NOIDA INSTITUTE OF ENGG. & TECHNOLOGY, GREATER NOIDA, GAUTAM BUDDH NAGAR (AN AUTONOMOUS INSTITUTE)



Affiliated to

DR. A.P.J. ABDUL KALAM TECHNICAL UNIVERSITY UTTAR PRADESH, LUCKNOW



Evaluation Scheme & Syllabus

For

Master of Technology Biotechnology First Year

(Effective from the Session: 2023-24)

NOIDA INSTITUTE OF ENGG. & TECHNOLOGY, GREATER NOIDA, GAUTAM BUDDH NAGAR (AN AUTONOMOUS INSTITUTE)

Master of Technology Biotechnology Evaluation Scheme SEMESTER-I

S. No.	Subject Subject		Periods		Evaluation Schemes				End Semester		Tota I	Credit	
	Codes		L	т	Ρ	СТ	ТА	TOTAL	PS	TE	PE		
1	AMTBT0101	Applied Biochemistry & Molecular Biology	3	0	0	20	10	30		70		100	3
2	AMTBT0102	Bioprocess Engineering & Technology	3	0	0	20	10	30		70		100	3
3	AMTCC0101	Research Process and Methodology	3	0	0	20	10	30		70		100	3
4		Departmental Elective-I	3	0	0	20	10	30		70		100	3
5		Departmental Elective-II	3	0	0	20	10	30		70		100	3
6	AMTBT0151	Applied Biochemistry & Molecular Biology Lab	0	0	4				20		30	50	2
7	AMTBT0152	Bioprocess Engineering & Technology Lab	0	0	4				20		30	50	2
		TOTAL										600	19

Departmental Elective-I

1.AMTBT0111 Immunology & Vaccine Technology

2.AMTBT0112 Quality Assurance and Quality Control

3.AMTBT0113 Applied Clinical Research

Departmental Elective-II

1.AMTBT0114 Biological Treatment of Wastewater

2.AMTBT0115 Nano Biotechnology & Toxicology

3.AMTBT0116 Industrial Biotechnological Products

Abbreviation Used:-

L: Lecture, T: Tutorial, P: Practical, CT: Class Test, TA: Teacher Assessment, PS: Practical Sessional, TE: Theory End Semester Exam., PE: Practical End Semester Exam.

NOIDA INSTITUTE OF ENGG. & TECHNOLOGY, GREATER NOIDA, GAUTAM BUDDH NAGAR

(AN AUTONOMOUS INSTITUTE)

Master of Technology Biotechnology Evaluation Scheme SEMESTER-II

	1st Year, I						I nd			Semester			
SI. No	Subject Codes	Subject	Periods		Eva	aluati	on Schem	n Schemes		End Semester		Credit	
110	Coues		L	Τ	Р	СТ	TA	TOTAL	PS	TE	PE		
1	AMTBT0201	Bioinformatics	3	0	0	20	10	30		70		100	3
2	AMTBT0204	Cell & Tissue Culture Techniques	3	0	0	20	10	30		70		100	3
3		Departmental Elective-III	3	0	0	20	10	30		70		100	3
4		Departmental Elective-IV	3	0	0	20	10	30		70		100	3
5		Departmental Elective-V	3	0	0	20	10	30		70		100	3
6	AMTBT0251	Bioinformatics Lab	0	0	4				20		30	50	2
7	AMTBT0254	Cell & Tissue Culture Techniques Lab	0	0	4				20		30	50	2
8	AMTBT0253	Seminar-I	0	0	2				50			50	1
		TOTAL										650	20

NOIDA INSTITUTE OF ENGG. & TECHNOLOGY, GREATER NOIDA, GAUTAM BUDDH NAGAR (AN AUTONOMOUS INSTITUTE)

Master of Technology Biotechnology

Departmental Elective-III

1.AMTBT0211 Genetic Engineering

2.AMTBT0212 Applied Food Biotechnology

3.AMTBT0213 Molecular Modelling & Industrial Application

Departmental Elective-IV

1.AMTBT0214 Bioreactor Analysis & Design

2.AMTBT0215 Enzyme Technology & Industrial Application

3.AMTBT0216 Applied Bioenergy

Departmental Elective-V

1.AMTBT0218 Diagnostic Techniques in Biotechnology

2. AMTBT0219 3-D Printing Technology

3. AMTBT0220 Entrepreneurship, IPR & Biosafety

		M. TECH FIRST YEAR				
Course	Code	AMTBT0101 L T I	P Credit			
Course	Title	Applied Biochemistry & Molecular Biology3 0 0) 3			
Course	objective:					
<u>1</u>		erstand the various concepts of molecular biology and biochemistry				
-		ine the structure and function of biomolecules and evaluate the complexity	of			
2		arious biomolecules.				
3	Underst	and the principles of bioenergetics to learn the various pathways.				
4	Evaluate	Evaluate the concept of metabolisms of various types.				
5	Evaluate	e structure of genetic material and the central dogma of molecular biology.				
Pre-req	uisites:					
		Students are expected to have knowledge of basic biology, cell biology ar biochemistry	ıd			
Course	Contents	/ Syllabus:				
Unit 1	Struc	tures and functions of Bio-molecules:	8 hr			
	acids, function	hydrates: classification, mono, di, oligo and polysaccharides. Lipids: fat simple, complex & derived lipids. Protein: Amino Acids Structure an on, Protein Structure Hierarchy. Nucleic acids: nucleosides, nucleotides, & RNA.				
Unit 2	Bioen	ergetics:	8 hr			
	Energ	iew of principles of bioenergetics (free energy, enthalpy and entropy y relationships between catabolic and anabolic pathways. Phosphoryl grou ers and ATP, Free-energy change for ATP hydrolysis.				
Unit 3	Metal	bolism:	8 hr			
	Cycle	lysis, Gluconeogenesis, Respiration and Introduction to the Citric Acid, Electron Transport, Oxidative phosphorylation, Fatty Acid Catabolis acid oxidation, Protein Metabolism: The Urea Cycle	m:			
Unit 4	-	Structure and Function	10 hr			
	DNA replica	structure, DNA & RNA as a genetic material, RNA World, packaging of as chromosome, DNA replication- Prokaryotic and eukaryotic DN ation, Mechanism of replication. Telomeres, telomerase and end replication. of telomerase in aging and cancer.	A			
Unit 5	Centr	al Dogma	10 hr			
		cription, genetic code, reverse transcription, mRNA processing. Translation regulation, operons: Lac operon, Trp operon, transposons.	on,			
Course	outcome					
CO1		After completion of the course, students will understand about the structure and function of biomolecules	ure			
CO2		They will learn about principles of bioenergetics.				
CO3		They will understand the different types of metabolisms.				
CO4		Students will learn the overall gene structure and function.				
CO5		Students will be able to understand the molecular functioning of cells.				
Text bo	1					
1		nistry- L.Stryer, Third Edition				
2		nistry- Voet&Voet.				
3	Principl	es of Biochemistry- A.Lehninger, CBS Publishers and Distributors, 1987.				

Referen	ce Books				
1	Watson. J. D, Baker. T. A, Bell. S. P, Gann. A, Levine. M, Losick. R. Molecular Biology of Gene. 6th The Benjamin / Cummings Pub. Co. Inc, 2008.				
2	2 Darnell, Lodish and Baltimore. Molecular Cell Biology, Scientific American Publishing Inc, 2000.				
3	Bruce Alberts, Alexander Johnson, Julian Lewis, Martin Raff, Keith Roberts, Peter Walter. Molecular biology of the Cell. 4th ed. Garland publishing Inc, 2002				
Journal/Research Paper Link:					
	As suggested by concern subject faculty				

		M. TECH FIRST YEAR	
Course	Code	AMTBT0102 L T P	Credit
Course	Title	Bioprocess Engineering & Technology3 0 0	3
<u> </u>			
Course o	-		
1		vide basic concepts of bioprocess engineering.	
2	enzym	rn engineering principles that can be applied to processes involving cell e.	or
3	To lear	rn the basics of bioreactor design and operation control.	
4	To ana	lyze variety of bioprocess techniques and also conduct related experiments.	
5	To und	lerstand various unit operations in bioprocess.	
Pre-requ	isites:		
		ents are expected to have knowledge of basic biology, cell biology and nemistry	
Course C	Contents	s / Syllabus :	
UNIT I	Intro	oduction to Bioprocess Technology	8 Hr.
	ferme ancill Diffe Medi comr	ern applications of biotechnological processes, General requirements entation processes, Basic design and construction of fermenter and laries, Main parameters for monitoring & control of fermentation processes erent raw materials used in fermentation industry and their pretreatmer ium for plant cell culture and animal cell culture, Medium design mercial media for industrial fermentations-Plackettburman design, respon ce methodology, simplex design.	nd s, nt, of
UNIT II	Stoic	chiometry of Cell growth	8 Hr.
	of ro coeff analy	chiometry of Cell growth and product formation, elemental balances, degree eduction of substrate and biomass, available electron balances, yie ficients of biomass and product formation, maintenance coefficients Energet visis of microbial growth and product formation, oxygen consumption and he ation in aerobic cultures, thermodynamic efficiency of growth.	ld ic
UNIT III	Mass	s Transfer in Bioreactors	8 Hr.
	oxyg oxyg proce	s transfer includes transport phenomena in bioprocesses, Factors affecting en transfer rate in bioreactors, Techniques for measurement of volumetr en transfer coefficient, Fluid rheology and factors affecting bioreactor esses, Flow Patterns in agitated tanks, Mechanism & Power requirements ong, Scale up of mixing systems.	ic or
UNIT IV	Meta	abolic Regulation	10 Hr.
	anabo catab Conc acid,	erent regulatory mechanisms involved in controlling the catabolic ar olic processes of microbes, Induction, nutritional repression, carbo polite repression, Crabtree effect, feedback inhibition and feedback repression cept of Overproduction of metabolites, Case studies on production of Lact Glutamic acid, Penicillin, Microbial Lipase and Protease, Recombina in, Interferons, Hepatitis Vaccines etc. Case studies should deal with stra	n, ic nt
	impro	ovement, medium designs, process optimization technology.	

