biotechnology	
Semester		III	
Name of the Course		Molecular Genetics	
Course Code		M24-BTY-303	· ·
Course Type		DEC-1	
Level of the course		500-599	
Pre-requisite for the course (if any) Course Learning Outcome:		NA	
(CLO). After completing	- 1	J. Scholoxicity assays, Iranso	TIDLION roomleties !
be able to:	genome manipulation. CLO-2: Learn advanced comparative Genomics and CLO-3: Know fundamental CLO-4: Get acquainted was	s and applications of Metabol	nd its applications in ping & Sequencing
be able to:	genome manipulation. CLO-2: Learn advanced comparative Genomics and CLO-3: Know fundamental CLO-4: Get acquainted was	techniques of Genome Map Transcriptome analysis.	nd its applications in ping & Sequencing. ic Engineering. and tools needed to
be able to: Credits	genome manipulation. CLO-2: Learn advanced comparative Genomics and CLO-3: Know fundamental CLO-4: Get acquainted wacquire top level skills in the Theory	techniques of Genome Map Transcriptome analysis. Is and applications of Metabol ith Methodological concepts of field of Molecular Genetics.	nd its applications in ping & Sequencing ic Engineering. and tools needed to Total
Credits Teaching Hours per week	genome manipulation. CLO-2: Learn advanced comparative Genomics and CLO-3: Know fundamental CLO-4: Get acquainted wacquire top level skills in the Theory  4	techniques of Genome Map Transcriptome analysis. Is and applications of Metabol ith Methodological concepts of field of Molecular Genetics.	nd its applications in ping & Sequencing ic Engineering. and tools needed to total
Credits  Teaching Hours per week  Internal Assessment Marks	genome manipulation. CLO-2: Learn advanced comparative Genomics and CLO-3: Know fundamental CLO-4: Get acquainted wacquire top level skills in the Theory  4  4  30	techniques of Genome Map Transcriptome analysis. Is and applications of Metabol ith Methodological concepts of field of Molecular Genetics.  Practical	ond its applications in ping & Sequencing ic Engineering. and tools needed to tool and tool a
Credits  Teaching Hours per week Internal Assessment Marks End Term Exam Marks	genome manipulation. CLO-2: Learn advanced comparative Genomics and CLO-3: Know fundamental CLO-4: Get acquainted wacquire top level skills in the Theory  4  4  30  70	techniques of Genome Map Transcriptome analysis. Is and applications of Metabol ith Methodological concepts of field of Molecular Genetics.  Practical  0 0	ond its applications in ping & Sequencing ic Engineering.  and tools needed to
this course, the learner will be able to:  Credits  Teaching Hours per week Internal Assessment Marks End Term Exam Marks Max. Marks Examination Time	genome manipulation. CLO-2: Learn advanced comparative Genomics and CLO-3: Know fundamental CLO-4: Get acquainted wacquire top level skills in the Theory  4  4  30	techniques of Genome Map Transcriptome analysis. Is and applications of Metabol ith Methodological concepts of field of Molecular Genetics.  Practical  0 0 0	nd its applications in ping & Sequencing ic Engineering. and tools needed to tools

Instructions for Paper- Setter: The examiner will set 9 questions asking two questions from each unit and one compulsory question by taking CLOs into consideration. The compulsory question (Question No. 1) will consist at least 4 parts covering entire syllabus. The examinee will be required questions will carry equal marks.

Juits	Topics	Contact
1	Eukarvotic Genome Structure 8 O	Hours
11	Eukaryotic Genome Structure & Organization: Genome Sequence & Chromosome Diversity, Variation in Chromosome Number, Special features of Metaphase Chromosomes, Chromosome Banding, Genome Size & Complexity, Organization & Content of Human Genome, Repetitive DNA, Microsatellites, Genome Wide Repeats, Split Genes, Overlapping Genes, Cryptic Genes, Retro-Genes, Multi-Gene Families, Pseudo-Genes.  Nucleosome: Basic Structure, Spatial arrangements of Histones, Chromatosome, Solenoid Model, Chromatin domains & Modifications.  Mutability of DNA: An overview of Mutation and Polymorphism, VNTR Polymorphism, DNA damage-Spontaneous, Induced (Alkylation, Oxidation, Radiation), Genotoxicity/Mutagenicity test systems-Ames test, Sister Transcription (Chromatid exchanges, Micronucleus, Comet assay.	15
	Transcription Regulation in Prokaryotes: Positive & Negative Control of Transcription, Repression & Activation, Organization & Regulation of Lac, Trp & Ara Operon in E. coli, Organization of Genome in Lambda phage	18

Repressor Prote binding of Repressor Prote Franscription A

1				
	(Early, Middle & Late Genes), Regulation of Ly Repressor proteins (c1,c11,c111,cro), Establishm binding of Repressor, Maintenance of Autogenou <b>Transcription Regulation in Eukaryotes:</b> binding domains, Transcriptional repressors, pos Yeast Galactose utilizing genes. Signal Transcriptional regulators, Gene Silencing, Epigen	ent of L s circuit Eukaryo itive & ansduct	by c1 repressor.  tic activators, DNA negative regulation of ion and control of	
111	Regulatory RNAs: Ribo-switches, Interfering expression, Short interfering RNA (siRNA) and its functions, Antisense RNA and Gene expression Cas9 gene editing technology.  Site-Specific Recombination: Concept, Recomb lox recombination, Biological role and a recombination in genome manipulation.  Genome Mapping: DNA markers for genetic in Physical Mapping, Restriction mapping, Flor (FISH), Sequence tagged sites (STS) mapping.	RNA ts funct n, An or inases a oplication	(RNAi) and gene ions, Micro RNA and verview of CRISPER- nd their function, cre- ons of site-specific g, RFLP, SSP, SNPs,	15
IV	Genome Sequencing: Types-Whole genome sequencing, targeted sequencing, metagenomic approach or map-based sequencing, shot gun sequencing of genome sequencing. Sequencing of Pyro-sequencing, Next generation sequencing. Applications of genome sequencing.  Comparative Genomics: Concept, Orthologs and Horizontal gene transfer, Genome similarity, prokaryotes and eukaryotes, Genomic synteny, pherunctional Genomics: Expression profiling, The Gene function determination (Gene knockout strate Metabolic Engineering: Principle and method Directed production of molecules, production studies on rerouting of metabolic pathways; engineering.	sequence equencionethods High the paralog Compa ylogene Transcrip egy, Ins s of move	cing; Clone by clone ng; Technologies for (Sanger sequencing, roughput sequencing; gs, exon shuffling, arative Genomics in tic foot-printing otome, DNA Arrays, sertional mutagenesis) etabolic engineering; el compounds, Case	12
	engineering.		Total Contact Hours	60
	Suggested Evaluation	Method.		
	Internal Assessment: 30		End Term Examina	
	Theory	30	Theory	70
	<ul> <li>Class Participation</li> </ul>	05	Written examination	
	Seminar/Presentation/Assignment/Quiz or Class test, etc.	10	1	
	❖ Mid-Term Exam	15		

Part C-Learning Resources

## Recommended Books:

- 1. Essential Genes. Benjamin Lewin (2007). Pearson Edu. International.
- 2. Genomes. TA Brown (2017). Garland Science, Taylor & Francis, NY.

30

3. Principles of Gene Manipulation & Genomics. Primrose & Twyman (2006). Blackwell

Publishing.

- 4. Molecular Biotechnology. Bernard and Pasternak (2017). 5th Edi. ASM press, Washington.
- 5. Human Molecular Genetics. Tom Strachan & Andrew P Read (2011), 4th Edi. Garland Science.
- 6. Molecular Biology of Gene. Watson Baker et al. (2007). Levine & Losick, Pearson Education Inc.
- 7. Principles of Genetics. Gardener et al. (2006). 8th Edi. John Wiley, NY.
- 8. Genes XII. Lewin B (2017). 12th Edi. Jones and Bartlett Publishers.
- 9. Principles of Genetics. Snustad and Simmons (2006), 8th Edi. Wiley
- 10. Analysis of Genes and Genomes. Daniel and Bruce (2017). Cochrane, Jones and Bartlett Publishers.
- 11. Biotechnology and Genomics. Gupta PK (2013). 1st Edi. Rastogi Publishers.

	Session: 2025	5-26	
	Part A-Introdu	ection	
Name of Programme	N	1.Sc. Biotechnology	
Semester	I	II	
Name of the Course	I	mmunology	
Course Code	I	M24-BTY-305	
Course Type		DEC-2	
Level of the course	1	500-599	
Pre-requisite for the course (if any)		NA	
(CLO). After completing	coordinate to fight invading		
this course, the learner will be able to:	CLO-2: Understand and of generation of immune cell the production of monocle different types of vaccines. CLO-3: Know about probsolve them with the knowle CLO-4: Learn about different treatment of diseases.	lescribe antigen, antibodies is responses, and hybridon and antibodies, recombinated blems emerging in health sedge of this subject.	na technology fo nt antibodies, and sector and how to outic techniques in
this course, the learner will	CLO-2: Understand and of generation of immune cell the production of monocle different types of vaccines. CLO-3: Know about probsolve them with the knowle CLO-4: Learn about diffe treatment of diseases.  Theory	lescribe antigen, antibodies is responses, and hybridon onal antibodies, recombinated blems emerging in health seedge of this subject.  The rent diagnostic and therape of the seedge of the subject.	na technology for nt antibodies, and sector and how to the techniques in Total
this course, the learner will be able to:  Credits	CLO-2: Understand and of generation of immune cell the production of monocle different types of vaccines. CLO-3: Know about probsolve them with the knowle CLO-4: Learn about diffe treatment of diseases.  Theory	lescribe antigen, antibodies ls responses, and hybridon and antibodies, recombinated blems emerging in health sedge of this subject. The rent diagnostic and therape are the rent diagnostic and the repersist of the rent diagnostic and diagnostic	na technology fo nt antibodies, and sector and how to sector techniques in Total
this course, the learner will be able to:  Credits  Teaching Hours per week	CLO-2: Understand and of generation of immune cell the production of monocle different types of vaccines. CLO-3: Know about probsolve them with the knowle CLO-4: Learn about diffe treatment of diseases.  Theory  4  4	lescribe antigen, antibodies ls responses, and hybridon onal antibodies, recombinate olems emerging in health sedge of this subject. The rent diagnostic and therape of the rent diagnostic and the reperturbation of the rent diagnostic and diagnostic an	na technology fo nt antibodies, and sector and how to sector techniques in Total 4
this course, the learner will be able to:  Credits  Teaching Hours per week Internal Assessment Marks	CLO-2: Understand and of generation of immune cell the production of monocle different types of vaccines. CLO-3: Know about probsolve them with the knowle CLO-4: Learn about diffe treatment of diseases.  Theory  4  4  30	lescribe antigen, antibodies ls responses, and hybridon onal antibodies, recombinate olems emerging in health sedge of this subject. The rent diagnostic and therape of the sedge of the subject.  Practical  O  O  O	ra technology foont antibodies, and sector and how to sector and h
Credits  Credits  Teaching Hours per week Internal Assessment Marks End Term Exam Marks	CLO-2: Understand and of generation of immune cell the production of monocle different types of vaccines. CLO-3: Know about probsolve them with the knowle CLO-4: Learn about different treatment of diseases.  Theory  4  30  70	lescribe antigen, antibodies ls responses, and hybridon onal antibodies, recombinated blems emerging in health seedge of this subject.  The rent diagnostic and therape  Practical  0 0 0 0 0 0	ra technology foont antibodies, and sector and how to sector and h
this course, the learner will be able to:  Credits  Teaching Hours per week Internal Assessment Marks	CLO-2: Understand and of generation of immune cell the production of monocle different types of vaccines. CLO-3: Know about probsolve them with the knowle CLO-4: Learn about diffe treatment of diseases.  Theory  4  4  30	lescribe antigen, antibodies ls responses, and hybridon onal antibodies, recombinate olems emerging in health sedge of this subject. The rent diagnostic and therape of the sedge of the subject.  Practical  O  O  O	ra technology foont antibodies, and sector and how to sector and how to sector techniques in Total 4 4 30

Instructions for Paper- Setter: The examiner will set 9 questions asking two questions from each unit and one compulsory question by taking CLOs into consideration. The compulsory question (Question No. 1) will consist at least 4 parts covering entire syllabus. The examinee will be required to attempt 5 questions, selecting one question from each unit and the compulsory question. All questions will carry equal marks.

