MOTHER TERESA WOMEN'S UNIVERSITY KODAIKANAL – 624 102

M.SC BIOTECHNOLOGY

Syllabus (With Effect from 2021)



DEPARTMENT OF BIOTECHNOLOGY

Mother Teresa Women's University, Kodaikanal Department of Biotechnology Choice Based Credit System (CBCS) (2021-2022 onwards) M.Sc. Biotechnology

1. About the Programme:

M.Sc., Biotechnology is a 2 year postgraduate program that is divided into 4 semesters. This programme is to develop the students theoretically knowledgeable and experimentally competent in the field of Biotechnology. This programme is designed in a way that it provides adequate knowledge of advanced Biotechnology and related subjects such as Advanced Biochemistry, Applied Microbiology, Molecular Biology and Genetics, Bioprocess Technology, Pharmaceutical Biotechnology, Omics and Genome Editing etc. The programme will facilitate students get skills and learn techniques in biological science. This advanced programme can help students in taking a career in Research as well as getting employed in companies like pharma, healthcare, agri-based and many other life science sectors.

2. Programme Educational Objectives (PEOs)

- **PEO1:** To train the students in advanced areas of biotechnology and other related subjects and sensitizing them with all possible scopes.
- **PEO2:** To endow the students with analytical and research skills, to enhance entrepreneurial accomplishments
- **PEO3:** To prepare a knowledgeable generation of biotechnologists with proficient skills to excel in their careers.
- **PEO4:** To enrich them with good communicative and technical skills to perform efficiently as an individual and as a team member in a professional environment.
- **PEO5:** To develop biotechnologists with professional ethics in order to address socioeconomic challenges and global issues logically.

3. Eligibility:

- A candidate who has passed Graduate in Life Sciences (Biotechnology/ Botany/Zoology/Microbiology/Biochemistry/EnvironmentalScience/Food Science and Herbal Sciences) and other Relevant Subject
- Candidate should have secured at least 55% in the above subject from any recognized University.

4. General Guidelines for PG Programme

- i. **Duration:** The programme shall extend through a period of 4 consecutive semesters and the duration of a semester shall normally be 90 days or 450 hours. Examinations shall be conducted at the end of each semester for the respective subjects.
- ii. **Medium of Instruction:** English
- iii. **Evaluation:** Evaluation of the candidates shall be through Internal Assessment and External Examination.

Evaluation Pattern

Evaluation	The	eory	Practical		
Pattern	Min	Max	Min	Max	
Internal	13	25	13	25	
External	38	75	38	75	

- Internal (Theory): Test (15) + Assignment (5) + Seminar/Quiz(5) = 25
- External Theory: 75

• Question Paper Pattern for External examination for all course papers.

Max. Marks: 75 Time: 3 Hrs.

S.No.	Part	Туре	Marks
1	A	10*1 Marks=10	10
		Multiple Choice Questions(MCQs): 2 questions from each Unit	
2	В	5*4=20	20
		Two questions from each Unit with Internal Choice (either / or)	
3	C	3*15=45	45
		Open Choice: Any three questions out of 5 : one question from	
		each unit	
		Total Marks	75

^{*} Minimum credits required to pass: 90

• Project Report

A student should select a topic for the Project Work at the end of the third semester itself and submit the Project Report at the end of the fourth semester. The Project Report shall not exceed 75 typed pages in Times New Roman font with 1.5 line space.

• Project Evaluation

There is a Viva Voce Examination for Project Work. The Guide and an External Examiner shall evaluate and conduct the Viva Voce Examination. The Project Work carries 100 marks (Internal: 25 Marks; External (Viva): 75 Marks).

5. Conversion of Marks to Grade Points and Letter Grade (Performance in a Course/Paper)

Range of	Grade Points	Letter Grade	Description
Marks			
90 - 100	9.0 - 10.0	O	Outstanding
80-89	8.0 - 8.9	D+	Excellent
75-79	7.5 - 7.9	D	Distinction
70-74	7.0 - 7.4	A+	Very Good
60-69	6.0 - 6.9	A	Good
50-59	5.0 - 5.9	В	Average
00-49	0.0	U	Re-appear
ABSENT	0.0	AAA	ABSENT

6. Attendance

Students must have earned 75% of attendance in each course for appearing for the examination. Students with 71% to 74% of attendance must apply for condonation in the Prescribed Form with prescribed fee. Students with 65% to 70% of attendance must apply for condonation in the Prescribed Form with the prescribed fee along with the Medical Certificate. Students with attendance less than 65% are not eligible to appear for the examination and they shall re-do the course with the prior permission of the Head of the Department, Principal and the Registrar of the University.

7. Maternity Leave

The student who avails maternity leave may be considered to appear for the examination with the approval of Staff i/c, Head of the Department, Controller of Examination and the Registrar.

8. Any Other Information

In addition to the above mentioned regulations, any other common regulations pertaining to the PG Programmes are also applicable for this Programme.

9. PROGRAMME SPECIFIC OUTCOMES (PSOs):

On completion of M.Sc Biotechnology programme, students will be able to

PSO1: attain knowledge in the fundamentals and applications of biotechnology to solve problems.

PSO2: gain proficient and practical knowledge on advanced and modern techniques to be used in research and industries.

PSO3: apply their knowledge and the skills for the betterment and advancement of their professional career.

PSO4: apply the research skill to nurture Entrepreneurial Endeavor by various funding schemes of government

PSO5 understand the ever evolving need of biotechnologist professionals and their impact in finding solutions for global issues pertaining to environment, health, food and agriculture.

10. PROGRAMME OUTCOME (PO)

On completion of M.Sc Biotechnology programme, students will be able to

PO1: gain in-depth knowledge in the advanced concepts and principles of Biotechnology and apply in research.

PO2: apply the knowledge of bio-techniques to identify solutions to problems in a systemic way.

PO3: perform the advanced techniques in the field of biology and related fields.

PO4:acquire professional ethics, leadership qualities and team-building skills to accomplish a common goal.

PO5:apply their skills of Bioinformatics to offer new insight for design and discovery of Drug

PO6: apply the theoretical and practical knowledge in securing a successful career as researcher, product developer, employee in industries and bio-business sectors,

educator or pursue higher studies.

PO7: use the scientific skills acquired to develop into a successful women entrepreneur and set up bio-business.

PO8: use the scientific knowledge obtained to contribute to the scientific society and research of our country.



M. Sc. BIOTECHNOLOGY

Sl.	Course					CIA	ESE	Total
No.	Code	C		L	P			
1	P21BTT11	Semester I	1	5		25	75	100
1. 2.	P21BTT11	Core I - Applied Missakislassy	4	5	-	25 25	75 75	100
3.	P21BTT12 P21BTT13	Core II - Applied Microbiology	4	5	-	25	75	100
3.		Core III – Molecular Biology and Genetics	4	3	-	23		100
4.	P21BTT14	Core IV – Bioprocess Technology	4	5	-	25	75	100
5.	P21BTP11	Core V - Practical in Advanced	4	-	6	25	75	100
		Biochemistry, Microbiology and Molecular Biology						
6.	P21CSS11	Supportive Course I (Skill)-	2	4	_	25	75	100
0.	12100011	Computer skills for web designing					, 5	100
		and video editing						
		Total	22	3	80	-	-	600
		Semester	-07					
7.	P21BTT21	Core VI – Immunology and Immuno	4.	4	_	25	75	100
		Technology	豆 題					
8.	P21BTT22	Core VII Genetic Engineering	<u>4</u> 9	5	-	25	75	100
9.	P21BTT23	Core VIII – Pharmaceutical 4 5 - 25		75	100			
		Biotechnology	1					
10.	P21BTT24	Core IX- Bioethics, Biosafety and IPR	4	4	-	25	75	100
11.	P21BTP22	Core X Practical in Immunology	4 >	-	6	25	75	100
		and Immuno Technology and	· 0 S					
		Genetic Engineering	2/8					
12.		Non Major Elective	4	4	-	25	75	100
13.	P21BTS21	Supportive Course II(Skill) – Phytochemistry	5 2	2	-	25	75	100
		Total	26	3	80	_	_	700
		Semester						
14.	P21BTT31	Core XI – Plant Biotechnology	4	4	-	25	75	100
15.	P21BTT32	Core XII- Animal Biotechnology	4	4	-	25	75	100
16.	P21BTT33	Core XIII – Applied Environmental	4	4	-	25	75	100
1.7	DO 1 DEED 1	Biotechnology	4			25	7.5	100
17.	P21BTT34	Core XIV - Omics and Genome Editing	4	5	-	25	75	100
18.	P21BTT35	Core XV – Bioinstrumentation and	4	5	-	25	75	100
		Biostatistics				_	_	
19.	P21BTP33		Core XVI – Practical in Plant 4 - 6		6	25	75	100
		Biotechnology, Animal						
		Biotechnology, Applied						
20	DOINGGOO	Environmental Biotechnology	2	2		25	75	100
20.	P21WSS33	Supportive Course III-Women	2	2	_	25	75	100
		Empowerment	26	1	<u> </u>			700
		Total	26	3	80		1	700

		Semester IV						
21.	P21BTE411	Elective-I* / Any MOOC courses \$	4	4	ı	25	75	100
22.	P21BTE421	Elective-II* / Any MOOC courses \$	4	4	-	25	75	100
23.	P21BTR41	Project	8	22	-	25	75	100
		16	3	0			300	
	Total				20			2300

Non Major Elective

The candidates who have joined the PG Programme, can also undergo Non Major Elective offered by other Departments.

Non Major Elective (NME) offered by Department of Biotechnology

NME - I: Industrial Waste Management (P21BTN211)

Additional Credit Courses (Mandatory)

- 1. Value Added Program I- (P21BTV11)Two Credits (First Semester)
- 2. Internship/Industrial Training (P21BTI21)Two Credits- (Second Semester)
- 3. Online Courses-Two Credits- (P21BTO31) (Third Semester)
- 4. Value Added Program II- (P21BTV41)Two Credits (Fourth Semester)

Electives

- 1. Stem Cell Biology (P21BTE411)
- 2. Forest Conservation (P21BTE412)
- 3. NanoTechnology & Cancer Biology (P21BTE413)
- 4. Drug metabolism (P21BTE414)
- 5. Molecular Modelling and Drug designing (P21BTE421)
- 6. Wildlife Conservation (P21BTE422)
- 7. Human pathology (P21BTE423)
- 8. Biobusiness (P21BTE424)
- 9. Any MOOC^{\$}

Value Added Programme

- 1. Chromatographic Techniques (P21BTV11)
- 2. System Biology (P21BTV41)

Outside class hours

- Health, Yoga and Physical Fitness
- Library Information access and utilisation
- Employability Training

^{*} Those who have CGPA as 9, and want to do the project in industry/institution during IV semester, may opt for these two papers in III semester.

^{\$} Students can take one 4 credit course in MOOC as an elective or two 2 credit courses in MOOC as electives with the approval of Department committee.

SEMESTER - I

Course	P21BTT11			L	T	P	C
Code		ADVANCI					
COR	RE I			5	-	-	4
Cognitive	K1: Recall	K2: Understand	K4:Analyze				
Level							
Learning Objective	classi To ac and c To ur metal	fication and functions in equire knowledge on the ellular interactions and a derstand the biosynthes polism.	metabolic reactions, biochemical pa		way	'S	
Unit I	t I Biomolecules						

Atom, Molecules & chemical bonds, Introduction to metabolism – anabolism and catabolism. Carbohydrates – Occurrence, chemical properties, stereo and optical isomerism, structure and classification. Metabolism and its regulation – Glycolysis, TCA cycle, Oxidative phosphorylation, pentose phosphate pathway and gluconeogenesis, ATP synthesis, Photosynthesis, Glycogenolysis. Disorders of carbohydrate metabolism.

Unit II | Lipids

Occurrence, chemical properties and classification-biosynthesis of fatty acids triglycerides, phospholipids and cholesterol – Oxidation of fatty acids, lipid storage and membrane lipids and their organization, Lipoproteins. Disorders of lipid metabolism. Vitamins – classifications, derivatives, hormones.

Unit III Amino acids and Proteins

Amino acids: structure, classification and chemical properties, structure of peptide bond – protein: classification, amino acid composition. Protein structure – Primary structure, secondary structure – alpha helix and beta pleated structure, tertiary and quaternary structure. Protein metabolism and degradation: A.A oxidation & Urea cycle. Ramachandran plot. Model proteins myoglobin, hemoglobin and chymotrypsin. Disorders of amino acid metabolism.

Unit IV Nucleic acids

DNA & RNA – structure of purine and pyrimidine bases, nucleotides and nucleotide biosynthesis, its regulation & degradation of purine and pyrimidine nucleotides – Biosynthesis of deoxyribonucleotides. Sequencing of nucleotides. Disorders of nucleic acid metabolism.

Unit V Enzymes

Nomenclature and Classification – protein enzymes, coenzymes, prosthetic groups, cofactors, isoenzymes, ribozymes, abzymes: chemical properties of enzymes: types of specificity – absolute, group, stereochemical and geometrical; factors influencing enzyme activity – temperature, pH, concentration of enzyme, substrate and effect if ions; enzyme kinetics, types of enzyme inhibition – reversible, competitive, non-competitive, uncompetitive, irreversible inhibition; allosteric enzymes. Single substrate and multi substrate enzymes. Relevance of enzymes in metabolic regulation.

Text Books	1. Sowbhaghya Lakshmi, Textbook of Biochemistry, Paras Medical
	Publisher, 2015
	2. S.P. Singh, Textbook of Biochemistry, Publisher CBS Publishers & Distributors, 2015
References	1. Nelson, D.L and Cox, M.M. Lehninger Principles of Biochemistry, 8 th

	 Edition, Macmillan worth Publishers,2021. Voet, D,Voet,J.G and Pratt, C.W. Principles of Biochemistry.,4th Edition, Publisher Wiley,2013. Mathews, C.K. & Van Holde, K.E. & Ahern, K.G. Biochemistry. 4th
	Edition, Publisher AddisonWesley.2012. 4. U. Satyanarayana. Biochemistry, Publisher Books and Allied (P) Ltd., Calcutta,2017.
E- Reference links	 https://doi.org/10.1002/cbf.1216 https://www.pdfdrive.com/biochemistry-biochemistry-e19576202.html https://www.pdfdrive.com/textbook-of-biochemistry-e14983388.html
	 4. https://www.pdfdrive.com/biochemistry-genetics-molecular-biology-e18198970.html 5. https://www.pdfdrive.com/lehninger-principles-of-biochemistry-5th-edition-e164892141.html

Upon cor	mpletion of this course the students will be able to	
CO1	know the structure and classification of biomolecules and important metabolic pathways	K1
CO2	understand the structure and functions of lipid molecules, vitamins and hormones	K2
CO3	know the properties and disorders of amino acids and proteins	K1
CO4	understand nucleotides, their structure, biosynthesis, its regulation & degradation	K1
CO5	analyze the basic concept of Enzymes in nomenclature and classification, factors influencing enzyme activity and enzyme inhibition	K4

Mapping of COs with POs & PSOs:

						0.2							
\mathbf{CO}	PO					PSO							
	1	2	3	4	5	6	7	8	1	2	3	4	5
CO1	S	M	S	M	M	S	M	S	S	S	S	M	S
CO2	S	M	S	M	M	S	M	S	S	S	S	M	S
CO3	S	M	S	M	M	S	S	S	S	S	S	M	S
CO4	S	M	S	M	M	S	M	S	S	S	S	M	S
CO5	S	M	S	M	M	S	M	S	S	S	S	M	S

Course Code	P21BTT12	APPLII	APPLIED MICROBIOLOGY				P	C
COF	RE II				5	-	-	4
Cognitive	K1: Recall	K2: Understand	K3: Analyze	K4: Apply				
Level								
Learning Objective	molect To un epider To lea	ain knowledge on bullar taxonomy. Inderstand the key print Indication of microbial of arn the applications of try, food, environment	nciples behind the midiseases.	icrobial world and	d le	arr	ı th	ie
Unit I	Medical Mic	robiology						

Introduction to Infectious diseases - Methods of transmission. Host parasite relationship. Causative agent, Epidemiology, Pathogenesis, Prophylaxis and Treatment - Staphylococcosis, Salmonellosis Aspergillosis, Cadidiasis, Giardiasis, Mycoplasmosis, Malaria, Rickettsiosis, AIDS, Influenza, flu (H1N1) and Covid-19. Importance of nosocomial infections (hospital borne), mode of transmission of airborne pathogens.

Food Microbiology Unit II

Production of distilled beverage - alcohol, wine, brandy and beer. Single cell protein and Baker's yeast. Food industry enzymes -source and application. Contamination and spoilage of meat, fish, milk, vegetables and fruits. - Principle of food preservation methods, Food quality and control. Determination of microorganisms in food -culture, microscopy and sampling methods

Agriculture Microbiology

Microbial flora of soil – bacteria, fungi, algae and protozoa. Microbial interactions among soil microorganisms. Plant growth promoting bacteria. Introduction to Nitrogen fixing bacteria-Rhizobium. Phosphorus solubilizing bacteria -VAM, Anabaena -importance in agriculture. Disease causing microbes- Xanthomonas oryzae, Puccinia spp., Banana bunchy top virus

Unit IV **Environmental Microbiology**

Microbial degradation of xenobiotics (DDT, PCB). Sewage and wastewater treatment. Microbial insecticides: NPV, Bacillus thuringiensis, B. sphaericus. Microbial removal of heavy metals: precipitation of metal sulphides by SRB. Bioleaching-recovery of metals from ores. Solid Waste Management-composting and Biogas. Plastic degrading organisms.

