NOIDA INSTITUTE OF ENGINEERING AND TECHNOLOGY, GREATER NOIDA

(An Autonomous Institute)



Affiliated to

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



Evaluation Scheme & Syllabus

For

Master of Technology in Biotechnology (BT) First Year

(Effective from the Session: 2021-22)

NOIDA INSTITUTE OF ENGINEERING & TECHNOLOGY, GREATER NOIDA (An Autonomous Institute)

M.TECH. BIO-TECHNOLOGY

Evaluation Scheme SEMESTER-I

SI.	Subject	Subject	Pei	riods	5	Eval	uation	Scheme	es	End Semester		Tota I	Credit
No.	Codes	•	L	т	Р	СТ	TA	TOT AL	PS	TE	PE		
1	АМТВТО101	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
5		Elective -I*	3	0	0	20	10	30		70		100	3
6		Elective -II*	3	0	0	20	10	30		70		100	3
7	AMTBT0151	Applied Biochemistry & Molecular Biology Lab	0	0	4				20		30	50	2
8	AMTBT0152	Bioprocess Engineering & Technology Lab	0	0	4				20		30	50	2
		TOTAL										600	19

L: Lecture T: Tutorial P: Practical/Project CT: Class Test TA: Teacher's Assessment, ESE: End Semester Examination

(*) Refer the Electives list

Elective-I*

- 1.AMTBT0111 Immunology & Vaccine Technology
- 2.AMTBT0112 Quality Assurance and Quality Control
- 3.AMTBT0113 Applied Clinical Research

Elective-II*

- 1.AMTBT0114 Biological Treatment of Wastewater
- 2.AMTBT0115 Nano Biotechnology & Toxicology
- 3.AMTBT0116 Industrial Biotechnological Products

Abbreviation Used:-

NOIDA INSTITUTE OF ENGINEERING & TECHNOLOGY, GREATER NOIDA (An Autonomous Institute)

M.TECH. BIO-TECHNOLOGY

Evaluation Scheme SEMESTER-II

Sl. No	Subject Codes	Subject	Per	riod	S	Evalu	ıation	Schen	nes	Enc Seme		Tot al	Cre dit
			L	Т	P	CT	TA	TO TAL	PS	TE	PE		
1	AMTBT0201	Bioinformatics	3	0	0	20	10	30		70		100	3
2	AMTBT0202	Entrepreneurship, IPR &Biosafety	3	0	0	20	10	30		70		100	3
3		Elective – III*	3	0	0	20	10	30		70		100	3
4		Elective- IV*	3	0	0	20	10	30		70		100	3
5		Elective- V*	3	0	0	20	10	30		70		100	3
6	AMTBT0251	Bioinformatics Lab	0	0	4				20		30	50	2
7	AMTBT0252	Entrepreneurship, IPR & Biosafety Lab	0	0	4				20		30	50	2
8	AMTBT0253	Seminar-I	0	0	2				50			50	1
		TOTAL										650	20

L: Lecture T: Tutorial P: Practical/Project CT: Class Test TA: Teacher's Assessment, ESE: End Semester Examination

(*) Refer the Electives list

Abbreviation Used:-

NOIDA INSTITUTE OF ENGINEERING & TECHNOLOGY, GREATER NOIDA (An Autonomous Institute)

M.TECH. BIO-TECHNOLOGY

Elective-III*

- 1.AMTBT0211 Genetic Engineering
- 2.AMTBT0212 Applied Food Biotechnology
- 3.AMTBT0213 Molecular Modelling & Industrial Application

Elective-IV*

- 1.AMTBT0214 Bioreactor Analysis & Design
- 2.AMTBT0215 Enzyme Technology & Industrial Application
- 3.AMTBT0216 Applied Bioenergy

Elective-V*

- 1.AMTBT0217 Cell & Tissue Culture Techniques
- 2.AMTBT0218 Diagnostic Techniques in Biotechnology
- 3. AMTBT0219 3-D Printing Technology

		M. TECH FIRST YEAR				
Course	Code	AMTBT0101 L T P	Credit			
Course	Title	Applied Biochemistry & Molecular Biology 3 0 0	3			
Course	objective:					
1	To unde	erstand the various concepts of molecular biology and biochemistry				
2	Determi	ne the structure and function of biomolecules and evaluate the complexity o	f			
2	various	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-requ	uisites:					
		Students are expected to have knowledge of basic biology, cell biology and biochemistry				
Course	Contents	/ Syllabus:				
Unit 1		tures and functions of Bio-molecules:	8 hr			
		hydrates: classification, mono, di, oligo and polysaccharides. Lipids: fatty				
	acids,	simple, complex & derived lipids. Protein: Amino Acids Structure and				
		on, Protein Structure Hierarchy. Nucleic acids: nucleosides, nucleotides,				
TI 0		& RNA.	0.1			
Unit 2		ergetics:	8 hr			
	Energ	iew of principles of bioenergetics (free energy, enthalpy and entropy). y relationships between catabolic and anabolic pathways. Phosphoryl groupers and ATP, Free-energy change for ATP hydrolysis.				
Unit 3		polism:	8 hr			
CINC		lysis, Gluconeogenesis, Respiration and Introduction to the Citric Acid	O III			
	Cycle,	Electron Transport, Oxidative phosphorylation, Fatty Acid Catabolism acid oxidation, Protein Metabolism: The Urea Cycle	:			
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 DNA ation, Mechanism of replication. Telomeres, telomerase and end replication. of telomerase in aging and cancer.				
Unit 5	Centr	al Dogma	10 hr			
		cription, genetic code, reverse transcription, mRNA processing. Translation regulation, operons: Lac operon, Trp operon, transposons.	,			
Course	outcome					
CO1		After completion of the course, students will understand about the structure and function of biomolecules	2			
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 boo	oks					
1	1	nistry- L.Stryer, Third Edition				
2		nistry- Voet&Voet.				
	Diochen	1001 1000 1000				

3	Principles of Biochemistry- A.Lehninger, CBS Publishers and Distributors, 1987.		
Reference	ee 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	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	LTP	Credit
Course	Title	Bioprocess Engineering & Technology	3 0 0	3
Course o	bjective			
1	To pro	vide basic concepts of bioprocess engineering.		
2		rn engineering principles that can be applied to processes involve	ing cell or	
	enzymo			
3		n the basics of bioreactor design and operation control.		
4		lyze variety of bioprocess techniques and also conduct related exper	riments.	
5		erstand various unit operations in bioprocess.		
Pre-requ			1	
		ents are expected to have knowledge of basic biology, cell biology a memistry	ina	
Course (s/Syllabus:		
UNIT I		eduction to Bioprocess Technology		8 Hr.
	mode ferme ancill Diffe Medi comm	prical development of bioprocess technology, An overview of tradi- tern applications of biotechnological processes, General require entation processes, Basic design and construction of ferme laries, Main parameters for monitoring & control of fermentation rent raw materials used in fermentation industry and their pre- um for plant cell culture and animal cell culture, Medium nercial media for industrial fermentations-Plackettburman design ce methodology, simplex design.	ements of enter and processes, treatment, design of	
UNIT II	Stoic	hiometry of Cell growth		8 Hr.
	of recoeff analy	hiometry of Cell growth and product formation, elemental balance eduction of substrate and biomass, available electron balance icients of biomass and product formation, maintenance coefficients as is of microbial growth and product formation, oxygen consumption in aerobic cultures, thermodynamic efficiency of growth.	ces, yield Energetic	
UNIT II	Mass	Transfer in Bioreactors		8 Hr.
	oxygo oxygo proce	transfer includes transport phenomena in bioprocesses, Factors en transfer rate in bioreactors, Techniques for measurement of ven transfer coefficient, Fluid rheology and factors affecting esses, Flow Patterns in agitated tanks, Mechanism & Power requiring, Scale up of mixing systems.	olumetric bioreactor	
UNIT IV	Meta	bolic Regulation		10 Hr.
	anabo catab Conc acid, Insul	rent regulatory mechanisms involved in controlling the catalolic processes of microbes, Induction, nutritional repression olite repression, Crabtree effect, feedback inhibition and feedback rept of Overproduction of metabolites, Case studies on production Glutamic acid, Penicillin, Microbial Lipase and Protease, Regin, Interferons, Hepatitis Vaccines etc. Case studies should deal vovement, medium designs, process optimization technology.	epression, of Lactic	
	1			