Units	Topics	Contact Hours
1	Introduction and Overview: Introduction and overview of Immunology, Cells of Immune System, Innate & Cellular Immunity, Physical & Chemical barriers, Cellular defences, Inflammation, Receptors involved in Innate Immune System, Cells and Organs Involved in Adaptive Immune Response, Fate of Antigen after Penetration, Interrelationship between Innate & Acquired Immunity.	15
11	Antigens, Antibodies & their interactions: Requirements of Immunogenicity, Primary and Secondary responses, Major Classes of Antigens, Basic Structure of Antibodies, Antibody Classes and Biological Activity, Antigenic determinants on Immunoglobulins (Ig), Ig Super Family, Organization and expression of Ig Genes.  Antigen-Antibody Interactions: Immunoprecipitation, Agglutination, ELISA, Immunofluorescence, Flow Cytometry.	15

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III	Generation of B-cell & T-cell Responses: Complement system & its activation, Structure & role of Major Histocompatibility Complex, T-cell receptor-structure, Complex & Accessory membrane molecules, Thymic Selection of T-cells, T-cell activation & differentiation, B-cell maturation, activation & proliferation, Humoralresponse, Cytokines properties & receptors.				
IV	Immune System in Health and Disease: Hypersensitivity reactions & their types and mechanism, Cancer and Immune System, Cancer Immunotherapy.  Hybridoma Technology: Commercial production of antibodies using monoclonal antibodies.  Vaccines: Live attenuated, killed, subunit, conjugate and DNA vaccines.  Production of recombinant antibodies and edible vaccines, development of diagnostics using biotech and nanotech tools.				
	Total Contact Hours				
	Suggested Evaluation	Method	End Term Examina	tion: 70	
	Internal Assessment: 30	30	Theory	70	
		1511	Incorv		
	Theory	_			
	❖ Class Participation	05	Written examination	I seems	
	Validation desired to the second seco	_			

- 1. Immunology: A short course. Benjamin E. 4th Edi. John Wiley, New York.
- 2. Immunology. Kuby J (8th Edi.). WH Freeman & Co. New York
- 3. Essential Immunology. Roitt IM (12th Edi.). Oxford Black Well Science, London
- 4. Immunology: An introduction. Tizardd IR (9th Edi.). Philadelphia Saunders College press.
- 5. Biotechnology and Genomics. Gupta PK. Rastogi Publications Meerut, India
- 6. Fundamentals of Microbiology. Jeffery C Pommerville et al. (2013). Jones and Bartcett Publishers.

13

	Session: 202	5-26				
	Part A-Introd	uction				
Name of Programme		M.Sc. Biotechnology				
Semester		III				
Name of the Course	Molecular I	Molecular Medicine & Diagnostic Techniques				
Course Code		M24-BTY-306				
Course Type		DEC-2				
Level of the course		500-599				
Pre-requisite for the course (if any)		NA inderstanding of various chro	and gane			
Course Learning Outcomes (CLO). After completing this course, the learner will	and mitochondrial disor	ders, different approaches	to detect these			
	role of gene therapy and medicine, role of free radi CLO-3: Have a broad ubiotechnological applicat aspects of Biotechnology. CLO-4: Get a springbo explore their ideas of Mol	ard to develop their creative ecular Medicine and Diagnos	ne. cal research for ht in to clinical ve thinking and tics.			
	role of gene therapy and medicine, role of free radical CLO-3: Have a broad ubiotechnological applicat aspects of Biotechnology.	recombinant molecules as a cals and metal ions in medicin nderstanding of the biomeditions. They would gain insignard to develop their creative cular Medicine and Diagnost Practical	cal research for the in to clinical ve thinking and tics.			
be able to:	role of gene therapy and medicine, role of free radi CLO-3: Have a broad ubiotechnological applicat aspects of Biotechnology. CLO-4: Get a springbo explore their ideas of Mol	recombinant molecules as a cals and metal ions in medicin nderstanding of the biomeditions. They would gain insignard to develop their creative cular Medicine and Diagnost Practical	cal research for the intervention of the inter			
Credits	role of gene therapy and medicine, role of free radi CLO-3: Have a broad ubiotechnological applicat aspects of Biotechnology. CLO-4: Get a springbo explore their ideas of Mol Theory  4	recombinant molecules as a cals and metal ions in medicin nderstanding of the biomeditions. They would gain insignard to develop their creative cular Medicine and Diagnost Practical  0 0	re thinking and tics.  Total			
be able to:	role of gene therapy and medicine, role of free radic CLO-3: Have a broad unbiotechnological applicat aspects of Biotechnology. CLO-4: Get a springbout explore their ideas of Mol Theory  4  4  30	recombinant molecules as a cals and metal ions in medicin nderstanding of the biomeditions. They would gain insignard to develop their creative cular Medicine and Diagnost Practical  0 0 0	re thinking and tics.  Total  4  30			
Credits  Teaching Hours per week	role of gene therapy and medicine, role of free radic CLO-3: Have a broad unbiotechnological applicat aspects of Biotechnology. CLO-4: Get a springbo explore their ideas of Mol Theory  4  30  70	recombinant molecules as a cals and metal ions in medicin nderstanding of the biomeditions. They would gain insignard to develop their creative cular Medicine and Diagnost Practical  0 0 0 0	re thinking and tics.  Total  4  30  70			
Credits  Teaching Hours per week Internal Assessment Marks	role of gene therapy and medicine, role of free radic CLO-3: Have a broad unbiotechnological applicat aspects of Biotechnology. CLO-4: Get a springbout explore their ideas of Mol Theory  4  4  30	recombinant molecules as a cals and metal ions in medicin nderstanding of the biomeditions. They would gain insignard to develop their creative cular Medicine and Diagnost Practical  0 0 0	re thinking and tics.  Total  4  30			

Instructions for Paper- Setter: The examiner will set 9 questions asking two questions from each unit and one compulsory question by taking CLOs into consideration. The compulsory question (Question No. 1) will consist at least 4 parts covering entire syllabus. The examinee will be required to attempt 5 questions, selecting one question from each unit and the compulsory question. All questions will carry equal marks.

Units	Topics			
I	Chromosomes Anomalies and Disorders: Numerical (Polyploidy, Ancuploidy, Autosomal, Sex-chromosomal) & Structural (Deletion, Duplication, Translocation, Inversion, Iso-chromosome, Ring-chromosome). Single Gene Disorders: Sickle Cell Anaemia, Haemophilia, Cystic Fibrosis, Tay-Sachs disease.  Huntington disease: Genetics, Prevalence, Diagnosis and Prognosis, Polygenic disorders: Type-1 Diabetes, Breast Cancer.  Alzheimer disease: Genetics, Prevalence, Diagnosis and Prognosis.  Mitochondrial disorders: Mitochondrial Homeostasis and Parkinson disease.	15		
II	Immunological Approaches: To detect protein biomarkers of disease by ELISA, Sandwich ELISA for measuring disease associated proteins, diagnosing Autoimmune diseases by indirect ELISA, Immunoassays for Infectious disease, Protein arrays to detect Polygenic disorder.	15		

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Name of Proorage	Stain	
e of P	ster	Jame of the
Nan	Semester	Name

	DNA based approaches to disease diagnosis:	: Hybri	dization probes -11 t	7	
	specific hybridization, Oligonucleotide ligation specific PCR, Real-time PCR to detect infectious disease associated mutations using Microarrays.	acan.	Day 11		
111	Introduction to Metabolic disambana I.B.		90		
	Introduction to Metabolic disorders and Profit Disorders in Hormonal Action. Insulin dependent Ligand induced signalling and gene expression in of Intracellular Trafficking and its related pathogon Gene therapy: As a potential tool to cure endocrinology in health and disease. Cancer a molecules in medicine.	ent and cukary enesis.	yotic cells. Importance	15	
IV	molecules in medicine.	and ce	Il cycle, Recombinant		
	Free Radicals and Metal ions in Medicine: Mechanisms of lipid, protein and DNA oxidation, Antioxidants-small molecules and enzymes, Reactive Oxygen Intermediates (ROI), Transition metals in oxidative processes, Involvement of oxidative processes in ageing, cancer and atherosclerosis, Metal Ions in Gene Regulation: Iron in human diseases-anaemia, and thalassemia, Metals and free radicals in Alzheimer's disease and other neurodegenerative diseases (NDDs).				
	Suggested Evolution 1 Total Contact Hours			60	
	Suggested Evaluation Internal Assessment: 30	Method	ds	100	
	Theory		End Term Examina	tion: 70	
	* Class Participation	30	Theory	70	
	Seminar/Presentation / A	05	Written examination	1,0	
	Seminar/Presentation/Assignment/Quiz or Class test, etc.	10	- Indiana		
	❖ Mid-Term Exam				
	T Wild I Cliff Exam	15			

- 1. Medical Biotechnology. Glick BR, Delovitch TL and Pattern CL (2014). ASM press.
- 2. Molecular Diagnosis of Genetic Diseases. Rob Elles (2003). (2nd Edi), Humana Press (2003).
- 3. Introduction to Molecular Medicine. Dennis Ross (2002). (3rd Edi.), Springer.
- 4. Molecular Medicine: Genomics to Personalized Healthcare. Tent RJ (2012). Academic Press.
- 5. Principles of Molecular Medicine. Runge, Marschall S, Patterson (2006). (2nd Edi.), Humana Press.
- 6. Medical Biotechnology. Judit P & M Keen (2009). 1st Edi. Elsevier Pub.
- 7. Medical Biotechnology. Jogdand SN (2011). 2<sup>nd</sup> Edi. Himalaya Publishers.
- 8. Biotechnology-Applying the genetic Revolution. Clark and Pazdernik (2009). Academic Press
- 9. Basic & Clinical Pharmacology. Bartram G Katzung (2004). 9th Edi. Mc Graw Hill Publications.
- 10. Text book of biochemistry with Clinical Correlations. Devlin TM (2002). 5th Edi.