Unit V **Industrial microbiology**

Production of alcohol (ethanol), acids (citric acid, lactic acid), solvents (ethanol, butanol), antibiotics (penicillin, cephalosporine), amino acids (lysine, aspartate), Statins, therapeutic products, Commercial production of fructose. Enzymes used for commercial purposes and their industrial production. Whole cell immobilization and industrial applications

maastrar pro	duction. Whole cen immoonization and industrial applications						
Text	1. Sarafaraz Ahmad, A Textbook of Applied Microbiology, Publisher						
Books	2. Anmol Publications Pvt Ltd,2011						
	3. R.C.Dubey, A Textbook Of Microbiology, Publisher SChand 2011						
	4. V.S. Randhawa, Textbook Of Microbiology, Peepee Publishers and						
	Distributors,2019						
References	1. Jeffrey C. Pommerville, Fundamentals of Microbiology. 15 th Edition,						
	Publisher Jones and Bartlette. 2018.						
	2. Madigan Michael T, Martinko John M., Bender Kelly S. 2017.Biology of						
	Microorganisms. 14 th Edition, Publisher Pearson Educatio, 2017.						
	3. Gerald J.Tortora, Microbiology,11 th Edition, Publisher Pearson Education.						
	2016.						

	4. Greenwood D, Slack R and Peutherer J. Medical Microbiology, Publisher Churchill Livingstone, Hong Kong, 2012.
	5. Ian L. Pepper, Charles P.Gerba, Terry J.Gentry, Environmental Microbiology, publisher Academic Press, 2014.
E-	1. https://www.pdfdrive.com/medical-microbiology-e18737002.html
Reference	2. https://www.pdfdrive.com/microbiology-and-immunology-textbook-of-
links	2nd-edition-e33405391.html
	3. https://www.pdfdrive.com/prescotts-microbiology-e166597880.html
	4. https://www.pdfdrive.com/food-microbiology-fundamentals-and-frontiers-
	e175273799.html

Upon c	ompletion of this course the students will be able to	
CO1	understand the epidemiology and pathogenesis of microbes	K1
CO2	understand the role of microbes in food industry and gain knowledge on food spoilage	K2
CO3	acquire knowledge on soil microbes, learn the techniques in agricultural microbiology and able to apply it.	К3
CO4	know to distinguish microorganisms beneficial to environment and their applications	K4
CO5	gain knowledge in evaluating the role of micro-organisms in specific biotechnological processes in industries	К3

Mapping of COs with POs & PSOs:

				79	100								
CO				$\leq P$	O	7 12	L as				PSO		
	1	2	3	4-1	5	6	7	8 0	1	2	3	4	5
CO1	S	S	S	Sm	S	S	S	S	/S	S	S	M	S
CO2	S	S	S	M	M	S	STO	S	S	S	M	M	S
CO3	S	S	M	S	M	S	S	S	S	S	M	M	S
CO4	S	S	M	S	MS	SVON	M	S	S	S	S	S	M
CO5	S	S	S	S	S	S	S	S	S	S	S	S	M

Course Code COR	P21BTT13 E III	MOLECULAR E	SIOLOG	GY AND GEN	ETICS	L 5	T -	P -	C 4
Semester		Semester-I		Credits:4	Hours	/we	eks	: 5	
Cognitive Level	K1: Recall	K2: Understand	K3:	Apply	K4: Analyzo	e			
Learning Objective	of lift To oreplicate the regularity of the reg	nderstand various concere. develop a comprehent cation, transcription and gain extensive knowled atory pathways. cocure knowledge on M	sive ur translat	nderstanding i ion DNA repair	n the med	char	nism	ıs	of
Unit I	Genetic Ma	terial / DNA Replication	n						

DNA as genetic material. The Geometry of DNA replication – Semi conservative replication of double– stranded DNA and Circular DNA molecules. Enzymes in DNA replication -prokaryotic and eukaryotic. DNA Polymerases, DNA ligase and DNA gyrase. Events in the replication fork – Continuous and discontinuous. Okazaki fragments. Initiation, Elongation, termination of replication. Eukaryotic DNA replication. Inhibitors of replication.

Unit II Transcription

Basic factors of RNA Synthesis - RNA ploymerases - I, II and III - Transcription Mechanisms in prokaryotes and eukaryotes - chain Initiation, elongation and termination. Significance of pribnow box, TATA box, CAAT box and enhancers in transcription initiation. Rho dependent and Rho independent termination of transcription. Classes of RNA Molecules -Messenger, ribosomal and transfer RNA. Post -transcriptional and modification - RNA splicing -role of lysozyme - Spliceosomes, Group I and Group II introns Self-splicing. Capping and tailing of 5' and 3' termini of Eukaryotic mRNA molecules.

Unit III Translation

Genetic code – Definition, deciphering of codons – Universality of the code – Wobble hypothesis and codon degeneracy - codon dictionary. Mechanism of protein synthesis - importance of Initiation (IF), elongation (EF) and releasing factors (RF) - post translational modifications – protein splicing and folding – role of molecular chaperones. Regulation of gene expression in prokaryotes – the operon model. Lactose, galactose and tryptophan operon. Feedback inhibition and Allosteric enzymes.

Unit IV Gene Transfer Mechanism

Mechanisms of Genetic Exchange- Lateral and Horizontal gene transfer. Bacterial Conjugation - Hfr and F' strains, DNA Transformation- Lytic and Lysogenic infection. Transduction - Generalized transduction, specialized transduction, LFT & HFT lysates.

Unit V Mutation and Gene arrangement

Classes of mutations, spontaneous and induced mutation, mutagens, Reversion and suppression mutations, Ames test. Genetic characterization of mutants. DNA damages – DNA repair mechanism –photoreactivation, excision repair, recombinant repair and SOS function.

Transposition-Transposons, structure, types and mechanism.

Genetic Recombination - Homologous Recombination, enzymes, and models (Double-stranded invasion model and Meselson and Radding model). Site-specific recombination (Bacteriophage lambda). Short sequence recombination

Text	1. Tom Strachan, Andrew Read, Human Molecular Genetics, Garland Science,
Books	2018

	2. Verma P.S., Agarwal V.K. Molecular Biology, Publisher S Chand, 2010
	3. James D. Watson, Molecular Biology of the Gene, Publisher Pearson
	Education, 2017
References	Gildroy Swan, Textbook of Molecular Biology. Syrawood Publishing
	House, 2017.
	2. David P. Clark, Nanette J. Pazdernik and Michelle R. McGehee.
	Molecular Biology. 3 rd Edition, Elsevier, 2018.
	3. Kensal Holde Jordanka ZlatanovaThe Evolution of Molecular
	Biology, 1 st Edition, Academic Press, 2018.
	4. Krebs JE Lewin B, Goldstein ES and Kilpatrick ST. Lewin's GENES XI
	Jones & Bartlett Publishers, 2014.
	5. B. Alberts, A. Johnson, J. Lewis, M. Raff, K. Roberts and P Walter,
	Garland Molecular Biology of the Cell, 6 th Edition, Publishing (Taylor &
	Francis Group), New York & London, 2014.
	6. Harvey Lodish, Molecular Cell Biology. 7 th Edition, W.H.Freeman and
	Company, New York, 2014.
E-	1. https://www.pdfdrive.com/cell-division-genetics-and-molecular-biology-
Reference	cell-division-genetics-and-molecular-biology-e22406140.html
links	2. https://www.pdfdrive.com/cell-biology-genetics-molecular-biology-
	evolution-and-ecology-e132225829.html
	3. https://www.pdfdrive.com/molecular-cell-biology-molecular-cell-biology-
	e7302545.html
	4. https://www.pdfdrive.com/biochemistry-genetics-molecular-biology-
	e18198970.html
	5. https://www.pdfdrive.com/molecular-cell-biology-lodish-5th-ed-
	e15674865.html
	6. https://www.pdfdrive.com/karps-cell-and-molecular-biology-
	e176035175.html

Upon c	ompletion of this course the students will be able to	
CO1	know the structure, types, replication process and function of both prokaryotic & Eukaryotic nucleic acids	K1
CO2	understand the RNA synthesis and processing, Protein synthesis and processing inside the cell	K2
CO3	illustrate the mechanisms behind control of gene expression and molecular recombination inside the cell	К3
CO4	differentiate methods of DNA repair mechanisms in the cell, know Gene mapping techniques and cellular signal transduction pathways	K4
CO5	learn the basic concept of Quorum sensing, recognize oncogenes and anti-oncogenes.	K2

Mapping of COs with POs & PSOs:

CO]	PO						PSC	•	
	1	2	3	4	5	6	7	8	1	2	3	4	5
CO1	S	M	S	S	S	S	M	S	S	S	S	S	S
CO2	S	M	M	S	S	S	M	S	S	S	S	M	S
CO3	S	M	M	M	S	S	M	S	S	S	S	M	S
CO4	S	M	S	M	S	S	M	S	S	S	S	S	S
CO5	S	M	S	M	S	S	M	S	S	S	S	M	S



Course	P21BTT14	BIOPRO	CESS TECHNOLOG	L T P C		
	E IV			5 4		
		K2:Understand	K3:Apply			
Level			·FF-J			
Learning	• To ga	in knowledge about th	e importance of microb	oes in industries		
Objective	• To a	ttain knowledge on	fermentation technique	es and different types of		
	Code CORE IV K1:Recall K2:Understand K3:Apply K6: Create Level Learning Objective • To gain knowledge about the importance of microbes in industries • To acquire knowledge on fermentation techniques and different types of fermentation and fermentors. • To acquire knowledge on upstream process and downstream processes for product recovery after fermentation. • To understand the basics of primary and secondary metabolites production. Unit I Milestones of fermentation technology Identification of industrially important microorganism, primary and secondary screening, strain development and improvement for increase yield, product assays. Microbial growth and death kinetics, mathematical expression of bacterial growth. Unit II Fermentor Designing and types of fermentor: liquid, solid state and immobilized, Media and ingredients for industrial fermentation, industrial sterilization of fermentor media and air. Types of heat exchangers, immobilization techniques, Bioreactor for cell cultures. Diauxic growth and factors affecting microbial growth. Unit III Key Factors in Bioreactor Instrumentation for monitoring bioreactor and fermentation process – PH, temperature pressure dissolved O2, air flow rate, shaft speed, foaming, viscosity and controlling. Batch, fed and continuous fermentation, large scale cultivation of plant and animal cells. Up-streaming process in product production. Unit IV Downstream processing Recovery and purification of fermentation products—filtration, flocculation, centrifugation, cell disruption, liquid—liquid—extraction, Solvent and super-critical extraction, precipitation, chromatography, ultra filtration, drying, crystallization, lyophilization. Storage and packing of products. Unit V Metabolite Production Production of primary metabolites such as organic acids like citric acid, glucamic acid, Lysine, Protease. Alcohols: Beer and Wine production. Production of Bioethanol. Secondary metabolites Protuction. 1. Vinay Sharma, Arindam Kuila, Principles and Applications of Fermentation Tec					
				downstream processes for		
		•				
TT *4 T			• • •	metabolites production.		
				1		
				icrobiai growth and death		
		lession of bacterial gro) w u1.			
		rmentor: liquid, solid	state and immobilized.	Media and ingredients for		
affecting mic	crobial growth	6 20° 1	2 8.			
Unit III			P E			
		arge scale cultivation	of plant and animal cell	s. Up-streaming process in		
	ı		43			
			Claudian Class	alation contributation call		
•	-					
-			-1-0-1			
	pny, unu m	ration, or ying, or you	inzation, Tyophinzation	i. Storage and packing of		
Unit V	Metabolite l	Production	W. J.			
Production	of primary me	etabolites such as org	anic acids like citric ac	eid, glucamic acid, Lysine.		
				•		
	Penicillin V,	Streptomycin and A	ampicillin sodium salt	. Flavouring and colour		
production.	T					
				plications of Fermentation		
Textbooks		~ .	=			
		-	-	rermentation Technology,		
				ess Publications 2015		
References		K2:Understand K3:Apply K6: Create knowledge about the importance of microbes in industries in knowledge on fermentation techniques and different types of ation and fermentors. uire knowledge on upstream process and downstream processes for recovery after fermentation. erstand the basics of primary and secondary metabolites production. fermentation technology important microorganism, primary and secondary screening, strain ent for increase yield, product assays. Microbial growth and death ssion of bacterial growth. lentor: liquid, solid state and immobilized, Media and ingredients for lustrial sterilization of fermentor media and air. Types of heat techniques, Bioreactor for cell cultures. Diauxic growth and factors a Bioreactor ing bioreactor and fermentation process – PH, temperature pressure extraction, of plant and animal cells. Up-streaming process in ge scale cultivation of plant and animal cells. Up-streaming process in sorocessing of fermentation products – filtration, flocculation, centrifugation, cell extraction, Solvent and super critical extraction, precipitation, tion, drying, crystallization, lyophilization. Storage and packing of soduction bolites such as organic acids like citric acid, glucamic acid, Lysine. I Wine production. Production of Bioethanol. Secondary metabolites - treptomycin and Ampicillin sodium salt. Flavouring and colour sharma, Arindam Kuila, Principles and Applications of Fermentation logy, Publisher Wiley-Scrivener, 2019 y, Peter F.; Hall, Steve, Principles of Fermentation Technology, erButterworth-Heinemann, 2015. Death Jermentation Technology – II, Success Publications, 2015 IL. Shuler, Fikret Kargi, Bioprocess Engineering: Basic Concepts. er Pearson Education India, 2015. Bioprocess Engineering Principles. Publisher Elsevier, 2012. ata Das and Debayan Das. Biochemical Engineering: An Introductory by L. K. Niazi, Justin L. Brown. Fundamentals of Modern Bioprocessing. er CRC Press. 2015.				
References				meering. Danc Concepts.		
			, , , , , , , , , , , , , , , , , , ,	er Elsevier, 2012.		
		•	•			
				of Modern Bioprocessing.		
				- Diames - E ' '		
	5. Pau I	Loke Show, Chien Wo	ei Ooi, Tau Chuan Lin	g, Bioprocess Engineering		

	:Downstream Processing, Published CRC Press, 2021 6. Essentials in Fermentation Technology, Berenjian, Aydin, Publisher Springer, 2019.
E- Reference links	 https://www.pdfdrive.com/bioprocess-technology-d27110100.html https://www.pdfdrive.com/advances-in-bioprocess-technology-d186651074.html https://www.pdfdrive.com/biotechnology-bioprocessing-d158764194.html

Upon o	completion of this course the students will be able to	
CO1	choose the industrially important organisms and create new designs for application	K6
CO2	understand the principles and techniques in different types and designs of fermenter	K1
CO3	know the bioreactor usage and fermentation process	K2
CO4	employ the knowledge in fermentation product purification and characterization	К3
CO5	discuss the industrial production of valuable bio-products	K2

Mapping of COs with POs & PSOs:

СО				P	O						PSO)	
	1	2	3	4	5	6	7	8	1	2	3	4	5
CO1	S	M	S	S	S	S	M	S	S	S	S	M	S
CO2	S	M	S	M	S	S	M .	S	S	S	S	M	S
CO3	S	M	S	M	S	S	S	S	S	S	S	M	S
CO4	S	S	S	S	\$ -39	S	Mo	S	S	S	S	S	S
CO5	S	S	S	M	S	ST &	S	S	S	S	S	S	S