Course outco	Centrifugation process and its equipments, Cell disruption, Aqueous Two-Phase Liquid Extraction. Adsorption process and its operations, Chromatography: Theory and mechanism, Scaling-up chromatography. Describe the underlying principles of main bioprocess unit operations like fermentation, downstream processing. Demonstrate Stoichiometry of Cell growth and product formation. Design or Select appropriate bioreactor models based upon bioproducts and cell lines and other process criteria. Develop a basic understanding of regulatory mechanisms involved in controlling the catabolic and anabolic processes of microbes. Acquire a basic understanding of various unit operations in bioprocess engineering.					
CO1 CO2 CO3 CO4	Describe the underlying principles of main bioprocess unit operations like fermentation, downstream processing. Demonstrate Stoichiometry of Cell growth and product formation. Design or Select appropriate bioreactor models based upon bioproducts and cell lines and other process criteria. Develop a basic understanding of regulatory mechanisms involved in controlling the catabolic and anabolic processes of microbes. Acquire a basic understanding of various unit operations in bioprocess					
CO2 CO3 CO4 CO5	fermentation, downstream processing. Demonstrate Stoichiometry of Cell growth and product formation. Design or Select appropriate bioreactor models based upon bioproducts and cell lines and other process criteria. Develop a basic understanding of regulatory mechanisms involved in controlling the catabolic and anabolic processes of microbes. Acquire a basic understanding of various unit operations in bioprocess					
CO3 CO4 CO5	Design or Select appropriate bioreactor models based upon bioproducts and cell lines and other process criteria. Develop a basic understanding of regulatory mechanisms involved in controlling the catabolic and anabolic processes of microbes. Acquire a basic understanding of various unit operations in bioprocess					
CO4 CO5	cell lines and other process criteria. Develop a basic understanding of regulatory mechanisms involved in controlling the catabolic and anabolic processes of microbes. Acquire a basic understanding of various unit operations in bioprocess					
CO5	controlling the catabolic and anabolic processes of microbes. Acquire a basic understanding of various unit operations in bioprocess					
Text books						
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.					
Reference Bo	ooks					
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/Reso	Journal/Research Paper Link:					
	curen a upor minis					

M. TECH FIRST YEAR						
Course Code	AMTCC0101 LT	P Credit				
Course Title	Research Process & Methodology 3 0 0					
Course Objec	tive:					
1	To explain the concept / fundamentals of research and their types					
2	To study the methods of research design and steps of research process	ch				
3	To explain the methods of data collection and procedure of samplin techniques	ng				
4	To analyze the data, apply the statistical techniques and understarthe concept of hypothesis testing	nd				
5	To study the types of research report and technical writing.					
	Basics of Statistics					
11c-requisites						
TINITO T	Course Contents / Syllabus INTRODUCTION TO RESEARCH	0 h a				
UNIT-I		8 hours				
	tive and motivation of research, types and approaches of research, lied vs. Fundamental, Quantitative vs. Qualitative, Conceptual					
• • • •	s versus Methodology, significance of research, criteria of good research	-				
UNIT-II	RESEARCH FORMULATION AND DESIGN	8 hours				
	and steps involved, Definition and necessity of research problem. I					
_	rature review, Locating relevant literature, Reliability of a source, W	-				
	the research problem, Literature Survey, Research Design, Metho					
design.	the research problem, Entertaine Survey, Research Design , Metho	ds of rescaren				
UNIT-III	DATA COLLECTION	8 hours				
	Data, accepts of method validation, Methods of Data Collection,					
	ondary data, sampling, need of sampling, sampling theory and Techr					
· •	different types of sample designs, ethical considerations in research.	1				
UNIT-IV	DATA ANALYSIS	8 hours				
Processing Opera appropriate statis statistical infere Visualization – M	ations, Data analysis, Types of analysis, Statistical techniques an stical technique, Hypothesis Testing, Data processing software (e.nce, Chi-Square Test, Analysis of variance(ANOVA) and contoring Research Experiments, hands-on with LaTeX.	d choosing an g. SPSS etc.), variance, Data				
UNIT-V	TECHNICAL WRITING AND REPORTING OF RESEARCH	8 hours				
Types of research report: Dissertation and Thesis, research paper, review article, short communication, conference presentation etc., Referencing and referencing styles, Research Journals, Indexing, citation of Journals and Impact factor, Types of Indexing-SCI/SCIE/ESCI/SCOPUS/DBLP/Google Scholar/UGC-CARE etc. Significance of conferences and their ranking, plagiarism, IPR- intellectual property rights and patent law, commercialization, copy right, royalty, trade related aspects of intellectual property rights (TRIPS); scholarly publishing-IMRAD concept and design of research paper, reproducibility and accountability.						
	ne: Upon completion of the course, the student will be able to					
CO 1	Explain concept / fundamentals for different types of research	K1				
CO 2	Apply relevant research Design technique	К3				
CO 3	Use appropriate Data Collection technique	К3				
CO 4	Evaluate statistical analysis which includes various parametric te and non-parametric test and ANOVA technique	est K5				

CO 5	Prepare research report and Publish ethically.	K6
Text books		
1. C. R. Ko	othari, Gaurav Garg, Research Methodology Methods and Techniques	, New Age
Internation	onal publishers, Third Edition.	
2. Ranjit K	umar, Research Methodology: A Step-by-Step Guide for Beginners,	2 nd Edition,
SAGE 20	005.	
3. Deepak C	hawla, NeenaSondhi, Research Methodology, Vikas Publication	
Reference Bo	oks	
1. Donald Co	oper & Pamela Schindler, Business Research Methods, TMGH, 9 th edition	on
2. Creswell,	John W., Research design: Qualitative, quantitative, and mixed methods a	pproaches
sage publications	5,2013	

M. TECH FIRST YEAR						
Course Code	AMTBT0151 LTP	Credit				
Course Title	Applied Biochemistry & Molecular Biology Lab 0 0 4	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	Experiment:					
Sr. No.	Name of Experiment	CO				
1	Quantitative estimation of amino acids by ninhydrin reaction.	1 2				
2	Quantitative estimation of proteins.	1 2				
3	To separate lipids with the help of thin layer chromatography (TLC).	1 2				
4	To verify the Lambert Beer's law with the help of UV absorption spectra of proteins.	1 2				
5	Protein purification by ammonium sulfate precipitation.	1 2				
6	Isolation of DNA and RNA from animal tissue and plant tissue.	1 2				
7	Gel electrophoretic analysis of various DNA and their restriction digests	1				
8	Transformation with plasmid and bacteriophage DNA	1 3				
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.					