	Unit Operation: Filtration, filter aids, filtration Equipment and filtration theory, Centrifugation process and its equipments, Cell disruption, Aqueous Two-Phase	
	Liquid Extraction. Adsorption process and its operations, Chromatography: Theory and mechanism, Scaling-up chromatography.	
Course	outcome:	
CO1	Describe the underlying principles of main bioprocess unit operations like fermentation, downstream processing.	
CO2	Demonstrate Stoichiometry of Cell growth and product formation.	
CO3	Design or Select appropriate bioreactor models based upon bioproducts and cell lines and other process criteria.	
CO4	Develop a basic understanding of regulatory mechanisms involved in controlling the catabolic and anabolic processes of microbes.	
CO5	Acquire a basic understanding of various unit operations in bioprocess engineering.	
Text bo	oks	
1	Principles of fermentation technology" by P F Stanbury and A Whitaker, Pergamon press.	
2	Bioprocess Technology - Kinetics & Reactors" by A Moser, Springer- Verlag.	
3	Biochemical Engineering and Biotechnology Handbook" by B. Atkinson & F. Mavituna, 2nd Ed. Stockton Press.	
4	Bioprocess Engineering Principles" by Pauline M. Doran, Academic Press.	
5	Biochemical Engineering- S. Aiba , A.E. Humphray, University of Tokyo Press.	
Referen	ce Books	
1	Lee J.M, Biochemical Engineering 2nd ed, Prentice Hall, 2000.	
2	Principles of Cell Energetics": BIOTOL series, Butterworth - Heinemann.	
3	Biotechnology" Vol.4 Meaning Modelling and Control Ed. K.Schugerl, VCH (1991).	
4	Unit operations of Chemical Engineering" 5th ed. by W L McCabe, J C Smith and P. Harriot Mc Graw-Hill (1993).	
5	Diffusion" by E L Cussler, Cambridge University Press (1984).	
6	Bioprocess Engineering Principles" by Pauline M.Doran, Academic Press.	
Journal	/Research Paper Link:	
	As suggested by concern subject faculty	

Course Code	AMTCC0101	L T P	Credit
Course Title	Research Process & Methodology	300	3
Course Object			
1	To explain the concept / fundamentals of research and their ty		
2	To study the methods of research design and steps of n process	research	
3	To explain the methods of data collection and procedure of sate techniques	ampling	
4	To analyze the data, apply the statistical techniques and und the concept of hypothesis testing	derstand	
5	To study the types of research report and technical writing.		
Pre-requisites	Basics of Statistics		
	Course Contents / Syllabus		
UNIT-I	INTRODUCTION TO RESEARCH		8 hours
	tive and motivation of research, types and approaches of rese	earch. Des	
	lied vs. Fundamental, Quantitative vs. Qualitative, Concer		
	s versus Methodology, significance of research, criteria of good		
UNIT-II	RESEARCH FORMULATION AND DESIGN		8 hours
objective of Liter	s and steps involved, Definition and necessity of research prob rature review, Locating relevant literature, Reliability of a sour the research problem, Literature Survey, Research Design, 1	rce, Writi	ng a survey
design.	DATA COLLECTION		8 hours
design. UNIT-III		ection, C	8 hours
design. UNIT-III Classification of primary and seco	DATA COLLECTION Data, accepts of method validation, Methods of Data Collected ondary data, sampling, need of sampling, sampling theory and different types of sample designs, ethical considerations in rese	Techniqu	ollection of
design. UNIT-III Classification of primary and seco	Data, accepts of method validation, Methods of Data Colle ondary data, sampling, need of sampling, sampling theory and	Techniqu	ollection of
design. UNIT-III Classification of primary and seco sampling design, UNIT-IV Processing Opera appropriate statis statistical inferent Visualization – M	 Data, accepts of method validation, Methods of Data Collected ondary data, sampling, need of sampling, sampling theory and different types of sample designs, ethical considerations in rese DATA ANALYSIS ations, Data analysis, Types of analysis, Statistical technique stical technique, Hypothesis Testing, Data processing software nee, Chi-Square Test, Analysis of variance(ANOVA) an Monitoring Research Experiments, hands-on with LaTeX. 	Techniquearch.	ellection of tes, steps in 8 hours shoosing an SPSS etc.), ance, Data
design. UNIT-III Classification of primary and seco sampling design, UNIT-IV Processing Opera appropriate statis statistical inferent Visualization – M UNIT-V	 Data, accepts of method validation, Methods of Data Collected ondary data, sampling, need of sampling, sampling theory and different types of sample designs, ethical considerations in rese DATA ANALYSIS ations, Data analysis, Types of analysis, Statistical technique stical technique, Hypothesis Testing, Data processing software nee, Chi-Square Test, Analysis of variance(ANOVA) an Ionitoring Research Experiments, hands-on with LaTeX. TECHNICAL WRITING AND REPORTING OF RESEA 	Techniquearch. ues and care (e.g. ud covari	ollection of ies, steps in 8 hours thoosing an SPSS etc.), ance, Data 8 hours
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Text books

- **1.** C. R. Kothari, Gaurav Garg, Research Methodology Methods and Techniques, New Age International publishers, Third Edition.
- Ranjit Kumar, Research Methodology: A Step-by-Step Guide for Beginners, 2nd Edition, SAGE 2005.
- 3. Deepak Chawla, NeenaSondhi, Research Methodology, Vikas Publication

Reference Books

- 1. Donald Cooper & Pamela Schindler, Business Research Methods, TMGH, 9th edition
- **2.** Creswell, John W.,Research design: Qualitative, quantitative, and mixed methods approaches sage publications,2013

	M. TECH FIRST YEAR	
Course Code	AMTBT0151 L T P	Credit
Course Title	Applied Biochemistry & Molecular Biology Lab0 04	2
Course objectiv	e:	
1	To understand the various concepts of molecular biology and biochemistry	
2	Determine the structure and function of biomolecules and evaluate the complexity of various biomolecules.	
3	Understand the principles of bioenergetics to learn the various pathways.	
4	Evaluate the concept of metabolisms of various types.	
5	Evaluate structure of genetic material and the central dogma of molecular biology.	
Pre-requisites:		
	Students are expected to have knowledge of basic biology, cell biology and biochemistry	
Suggested list of	f Experiment :	
Sr. No.	Name of Experiment	СО
1	Quantitative estimation of amino acids by ninhydrin reaction.	12
2	Quantitative estimation of proteins.	12
3	To separate lipids with the help of thin layer chromatography (TLC).	12
4	To verify the Lambert Beer's law with the help of UV absorption spectra of proteins.	12
5	Protein purification by ammonium sulfate precipitation.	12
6	Isolation of DNA and RNA from animal tissue and plant tissue.	12
7	Gel electrophoretic analysis of various DNA and their restriction digests	1
8	Transformation with plasmid and bacteriophage DNA	13
9	Restriction mapping of plasmid DNA	3
10	Blotting: northern blotting, southern blotting	3
11	PCR technique	3
Lab Course Ou	tcome:	
CO 1	Students will be able to understand the various biomolecules.	
CO 2	Students will learn through demonstration the process of isolation and analysis of different biomolecules.	
CO 3	They will learn about the structure and function of DNA, RNA and Protein.	
CO 4	Students will learn advanced molecular methods.	

	M. TECH FIRST YEAR		
Course Code	AMTBT0152	L T P	Credit
Course Title	Bioprocess Engineering & Technology Lab	0 0 4	2
Course objectiv	e:		
1	To understand the various concepts of microbial culturing.		
2	To learn the activation energy, volumetric oxygen transfer coefficient	ient etc.l	
3	To Understand the principles and various pathways of enzyme pro-	duction.	
4	Evaluate the concept of separation and purification of microbial pu	roduce.	
5	To understand the process of fermentation.		
Pre-requisites:			
	Students are expected to have knowledge of basic biology, cell bio biochemistry	ology and	
Suggested list of	f Experiment :		
Sr. No.	Name of Experiment		
1	Determination of kinetic parameters for batch cultivation of yeast flask conditions.	under shake	
2	Determination of volumetric oxygen transfer coefficient (KLa)		
3	Determination of activation energy (Ea) of microbial strains.		
4	Process optimization for enzyme production using specific experindesign.	mental	
5	Preparation of immobilized enzymes & cells and evaluation of kin parameters.	ietic	
6	Computational Design of Fermentative Process.		
7	Fermenter designing and the study of various parts of ferment function for microbial cell culture.	er and their	
8	Fermentative production of Penicillin by using Peniciliumchrysog	enum.	
9	Microbial production of enzymes Cellulase & Protease.		
10	Ethanol production from molasses or starchy raw material.		
11	Fermentative production of Wine from grapes.		
12	Separation and purification of microorganisms from yogurt and ch	neese.	
13	Fermentative production of alpha amylase under solid & submerge conditions	ed	
14	Protein profiling of fermentation broth through dialysis procedure		
15	To study the Scale-up and Sterilization in Bioreactors		
Lab Course Ou	tcome:		
CO 1	Student will be able to understand the various concepts of microbi	al culturing.	
CO 2	Student will learn the activation energy, volumetric oxygen transfer coefficient etc.	er	
CO 3	Student will Understand the principles and various pathways of enproduction.	zyme	
CO 4	Student will be able to evaluate the concept of separation and pu microbial produce.	rification of	
CO 5	Student will be able to understand the process of fermentation.		

Course Title Immunology & Vaccine Technology 3 0 0 3 Course title Immunology & Vaccine Technology 3 0 0 3 Course objective: I Icarn the concept and components of the Immune system. 1 2 Understand the kinetics and mechanisms of immune response. 3 1 3 Evaluate the concept of vacious vaccine types viz. viral vaccines, bacterial vaccines and parasitic vaccines etc. 1 4 Understand the vaccine industry and the safety and legal issues related to its production. 1 Pre-requisites: Students are expected to have knowledge of basic Cell and Molecular biology, knowledge of the various diseases and causative agents will be an edge. 8 hr Course Contents / Syllabus : UNIT-1 Fundamental of Immune System 8 hr Immunological Processes 8 hr 1 1 Indamental concepts and anatomy of the immune system. Components of immune oplobulins, antigens intrunone genose, memory; B cell maturation, activation and differentiation. Generation of antibody diversity. Antigen processing and presentationendogenous antigens and exogenous antigens. 8 hr UNIT-1 Immunological Processes 8 hr Immunolobulini, sinteites of immune responsy. Beelin maturation, activation and differentiation. Generation		M. TECH FIRST YEAR	
Course objective: Image: Constraint of the concept and components of the Immune system. 2 Understand the kinetics and mechanisms of immune response. 3 Evaluate the concept of vaccination and various types of vaccines. 4 Understand the concept of various vaccine types viz. viral vaccines, bacterial vaccines and parasitic vaccines etc. 5 Understand the vaccine industry and the safety and legal issues related to its production. Pre-requisites: Students are expected to have knowledge of basic Cell and Molecular biology, knowledge of the various diseases and causative agents will be an edge. Course Contents / Syllabus : UNIT-I Fundamental concepts and anatomy of the immune system, Components of imnate and acquired immunity, Humoral and Cell mediated immunity, Ucomplex - MHC genes, MHC and immune responsiveness and disease susceptibility, HLA typing. 8 hr UNIT-II Immunological Processes 8 hr Immunolobulins, antigenic determinants, Multigene organization of immunoglobuling sense, Immunological basis of self -non-self discrimination; Kinetics of immune response, memory; B cell maturation, activation and differentiation; Generation of antibody diversity, Antigene processing and presentationendogenous antigens and exogenous antigens. UNIT-III Basic Introduction to Vaccines; 8 hr A short history of vaccination of immunocompromised hosts, Vaccines, vaccines, vaccines, vaccines; vaccines, vaccines; vaccines, plan	Course Code	AMTBT0111 L T P	Credit
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After completion of the course, students will understand the fundamentals of		across the centuries, Vaccine additives and manufacturing residuals, Regulation and testing of vaccines, Regulation of vaccines in developing countries, Vaccine safety and Legal issues.	
-	Course outcome		
	CO 1	-	

CO 2	They will learn about immunological processes.	
CO 3	They will understand the different types of immunization and vaccines.	
CO 4	Students will learn the different types of advanced vaccines.	
CO 5	Students will be able to understand the vaccine industry and their production process.	
Text books		
1	Kuby, RA Goldsby, Thomas J. Kindt, Barbara, A. Osborne Immunology, 6th Edition, Freeman, 2002.	
2	Brostoff J, Seaddin JK, Male D, Roitt IM., Clinical Immunology, 6th Edition, Gower Medical Publishing, 2002.	
3	Janeway et al., Immunobiology, 4th Edition, Current Biology publications., 1999. 4. Paul,	
4	Fundamental of Immunology, 4th edition, Lippencott Raven, 1999.	
Reference Books	S S	
1	Stanley A. Plotkin & Walter Orenstein & Paul A. Offit, Vaccines, 6th Edition 2013 BMA Medical Book Awards Highly Commended in Public Health! Elsevier Publication.	
2	Roitt's Essential Immunology. 11th ed. P. Delves, et al., ed., Blackwell Publishing, 2006.	
3	Bruce Alberts, Alexander Johnson, Julian Lewis, Martin Raff, Keith Roberts, Peter Walter. Molecular biology of the Cell. 4th ed. Garland publishing Inc, 2002	
Journal/Researc	h Paper Link:	
	As suggested by concern subject faculty	