Course Code	P21BTP11	PRACTICAL	L	T	P	C
CORE		ADVANCED BIOCHEMISTRY, MICROBIOLOGY AND MOLECULAR BIOLOGY	-	-	6	4
Cognitive Level	K1: Recall	K2: Understand K4: Analyze				
Learning	• To le	arn how to estimate biomolecules such as DNA,RNA prec	isel	у		
Objective		tain knowledge on isolation, characterization and identi	fica	tio	n (of
	micro	rform isolation of plasmid DNA				
	-	arn the basics of protein isolation and purification method				
Experiments		ation of DNA by Diphenylamine				_
in		ation of RNA by Orcinol method.				
Biochemistry		ction and Estimation of starch from potato/ tapioca				
		ation of protein by Lowry's method		. 1	4	
		me assay: Estimation of salivary amylase from saliva & pl	iosp	ma	ıas	е
		ation of amino acids by Paper chromatography				
		ation of amino acids by Thin layer chromatography				
		ation of pigments by Column chromatography				
Experiments		ion of microbes from different sources-water, food, sewage	, so	il		
in		urement of microbial Size – Micrometry				
Microbiology and		nemical Characterization of Bacteria				
Molecular		ction of Hydrolytic enzymes – Protease, Amylase and Liprobic Cultivation- Fluid Thioglycolate broth	ase			
Biology		iotic sensitivity assay- Disc and Well diffusion				
		PAGE				
	8. PCR	T 1 1 - 1				
	1	ion of Plasmid DNA				
		ion and precipitation by Ammonium sulphate and acetone		<i>r</i> .	1.	1
Text Books	1. Damo	odaran Geetha K. Practical Biochemistry. Jaypee Brothe	rs N	/iec	11C	aı
Text Dooks	2. Apur	shers,2016. Da S Sastry, Essentials Of Practical Microbiology 2	nd ·	Edi	itio	n
		shr Jaypee, 2021.	-			
References	1. Gupta P	rem Prakash. Essentials Of Practical Biochemistry. Jaype	е В	rot	hei	rs
		Publishers, 2017.				
		Sankar Sastry, Sandhya Bhat K. Essentials of	P	ract	tica	al
		ology. Jaypee Brothers Medical Publishers, 2018. es in Molecular Techniques: Rakesh S. Sengar, Amit K	um	or		
		Chaudhary, Ashu Singh, CRC Press, 1 st Edition, 2018.	Luii	ıaı,		
		apley, David White. House Molecular Biology and Biotec	hno	log	y,	
	_	r Royal Society of Chemistry.2021.				
Е-	-	www.labome.com/method/Protein-Quantitation.html				
Reference	-					
Links	_		001			
	_	1	cai-	•		
	1. https:// 2. http://n 3. https:// 4. https:// engined	www.labome.com/method/Protein-Quantitation.html nbvi-au.vlabs.ac.in/ vlab.amrita.edu/?sub=3&brch=77 www.vlab.co.in/ba-nptel-labs-biotechnology-and-biomedi	cal-	-		

Upon c	Upon completion of this course the students will be able to							
CO1	gain practical knowledge on estimation of biomolecules (DNA,RNA,	K1						
	Protein, Starch)							
CO2	experiment on various chromatography techniques such as Paper, Thin	K4						
	layer and Column chromatography							
CO3	gain hands on skill in isolation, identification and biochemical	K1						
	characterization of microbes from water, food, sewage, soil							
CO4	learn the method of isolation of Plasmid DNA and SDS PAGE	K2						
CO5	know the techniques to isolate and precipitate protein	K 1						

Mapping of COs with POs & PSOs:

CO				P	PO				PSO				
	1	2	3	4	5 811	68611	7-1600	8	1	2	3	4	5
CO1	S	M	M	M/	S	S	M	S	S	S	S	M	S
CO2	S	S	M	M	M	S	M	S an	S	S	S	M	S
CO3	S	S	S	M	M	S	M	SB	S	S	S	S	S
CO4	S	S	S	M	S	S	S	S	S	S	S	S	S
CO5	S	S	S	M	S	S	M	S	S	S	S	S	M

Strongly Correlating (S) Weakly Correlating

- 3 marks;

Moderately Correlating

(M) - 2 marks

(W)

- 1 mark;

No Correlation

(N) - 0 mark

SEMESTER -II

Course Code	P21BTT21	IMMUNOLOGY AN	D IMMUNO TECH	INOLOGY	LT	P	C					
COR	E VI				4 -	-	4					
Cognitive Level	K1: Recall	K2: Understand	K3: Apply	K4: Anal	yze							
Learning	• To un	derstand the basic conce	ots of the various im	mune systems	of hu	ıma	ın					
Objective	body.											
	media	ain knowledge on antig ted immunity.	•									
		ow the role of Immune		d Disease - Co	omple	mer	at					
	_	n and its pathways, Autoi										
		To obtain knowledge on the various immune techniques in diagnosis and Immunotherapy and various other applications.										
Unit I	Immune Sys	tem										
		Immunology, Cells and										
	1 0	s, innate and acquired in city/Immunogenicity and			•	_						
		unction, Immunoglobulir										
	l anti-idiotytop		classes and subclas	sses (isotypie,	anoty	/pc	3,					
Unit II		d Cell mediated immuni	tv E 9									
		ctivation, Class switching	- /A - O -	dv genes and	genera	atio	n					
		ymus derived lymphocy	7		_							
•	•	omplex (MHC) Complex	· · · · · · · · · · · · · · · · · · ·									
I and II n	nolecules. An	tigen processing and	presentation proces	ss. Immune	regula	atio	n					
mechanisms-	-immuno-indu	ction, immunosuppression	on. Immuno-toleran	ce, Role of	cytok	ine	s,					
lymphokines	and chemokin	es. – A	·9 S									
Unit III	•	em in Health and Disea										
-	•	its pathways, Gell and		• •			-					
	_	and treatment. Autoimn										
	-	se to infections: immuni	-	_	-							
		assification and immuniz										
		ondary immunodeficiend	ey; Autoimmune di	sorders; Hyp	ersens	1t1V	'e					
	tokine related											
Unit IV		nmunology & Immuno			DO 5		_					
typing princ	iples. Princip	on – Precipitation react les and applications of escence activated cell so	f ELISA, Radio	Immuno Ass	ay (R	RIA),					

technology in immunology, Production of humanized monoclonal antibodies, immunotherapy with genetically engineered antibodies. Recombinant Vaccines - recombinant vector vaccines,

DNA vaccines ,Multivalent subunit vaccines, minicell vaccines and conjugate vaccines.

Unit V **Transplantation and Tumor immunology**

Transplantation and its classification, Immunologic basis of graft rejection and its mechanism, Transplantation antigens, tissue typing role of MHC molecules in allograft rejection, Clinical transplantations, bone marrow, HSC transplantation and immune suppressive therapy. Tumors of the immune system, tumor antigens and immune response to tumors, tumor immunotherapy.

1. Sunil Kumar Mohanty, K Sai Leela, Textbook of Immunology, Publisher **Text Books**

	2. Jaypee Brothers Medical Publishers,2014
	3. Basir F, Textbook Of Immunology, Publisher Prentice Hall India
	Learning Private Limited,2012
	4. Latha P.Madhavee, Textbook Of Immunology, Publisher S Chand &
	Company 2012
References	1. Kuby J. Immunology, 6th Edition. W.H. Freeman and Company, New
	York. 2006.
	2. Roitt I. 2017, Essential Immunology Blackwell Scientific Publications,
	Oxford 13th Edition
	3. Geoffrey Sunshine, Immunology: A Short Course Richard Coico, Wiley-
	Blackwell 7th Edition,2015.
	4. Abul K. Abbas and Andrew H. Lichtman, Cellular and Molecular
	Immunology, 2014,
	5. David K. Male, Jonathan Brostoff, David E. Roth, and Ivan M. Roitt,
	Immunology, 8th revised edition, Elsevier, 2012.
	6. Thao Doan, Roger Melvold, Susan Viselli, Carl Waltenbaugh,
	Immunology, Lippincott Illustrated Reviews Series, 2012.
E -	1. https://www.pdfdrive.com/microbiology-and-immunology-textbook-of-
Reference	2nd-edition-e33405391.html
links	2. https://www.pdfdrive.com/cellular-molecular-immunology-7th-edition-
	e157242744.html
	3. https://www.pdfdrive.com/basic-immunology-e21670961.html
	4. https://www.pdfdrive.com/medical-microbiology-virology-immunology-
	e43491517.html

Upon c	ompletion of this course the students will be able to	
CO1	thoroughly understand the basic concepts of immunology, cells and organs	K1
CO2	of immune system and types of immunity acquire knowledge on various immunological phenomena like activation,	K1, K2
	regulation and processing of various immune components in humoral and cell mediated immunity	·
CO3	interpret the role of Immune function in Health, vaccine and Disease	К3
CO4	analyze the immunological problems, understand various immuno diagnosis testing principles	K1,K4
CO5	explain the transplantation and its classification, tumors of the immune system - tumor antigens	K2

Mapping of COs with POs & PSOs:

CO						PSO							
	1	2	3	4	5	6	7	8	1	2	3	4	5
CO1	S	M	M	S	M	M	M	M	S	M	M	M	M
CO2	S	M	M	S	M	M	M	M	S	M	S	M	M
CO3	S	M	S	S	S	S	M	S	S	S	S	S	S
CO4	S	S	S	S	M	S	S	S	S	S	S	S	S
CO5	S	S	S	S	M	S	M	S	S	S	S	S	S

Course Code	de P21BTT22 GENETIC ENGINEERING											
COR	E VII		5	-	-	4						
Cognitive	K1: Recall	K2: Understand										
Level	K3: Apply	3: Apply K4: Analyze										
Learning	 To acqu 	To acquire knowledge on tools and techniques in genetic engineering										
Objective	and screeTo und sequenceTo learn	knowledge in choice of host for cloning, methods of the ening procedure derivation and application of PCR techning methods in about genetically modified organisms and their benefit ze ethical issues on genetic engineering.	nniq									
Unit I	Tools for Ge	ene cloning: Nucleases		12 h	our	'S						

Exonucleases and Endonucleases, Restriction Enzymes and its types, RNases, Methylases: CpG Methylase, Dam Methylase, Dcm Methylase, Polymerases: DNA Pol I, Klenow Fragments, Reverse Transcriptase, Taq & Pfu Polymerases, Ligases: T4DNA Ligase, *E.coli* DNA Ligase, T4 RNA Ligase, Topoisomerases: Type I&II, End Modifying Enzymes: Terminal Transferase, T4 Polynucleotide Kinase, Alkaline Phosphatases.

Unit II Cloning vectors

12 hours

Plasmid Vectors (pBR322 and pUC), phage vectors (λ), cosmids. Phagemids. Expression vectors: (pRT plasmids), Shuttle vector. Viral vectors: Lentiviral Vectors, Retroviral Vectors, Adenoviral Vectors. Plant Vectors (Ti plasmids). Artificial vectors: BACs, YACs, HACs, PACs. Methods of Gene transfer - Physical methods Microinjection, Electrophoration, lipofection, Gene gun, Biolistics transformation. Chemical methods: Liposome mediated gene transfer, Calcium phosphate mediated gene transfer, DEAE- Dextran and polyethylene glycol mediated gene transfer. Choice of host organisms for cloning (*E.coli* and Yeast). Screening of recombinant clones - blue-white screening, Screening of transfected mammalian cells - Reporter Gene Assay, Luciferase reporter gene system.

Unit III Construction of DNA Libraries

12 hours

Genomic library and c DNA library. Screening of Libraries and preservation of libraries - Methods for screening based on detecting a DNA sequence. Screening by hybridization - Colony hybridization and Plaque hybridization. Probes used for hybridization, Screening by PCR. Screening methods based on gene expression - Immunological screening, functional complementation. Protein Engineering: Site Directed Mutagenesis.

Unit IV PCR 12 hours

Principle, Components and applications of PCR. Types of PCR: RT-PCR and real-time, and PCR. PCR based microbial typing: Bacterial identification based on 16S rRNA sequences-Amplified Ribosomal DNA Restriction analysis (ARDRA), Randomly Amplified Polymorphic DNA (RAPD). DNA sequencing Methods: Maxam and Gilbert method. Chain termination method, semiautomated method, automated method, Pyrosequencing and whole-genome shotgun sequencing method. Applications of DNA sequencing.

Unit V Applications of genetic engineering

12 hours

Gene therapy, Types of gene therapy - Somatic gene therapy and Germ line gene therapy. Gene Therapy Strategies - Gene Augmentation Therapy (GAT), Targeted Killing of Specific Cells. Gene Therapy for Severe Combined Immunodeficiency Syndrome (SCID). Gene knockout mice, disease model, Transgenic animals, production of recombinant pharmaceuticals. Genetically modified foods - Production of extra nutrients in the food, Disease resistance and herbicide

resistance. D	egradation of toxic wastes – GMO. Ethical consideration of genetic engineering.
Text	1. Desmond S. T. Nicholl ,An Introduction to Genetic Engineering,
Books	Cambridge University Press,2018.
	2. Mariana Ianello Giassetti, Tatjana Brankov, Genetic Engineering Principles
	And Methods, Scitus Academics, 2019
	3. T.A.Brown. An Introduction to Gene cloning & DNA analysis, 7th edition,
	Wiley balckwell, US. 2016.
References	1. Bernard R.Glick and Cheryl L.Patten, Principles and Applications of
	Recombinant DNA - Molecular Biotechnology,5 th Edition, ASM Press,
	United states, 2017
	2. T.A.Brown, Genomes 4, Publisher Taylor and Francis, New York,2018.
	3. Bernard R. Glick, Cheryl L. Patten. Molecular Biotechnology: Principles
	and Applications of Recombinant DNA, 5th Edition Publisher,2017.
	4. Isil Aksan Kurnaz, Techniques in Genetic Engineering, Publisher CRC
	Press, 2021.
	5. Walter E. Hill, Genetic Engineering A Primer, Publisher CRC Press, 2019.
E-	1. https://www.pdfdrive.com/molecular-biotechnology-principles-and-
Reference	applications-of-recombinant-dna-4th-edition-e162050162.html
links	2. https://www.pdfdrive.com/molecular-biotechnology-principles-and-
	applications-of-recombinant-dna-e156918014.html
	3. https://www.pdfdrive.com/recombinant-dna-technology-molecular-biology-
	and-paradigm <mark>s-e11385991.html </mark>
	4. https://www.pdfdrive.com/recombinant-dna-principles-and-methodologies-
	e185941491.html

	Upon completion of this course, the students will be able to	
CO1	acquire deep knowledge on the tools and techniques of gene cloning	K1,K2
CO2	know suitable eukaryotic and prokaryotic cloning vectors, transfer technique and screening techniques for cloning	K2
CO3	understand the procedure for construction of genomic libraries	K1
CO4	learn the principle of PCR techniques, its application and methods of DNA Sequencing	К3
CO5	analyze the important strategies of Gene therapy-types, production of recombinant pharmaceuticals and Genetically modified organisms by understanding the concepts and ethics	K2,K4

Mapping of COs with POs & PSOs:

CO					PO		PSO						
	1	2	3	4	5	6	7	8	1	2	3	4	5
CO1	S	S	S	S	S	S	M	S	S	S	S	S	S
CO2	S	S	S	S	S	S	M	S	S	S	S	S	S
CO3	S	S	S	S	S	S	M	S	S	S	S	S	S
CO4	S	S	S	S	S	S	S	S	S	S	S	S	S
CO5	S	S	S	S	S	S	M	S	S	S	S	S	S

Course	P21BTT23			TF	C							
Code	CODE VIII	PHARMACEUTICAL PLOTECHNOLOGY	_		1							
	CORE VIII	BIOTECHNOLOGY	5 .	-	4							
Cognitive	K2: Understand											
Level	K3: Apply											
	K4: Analyze											
Learning	 To know the basic c 	oncepts in Pharmaceutical Biotechnology and so	ource	es o	f							
Objective	biopharmaceuticals,	drug isolation and evaluation										
	*	lge on drug metabolism and principles of drug										
	manufacturing											
	<u> </u>	To understand the process of drug development, approval process and										
	manufacturing of bio	opharmaceuticals.										
Unit 1	Introduction											
		opharmaceuticals. Sources of biopharmaceutica		•								
		plants, animals, transgenic plants. Drug iso										
	cals. Site specific delivery	rmaceutical products. Shelf life of prote	31II	bas	sea							
Unit II	Pharmacokinetics and I											
		orption of drugs. Bioavailability - factors i	nflu		ina							
		distribution - plasma protein binding, placenta			_							
		ig action, receptor theory, adverse effects of d										
interactions.	barrier: Weenamsin or are	ag action, receptor theory, adverse effects of d	rugs	, ui	us							
Unit III	Drug Metabolism and M	Manufacturing										
		eamination, oxidation, disulfide exchange),	redi	ıcti	on.							
		retion Manufacturing principles - compresso										
		formsenteric coated tablets and capsules.			ĺ							
Unit IV	Biopharmaceuticals	2 2 2										
Vaccines, r	nodern vaccine technolog	gies, pharmaceutical aspects. Recombinant p	rotei	ins	as							
		ering, peptide chemistry and peptidomimetics										
antibodies. N	Monoclonal antibody based	l pharmaceuticals. Hematopoietic growth factor	rs. N	ucle	eic							
	•	utical enzymes. Development of adhesion mole	cules	3.								
Unit V	Drug development and a		<u> </u>									
		d compound, combinatorial approaches to drug										
_		I, II and III. Regulatory authorities - Food			_							
		lations- National security authorities, Europear	ı me	aici	ine							
	new EU drug approval system											
Text		ceutical Biotechnology, Publisher CBS,2019.	ootic	.n n	× 74							
Books	2. Pankaj verma S ltd.2019	Jayaraman, Richa Ohri , Publisher Thakur publi	Cauc	шр	٧١.							
		maceutical Biotechnology: Concepts and Applic	ratio	ne								
	Publisher Wiley In		au.	110,								
References	-	oodman and Gilman's The Pharmacological	Ba	sis	of							
		GrawHill.12th ed. 2011.			~1							
	-	kar and Rege. Pharmacology and Pharmacother	apeu	tics								
	3. Poular Prakashan.		1									
		ABC, Wu-Pong S. Applied Biopharmaceu	tical	s a	ınd							
	_	. McGraw-Hill. 6th ed. 2012.										