Part A-Intro					
	duction				
	M.Sc. Biotechnology	1			
	III				
Plant d	& Microbial Biotechnolo	gy (Lab)			
Course Code M24-BTY-307					
Course Type PC-5					
evel of the course 500-599					
	NA				
and techniques in fid will be able to und organismal, cellular, CLO-2: Analyse and and plant biotechnolog for various products a CLO-3: Understand plant biotechnology to CLO-4: Imbibe the	eld of microbial & plant erstand various biologic biochemical and moleculed solve various problems ogy, launch start-ups and processes.  bio-safety measures related biological processes and processes with the processes of the processes of the processes of the plant	biotechno al aspects ar biologica ar related to become ent ated to mice.	related to al. microbia repreneurs crobial and		
-110019			4		
			8		
0.55	30	30			
0	70	70			
0	100	100			
	nts of the Course				
Practical's			Contact Hours		
Practical Exerc	eises	*	120		
, Fermentation.			120		
eer, ethanol.					
y important micro-or	ganisms.				
lulolytic and pectinol	ytic micro-organisms.				
		*			
		activity			
tion system for transfe	ormants.				
			2		
	leaves tissue				
	CLO-1: Develop pra and techniques in fie will be able to und organismal, cellular, CLO-2: Analyse and and plant biotechnolog for various products at CLO-3: Understand plant biotechnology to CLO-4: Imbibe the independently to write Theory  0  0  0  Part B-Content Practical Exerce Fermentation.  cer, ethanol.  y important micro-or lulolytic and pectinol ipase/amylase product producing micro-org lanase/Cellulase/Pectin stick transformation.  So assay.  nomic DNA from the	Plant & Microbial Biotechnolo  M24-BTY-307  PC-5  500-599  NA  CLO-1: Develop practical skill & acquaint wand techniques in field of microbial & plant will be able to understand various biologic organismal, cellular, biochemical and molecular clooped and plant biotechnology, launch start-ups and for various products and processes.  CLO-2: Analyse and solve various problems and plant biotechnology techniques.  CLO-3: Understand bio-safety measures relaplant biotechnology techniques.  CLO-4: Imbibe the value of team spirit independently to write and manage their researed to the company of the company	Plant & Microbial Biotechnology (Lab)  M24-BTY-307  PC-5  500-599  NA  CLO-1: Develop practical skill & acquaint with recent and techniques in field of microbial & plant biotechno will be able to understand various biological aspects organismal, cellular, biochemical and molecular biological cand plant biotechnology, launch start-ups and become ent for various products and processes.  CLO-2: Analyse and solve various problems related to and plant biotechnology, launch start-ups and become ent for various products and processes.  CLO-3: Understand bio-safety measures related to mice plant biotechnology techniques.  CLO-4: Imbibe the value of team spirit & as well independently to write and manage their research experim Theory Practical To 0 4 5 5 6 7 7 6 7 7 7 7 7 7 7 7 7 7 7 7 7 7		

13. Restriction digestion of plant genomic DNA.			T Z
14. Transgene detection by amplification.			
15. Southern blotting of DNA.			
16. Plants crude extracts preparation from plant ti	issues.		
17. Isolation of essential oils from plant tissues.			
		<b>Total Contact Hours</b>	120
Suggested Evaluation	Method	ds	
Internal Assessment: 30		End Term Examina	tion: 70
Practicum	30	Practicum	70
❖ Class Participation	05	Lab record, Viva-Voc	
Seminar/Demonstration/Viva-Voce/Lab records, etc.	10	and Execution of the I	Practical
Mid-Term Exam	15		

- 1 Introduction to Biotechnology. HS Chawla (2009), 3rd Ed., Science Pub.
- 2. Plant Tissue Culture Manual. Lindsey (2007). Springer (India) Publication.
- Molecular Cloning: A Laboratory Manual. Sambrook & Maniatis (2002). Cold Spring Harbor Laboratory Press, New York.

Part C-Learning Resources

- 4. Plant Biotechnology: Laboratory Manual. Chawla HS (2008). Oxford & IBH Publishing Co. Pvt. Ltd. India.
- Molecular Cloning: A Laboratory Manual. Sambrook & Russell (2007) 3<sup>rd</sup> Edi. (Vol.1-3). Cold Spring Harbor Laboratory Press, New York.
- 6. Manual of Industrial Microbiology & Biotechnology. Demain & Davies (1999) 2<sup>nd</sup> Edi. ASM Press, Washington D.C.
- 7. Practical Manual of Biotechnology: Applied Sciences. Mahajan R, et al. (2010). Vayu Education of India.
- 8. Microbiology: A Laboratory Manual. Cappuccino & Welsh (2016). 11th Edi. Pearson Education Ltd.

	Session:	2025-26				
Name of Programme	Part A-Int	roduction				
Semester		Biotechnology				
Name of the Course		III				
Tame of the Course	Molecular Ger	netics & Immunology/M	olecular Me	edicine &		
Course Code		Diagnostic Techniques	(Lab)			
Course Type		M24-BTY-308				
Level of the course	el of the games					
Pre-requisite for the course		500-599				
(if any) Course Learning Outcomes		NA				
(CLO). After completing this course, the learner will be able to:	plating, metaphase various assays such of genotoxicity to de CLO-2: Work will detection, DNA Flymphocytes, detern medical fields. CLO-3: Get trained different diseases. CLO-4: Get acquaestimation of antiger	Practical	n, banding onucleus as  PCR-RFLP of periphe for use in concept the detailed and ques and questive and questiv	techniques, biomarkers for SNP eral blood clinical and etection of quantitative		
	0	4		4		
	U			4		
eaching Hours per week	0	8		8		
nternal Assessment Marks	0	8 30		-17.		
nternal Assessment Marks and Term Exam Marks	0 0 0	8 30 70		8 30 70		
nternal Assessment Marks and Term Exam Marks ax. Marks	0	8 30 70 100		8		
nternal Assessment Marks and Term Exam Marks	0 0 0 0	8 30 70 100 4 Hours		8 30 70		
nternal Assessment Marks and Term Exam Marks ax. Marks	0 0 0 0 Part B-Conter	8 30 70 100		8 30 70 00		
nternal Assessment Marks and Term Exam Marks ax. Marks	0 0 0 0	8 30 70 100 4 Hours		8 30 70		
nternal Assessment Marks and Term Exam Marks ax. Marks	0 0 0 0 Part B-Conter	8 30 70 100 4 Hours ats of the Course		8 30 70 00 Contact Hours		
nternal Assessment Marks and Term Exam Marks ax. Marks amination Time	0 0 0 0 Part B-Conter Practicals	8 30 70 100 4 Hours ats of the Course		8 30 70 00 Contact		
nternal Assessment Marks and Term Exam Marks ax. Marks amination Time	Part B-Conter Practicals  Practical Exerced mutations.	8 30 70 100 4 Hours nts of the Course	1	8 30 70 00 Contact Hours		
nternal Assessment Marks and Term Exam Marks ax. Marks amination Time  1. Spontaneous and induce 2. Metaphase chromosome	Part B-Conter Practicals  Practical Exerced mutations.	30 70 100 4 Hours nts of the Course ises	ues.	8 30 70 00 Contact Hours		
nternal Assessment Marks and Term Exam Marks ax. Marks amination Time  1. Spontaneous and induce 2. Metaphase chromosome 3. Sister chromatid excha	Part B-Conter Practicals  Practical Exerced mutations.	30 70 100 4 Hours nts of the Course ises	ues.	8 30 70 00 Contact Hours		
aternal Assessment Marks and Term Exam Marks ax. Marks amination Time  1. Spontaneous and induce 2. Metaphase chromosome 3. Sister chromatid excha	Part B-Conter Practicals  Practical Exerced mutations.  e preparation, chrominge assay using points.	8 30 70 100 4 Hours ats of the Course  ises  osome banding techniqueripheral blood lympho	ues.	8 30 70 00 Contact Hours		
1. Spontaneous and induce 2. Metaphase chromosome 3. Sister chromatid excha genotoxicity studies. 4. Single Cell Gel Electrop	Part B-Conter Practicals  Practical Exerced mutations.  Preparation, chromonge assay using positions to detect DN	8 30 70 100 4 Hours ats of the Course sees sees osome banding technique eripheral blood lympho	ues.	8 30 70 00 Contact Hours		
1. Spontaneous and induce 2. Metaphase chromosome 3. Sister chromatid excha genotoxicity studies. 4. Single Cell Gel Electrop	Part B-Conter Practicals  Practical Exerced mutations.  Preparation, chromonge assay using positions to detect DN	8 30 70 100 4 Hours ats of the Course sees sees osome banding technique eripheral blood lympho	ues.	8 30 70 00 Contact Hours		
1. Spontaneous and induce 2. Metaphase chromosome 3. Sister chromatid excha	Part B-Conter Practicals  Practical Exerced mutations.  Preparation, chromonge assay using positions to detect DN	8 30 70 100 4 Hours ats of the Course sees sees osome banding technique eripheral blood lympho	ues.	8 30 70 00 Contact Hours		
1. Spontaneous and induce 2. Metaphase chromosome 3. Sister chromatid excha genotoxicity studies. 4. Single Cell Gel Electrop 5. Analysis of Micronuc epithelial cells.	Part B-Conter Practicals  Practical Exerced mutations.  Preparation, chromonge assay using position properties to detect DN cleus as biomarker	8 30 70 100 4 Hours ats of the Course sees sees osome banding technique eripheral blood lympho	ues.	8 30 70 00 Contact Hours		
1. Spontaneous and induce 2. Metaphase chromosome 3. Sister chromatid excha genotoxicity studies. 4. Single Cell Gel Electrop 5. Analysis of Micronuc epithelial cells. 6. To determine IC <sub>50</sub> of a to	Part B-Conter Practicals  Practical Exerced mutations. The preparation, chromonge assay using position process to detect DN cleus as biomarker exic compound.	30 70 100 4 Hours  At sof the Course  ises  osome banding technique ripheral blood lympho	ues.	8 30 70 00 Contact Hours		
1. Spontaneous and induce 2. Metaphase chromosome 3. Sister chromatid excha genotoxicity studies. 4. Single Cell Gel Electrop 5. Analysis of Micronuc epithelial cells. 6. To determine TLC and I	Part B-Conter Practicals  Practical Exerced mutations.  Preparation, chromonge assay using position properties to detect DN celeus as biomarker exic compound.  DLC in human blood	8 30 70 100 4 Hours  ats of the Course  ises  osome banding technique ripheral blood lympho  NA damage.  of genotoxicity usin	ues.	8 30 70 00 Contact Hours		
1. Spontaneous and induce 2. Metaphase chromosome 3. Sister chromatid excha genotoxicity studies. 4. Single Cell Gel Electrop 5. Analysis of Micronuc epithelial cells. 6. To determine IC <sub>50</sub> of a to 7. To determine TLC and II 8. Isolation of Lymphocyte	Part B-Conter Practicals  Practical Exerced mutations. The preparation, chromonge assay using position process to detect DN cleus as biomarker oxic compound. The process of the process o	8 30 70 100 4 Hours  Its of the Course  ises  osome banding technique ripheral blood lympho  NA damage.  of genotoxicity using the series of genotoxicity using the s	ues. Deytes for	8 30 70 00 Contact Hours		
1. Spontaneous and induce 2. Metaphase chromosome 3. Sister chromatid exchargenotoxicity studies. 4. Single Cell Gel Electrop 5. Analysis of Micronuce epithelial cells. 6. To determine IC50 of a tellocation of Lymphocyte 9. Serum preparation and serial cells.	Part B-Conter Practicals  Practical Exerced mutations.  Preparation, chromoninge assay using positions to detect DN cleus as biomarker exic compound.  DLC in human blooders from peripheral blooders from peripheral blooders of the periphe	8 30 70 100 4 Hours  Its of the Course  ises  osome banding technique ripheral blood lympho  A damage.  of genotoxicity using the smear.  od.  Agglutination and Preceived.	ues. Deytes for	8 30 70 00 Contact Hours		
1. Spontaneous and induce 2. Metaphase chromosome 3. Sister chromatid excha genotoxicity studies. 4. Single Cell Gel Electrop 5. Analysis of Micronuc epithelial cells. 6. To determine IC <sub>50</sub> of a to 7. To determine TLC and II 8. Isolation of Lymphocyte	Part B-Conter Practicals  Practical Exerced mutations.  Preparation, chromoninge assay using positions to detect DN cleus as biomarker exic compound.  DLC in human blooders from peripheral blooders from peripheral blooders of the periphe	8 30 70 100 4 Hours  Its of the Course  ises  osome banding technique ripheral blood lympho  A damage.  of genotoxicity using the smear.  od.  Agglutination and Preceived.	ues. Deytes for	8 30 70 00 Contact Hours		