		M. TECH FIRST YEAR		
Cour	se Code	AMTBT0112	L T P	Credit
Cour	se Title	Quality Assurance and Quality Control	300	3
Cours	se objective	:		
1	To learn t	ne basics of GLP		
2	To learn t	ne manufacturing process and its audit.		
3		tand the clinical trial process		
4		he statistical tools to the various QC events		
5		tand the tools and softwares used in QC and QA.		
Pre-re Stude	equisites:	e expected to have knowledge of basic biology, cell biology and bio	ochemistry	
Cours	se Contents	/ Syllabus :		
UNIT		cept of Quality control and quality assurance		
	Man resp data	cept and evolution of quality control and quality assurance. Total Q agement, Philosophy of GMP and CGMP. Quality control laborato onsibilities: GLP protocols on nonclinical testing control on animal generation, integration and storage, standard test procedure, CPCS elines.	ry house,	
UNIT		umentation practices and root cause analysis		
	finis	ntion of sample records, Quality review and batch release documented products, Good documentation practices, route cause analysis, on preventive action (CAPA), out of specifications (OOS) and out of T)	corrective	
UNIT	III Con	cept of Audits		
	audi qual	ual product quality review and parametric release, Audits,Preparati t, conducting audit, Audit Analysis, Audit Report and Audit follow ity audits of manufacturing processes and facilities, audits of qualit e Studies of Audit reports.	up,	
UNIT	IV Qua	lity agreements and risk management		
	qual	cepts and management of contract manufacturing guidelines,principity risk management, ICH guidance for industry, BABE (bioavailabquivalence) studies, post marketing surveillance, Pharmacovigilance	bility and	
UNIT	V Too	s and softwares in QC and QA		
	Con appl solu	stical Tools for Quality Control and Precision, Tools of Problem Section tinuous Improvement. Softwares for inspection and quality testing a ications. concept of automation of procedure through Digital, IoT tions. Systematic approach to scale-up and technology transfer in echnology quality systems: Applications and challenges.	and their	
Cours	se outcome			
CO1		ognize the importance of quality control and assurance and understate of GMP, CGMP and GLP.	and the	
CO2	prev	ognize the importance of good documentation practices and reframe entive actions.		
CO3		yse, develop, follow and audit the quality standards and guidelines wed in a biotechnology industry.	, being	
CO4	Und	erstand the contract guidelines to effectively manage the quality ag	reements.	

	Apply statistical tools and modern software to evaluate and ensure quality control,	
CO5	assurance and precision.	
Text boo	oks	
1	Sharp J. Good Pharmaceutical Manufacturing Practice: Rationale and Compliance. CRC Press; 2005.	
2	Gad SC. Pharmaceutical Manufacturing Handbook: Production and Processes. John Wiley & Sons; 2008.	
3	Steinborn L. GMP/ISO Quality Audit Manual for Healthcare Manufacturers and Their Suppliers, Sixth Edition, (Volume 1 - With Checklists and Software Package). Taylor & Francis; 2003.	
4	Kolman J, Meng P, Scott G. Good Clinical Practice: Standard OperatingProcedures for Clinical Researchers. Wiley; 1998.	
5	Waller P. An Introduction to Pharmacovigilance. John Wiley & Sons; 2011.	
Reference	ce Books	
1	Niazi S. Handbook of Bioequivalence Testing. CRC Press; 2007.	-
2	Chalmers AA. International Pharmaceutical Registration. Interpharm Press; 2000.	
3	Edwards AJ. ISO 14001 Environmental Certification Step- by-Steps: Revised Edition. Butterworth-Heinemann; 2003.	
4	Mantus D. FDA Regulatory Affairs: A Guide for Prescription Drugs, Medical Devices, and Biologics. Informa Healthcare USA; 2008.	
5	Chalmers AA. International Pharmaceutical Registration. Interpharm Press; 2000.	
6	Contract manufacturing arrangement for drugs, quality agreements: guidance for industry,November 2016.	
Journal/	Research Paper Link:	
	As suggested by concern subject faculty	

Course Title Applied Clinical Research 3 00 3 Course objective:		M. TECH FIRST YEAR		
Course objective: Image: Course objective: 1 To learn the basic of drug development process 1 2 To learn the basic step involve in clinical trial of drug. 1 3 To understand the crinciples of controlled clinical trials 1 5 To apply the statistical tool for data management. Pre-requisites: 8 To curse Contents / Syllabus : 1 Course Contents / Syllabus : 8 Hr. UNIT I: Introduction to clinical research 8 Hr. Basic pharmacology and drug development process, clinical research/definition, Basic terminology used in clinical research, preclinical studies, Introduction to pharmacoconomics, Types of clinical trials, single blinding, double blinding, open access, randomized trials and their examples, interventional study, Good Clinical Practices, Types and Scope of Clinical Research. 8 UNIT II: Clinical trials 8 Hr. New drug discovery process- purpose, main steps involved in new drug discovery process, timelines of cach steps, advantages and purposes of cach steps, Per clinical toxicology: General principles, Systemic toxicology (Single dose and repeat dose toxicity studies), Carcinogonicity, Matagenicity, Teratogenicity, Reproductive toxicity, Local toxicity, Genotoxicity, animal toxicity requirements, Phase-1, II, II, Vi trials: Introduction and designing, Various phases of clinical trials, Post Marketing survicu	Course Code	AMTBT0113 L T	' P	Credit
1 To learn the basic of drug development process 2 To learn the basic step involve in clinical trial of drug. 3 To understand the chics involved in clinical research 4 To understand the principles of controlled clinical trials 5 To apply the statistical tool for data management. Pre-requisites: Students are expected to have knowledge of basic biology, cell biology and biochemistry Course Contents / Syllabus : 8 Hr. Basic pharmacology and drug development process, clinical research/effinition, Basic terminology used in clinical research, preclinical studies, Introduction to pharmacoeconomics, Types of clinical trials, single blinding, double blinding, open access, randomized trials and their examples, interventional study, Good Clinical Practices, Types and Scope of Clinical Research. UNIT II: Clinical trials 8 Hr. New drug discovery process- purpose, main steps involved in new drug discovery process- timelines of each steps, advantages and purposes of each steps, Pre clinical toxicology: General principles, Systemic toxicology (Single dose and repeat dose toxicity studies), Carcinogenicity, Mutagenicity, Teratogenicity, Reproductive toxicity, Local toxicity, Genotoxicity, animal toxicity requirements, Phase-I, II, III, Vi trials: Introduction and designing, Various phases of clinical trials. Post Marketing surveillance, methods & Principles of sampling, Inclusion and exclusion criteria, Methods of allocation and randomization, Informed consent process in brief monitoring, treatment outcome, Termination of trial, Safety monitoring in clinical trials	Course Title	Applied Clinical Research3 0	0	3
1 To learn the basic of drug development process 2 To learn the basic step involve in clinical trial of drug. 3 To understand the chics involved in clinical research 4 To understand the principles of controlled clinical trials 5 To apply the statistical tool for data management. Pre-requisites: Students are expected to have knowledge of basic biology, cell biology and biochemistry Course Contents / Syllabus : 8 Hr. Basic pharmacology and drug development process, clinical research/effinition, Basic terminology used in clinical research, preclinical studies, Introduction to pharmacoeconomics, Types of clinical trials, single blinding, double blinding, open access, randomized trials and their examples, interventional study, Good Clinical Practices, Types and Scope of Clinical Research. UNIT II: Clinical trials 8 Hr. New drug discovery process- purpose, main steps involved in new drug discovery process- timelines of each steps, advantages and purposes of each steps, Pre clinical toxicology: General principles, Systemic toxicology (Single dose and repeat dose toxicity studies), Carcinogenicity, Mutagenicity, Teratogenicity, Reproductive toxicity, Local toxicity, Genotoxicity, animal toxicity requirements, Phase-I, II, III, Vi trials: Introduction and designing, Various phases of clinical trials. Post Marketing surveillance, methods & Principles of sampling, Inclusion and exclusion criteria, Methods of allocation and randomization, Informed consent process in brief monitoring, treatment outcome, Termination of trial, Safety monitoring in clinical trials	Course objectiv	e:		
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5 To apply the statistical tool for data management. Pre-requisites: Students are expected to have knowledge of basic biology, cell biology and biochemistry Course Contents / Syllabus : 8 Hr. Basic pharmacology and drug development process, clinical research. 8 Hr. Basic pharmacology and drug development process, clinical studies, Introduction to pharmacoeconomics, Types of clinical trials, single blinding, double blinding, open access, randomized trials and their examples, interventional study, Good Clinical Practices, Types and Scope of Clinical Research. UNIT II: Clinical trials 8 Hr. New drug discovery process- purpose, main steps involved in new drug discovery process, timelines of each steps, advantages and purposes of each steps, Pre clinical toxicology: General principles, Systemic toxicology (Single dose and repeat dose toxicity, studies), Carcinogenicity, Mutagenicity, Teratogenicity, Reproductive toxicity, Local toxicity, Genotoxicity, animal toxicity requirements, Phase-I, II, III, IV trials: Introduction and designing, Various phases of clinical trials, Post Marketing surveillance, methods & Principles of sampling, Inclusion and exclusion criteria, Methods of allocation and randomization, Informed consent process in brief monitoring, treatment outcome, Termination of trial, Safety monitoring in clinical trials, thalidomide tragedy, Conflicts of Interest, Evolution and History of Regulations in Clinical Research, Study of various clinical trials, completed or ongoing),Patents US Regulatory Structure, Clinical Trial Application in India Import & Export of Drug in India, Investigational New Drug Application (NDA), Abbreviated New Drug Appliciation (NDA), Abot Strug Application and F	4	To understand the principles of controlled clinical trials		
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	UNIT IV:			10 II