	5. Vivekanand Kisan Chatap , Pawan Tiwari, Ashish Dixit, Textbook of
	Pharmaceutical Biotechnology, Paging Publishers, 2019.
E -	1. https://www.pdfdrive.com/pharmaceutical-biotechnology-fundamentals-
Reference	and-applications-e164753639.html
links	2. https://www.kobo.com/us/en/ebook/pharmaceutical-biotechnology-2
	3. https://www.pdfdrive.com/pharmaceutical-biotechnology-concepts-and-
	applications-d38535075.html

Upon completi	Upon completion of this course the students will be able to					
CO1	understand the scope of pharmaceutical biotechnology.	K2				
CO2	know pharmacokinetics, metabolism, dynamics of	K2				
	drugs and the steps involved in drug discovery process					
CO3	illustrate the manufacturing principles in formulation of drugs	K3				
	and biopharmaceuticals.					
CO4	compare the production of recombinant proteins, enzymes and	K4				
	carbohydrate and nucleic acid based biopharmaceuticals.					
CO5	acquire knowledge on regulatory aspects in drug development	K2				
	and drug approval					

Mapping of COs with POs & PSOs:

CO		PO A							PSO					
	1	2	3	4	5	6	7	8	1	2	3	4	5	
CO1	S	M	M	M	M	M	M	M	S	S	M	M	M	
CO2	S	M	S	M	S	S	M	M >	S	S	M	M	M	
CO3	S	S	S	S	S	S	S	S	S	S	S	S	S	
CO4	S	M	S	SI	S	S	S	S	S	S	S	S	S	
CO5	S	M	M	S	M	M	M. O	S	S	S	S	M	S	

Course Code COR	P21BTT24 RE IX	BIOETHICS, BIOSAFETY AND IPR	4	T	P -	C 4
Cognitive Level	K1: Recall K2: Understa K3: Apply	nd				
Learning Objective	modif To pro therap To lead	quire knowledge on concepts of bioethics, emerging issues of cation and recombinant DNA technology ocure knowledge on ethical, legal and socio economic aspectly and reproductive cloning arn about Intellectual property rights.— types, patentable and table—PCT and patent drafting. quire knowledge in biohazard and bio-safety level	ets (of g		
Unit I	Introduction	to bioethics				

concepts, ethical terms, issues on genetic modification and recombinant DNA technologies, ethics in agriculture and Environment benefits, risks, trans humanism and bioweapons. GM crops, Release of GMO to the environment. Special procedures for r-DNA based product production. Risk of genetic engineering, Ecocide-Eco terrorism. Emerging issues of biotechnology's impact on society.

Unit II Animal Ethics

Animal rights, ethics of human cloning, Reproductive cloning, Ethical legal and Socio economic aspects of Gene therapy, Somatic, Embryonic and Adult stem cell research, ELSI of human genome project. Transgenic plants and animals. Challenges to public policy and regulations. CCAC Guidelines on Transgenic Animals (1997), CCAC Guidelines on Animal Welfare, Laboratory Animal Management, The Need for Ethical Review

Unit III Biohazards & Biosafety

Primary containments for biohazards, Biosafety levels, recommended biosafety levels for specific microorganism, infectious agents and Infected animals. Environmental release of GMO and risk assessment. Biosafety regulations, r-DNA guidelines- National and international, levels of containment. Role of Intuitional biosafety committee, GEAC, RCGM, Cartagena protocol. CPCSEA Guidelines. Hazardous Materials Used in Biotechnology—Handling and Disposal, Good Manufacturing Practices, Good Laboratory Practices.

Unit IV IPR

Introduction to IPR – types; copy rights, patents, trademarks, trade secret design rights, geographical indication, PVPR, patentable and non-patentable – PCT, importance of IPR, Types of Patent applications, PCT cost, procedure and requirements for international patenting- patent infringement – scope, litigation, meaning, case studies & examples. TKDL, Biopiracy. Patenting of biological material. Precautions to be taken before patenting.

Unit V Patent

Introduction to WTO, GATT, WIPO, TRIPS, Patenting in India, Indian patent act, WIPO treaty budaspest treaty, publication of patents-Gazette of India, Patenting by research students, lectures and scientist University/Organizational rules in India and aboard. Global scenario of patents and Indian position. IP as a determining factor in biotechnology.

Text Books

1. Princy Louis Palatty, Ashish Kumar U, Russell Souza..A Textbook of Bioethics for Healthcare Professionals. Jaypee Brothers Medical Publishers (P) Ltd. 2017.

	2. S.V. Damodar Reddy. Intellectual Property Rights Law and Practice,
	Publisher Asia Law House,2019.
References	1. Lewis Vaughn.Bioethics: Principles, Issues, and Cases. Oxford University
	Press,2016.
	2. Nithyananda. K. V. Intellectual Property Rights: Protection and Management
	Publisher Cengage Learning India Pvt. Ltd. 2019.
	3. Sateesh MK, Bioethics and Biosafety, IK International, 2012
	4. Ramesh Shahabadkar, S Sai Satyanarayana Reddy, Intellectual Property
	Rights, Publisher Notion Press,2019.
E-	1. https://www.pdfdrive.com/bioethics-and-biosafety-in-biotechnology-
Reference	e52867075.html
links	2. https://www.pdfdrive.com/bioethics-medicine-and-the-criminal-law-volume-
	1-the-criminal-law-and-bioethical-conflict-walking-the-tightrope-
	e176230762.html
	3. https://www.pdfdrive.com/patents-and-standards-a-modern-framework-for-
	ipr-based-standardisation-e45986739.html

Upon co	ompletion of this course, the students will be able to	
CO1	get exposed to bioethics in gene cloning and emerging issues of	K1
	biotechnology's impact on society.	
CO2	know the ethical, legal and socio economic aspects in latest	K2
	biotechnological advancements and guidelines to be followed during	
	animal experimentation	
CO3	illustrate bio-safety levels required in the laboratory and understand	K2
	biohazards	
CO4	understand and inculcate knowledge on the principles of IPR types,	K2
	patentable and non-patentables and infringement	
CO5	apply the learned patenting procedure in India and abroad	К3

Mapping of COs with POs & PSOs:

								3							
CO		PO									PSO				
	1	2	3	4	5	6	7	8	1	2	3	4	5		
CO1	S	S	M	M	M	M	M	M	S	M	S	M	M		
CO2	S	S	M	S	M	M	M	M	S	M	S	M	M		
CO3	S	S	M	M	M	M	M	M	S	M	S	M	M		
CO4	S	S	M	S	M	S	M	S	S	M	S	S	S		
CO5	S	S	M	M	M	S	S	S	S	M	S	S	S		

Course Code	P21BTP22	PRACTICAL	L	T	P	C
CORI	E - X	IMMUNO TECHNOLOGY AND GENETIC ENGINEERING	-	-	6	4
Cognitive Level	K1: Recall K2: Understa K3: Apply K6: Create					
Learning Objective	To unTo obTo leaTo ol exper	derstand and develop skill sets in immunotechnology tain skill in Blotting techniques arn the techniques in isolation and amplification of DNA otain skill to prepare competent cell and perform transferents derstand molecular assays to solve biological problems	sfor	ma	atio	n
Experiments in Immuno Technology	 Preparation count. ELISA FACS Blotting terms. Immune di 	chniques (Southern Blotting, Northern Blotting, Western Bl ffusion methods (Radial immunodiffusion, single immunod nodiffusion), Rocket electrophoresis, Immuno electrophore	otti liffu	ng))	
Experiments in Genetic Engineering	1.Isolation of 2. DNA ampl 3. Restriction 4. Ligation	DNA from Bacteria ification by PCR digestion of genomic or lambda DNA of competent cells, transformation of E.coli and screening				
Text Books	1. Asim Public 2. M.R.C Manu 3. Talwa	Kumar Roy. Immunology Theory and Practical. cations, 2019. Green and J. Sambrook (2012) Molecular cloning, A lal Vol. IIII. Fourth edition, Cold Spring Harbor Laboratory ar, A Handbook of Practical and Clinical immunology, Public 2017	Lab Pre	ora ess	yaı	
References	1. Abbas 2. Karth Balak Publi 3. Fred John 4. M.R.G Vol. I 5. Fred I Sons,	s et al. Cellular and Molecular Immunology. Elsevier. 9th edik Kaliaperumal und Senbagam Duraisamy Serrishnan. Practical Immunology A Laboratory Manual. Sher: LAP LAMBERT Academic Publishing, 2017. M. Ausubel et al. editors Current Protocols in Molecula Wiley and Sons, Inc,2017. Green and J. Sambrook. Molecular cloning, A Laborator III. 4 th Edition, Cold Spring Harbor Laboratory Press,2012. M. Ausubel. Current Protocols in Molecular Biology. John Inc,2017.	nthi 1 st 1 r B ry 1	lku Edi iol Ma	ima itio ogy nua	y. al
E-Reference links	immu	//www.immunology.org/public-information/bitesized- nology/experimental-techniques //www.nature.com/subjects/immunological-techniques				

- 3. https://currentprotocols.onlinelibrary.wiley.com/hub/journal/1934368x/aim sandscope/
- 4. https://www.apsnet.org/edcenter/disimpactmngmnt/labexercises/PlantBiote chnology/Pages/Activity4.aspx
- $5. \ http://www2.southeastern.edu/Academics/Faculty/jtemple/486/experiment\\ \% 202.pdf$
- 6. https://www.life.illinois.edu/molbio/geldigest/digest.html

Upon cor	mpletion of this course the students will be able to	
CO1	gain practical knowledge on immunological techniques	K 1
CO2	perform immunodiagnosis tests in laboratories	K2
CO3	apply the immunodiagnosis knowledge in research	К3
CO4	choose and compare different PCR techniques for amplification of multiple copies of DNA	K6
CO5	procure skill in preparation of competent cells, transformation of <i>E.coli</i> and screening of transformants	K6

Mapping of COs with POs & PSOs:

CO				9 p	0- <			S	5		PSO		
CO	1	2	3	4	5	6	7	8	1	2	3	4	5
CO1	S	S	S	S	S	S	S	S	S	S	S	M	S
CO2	S	S	S	M	M	S	S	S	S	S	S	M	M
CO3	S	S	S	M	M	S	S	S	S	S	S	M	S
CO4	S	S	S	M	M	S	S	S	S	S	S	M	S
CO5	S	S	S	M	M	S	S	S	S	S	S	M	S

Strongly Correlating (S)
Weakly Correlating (W)

- 3 marks; - 1 mark; Moderately Correlating
No Correlation

(M) - 2 marks

SA WOMEN'S

(N) - 0 mark

Course Code	P21BTS21 PHYTOCHEMISTRY		L	T	P	C
SUPPO COUR		PHYTOCHEMISTRY	2	-	-	2
Cognitive Level	K1: Recall K2: Understa K4: Apply	and				
Learning Objective	To lead compTo un	arn the effective procedures in extraction and purification of ounds aderstand the structural analysis of bioactive compounds in knowledge on herbal medicine and phyto pharmaceutica		oad	 ctiv	re
Unit 1	Extraction N					
		ents for extraction. Extraction – purification of bio-active c extract. Soxhlet extraction - crude extracts purification l				
liquid chroma	atography - HF	npounds- chromatographic techniques - thin layer chromately chroma	ato	gra	phy	y -
Unit III		nalysis of bioactive compounds				
		ectrosc <mark>opy – NMR spectroscopy.</mark>				
Unit IV	Herbal medi					
Pharmacolog <i>Eclipta alba</i> ,	ical action - <i>Gymnema sylv</i>	e - different types of herbal medicine - Ayurveda, Siddha and clinical research and traditional uses of Indian medicinal vestre, Ocimum sanctum, Curcuma longa.				
Unit V	Phytopharn					
		their health benefits - anthocyanins, carotenoids, lycopene,				
		mega 3 - fatty acids, biological effects of resveratrol.				
Text Books	 Vaibby Pharm House Deep Extract Limit 	B. Pharmacognosy and Phytochemistry, Publisher CBS,201 nav Darvhekar Rageeb, Lodhi, Vadnere, A Textbook of nacognosy & Phytochemistry, Publisher Everest Publishing e,2019 Panhekar, Ms. Trupti P. Sawant, D. P. Gogle, Phytochemic ction, Separation & Analysis, Publisher Global Education ed,2019	als			
References	and P 2. Kausa Publis 3. Sapna	tini Shukla, Dr. Shashi Alok, Dr. Prabodh Shukla, Pharmaco Phytochemistry, Publisher Nirali Prakashan, 2019. Far Jabeen, Pharmacognosy And Phytochemistry – II, Publish Shers & Distributors Pvt Ltd, 2020. Far Malviya, Swati Rawat, Pharmacognosy and Phyto Sher: Oxford and IBH Publishers, 2020.	ner:	SI	A	y,
E- Reference links	phyto 2. https: &sou 3. https:	//www.pdfdrive.com/textbook-of-pharmacognosy-and-ochemistry-d184620437.html //books.google.co.in/books?id=satDwAAQBAJ&printsec=farce=gbs_ge_summary_r&cad=0 //www.pdfdrive.com/trease-and-evans-pharmacognosy-33029.html	ron	tco	vei	r

Upon	Jpon completion of this course the students will be able to					
CO1	know the extraction and purification of bioactive compounds	K2				
CO2	understand the principles of various chromatographic techniques	K2				
CO3	acquire knowledge on the structural analysis of bioactive compounds using	K2				
	spectroscopy					
CO4	compare the medicinal properties of important medicinal plants	K4				
CO5	know the importance and health benefits of phytopharmaceuticals	K1				

Mapping of COs with POs & PSOs:

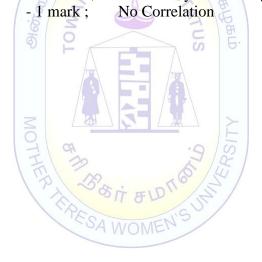
CO		PO								PSO				
	1	2	3	4	5	6	7	8	1	2	3	4	5	
CO1	S	S	S	S	M	S	M	S	S	S	S	S	M	
CO2	S	S	S	S	M	S	M	S	S	S	S	S	S	
CO3	S	S	S	S	S	Seem	M	S	S	S	S	S	S	
CO4	S	S	M	M	M	S	Mode	S	S	M	S	M	M	
CO5	S	S	M	M	M	MQU	M	S	S	M	S	M	S	

Strongly Correlating (S) Weakly Correlating (W)

- 3 marks ;

Moderately Correlating
No Correlation

- 2 marks (M) - 0 mark (N)



SEMESTER-III

Course Code	P21BTT31	PLANT BIOTECHNOLOGY	L T P C
	E XI	12.11(121012011(02001	4 4
Cognitive	K1: Recall	K2: Understand K3: Apply K6: C	reate
Level		rr J	
Learning	• To imbib	e knowledge in plant biotechnology and its application f	or increasing
Objective		al production, environment improvement, human nutrition	_
	_	with theoretical knowledge regarding the techniques and	
		siotechnology, preservation of plant cells and Genetic Eng	
	• To acqui	re knowledge on plant molecular biology, Genetic trans	formation in
	plants, m	etabolic engineering and molecular pharming	
	• To learn	about genome organization in plants, basic techniques in	tissue culture
	and its ap	plications.	
	• To be aw	are of various ethical issues and guidelines for GMOs	
Unit I		d Tissue culture	12 hours
•		Il and tissue culture-culture media; composition and prep	' 1
		allus culture, Micropropagation, suspension culture, roo	
	e pollen cultur	e, ovary <mark>culture, embryo cultur</mark> e, embryo rescue, large sc	ale culture of
plant cells		6 3 9 1	
Unit II	Regeneration		12 hours
		natic hybridization – protoplast isolation fusion and culti	
1		ation hardening and green house technology. Use of hap	loids in plant
	•	id seeds and regeneration of hybrid plants.	10 1
Unit III		ene transfer in plant cells	12 hours
		rium mediated, viral vector and their application, Caulio clear transformation methods, Promoters, reporter genes	
		ree gene targetting. Gene silencing.	and marker
Unit IV	Transgenic		12 hours
	U	y – delayed fruit ripening, transgenic plants-plantibodies	
		ant-Bt, cry genes of Bt and their gene expression, heribicion	
		ant-antifungal proteins, Virus resistance-coat protein & n	
		e stress tolerant.	deles capsia,
Unit V	Plant as bion		12 hours
Green & red		otein, starch and fructans. Nitrogen fixation and genes. A	
	-	nger printing in plant biotechnology. Biosafety guidelines	
		and risks. IPR related to plants, IPP.	
	1. S. Uı	nesha, Plant Biotechnology. Publisher CRC Press.2019.	
Text		Ranabhatt, Renu Kapor. Plant Biotechnology. 1st Edition,	Publiser WPI
Books		thing. 2018.	
		la H S. Introduction To Plant Biotechnology. Publisher O hing. 2020.	xford & IBH
References		hoff Peter M. Plant Biotechnology and Development. Publ ncis Inc. 2020.	isher: Taylor
		, M.Z., Kiran, U., Kamaluddin, M., Ali, A. Plant Biotechn	ology:
		ples and Applications, Publisher Springer. 2017.	
	3. Halfo	rd NPlant Biotechnology: Current And Future Application	ns of

	Genetically Modified Crops, Publisher Wiley,2015. 4. Smith RH. Plant Tissue Culture. 3 rd Edition, Publisher Elsevier. 2013.
E -	1. https://www.pdfdrive.com/plant-biotechnology-and-genetics-principles-
Reference	techniques-e15853574.html
links	2. https://www.pdfdrive.com/plant-cell-and-tissue-culture-a-tool-in-
	biotechnology-e20389188.html
	3. https://www.pdfdrive.com/principles-of-plant-biotechnology-
	e33514134.html
	4. https://www.pdfdrive.com/plant-genomics-e28703875.html

	Upon completion of this course the students will be able to	
CO1	learn basic techniques and setup of plant tissue culture laboratory	K1
CO2	know the application and techniques of germplasm conservation, hardening and green house technology.	K2
CO3	get updated with the research in Plant transformation techniques	К3
CO4	gain knowledge about terminator seed technology, and research advancement and its production of plantibodies edible vaccine, in transgenic plants.	K6
CO5	acquire knowledge on application of techniques in plant biotechnology and describe biosafety guidelines for research involving GMO's and IPR	K2

Mapping of COs with POs & PSOs:

CO				$\leq P$	≥ PO				PSO				
	1	2	3	49	5	6	7	8 =	1	2	3	4	5
CO1	S	S	S	SI	M	S	S	S	S	S	S	S	S
CO2	S	M	S	S	M	S	Sago	S	S	S	S	S	S
CO3	S	M	S	S	M	ST &	M	S	S	S	S	S	S
CO4	S	M	S	M	SES	S	MS	M	S	S	S	S	S
CO5	S	M	M	M	S	M	M	M	S	M	M	M	M

Course Code	P21BTT32	ANIMAL BIOTECHNOLOGY L T P C					
CORE XII		4 4					
Cognitive	K1: Recall						
Level	K2: Understa	nd					
	K3: Apply						
	K4:Analyze						
	K6: Create						
Learning Objective	molec	in knowledge on the basics of Animal cell culture, transgenic animals, cular markers and their applications.					
		 To obtain familiarity to biology and characteristics of cell culture and maintenance 					
		nderstand the principles of gene knock out, molecular pharming and yonic preservation					
		now about the stem cell technology and advanced techniques in animal chnology					
Unit I	Animal cell	culture 12 hours					

Structure and organization of animal cell. History of animal cell culture technique. Constituents of culture medium; serum and supplements; Facilities for animal cell culture-infrastructure, equipment, culture vessels. Biology and characterization of cultured cells-cell adhesion, proliferation, differentiation, morphology of cells and identification. Animal cell culture-merits and demerits.