or double).  12. To perform immuno-electrophoresis with a g 13. To perform DNA fingerprinting analysis.  14. PCR-RFFLP for SNP detection.	iven an	tigen-antibody system.	
Constant I Day		<b>Total Contact Hours</b>	30
Suggested Evaluation Internal Assessment: 30	Metho		
Practicum Practicum		End Term Examinat	ion: 70
	30	Practicum	70
Class Participation	05	Lab record, Viva-Voc	e Write-III
<ul> <li>Seminar/Demonstration/Viva-Voce/Lab records, etc.</li> </ul>	10	and Execution of the I	Practical
<ul> <li>Mid-Term Exam</li> </ul>	15		
Part C-Learning Re		s	

- 1. Introductory Practical Biochemistry. Sawhney & Singh (2005). Alpha Science Publishing Pvt. Ltd. India.
- 2. Principles & Techniques of Biochemistry & Molecular Biology. Wilson & Walker.
- 3. Practical Manual of Biotechnology: Applied Sciences. Mahajan R, et al. (2010). Vayu Education of India.
- 4. Molecular Biology Techniques: Susan (2nd Edi.) Academic Press
- 5. Molecular Biology: Principles and Practice. Michael et al., (3rd Edi.). WH Freeman.
- 6. Experiments in Microbial Genetics. Tortora (4th Edi.). John Wiley & Sons, Inc.
- 7. Medical Biotechnology: Techniques & Applications. Raymond & Wegner (2nd Edi.). Academic Press.
- 8. Cytogenetic Laboratory Management: Chromosomal, FISH and Microarray-Based Best Practices and Procedures. Susan Mahler Zneimer.



	Session:	2025-26			
Name of Programme	Part A-Int				
Semester	M.Sc. Biotechnology				
Name of the Course					
Course Code	Bioto	chnology & Human W	'elfare		
Course Type		M24-OEC-303			
Level of the course		OEC			
Pro requisite for all		500-599			
Pre-requisite for the course (if any)		NA			
Course Learning Outcomes (CLO). After completing this course, the learner will be able to:	biotechnology. Lear processing, uses of enzyme engineering CLO-2: Learn about and bioremediation biosensors and biofu CLO-3: Understand vaccines, gene therapy CLO-4: Get acquaints	n about basics of ferme microbes, industrial app at role of biotechnology Describe various con- els. d basic concepts of by, DNA fingerprinting, inted with the uses of ted with the latest knowled will be able to so	vledge of different areas		
Credits	Theory	Practical	Total		
Cicuits	2	0	2		
Teaching Hours per week	2	0	2		
Internal Assessment Marks	15	0	15		
End Term Exam Marks	35	0	35		
Max. Marks	50	0	50		
Examination Time	3 Hours				
ZAMITIMETICII A MATE	Part B-Conten	ts of the Course			

Instructions for Paper- Setter: The examiner will set 9 questions asking two questions from each unit and one compulsory question by taking CLOs into consideration. The compulsory question (Question No. 1) will consist at least 4 parts covering entire syllabus. The examinee will be required to attempt 5 questions, selecting one question from each unit and the compulsory question. All questions will carry equal marks.

Units	Topics			
I	Industrial Biotechnology: Introduction, Isolation and screening of microbes, approaches for strain development, Production of organic compounds, enzymes and antibiotics by microbes, Types of Fermentation, Downstream processing Enzyme immobilization, Industrial applications of enzymes, Protein and enzyme engineering			
11	Environmental Biotechnology: Role of Biotechnology in the treatment of waste water, Solid waste management using biotech approaches, Bioremediation: Concept and principles, Bioremediation using microbes and plants, Biofuels, Biosensors.	06		

Ш	Animal and Medical Biotechnology: Molecu probes, PCR to detect infectious diseases, N	lar Dia Ionoclo	agnostics- DNA/RNA onal antibodics- their	08	
	Production and applications.  Vaccines: live, attenuated, killed, subunit, conjugated the Conference of the Conference o	ate and nentations. Tran	DNA vaccines. on & Targeted Gene asgenic animals- mice,		
IV	Agricultural Biotechnology: Transgenic plants for biotic (insects, herbicide, fungal and viral resistance) and abiotic stress tolerance. Nutritional quality modifications in crop plants, Molecular Farming (Medically Related Proteins - edible vaccines, plantibodies, etc.), Plant secondary metabolites.				
	Total Contact Hours				
	Suggested Evaluation Methods				
	Internal Assessment: 15		End Term Examinat	tion: 35	
	Theory	15	Theory	35	
	<ul> <li>Class Participation</li> </ul>	04	Written examination	*	
	<ul> <li>Seminar/Presentation/Assignment/Quiz or Class test, etc.</li> </ul>	04			
	<ul> <li>Mid-Term Exam</li> </ul>	07			
	Part C-Learning Res				

- 1. Biotechnology: Expanding Horizon. BD Singh (2010). 3rd Edi. Kalyani Publishers.
- 2. Biotechnology and Genomics. PK Gupta (2013). 1st Edi. Rastogi Publishers.
- 3. Applying Genetic Revolution. Clark DV and Pazdernik NJ (2009). Academic Press.
- 4. Hand book of Plant Biotechnology. Gistou P and Klu H (2004). John Publication.
- 5. Plant Biotechnology: Halford NG (2006). John Wiely Publishers.
- 6. Animal Biotechnology. Ballinic, Philips JP & Moo Young M (1989). Pergamon Press, NY.
- 7. Molecular Biology of Gene. Watson JD et al. (2007) 6th Edi. Publisher Benjamin Cummings.
- 8. Basic Biotechnology. Ratlege C & B Kristiansen (2001). Cambridge Univ. Press, London.
- 9. Microbial Biotechnology. Glazer and Nikaido, WH Freeman & Company, New York.
- 10. Biotechnology in crop improvement. HS Chawla. International Book Distributing Company.
- 11. Genetics to Gene Therapy: Mol. Pathology of Human Disease. 1st Edi (1994). BIOS Scientific Pub.
- 12. Medical Biotechnology. Jogdand SN (2011). 2nd Edi. Himalaya Publishers.
- 13. Introduction to Nanobiotechnology. Niemeyer and Mirkin (2003). Wiley, VCH-Publishers.
- 14. Medical Biotechnology. Glick BR, Delovitch and Patten (2014). ASM Press.
- 18. Lehninger Principles of Biochemistry. Nelson & Cox. Freeman & Company, NY.
- 19. Principles of Fermentation Technology. Stansbury PF et al. (1997). Pergmon Press, Oxford.
- 20. Biotechnology: A Textbook of Industrial Microbiology. Cruger & Cruger (2002), 2nd Edition.



## Chaudhary Ranbir Singh University, Jind

Syllabus for M.Sc. Biotechnology (CBCS) (IV-Semester)

	Session:	2025-26	
	Part A-Inti	roduction	
Name of Programme		M.Sc. Biotechnology	
Semester		IV	<b>C</b> arriery
Name of the Course		Animal & Medical Biotechi	nology
Course Code		M24-BTY-401	
Course Type		CC-11	
Level of course (Anx-1)		500-599	
Pre-requisite for the course (if any)	la l	NA s of animal cloning, embryo tran eir applications for human welfare.	sfer, production
Course Learning Outcomes	is animals and the	oir applications for human	theranculus
course, the learner will be	CLO-2: Gain thorough us role of stem cells in biome CLO-3: Learn advance	nderstanding of Nucleic acid & pedical research, gene therapy & DN ed techniques such as nanob	NA fingerprinting intechnology and intechnology.
course, the learner will be	CLO-2: Gain thorough use role of stem cells in biomed CLO-3: Learn advance pharmacogenomics and gas CLO-4: Have a broad uncomplete the classification of the complete the classification of the classi	nderstanding of Nucleic acid & pedical research, gene therapy & DN ed techniques such as nanob ain insight into clinical aspects of Ederstanding of the animal and biom ons and explore their ideas of new or an explore the explore their ideas of new or an explore their ideas of new or an e	NA fingerprinting intechnology and intechnology.
course, the learner will be able to:	CLO-2: Gain thorough us role of stem cells in biome CLO-3: Learn advance	nderstanding of Nucleic acid & pedical research, gene therapy & DN ed techniques such as nanobain insight into clinical aspects of Ederstanding of the animal and biomons and explore their ideas of new y.  Practical	NA fingerprinting iotechnology and Biotechnology edical research for wision of anim  Total
course, the learner will be able to:	CLO-2: Gain thorough use role of stem cells in biomed CLO-3: Learn advance pharmacogenomics and gas CLO-4: Have a broad uncomposed biotechnological application and medical biotechnology.	nderstanding of Nucleic acid & pedical research, gene therapy & DN ed techniques such as nanobain insight into clinical aspects of Ederstanding of the animal and biomons and explore their ideas of new practical	JA fingerprinting intechnology and intechnology. edical research for vision of anim  Total  4
course, the learner will be able to:  Credits	CLO-2: Gain thorough use role of stem cells in biomes CLO-3: Learn advance pharmacogenomics and gas CLO-4: Have a broad unch biotechnological application and medical biotechnology.  Theory  4	nderstanding of Nucleic acid & pedical research, gene therapy & DN ed techniques such as nanobain insight into clinical aspects of Ederstanding of the animal and biomons and explore their ideas of new y.  Practical	NA fingerprinting intechnology and intechnology and intechnology, edical research for vision of anim  Total  4  30
Credits  Teaching Hours per week Internal Assessment Marks	CLO-2: Gain thorough use role of stem cells in biomes CLO-3: Learn advance pharmacogenomics and gase CLO-4: Have a broad uncomposed biotechnological application and medical biotechnology.  Theory  4  4  30	nderstanding of Nucleic acid & Pedical research, gene therapy & DNeed techniques such as nanobain insight into clinical aspects of Ederstanding of the animal and biomons and explore their ideas of new Practical  O  O	NA fingerprinting intechnology and intechnology are dical research for wision of anim.  Total  4  30  70
Credits  Teaching Hours per week Internal Assessment Marks	CLO-2: Gain thorough use role of stem cells in biomed CLO-3: Learn advance pharmacogenomics and gase CLO-4: Have a broad uncomposed biotechnological application and medical biotechnology.  Theory  4  30  70	nderstanding of Nucleic acid & Pedical research, gene therapy & DNeed techniques such as nanobain insight into clinical aspects of Ederstanding of the animal and biomons and explore their ideas of new Practical  O  O  O	NA fingerprinting intechnology and intechnology are dical research for vision of animal states of the states of th
Credits  Teaching Hours per week Internal Assessment Marks End Term Exam Marks Max. Marks Examination Time	CLO-2: Gain thorough use role of stem cells in biomes CLO-3: Learn advance pharmacogenomics and gase CLO-4: Have a broad uncomposed biotechnological application and medical biotechnology.  Theory  4  4  30	nderstanding of Nucleic acid & Pedical research, gene therapy & DNeed techniques such as nanobain insight into clinical aspects of Ederstanding of the animal and biomons and explore their ideas of new Practical  O  O  O	NA fingerprinting intechnology and intechnology are dical research for wision of anim.  Total  4  30  70