	Clinical trial design (observational and interventional) protocol, consent in clinical trials, placebo, bias and methods to prevent bias, monitoring. Multicentre clinical trials, Requirements, regulations and feasibility, Designing of Protocol, CRF, eCRF, IB, ICF, SOP BA/BE Studies Report writing, Publication, Improving patient enrolment and retention in Clinical Trials. ADR monitoring,Pharmacovigilance Training in clinical research.	
UNIT V	Biostatistics and data management	10 Hr.
	 Preparation of a successful clinical study, Study management, Project management Documentation, Monitoring, Audits and Inspections. Budgeting in clinical research, Supplies and vendor management. Importance of statistics in clinical research Statistical considerations at the design, analysis and reporting stage. Data management, Data validation, SAE reconciliation, query management Software considerations. Clinical Trial studies: Cancers and Other Neoplasms , Behaviours and mental Disorders, Immune System studies, Urinary Tract, Sexual Organs and pregnancy condition. 	
Course	outcome	
CO1	Describe the process of drug development and principles of clinical pharmacology.	
CO2	Develop a clear understanding of why ethics are important in clinical research and be familiar with the regulatory practices in place to protect both the researcher and the subject	
CO3	Effectively manage the regulatory process from Innovation \rightarrow Discovery \rightarrow Approval \rightarrow Commercialization to bring the product to the market globally.	
CO4	Communicate ideas and data in writing, including of scientific concepts and research design of clinical trials	
CO5	Describe the various types of clinical studies and the methods used to choose the appropriate design, evaluation and interpretation of clinical trial results.	
Text bo	oks	
1	Basic and Clinical Pharmacology, Prentice hall, International, Katzung, B.G.	
2	Clinical Pharmacology, Scientific book agency, Laurence, DR and Bennet PN.	
3	Clinical pharmacokinetics, Pub. Springer Verlab, Dr. D.R Krishna, V. Klotz	
4	Remington Pharmaceutical Sciences, Lippincott, Williams and Wilkins	
5	Drug interaction, Kven Stockley. Hamsten	
1	Clinical pharmacology and drug therapy Grahame smith and Aronson,	
2	Text Book of Therapeutics Drug and Disease Management Hardbound. Richard A Helms,	
3	Clinical Pharmacy and therapeutics Herfindal E T and Hirschman JL, Williams and Wilkins,	
4	Methodology of Clinical Drug Trials, 2nd Edition. Spriet A., Dupin-Spriet T., Simon P. Publisher: Karger.	
Journal	/Research Paper Link:	
	As suggested by concern subject faculty	

Course	Code	AMTBT0114 L T	P Credit
Course	Title	Biological Treatment of Waste Water 3 0	0 3
Course o	bjective:		
1	To learn	about the mass balance involved in waste water treatment	
2	To under	stand the anaerobic treatment process.	
3		about the various chemical and physical processes involved in waste wat	er
	treatment		
4		stand the basic of phosphorus and nitrogen removal	
5		about the recycling of waste	
Pre-requ			
	Students biochemi	are expected to have knowledge of basic biology, cell biology and	
Cauraa		•	
Course		Syllabus: IVATED SLUDGE PROCESS-PROCESS ANALYSIS A	ND a
UNIT I-	SELI	ECTION	8 Hr.
		acteristics of Activated Sludge (aerobic and anaerobic); Analysis of Data	-
		Balance Analysis. Reactors used in waste water treatment- Up Flow	
		robic Sludge Blanket (UASB), Two-stage, Aerobic UNI Tank System System, Route Zone Treatment, Submerged Aerobic Fixed Film (SAFF)	
		tor, and Fluidized Aerobic Bioreactor (FAB).	
UNIT II-		OBIC FIXED-FILM & ANAEROBIC TREATMENT PROCESSES	8 Hr.
011111		Im process considerations; Trickling Filters and Biological Towers;	
		ting Biological Contactors; Granular – Media Filters; Fluidized – Bed &	
		llating Bed- Biofilm reactors. Hybrid Biofilm/suspended growth processe	es.
	Anae	robic Processes: Methanogenesis, process chemistry and microbiology;	
		ess kinetics and factors for the design of anaerobic digestors.	
UNIT III	I- ADV	ANCED WASTE WATER TREATMENT	8 Hr.
	Tech	nologies used in advanced treatment-Classification of technologies;	
		oval of Colloids and suspended particles-Depth Filtration, Surface	
		tion, Membrane Filtration Absorption, Ion Exchange, Advanced oxidation	on 🛛
	1 4	ess, Activated Carbon, Air Stripping, Heavy Metals Removal, Steam	
		ping, Chemical Precipitation, and Electrolysis.	
UNIT IV		LOGICAL PHOSPHORUS AND NITROGEN REMOVAL	10 Hr.
		fication & Denitrification Processes: Biochemistry and Physiology of	
		fying Bacteria; Common process considerations; One sludge versus two	
	U U	e nitrification. Physiology of Denitrifying Bacteria; Tertiary	
		trification; One- sludge denitrification, Normal Phosphorus Uptake into ass; Mechanism for Biological Phosphorus Removal; Enhanced Biologic	vo1
		phorus Removal by Bacteria and Algae.	
UNIT V	-	IRONMENTAL CONCERNS & RECYCLING OF WASTES	10 Hr.
		conmental regulations and technology- Regulatory Concerns, Technology	
		, regulations and permits, Air, Water, Solid Waste, Environmental	,
		ting, National Environmental Policy act, Occupational Safety and Health	
		OSHA), Storm Water Regulations; Technology (waste water); Recycling	
	Indus	strial wastes: paper, plastics, leather and chemicals.	
Course o			
CO1		er completing the course students will able to perform mass balance for t reactor	he
COI	D101		

CO3	After completing the course students will able to categorize various chemical and physical processes involved in waste water treatment.
CO4	After completing the course students will able to describe the basic of phosphorus and nitrogen removal
CO5	After completing the course students will able to perform recycling of waste
Text books	
1	Wastewater Engineering: Treatment Disposal Reuse by Metcalf & Eddy
2	Environmental Biotechnology : Principles and Applications by Bruce E. Rittmann
3	Waste water Engineering Treatment and Reuse: McGraw Hill, G. Tchobanoglous, FIBiston, 2002.
4	Industrial Waste Water Management Treatment and Disposal by Waste Water McGraw Hill III Edition 2008.
5	Biological Wastewater Treatment", Second Edition, Marcel Dekker, Inc., New York,
Reference Bo	ooks
1	Introduction to Waste Water Treatment- R. S. Ramalho, Academic Press.
2	Environmental Biotechnology, B.C. Bhattacharya &Ritu Banerjee, Oxford Press, 2007.
Journal/Rese	arch Paper Link:
	As suggested by concern subject faculty

~		M. TECH FIRST YEAR	• •• =	~ -
Course (AMTBT0115	L T P	Credit
Course [Fitle	Nano Biotechnology & Toxicology	3 0 0	3
Course of	ojective:			
1		To understand the fundamentals concepts of nanotechnology		
2		To learn about the different types of nanoparticles		
3		To understand the principle behind the different characteristic techniques involved in nanotechnology	acterization	
4		To understand the applications of nanotechnology		
5		To learn the toxicology of nanomaterials		
Pre-requi	sites:			
		Students are expected to have knowledge of basic biology, cell and biotech	biology	
Course C				
UNIT-I		duction to Nanobiotechnology:		8hr.
	Botto Curre	ition of Nanobiotechnology, History, Origin, Fundamental m-up versus Top-down approaches, Discussion on Nanof nt research, Tool and Techniques, Applications and Implication fabrication.	abrication,	
UNIT-II	Nano	materials and Nanoparticles:		8hr.
UNIT-III	Silver biome Nano UV-v	embrane and genes. charecterization Tool and Techniques: isible spectrophotometry, Fourier transform infrared spectrosco	py (FTIR),	8hr.
	Scanr (STM	ning Electron Microscopy (SEM), Scanning tunnelling n I), Transmission electron microscopy (TEM), Atomic force n I), Zeta Potential size analyser etc.	nicroscopy	
UNIT-IV	```	medicine and Sensor Technology:		10 hr.
	Drug	delivery tools, Bioavailability, Nano imaging agents, Protein a ery (Cancer and Surgery) and Nano sensors technology with app		
UNIT-V	Toxic	cology:		10 hr.
	Toxic toxici mech	ition of toxicology, History and origin of toxicology, Pri cology, Concept of Toxicology, Types of toxicology, Na ty evaluation mechanism as in vitro, Nanomaterial toxicity anism as in vivo, Assessment of nanoparticles toxicity: A toxicity, Genotoxicity, Hepatotoxicity, Neurotoxicity, Nep	nomaterial evaluation case study	
Course ou				
CO1	concepts	npleting this course, the students will be able to learn the fu of nanotechnology		
CO2	and differ	npleting this course, the students will be able to ability for under rentiate the various nano materials		
CO3	After completing this course, the students will be able to understand the principal behind the different characterization techniques involved in nanotechnology			
CO4		npleting this course, the students will be able to get insight the chnology in drug delivery system	application	

CO5	After completing this course, the students will be able to evaluate the toxicology of nanomaterials	
Text boo	ks	
1	Nanomedicine: Biocompatibility- Robert A. Freitas; Landes Biosciences	
2	The Nanobiotechnology Hand Book- YobingXie, CRC Press.2012	
3	Nanobiotechnology: Christof M. Niemeyer, Chad A. Mirkin, John Wiley & Sons, 2004	
Reference	ee Books	
1	Nancy A. Monteiro-Riviere, C. Lang Tran., 'Nanotoxicology: Characterization, Dosing and Health Effects', Informa Healthcare publishers, 2007.	
2	P. Houdy, M. Lahmani, F. Marano, 'Nanoethics and Nanotoxicology', SpringerVerlag Berlin Heidelberg 2011.	
Journal/	Research Paper Link:	
	As suggested by concern subject faculty	

		M. TECH FIRST YEAR		
Course	Code	AMTBT0116	L T P	Credit
Course	Title	Industrial Biotechnological Products	300	3
Course o	hioctive	•		
1	-	• n about the different media for the growth of microbes		
2		erstand the production process of Primary and Secondary meta	abolites	
3		ign and deliver useful modern biotechnology products to the S		
4	Unders	tand the methods to obtain enzymes of industrial importance product development Research &Development	-	
5		erstand the manufacturing of various organic and alcoholic pro-	oducts	
Pre-requ				
	Stude	ents are expected to have knowledge of basic biology, miggy and biochemistry	crobiology, cell	
Course C	Contents	/ Syllabus :		
Unit I	Fund	amentals of Fermentation		8 hr.
	Indus	rent types of culture media; Substrates for industrial micro trially important micro-organisms: Isolation, screening, Select ess optimization techniques.		
Unit II	Prod	uction of Metabolites		8 hr.
	wine produ	ol, citric acid, vinegar and amino acid; Production of alcoh and beer; Secondary metabolites: Antibiotics; Process tech action of microbial biomass.		
Unit III	Biop	roducts		8 hr.
	Produ	duction and production of secondary metabolites with sor action of bioplastics (PHB, PHA), bioinsecticides, alymers, Biofertilizers and biological weapons with reference t	bioherbicides,	
Unit-IV	Prod	uction of industrially important enzymes		8 hr.
	ferme Prote	action of industrially important enzymes: Solid state fermentat entation, Extraction, Purification and characterization of indu ases, Cellulase, Lipase, Amylase and Pectinase, industrial mes for production of drugs and fine chemicals, Enzyme based	strial enzymes: process using	
Unit V	Prod	uction of Fermented Food Products		8 hr.
	comm prepa bever	nological processes for industrial manufacture of select nercial importance from plants and animal sources. Proce ration of Yoghurt, acidophilus milk, Koumis, kefir, cheese, b rage, vinegar and oriental fermented food. Food packagin ved in the commercially important food processing methods.	ess involved in pread, alcoholic	
Course o				
CO1		p key practical skills in fermenting biotechnology and be ons and commercial opportunities in fermentation-based biote		
CO2		e their understanding that 'industrial biotechnology' is b es to control the growth of microorganisms	based on using	
CO3	Develo	p knowledge of a variety of fermentation strategies		
CO4	Analys	e potential business opportunities in fermentation-based biotec	chnology	