Unit II Primary cell culture

12 hours

Primary cell culture techniques - aggregation, Cell growth & viability determination. Measurement of cell death, Transformation and Cytotoxicity assays. chromosome analysis and antigenic markers, selectable markers for animal cells. Mass culture of cells - manipulation of cell line selection - types of cell lines - maintenance of cell lines - immobilization of cells and its application - synchronization of cell - cryopreservation - germplasm conservation and establishment of gene banks. Hazards and safety aspects of cell culture techniques.

Unit III Transgenic animals and Molecular pharming

12 hours

Knock out and Knock in, Suicide gene therapy Gene silencing. Animal Biotechnology for the production of regulatory proteins, blood products, cell culture based vaccines and hormones and other therapeutic proteins. Embryonic preservation and its uses in endangered animals.

Unit IV Gene therapy and Diagnostics methods

12 hours

Gene therapy – IVF & Embryo transfer, Gene transfer techniques, Tissue engineering, Organ transplant. Synthetic viral vectors in gene transfer. Biotechnological applications for HIV. Diagnostics and therapy. DNA based diagnosis of genetic diseases, DNA barcoding. Oncogenes and anti-oncogenes. Genetic engineering approaches for genetic disorder correction. Transgenic animals as models for human disease

Unit V Stem cells

12 hours

types – Hematopoietic stem cells, Mesenchymal stem cells, embryonic stem cells, fetal stem cells, Adult stem cells- characterization, isolation, cultures. Stem cells as vector for cancer therapthy. Collection, processing, preservation and banking of Umbical cord blood stem cells. 3D culture, human cloning, ethical limits and mapping of human genome. Commercial application of animal cell culture

Text Books

- 1. A.K. Srivastava, RK. Singh. Animal Biotechnology, CBS Publishers & Distributors Pvt Ltd, India,2018.
- 2. M.M. Ranga..Animal Biotechnology,3rd Edition, Agrobios, India. 2017.

	3. B. Singh, S.K. Gautam, Textbook of Animal Biotechnology, Publisher The
	Energy and Resources Institute, TERI,2013
References	1. Birbal Singh, Gorakh Mal, Sanjeev K. Gautam, Manishi Mukesh. Advances
	in Animal Biotechnology. Springer, 2019.
	2. Singh, B., Mal, G., Gautam, S.K., Mukesh, M. Advances in Animal
	Biotechnology.Publisher Springer,2019.
	3. Rodrigues, Gabriela, Roelen, Bernard A. J, Concepts and Applications of
	Stem Cell Biology, Publisher Springer,2020.
E -	1. https://www.pdfdrive.com/animal-cell-biotechnology-e22743665.html
Reference	2. https://www.pdfdrive.com/animal-biotechnology-1-reproductive-
links	biotechnologies-e187110512.html
	3. https://www.pdfdrive.com/animal-cell-biotechnology-e177857548.html
	4. https://www.pdfdrive.com/molecular-biotechnology-principles-and-
	applications-of-recombinant-dna-4th-edition-e162050162.html

Upon o	Upon completion of this course, the students will be able to				
CO1	know the requirements to establish the cell culture laboratory	K1			
CO2	understand the methods to perform primary cell culture techniques, mass production, storage methods, germplasm conservation and establishment of gene banks.	K2			
CO3	interpret the practical difficulties in sources of contamination in cell culture and importance of transgenic animals and Molecular pharming	К3			
CO4	apply the precise gene therapy treatment for advanced medical treatment of human diseases	K4			
CO5	learn the concept of collection, processing and preservation of stem cells and to create stem cell banks that will ensure a future generation free from genetic disorders	K6			

Mapping of COs with POs & PSOs:

					1.00	4 1000	BAEN!) /					
CO	PO							PSO					
	1	2	3	4	5	6	7	8	1	2	3	4	5
CO1	S	S	S	S	M	S	S	S	S	S	S	S	S
CO2	S	S	S	S	M	S	S	S	S	S	S	S	S
CO3	S	M	S	M	S	S	M	S	S	S	S	M	S
CO4	S	S	S	S	S	M	M	S	S	M	S	M	S
CO5	S	S	S	M	M	S	S	S	S	S	S	S	S

Course	P21BTT33	L T P C								
Code	e viii	APPLIED ENVIRONMENTAL BIOTECHNOLOGY								
	E XIII K1: Recall	4 4								
Cognitive Level		nd								
Level	K2: Understand K3: Apply									
		K4: Analyze								
Learning	To understand the concept of natural resources, environmental pollution and									
Objective	remediation using biotechnology.									
Objective		9								
		• To know about the environmental hazards, solutions to protect the environment and sustainable development.								
		knowledge on remediation of contaminated environments (land, air,								
	_	and environment-friendly processes such as green manufacturing								
		ogies and sustainable development.								
		erstand the most important environmental problems such as global								
		g, ozone depletion, waste disposal and to acquire skills								
		e the environmental problems through biotechnological approach and to								
		environmental conscious.								
Unit I		n to Environment								
Environmen	t components	, Role of Biotechnology in Environmental protection, Biodiversity,								
		traditions for the conservation of Biodiversity- Sacred groves -								
Sthalavriksha	as. Insitu cons	servation- biosphere reserves, sanctuaries and national parks – Exsitu								
conservation	- Botanical (Gardens, Zoos, Gene bank, seed bank, Plant tissue culture, and								
environment	al Protection a	ct, Mass movements.								
Unit II	Pollution									
		ollution. Sources and effects of water, air, soil, thermal, noise and oil								
		nd biosensors for detection of pollution. Biomagnification. Global								
	_	Green house effect, global warming and climate change, Ozone								
		mog and acid rain.								
Unit III		tion and control								
		eters of water-water pollutants, Marine Pollution-minamata disease,								
		and Cauvery; Eutrophication; Water Quality standards; Waste water								
		ry-ASP, RBC, OP, anaerobic digestion.								
Unit IV	Solid waste r	8								
I		sources and effects, solid waste disposal-land filling, composting,								
-	-	ation, Biogas production, 5R concepts, Radioactive wastes, types of								
		ects of radiation, Disposal - Deinococcus radiodurans. Chenobyl disaster o – mark, Biodegradable plastics and Ecomark								
Unit V		ion and Biodegradation 12 hours								
		l mechanism, Bioremediation of dyes, Biodegradation of recalcitrant								
		cocarbons, degradative plasmids. GEMs, Microbial Transformation of								
	•	Bioaccumulation, Biosorption, and Bioprecipitation of Heavy Metals.								
_	_	cides and Biofertilizers.								
Text		harma.2019.Environmental Chemistry, Krishna Prakashan Media (P)								
Books	Limite									
	2. Pramo									
		chnology, Publisher WPI Publishing,2019								
References		oor Adejumoke A., Adebesin Babatunde O., Oluyori Abimbola P.,								
	2. 111,1110	2								

	Adelani-Akande Tabitha A., Dada Adewumi O. and Oreofe Toyin A.							
	,							
	2018. Water Pollution: Effects, Prevention and Climatic impact.							
	2. Daniel Vallero, Environmental Biotechnology: A Biosystems							
	Approach, Publisher Elvesier, 2015.							
	3. Pramod Kumar, Vipin Kumar, Environmental Biotechnology, Publisher							
	Woodhead Publishing India,2018							
E-	1. https://www.open.edu/openlearncreate/mod/oucontent/view.php?id=80588&							
Reference	printable=1							
links	2. http://mjcetenvsci.blogspot.com/							
	3. https://nptel.ac.in/courses/120/108/120108005/							

Upon c	Upon completion of this course, the students will be able to				
CO1	be aware on need for Environmental protection and conserve the floral and faunal diversity	K1			
CO2	understand and identify the environmental issues, gain knowledge on the types of pollution and ways to preserve the environment	K2			
CO3	apply the knowledge of water chemistry to prevent water pollution	К3			
CO4	inculcate knowledge about solid wastes management and its importance	K2			
CO5	compare the eco-friendly techniques and apply the methods for environmental protection	K4			

Mapping of COs with POs & PSOs:

CO	PO PO						PSO						
	1	2	3	41	500	6	7	8 2	/1	2	3	4	5
CO1	S	M	M	S	M	S	M 6	S	S	M	S	M	S
CO2	S	M	M	S	M	ST &	M	S	S	M	S	M	S
CO3	S	S	S	S	M	S	SNS	S	S	S	S	S	S
CO4	S	S	S	S	M	SVON	S	S	S	S	S	S	S
CO5	S	S	S	S	M	S	S	S	S	S	S	S	S

Course	P21BTT34		L T P C											
Code		OMICS AND GENOME EDITING												
CORI	E XIV		5 - 4											
Cognitive	K1: Recall													
Level	K2: Understa	nd												
	K3: Apply													
Learning		rn about prokaryotic and eukaryotic genomes, general												
Objective	_	e sequencing techniques, genome analysis and annot	ations and											
	* *	applications of genomics.To acquire knowledge in the contemporary computational tools for gene												
	_	analysis												
	-	 analysis To know about molecular systems biology, molecular systems network and its applications To learn about proteomics, metabolomics, interactomics and its applications in the field of medicine, agriculture and industries 												
	* *													
Unit I		Genome sequencing	12 hours											
		ganization - Eukaryotic genome - Organelle genome- G												
	1 Microbiome	s – Genome sequencing technologies –Comparative genor	nics and its											
applications.		8 57												
Unit II	Functional g		12 hours											
		n analysis –Experimental methods - Computational tools for												
		ering – Gene expression analysis– STS-EST-GSS-Assessing												
	_	Transcriptome analysis and a	pplications.											
Metagenomic		B - A												
Unit III	- v	stems biology	12 hours											
		ries - constraint and kinetic modeling - Biomass objective												
		echnological applications – Molecular network biology – N												
_	omics - Pharn	nacoge <mark>nomics and drug discovery – Agriculture genom</mark>	ics and its											
applications	D .	1 2 Signatural Signatura Si	101											
Unit IV	Proteomics 1.6		12 hours											
		eatures – Qualitative proteome technology (Gel-based and												
-	-	nology – Functional proteome technology – Methods, algorithms – Proteome detabases – Protein engineering												
Metabolomic		roteomics - Proteome databases - Protein engineering	resources.											
			12 1-											
Unit V	Interactomic		12 hours											
		n-protein interactions - Modelling of proteomic networks — otechnologies in proteomics — Modificomics — Proteomics												
		- Application of proteomics in agricultural biotechnology	* *											
	nd its applicati		– mausirar											
proteomics at		acharya, Anjanabha, Parkhi, Vilas, Char, Bharat. CRISPR/C	as Genome											
		g. Strategies And Potential For Crop Improvement. Published												
Text Books	2020.	5. Stategles That I defined I of Crop improvement. I donsing	or opinigor.											
2 CAU DOOMS		Banerjee, Garlapati Vijay Kumar, S.P. Jeevan Kumar. OM	IICS-Based											
		paches in Plant Biotechnology. Publisher Wiley. 2019.												
		Davies, Editing Humanity: The CRISPR Revolution and t	he New Era											
		nome Editing, Publisher Pegasus Books,2020	- · · · —- •											
References		g Qi. Plant Genome Editing with CRISPR Systems. Methods	and											
	_	cols. Publisher Springer. 2019.												

	 George M. Church, Krishnarao AppasaniGenome Editing and Engineering: From TALENs, ZFNs and CRISPRs to Molecular Surgery. Publisher Cambridge University Press.2018 Debmalya Barh, Vasudeo Zambare, Vasco Azevedo. OMICS,Applications in Biomedical, Agricultural, and Environmental Sciences.Publisher CRC Press. 2013. Yu LiuOmics in Clinical Practice: Genomics, Pharmacogenomics, Proteomics, and Transcriptomics in Clinical Research. Publisher Apple Academic Press. 2014 Turksen, Kursad. Genome Editing. Publisher Springer. 2016.
E-	www.genomic.org.uk/
Reference	2. https://www.britannica.com/science/genomics
Links	3. https://www.sciencedirect.com/journal/genomics

On con	npletion of this course, student would be able to	
CO1	understand the structure of prokaryotic and eukaryotic genomes	K1
CO2	employ the computational tools for gene analysis	K3
CO3	know the key concepts in molecular systems biology and list its applications	K2
CO4	understand the functional proteome technology and metabolomics.	K2
CO5	gain knowledge about the modelling of proteomic networks and identify its applications in clinical biomedicine.	K1,K2

Mapping of COs with POs & PSOs:

00				1/2	-40	DGO.								
CO	1	2	3	1						PSO PSO				
CO1	S	M	M	M	S	M/ON	M	S	S	S	M	M	M	
CO2	S	S	S	S	S	S	S	S	S	S	S	S	S	
CO3	S	S	S	S	S	S	M	S	S	S	S	S	S	
CO4	S	S	S	S	S	S	S	S	S	S	S	S	S	
CO5	S	S	S	S	S	S	M	S	S	S	S	S	S	

Course	P21BTT35		L T P C
Code		BIOINSTRUMENTATION AND BIOSTATISTICS	
	E XV		5 4
Cognitive	K1: Recall	J	
Level	K2: Understa K3: Apply	ind	
	K6: Create		
Learning		inderstand the principles and working methods of m	icroscopes
Objective		fuges, spectrometers, electrophoresis.	icroscopes,
3		gain knowledge in experimental designing and data	collection
	_	iques.	
	 To ac 	quire knowledge on applications of statistics in research.	
Unit I	Microscopy		
Principle ar	nd application	s of light, phase contrast, fluorescence, inverted, sca	nning and
		roscopy, scanning tunneling microscopy, atomic force n	
confocal la	-	microscopy, field emission scanning electron n	-
		cytometry. Micrometry, lyophilizer, Preparation of microb	ial, animal
	_	oscopy. Principles of colorimetry and spectroscopy.	1
Unit II	Centrifugati	ations—gradient and density centrifuge, Ultracentrifugation	on velocity
		ciple methodology and applications of gel – filtration, ion	
		why; Thin layer, liquid and gas chromatography; High po	
-		ra sonicator, pH meter, FT-IR.	
Unit III		biophysical method	
X ray diffr		scence, UV, visible, IR. Atomic absorption and plasma	a emission
spectroscopy	, NMR, MS,	ELISA reader, Electrophoresis: Principle and applications	of Native,
		phoresis, isoelectric focusing, isotachophoresis, MADI-	TOF-TOF,
Microarray 7	Techniques.	工力	
Unit IV		ion and Presentation of data	
		n of experimental data. Brief description and tabulation of	
		Measures of central tendency: arithmetic mean, medi	
_	lean, Harmoni d standard erro	c mean. Measures of dispersion: range, interquartile range.	e, standard
Unit V	1		<u> </u>
	Hypothesis 1	of two types of errors and level of significance. Tests of si	gnificance:
* *	-	Non parametric: Chi square tests. Simple linear regre	-
	Analysis of var		ession una
	, , , , , , , , , , , , , , , , , , , ,	y Diwedi, Usman, Srivastava. Biostatistics and	Research
		odology.Publisher S VikaS and Company. 2019.	
Text		erakumari. Bioinstrumentation. 1 st Edition Mjp Publishers.	2011.
Books	3. B An	nadurai, A Textbook of Biostatistics, Publisher New Age In	ternational
	Priva	te Limited,2017	
D.C.	1 7 1	CWI DILL WILLOW	
References		G Webster . Bioinstrumentation .Publisher Wiley.2021.	r Thoulan
		h Ved. Biostatistics & Research Methodology. Publishe cation.2019.	a maukuf
		eily. 2019. Bioinstrumentation. CBS Publishers & Distribut	ors
		nadurai. A Textbook of Biostatistics. Publisher New Age In	
L			