Course Title Course objective: To understand the various concepts of microbial culturing. To learn the activation energy, volumetric oxygen transfer coefficient etc.! To Understand the principles and various pathways of enzyme production. Evaluate the concept of separation and purification of microbial produce. To understand the principles and various pathways of enzyme production. Evaluate the concept of separation and purification of microbial produce. To understand the process of fermentation. Pre-requisites: Students are expected to have knowledge of basic biology, cell biology and biochemistry Suggested list of Experiment: Sr. No. Name of Experiment Determination of kinetic parameters for batch cultivation of yeast under shake flask conditions. Determination of volumetric oxygen transfer coefficient (KLa) Determination of activation energy (Ea) of microbial strains. Process optimization for enzyme production using specific experimental design. Preparation of immobilized enzymes & cells and evaluation of kinetic parameters. Computational Design of Fermentative Process. Fermenter designing and the study of various parts of fermenter and their function for microbial cell culture. Fermenter designing and the study of various parts of fermenter and their function for microbial cell culture. Ethanol production for praymes Cellulase & Protease. Detanol production for molasses or starchy raw material. Fermentative production of Wine from grapes. Separation and purification of microorganisms from yogurt and cheese. Ethanol production of alpha amylase under solid & submerged conditions Fermentative production of alpha amylase under solid & submerged conditions To study the Scale-up and Sterilization in Bioreactors Lab Course Outcome: CO 1 Student will be able to understand the various concepts of microbial culturing. Student will be able to understand the principles and various pathways of enzyme production. Student will be able to evaluate the concept of separation and purification o	M. TECH FIRST YEAR						
Course objective: To understand the various concepts of microbial culturing. To learn the activation energy, volumetric oxygen transfer coefficient etc.! To Understand the principles and various pathways of enzyme production. Evaluate the concept of separation and purification of microbial produce. To understand the process of fermentation. Pre-requisites: Students are expected to have knowledge of basic biology, cell biology and biochemistry Suggested list of Experiment: Sr. No. Name of Experiment Determination of kinetic parameters for batch cultivation of yeast under shake flask conditions. Determination of volumetric oxygen transfer coefficient (KLa) Determination of activation energy (Ea) of microbial strains. Process optimization for enzyme production using specific experimental design. Preparation of immobilized enzymes & cells and evaluation of kinetic parameters. Computational Design of Fermentative Process. Fermenter designing and the study of various parts of fermenter and their function for microbial cell culture. Be Fermentative production of Penicillin by using Penicillumchrysogenum. Microbial production of enzymes Cellulase & Protease. Microbial production from molasses or starchy raw material. Fermentative production of Wine from grapes. Separation and purification of microorganisms from yogurt and cheese. Separation and purification of microorganisms from yogurt and cheese. Separation and purification of microorganisms from yogurt and cheese. To study the Scale-up and Sterilization in Bioreactors Lab Course Outcome: CO 1 Student will be able to understand the various concepts of microbial culturing. CO 2 Student will Understand the principles and various pathways of enzyme production. Student will be able to evaluate the concept of separation and purification of microbial produce.	Course Code	AMTBT0152	LTP	Credit			
To understand the various concepts of microbial culturing. To learn the activation energy, volumetric oxygen transfer coefficient etc.l To Understand the principles and various pathways of enzyme production. Evaluate the concept of separation and purification of microbial produce. To understand the process of fermentation. Pre-requisites: Students are expected to have knowledge of basic biology, cell biology and biochemistry Suggested list of Experiment: Sr. No. Name of Experiment Determination of kinetic parameters for batch cultivation of yeast under shake flask conditions. Determination of volumetric oxygen transfer coefficient (KLa) Determination of activation energy (Ea) of microbial strains. Process optimization for enzyme production using specific experimental design. Preparation of immobilized enzymes & cells and evaluation of kinetic parameters. Computational Design of Fermentative Process. Fermenter designing and the study of various parts of fermenter and their function for microbial cell culture. Fermentative production of Penicillin by using Penicillumchrysogenum. Microbial production for microbiases or starchy raw material. Fermentative production form molasses or starchy raw material. Fermentative production of Wine from grapes. Separation and purification of microorganisms from yogurt and cheese. Separation and purification of microorganisms from yogurt and cheese. Separation and purification of microorganisms from yogurt and cheese. Fermentative production of alpha amylase under solid & submerged conditions Protein profiling of fermentation broth through dialysis procedure. To study the Scale-up and Sterilization in Bioreactors Lab Course Outcome: Co 1 Student will be able to understand the various concepts of microbial culturing. Student will be able to evaluate the concept of separation and purification of microbial produce.	Course Title	Bioprocess Engineering & Technology Lab	0 0 4	2			
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Evaluate the concept of separation and purification of microbial produce.	2	To learn the activation energy, volumetric oxygen transfer coeffic	ient etc.l				
To understand the process of fermentation.	3	To Understand the principles and various pathways of enzyme pro	oduction.				
Students are expected to have knowledge of basic biology, cell biology and biochemistry	4	Evaluate the concept of separation and purification of microbial p	roduce.				
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Biochemistry Suggested list of Experiment : Sr. No. Name of Experiment Determination of kinetic parameters for batch cultivation of yeast under shake flask conditions. Determination of volumetric oxygen transfer coefficient (KLa) Determination of activation energy (Ea) of microbial strains. Process optimization for enzyme production using specific experimental design. Preparation of immobilized enzymes & cells and evaluation of kinetic parameters. Preparation of immobilized enzymes & cells and evaluation of kinetic parameters. Permenter designing and the study of various parts of fermenter and their function for microbial cell culture. Permentative production of Penicillin by using Peniciliumchrysogenum. Permentative production of enzymes Cellulase & Protease. Permentative production of enzymes Cellulase & Protease. Permentative production of microorganisms from yogurt and cheese. Permentative production of Wine from grapes. Permentative production of Wine from grapes. Permentative production of alpha amylase under solid & submerged conditions Pertentative production of alpha amylase under solid & submerged conditions Protein profiling of fermentation broth through dialysis procedure. Protein profiling of fermentation broth through dialysis procedure. To study the Scale-up and Sterilization in Bioreactors Lab Course Outcome: CO 1 Student will be able to understand the various concepts of microbial culturing. CO 2 Student will learn the activation energy, volumetric oxygen transfer coefficient etc. Student will Understand the principles and various pathways of enzyme production production. Student will be able to evaluate the concept of separation and purification of microbial produce. Production of produce Production of produce Production of produce Production of produce P	Pre-requisites:						
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CO 2 Student will learn the activation energy, volumetric oxygen transfer coefficient etc. CO 3 Student will Understand the principles and various pathways of enzyme production. CO 4 Student will be able to evaluate the concept of separation and purification of microbial produce.	Lab Course Ou	tcome:					
coefficient etc. Student will Understand the principles and various pathways of enzyme production. CO 4 Student will be able to evaluate the concept of separation and purification of microbial produce.	CO 1	Student will be able to understand the various concepts of microb	ial culturing.				
co 3 production. Student will be able to evaluate the concept of separation and purification of microbial produce.	CO 2	coefficient etc.					
microbial produce.	CO 3	production.	·				
CO 5 Student will be able to understand the process of fermentation.	CO 4		rification of				
*	CO 5	Student will be able to understand the process of fermentation.					

	M. TECH FIRST YEAR	
Course Code	AMTBT0111 LTP	Credit
Course Title	Immunology & Vaccine Technology 3 0 0	3
Course objective	:	
1	Learn 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	
4	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	<u> </u>	0 1
UNIT-I	Fundamental of Immune System	8 hr
	Fundamental concepts and anatomy of the immune system, Components of innate and acquired immunity, Humoral and Cell mediated immunity, Hematopoiesis, Antigens, immunogens, haptens, Major Histocompatibility Complex - MHC genes, MHC and immune responsiveness and disease susceptibility, HLA typing.	
UNIT-II	Immunological Processes	8 hr
	Immunoglobulins-basic structure, classes and subclasses of immunoglobulins, antigenic determinants, Multigene organization of immunoglobulin genes, Immunological basis of self —non-self discrimination; Kinetics of immune response, memory; B cell maturation, activation and differentiation; Generation of antibody diversity, Antigen processing and presentationendogenous antigens and exogenous antigens.	
UNIT-III	Basic Introduction to Vaccines	8 hr
	A short history of vaccination, Active and passive immunization, General immunization practices, Vaccination of immunocompromised hosts, Vaccination of human immunodeficiency virus infected persons, Vaccines, Live, killed, attenuated, subunit vaccines; Vaccine technology- Roleand properties of adjuvants, recombinant DNA and protein-based vaccines, plant-based vaccines, reverse vaccinology; Peptide vaccines, conjugate vaccines.	
UNIT-IV	Recent Advances in Vaccines	10 hr
	Licensed vaccines, Viral Vaccine (Poliovirus vaccine-inactivated & Live, Rabies vaccines Hepatitis A & B vaccines), Bacterial Vaccine (Anthrax vaccines, Cholera vaccines, Diphtheria toxoid), Parasitic vaccine (Malaria Vaccine).	
UNIT-V	Vaccine Industry (Production & Regulations)	10 hr
	The vaccine industry, Vaccine manufacturing, Evolution of adjuvants 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		