CO5	Explore the biological and technological principles which govern actual and potential bio-business	
Text bo	oks	
1	Industrial Microbiology, Casida Jr. L. E. 1968) new Age International (P) Ltd. New Delhi.	
2	Presott& Dunn's Industrial Microbiology. Ed. E.G. Reed (1987). CBS Publishers, New Delhi.	
3	Biotechnology: A Text book of Induxctrial Microbiology 2 nd Edition. Crueger, W. and Cruger, A. (2000) Panima Publishing Corporation, New Delhi.	
Referen	ce Books	
1	Enzmes: Biochemistry, Biotechnology, Clinical Chemistry, Palmer, T. (2000) Horwood Publishing Colphon.	
2	Manual of Industrial Microbiology and Biotechnology 2nd Edition. Ed. Arnold L. Demain and Julian E. Davies (1999) ASM Press Washington D.C.	
3	Microbiology, Pelzar Jr. M.J.: Chan E.C.S. and Krieg, N. R. (1993) Tata Mc Graw Hill, New Delhi.	
Journa	/Research Paper Link:	
	As suggested by concern subject faculty	

	M. TECH FIRST YEAR		
Course C	ode AMTBT0201	LTP	Credit
Course T	itle Bioinformatics	3 0 0	3
Course obj			
1	To learn the various online databases		
2	To learn the online tools for analysing various methods of alignment	f sequence	
3	To understand the phylogenetic analysis and related conc	lusions	
4	To understand the concepts of system biology.		
5	To understand the various methods of genome sequencing	Ŭ.	
Pre-requisi	ites:		
1	Students are expected to have knowledge of basic biology biochemistry	, cell biology and	
Course Co	ntents / Syllabus		
UNIT I	Biological Databases		8 Hr.
	databases towards informatics projects. Primary and Seco GenBank, DDBJ, EMBL, PIR, Uniprot-KB, SWISS-I Specialized databases: Pubmed, OMIM, Medical database databases; Genome databases at NCBI, EBI, TIGR, SANC other popular tools for various bioinformatics exercises.	PROT, TrEMBL. ses, KEGG, EST	
UNIT II	Sequence Alignment		8 Hr.
	Introduction to sequence alignment, Optimal Alignment Met scores, substitution matrices, PAM, BLOSUM, Gap pe significance of Alignments, Pair wise sequence alignment alg Aspect of Multiple Sequence Alignment, Progressive and It Methods, CLUSTALW, Database similarity searching, LowComplexity Regions. PSIBLAST, PHI-BLAST.	nalties, Statistical gorithms, Practical terative Alignment	
UNIT III	Phylogenetic Analysis and Primer Design		8 Hr.
	Introduction to Phylogenetic analysis, Elements of phylogenetic Data Analysis, Phylogenetic Tree -construction Methods, Case studies related to phylogenetic analysis, Re Utilities, DNA strider, MacVector and OMIGA, gene constru NTI, Web based tools (MAP, REBASE); Primer design programs and software (PRIME3)	steps and Building striction mapping, action KIT, Vector	
UNIT IV	System Biology		8 Hr.
	Introduction to system biology, application related t introduction to different data types: PRIDE (Protein Identific Post translational modification, P-P interaction, Rotameric Str (Conformational Flexibility), Canonical DNA Forms (DNA Se	cations) databases, uctures of Proteins	

	Genome sequencing technology and analysis methods, Bioinformatics tools and automation in Genome Sequencing, analysis of gene expression data, Utility of EST database in sequencing, Bioinformatics in detection of Polymorphisms, SNPs and their relevance, Bioinformatics tools in microarray data analysis. Usages of visualization software available in public domain like VMD, Rasmol, Pymol, SpdbViewer, Chime, Cn3D and GRASP.	
Course	outcome	
CO1	The students shall get an adequate knowledge on the various online databases	
CO2	Students will be able to use the online tools for analysing various macromolecules of the cells	
CO3	The students shall Identify the role of phylogenetic analysis and related conclusions	
CO4	To learn the use of various tools for molecular analysis	
CO5	To understand the various methods for macromolecular sequencing	
Text bo	oks	
1	Bioinformatics (Sequence and Genome Analysis)- David W. Mount, Cold Spring Harbor Laboratory Press, 2001.	
2	Bioinformatics- Zoe Lacroix, Terence Critchlow, Morgan Kaufmann Publishersm, 2004.	
3	Bioinformatics – From Genomics to Drugs, Violume 1; Basic Technoliges, Thomas Lengauer, Wiley- VCH, 2001.	
4	Bioinformatics (Practical Approach): Sequence, Structure and Databanks – Des Higgins, OXFORD Univ. Press, 2003.	
5	Bioinformatics Computer Skills – Gibas&Jambeck, O' Reilly, 2001, I Ed.	
Referen	ice Books	
1	Bioinformatics Computing- Bryan Berjeron, Prentice-Hall of India, Private Ltd., 2003.	
2	Computational Molecular Biology (An Algorithmic Approach)- Pavel A. Pevzner, PrenticeHall of India, Private Ltd., 2004.	
3	11. Introduction to bioinformatics- T K Attwood, D J Parry-Smith, Pearson Education, 2004.	
4	Sequence Analysis (In A Nutshell)- Scott Market & Darryl Leon, O' Reilly, Ist Edition, 2003.	
5	Scolnick. J.; Drug Discovery and Design, Academic Press, London, 2001.	
6	N. R. Cohen, Editor, Guidebook on Molecular Modeling in Drug Design. Academic Press, San Diego, 1996.	
Journa	/Research Paper Link:	
	As suggested by concern subject faculty	

Com	Code AMTBT0220	LTP	Credit
Course [Fitle Entrepreneurship, IPR & Biosafety	300	3
Course of			
1	To learn the basics of accounting and finance in business		
2	To learn about the various policies of marketing		
3	To understand the use of IT in business development		
4	To learn about the IPR and its legal provisions.		
5	To learn about the various biosafety in various biological systems.		
Pre-requi	sites: Students are expected to have knowledge of basic biology, cell biolog biochemistry	gy and	
Course C	ontents / Syllabus		
Unit I	Accounting and Finance:		8 Hr.
	Taking decision on starting a venture; Assessment of feasibility of a new Approach a bank for a loan; Sources of financial assistance; Making a b proposal/Plan for seeking loans from financial institution and Banks; Fun bank for capital expenditure and for working; Statutory and legal requirem starting a company/venture; Budget planning and cash flow management in accounting practices: concepts of balance sheet, P&L account, and doub bookkeeping; Estimation of income, expenditure, profit, income tax etc.	business ids from nents for ; Basics	
Unit II	Marketing:		8 Hr.
	Assessment of market demand for potential product(s) of interest; conditions, segments; Prediction of market changes; Identifying ne customers including gaps in the market, packaging the product; Market li branding issues; Developing distribution channels; Pricing/Policies/Comp Promotion/Advertising; Services Marketing Negotiations/Strategy: financiers, bankers etc.; With government/law enforcement authorities companies/Institutions for technology transfer; Dispute resolution skills; I environment/changes; Crisis/ Avoiding/Managing; Broader vision- thinking	eeds of inkages, petition; With s; With External	
Unit III	Information Technology:		8 Hr.
Unit III	How to use IT for business administration; Use of IT in improving be performance; Available software for better financial management; E-bestup, management. Human Resource Development (HRD): Leadership Managerial skills; Organization structure, pros & cons of different structure building, teamwork; Appraisal; Rewards in small scale set up.	business p skills;	8 Hr.
Unit III Unit IV	How to use IT for business administration; Use of IT in improving be performance; Available software for better financial management; E-b setup, management. Human Resource Development (HRD): Leadership Managerial skills; Organization structure, pros & cons of different str	business p skills;	8 Hr. 8 Hr.
	How to use IT for business administration; Use of IT in improving be performance; Available software for better financial management; E-b setup, management. Human Resource Development (HRD): Leadership Managerial skills; Organization structure, pros & cons of different str Team building, teamwork; Appraisal; Rewards in small scale set up.	business p skills; ructures; opyright graphical rotection ew Case	

	An Introduction; Historical Backround; Introduction to Biological Safety Cabinets; Primary Containment for Biohazards; Biosafety Levels; Biosafety Levels of Specific Microorganisms; Recommended Biosafety Levels for Infectious Agents and Infected Animals; Biosafety guidelines - Government of India; Roles	
	of Institutional Biosafety Committee, Risk Analysis; Risk Assessment; Risk management and communication; Overview of National Regulations and relevant	
	International Agreements including Cartagena Protocol.	
Course out		
CO1	The students shall get an adequate knowledge on Accounting and Finance and will be able to do budget planning for any new venture	
CO2	Students will be able to Assessment of market demand for potential product(s) of interest and External environment/changes; Crisis/ Avoiding/Managing Broader vision–Global thinking	
CO3	The students shall Identify the role of Information Technology for business growth	
CO4	To disseminate knowledge on patents, patent regime in India and abroad and registration aspects and to make students aware about current trends in IPR and Govt. supports in promoting IPR	
CO5	The students shall Identify the role of regulatory committees in controlling the risk. Students should get enough information on ethical issues linked to research on animal models, transgenic, clinical trials.	
Text books	S	
1	Selected papers from scientific journals.	
2	Nithyananda, K V. (2019). Intellectual Property Rights: Protection and Management. India, IN: Cengage Learning India Private Limited.	
3	Neeraj, P., &Khusdeep, D. (2014). Intellectual Property Rights. India, IN: PHI learning Private Limited.	
4	V Sreekrishna, 2017. Bioethics and Biosafety in Biotechnology by New Age International publishers.	
E Reference	ce resources	
	https://kclau.com/wealth-management/best-budgeting-tools-online-softwares/ https://www.ccl.org/articles/leading-effectively-articles/fundamental-4- coreleadership-skills-for-every-career-stage/http://www.yourarticlelibrary.com/ organization/8-types-of-organisationalstructures-their-advantages-and- disadvantages/22143 https://opentextbc.ca/organizationalbehavioropenstax/chapter/reward-systems- inorganizations/#ch08rfin-9 https://online.hbs.edu/blog/post/accounting-skills-for-entrepreneurshttps:// www.investopedia.com/terms/f/feasibility-study.asphttps:// www.extension.iastate.edu/agdm/wholefarm/html/c5-92.htmlhttps:// economictimes.indiatimes.com/wealth/tax/how-to-compute-your-totaltaxabl e-income/articleshow/52956796.cms?from=mdr	
	•Subramanian, N., &Sundararaman, M. (2018). Intellectual Property Rights – An Overview. Retrieved from http://www.bdu.ac.in/cells/ipr/docs/ipr-eng-ebook.pdf	
	•World Intellectual Property Organization. (2004). WIPO Intellectual Property Handbook. (https://www.wipo.int/edocs/pubdocs/en/intproperty/489/wipo_pub	
	•489.pdf)	
Journal L	ink:	

https://www.ip.mpg.de/en/publications/journals/iic-international-review- ofintellectual-property-and-competition-law.html https://onlinelibrary.wiley.com/journal/15406261	https://www.springer.com/journal/10961	
https://onlinelibrary.wiley.com/journal/15406261		
	https://onlinelibrary.wiley.com/journal/15406261	