Instructions for Paper- Setter: The examiner will set 9 questions asking two questions from each unit and one compulsory question by taking CLOs into consideration. The compulsory question (Question No. 1) will consist at least 4 parts covering entire syllabus. The examinee will be required to attempt 5 questions, selecting one question from each unit and the compulsory question. All questions will carry equal marks.

questio Units	ns will carry equal marks.  Topics	Contact Hours
1	Introduction to Animal Biotechnology: Scope, global perspective & new horizons, economically important livestock breeds, Model animals in animal biotechnology & genetic engineering; An overview of animal cell culture techniques-cell lines, cell culture, cell viability assays, cryopreservation of cells Transgenic Animals: Principles of transgenesis; Methods of gene transfer-DNA microinjection, Retroviral & embryonic stem cell methods, Electroporation, Biolistic, lipofection; selectable markers, Application of transgenic animals-mice, sheep, pigs, goats, cows, fish; Molecular pharming, Case studies of transgenic animal models.	
11	Animal Cloning: Concept of animal cloning, cloning from embryonic and adult cells, Somatic cell nuclear transfer technique, Embryo splitting, Creation of Dolly, Molly and Polly, challenges and limitations; applications of animal cloning.  Embryo Transfer Technology: Definition, Super-ovulation, Artificial Insemination, In-vitro fertilization, embryo evaluation, embryo transfer in cattle,	16

	hematopoietic, mesenchymal and neural stem ce of pluripotent and multipotent stem cells, i	induced	I pluripotent stem cells.	
	therapeutic cloning for embryonic stem cells, clinical applications.			
III	Nucleic Acid Therapeutics: Anti-sense RNA, R RNAi, Zinc Finger Nucleases			1
	Growth Hormone, Interferon. etc.), Recombinant Antibodies, Hybrid Human Mouse Mond Antibodies), Enzymes (DNase, Alginate Lyase Ammonia Lyase, Glycosidases)	clonal e, Alph	Antibody, Anticancer na 1 Antitrypsin, Phenyl	
	Gene Therapy: Definition, Types of gene th inhibition, Gene editing; <i>in-vitro</i> and <i>in-vivo</i> g vectors for gene transfer, gene therapy for SCID Ethical issues.	, Cance	erapy, viral and nonviral er, Neurological disorders,	
IV	Nanobiotechnology: Introduction, Types and Sy based Nanostructures, applications of Nanopar Gene Delivery, Disease Diagnostics & Therapy, I Pharmacogenomics: Concept, Role of Genetic of individuals to drugs, Pharmacogenomics and DNA Fingerprinting in Forensic Sciences.	nthesis ticles, Risk Po	of Nanoparticles, Protein Nanobiosensors, Drug & stential of Nanomaterials	13
	Suggested Evaluation	14.1	Total Contact Hours	60
	Assessment: 30	Method		
	Theory	20	End Term Examination:	70
	Class Participation	30	Theory 70	
	<ul> <li>Seminar/Presentation/Assignment/Quiz or Class test, etc.</li> </ul>	05 10	Written examination	
	Ul Class test oto			

- 1. Culture of Animal Cells: A Manual of Basic Technique. Freshney (2016). 7th Edi. Wiley-Blackwell.
- 2. Animal Biotechnology: Ranga MM (2018). 3rd Edi. Agrobios (2018).
- 3. Medical Biotechnology. Glick BR, Delovitch and Patten (2014). ASM Press.
- 4. Stem Cell Biology. Marshak L (2001). Cold Spring Harbor.
- 5. Medical Biotechnology. Judit and Keen (2009). 1st Edi. Elsevier Publications.
- 6. Medical Biotechnology. Jogdand SN (2011). 2nd Edi. Himalaya Publishers.
- 7. Biotechnology: Applying the Genetic Revolution. Clark and Pazdernik (2009). Academic Press.
- 8. Concepts in Biotechnology. Balasubramanian & Dharmalingam (2004). 2nd Edi. University Press.
- 9. Biotechnology. Satyanarayan (2008). Books and Allied (Pvt.) Ltd.
- 10. Biotechnology: Expanding Horizon. BD Singh (2010). 3rd Edi. Kalyani Publishers.
- 11. Biotechnology and Genomics. Gupta PK (2013). Rastogi Publications Meerut, India
- 12. Introduction to Nanobiotechnology. Niemeyer and Mirkin (2003). Wiley VCH, Publishers.
- 13. Principles of Gene manipulation and Genomics. Primose SB and Twyman RM (2006). 7th Edi. Blackwell
- 14. Basic & Clinical Pharmacology. Bartram Katzung (2004). 9th Edi. Mc Graw Hill Publications.



	Session: 20	25-26	
Name of Programme	Part A-Intro		
Semester		M.Sc. Biotechnology	
Name of the Course		IV	
Course Code		Environmental Biotechnolog	gy
Course Type		M24-BTY-402	
evel of the course		CC-12	
re-requisite for the		500-599	
Course Learning Outcomes		NA	
(CLO). After completing	Biotechnology with spe	cial emphasis on the role	of microbes if
this course, the learner will be able to:	mitigating environment particles and particles are management, gaining known sewage treatment and solic CLO-3: Understand and environmental pollutants recovery.  CLO-4: Understand appropriation of the particles are management and pollutants recovery.	ollution as well as potability ole of microbes in solid an nowledge of various method d waste treatment. ole of microbes in bio & also utility of microbes in olications of Biotechnology	of water and its  nd liquid waste ds employed in  oremediation of in mineral & of
inis course, the learner will	mitigating environment particles and particles are management, gaining known and control a	ollution as well as potability ole of microbes in solid an nowledge of various method d waste treatment. role of microbes in bio & also utility of microbes in	of water and its  nd liquid waste ds employed in  remediation of in mineral & of  in environmen
Credits	mitigating environment particles and particles are management, gaining known sewage treatment and solution of the control of t	ollution as well as potability ole of microbes in solid an nowledge of various method d waste treatment. Tole of microbes in bio & also utility of microbes in olications of Biotechnology  Practical	of water and it nd liquid wast ds employed in remediation of in mineral & of in environment  Total 4
Credits  Teaching Hours per week	mitigating environment particles and particles are management, gaining known and solutions are management, gaining known as a sewage treatment and solutions.  CLO-3: Understand a particle covery.  CLO-4: Understand appropriate appropriate and solutions.  Theory  4	ollution as well as potability ole of microbes in solid and anowledge of various method distributed waste treatment. Tole of microbes in bio & also utility of microbes in bio blications of Biotechnology  Practical	of water and it  nd liquid wast ds employed i  oremediation of in mineral & o  in environmer  Total  4  30
Credits  Teaching Hours per week Internal Assessment Marks	mitigating environment particles and particles are management, gaining known sewage treatment and solution of the control of t	ollution as well as potability ole of microbes in solid and anowledge of various method waste treatment. Tole of microbes in bio & also utility of microbes in bio olications of Biotechnology  Practical  0 0 0 0	of water and it  nd liquid wast ds employed i  oremediation of in mineral & o  in environmer  Total 4 4 30 70
Credits  Teaching Hours per week	mitigating environment particles and particles are management, gaining know sewage treatment and solic CLO-3: Understand and environmental pollutants recovery.  CLO-4: Understand appropriate appropriate and particles are monitoring.  Theory  4  4  30	ollution as well as potability ole of microbes in solid and anowledge of various method distributed waste treatment. Tole of microbes in bio & also utility of microbes in bio olications of Biotechnology  Practical  0 0 0	of water and it nd liquid wast ds employed in oremediation of in mineral & of in environment  Total 4 4 30

Instructions for Paper- Setter: The examiner will set 9 questions asking two questions from each unit and one compulsory question by taking CLOs into consideration. The compulsory question (Question No. 1) will consist at least 4 parts covering entire syllabus. The examinee will be required to attempt 5 questions, selecting one question from each unit and the compulsory question. All questions will carry equal marks.