After completion of the course, students will understand the fundamentals of the immune system.	
They will learn about immunological processes.	
They will understand the different types of immunization and vaccines.	
Students will learn the different types of advanced vaccines.	
Students will be able to understand the vaccine industry and their production process.	
Kuby, RA Goldsby, Thomas J. Kindt, Barbara, A. Osborne Immunology, 6th Edition, Freeman, 2002.	
Brostoff J, Seaddin JK, Male D, Roitt IM., Clinical Immunology, 6th Edition, Gower Medical Publishing, 2002.	
Janeway et al., Immunobiology, 4th Edition, Current Biology publications., 1999. 4. Paul,	
Fundamental of Immunology, 4th edition, Lippencott Raven, 1999.	
Stanley A. Plotkin & Walter Orenstein & Paul A. Offit, Vaccines, 6th Edition 2013 BMA Medical Book Awards Highly Commended in Public Health! Elsevier Publication.	
Roitt's Essential Immunology. 11th ed. P. Delves, et al., ed., Blackwell Publishing, 2006.	
Bruce Alberts, Alexander Johnson, Julian Lewis, Martin Raff, Keith Roberts, Peter Walter. Molecular biology of the Cell. 4th ed. Garland publishing Inc, 2002	
h Paper Link:	
As suggested by concern subject faculty	
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Course 1 2 3 4 5 Pre-ree	se Code se Title e objective: To learn th	AMTBT0112 Quality Assurance and Quality Control	LTP	Credit
Course 1 2 3 4 5 Pre-ree Studer	e objective:		2 0 0	
1 2 3 4 5 Pre-rec			3 0 0	3
2 3 4 5 Pre-ree	To learn th			
3 4 5 Pre-ree		e basics of GLP		
5 Pre-ree Studer	To learn th	earn the manufacturing process and its audit.		
5 Pre-ree Studer	To underst	and the clinical trial process		
Pre-ree	To apply th	ne statistical tools to the various QC events		
Studer	To underst	and the tools and softwares used in QC and QA.		
	quisites:			
Course	nts are	expected to have knowledge of basic biology, cell biology and bio	chemistry	
	e Contents	/ Syllabus :		
UNIT	I Conc	ept of Quality control and quality assurance		
U1111 .		ept and evolution of quality control and quality assurance. Total Q	uality	
		gement, Philosophy of GMP and CGMP. Quality control laborator	•	
		nsibilities: GLP protocols on nonclinical testing control on animal		
		generation, integration and storage, standard test procedure, CPCSI	EA	
TINITO	guide			
UNIT		mentation practices and root cause analysis		
		ation of sample records, Quality review and batch release documented products, Good documentation practices, route cause analysis,		
		preventive action (CAPA), out of specifications (OOS) and out o		
	(OO)		i ticila	
UNIT	,	ept of Audits		
		al product quality review and parametric release, Audits, Preparation		
		conducting audit, Audit Analysis, Audit Report and Audit follow		
		y audits of manufacturing processes and facilities, audits of quality	y control.	
UNIT		Studies of Audit reports. ity agreements and risk management		
UNII			lag of	
		epts and management of contract manufacturing guidelines, princip ry risk management, ICH guidance for industry, BABE (bioavailab		
	1	uivalence) studies, post marketing surveillance, Pharmacovigilance	-	
UNIT		s and softwares in QC and QA	,	
		tical Tools for Quality Control and Precision, Tools of Problem So	olving and	
		nuous Improvement. Softwares for inspection and quality testing a		
		cations. concept of automation of procedure through Digital, IoT a		
		ons. Systematic approach to scale-up and technology transfer in		
	biote	chnology quality systems: Applications and challenges.		
Course	e outcome			
CO1	Recognize the importance of quality control and assurance and understand the concept of GMP, CGMP and GLP.			
CO2	Recognize the importance of good documentation practices and reframe the preventive actions.			
CO3	Analyse, develop, follow and audit the quality standards and guidelines, being			

CO4	Understand the contract guidelines to effectively manage the quality agreements.	
CO4		
CO5	Apply statistical tools and modern software to evaluate and ensure quality control, 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 Operating Procedures 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	

	M. TECH FIRST YEAR		
Course Code	AMTBT0113	LTP	Credit
Course Title	Applied Clinical Research	3 0 0	3
Course objective	/e:		
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 ethics 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 biobiochemistry	logy and	
Course Content	ts / Syllabus :		
UNIT I:	Introduction to clinical research		8 Hr.
	Basic pharmacology and drug development process, researchdefinition, Basic terminology used in clinical research, studies, Introduction to pharmacoeconomics, Types of clinical tr blinding, double blinding, open access, randomized trials and their interventional study, Good Clinical Practices, Types and Scope Research.	preclinical ials, single examples,	
UNIT II:	II: Clinical trials		8 Hr.
	New drug discovery process- purpose, main steps involved in discovery process, timelines of each steps, advantages and purposteps, Pre clinical toxicology: General principles, Systemic (Single dose and repeat dose toxicity studies), Carcinogenicity, McTeratogenicity, Reproductive toxicity, Local toxicity, Genotoxic toxicity requirements, Phase-I, II, III, IV trials: Introduction and Various phases of clinical trials, Post Marketing surveillance, Principles of sampling, Inclusion and exclusion criteria, Mallocation and randomization, Informed consent process in brief treatment outcome, Termination of trial, Safety monitoring in clinical	ses of each toxicology itagenicity, ity, animal designing, methods & Iethods of monitoring,	
UNIT III:	Ethics & Regulations in Clinical research		8 Hr.
	Ethical Theories and Foundations, Ethics Review Committee and Consent Process, Integrity & Misconduct in Clinical Research, unet thalidomide tragedy, Conflicts of Interest, Evolution and Regulations in Clinical Research, Study of various clinical trials or ongoing), Patents US Regulatory Structure, Clinical Trial Applindia Import & Export of Drug in India, Investigational New Drug application (IND), Application (NDA), Abbreviated New Drug Application (ANDA), Approval Activities, PMS, FDA Audits and Inspections EU Affairs, EMEA Organization and Function, INDIAN Regulato Schedule Y- Rules and Regulations.	hical trials, History of (completed plication in New Drug , Post Drug Regulatory	
UNIT IV:	Principles of controlled clinical trials		10 Hr.

UNIT V	:	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. Biostatistics and data management 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	10 Hr.	
Course	outcom	pregnancy condition.		
CO1		ibe the process of drug development and principles of clinical pharmacology.		
CO2	Devel	op a clear understanding of why ethics are important in clinical research and be ar with the regulatory practices in place to protect both the researcher and the		
CO3		tively manage the regulatory process from Innovation →Discovery → oval → 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		ibe the various types of clinical studies and the methods used to choose the priate design, evaluation and interpretation of clinical trial results.		
Text boo	oks			
1	Basic	and Clinical Pharmacology, Prentice hall, International, Katzung, B.G.		
2		cal Pharmacology, Scientific book agency, Laurence, DR and Bennet PN.		
3		eal pharmacokinetics, Pub. Springer Verlab, Dr. D.R Krishna, V. Klotz		
4		ngton Pharmaceutical Sciences, Lippincott, Williams and Wilkins		
5	Drug interaction, Kven Stockley. Hamsten			
Referen				
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		
L				