	M. TECH FIRST YEAR	
Course Code	AMTBT0251 L T P	Credit
Course Title	Bioinformatics Lab 0 0 4	2
Course objective		
1	To learn the various online databases	
2	To learn the online tools for analyzing various macromolecules of the cells	
3	To understand the phylogenetic analysis and related conclusions	
4	To learn the use of various tools for molecular analysis	
5	To understand the various methods for macromolecular sequencing	
Pre-requisites:		
	Students are expected to have knowledge of basic biology, cell biology and biochemistry	
Course Content		
1	To perform pair wise local and global sequence alignment for any two proteins and DNA sequences.	
2	To perform multiple sequence alignment for any five sequences and predicts the Phylogenetic relationship among them.	
3	Phylogenetic Analysis using PHYLIP - Rooted trees and Unrooted trees	
4	To predict secondary structure for any given protein sequence using Chou- Fasman, GOR and Neural network algorithms.	
5	To visualize tertiary structure of any given protein sequence using Rasmol/PyMol/PMV.	
6	To visualize the genomic map of Human genome and find out the size, number of genes and number of proteins encoded on Chr-Y.	
7	Homology Modelling using Modeller	
8	To find out the RMSD value from any two-protein structure alignment.	
9	Construction of Cladogram	
10	Different interactions using CYTOSCAPE	
11	Primary Structure Analysis of a Protein Using ProtParam	
12	Finding the Active Site Pockets of a given Protein Molecule	
Course outcome		
CO1	The students will learn the various online databases	
CO2	Students will learn the online tools for analysing various macromolecules of the cells	
CO3	They will understand the phylogenetic analysis and related conclusions	
CO4	The students will learn the use of various tools for molecular analysis	
CO5	The students will understand the various methods for macromolecular sequencing	

Course Code	AMTBT0254	LTP	Credit
Course Title	Cell & Tissue Culture Techniques Lab	0 0 4	2
Course objectiv	/e:		
1	To equip students with the skills to perform basic cell and tissue cu cultivate plant cells and tissues, induce callus tissue formation and regenerate plants, and conduct molecular biology techniques such	somatic embr	yos,
Pre-requisites:			
Students are ex	pected to have knowledge of basic biology, cell biology and bioch	nemistry	
Course Content	ts / Syllabus		
1	Preparation of a basic cell culture medium using common ingredien amino acids, vitamins, and salts.	nts such as	CO1
2	Growth curve analysis by measure their growth over time using a spectrophotometer.		CO1
3	Trypsinization to detach adherent cells from a culture dish using trypsin and then separate the cells using a cell strainer.		CO1
4	Suspension culture using a spinner flask and monitor their growth	and viability.	CO1
5	Transfection of cells with a plasmid DNA encoding a fluorescent pro then visualize the expression of the protein using a fluorescence m		CO3
6	Production of plant callus by explants on a callus-inducing medium the development of callus tissue over time.		CO2
7	Induction of somatic embryogenesis on a medium containing plant regulator to induce the formation of somatic embryos.	growth	CO2
8	Plant regeneration from callus tissue by transferring the tissue onto a regeneration medium.		CO2
9	Androgenesis, anthers or pollen grains culture on a medium containing plant growth regulator to induce the formation of haploid plants.		CO2
10	Field propagation of regenerated plants by the transfer of regenerated plants to soil and monitor their growth and development under field conditions.		CO1
Course outcom	e		
CO1	Students will be able to perform basic cell and tissue culture techniques, including media preparation, trypsinization, cell separation, and suspension culture.		K1-K2
CO2	Students will be able to culture plant cells and tissues using various techniques, induce the formation of callus tissue and somatic embryos, and regenerate plants from callus tissue.		К1-К2
CO3	Students will be able to perform molecular biology techniques, in DNA isolation.	ncluding	К2-КЗ

	M. TECH FIRST YEAR		
Course Code	AMTBT0211	LTP	Credit

Course Title	Genetic Engineering 3 0 0	3
Course objectiv	e:	
1	It is intended to impart basic undergraduate-level knowledge in the area of molecular biology and recombinant DNA technology.	
2	The student would be able to understand the working details of the cloning of a gene.	
3	They would also be able to assimilate recent research findings, advancement and development in the rDNA technology.	
4	The use of virtual lab and computational tools would enable them to perform in silico cloning of the selected DNA.	
5	To understand the DNA sequencing methods	
Pre-requisites:		
Students are exp	ected to have knowledge of basic biology, cell biology and biochemistry	
Course Content	s / Syllabus	
UNIT-I	Molecular Tools	8 Hr.
	DNA Structure and properties; Enzymes used in Genetic Engineering; Cohesive and blunt end ligation; Linkers; Adaptors; Homopolymeric tailing; Labelling of DNA: Nick translation, Random priming, Radioactive and non- radioactive probes, Hybridization techniques, Hybridization techniques; Chromatin Immunoprecipitation; DNA-Protein Interactions-Electromobility shift assay; DNaseIfootprinting; Methyl interference assay	
UNIT-II	Vectors	8 Hr.
	Plasmids; Bacteriophages; M13 mp vectors; PUC19 and Bluescript vectors, Phagemids; Lambda vectors, Insertion and Replacement vectors; Cosmids; Artificial chromosome vectors (YACs; BACs); Animal Virus derived vectors; Expression vectors; Inclusion bodies; Methodologies to reduce formation of inclusion bodies; Baculovirus and pichia vectors system, Plant based vectors, Ti and Ri as vectors, Yeast vectors, Shuttle vectors	
UNIT-III	Techniques in Genetic Engineering	8 Hr.
	Insertion of Foreign DNA into Host Cells; Transformation; Isolation of mRNA and total RNA; cDNA and genomic libraries and its construction; cDNA and genomic cloning; Expression cloning; Jumping and hopping libraries; Southwestern and Far-western cloning; Protein-protein interactive cloning and Yeast two hybrid system; Phage display; Principles in maximizing gene expression	
UNIT-IV	PCR and its applications	8 Hr.
	Primer design; Fidelity of thermostable enzymes; DNA polymerases; Concept of PCR, Types of PCR, Gene specific and degenerate primer design, linkers, adaptors, Fidelity of uDNA polymerase. Application of PCR. Chimeric protein engineering by PCR	
UNIT-V	Sequencing Methods	8 Hr.

	· · · · · · · · · · · · · · · · · · ·	
	Sequencing methods; Enzymatic DNA sequencing; Chemical sequencing of DNA; Automated DNA sequencing; RNA sequencing; Introduction of DNA into mammalian cells; Transfection techniques; Gene silencing techniques; siRNA technology; Micro RNA; Construction of siRNA vectors; Principle and application of gene silencing; Gene Therapy; Suicide gene therapy; Gene replacement; cDNA and intragenic arrays; Differential gene expression and protein array, Genome editing-CRISPR and other genome editing tools.	
Course outco	me	
CO 1	Understand the basic concept and procedure of gene cloning and the role of enzymes and vectors used for genetic manipulation and genetic engineering	
CO 2	Acquired theoretical knowledge of vectors, their different types and applications in genetic engineering.	
CO 3	Getting detailed knowledge of construction of gene libraries and their screening methods.	
CO 4	Have knowledge of PCR technique, their different types and applications.	
CO 5	Understand the basic concept of genetic engineering techniques for selection of recombinants.	
Text books		
1	Winnacker, Ernst L. (1987), From genes to clones: introduction to genetechnology [Gene und Klone] (in German), Horst Ibelgaufts (trans.),Weinheim, New York: VCH, ISBN 0-89573-614-4.	
2	Genetic Engineering by Dr Smita Rastogi & Dr Neelak Pathak, Oxford University Press	
3	Genetic Engineering, Principles& Practice by Sandhya Mitra, McGraw Hill Education.	
Reference Bo	ooks	
1	Principles of Gene Manipulation and Genomics, Primrose & Twyman.	
2	Molecular Biology of the Cell. 4th edition. Alberts B, Johnson A, Lewis J, et al. New York: Garland Science; 2002.	
3	Modern Genetic Analysis. Griffiths AJF, Gelbart WM, Miller JH, et al. New York: W. H. Freeman; 1999.	
Journal/Rese	arch Paper Link:	
	As suggested by concern subject faculty	

Course Code	AMTBT0212	L T P	Credit
Course Title	Applied Food Biotechnology	300	3
Course objectiv			
1	To learn about the various microbiological examination of food born diseases	s and food	
2	To learn about the development and production of novel products		
3	To understand GM foods and the legal issues associated with them		
4	To learn about the industrial production of various food products		
5	To learn the methods of production of vitamins and enzymes.		
Pre-requisites:			
	Students are expected to have knowledge of basic biology, cell bio biochemistry	logy and	
Course Content	s / Syllabus		
Unit I	Food Biotechnology		8 Hr.
	Introduction & Applications; Methods for the microbiological ex of water and foods; Control of Microbiological quality and sat borne illnesses and diseases; Microbial cultures for food fermenta maintenance, strain development	fety; Food	
Unit II	Biosensors in food technology		8 Hr.
	Starter cultures-types, designing and development, micro encapsu packaging, scopes and challenge; Development and formulation products such as probiotic foods. Nutrigenomics-concept, significance and relevance. Biosensors and novel tools and their a in food science & Technology	n of novel working,	
Unit III	GM foods		8 Hr.
	Introduction and controversies related to GMOs. Ethical issues of GM foods; testing for GMOs; current guidelines for the production and movement of GMOs; labelling and traceability; trade related biosafety; risk assessment and risk management. Public perception foods. IPR. GMO Act-2004. New products and processes in value commodities including plant and animal products.	on, release ed aspects; on of GM	
Unit IV	Industrial Food Biotechnology I		10 Hr.
	Industrial production of organic acids (vinegar, lactic acid), beverages (beer, wine, and distilled alcoholic beverages such as rum, vodka), glycerol; Propagation of baker's yeasts; Fermer products such as cheese, yoghurt, sweet curd, paneer, sh Fermented pickles.	s whiskey, nted dairy	
Unit V	Industrial Food Biotechnology II		10 Hr.
	Industrial production of important primary and secondary metabolistic, vitamins, biosurfantants, polysaccharides. Enzyme a in food industry. Advantages and constraints of immobilized enzymicrobial cells. Types of enzyme reactors. Aerobic and anaerobic of effluents from food processing industry	application zymes and	
Course outcome			
CO1	To identify microorganism responsible for food spoilage.		