Jnits	Topics Topics					
I	Environmental Biotechnology: An Overview, Concept, Scope & Market Biological Control of Air-Pollution, Bacterial examination of H <sub>2</sub> O for Potability, testing of H <sub>2</sub> O for Physiochemical Parameters including BOD & COD.  Solid Waste: Sources and Management (Composting, Vermi-composting and Methane Production).	15				
П	Waste Water: Origin, Composition and Treatment. Physical, Chemical and Biological Treatment of Waste Water.  Aerobic Processes: Activated Sludge, Oxidation Ponds, Trickling Filter Towers and Rotating Discs.	16				

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Old.	10 2	100	5/
-	24	IPN/	4
	_	V	100

	Control of the contro		
	Sludge Blanket Reactors. Microbiology and Biochemistry of Acrobic and	nemistry of Acrobic an	
	Anacrobic Waste Water Treatment Processes.		;
	Treatment of Industrial Effluents: Distillery Effluent, Paper and Pulp Mill	ient, Paper and Pulp M	=
	Effluent, Tannary Effluent, Textile Dye Effluent, Removal of Heavy Metals	cmoval of Heavy Meta	S
	from Waste Waters.		
Ξ	Bioremediation: Introduction of Bioremediation, Advantages & Applications,	Ivantages & Application	
	Types of Bioremediation, Natural (Attenuation), Ex-situ & In-situ,	n), Ex-situ & In-sit.	(, 1)
	Bioaugmentation and Biostimulation, Solid Phase and Slurry Phase	ase and Slurry Phas	9
	Bioremediation.		
	Biodegradation: Acrobic versus Anacrobic Degradation Microbial basis of	lation Microbial basis of	4
	Biodegradation, Biodegradation of Xenobiotics. Microbial Degradation of	dicrobial Degradation of	
	Pesticides.	a manning a management	
	Biotechnological Methods of Pollution Detection: General Biogenie in	n. Gonoral Bioaccase	
	Pollution Monitoring, Cell Biology in Environmental Manifornia, Material	al Monitoring Melecular	
	Biology in Environmental Monitoring & Biosensors in Environmental Control of the	a Phylinoring, Molecula	
>	Microbial Insecticides: Bacteria, Funoi and Viruses The of Town	ir Erry II Official and Jysis	-
	Technology to enhance efficacy Microbial Insecticides Bio-fartificant	ecticides Bio-fortilizare	
	Microbes in oil Recovery and Bioleaching.	DIO-101 IIIZEIS	
	Bio-deterioration: Bio-deterioration of stored Plant Ecod Matarial I	Food Matanials 1	
	Wool, Metals, Textiles, Stone & Related Building Control of Misself Electher,	Control of Mignetical De	
	deterioration.	Control of Milcrobial Bio	
		Total Contact Hours	09
	Suggested Evaluation Methods	hods	1
	Internal Assessment: 30	End Torm Evamination 70	70
	Theory	30 Theory	70 70
	❖ Class Participation		0/
	tion/Assignment/Quiz		_
		*	
	❖ Mid-Term Exam		

- 1. Environmental Biotechnology: Principles and Applications. Bruce & McCarty (2020), 2nd Edi. Mc Graw Hills Publisher.
- 2. Introduction to Biodeterioration. Allsopp and Scal. ELBS/Edward Arnold Publisher.
- 3. Advanced Environmental Biotechnology. SK. Agarwal (2005). APH Publishing, New Delhi.
  - 4. Environmental Biotechnology: Biodegradation, Bioremediation, Sustainable Development. Jeyabalan & Abdullah (2016). Apple Publisher Academic Press.
- 5. Environmental Science and Technology. Stankey EM (1997). Lewis Publishers, New York.
- 6. Microbial Biotechnology: Basic Research and Applications Singh et al. (2020). Pub. Springer
  - 7. Biodegradation and Bioremediation: Soil Biology. Singh & Ward (2004). Pub. Springer

	Session: 2	2025-26					
21	Part A-Intr	oduction					
Name of Programme		M.Sc. Biotechnolog	***				
Semester		IV	gy				
Name of the Course	ame of the Course Food Biotechnology						
Course Code			<u> </u>				
Course Type		M24-BTY-403					
Level of the course		DEC-3					
Pre-requisite for the		500-599					
course (if any)		NA					
Course Learning Outcome (CLO). After completing this course, the learner will be able to:	recent theoretical knowled processing of biotech foods CLO-2: Comprehend about food industry for shelf-life CLO-3: Gain the knowledge interaction with food production of the process of the communication of the communica	edge and techniques research and supplements.  The food additives the extension, processing an edge of food packaginucts. They would be all huge different types on the edge of preservation techniques of food safetiances related to food be savely integrated per savely integrated.	at are relevant to processed and sensory appeal.  ag, its importance and its ole to launch start-ups and processes are products are products and processes are products and processes are products are products are products and processes are products and processes are products and processes are products and processes are products are products and processes are products are products and processes are products and processes are products are products are products and processes are products are				
Credits	Theory		4				
	4	0	The state of the s				
Feaching Hours per week	4	0	4				
Teaching Hours per week	30	0	30				
nternal Assessment Marks	30 70	0	30 70				
Teaching Hours per week Internal Assessment Marks and Term Exam Marks Max. Marks	30	0	30				
nternal Assessment Marks and Term Exam Marks	30 70	0	30 70				

Instructions for Paper- Setter: The examiner will set 9 questions asking two questions from each unit and one compulsory question by taking CLOs into consideration. The compulsory question (Question No. 1) will consist at least 4 parts covering entire syllabus. The examinee will be required to attempt 5 questions, selecting one question from each unit and the compulsory question. All questions will carry equal marks.

Units	Topics	Contact Hours
1	Biotech Foods and Supplements: Introduction to Food Biotechnology and related Industries.  Transgenic Plant Foods: Carbohydrates, Proteins, Vitamins Nutritional Quality Improvement of Food Crops by Genetic Engineering, Safety of GM Food Crops.  Dietary Supplements: Single Cell Protein (SCP) Production, Mushrooms Production Technology, Large Scale Production of Algae and Yeast.	
II	Food Additives & Preservation Techniques: Food Additives-Definitions, Need for Food Additives.  Classification & Functions of Different Additives: Thickeners, Antioxidants,	15



	<ul> <li>❖ Seminar/Presentation/Assignment/Quiz or Class test, etc.</li> <li>❖ Mid-Term Exam</li> </ul>	10		
	Class Participation	05	Written examination	1,0
	Theory	30	Theory	70
	Assessment: 30		End Term Examina	4:
	Internal Assessment 20	Method	s contact Hours	60
			<b>Total Contact Hours</b>	60
	Critical Control Point (HACCP).  Role of International Regulatory Agencies: Organization for Standards (ISO).  Indian Food Laws & Standards: Prevention of Fruit Products Order (FPO), Meat Products Order (CSO), Role of AGMARK Standard, Bureau of Food Safety & Standards Authority of India (FSS)	US-FI Food A	OA and International dulteration (PFA) Act, O, Cold Storage Order of Standards (BIS) and	
	Food Quality Assurance, Food Adulteration, Natu Evaluation of Food Adulterants & Toxic C	to Conc	cpts of Food Safety &	15
IV	Aseptic Packaging of Foods: Sterilization Techn Methods for Microbiological Examination of Food Advantages/functions & Disadvantages associated Food Safety and Quality Control: Introduction	us.		
	Introduction to Food Packaging: Definition, F and Selection of a Food Package, Types of F Functioning Properties	actors i	nvolved in Evolution ng Materials & their	
111	Fermented Foods & Food Packaging: Chees Fermented Foods of India: Dairy Products, Vegetables/Fruits, Meat & Fish.			15
111	Colouring Agents, Flavouring Agents, Sweet Improvers.  Probiotics: Production & Importance of Probiotics: Preservation Techniques: Refrigeration & Freet Irradiation, Antimicrobial Agents used in Food Professional Foods & Freet Production & Freedom Production &	s. ezing, [	Ochydration, Heating,	

- 1. Essential Genes. Benjamin Lewin (2007). Pearson Edu. International.
- 2. Genomes. TA Brown (2017). Garland Science, Taylor & Francis, NY.
- 3. Introduction to Food Biotechnology. Skariyachan & Abhilash (2012). CBS, Publishers, New Delhi.
- 4. Food Processing and Preservation. Sivasankar B (2002). Prentice Hall of India Pvt. Ltd., New Delhi.
- 5. Food Processing and Preservation. Khetarpaul N (2005). Dya Publishing House, New Delhi.
- 6. Food Packaging: Principles and Practice. Robertson GL (2012). 3rd Edi. Taylor & Francis Pub.
- 7. Novel Food Packaging Techniques. Ahvenainen R (2003). CRC Press.
- 8. Innovations in Food Packaging. Han JH (2005). Elsevier Academic Press.
- 9. Total Quality Assurance for the Food Industries. Gould & Gould (2001). CTI Pub. Inc. Baltimore.
- 10. Biotechnology: Food Fermentation in Microbiology, Biochemistry & Technology. Josh VK (2009).
- 11. Food Microbiology. Adams & Moss (2008). RSC Publishing Cambridge, UK.
- 12. Food Microbiology. Frazier & Westhoff (2013). TATA McGraw-Hill Pub. Pvt. Ltd. New Delhi.



1		Session: 20	25-26		
		Part A-Intro	duction		
I of Programme			M.Sc. Biotechnology		
Name of Programme			IV		
Semester	t D / Matabalamics				
Name of the Course		Genomics, 11			
Course Code			M24-BTY-405		
Course Type			DEC-4		
Level of the course			500-599		
Pre-requisite for th	e		NA		
course (if any)			oncept of genome, proteome and meta	abolome and	
(CLO). After complethis course, the learn be able to:	and and CL met	molecular phylogenetics (O-3: Conceptualize abolabolomics.	ther. ic organization of nuclear genomes of eukaryotic organelle genomes, genores. but different techniques used for properties of techniques for further research detabolomics.	teomics and	
	Gei	nomics, Proteomics and N		Total	
Credits		Theory Practical		4	
Cicuito		4	0	4	
Teaching Hours pe	r week	4 0		30 70	
Internal Assessment	Marks	30			
End Term Exam Ma	arks	70	0	100	
Max. Marks		100			
Time		3 Hours Part B-Contents of	the Course	1	
(Question No. 1) will to attempt 5 question questions will carry e	I consist at lons, selecting qual marks.	east 4 parts covering ender one question from the contract of	9 questions asking two questions nto consideration. The compulson tire syllabus. The examinee will each unit and the compulsory questions.	ry question be required uestion. All Contact Hours	
I Genetic Fea	ome! How a	are Genes of garnzea	enomes: Where are Genes in a a Nuclear Genome? How many	14	
Genetic Fea Prokaryotic	ere and What tures of Pr Genome? Ho	okaryotic Genomes:  ow many genes are the	How are Genes organized in a ere and what are their Functions?		
features of C	organelle Ger	nomes, Genetic Conte	f Organelle Genomes, Physical nt of Organelle Genomes. ion Years, Origins of Genomes,		
Acquisition Non-coding Evolution.	of new Gene DNA & Go	s by Duplication even enome Evolution: Tra	ts from other Species.  Ansposable Elements & Genome  Colecular Phylogenetics-Origin of  Cos, Key features of DNA based		