Course objective: To learn about the mass balance involved in waste water treatment			M. TECH FIRST YEAR		
Course objective: 1 To learn about the mass balance involved in waste water treatment 2 To understand the anaerobic treatment process. 3 To learn about the various chemical and physical processes involved in waste water treatment. 4 To understand the basic of phosphorus and nitrogen removal 5 To Learn about the recycling of waste Pre-requisites: Students are expected to have knowledge of basic biology, cell biology and biochemistry Course Contents / Syllabus: UNIT I- ACTIVATED SLUDGE PROCESS-PROCESS ANALYSIS AND SELECTION Characteristics of Activated Sludge (aerobic and anaerobic); Analysis of Data Mass Balance Analysis. Reactors used in waste water treatment- Up Flow Anaerobic Sludge Blanket (UASB), Two-stage, Aerobic Unit Tank System (TSUSystem, Route Zone Treatment, Submerged Aerobic Fixed Film (SAFF) Reactor, and Fluidized Aerobic Bioreactor (FAB). UNIT II- AEROBIC FIXED-FILM & ANAEROBIC TREATMENT PROCESSES Biofilm process considerations; Tricking Filters and Biological Towers; Rotating Biological Contactors; Granular Media Filters; Fluidized Bed & Circulating Bed-Biofilm reactors, Hybrid Biofilm/suspended growth processes. Anaerobic Processes: Methanogenesis, process chemistry and microbiology; process kinetics and factors for the design of anaerobic digestors. UNIT III- ADVANCED WASTE WATER TREATMENT Technologies used in advanced treatment-Classification of technologies; Removal of Colloids and suspended particles-Depth Filtration, Surface Filtration, Membrane Filtration Absorption, Ion Exchange, Advanced oxidation process, Activated Carbon, Air Stripping, Heavy Metals Removal, Steam Stripping, Chemical Precipitation, and Electrolysis. UNIT IV- BIOLOGICAL PHOSPHORUS AND NITROGEN REMOVAL Nitrification & Denitrification Processes: Biochemistry and Physiology of Nitrifying Bacteria; Common process considerations; One sludge versus two sludge nitrification, Ohysiology of Denitrifying Bacteria; Tertiary Denitrification; One-sludge denitrification, Normal Phosphorus Uptake into Biomass; Mechanism	Course	Code	AMTBT0114	LTP	Credit
To learn about the mass balance involved in waste water treatment To understand the anaerobic treatment process. To learn about the various chemical and physical processes involved in waste water treatment. To understand the basic of phosphorus and nitrogen removal To Learn about the recycling of waste Pre-requisites: Students are expected to have knowledge of basic biology, cell biology and biochemistry Course Contents / Syllabus: UNIT I- ACTIVATED SLUDGE PROCESS-PROCESS ANALYSIS AND SELECTION Characteristics of Activated Sludge (aerobic and anaerobic); Analysis of Data-Mass Balance Analysis, Reactors used in waste water treatment- Up Flow Anaerobic Sludge Blanket (UASB), Two-stage, Aerobic UNI Tank System (TSUSystem, Route Zone Treatment, Submerged Aerobic Fixed Film (SAFF) Reactor, and Fluidized Aerobic Bioreactor (FAB). UNIT II- AEROBIC FIXED-FILM & ANAEROBIC TREATMENT PROCESSES Biofilm process considerations; Trickling Filters and Biological Towers; Rotating Biological Contactors; Granular – Media Filters; Fluidized Bed & Circulating Bed-Biofilm reactors. Hybrid Biofilm/suspended growth processes, Anaerobic Processes: Methanogenesis, process chemistry and microbiology; process kinetics and factors for the design of anaerobic digestors. UNIT III- ADVANCED WASTE WATER TREATMENT Technologies used in advanced treatment-Classification of technologies; Removal of Colloids and suspended particles-Depth Filtration, Surface Filtration, Membrane Filtration Absorption, Ion Exchange, Advanced oxidation process, Activated Carbon, Air Stripping, Heavy Metals Removal, Steam Stripping, Chemical Precipitation, and Electrolysis. UNIT IV- BIOLOGICAL PHOSPHORUS AND NITROGEN REMOVAL Nitrification, One-sludge dentirification, Normal Phosphorus Uptake into Biomass; Mechanism for Biological Phosphorus Removal; Enhanced Biological Phosphorus Removal; Enhanced Biological Phosphorus Removal by Bacteria and Algae. UNIT V ENVIRONMENTAL CONCERNS & RECYCLING OF WASTES Environmental regulations and technology-Regulat	Course	Title	Biological Treatment of Waste Water	3 0 0	3
To learn about the mass balance involved in waste water treatment To understand the anaerobic treatment process. To learn about the various chemical and physical processes involved in waste water treatment. To understand the basic of phosphorus and nitrogen removal To Learn about the recycling of waste Pre-requisites: Students are expected to have knowledge of basic biology, cell biology and biochemistry Course Contents / Syllabus: UNIT 1- ACTIVATED SLUDGE PROCESS-PROCESS ANALYSIS AND SELECTION Characteristics of Activated Sludge (aerobic and anaerobic); Analysis of Data-Mass Balance Analysis. Reactors used in waste water treatment- Up Flow Anaerobic Sludge Blanket (UASB), Two-stage, Aerobic UNI Tank System (TSUSystem, Route Zone Treatment, Submerged Aerobic Fixed Film (SAFF) Reactor, and Fluidized Aerobic Bioreactor (FAB). UNIT II- AEROBIC FIXED-FILM & ANAEROBIC TREATMENT PROCESSES Biofilm process considerations; Trickling Filters and Biological Towers; Rotating Biological Contactors; Granular – Media Filters; Fluidized – Bed & Circulating Bed-Biofilm reactors. Hybrid Biofilm/suspended growth processes. Anaerobic Processes: Methanogenesis, process chemistry and microbiology; process kinetics and factors for the design of anaerobic digestors. UNIT II- ADVANCED WASTE WATER TREATMENT Technologies used in advanced treatment-Classification of technologies; Removal of Colloids and suspended particles-Depth Filtration, Surface Filtration, Membrane Filtration Absorption, Ion Exchange, Advanced oxidation process, Activated Carbon, Air Stripping, Heavy Metals Removal, Steam Stripping, Chemical Precipitation, and Electrolysis. UNIT IV- BIOLOGICAL PHOSPHORUS AND NITROGEN REMOVAL Nitrification, One-shudge dentirification, Normal Phosphorus Uptake into Biomass; Mechanism for Biological Phosphorus Removal; Enhanced Biological Phosphorus Removal; Enhanced Biological Phosphorus Removal; Bacteria; Tertiary Dentirification, One-shudge dentirification, Normal Phosphorus Uptake into Biomass; Mechanism for Biolo					
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Students are expected to have knowledge of basic biology, cell biology and biochemistry	4	To under	stand the basic of phosphorus and nitrogen removal		
Students are expected to have knowledge of basic biology, cell biology and biochemistry	5	To Learn	about the recycling of waste		
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UNIT V ENVIRONMENTAL CONCERNS & RECYCLING OF WASTES Environmental regulations and technology- Regulatory Concerns, Technology; Laws, regulations and permits, Air, Water, Solid Waste, Environmental Auditing, National Environmental Policy act, Occupational Safety and Health Act (OSHA), Storm Water Regulations; Technology (waste water); Recycling of		Nitrif sludg Denit Biom	Tying Bacteria; Common process considerations; One sludge versue nitrification. Physiology of Denitrifying Bacteria; Tertiary crification; One-sludge denitrification, Normal Phosphorus Uptakass; Mechanism for Biological Phosphorus Removal; Enhanced I	as two te into	
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r		Laws Audit Act (, regulations and permits, Air, Water, Solid Waste, Environmentating, National Environmental Policy act, Occupational Safety and	l Health	
Course outcome	Course o	utcome			

CO1	After completing the course students will able to perform mass balance for the bioreactor			
CO2	After completing the course students will able to design an anaerobic system			
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.			
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	oks			
1	Introduction to Waste Water Treatment- R. S. Ramalho, Academic Press.			
2	Environmental Biotechnology, B.C. Bhattacharya &Ritu Banerjee, Oxford Press, 2007.			
Journal/Resea	arch Paper Link:			
	As suggested by concern subject faculty			