CO2	Demonstrate knowledge methods of packing, and the application of biosensors in food industries
CO3	To understand the ethical issues lined with GM food production
CO4	Demonstrate the industrial production of various food products
CO5	To explain the industrial application of various enzymes
Text books	·
1	Industrial Microbiology Prescott & Dunn, CBS Publishers
2	Modern Food Microbiology by Jay JM, CBS Publishers
3	Comprehensive Biotechnology by Murray & Mooyoung, Academic press
4	Industrial Microbiology by Casida L.R., New Age International Pvt. Ltd.
5	Food Microbiology; Frazier WC; 4th ed, Tata-McGrowhill Pub.
Reference Bool	KS I I I I I I I I I I I I I I I I I I I
1	Microbiology by Pelczar, Chan, and Krieg, TMH
2	Fermentation Biotechnology, Principles, Processed Products by Ward OP, Open
3	University Press.
4	Lee, B. H. Fundamentals of Food Biotechnology.VCH. 2006
Journal/Resear	ch Paper Link:
	As suggested by concern subject faculty

	M. TECH FIRST YEAR	
Course Code	AMTBT0213 L T P	Credit
Course Title	Molecular Modelling & Industrial Application3 0 0	3
Course objectiv	۵.	
1	To learn about the basics of molecular modelling	
2	To understand the usage of computer simulation	
3	To understand the basic of drug development.	
4	To learn about the herbal drug and its trade scenario.	
5	To understand the method of vaccine production.	
Pre-requisites:	1	
	Students are expected to have knowledge of basic biology, cell biology and biochemistry	
Course Content	s / Syllabus	
Unit I	Molecular Modelling	8 Hr.
	Introduction; Useful Concepts in Molecular Modelling; The Molecular Modelling Literature; Molecular Modelling software: BIOSUITE; Force Fields	
Unit II	Init II Energy Minimisation and Computer Simulation	
	Minimisation and Related Methods for Exploring the Energy Surface. Non- Derivative method, 1st and 2nd order minimisation methods. Results of a Simulation and Estimating Errors. GROMACS and CNS. Molecular Dynamics & Monte Carlo Simulation.	
Unit III	Drugs	8 Hr.
	An introduction, Overview of drug discovery process, Trends in drug discovery process. Rationale of Drug Discovery: Medical needs, Target identification, Target validation, Receptors and assay development.	
Unit IV	Herbal Drugs	8 Hr.
	Definition, Trade scenario, Phytochemical standardization and fingerprinting, Marker compounds, Polyherbal formulations. Drug Development and Pre-Clinical Studies: Drug receptor interactions; enzyme inhibition and inactivation, In-vitro and in-vivo pharmacodynamic models, Therapeutic index, Pharmacokinetics - Microbial and animal models, In- vitro and insilico toxicological models, Drug formulations.	
Unit V	Applications of microbes for designing vaccines	8 Hr.
	Applications of microbes for designing vaccines: case study.	
Course outcome		
1	Students will learn about the basics of molecular modelling	
2	Students will understand the usage of computer simulation	
3	Students will understand the basic of drug development.	
4	Students will learn about the herbal drug and its trade scenario.	
5	Students will be able to understand the method of vaccine production.	
Text books		

1	A.R.Leach, Molecular Modelling Principles and Application, Longman, 2001.
2	J.M.Haile, Molecular Dynamics Simulation Elementary Methods, John Wiley and Sons, 1997.
3	Satya Prakash Gupta, QSAR and Molecular Modelling, Springer - Anamaya Publishers, 2008.
4	Patwardhan B, Drug Discovery and Development-Traditional Medicine and Ethnopharmacology, New India Publishing (2007)
5	Larsen PK, Leljifore T and Medsan U, Text Book of Drug Design and Discovery, CRC Press (2009)
Reference Bool	KS I I I I I I I I I I I I I I I I I I I
1	Hillisch A and Hilgenfeld R, Modern Methods of Drug Discovery, Birkhauser (2003).
Journal/Research Paper Link:	
	As suggested by concern subject faculty

	M. TECH FIRST YEAR	
Course Code	AMTBT0214 L T P	Credit
Course Title	Bioreactor Analysis and Design3 0 0	3
Course objectiv	٥.	
1	To learn about the designing of bioreactor systems	
2	To learn about the control involved in bioreactor system	
3	To learn about the various types of bioreactor processes	
4	To understand the reactor dynamics	
5	To learn the design aspect and safety issues.	
Pre-requisites:		
-	Students are expected to have knowledge of basic biology, cell biology and biochemistry	
Course Content	s / Syllabus	
UNIT I	Material balance and design	8 Hr.
	Introduction; General design information; Material and energy balance calculations; Process Flow sheeting,Selection of bioprocess equipment (upstream and downstream); Specifications of bioprocess equipment; Mechanical design of reactors, heat transfer and mass transfer equipment; Design considerations for maintaining sterility of process streams and process equipment; Piping and instrumentation; Materials of construction for bioprocess plants.	
UNIT II	Control of bioreactor	8 Hr.
	Basic aspects of bioreactor designing, Physical, chemical and biological sensors and control, Advanced control strategies viz. PID controllers, Fuzzy logic based controllers and Artificial Neural Network (ANN) based controllers, Basic concepts of computer modelling and optimization in bioprocess applications.	
UNIT -III	Ideal Bioreactor and its working	8 Hr.
	Ideal bioreactors: Batch reactors, Fed-batch reactors, enzyme-catalyzed reaction in CSTRs, CSTR reactors with recycle and wall cell growth, the ideal plug-flow tubular reactor, Reactors with nonideal mixing: Mixing times in agitated tanks, residence time distribution, models for nonideal reactors, Mixing-bioreaction interactions.	
UNIT -IV	Types of Bioreactors	8 Hr.
	Reactor dynamics and stability, Multiphase bioreactors: conversion of heterogeneous substrates, packed-bed reactors, bubble column bioreactors, fluidized bed bioreactors, trickle-bed reactors, airlift reactor, Immobilized Enzyme reactors, Photo bioreactors, Hollow fibre membrane bioreactors. Scale up and scale down issues: Effect of scale on oxygenation, mixing, sterilization, pH, temperature, inoculum development, nutrient availability and supply; Bioreactor scale-up based on constant power consumption per volume, mixing time, impeller tip speed (shear), mass transfer coefficients.	

UNIT V	Downstream Processing	8 Hr.
	Facility design aspects; Utility supply aspects; Equipment cleaning aspects; Culture cell banks; cGMP guidelines; Validation; Safety. Process economics; Case studies, Scale up of downstream processes: Adsorption	
	(LUB method); Chromatography (constant resolution etc.); Filtration (constant resistance etc.); Centrifugation (equivalent times etc.); Extractors	
Course outco	(geometry based rules).	
CO1	After completing the course students will able to design the bioreactor system	
CO2	After completing the course students will able to illustrate the control involved in bioreactor system	
CO3	After completing the course students will able to identify the various types of bioreactor processes	
CO4	After completing the course students will able to analyse the reactor dynamics	
CO5	After completing the course students will able to evaluate the design aspect and safety issues associated with reactor system.	
Text books		
1	Moser, Anton, Bioprocess Technology: Kinetics and Reactors, Springer Verlag, 1988.	
2	Bailey J.E. &Ollis, D.F. Biochemical Engineering Fundamentals, 2nd ed., McGraw Hill, 1986	
3	Lee, James M. Biochemical Engineering, PHI, USA.	
4	Atkinson, Handbook of Bioreactors, Blanch, H.W. Clark, D.S. Biochemical Engineering, Marcel Decker, 1999	
5	Biochemical Engineering fundamentals" 2nd edJ E Bailey and D F Ollis, McGraw-Hill (1986) Chapters 8,9&10.	
6	Biochemical Engineering" -S Aiba, A E Humphrey and N Millis , 1978, University of Tokyo Press.	
7	Biotechnology" Vols. 3 & 4 Eds., S Rehm and G Reed. VCH (1991).	
Reference B		
1	Biochemical Engineering and Biotechnology Handbook" 2nd Ed.,.Atkinson &F.Mavituna, Stockton Press (1991).	
2	Biorector Design & Product Yield", BIOTOL series, Butterworth - Heinemann (1992).	
3	Principles of fermentation technology" - F Stanbury and A Whitaker, Pergamon press (1984)	
4	Unit operations of Chemical Engineering" 5th ed. by W L McCabe, J C Smith and P. Harriot Mc Graw-Hill (1993).	
5	Bioprocess Engineering Principles" by Pauline M.Doran, Academic Press.	
6	Feedback and Control systems- Schaum's outline series, McGraw-Hill Book Comp., 1967	
7	Unit Operations of Chemical Engineering- Mc Caba Smith, Harriott, Mc Graw – Hill Chemical Engg. Series., V Ed., 1985.	
Journal/Rese	earch Paper Link:	
	As suggested by concern subject faculty	

	M. TECH FIRST YEAR	
Course Code	AMTBT0215 L T P	Credit
Course Title	Enzyme Technology & Industrial Application 3 0 0	3
Course objectiv		
1	To learn about the kinetics involved in enzymatic reactions.	
2	To learn about the various biochemical processes involved in the microbial growth	
3	To learn about the various processes in bioreactor	
4	To understand the various separation methods involved in bioprocess	
5	To analyze the different bioprocess steps in industrial production.	
Pre-requisites:		
	Students are expected to have knowledge of basic biology, cell biology and biochemistry	
Course Content		
UNIT I-	ENZYME TECHNOLOGY	8 Hr.
	Introductions: Enzymes- Michaelis-Menten kinetics. Kinetics and StatisticsInhibition- Effect of pH and temperature- Enzymology-Immobilized enzymes: Methods, Mass transfer considerations and Industrial enzymes.	
UNIT II-	METABOLISM, STOICHIOMETRY AND MICROBIAL GROWTH KINETICS	8 Hr.
	Introduction to metabolism- Nutrient transport- Glycolysis - TCA cycle and other pathways - Control of metabolism. Factors affecting microbial growth – Stoichiometry- mass balances and energy balances. Growth kinetics, Measurement of growth.	
UNIT III-	BIOREACTORS, STERILIZATION, SENSORS AND INSTRUMENTATION	8 Hr.
	Introduction to bioreactors - Batch and Fed-batch bioreactors, Continuous bioreactors, Immobilized cells, Bioreactor operation, Sterilization, Aeration, Sensors. Instrumentation, Culture - specific design aspects: plant/mammalian cell culture reactors.	
UNIT IV-	PRIMARY & SECONDARY SEPARATION PROCESS	8 Hr.
	Biomass removal - Biomass disruption – Membrane based techniques. Extraction -solvent, aqueous two phases, super critical, and Adsorption. Chromatography, Precipitation (Ammonium Sulfate, solvent), Electrophoresis (capillary), Crystallization, Drying and Freeze drying.	
UNIT V-	INDUSTRIAL APPLICATION	8 Hr.
	White Biotechnology: Few industrial process using enzymes for production of drugs and fine chemicals, Enzyme based biosensors, Enzyme in organic catalysis, Analytical applications, Applications in food	

	M. TECH FIRST YEAR		
Course Code	AMTBT0216	L T P	Credit

Course Title	Applied Bioenergy 3 0 0	3
Course objective	2:	
1	To understand the basics of bioenergy	
2	To learn the principals of biofuel production	
3	To learn about the current application of bioenergy	
4	To understand the impact of energy on economy	
5	To understand production of biofuels in real life.	
Pre-requisites:	1	
*	Students are expected to have knowledge of basic biology, cell biology and biochemistry	
Course Contents	s / Syllabus	
Unit I	Introduction to applied bioenergy	8 Hr.
	Introduction to applied bioenergy, Types of bioenergy, Energy scenario-role of energy in economic development and social transformation, Commercial and non-commercial forms of energy, Present and future global projections of energy consumptions.	
Unit II	Biomass and Energy Conservation	8 Hr.
	Principles of biomass energy conversion processes, biological, chemical and thermo-chemical technologies for biomass conversion and their utilization covering: Biogas, Produces gas, Alcohol and Biodiesel, Second generation biofuel from high efficiency algal-derived biocrude, Biobased fats (Lipids) and oils from biomass for energy production, Biorefinery systems: An Overview. Microbial fuel cell and their application	
Unit III	Bioenergy	8 Hr.
	Current bio-energy applications and conversion technologies, Advantages of applied bioenergy over other sources of energy, Advances in bio-energy research: An overview of technological developments, bioenergy value chain, Databases of bioenergy related enzymes, Sustainable farming of bioenergy crops.	
Unit IV	Impact of Energy on Economy and Environment	8 Hr.
	Impact of Energy on Economy, Development and Environment, Energy for Sustainable Development, Energy and Environmental policies, Need for use of new and renewable energy sources, Energy Policy Issues: Fossil Fuels, Renewable Energy, Power sector reforms, restructuring of energy supply sector, energy strategy for future, Status of Nuclear and Renewable Energy: Present Status and future promise.	
Unit V	Case study	8 Hr.
	Case study 1: Biodiesel from Jatropha plant as transport fuel, A case study of UP State (India) 2. Generation of Bio-fuel by Using Waterweeds: A Case Study in Solapur City	
Course outcome		
CO1	Demonstrate different types of bioenergy.	
CO2	Demonstrate the production of various types of biofuel using different substrates.	