	❖ Mid-Term Exam			
	Seminar/Presentation/Assignment/Quiz. or Class test, etc.	10		
	Class Participation Seminar/Presentation/A	05	Written examination	
		30	Theory 70	
	Theory		End Term Examinatio	n: 70
	Suggested Evaluation Internal Assessment: 30	Methoa	s	00
			Total Contact II	60
	Altering/redirecting Metabolite Flow, Limitations	in Met	abolic Engineering for	
	Biosynthetic Pathways for creating new Product Altering/redirecting Metabolite Flow, Limitations	, Parti	al Pathways, Entire-	
	Metabolic Engineering, Tour			
	Example of Metabolic Regulation	Grycory	sis in Muscle as an	
	Metabolic Flux, Metabolic Control Analysis Mechanisms of Flux Control, Regulation of	, Dem	and-Supply Applysis	
	Metabolic Regulation & Control: Homeos Metabolic Flux, Metabolic Control Analysis	tasis &	Metabolia C	
	Methods for Measurement of Matal 11	110160	me and Metabolome	
	Metabolomics: Methods/Approaches Employed relationship between Genome, Transcriptome, Methods for Measurement of Metabolites Level/6	to Stu	idy Metabolism, Inter-	
	Metabolomics: Methods/Annual Metabolic		hways, Metabolite,	15
V	Introduction to Motel !			
	Affinity Purification (TAP) Tag Method, PPI Manalytical, Reverse phase, Functional.	Mapping	, Protein Microarrays	
	Protein Interaction (PPI) using the	TION O	a Protein: Protein-	
	Various Approaches for Date 101 Que	unnany	e Proteome Analysis	
	Coded Affinity Tag (ICAT) M.	1 Gel E	lectrophoresis, Isotope-	
	Protein Expression Profit: 213 um	K INIVIK	-Spectroscopy.	
	Protein Sequence Databases: Peptide Fingerpri of 3D-Structure of Protein by X-ray Diffraction 6	nting/N	Sapping, Determination	
	Proteomics.	iass-sp	ectrometry, Shot-gun	
	Sequencing of Protein by Edman Methological Reproteomics.	od (T	raditional Approach),	
	Sequencing of Protein by Edward 18	and N	1ALDI), Amino-Acid	
	Pertuacites for Study of Protoomies C		0 10 1 1	
	1 Whence to a	Juntan		1 10
11	Introduction to Proteomics Proteoms: A		C Dandanasian Communication	
,,	Phylogenetic as a tool in the study of Human Pre	hietory		
	The strips between Humans & other Primates	Origin	is of AIDS. Molecular	

- 1. Principles of Gene manipulation and Genomics. Primrose & Twyman. 7th Edi. Blackwell Publisher
- 2. Biochemistry. Voct & Voct. John Wiley and Sons, USA
- 3. Biochemistry. Satyanarayana & Chakrapani. Books and Allied (Pvt.) Ltd. India.
- 4. Lehninger Principles of Biochemistry. Nelson & Cox. Freeman & Company, NY.
- 5. Introductory Practical Biochemistry. Sawhney & Singh. Narosa Publishing Pvt. Ltd. India.

- J. Text book of Biotechnology. Dubey. S Chand & Company Ltd. India.
- 7. Principles & Techniques of Biochemistry & Molecular Biology. Wilson & Walker.
- 8. Molecular Biotechnology. Glick & Pasternak. ASM Press, Washington DC.
- 9. Genomes. TA Brown (2017). Garland Science, Taylor & Francis, NY.
- 10. Human Molecular Genetics. Tom Strachan & Andrew P Read (2011), 4th Edi. Garland Science.
- 11. Elements of Biotechnology. PK Gupta, 4th Reprint (2nd Edition): 2019-2020, Rastogi Pub.
- 12. Fundamentals of Enzymology. Devasena. Oxford University Press.
- 13. Enzymology. Devasena. Oxford University Press.

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	Session: 20	25-26	
	Part A-Intro	duction	
Name of Programme	M.Sc. Biotechnology		
Semester	IV		
Name of the Course	Biosafety, Bioethics & IPR issue of Biotechnology		
Course Code	Dissarcty, Blocti		lology
Course Type		M24-BTY-406	
Level of the course		DEC-4	
Pre-requisite for the		500-599	
Course (if any) Course Learning Outcomes		NA	
be able to:	CLO-3: Understand so biotechnology. CLO-4: Perform project	ocial, economic and legal i	ssues related to
Credits	Theory	for protection of their research Practical	/end product.
Paret: 1	4	Tractical	Total
Teaching Hours per week	4	0	4
Internal Assessment Marks End Term Exam Marks	30	0	4
Max. Marks	70	0	30
	100	0	70
- Xaiiiination I ime	3 Hours		100
Examination Time	Part B-Contents of		100

Instructions for Paper- Setter: The examiner will set 9 questions asking two questions from each unit and one compulsory question by taking CLOs into consideration. The compulsory question (Question No. 1) will consist at least 4 parts covering entire syllabus. The examinee will be required questions will carry equal marks.

Units	Topics	Contact
1	Biosafety: Introduction Historical background Di	Hours
	Biosafety: Introduction, Historical background, Biosafety in the Laboratory, Laboratory Associated Infections and other Hazards, Biosafety Management for Environmentally Safe use of Biotechnology, Biosafety Guidelines, Recommended Biosafety Levels for Infectious Agents and Infected Animals. Definition of GMOs & LMOs: Good Manufacturing Practices (GMP) and Good Lab Practices (GLP), Overview of National Regulations and Relevant International Agreements including Cartagena Protocol, Roles of Institutional Biosafety Committee (IBSC), RCGM, GEAC, MEC, SBCC, DLC and RDAC, Guidelines for Research in Transgenic Sciences and Release of GMOs to	15
II	Monney, Biological Weapons	
	associated with Consumptions of Genetically Modified Foods and other Products, Ethical Implications of Human Genome Project, Social and Ethical Implications of Biological Weapons.	15
	Bioremediations & Environmental Impacts of using GMOs: Ethics of	

	or Class test, etc.  ❖ Mid-Term Exam	15			
	❖ Seminar/Presentation/Assignment/Quiz	10			
	♣ Class Participation	05	Written examination		
-	Theory	30	Theory	/0	
	Internal Assessment: 30		End Term Examinat	70	
	Suggested Evaluation	Method	ls	: 70	
_	Related Areas-GATT, TRITS, Blodiversity Contact Hours		60		
	Breeder's Rights, International & National Conventions on Biotechnology				
	Patent and Rights: Patent Search, Patent Cooperation Treaty (PCT), Traditional Knowledge Commercial Exploitation, Farmers Rights, Plant				
	India				
	Protection, Patenting & Procedure involved in Application of Patents & Granting of a Patent, Compulsory Licenses, Legislations covering IPR's in				
V	Intellectual Property Rights: Intellectual Property Rights (IPR) & its			15	
	Transplantation-Ethical & Legal Implications, Research Focus to Address the Need of Poor Environment.				
	Human Cloning & Gene Therapy: Ethic				
	Life for Informed Decision Making, Testing of				
H	Social, Economic & Legal Issues Related to Biotechnology: Public Education of Processes of Biotechnology involved in Generating new Forms of				
	Patenting & its Impact on Biodiversity Rich Animals for Research & Testing and Alternative	s for A	nimals in Research.		

- 1. Biotechnology and Safety Assessment. Thomas & Fuch (2002). Academic Press.
- 2. Biological Safety Principles and Practices. Fleming & Hunt (2000). ASM Press.
- 3. Bioethics & Biosafety. Sateesh MK (2008). IK Publishers.
- 4. Intellectual Property Rights on Biotechnology. Singh K (2008). BCIL, New Delhi.
- 5. Dynamics of Entrepreneurial Development and Management. Desai V (2007). Himalaya Publishing.
- 6. Patent law and Entrepreneurship. Singh & Kaur (2006). Kalyani Publishers.
- 7. IPR, Biosafety and Bioethics. Goel and Prashar (2013). Pearson Education, India.
- 8. Important Web Links: <a href="http://www.w3.org/IPR/">http://www.wipo.int/portal/index.html.en</a>;

http://www.ipr.co.uk/IP\_conventions/patent\_cooperation\_treaty.html; www.patentoffice.nic.in;

www.iprlawindia.org; http://www.cbd.int/biosafety/background.shtml

my.

	Session: 202	25-26		
	Part A-Introd	M.Sc. Biotechnol	ngv	
	T	M.Sc. Biotechno	95	
Name of Programme		IV	hnology (La	ab)
Semester	Food & E	nvironmental Biotec	IIIOIOBJ (=	
Name of the Course		M24-BTY-407		
Course Code		PC-7		
Course Type		500-599		
Level of the course  Pre-requisite for the course		NA		
(if any) Course Learning Outcomes (CLO). After completing this		edge and hands-on transhrooms. cal knowledge of meth		
course, the learner will be able to:	of different water san CLO-3: Have practic qualitative aspects of CLO-4: Choose mo	aples. cal understanding of te diverse water & food s st appropriate technic e value of team spirit ons.	sehniques to samples. que for food while work	test various and water ing together
Credits	Theory	Practical	T	'otal
	0	4		4
Teaching Hours per week	0	8		8
Internal Assessment Marks	0	30		30
End Term Exam Marks Max. Marks	0	70 100	-	70 100
Examination Time	U	4 Hours		100
Examination 1 line	Part R-Conter	nts of the Course		
	Practical's			Contact Hours
<ul> <li>2. To study the product</li> <li>3. To study the cultive</li> <li>4. To study the various</li> <li>5. Estimation of (a) Io</li> <li>6. Determination of m</li> </ul>	ation of mushrooms.  Is sterilization and food podine (b) Saponification and state (b) state (b) saponification and state (b) saponification (c) state (c) saponification (c) saponifica	oreservation techniques (c) Acid values of fats (a given food sample.		120
8. Determination of to 9. To determine TDS, 10. Total bacterial pecount (SPC) technique 11. To check the potal 12. To check present	cidity & pH in food samulatal, non-reducing and reducing and reducing and reducing and reducing properties.  DO, COD, BOD of given population of given samulation of given samulation of given water samulation of coliform in given most probable number of the properties.	educing sugars.  en water sample.  nples of water by stan  nple.  n water sample by M	ultiple-tube	

14. wate	To check the presence of coliforms using ments of check the presence of faecal and non- er sample and confirmation of faecal coliform. To determine the quality of given milk sample Microbial production of Sauerkraut.	faecal	NO. 11 CONTRACTOR OF THE STATE		
10.	viictobiai production of buderkidat.	-	Total Contact Hours	120	
	Suggested Evaluation Methods				
Inte	Internal Assessment: 30		End Term Examinati	ion: /U	
	ticum	30	Practicum	70	
	Class Participation	05	Lab record, Viva-Voc	e, write-u	
	Seminar/Demonstration/Viva-Voce/Lab records, etc.	10	and Execution of the F	lactical	
-	Mid-Term Exam	15			

## Part C-Learning Resources

## Recommended Books:

- 1. Food and Agricultural Organization: Manuals of Food Quality Control.
- 2. Introductory Practical Biochemistry. Sawhney & Singh (2005). Alpha Science Publishing Pvt.

3. Practical Manual of Biotechnology: Applied Sciences. Mahajan R, et al. (2010). Vayu Education of

India.