		M. TECH FIRST YEAR		
Course Code		AMTBT0115	LTP	Credit
Course Title		Nano Biotechnology & Toxicology	3 0 0	3
Course ob	jective:			
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 char techniques involved in nanotechnology	acterization	
4		To understand the applications of nanotechnology		
5		To learn the toxicology of nanomaterials		
Pre-requis	sites:			
		Students are expected to have knowledge of basic biology, cell and biotech	l biology	
Course Co	ontents / S	Syllabus:		
UNIT-I		eduction to Nanobiotechnology:		8hr.
		nition of Nanobiotechnology, History, Origin, Fundamental		
		om-up versus Top-down approaches, Discussion on Nanof ent research, Tool and Techniques, Applications and Implication	· ·	
		of sprication.	is allu	
UNIT-II		omaterials and Nanoparticles:		8hr.
		on nanotubes and related structures, Properties, Synthesis, Ap	nlications	om.
		y balls, Nanoparticles types and their synthesis, Application		
		r and Zinc oxide nanoparticles, Interaction of nanopart		
	biom	embrane and genes.		
UNIT-III		Nanocharecterization Tool and Techniques:		
		visible spectrophotometry, Fourier transform infrared spectrosco		
		ning Electron Microscopy (SEM), Scanning tunnelling r		
	`	(I), Transmission electron microscopy (TEM), Atomic force r (I), Zeta Potential size analyser etc.	nicroscopy	
UNIT-IV	`	Nanomedicine and Sensor Technology:		10 hr.
01111-11		delivery tools, Bioavailability, Nano imaging agents, Protein	and partida	TO III.
		ery (Cancer and Surgery) and Nano sensors technology with app		
UNIT-V		cology:		10 hr.
01(11)		nition of toxicology, History and origin of toxicology, Pri	inciples of	10 111 0
		cology, Concept of Toxicology, Types of toxicology, Na		
		ity evaluation mechanism as in vitro, Nanomaterial toxicity		
		anism as in vivo, Assessment of nanoparticles toxicity: A	•	
		otoxicity, Genotoxicity, Hepatotoxicity, Neurotoxicity, Nep	hrotoxicity	
Carrage	etc.)			
Course ou		mplating this course the students will be able to learn the for	undomentale	
CO1		After completing this course, the students will be able to learn the fundamentals oncepts of nanotechnology		
		npleting this course, the students will be able to ability for under	retanding	
CO2		rentiate the various nano materials	istanumg	
After completing this course, the students will be able to understand the p		ne principal		
behind the different characterize				

CO4	After completing this course, the students will be able to get insight the application of nanotechnology in drug delivery system		
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 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	LTP	Credit
Course	Title	Industrial Biotechnological Products	3 0 0	3
Course o	bjective	•		
1	To lear	n about the different media for the growth of microbes		
2	To und	erstand the production process of Primary and Secondary meta-	abolites	
3	To desi	gn and deliver useful modern biotechnology products to the S	ociety.	
4		Understand the methods to obtain enzymes of industrial importance and in general		
		product development Research & Development		
5		erstand the manufacturing of various organic and alcoholic pro-	oducts	
Pre-requ			1 1 1	
		ents are expected to have knowledge of basic biology, migy 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, Selectors optimization techniques.	-	
Unit II	Prod	uction of Metabolites		8 hr.
	ethan wine	ess technology for the production of various Products: Prima ol, citric acid, vinegar and amino acid; Production of alcoh and beer; Secondary metabolites: Antibiotics; Process tech action of microbial biomass.	olic beverages:	
Unit III	Biop	roducts		8 hr.
	Produ	duction and production of secondary metabolites with sorticion of bioplastics (PHB, PHA), bioinsecticides, plymers, Biofertilizers and biological weapons with reference to	bioherbicides,	
Unit-IV	Prod	uction of industrially important enzymes		8 hr.
	ferme Prote	action of industrially important enzymes: Solid state fermentate entation, Extraction, Purification and characterization of industrial 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 selection importance from plants and animal sources. Procestation of Yoghurt, acidophilus milk, Koumis, kefir, cheese, bage, vinegar and oriental fermented food. Food packaging wed in the commercially important food processing methods.	ess involved in oread, alcoholic	
Course o	utcome			
CO1		p key practical skills in fermenting biotechnology and be ons and commercial opportunities in fermentation-based biote		
CO2	Increase their understanding that 'industrial biotechnology' is based on using machines to control the growth of microorganisms			
CO2	macmi	Develop knowledge of a variety of fermentation strategies		
CO2				

CO5	Explore the biological and technological principles which govern actual and potential bio-business	
Text boo	ks	
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.	
Reference	ee 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.	
Journal/Research Paper Link:		
	As suggested by concern subject faculty	

	M. 1	TECH FIRST YEAR	
Course C	de AMTBT0201	LT	P Credit
Course Ti	le Bioinformatics	3 0	0 3
Course obje	ntivo.		
1	To learn the various online	datahases	
2		r analysing various methods of sequence	
_	alignment	a unarysing various methods of sequence	
3	To understand the phyloger	netic analysis and related conclusions	
4	To understand the concepts	of system biology.	
5	To understand the various r	methods of genome sequencing.	
Pre-requisi	es:		
•	Students are expected to ha biochemistry	ve knowledge of basic biology, cell biology and	nd
Course Cor	ents / Syllabus		
UNIT I	Biological Databases		8 Hr.
	GenBank, DDBJ, EMBL, I Specialized databases: Pubme	projects. Primary and Secondary Database PIR, Uniprot-KB, SWISS-PROT, TrEMING, OMIM, Medical databases, KEGG, Eat NCBI, EBI, TIGR, SANGER. Overview bioinformatics exercises.	BL. ST
UNIT II	Sequence Alignment		8 Hr.
	scores, substitution matrices, significance of Alignments, Pai Aspect of Multiple Sequence	nent, Optimal Alignment Methods, Substitut PAM, BLOSUM, Gap penalties, Statistic r wise sequence alignment algorithms, Practic Alignment, Progressive and Iterative Alignments base similarity searching, FASTA, BLAST, PHI-BLAST.	cal cal ent
UNIT III	Phylogenetic Analysis and Prim	er Design	8 Hr.
	Phylogenetic Data Analysis, Ph	analysis, Elements of phylogenetic Modeylogenetic Tree -construction steps and Build to phylogenetic analysis, Restriction mappitor and OMIGA, gene construction KIT, Vec 2, REBASE); Primer design – Primer des 3)	ing ng, ctor
UNIT IV	System Biology		8 Hr.
	introduction to different data ty Post translational modification,	ogy, application related to bioinformatives: PRIDE (Protein Identifications) database P-P interaction, Rotameric Structures of Protein IDNA Forms (DNA Sequence Effects	ses, ins
UNIT V	Genome Sequence Analysis		ı

	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	nce 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.		
Journal	/Research Paper Link:		
	As suggested by concern subject faculty		
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		M. TECH FIRST YEAR		
Course (Code	AMTBT0202	LTP	Credit
Course 7	Γitle	Entrepreneurship, IPR & Biosafety	3 0 0	3
Course ob	jectiv	e:		
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-requis	sites:			
		Students are expected to have knowledge of basic biology, cell bio biochemistry	ology and	
Course Co	ontents	s / Syllabus		
Unit I	Acco	ounting and Finance:		8 Hr.
	Appr prope bank starti in ac	ng decision on starting a venture; Assessment of feasibility of a network a bank for a loan; Sources of financial assistance; Making osal/Plan for seeking loans from financial institution and Banks; I for capital expenditure and for working; Statutory and legal requiring a company/venture; Budget planning and cash flow managem counting practices: concepts of balance sheet, P&L account, and dekeeping; Estimation of income, expenditure, profit, income tax etc.	a business Funds from rements for ent; Basics buble entry	
Unit II	Mar	keting:		8 Hr.
	cond custo brand Prom finan comp	ssment of market demand for potential product(s) of interestions, segments; Prediction of market changes; Identifying omers including gaps in the market, packaging the product; Market ding issues; Developing distribution channels; Pricing/Policies/Contoion/Advertising; Services Marketing Negotiations/Stratesticiers, bankers etc.; With government/law enforcement authority panies/Institutions for technology transfer; Dispute resolution skill comment/changes; Crisis/ Avoiding/Managing; Broader visiting	needs of et linkages, empetition; gy: With ties; With s; External	
Unit III	Info	rmation Technology:		8 Hr.
	perfo setup Mana	to use IT for business administration; Use of IT in improving primance; Available software for better financial management; on, management. Human Resource Development (HRD): Leaders agerial skills; Organization structure, pros & cons of different in building, teamwork; Appraisal; Rewards in small scale set up.	E-business ship skills;	
Unit IV	IPR:			8 Hr.
	& R Indic of IF Studi	oduction to Intellectual Property, Types of IP: Patents, Trademarks, Related Rights, Industrial Design, Traditional Knowledge, Gerations, Protection of New GMOs; International framework for the P IP as a factor in R&D IPs of relevance to Biotechnology and ies; Introduction to History of GATT, WTO, WIPO and TRIPS, at application and Process of Technology Transfer	eographical protection few Case	
Unit V	Bios	afety:		
	1	v		