	Private Limited.2017. 5. Norman T.S. Bailey. Statistical Methods in Biology. Cambridge University Press, UK. 2012.
E-	1. https://www.pdfdrive.com/biostatistical-methods-biostatistical-
Reference	methods-e15213717.html
links	2. https://www.pdfdrive.com/biostatistics-e42988735.html
	3. https://www.pdfdrive.com/introductory-biostatistics-e15112721.html
	4. https://www.pdfdrive.com/introductory-biostatistics-e176105301.html
	5. https://www.pdfdrive.com/bioinstrumentation-instructional-resources-
	technology-austin-e15581883.html

Upon co	empletion of this course the students will be able to							
CO1	know the principle and applications of different types microscopes, cytometer and calorimeter							
CO2	understand the importance centrifugation and chromatographic techniques	K2						
CO3	know to apply the knowledge in identification of biopolymer structures, understand the principles of spectroscopic and electrophoretic techniques	К3						
CO4	develop skill in collection and presentation of biological data through biostatistics	К3						
CO5	bring out solutions to solve the biological research problems trough statistics	К6						

Mapping of COs with POs & PSOs:

CO				P	0 10		PSO						
	1	2	3	4	5	67 8	7	8	1	2	3	4	5
CO1	S	M	S	M	M	S	Sis	S	S	S	S	M	S
CO2	S	M	S	M	M	SVOI	S	S	S	S	S	M	S
CO3	S	M	S	M	M	S	S	S	S	S	S	M	S
CO4	S	M	M	M	M	S	M	S	S	S	S	M	S
CO5	S	M	M	M	M	S	M	S	S	S	S	M	S

Course Code	P21BTP33	PRACTICAL	L	T	P	C
CORE	XVI	PLANT BIOTECHNOLOGY & ANIMAL BIOTECHNOLOGY, APPLIED ENVIRONMENTAL BIOTECHNOLOGY	-	-	6	4
Cognitive	K1: Recall					
Level	K2: Understa K4: Analyze	and				
	K6: Create					
Learning		arn to set up a plant cell culture and animal cell culture la	ıboı	ato	orv	
Objective		equire knowledge on culturing callus and root tip, and also				1
		insformation techniques	Ü			
	_	in skills on handling and maintenance of animal cell cult				
	• To ac	equire knowledge on analysis and estimation of different	para	ame	eters	,
		ste water				
Experiments		duction to the laboratory and general safety practices for p	olan	t co	ell	
in Plant Biotechnology	cultur 2. Prepa	re. Tration of media, stock preparation and sterilization techni	and	.		
Diotechnology		genomic DNA extraction.	que	· S.		
		opropagation using shoot tip.				
		sculture				
		tip culture				
		netic seed preparation				
		plast isolation				
		formation using Agrobacterium tumefaciens.				
		analysis of DNA by agarose gel electrophoresis. Induction to the laboratory and general safety practices for a	nin	201	co11	
Experiments	cultur	, , , , ,	umm	Iai	cen	
in Animal		ion of fibroblast from chick embryo				
Biotechnology		inoculation methods				
3.0	4. Isolat	ion of genomic DNA from Animal cells				
	_	tification of DNA by spectroscopic method				
		growth analysis				
		viability test – MTT				
		val of frozen cell lines eulture of Adherence cell lines				
		ulture of Adherence cen lines lling of Animal (Mice) – Different routes of drug Admini	stra	tio	n	
		ling techniques of water		0		
Experiments	_	nation of total alkalinity				
in Applied		nation of chloride				
Environmental		nation of total hardness				
Biotechnology		nation of Calcium				
		nation of DO, BOD and COD				
		nation of phosphate nation of Nitrate				
		nation of chromium				
		nation of ferrous ion				
		navi Adhav. Practical Book of Biotechnology & Pla	ant	Ti	ssue	,
Text Books		re.Publisher Chand,2010.				

	2. Kiran Musunuru. Genome Editing: A Practical Guide to Research and
	Clinical Applications. 1 st Edition.Publisher Academic Press Inc.2021.
	3. G Lakshmi Swarajya, P Prabhu Prasadini, Environmental Science: A
	Practical Manual, Publisher BS Publications, 2018
References	1. Vargas V M L, Plant Cell Culture Protocols 4 th Edition, Publisher
	Springer. 2018.
	2. Jayanta Kumar Patra, Gitishree Das, Swagat Kumar Das, Hrudayanath
	Thatoi A Practical Guide to Environmental Biotechnology. Publisher
	Springer. 2020.
	3. Patra, J.K., Das, G., Kumar Das, S., Thatoi, H. A Practical Guide to
	Environmental Biotechnology.Publisher Springer. 2020.
	4. American Public Health Association, American Water Works
	Association and Water Pollution Control Federation, Standard Methods
	for the Examination of Water and Wastewater, 22 nd edition, American
	Public Health Association, Inc, Washington DC. 2012
	5. Turksen, Kursad. Genome Editing. Publisher Springer.2016.
E-Reference	1. https://www.jove.com/science-education/11112/plant-tissue-culture
links	2. https://www.plantcelltechnology.com/blog/meristem-and-shoot-tip-
	culture
	3. https://phytocultures.com/plant-tissue-culture/the-procedure
	4. https://www.microscopemaster.com/cell-culture.html
	5. https://www.youtube.com/watch?v=Fl7aAmzZdMw
	6. https://www.youtube.com/watch?v=9BvTFowr0rI
	7. https://madhavuniversity.edu.in/animal-tissue-culture.html
	8. https://vlab.amrita.edu/index.php?sub=3&brch=258∼=1450&cnt=5
	9. https://www.mpcb.gov.in/sites/default/files/water-quality/reports/LSD-
	NEERI-%20Water%20Quality%20Analysis.pdf

Upon c	ompletion of this course the students will be able to	
CO1	imbibe practical knowledge in plant cell culture techniques	K 1
CO2	understand and gain skills in transformation techniques for plant cells	K2
CO3	learn procedures of culture media preparation, animal cell culture, analysis of cell viability and storage of cells	K4
CO4	gain knowledge about analysis of water quality and solve the problem to the society	K6
CO5	identify the environmental problems, find solution using biotechnique	K6

Mapping of COs with POs & PSOs:

CO	PO									PSO					
	1	2	3	4	5	6	7	8	1	2	3	4	5		
CO1	S	S	S	S	M	S	S	S	S	S	S	S	S		
CO2	S	S	S	S	M	S	S	S	S	S	S	M	S		
CO3	S	S	S	S	M	S	S	S	S	S	S	M	S		
CO4	S	S	S	S	M	S	S	S	S	S	S	M	S		
CO5	S	S	S	S	M	S	S	S	S	S	S	M	S		

SEMESTER IV

Course Code	P21BTE411	CHOICE -I	L	T	P	C					
ELEC	CTIVE -I	STEM CELL BIOLOGY	4	-	-	4					
Cognitive Level	K1: Recall K2: Understand K4:Analyze K6:Create										
Learning Objective	 To learn the characteristics of embryonic and adult stem cells To understand the use of stem cells in organ regeneration To imbibe the ethics in the usage of stem cells in therapy and research 										
Unit I	Stem cell										
multipotent a division, and	and Induced plur differentiation	em cells, Blastula, Inner cell mass, Totipotent, pripotent stem cells characterization, potency, self-ren									
Unit II		on and differentiation		<u> </u>		_					
transduction	pathways and sig	cell proliferation, differentiation, and dedifferentiation nalling molecules involved cellular proliferation, differentiation between cellular proliferation and differentiation of	ren	tia	ior	1,					
Unit III	Embryonic sten										
manipulation from adults (technology (and nuclear trans (Amniotic fluid, o (IPS), genes, and and disadvantages	re obtained, in vitro multiplication: embryonic stem sfer technology. Adult stem cells - Methods to obtain cord blood cells, Mesenchymal stem cells.). Induced placed their mode of action in inducing stemness in a of IPS technology.	ste olur	m o	cel] oter	ls nt					
		sis, kidney regeneration, a neurodegenerative disorder, s	nin	na1 /	or	·d					
injury, tissue Ethical consi	engineering. Eth	ics in using Embryonic stem cells - Human stem cell religion consideration; Stem cell-based theories:	l re	sea	rcł	1:					
Unit V	Application of s	tem Cells									
Parkinson's,	Alzheimer's, Spi betes; Cardiomyon	adult stem cells for therapy in Neurodegenerative nal Code Injuries and other brain Syndromes; Tisspathy; Kidney failure; Liver failure; Cancer; Hemophilia	ue								
Text Book	2. Robert La 2013	E, Stem cells JP brothers medical publishers, 2011. anza, Essentials of Stem Cell Biology, Publisher Acader arke, and Jonathan Frampton, (2020), Stem Cells Biolog			ess	,					
	_	on, Taylor and Francis. 2020	, ai	.14							
References	1.1	Progenitor and Stem Cell Technologies and Therapies V	Voc	od ł	iea	d					
		DG,Adult stem cells: Biology and methods of analysi	s E	Iun	nan	a					
	3. Marek J.	Łos, Andrzej Hudecki and Emilia Wiechec. Publisher ls and Biomaterials for Regenerative Medicine. 2019.	r E	lse	vie	r,					

	4. El-Badri Abdelkodous, Nagwa. Regenerative Medicine and Stem Cell Biology.2020.
E- Reference links	1. https://www.law.berkeley.edu/files/stem_cell_day1_part2_shelanski.pdf 2. https://www.bjcancer.org/Sites_OldFiles/_Library/UserFiles/pdf/stem_cell_h andbook.pdf
	3. https://go.openathens.net/redirector/tulane.edu?url=http://www.sciencedirect.com/science/book/9780123815354

Upon o	completion of this course, the students will be able to	
CO1	know the potency and characteristics of stem cells	K 1
CO2	understand stem cell proliferation, differentiation and dedifferentiation	K2
CO3	compare and contrast the characteristics of embryonic stem cells and adult stem cells	K4
CO4	acquire knowledge on the methods of organ regeneration using stem cells and understand the ethics in stem cell research	K2
CO5	evaluate the applications of stem cells in treating neurodegenerative diseases	K1,K6

Mapping of COs with POs & PSOs:

							/ 9 / 7						
CO			PO PO						PSO				
	1	2	3	40	5	6	7	8	1	2	3	4	5
CO1	S	M	M	M	M	S	M	S	S	S	S	M	M
CO2	S	M	M	S	M	S	S	SW	/S	S	S	M	M
CO3	S	M	S	S	M	STE	M	S	S	S	S	M	M
CO4	S	S	S	S	M	S	Sis	S	S	S	S	S	S
CO5	S	S	S	S	M	SVOI	S	S	S	S	S	S	S

Course Code	P21BTE412	CHOICE -II	L	T	P	C						
	TIVE -I	FOREST CONSERVATION	4	_	_	4						
Cognitive	K1: Recall	TOREST CONSERVATION	•									
Level	K2: Understand	d										
	K4:Analyze											
Learning Objective	To know th vegetation.To understateTo learn ab	To understand the role of forests in environmental sustenance. To learn about the manipulations in management and establishment of forest vegetation.										
Unit I	Silviculture											
	10 1 - 0 0 0	ope of Silviculture. Status of forests in India and their rol	le. (Ger	nera	1						
		hods of propagation, grafting techniques; site factors; n										
planting tech	niques-nursery	beds, polybags and maintenance, water budgeting, gr										
hardening of	seedlings; speci	al approaches; establishment and tending.										
Unit II	Forest Types	8 20 1 37 8										
		d th <mark>eir general classification under different for</mark>										
		their peculiar characters. Types of trees and canopy										
	nd broad leaved	I tree species. Trees in tropical, sub-tropica, temperate	and	al	pin	e						
regions.			ı									
Unit III	Forest soil and											
properties. S conservation	oil conservation	factors affecting soil formation; physical, chemical and definition, causes for erosion; types - wind and water of eroded soils/areas, wind breaks, shelter belts; sand definition.	er e	ros	sion	1;						
Unit IV		ement and Management Systems				-						
rotation, nor commercial strategic plar governance.	mal forest, grow forests, forest c nning, (iii) Appr	chniques; stand structure and dynamics, sustained yield ying stock; regulation of yield; management of forest prover monitoring. Approaches viz., (i) site-specific platoval, sanction and expenditure, (iv) Monitoring (v) Rep	lan nni	tati ng,	ons (ii	s, i)						
Unit V	Forest Disturb	pance										
afforestation	and forest rege human impacts;	nd biotic, destructive agencies, insect-pests and disease neration in absorption of CO2. Effect of wild animals encroachment, poaching, grazing, live fencing, shifting	or cult	iva	ores	st n						
Text Books	bandhu,201 2. Nyland R.D 3. Sagwal S.S. 4. Mark S. As	t, Principles and Practice Of Silviculture, Publisher 5 D.Silviculture concepts and Applications, Publisher CBS,2 D.A Textbook of Silviculture, Publisher Kalyani, 2017 The hton, Matthew J. Kelty. The Practice of Silviculture: Appublisher Wiley, 2018	015	í								

References	1. Francesco Ferrini, Cecil C. Konijnendijk van den Bosch, Alessio Fini,
	Routledge Handbook of Urban Forestry, Routledge,2017
	2. Sergius Alexander Wilde ,Forest Soils: Their Properties and Relation to
	Silviculture, Publisher Forgotten Books,2019
	3. Kevin Laughlin O'Hara, Multiaged Silviculture: Managing for Complex
	Forest Stand Structures, Oxford University Press, 2014
E-	1.https://forestrypedia.com/download/general-silviculture-notes-by-naeem-
Reference	javid-muhammad-hassani/
links	2.http://www.westbengalforest.gov.in/upload/development/cm24.pdf
	3.https://www.forestrynotes.in/
	4.http://www.fao.org/3/ap467e/ap467e00.pdf
	5.https://www.geospatialworld.net/article/forest-management-information-
	system-fmis-2/

Upon	Upon completion of this course, the students will be able to									
CO1	acquire knowledge on forests in India and their role	K 1								
CO ₂	understand the trees and their general classification under different	K2								
	forest types									
CO3	compare the role of forests in conserving soils	K4								
CO4	know the Forest Management and Management Systems	K1								
CO5	gain knowledge about the causes, importance of afforestation	K2								
	deforestation and reforestation									

Mapping of COs with POs & PSOs:

CO	I PO						PSO							
	1	2	3	4	5 23	6	7 16	8	1	2	3	4	5	
CO1	S	M	M	M	M	M	M	S	S	S	S	M	S	
CO2	S	S	M	M	MS	S	SNS	S	S	S	S	M	S	
CO3	S	M	M	M	M	S	S	S	S	S	S	S	S	
CO4	S	M	M	S	M	S	M	S	S	S	S	M	S	
CO5	S	M	M	S	M	S	M	S	S	S	S	M	S	

Course	P21BTE413	CHOICE -3	L	T	P	C							
Code													
ELEC	TIVE -I	NANOTECHNOLOGY AND CANCER BIOLOGY	4	-	-	4							
Cognitive	K1: Recall	BIOLOGI											
Level	K2: Understand	1											
Learning		owledge and basic understanding of nanotechnology and	can	cer									
Objective	_	ne properties of nanomaterials, and the principles behind				ed							
		al and computational techniques for studying nanomateria											
	 To attain k 	nowledge on synthesis of nanomaterials, characterization	n ai	nd	the	ir							
	application	pplication in almost all the field to the benefit of humankind.											
Unit I	Nanotechnolog	50											
		Nanowires & properties, 2D films. Nano scale materials.	Van	opo	ore	s.							
		cles and Nanomaterials.											
Unit II		nanotechnology		r·	11								
		plications, Nano carriers for drug delivery-polymeric NP											
		ins as pharmaceutical carriers. Solid lipid NP as dru Characterization & therapeutic applications. Nano											
		able materials, Devices, Surgical aids, diagnostic tool											
testing, Imag		allo materials, Devices, Surgicul alas, diagnostic tool	σ,	GCI	.101.	10							
Unit III		f Nanotechnology											
Nanotechnol		research & therapy. Environmental nano remediation to	ech	nol	og	у.							
		and Biological methods. Nano filtration for the treatme											
removal of or	rganics, Inorgani	cs and pathogens. Nanotechnology for water purification.											
Unit IV	Cancer												
		ncer types, characteristics of cancer cells, carcinogenes											
_	_	rogression, termination. Factors responsible for Carci	nog	gen	esi	s;							
	emical and Biolo												
Unit V	Tumour immu			Гот									
		vaccine development, immunotherapy and its limitation nses. Principles of chemotherapy and chemoprevention.	is,	I UI	not	JL							
cerrevasions		nith, Principles of Cancer Biology, Publisher Pearson	Fd	1102	atic)n							
Text Books	India,20		Lu	uci	ııı	/11							
	2. Sunipa		r	Sa	rka	ır,							
	Nanoted	chnology:Synthesis to Applications, Publisher CRC Press	,20	18									
	3. Ann-Mai	rie Broome, Cancer Nanotechnology, Academic Press. 20	18										
Defe	1 0 00	- M.C. and D.L. Ell Ell Ell Clink	1.1	. 1	1								
References		y M.Cooper and Robert E.Hausman. The Cell: A											
	* *	ch 7th Edition, ASM Press, Washington D.C. & Sinauer Anderland, Massachusetts. 2016.	155	oci	ate	٥,							
	· · · · · · · · · · · · · · · · · · ·	Karp, Harris, D. Cell and Molecular Biology – Cor	icei	ots	ar	nd							
		nents (ed), John Wiley & Sons Inc, New York,2016	_]		ui.								
		Jain Kaisar Raza Ashish Kumar Agrawal Ankur Vaid	ya	1 st	E	d,							
		chnology Applications for Cancer Chemotherapy, Elsevie											
		·											