CO3	To explain the advantages of applied bioenergy over other sources of energy and advances in bio-energy research.	
CO4	To describe the Impact of Energy on Economy.	
CO5	To describe the application of of biofuel in real life.	
Text books		
1	Anthony San Pietro (1980); Biochemical and Photosynthetic aspects of Energy Production, Academic Press, New York.	
2	Berman, ER Geothermal Energy, Noyes Data Corporation, New Jersey	
3	Parker, Colin & Roberts, (1985); Energy from Waste- An Evaluation of Conversion Technologies, Elsevier Applied Science London	
Reference Bool	ks	
1	Ralph E.H. Simsed. (2004); Bioenergy options for cleaner environment by World Renewable Energy Network.	
2	Ravindranath N.H. and Hall D.O. (1995); Biomass, Energy and Environment, A developing country perspective from India by, Oxford University Press,	
3	Brown Robert C. (2003); Biorenewable Resources: Engineering New Products from Agriculture, Iowa State University Press, USA	
4	Boyle Godfrey ed. (1996): Renewable Energy: Power for a sustainable future, Oxford, OUP	
Journal/Resear	ch Paper Link:	
	As suggested by concern subject faculty	

Course	Code	AMTBT0204	L T P	Credit
Course	Title	Cell & Tissue Culture Techniques	3 0 0	3
Course of	bjectiv			
1		To learn the basics of animal cell culturing technique.		
2		To understand the various methods and advancements of culture tee	chniques.	
3		To analyse the applications of animal cell culturing.		
4		To learn the basics of plant cell and tissue culture technique.		
5		To understand the various methods and advancements in plant cell culture.	and tissue	
Pre-requi	isites:			
		Students are expected to have knowledge of basic cell and molecula	ar biology.	
1		s / Syllabus		
Unit 1		z Tissue Culture Technology Basics		8 hr
	Physic Tempe	cell culture techniques, Types of cell culture media; Ingredients ochemical properties; CO ₂ and bicarbonates; Buffering; Oxygen; O erature; Surface tension and foaming; Balance salt solutions; A n supplements;	smolarity;	
Unit 2	Metho	ods of Cell & Tissue Culture		8 hr
	Trypsi culture	last culture; Chicken liver and kidney culture; Secondary nization; Cell separation; Continuous cell lines; Suspension cultu e etc.; Behaviour of cells in culture conditions: division, growt olism of estimation of cell number; Development of cell lines	re; Organ	
Unit 3	Applic	cations of Cell and Tissue Culture Technique		8 hr
	scale j animal polluta	loning and selection; Transfection and transformation of cells; Co production of animal cells, stem cells and their application; Appl cell culture for <i>in vitro</i> testing of drugs; Testing of toxicity of envi ants in cell culture; Application of cell culture technology in proc and animal viral vaccines and pharmaceutical proteins, Green Me	ication of ronmental luction of	
Unit 4	Plant	Cell & Tissue Culture Basics		10 hr
	embry improv disease and cl	mentals of plant tissue culture, plant regeneration: organogenesis ogenesis; somaclonal variation, its genetic basis and application vement. Cell/callus line selection for resistance to herbicide, s es.: Isolation, culture and plant regeneration, protoplast fusion, iden haracterization of somatic hybrids., Field techniques for propa- prated plants.	n in crop stress and ntification	
Unit 5		iques of Plant Cell & Tissue Culture		10 hr
	B5, S parame	nt selection, sterilization and inoculation; Various media preparat H PC L2; Callus and cell suspension culture; Induction an eters; Chromosomal variability in callus culture. Plant regenera o, meristem and callus culture. Androgenesis: Anther and pollen cul	d growth tion from	
Course o	utcome	:		
CO1	After c culturi	completion of the course, students will learn the basics of animal celling.	1	

CO2	They will understand about the various methods and protocols of cell culturing.	
CO3	They will analyse the different types of applications of animal cell culturing.	
CO4	Students will learn the basics of plant tissue culture.	
CO5	Students will be able to understand the different methods of plant tissue culture and their applications.	
Text be	ooks	
1	B. Hafez and E.S.E Hafez, Reproduction in farm animals, 7th Edition, Wiley Blackwell, 2000	
2	G.E. Seidel, Jr. and S.M. Seidel, Training manual for embryo transfer in cattle (FAO Animal Production and Health Paper-77), 1st Edition, W.D. Hoard and sons FAO, 1991	
3	I. Gordon, Laboratory production of cattle embryos, 2nd edition, CAB International, 2003.	
Refere	nce Books	
1	Louis-Marie Houdebine, Transgenic Animals: Generation and Use 5th Edition, CRC Press, 1997.	
2	Plant Tissue Culture: Theory and Practice, a Revised Edition by S.S. Bhojwani and M.K. Razda	
3	Plants from Test Tubes: An Introduction to Micropropagation by LydianeKyte	
Journa	I/Research Paper Link:	
	As suggested by concern subject faculty	

	M. TECH FIRST YEAR	
Course Code	AMTBT0218 L T P	Credit
Course Title	Diagnostic Techniques in Biotechnology3 0 0	3
Course objective		
1	To learn the basics of diagnostic techniques.	
2	To understand the different enzymes and related test methods.	
3	To learn the methods of immunodiagnostics.	
4	To understand the product development related to diagnostics.	
5	To learn the methods of DNA based diagnostics.	
Pre-requisites:	To fear the methods of DTVT based diagnostics.	
re-requisites.	Students are expected to have knowledge of basic biology, cell biology and biochemistry	
Course Contents	s / Syllabus	
Unit I	Analytical Methods	8 hr
	Volumetric analysis, Balancing & Weighing, Concept of solute & solvent, Units of measurement. Specimen Collection & Processing: Specimen collection (Blood, urine, spinal fluid, saliva synovial fluid, Amniotic fluid), Preservation, transportation	
Unit II	Clinical Enzymology	8 hr
	Principle of diagnostic enzymology, Digestive enzyme, Miscellaneous enzyme. General Function Tests: Liver function test, Cardiac Function Test, Renal Function Test, Thyroid Function test, Reproductive endocrine function test	
Unit III	Immunodiagnostics	8 hr
	Introduction, Antigen-Antibody Reactions, Conjugation Techniques, Antibody Production, Enzymes and Signal Amplification Systems, Separation and Solid-Phase Systems, Studies related to bacterial, viral and parasitic infections.	
Unit IV	Product Development	10 hr
	Immunoassay Classification and Commercial Technologies, Assay Development, Evaluation, and Validation, Reagent Formulations and Shelf Life Evaluation, Data Analysis, Documentation, Registration, and Diagnostics Start-ups.	
Unit V	DNA based diagnostics	10 hr
	PCRRT-PCR, qPCR, Hot start PCR, Nested PCR), RFLP, SSCP, Microarrays, FISH, In-situ hybridization, Studies related to bacterial, viral and parasitic infections, Cell based diagnostics: Antibody markers, CD Markers, FACS, HLA typing, Bioassays, Viral DNA detection using Rapid kits and PCR	
Course outcome		
CO1	The students will learn the basics of diagnostic techniques.	
CO2	The students will understand the different enzymes and related test methods.	
CO3	The students will learn the methods of immunodiagnostics.	
CO4	The students will understand the product development related to diagnostics.	

CO5	The students will learn the methods of DNA based diagnostics.	
Text books		
1	Tietz Textbook of Clinical Chemistry, Carl A. Burtis, Edward R. Ashwood,Harcourt Brace & Company AisaPvt. Ltd.	
2	Commercial Biosensors: Graham Ramsay, John Wiley & Son, INC. (1998).	
3	Essentials of Diagnostic Microbiology, Lisa Anne Shimeld.	
Reference Book	<u>(</u> \$	
1	Diagnostic Microbiology, Balley& Scott's.	
2	Tietz Text book of Clinical Biochemistry, Burtis& Ashwood. 6. The Scienceof Laboratory Diagnosis, Crocker Burnett.	
Journal/Resear	ch Paper Link:	
	As suggested by concern subject faculty	

		M. TECH FIRST YEAR		
Course	Code	АМТВТ0219	L T P	Credit
Course Title		3-D Printing Technology	300	3
Course of	ojective	:		
1		Able to know the fundamentals of RP Systems & its evolution and the Process in RP and association of RP Systems with 3D modelling & Mesh		
2	Able to	Able to know the RP Systems, Process, Materials & Classifications		K3, K4
3	Able to know and working with Mesh File & their formats like STL format, 3MF format, OBJ formats. Conversion to Mesh files, their properties, operations, storage, inspections & defects			K3, K4
4	Able to	ble to know the applications of RP Systems in various Fields		K3, K4
Pre-requi	sites:			
Basic und	erstandi	ng of Information Technology.		
Course C	ontents	/ Syllabus		
UNIT-I	Introduction:			4 hours
	differ	rical Developments, Fundamentals of RP Systems and its Classi rent basis, Rapid Prototyping Process Chains, 3D Modelling ration, Data Conversion and Transmission.		
UNIT-II	RP Systems:			12 hours
	Input Manu Rapio	 d Polymer Based Rapid Prototyping systems: SLA, Material Jet Materials Based Rapid Prototyping Systems: Laminate ifacturing (LOM) and Fused Deposition Modelling Systems, Po d Prototyping Systems: Selective Laser Sintering, Multi-Jet Fusi g Systems. 	ed Object wer Based	
UNIT-III	RP D	atabase & Design Optimization:		8 hours
		l Prototyping Data Formats, STL Format, STL file problems, STL A, Topology Optimization, Gcode for RP Systems	file repair,	
UNIT-IV	RP Applications:			8 hours
	produ	lopment of dies for Moulding, RP Applications in developing protects, application in medical fields, Development of bone replaces, etc., RP materials and their biological acceptability.	• 1	
Course ou	itcome	: After completion of this course students will be able to		
CO 1	Under	stand the fundamentals of RP Technologies and process involvement	ent in them	K1,K2
CO 2		Understand the methodology to manufacture the products using RP technologies and study their applications, advantages and case studies		K3, K4
CO 3		Understand the Design aspects and their respective challenges along with the resolution for them		K3, K4, K5
CO 4		Understand the various applications of various RP Systems with case studies & Materials		K3,K4
Text book	KS			
1		Rapid Prototyping: Principles an Applications: Chee Kai Chu Leong, Chu Sing Lim	a, Kah Fai	
2		Additive Manufacturing Technologies: 3D Printing, Rapid Proto Direct Digital Manufacturing: Brent Stucker, David W. Rosen, Ia	• I U	

Reference Books		
1	Rapid Manufacturing: The Technologies and Applications of Rapid Prototyping and Rapid Tooling: Pham, Duc, Dimov, S.S.	
2	Rapid Prototyping and Manufacturing: Fundamentals of Stereo Lithography: P. Jacobs	
3	Rapid System Prototyping with FPGAs: Accelerating the Design Process: R.C. Cofer, Benjamin F. Harding	
4	Rapid Prototyping of Digital Systems: Hamblen, James O., Hall, Tyson S., Furman, Michael D.	