	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 or	utcome	
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 book	ks	
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 Referen	nce 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)	

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

	M. TECH FIRST YEAR		
Course Code	AMTBT0251	LTP	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	g	
Pre-requisites:			
	Students are expected to have knowledge of basic biology, cell bio biochemistry	logy and	
Course Contents	 		
1	To perform pair wise local and global sequence alignment for any and DNA sequences.	two proteins	
2	To perform multiple sequence alignment for any five sequences ar Phylogenetic relationship among them.	nd predicts the	
3	Phylogenetic Analysis using PHYLIP - Rooted trees and Unrooted	trees	
4	To predict secondary structure for any given protein sequence usin Fasman, GOR and Neural network algorithms.	g Chou-	
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 of genes and number of proteins encoded on Chr-Y.	e size, number	
7	Homology Modelling using Modeller		
8	To find out the RMSD value from any two-protein structure alignr	nent.	
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 macromo cells	lecules of the	
CO3	They will understand the phylogenetic analysis and related conclusions	sions	
CO4	The students will learn the use of various tools for molecular analy	rsis	
CO5	The students will understand the various methods for macromolecus equencing	ılar	

M. TECH FIRST YEAR		
Course Code	AMTBT0252 L T P	Credit
Course Title	Entrepreneurship, IPR & Biosafety Lab 0 0 4	2
Course objective	:	
1	To make students aware of the process of patent registration.	
2	To learn to design Biosafety lab.	
3	To develop the entrepreneur and marketing skills.	
Pre-requisites:		
Students are	expected to have knowledge of basic biology, cell biology and biochemistry	
Course Contents	s / Syllabus	
1	Demonstration of Procedure for patent registration in India	
2	Writing a Patent Application	
3	3 Microbiological risk assessment	
4	Basic laboratories – Biosafety Levels 1 and 2 Basic laboratories – Biosafety Levels 1 and 2 Code of practice Laboratory design and facilities	
5	Laboratory equipment Health and medical surveillance	
6	6 Biosafety practices and procedures	
7	Development of project proposals - SWOT analysis	
8	SWOT analysis of selected enterprise	
9	Practical on developing distribution channels; Pricing/Policies/Competition; Promotion/Advertising through the use of social Media	
10	Preparation of Balance Sheet	
Course outcome	,	
CO1	Students will be able to understand the process of establishing Biosafety labs.	
CO2	Students will learn through demonstration the process Patent Registration.	
CO3	They will develop the skills of marketing and entrepreneur.	

	M. TECH FIRST YEAR	
Course Code	AMTBT0211 L T P	Credit
Course Title	Genetic Engineering 3 0 0	3
Course objectiv	۵۰	
Course objectiv	It is intended to impart basic undergraduate-level knowledge in the area of	2
1	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.	1
5	To understand the DNA sequencing methods	
Pre-requisites:		
Students are expe	ected to have knowledge of basic biology, cell biology and biochemistry	1
Course Content	s / Syllabus	
UNIT-I	Molecular Tools	8 Hr.
	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 outcom	ne	
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 gene technology [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 Boo		
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/Resea	rch Paper Link:	
	As suggested by concern subject faculty	

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	M. TECH FIRST YEAR		
Course Code	AMTBT0212	LTP	Credit
Course Title	Applied Food Biotechnology	3 0 0	3
Course objective	۵۰		
1	To learn about the various microbiological examination of food born diseases	ls 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	1.	
4	To learn about the industrial production of various food products		
5	To learn the methods of production of vitamins and enzymes.		
Pre-requisites:	5		
Î	Students are expected to have knowledge of basic biology, cell biochemistry	ology and	
Course Content	s / Syllabus		
Unit I	Food Biotechnology		8 Hr.
	Introduction & Applications; Methods for the microbiological exof water and foods; Control of Microbiological quality and sa borne illnesses and diseases; Microbial cultures for food ferment maintenance, strain development	fety; Food	
Unit II	Biosensors in food technology		8 Hr.
	Starter cultures—types, designing and development, micro encapsupackaging, scopes and challenge; Development and formulation products such as probiotic foods. Nutrigenomics-concept, significance and relevance. Biosensors and novel tools and their in food science & Technology	n of novel working,	
Unit III	GM foods		8 Hr.
	Introduction and controversies related to GMOs. Ethical issues 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 percept foods. IPR. GMO Act—2004. New products and processes in vaccommodities including plant and animal products.	on, release ed aspects; ion 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 a rum, vodka), glycerol; Propagation of baker's yeasts; Ferme products such as cheese, yoghurt, sweet curd, paneer, so Fermented pickles.	s whiskey, ented dairy	
Unit V	Industrial Food Biotechnology II		10 Hr.
	Industrial production of important primary and secondary metabas antibiotic, vitamins, biosurfantants, polysaccharides. Enzyme in food industry. Advantages and constraints of immobilized en microbial 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 Book	is S
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 LTP	Credit
Course Title	Molecular Modelling & Industrial Application 3 0 0	3
Course objectiv		
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:		
	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	Energy Minimisation and Computer Simulation	8 Hr.
	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 Book	XS .			
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 LTP	Credit	
Course Title	Bioreactor Analysis and Design 3 0 0	3	
Course objectiv			
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 outcome	(geometry based rules).	
CO1	After completing the course students will able to design the bioreactor system	
	After completing the course students will able to illustrate the control	
CO2	involved in bioreactor system	
CO3	After completing the course students will able to identify the various types of	
COS	bioreactor processes	
CO4	After completing the course students will able to analyse the reactor	
	dynamics After completing the course students will able to evaluate the design aspect	
CO5	and safety issues associated with reactor system.	
Text books		
1	Moser, Anton, Bioprocess Technology: Kinetics and Reactors, Springer	
1	Verlag, 1988.	
2	Bailey J.E. &Ollis, D.F. Biochemical Engineering Fundamentals, 2nd ed., McGraw	
2	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 Book		
1	Biochemical Engineering and Biotechnology Handbook" 2nd Ed.,.Atkinson &F.Mavituna, Stockton Press (1991).	
	Biorector Design & Product Yield", BIOTOL series, Butterworth -	
2	Heinemann (1992).	
2	Principles of fermentation technology" - F Stanbury and A Whitaker,	
3	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/Resear		
	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 objective			
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	s / Syllabus		
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.		
	BIOREACTORS, STERILIZATION, SENSORS AND		
UNIT III-	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	
	industry, Pharmaceuticals, Biochemical applications: Role of soluble and	
	immobilized enzymes in the synthesis and production of amino acids and	
	chiral compounds; use of enzymes as detergents. Molecular Imprinting;	
	Enzyme engineering: In vitro approaches to improve functional efficiency;	
	Recombinant enzymes, Case study.	
Course outcome		
CO1	Describe the fundamentals of enzyme properties, nomenclatures, characteristics and mechanisms & plot graphs based on kinetics data.	
CO2	Demonstrate metabolism, stoichiometry and microbial growth kinetics.	
CO3	Perform bioreactor operations as applicable in bioprocess industries.	
CO4	Discuss various separation and purification process of fermentation products.	
CO5	Predict the current and future trends of applying enzyme technology for the commercialization purpose of biotechnological products.	
Text books		
1	Michael Shuler and FikretKargi. "Bioprocess Engineering: Basic Concepts", 2nd Edition, Prentice Hall, and Englewood Cliffs, NJ, 2002.	
2	Pauline Doran. "Bioprocess engineering principles", Academic Press, 1995.	
3	Colin Ratledge, Bjorn Kristiansen, "Basic Biotechnology", 2nd Edition, Cambridge University Press, 2001.	
Reference Book	s	
1	Roger Harrison et al., "Bioseparation Science and Engineering", Oxford University Press, 2003.	
2	Harrison R.G. Todd P., Rudge S.R. "Bioseparation Science and Engineering", Oxford Press 2003.	
Journal/Researc	ch Paper Link:	
	As suggested by concern subject faculty	