E-	1. https://books.google.co.in/books?id=81vBBwAAQBAJ&printsec=front
Reference	cover&source=gbs_ge_summary_r&cad=0
links	2. https://www.pdfdrive.com/cancer-nanotechnology-methods-and-
	protocols-d158801917.html
	3. https://www.pdfdrive.com/introduction-to-cancer-biology-d58366931.html
	4. https://www.pdfdrive.com/nanotechnology-and-nanosensors-introduction-to-nanotechnology-d187619895.html

Upon c	ompletion of this course the students will be able to							
CO1	know nanotechnology, nanomaterial and nanoparticles	K1						
CO2	acquire knowledge on the application of nanotechnology in different field and use as problem solving solution							
CO3	update the research trends in Nanotechnology for cancer research and therapy	K2						
CO4	attain knowledge about epidemiology of cancer, cancer types, characteristics of cancer cells in molecular aspects	K2						
CO5	gain knowledge on vaccine development, chemotherapy and chemoprevention	K2						

Mapping of COs with POs & PSOs:

CO				P	O		/e i s				PSO	PSO				
	1	2	3	4	5	6	7	8 >	1	2	3	4	5			
CO1	S	S	M	M	M	M	M	S	S	S	M	M	M			
CO2	S	S	S	M	M	M	M	S	S	S	M	M	S			
CO3	S	S	S	S	M	S	S	S	S	S	S	S	M			
CO4	S	S	S	M	S	ST 8	M	S	S	S	S	S	S			
CO5	S	S	S	S	SEO	S	Sis	S	S	S	S	S	S			

Course Code	P21BTE414	CHOICE -4	L	T	P	C
	TIVE - I	DRUG METABOLISM	4	-	_	4
Cognitive	K1: Recall					
Level	K2: Understand	d				
	K3:Apply					
T	K4: Analyze			C 1		
Learning Objective		erstand the classification of drug and mechanism of actio	n o	i dr	ugs	S
Objective	_	knowledge on pharmacokinetics				
		w about the drugs for metabolic disorders and its toxicity	1			
Unit I	General Pharr	St .				
		gy, sources of drugs, Classification of drugs based or				
		nistration, site of action of drugs. Mechanism of action, of drugs, factors modifying drug action. Dose response cur				
and LD50.	momed effect of	drugs, factors mountying drug action. Dose response cu	ı vc-	. 151	J J(U
Unit II	Pharmacokine	etics DB6ffir Usi				
Absorption	and distribution	n of drugs, importance of drug – protein interact	ion.	Г)ru	g
		yay of drug metabolism, phase I and phase II reaction				
-		osomal reactions of drug metabolism, drug metabolizing	en	zyn	nes	S .
	tion of liver and	kidney				
Unit III	Therapeutics	₹ 5 E				
Biochemical	mode of action	of antibiotics- penicillin and chloramphenicol, actions of				
	d antimalarial					
		ncy and drug efficacy. General principles of chem				
_	-	infections, fungal infections, viral diseases. Introductherapy of cancer.	luct	юп	ι	O
Unit IV		pharmacological activity				
		0 4	<u> </u>			_
		y and antipyretic agents, gastrointestinal drugs, anti				
		ancer and anti-fertility agents. Drugs for metabolic disc	orae	ers	IIK(e
Unit V	Clinical Toxic	nic, anti-obesity and hepatoprotective agents				
			<u> </u>			
		toxicity – occupational, environmental and pharmaceuti		•	-	
		n of action. Factors affecting toxicity- Drug tolerance, in sitivity, antagonism and synergism. Methods of detect				
		ffects. Rational prescription of drugs. Toxicity of anticar				_
	_	and marker parameters.		GI.	~5°	•
Text Books		Uetrecht, William Trager, (2007)1st Ed, Drug Metabolism	nCh	iem	iica	al
	and Enz	zymatic Aspects, Taylor and Francis.				
		G. Gordon, Skett, Paul, (1986), 1 st Ed, Introduction	to	D	rug	g
		olism, Springer US.	, .			
		Mino R., Ionescu, Corina, (2005), 1 st , Ed, Drug Mo	etat)Oli	sm	l,
References		Concepts, Springer Netherlands 1 D. Coleman . Human Drug Metabolism, 3 rd Edition,201	8		—	
Acici ciices		n Katzung, Anthony TrevorBasic and Clinical Phar		colo	ροι	<i>J</i> .
		· · · · · · · · · · · · · · · · · · ·		. 510	-01	,
	McGrav	w Hill Professional. 2014				

	of pharmacology: the pathophysiologic basis of drug therapy. Lippincott Williams &Wilkins. 2011.
	4. Paul G. Pearson, Larry C. Wienkers, Handbook of Drug Metabolism, Publisher CRC Press, 2021
E-	1. https://www.pdfdrive.com/drug-metabolism-e-library-fakultas-kedokteran-
Reference	uwks-d3133731.html 2. https://www.pdfdrive.com/principles-of-pharmacology-the-
links	pathophysiologic-basis-of-drug-therapy-d157890965.html
	3. https://www.pdfdrive.com/pharmacology-d33542642.html
	4. https://www.pdfdrive.com/basic-clinical-pharmacology-e34443843.html

Upon	completion of this course, the students will be able to	
CO1	classify drugs and their mechanism of action	K2
CO2	illustrate the importance of experimental models and know drug	К3
	metabolism	
CO3	compare the action of antibiotics and distinguish between cancer	K4
	therapies for parasitic, fungal and viral disease	
CO4	know the pharmacological activity of the drugs used in different	K2
	metabolic disorders	
CO5	classify toxins, types, detection methods and the factors affecting toxicity	K2

Mapping of COs with POs & PSOs:

CO				> P	O	3		\ \			PSO			
	1	2	3	40	5	6	7	8	1	2	3	4	5	
CO1	S	M	M	M	S	S	M	SS	S	S	S	M	S	
CO2	S	S	S	M	S	S	S	SU	S	S	S	S	S	
CO3	S	M	S	M	S	STA	M	S	S	S	S	M	S	
CO4	S	M	S	M	S	S	Mis	S	S	S	S	S	S	
CO5	S	M	M	M	S	SVON	M	S	S	S	S	M	S	

Course Code	P21BTE421	CHOICE -1	L	T	P	C				
	CTIVE II	MOLECULAR MODELLING AND DRUG DESIGNING	4	-	-	4				
Cognitive Level	K2: Understand K3: Apply K4: Analyze									
Learning Objective										
Unit I		nics & concepts in molecular modelling								
Bond stretch interactions;	vstems, potential en ning; angle bendi Vander Waals intera	ergy surfaces. Introduction to quantum mechanics. ng. torsional terms; non-bonded interactions; actions								
Unit II		nics and Monte Carlo simulation								
	/ /	MD simulation, Molecular dynamics algorithms.								
Unit III	Analysis and Pro									
_	•	tional frequencies: potential energy surface, pint vibrational energies.	har	mon	ic	VS.				
Unit IV	Modelling									
	_	Protein Threading. Drug design - Structure-base g lead compounds by searching 3D databases; d								
Unit V	Molecular Docki									
Docking - mo	olecular modeling i	n d <mark>rug design – structure-</mark> based drug design – pha	rma	cop	hore	żs -				
Text Books	2. Schneider, 2008	A, Molecular Modelling And Bonding, C Publishin Molecular Design Concepts And Applications, umar, Anju Sharma, Tiwari, Introduction To Di	, Jo	hn						
	4. Cohen Cla Elsevier In									
References	education	R, Molecular Modeling, Principles & Applicat Ltd, UK. 2010. Drug Discovery, Design & Development Lamb 2013								
	3. Clark T, Computati chemistry,	Thurston DE, and Banting L,Drug Desigonal Techniques & Applications Royal	SC	ciet	У	of				
E- Reference	1. https://ww 2. https://ww	w.mdpi.com/books/pdfview/book/1187 w.kobo.com/us/en/ebook/molecular-modelling-and	-							
links	3. https://ww	w.ncbi.nlm.nih.gov/pmc/articles/PMC6539951/								

- 4. https://link.springer.com/article/10.1007/BF02834015
- 5. http://www.drugdiscoverytoday.com/view/25419/molecular-modeling/

Upon completion of this course the students will be able to							
CO1	know the concepts of Molecular modelling						
CO2	employ different designs and potentials in molecular dynamics simulation	К3					
CO3	illustrate the concept of optimization and vibrational frequencies	К3					
CO4	understand homology modelling and the methods to identify lead compounds	К3					
CO5	compare different drug designs for molecular modelling by docking	K4					

Mapping of COs with POs & PSOs:

	L DESTITUTE OF THE PROPERTY OF													
CO				P	OURU		प्राचित्र ।				PSO			
	1	2	3	4/	3	EQU.	7	8	1	2	3	4	5	
CO1	S	S	M	S	SO	S	M	S. S.	S	S	S	S	S	
CO2	S	S	S	S	S	S	M	S	S	S	S	S	S	
CO3	S	S	S	S	S	S	M	S S	S	S	S	S	S	
CO4	S	S	S	S	S	S	M	S	S	S	S	S	S	
CO5	S	S	S	S	S	S	S	S	S	S	S	S	S	

Strongly Correlating **(S)** (W) - 3 marks;

Moderately Correlating

(M)

- 2 marks

Weakly Correlating

- 1 mark;

No Correlation

- 0 mark (N)

Course Code	P21BTE422	P21BTE422 CHOICE - 2							
ELECT	TIVE - II	WILD LIFE CONSERVATION	4	-	-	4			
Cognitive Level	K1: Recall K2: Understand								
Learning Objective		erstand about habitat analysis, Human-wildlife interaction wabout the concepts of wildlife management	S						
Unit I	Introduction								
Values and eth	nics of wildlife c	onservation; importance of conservation. Conservation v	s pi	rote	ecti	on			

Values and ethics of wildlife conservation; importance of conservation. Conservation vs protection Concept of Buffer zones, Wildlife corridors, Strategies to reduce human-wildlife interactions.

Unit II Habitat analysis

Types of Habitats & their major ecological factors. Ecological Succession & climax ecosystems (e.g. Sholas). Maximizing usage of Habitat resources by populations. Insular habitats & insular flora & fauna Extreme. Habitats and their flora & fauna (Dark Caves, deep sea etc.). Evaluation and management of wild life - Physical parameters and Biological Parameters; Standard evaluation procedures: Faecal analysis of ungulates and carnivores. Geographical Information System (GIS), Global Positioning System (GPS) and Remote Sensing (RS).

Unit III Human-wildlife interactions

Poaching, illegal trading, conflict management and shifting from extraction to preservation; effect of extinction of a species on ecosystem; Forest landscape restoration. Conservation Vs protection. Concept of Buffer zones, Wildlife corridors Strategies to reduce human-wildlife interactions Role of Government and NGOs in controlling human-wildlife interactions. Socio-economic issues related to human-wildlife interaction.

Unit IV Concepts of Wildlife management

Protected Area Network (PAN), WWFN, IUCN, and CITES. Wild life Legislation – Wild life Protection act (1972), its amendments and implementation. IUCN Red data book and red list categories (only names), Protected areas National parks & sanctuaries, Community reserve; Important features of protected areas in India; Project Tiger and Project Elephant.

Unit V Sustainable wildlife management

Natural resource management. Eco tourism / wild life tourism in forests; various Environmental movements in India: Bishnoi movement, Chipko movement, Narmada bachao andolan, Silent valley movement, Baliyapal movement.

valley moveme	ent, Baliyapal movement.
Text Books	1. Reena Mathur, Wildlife Conservation and Management, Rastogi
	Publications,2018
	2. Singh S K, Textbook of Wildlife Management, Publisher CBS, 2020
	3. Goutam Kumar Saha, Subhendu Mazumdar, Wildlife Biology: An Indian
	Perspective, Publisher PHI Learning, 2017
	4. Jagbir Singh, Ecotourism, I.K. International Publishing House Pvt. Ltd. 2010
	5. S.K. Singh, Textbook Of Wildlife Management, Publisher CBS Publishers &
	Distributors Pvt Ltd, India,2018
D 6	
References	1. Sutherland, W.J. The Conservation Handbook: Research, Management and
	Policy. Blackwell Sciences 2010
	2. Woodroffe R., S. Thirgood and A. Rabinowitz. People and Wildlife,
	Conflict or Coexistence? Cambridge University Press, 2011
	3. McCleery, Robert A, Moorman, Christopher, Peterson, M. Nils, Urban

	Wildlife Conservation, Springer, 2014
	4. David A. Fennell, Ecotourism, Publisher Routledge, 2014
E-Reference	1. https://www.pdfdrive.com/wildlife-ecology-conservation-and-management-
links	2nd-edition-d184311905.html
	2. https://www.pdfdrive.com/comprehensive-wildlife-conservation-strategy-
	e38430632.html
	3. https://www.pdfdrive.com/wildlife-ecology-and-management-wildlife-
	producers-association-e9899184.html

Upon	completion of this course, the students will be able to							
CO1	understand the values and ethics of wildlife conservation K2							
CO2	gain knowledge on the salient features of habitats and their ecological factors and standard procedure for assessment and management of wild life	K2						
CO3	attain the basic knowledge of human-wildlife interactions and socio- economic issues related to human-wildlife interaction.	K1						
CO4	understand the concepts of wildlife management and wild life protection act	K2						
CO5	identify the value and importance of Eco tourism	K2						

Mapping of COs with POs & PSOs:

							//2005						
CO				P	O						PSO		
	1	2	3	4	5	6	7	8	1	2	3	4	5
CO1	S	M	S	M	M	S	Μ.	SS	S	S	S	M	S
CO2	S	M	S	M	M	S	M	S	S	S	S	M	S
CO3	S	M	S	M	M	S	Sal	S	S	S	S	M	S
CO4	S	M	S	M	M	STB	M	Z	S	S	S	M	S
CO5	S	M	S	M	MS	S	M	S	S	S	S	M	S

Course Code	P21BTE423	CHOICE -3	L	T	P	C
	TIVE - II	HUMAN PATHOLOGY	4	-	-	4
Cognitive Level	K1: Recall K2: Understand K3-Apply K4: Analyze	1				
Learning Objective	• To gain Gastrointe	a theoretical knowledge in general pathology and mycolo knowledge about systemic pathology and Tur stinal system, Respiratory tract and Breast and and acquire knowledge on Transfusion Medicine and	not	ırs		n al
Unit I	General Patho	logy				
Inflammation Neoplasia		thology, Cell injury and cell death, Cellular accurately disturbances, Immunological disorders, I				
Unit II	Mycology	(4)				
Superficial n infections. La	nycotic infection aboratory diagno	y and reproduction. Classification of fungi .Opportunism. Fungi causing subcutaneous mycoses. Fungi causing sis of fungal infections.			_	
Unit III	Systemic Path	ology O				
rectum and immunologic conditions. P	anal canal. I al and neoplastic athogenesis path	orders of mouth, salivary glands ,esophagus, stomach , Respiratory tract – infections, inflammations, envir e disorders and their identification. Breast - Tumors and to ology and diagnosis.	oni	nei	nta	l,
Unit IV	Haematology	7 9 8				
Anaemia – d Disorders of	leficiency, hemo leucocytes and p	gy of blood cells, bone marrow, general alterations in alysis and other causes. Disorders of hemostasis and coplatelets – quantitative, qualitative and in neoplastic proled disorders.	agu	ılat	ior	ı.
Unit V	Transfusion M	Iedicine				
principles and	d methods emplo cluding tissue m 1. Husain	serology and transfusion medicine. Clinical Pathology byed in tissue processing, paraffin and frozen sections an icroarrays. A. Sattar, Fundamentals of Pathology. Published by Path	d s	taiı	nin	g
	3. Vinay	Mohan. Textbook of Pathology, 7th Edition. s,Medical Publishers Pvt. Limited,2014. Kumar Abul Abbas Jon Aster. Pathologic Basis of D Elsevier. 2014.			pe	
References	Pvt. Ltd 2. L. Max Saunder 3. Inderbin	imilian Buja, Gerhard R. F. Krueger, Human Pathology.	Pu	bli	she	r

E-	1. https://www.pdfdrive.com/pathology-handbook-capital-pathology-
Reference	e36414786.html 2. https://www.pdfdrive.com/genitourinary-pathology-a-volume-in-
links	foundations-in-diagnostic-pathology-series-high-yield-e176374227.html 3. https://www.pdfdrive.com/harsh-mohan-textbook-of-pathology-e52206258.html
	4. https://www.pdfdrive.com/fundamentals-of-pathology-pathoma-2018-e185838619.html
	5. https://www.pdfdrive.com/pathology-usmle-step-1-volume-1-basic-pathology-e187109588.html

Upon c	Upon completion of this course, the students will be able to					
CO1	understand the basics of pathological disorders	K2				
CO2	gain knowledge on fungal diseases and diagnosis	K1				
CO3	acquire knowledge on Systemic pathology and disorders in blood	K2				
CO4	illustrate the developments in blood cells and explain their disorders	К3				
CO5	compare the different staining techniques in clinical pathology	K4				

Mapping of COs with POs & PSOs:

CO				P	PO				PSO					
	1	2	3	4	5	6	7	8	1	2	3	4	5	
CO1	S	M	M	M	M	M	M	\S >	S	S	S	M	S	
CO2	S	M	M	M	M	M	M	S	S	S	S	M	S	
CO3	S	M	S	M	M	M	M	S	S	S	S	M	S	
CO4	S	M	S	M	M	S	M.	S	S	S	S	M	S	
CO5	S	M	S	S	M	STE	M	S	S	S	S	M	S	
					PER		115	2.						