	Session: 202	25-26				
Vana CD	Part A-Introd	luction				
Name of Programme		M.Sc. Biotechnolog	.v			
Semester	IV					
Name of the Course	Biotechnology	Animal & Medical ; Biosafety, Bioethics &	& IPR/ Genomics	·,		
Course Code	Proteomics & Metabolomics (Lab) M24-BTY-408					
Course Type		PC-8				
Level of the course	500-599					
Pre-requisite for the course (if any) Course Learning Outcomes	NA NA					
course, the learner will be able to:	different cell culture cryopreservation tech CLO-2: Work with as single nucleotide and DNA Fingerprint CLO-3: Get acquair and metabolic en knowledge of IPR, B CLO-4: Get acquair Have knowledge of ILab. They will important the cryopreservation of ILab.	ted with different tools. I Biotechnology. Get techniques, cell viabilismiques, Transfection motechniques for genetic viabilismiques for genetic viabilismiques for use in forensic solution for use in forensic solution. Get acquational solutions and Bioethics. Interest with techniques/probability and safety means bibe the value of teams ical sessions.	ty/Proliferative assethods variation detection in pharmacogenor cience.  genomics, proteouinted with practices of patent for the process of patent for the practices are the practices.	says such mics omics ctica		
Credits	together during pract Theory	iononio.		KIIIE		
	0	Practical	Total			
Teaching Hours per week	0	4	4			
Internal Assessment Marks	0	8	8			
End Term Exam Marks	0	30 70	30			
Max. Marks	0	100	70			
Examination Time		4 Hours	100			
	Part B-Conte	nts of the Course				
	Practical's		Com			
			Con Hou			
1 7- 1	Practical Exer	cises		13		
The about Coll C	unture techniques	mal Biotechnology lab				
proliferation assays 3. Cryopreservation techniques		, cen viability/cytotox	cicity and			
4. Detection of Single	nucleotide polymorphi	sm				
5. To perform DNA fir	ngerprinting analysis					
6. Synthesis of nanopal	rticles and nanocompo	sites				
7. Engineering E. coli to 8. Protein Identification	for Enhanced Production I	on of a Metabolite Using Mass Spectrometr				

10. Whou

- 9. Genome Browsing and Annotation Using Bioinformatics Tools
- 10. Whole Genome Sequencing Data Analysis
- 11. Performance of GLP in Biotechnology laboratory.
- 12. Survey of different methods of Public Education on Biotech Processes involved in generating new forms of life for informed decision making.
- 13. Study of Indian Legislation on Protection of Product and Services in Agri-Biotech Sector.
- 14. Process of implementation of rDNA guidelines in India.
- 15. Perform patent search for specific category
- 16. Protocol for filling patent and other IPR
- 17. To study NCBI Homepage and virtual library via NCBI
- 18. To perform BLAST for Nucleotide Sequence and for protein sequence
- 19. To study phylogenetic analysis
- 20. To study PDB structure
- 21. Comparative study of: Gene Bank/ Genepept and FASTA

Suggested Evaluation	Memo	End Term Examination: 7
Internal Assessment: 30	30	Practicum 70
Practicum	05	Lab record, Viva-Voce, Wri
* Class Participation	10	and Execution of the Practic
Seminar/Demonstration/Viva-Voce/Lab records, etc.	10	
❖ Mid-Term Exam	15	

## Part C-Learning Resources

- 1. Manual of Animal Biotechnology. Ramadass & Kumanan (1st Edi.). New Age, International Pvt. Ltd.
- 2. Culture of Animal Cells: A Manual of Basic Technique. Freshney (2016). 7th Edi. Wiley-Blackwell.
- 3. Medical Biotechnology: Techniques & Applications. Raymond & Wegner (2<sup>nd</sup> Edi.). Academic Press.
- 4. The Proteomics Protocols Handbook. John Walker. Humana Press.
- 5. Plant Biotechnology: Principles & Applications. Zainul et al., (2017). Springer.
- 6. IPR, Biosafety and Bioethics. Goel and Prashar (2013). Pearson Education, India.
- 7. Animal Cell Culture: Practical Approach. John (2000) 3rd Edi. Oxford Press.
- 8. Animal Cell Culture Methods In: Methods in Cell Biology. Jenni Mather & David Barnes. Academic
- 9. Important Web Links: http://www.w3.org/IPR/
- 10. Important websites: http://www.wipo.int/portal/index.html; www.patentoffice.nic.in; http://www.ipr.co.uk/IP\_conventions/patent\_cooperation\_treaty.html; www.iprlawindia.org http://www.cbd.int/biosafety/background.shtml.



	Session: 202	5-26				
	Part A-Introd	uction				
Name of Programme		Biotechnology				
Semester	IV					
Name of the Course	Entrepreneurship & Diagnostic Techniques					
Course Code		M24-BTY-409				
Course Type		EEC				
Level of the course		500-599				
Pre-requisite for the course (if any)  Course Learning Outcomes		NA NA				
(CLO). After completing this course, the learner will be able to:	support for small busic CLO-2: Perform produce CLO-3: Understand identification methods CLO-4: Know the cobiomarkers and their	I about diagnostic s and biosensors for bio ollection and handling diagnosis, test metho	and assess sources ors. ensing and release of test strips, microbia of biosamples, cancer			
Credits	Theory	Practical	75			
Teaching House	2	0	Total			
Teaching Hours per week Internal Assessment Marks	2	0	2			
	15	0	2			
End Term Exam Marks						
End Term Exam Marks	35		15			
End Term Exam Marks Max. Marks Examination Time	35 50 3 Hours	0	35 50			

Instructions for Paper- Setter: The examiner will set 9 questions asking two questions from each unit and one compulsory question by taking CLOs into consideration. The compulsory question (Question No. 1) will consist at least 4 parts covering entire syllabus. The examinee will be required to attempt 5 questions, selecting one question from each unit and the compulsory question. All

I	Entrepreneurship: Entrepreneurship & principle	Contact Hours
	development, Qualities of an entrepreneur, Functions & types of entrepreneurs. Industrial licensing, venture capital. Biotechnological industries in India and measures to promote bio-entrepreneurship in India.	07
11	Project Management: Formulation, Identification and selection based on size, Technological assessment, Project cost and market potential. Process of drug development and licensing; Guidelines for release of GMOs and their derived products in India.	08

III	Immuno-chromatographic diagnostic test strip methods (biochemical and molecular) for confirmation. Sterile disk method to test antib detecting biomolecules and use in physiologic limitations of biosensors.	microl iotic ser	pial identification and nsitivity. Biosensors for	07
IV	Specimen Collection and handling of different types of clinical specimens.  An overview of cancer biomarkers: types, tests and importance in management of cancer disease. Hematological tests (e.g., complete blood count, erythrocyte sedimentation rate, blood group and Rh factor, coagulation tests and interpretation).  Biochemical analysis of various fluids and other samples (e.g., liver function tests, kidney function tests, thyroid function tests, lipid profile tests, sugar test). Applications of techniques such as ELISA, PCR, HPLC and Mass			08
	Spectroscopy in diagnostics.  Total Contact Hours			30
	Suggested Evaluation	Method	ls .	
	Internal Assessment: 15		End Term Examinat	
	And the control of th	15	Theory	35
		_		
	Theory  Class Participation	104	Written examination	
	Class Participation     Seminar/Presentation/Assignment/Quiz     or Class test, etc.	04 04 07	Written examination	

- 1. Biotechnology: Expanding Horizon. BD Singh (2010). 3rd Edi. Kalyani Publishers.
- Experiments in Microbiology: Plant Pathology & Biotechnology. Aneja KR (2007). New Age, International Pvt. Ltd.
- 3. Introductory Practical Biochemistry. Sawhney & Singh (2005). Alpha Sci. Pub. Pvt. Ltd.
- Practical Manual of Biotechnology: Applied Sciences. Mahajan R, et al. (2010). Vayu
  Education of India.
- 5. Handbook of Bioentrepreneurship. Thomas Brenner & Holger Patzelt (2008). Springer, NY.
- Basic Biotechniques for Bioprocess & Bioentrepreneurship. Bhatt et al., (2022). Elsevier Science Publishing Co. Inc. Λcademic Press Inc.
- 7. Dynamics of Entrepreneurial Development & Management. Desai (2007). Himalaya Publishing.
- 8. Indian Entrepreneurship in Biotechnology Comes of Age. Gupta et al., (2012).
- 9. Introduction to Bioentrepreneurship. Sinha et al., (2021).
- 10. Textbook of Medical Laboratory Technology. Praful, Godkar and Darshan.
- 11. Practical Clinical Biochemistry: Methods and Interpretations. Ashfaq Ahmed.
- 12. Clinical Pathology and Clinical Biochemistry. Abhijit Chaudhari.
- 13. Clinical Biochemistry: Theory and Practical. Ramesh, Nandini and Anuradha.

	Session: 20	25-26			
	Part A-Intro	duction			
Name of Programme	M.Sc. Biotechnology				
Semester	IV				
Name of the Course	Dissertation/Project work				
Course Code		M24-BTY-410			
Course Type	Dissertation/Project work				
Level of the course	500-599				
Pre-requisite for the course (if any)		NA			
Course Learning Outcomes (CLO). After completing this course, the learner will be able to:	clear researchable the CLO-2: Select and data collection, an knowledge to perfor CLO-3: Communic both written and ora future research direct CLO-4: Develop a literature that make	ating relevant literatureme and research plantapplying appropriate danalysis. Apply pure research and to solve at research findings all formats. Drawing contions.  Tresearch project by a meaningful control of the project of the pro	research methodologies		
Credits	Theory	Practical	Total		
Teaching Hours per week	*		12		
Evaluation of Dissertation					
Viva-Voce			200		
Max. Marks			100		
Examination Time			300		
	Part B-Conte	ents of the Course	ppic in the field of life		

rite a well-structured dissertation that would reflect critical thinking, methodology, results and analysis and scholarly engagement with primary

Suggested Evaluation Methods

The dissertation will be evaluated by an external examiner out of 300 marks. Evaluation of Dissertation: 200

Total: 200 + 100 = 300

Viva-Voce: 100

Part C-Learning Resources

- Introductory Practical Biochemistry. Sawhney & Singh (2005). Alpha Sci. Pub. Pvt. Ltd.
- Textbook of Medical Laboratory Technology. Praful, Godkar and Darshan.
- Practical Clinical Biochemistry: Methods and Interpretations. Ashfaq Ahmed.
- Clinical Pathology and Clinical Biochemistry. Abhijit Chaudhari. 4.
- Clinical Biochemistry: Theory and Practical. Ramesh, Nandini and Anuradha.



- 6. Principles & Techniques of Biochemistry & Molecular Biology. Wilson & Walker.
- 7. Plant Tissue Culture Manual. Lindsey (2007). Springer (India) Publication.
- Molecular Cloning: A Laboratory Manual. Sambrook & Maniatis (2002). Cold Spring Harbor Laboratory Press, New York.
- Molecular Cloning: A Laboratory Manual. Sambrook & Russell (2007) 3<sup>rd</sup> Edi. (Vol.1-3).
   Cold Spring Harbor Laboratory Press, New York.
- Manual of Industrial Microbiology & Biotechnology. Arnold I Demain & Julian E Davies (1999).
   2nd Edition, ASM Press, Washington DC.
- 11. Molecular Biology Techniques: Susan (2nd Edi.) Academic Press
- 12. Molecular Biology: Principles and Practice. Michael et al., (3rd Edi.). WH Freeman.
- 13. Experiments in Microbial Genetics. Tortora (4th Edi.). John Wiley & Sons, Inc.
- Medical Biotechnology: Techniques & Applications. Raymond & Wegner (2<sup>nd</sup> Edi.).
   Academic Press.
- Cytogenetic Laboratory Management: Chromosomal, FISH and Microarray-Based Best Practices and Procedures. Susan Mahler Zneimer.
- 16. How to Write a Thesis. Murray & Rowena (2017). 4th Edi. Open University Press.
- 17. Scientific Thesis Writing and Paper Presentation. Gurumani. MJP Publishers, Chennai, India