M. TECH FIRST YEAR			
Course Code	AMTBT0216 LTP	Credit	
Course Title	Applied Bioenergy 3 0 0	3	
Course objective			
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:			
	Students are expected to have knowledge of basic biology, cell biology and biochemistry		
Course Contents	/ 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 Books		
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/Researc	h Paper Link:	
	As suggested by concern subject faculty	

		M. TECH FIRST YEAR	
Course	Code	AMTBT0217 L T P	Credit
Course	Title	Cell & Tissue Culture Techniques 3 0 0	3
Course o	bjectiv		
1		To learn the basics of animal cell culturing technique.	
2		To understand the various methods and advancements of culture techniques.	
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 and tissu culture.	e
Pre-requ	isites:		
		Students are expected to have knowledge of basic cell and molecular biology	'.
Course C		s / Syllabus	
Unit 1	Cell &	a Tissue Culture Technology Basics	8 hr
	Physic Tempe	cell culture techniques, Types of cell culture media; Ingredients of media ochemical properties; CO ₂ and bicarbonates; Buffering; Oxygen; Osmolarity erature; Surface tension and foaming; Balance salt solutions; Antibiotics in supplements;	;
Unit 2	Metho	ods of Cell & Tissue Culture	8 hr
	fibrobl Trypsi culture	ent tissue culture techniques; Types of primary culture; Chicken embryo last culture; Chicken liver and kidney culture; Secondary culture nization; Cell separation; Continuous cell lines; Suspension culture; Organ e etc.; Behaviour of cells in culture conditions: division, growth pattern colism of estimation of cell number; Development of cell lines	;
Unit 3	Applio	cations of Cell and Tissue Culture Technique	8 hr
	scale panimal	loning and selection; Transfection and transformation of cells; Commercial production of animal cells, stem cells and their application; Application of cell culture for <i>in vitro</i> testing of drugs; Testing of toxicity of environmenta ants in cell culture; Application of cell culture technology in production of and animal viral vaccines and pharmaceutical proteins, Green Meat, Organing	f l f
Unit 4	Plant	Cell & Tissue Culture Basics	10 hr
	embry improv disease and cl	mentals of plant tissue culture, plant regeneration: organogenesis. Somation organogenesis; somaclonal variation, its genetic basis and application in crop vement. Cell/callus line selection for resistance to herbicide, stress and es.: Isolation, culture and plant regeneration, protoplast fusion, identification haracterization of somatic hybrids., Field techniques for propagation of crated plants.	1
Unit 5	Techn	iques of Plant Cell & Tissue Culture	10 hr
	B5, S parame	nt selection, sterilization and inoculation; Various media preparations; MS H PC L2; Callus and cell suspension culture; Induction and growth eters; Chromosomal variability in callus culture. Plant regeneration from o, meristem and callus culture. Androgenesis: Anther and pollen culture.	1

Course outcome:			
CO1	After completion of the course, students will learn the basics of animal cell culturing.		
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 boo	ks		
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.		
Reference	ee 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		
Journal/	Research Paper Link:		
	As suggested by concern subject faculty		
3	M.K. Razda Plants from Test Tubes: An Introduction to Micropropagation by LydianeKyte Journal/Research Paper Link:		

M. TECH FIRST YEAR			
Course Code	AMTBT0218	TP	Credit
Course Title	Diagnostic Techniques in Biotechnology 3	3 0 0	3
Course objective	e:		
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:	,		
	Students are expected to have knowledge of basic biology, cell biolog biochemistry	gy and	
Course Contents	s / Syllabus		
Unit I	Analytical Methods		8 hr
	Volumetric analysis, Balancing & Weighing, Concept of solute & s Units of measurement. Specimen Collection & Processing: Specollection (Blood, urine, spinal fluid, saliva synovial fluid, Amniotic Preservation, transportation	ecimen	
Unit II	Clinical Enzymology		8 hr
	Principle of diagnostic enzymology, Digestive enzyme, Miscell enzyme. General Function Tests: Liver function test, Cardiac Function Renal Function Test, Thyroid Function test, Reproductive englishment function test	on Test,	
Unit III	Immunodiagnostics		8 hr
	Introduction, Antigen-Antibody Reactions, Conjugation Tech Antibody Production, Enzymes and Signal Amplification Sy Separation and Solid-Phase Systems, Studies related to bacterial, vi parasitic infections.		
Unit IV	Product Development		10 hr
	Immunoassay Classification and Commercial Technologies, Development, Evaluation, and Validation, Reagent Formulations and Life Evaluation, Data Analysis, Documentation, Registration Diagnostics Start-ups.	d Shelf	
Unit V	DNA based diagnostics		10 hr
	PCRRT-PCR, qPCR, Hot start PCR, Nested PCR), RFLP, Microarrays, FISH, In-situ hybridization, Studies related to bacteria and parasitic infections, Cell based diagnostics: Antibody market Markers, FACS, HLA typing, Bioassays, Viral DNA detection using kits and PCR	ıl, viral rs, CD	
Course outcome		·	
CO1	The students will learn the basics of diagnostic techniques.		
CO2	The students will understand the different enzymes and related test m	ethods.	
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 Books	s	
1	Diagnostic Microbiology, Balley& Scott's.	
2	Tietz Text book of Clinical Biochemistry, Burtis& Ashwood. 6. The Science of Laboratory Diagnosis, Crocker Burnett.	
Journal/Researc	ch Paper Link:	
	As suggested by concern subject faculty	

Course		M. TECH FIRST YEAR AMTBT0219	LTP	Credit
Course		3-D Printing Technology	3 0 0	3
			3 0 0	3
Course of				
4		know the fundamentals of RP Systems & its evolution and the Prociation of RP Systems with 3D modelling & Mesh	ocess in RP	K1,K2
2	Able to	know the RP Systems, Process, Materials & Classifications		K3, K4
3	format,	o know and working with Mesh File & their formats like STL for OBJ formats. Conversion to Mesh files, their properties, operations & defects		K3, K4
4	Able to	know the applications of RP Systems in various Fields		K3, K4
Pre-requi	sites:			
		ng of Information Technology.		
Course C	ontents	/ Syllabus		
UNIT-I	Intro	duction:		4 hours
	differ	rical Developments, Fundamentals of RP Systems and its Classicent basis, Rapid Prototyping Process Chains, 3D Modelling ration, Data Conversion and Transmission.		
UNIT-II	RP S	ystems:		12 hours
	Input Manu Rapid Jettin	d Polymer Based Rapid Prototyping systems: SLA, Material Je Materials Based Rapid Prototyping Systems: Laminate facturing (LOM) and Fused Deposition Modelling Systems, Pol Prototyping Systems: Selective Laser Sintering, Multi-Jet Fuseg Systems.	ed Object ower Based	
UNIT-III	RP D	Patabase & Design Optimization:		8 hours
	_	l Prototyping Data Formats, STL Format, STL file problems, STI M, Topology Optimization, Gcode for RP Systems	L file repair,	
UNIT-IV	RP A	pplications:		8 hours
	produ	lopment of dies for Moulding, RP Applications in developing practs, application in medical fields, Development of bone replaces, etc., RP materials and their biological acceptability.	• •	
Course ou	itcome	After completion of this course students will be able to		
CO 1	Under	stand the fundamentals of RP Technologies and process involvementals	ent in them	K1,K2
CO 2		Understand the methodology to manufacture the products using RP technologies		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	ıa, Kah Fai	
2		Additive Manufacturing Technologies: 3D Printing, Rapid Proto Direct Digital Manufacturing: Brent Stucker, David W. Rosen, Ia	• 1	

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.	