Course	P21BTE424	CHOICE - 4	L	T	P	C
Code	CTIVE- II	BIOBUSINESS	4			4
Cognitive	K1: Recall	BIOBUSINESS	4	-	-	4
Level	K1. Recall K2: Understand					
Level	K3: Apply					
	K5: Evaluate					
Learning		owledge on different aspects of biobusiness.				
Objective	To acquire	information about various sector in bio business				
	_	and IPR related to biobusiness				
Unit I	Fundamentals of	Bio business				
History of evo	lution of Bio Busin	ess, Importance of Finance for Bio business –Se	ctor	ial s	supp	ort
	nt of India - policies	-				
Unit II		business in various sectors				
		nces, Agriculture and Agri-biotechnology, Env	'iroi	nme	nt a	ınd
Environmenta	l Biotechnology.	3511811				
Unit III	Business Models	in Bio business				
Product Based	d-Service Based-Su	bscription Based-Integrated Models.				
Unit IV	Best Practices	10 TO 10 10 10 10 10 10 10 10 10 10 10 10 10				
Current Good	Manufacturing Pra	ctices (cGMP), Current Good Laboratory Practice	es (c	GL	P).	
Unit V	IPR					
Determining	"patentability"; Ind	ustry-wise implications; use of patents – relevan	nt c	ase	stud	lies
		nportance of IPR in the Pharmaceutical In				
	_	enting- Marketing. Technology transfer, Licensin		•		Ü
Text Books		J. Gitman, Carl McDaniel, Amit Shah, Monique		eece	Lir,	ıda
		hann Talsma, James C. Hyatt Introduction				
	Publisher (OpenStax,2018.				
	2. Lenssen,	Gilbert G., Smith, Craig, Managing Sustaina	ble	Bu	sine	ess,
	Springer, 2	2019.				
References	1. Shahi, G.	BioBusiness in Asia: How Asian Countries Can	Ca	pita	lize	on
	the Life Sc	eience Revolution. Pearson Prentice Hall. 2004.		_		
	2. Xu, Xiaozl	hou ,Introduction to Entrepreneurship, Springer,20)20			
	3. Vashney,	Fundamentals of Entrepreneurship, Sahi	tya	В	hav	van
	Publication	ns, 2019.				
E-Reference	1. http	os://www.wur.nl/en/show/biobusiness.htm				
links	2. http	os://www.bio.org/save				
1111182	_	os://www.bio-				
		.com/webroot/web/pdf/lse/literature/Biobusiness.	pdf			
	-	os://www.the-scientist.com/tag/biobusiness	_			
	-	os://www.crg.eu/en/content/training-courses/biobu	ısin	ess-		
	ent	repreneurship				

Upon c	Upon completion of this course, the students will be able to					
CO1	know the history of bio business, support from government and the	K1				
	current scenario					
CO2	collect details of the various sectors of bio business	K5				
CO3	differentiate the types of business models viz. product, subscription	K2				
CO4	understand the importance of cGMP and cGLP	К3				
CO5	get knowledge on the role of IPR in bio business	K4				

Mapping of COs with POs & PSOs:

CO		PO							PSO				
	1	2	3	4	5	6	7	8	1	2	3	4	5
CO1	M	M	S	M	M	S	M	S	M	S	S	M	S
CO2	M	M	S	M	M	S	M	S	M	S	S	M	S
CO3	M	M	S	M	M	Seemle	M	S	M	S	S	M	S
CO4	M	M	S	M	M	S	Mode	S	S	S	S	M	S
CO5	M	M	S	M	M	SEQU	M	S	S	S	S	M	S

Strongly Correlating (S)- 3 marks Weakly Correlating (W) - 1 mark Moderately Correlating (M) - 2 marks No Correlation (N) - 0 mark

Course	P21BTN211		L	T	P	C
Code	NIME	INDUSTRIAL WASTE MANAGEMENT	4			4
	NME		4	-	-	4
Cognitive	K1: Recall					
Level	K2: Understand					
	K3-Apply					
	K4: Analyze	1 1 00 1 1 00	•			
Learning	_	owledge on effluent characteristics and effects on en	Viro	nme	nt	
Objective		ad the importance of industries for development				
TT . *4 T		skill for designing ETP for industries				
Unit I	Industries			L .	,•	
		tance of industries – Industrial pollution –cha				
		ndustrial effluents on streams and land. Envir	onm	enta	u 1a	lWS
	Ī	of industrial effluents. Waste audit.				
Unit II	Wastewater Treat					
		ed treatment: Classification and application of				
		rocess analysis, biological waste water treatm	nent	- 1	UAS	SB,
Wastewater d	isposal and Reuse. S					
Unit III		D <mark>istilleries, Tanneries,</mark> Fertilizer industries a	nd			
	pharmaceutical in					
Sources, cha	racteristics of waste	es, effects on receiving water bodies and Trea	tme	nt c	of th	ıeir
wastes and di	sposal.					
Unit IV	Cement industries					
Sources of po	ollution and wastes.	Effect of wastes. Control technique of pollution.	ther	mal	pov	ver
plants, Sourc	es of pollution, cha	aracteristics of pollutants and their effects. Po	lluti	on	cont	rol
techniques.	70					
Unit V	Hazardous waste	management S/2				
Biotechnolog	ical application of h	nazardous waste management and management	of 1	Reso	ourc	es:
bioremediatio	n, phytoremediation	, Use of microbial systems, Waste water treatn	nent	usii	ng r	oot
zone treatmen	nt by plants, Reclama	tion of wasteland: biomass production for Bioga	s.			
	 Lagrega, Hazard 	ous Waste Management, Medtech, 2015				
Text Books	-	ellman, Handbook of Water and Wastewater	Γreat	men	t Pl	lant
	Operations,CRO					
		D., Industrial Wastewater Treatment", Prentice Hall of	of Inc	dia, l	New	
	Delhi 2010.					
Defenences	•	onmental Management" Vikas Publications, 2010.	o du	ati a i	. En	0.122
References	1. Balagurusamy		odu			UIII
	_	gestion to a Sustainable Bioenergy Industry, Sprin Ekström , Waste Management and	_	2020 Susta		hla
		,	2	oust	ama	DIE
	· ·	Publisher Routledge,2014 Environmental Weste Management CRC Press	0015	:		
	· ·	Environmental Waste Management, CRC Press, 2			٠ <i>t</i> -	nd
		ebashish, Dubey, Brajesh K., Goel, Sudha, T	real	mer	ıı i	ıIIU
	Disposal of So	olid and Hazardous Wastes, springer, 2021				

E -
Reference
links

- https://www.mysciencework.com/publication/download/lecture-notes-cell-biology-1636c320/adc18b1228577d5353c56fdf7b69b6de
- https://gurukpo.com/Content/Bsc-biotech/Cell_Biology.pdf
- https://www.microscopemaster.com/cell-biology.html
- https://microbenotes.com/category/cell-biology/

Upon co	Upon completion of this course the students will be able to					
CO1	gain knowledge on effect of industries waste on environment and	K 1				
	environmental legislation					
CO2	understand the basic of common waste water treatment	K2				
CO3	acquire knowledge on effluent characteristic and treatment process of	K1				
	various industrial effluent					
CO4	analyse the characteristics of effluent and student can able to design	K4				
	treatment process for industries					
CO5	apply biotechniques to control the hazards waste pollution	К3				

Mapping of COs with POs & PSOs:

CO				/ P	PO S				PSO				
	1	2	3	4 6	5	6	7	8	1	2	3	4	5
CO1	S	S	S	S	M	M	S	S	S	M	S	S	S
CO2	S	S	S	M	M	S	S	US 6	S	S	S	S	S
CO3	S	S	S	M	M	S	S	S	S	S	S	S	S
CO4	S	S	S	M	M	S	S	S	S	S	S	S	S
CO5	S	S	S	M	M	S	S	S	S	S	S	S	S

Strongly Correlating (S)- 3 marks Weakly Correlating (W) - 1 mark Moderately Correlating (M) - 2 marks No Correlation (N) - 0 mark

Course	P21BTV11		Total	C						
Code		CHROMATOGRAPHIC TECHNIQUES	Hours							
Value Added	l Programme		30	2						
Cognitive	K1: Recall									
Level	K2: Understa	nd								
	K3: Apply									
	K4: Analyze									
Learning	 To learn 	• To learn the basics of advanced chromatographic techniques, their principles								
Objective	and appl									
	-	uire knowledge on data collection and data inter	rpretation	of						
		ographic techniques								
		w how to analyze and compare the chromatograms	with libr	ary						
	database									
Unit I		aphy techniques	•							
		Principles and application of various chromatography tecl	nnıques.							
Unit II	HPLC									
		nents- column and detector, working mechanism-isocratic	and gradi	ient						
	•	PLC. Safety considerations.								
Unit III	GC-MS		6.00.1							
		ents-column, detector, working mechanism and application	on of GC-I	VIS.						
Safety consid Unit IV	HPLC & GC	MC								
				4						
of HPLC an		on of instrument, Sample & solvent preparation, operation	ı –parame	ters						
Unit V		-MS Results interpretation								
		ysis with their mass and compared with library database.								
Text	_	rakumari, Bioinstrumentation, MJP Publishers, 2015.								
Books		undanes, Chromatography: Basic Principles, Sample Pre	narations :	and						
Books		ed Methods, Wiley, 2013	parations	um						
		er-Wilde, Katja, Engewald, Werner, Practical Gas Chr	omatograp	ohy,						
		ger, 2014	C I	3 /						
References	1. Reilly	M J,Bioinstrumentation, Publisher CBS,2018.								
	2. Hans-	Joachim Hübschmann, Handbook of GC-MS: Funda	amentals	and						
		eations, Wiley, 2015.								
E-		/microbenotes.com/types-of-chromatography/								
Reference	-	/nptel.ac.in/content/storage2/courses/103108100/module7		pdf						
links	-	/www.aweimagazine.com/article/chromatographic-technic	•							
	-	//www.sciencedirect.com/topics/engineering/chromatograp	ohic-							
	techni	que								

Course outcome

Upon co	Upon completion of this course, the students will be able to							
CO1	compare the method in chromatography	K4						
CO2	know the principle, components and working of HPLC	K1, K2						
CO3	perform confidentially the experimentation with GC-MS and apply	K3, K4						
	in research and analysis							
CO4	calibrate and operate HPLC and GCMS	K1, K2, K3						
CO5	learn and interpret the HPLC and GCMS data	K1, K2, K3, K4						

Mapping of COs with POs & PSOs:

CO	PO									PSO				
	1	2	3	4	5	6	7	8	1	2	3	4	5	
CO1	S	S	S	S	M	S	M	S	S	S	M	S	S	
CO2	S	S	S	M	Men	S	Moi	S	S	S	S	S	S	
CO3	S	S	S	S	M	SOI	M	S	S	S	S	S	S	
CO4	S	S	S	M	M	S	M	Sg.	S	S	S	M	S	
CO5	S	S	S	M	M	S	M	SB	S	S	S	S	S	

Strongly Correlating (S) Weakly Correlating (W) -3 marks; Moderately Correlating

(M) - 2 marks

- 1 mark; No Correlation

(N) - 0 mark

Course Code	P21BTV42	SYSTEM BIOLOGY	Total Hours	C
Value A	Added Programme	SISIEM BIOLOGI	30	2
Cognitive Level	K1: Recall K2: Understand			
	K3: Apply			
Learning	• To know the ba	asic concepts in pharmaceutical industry		
Objective		drug development, approval process and manu	ufacturing	of
	biopharmaceuti	icals.		
	 To know the st 	eps involved in drug discovery process		
Unit I	Introduction and Bio	ological networks		

Introduction - System-level Understanding of Biological Systems - Advanced Measurement Systems - Introduction to Biological Networks and Basic Concepts - Metabolic, Signaling and Regulatory networks - Why build and study models? - Characterizing dynamic states - Formulating and studying dynamic network models - Properties of dynamic states - Network structure versus dynamics

Unit II Standard models and Approaches in systems biology

Metabolism- enzyme kinetics and thermodynamics- Michaelis - Menten Kinetics - metabolic networks- metabolic control analysis - Signal transduction- introduction- function and structures-interactions- structural components - signaling selected biological processes - mathematical models - prediction of biological systems.

Unit III | E-Cell project

Organization - History - Research group - modeling methods - formalism - techniques numerical simulation algorithm-mathematical analysis methods-software environment-projects models-applications chemotaxis - molecular clock-circadian rhythms-oxidation stress-multi-enzyme systems.

Unit IV Systems biology software

Systems biology software project: About the project-model inter change-code use-bio-models-online services-SBML Layout viewer-SBML validation-simulation translator-model repository-SBW broker - Jurnac-J-designer- BioSpice - BioUMC - CellDesigner - Cytoscape - Dizzy-Oscillator- Virtual cell - virtual rice project.

Unit V Introduction to synthetic biology

Definition – Synthetic Biology versus Systems Biology - Synthesis and Engineering Tools - DNA Synthesis - Protein Engineering - Pathway Engineering - Genome Engineering - Computational and Theoretical Tools – Genomics, Proteomics and Metabolomics Tools - Applications in Synthetic Biology – Molecular, Pathway and Whole Cell Levels - Challenges and Future Perspectives.

Text Books	1.	Joseph DiStefano, Dynamic Systems Biology Modeling and Simulation,										
		PublisherAcademic Press,2015										
	2.	Vikram Singh, Pawan K. Dhar, Systems and Synthetic Biology, Publisher										
	3.	Springer Nature,2015										
	4.	Edda Klipp, Wolfram Liebermeister, Christoph Wierling, Axel										
		Kowald, Systems Biology, Publisher Wiley-Blackwell, 2016										
	5.	Eberhard Voit, Systems Biology, Publisher Garland Science,2017										
References	1.	Bernhard . Palsson, Systems Biology – Simulation of Dynamic Network										
		States, Cambridge Univ. Press, UK,2011.										
	2.	Huimin Zhao (Ed.), Synthetic Biology: Tools and Applications, Academic										

	Press, Elsevier, USA,2013. 3. Kayvan Najarian, Siamak Najarian,Shahriar Gharibzadeh,Christopher N. Eichelberger, Systems Biology and Bioinformatics A Computational Approach, CRC Press,2017
E-	1. www.systems-biology.org/
Reference	2. https://www.sysbiol.cam.ac.uk/
links	3. https://www.systemsbiology.org/

Upon con	Upon completion of this course the students will be able to								
CO1	understand the comprehensive measurements of biological systems.	K2							
CO2	know the factors involved in Biological System Design.	K1							
CO3	gain knowledge on the systems biology tools: E-Cell	K2							
CO4	know the networking of genes and protein interaction networks.	K1							
CO5	relate the engineering principles in Synthetic Biology and its applications.	К3							

Mapping of COs with POs & PSOs:													
~~				· Jo P	O)	7 6			PSO		
CO	1	2	3	49	5	6	7	8 5	1	2	3	4	5
CO1	S	S	S	M	S	M	M	S	S	M	S	M	M
CO2	S	S	S	M	S	S	S	S	S	S	S	M	S
CO3	S	S	S	S	S	S	S	S	S	S	S	M	S
CO4	S	S	S	S	S	S	S	S	S	S	S	M	S
CO5	S	S	S	S	S	S	S .	Sis	S	S	S	M	S

Strongly Correlating (S) - 3 marks; Moderately Correlating (M) - 2 marks
Weakly Correlating (W) - 1 mark; No Correlation (N) - 0 mark
