ACADEMIC CURRICULA

UNDERGRADUATE/ INTEGRATED POST GRADUATE DEGREE PROGRAMMES

(With exit option of Diploma)

(Choice Based Flexible Credit System)

Regulations 2021

Volume - 8
(Syllabi for Biotechnology Programming Courses)
(Revised on August 2024)



SRM INSTITUTE OF SCIENCE AND TECHNOLOGY

(Deemed to be University u/s 3 of UGC Act, 1956)

Kattankulathur, Chengalpattu District 603203, Tamil Nadu, India

ACADEMIC CURRICULA

Engineering Science Courses

Regulations 2021



SRM INSTITUTE OF SCIENCE AND TECHNOLOGY

(Deemed to be University u/s 3 of UGC Act, 1956)

Kattankulathur, Chengalpattu District 603203, Tamil Nadu, India

Course	21CHS251T	Course	BASIC CHEMICAL ENGINEERING	Course	c	ENGINEERING SCIENCES	L	Т	Р	С
Code	210032311	Name	BASIC CHEMICAL ENGINEERING	Category	9	ENGINEERING SCIENCES	3	0	0	3

Pre-requisite Courses	Nil	Co- requisite Courses	Nil	ogressive Courses	Nil
Course Offering Dep	artment	Chemical Engineering	Data Book / Codes / Standards		Nil

Course L	earning Rationale (CLR):	The purpose of learning this course is to:	-40	\overline{A}	<u> </u>		Progr	am Ou	itcome	s (PO)					rograi	
CLR-1:	describe the basic principl	es of process cal <mark>culation</mark>	1	2	3	4	5	6	7	8	9	10	11	12		pecifi utcom	
CLR-2:	explain the concepts of St	pichiometry <mark>equations a</mark> nd material balances	ge	work Work													
CLR-3:	demonstrate the behavior	of fluids a <mark>nd fluid flo</mark> w phenomena	w lec	sis sis oment of oment of oment of oment of oment of one of other sage am Work am Work or of other oth				Вu									
CLR-4:	IP-4: describe the principles of filtration, working of filtration equipment's and concept of adjustion					arni											
CLR-5:	illustrate the basic concep	ts and <mark>laws of t</mark> hermodynamics	ering	٩	n/deve		ě	engineel sty	Environment 8 Sustainability		<u>8</u>	mmunication	Mgt.	ong Le			
			<u>e</u>	Problem	rign/	onduct in f complex	Modern		iron tain	S	Individual	l III	Project		7)-2	53
Course C	Outcomes (CO):	At the end of this course, learners will be able to:	Eng	Pro	Des	55	Mod	The	Env Sus	Ethics	<u>lp</u>	Š	Proj	Life	PSO	PSO.	PSO
CO-1:	perform unit conversions a	nd <mark>stoichio</mark> metric calculations	2	3	F-10	1.7		7	-	-	-	-	-	-	-	-	-
CO-2:	interpret material balance	fo <mark>r non-rea</mark> ctive unit operations	3	3	File	-	-	7	-		-	-	-	-	-	-	-
CO-3:	apply fluid properties, cont	in <mark>uity and</mark> Bernoulli equation for fluid flow	2	3	17-	-	-	-	f -		-	-	-	-	-	-	-
CO-4:	formulate the concepts of	filt <mark>ration a</mark> nd agitation in processes	1	2	-45	الخز	-	-	-	- 1	-	-	-	-	-	-	-
CO-5:	comprehend the basic con	ce <mark>pts and</mark> laws of thermodynamics for different processes	2	2		6-	-		-	-	-	-	-	-	-	-	-

Unit-1 - Fundamental Concepts of Stoichiometry

9 Hour

Concept of units and dimensions, system of units, unit conversions, basis of calculation, concept of mole, expressing composition of mixture of solids, liquids and gases - percentage by weight, mole and volume and density calculation, concentrations - molality, molarity, normality, ppm, predicting P-V-T properties of gases using ideal gas law

Unit-2 - Material Balance in Unit Operations

9 Hour

Introduction to material balance, material balance for non-reactive chemical process systems - Mixing, Drying, Crystallization, Extraction, Chemical reactions and stoichiometric equations - limiting reactant, excess reactant, conversion, degree of completion, selectivity and yield, concept of recycle, purge and bypass stream

Unit-3 - Fluid Flow Phenomena

9 Hour

Fluid, properties of fluids, type of fluids and flow, Fluid statics - hydrostatic equilibrium, Pressure measurement by manometers - simple U-tube, differential U-tube, inclined differential manometers, Reynolds number, continuity equation, Bernoulli equation

Unit-4 - Filtration and Agitation

9 Hour

Concept of Filtration, Filter media, filter aid, principles of cake filtration, pressure drop through filter cake, Compressible and incompressible filter cakes, filter medium resistance, Constant pressure filtration, constant rate filtration, Filtration equipment's - principle and working of filter press, Vacuum leaf filter, rotary drum filters. Introduction to agitation, agitation equipment, impeller, turbines, flow patterns, prevention of swirling, draft tubes

Unit-5 - Basic Concepts in Thermodynamics

9 Hour

Chemical Engineering Thermodynamics- System, surrounding, boundary, Work, Energy, Heat, Internal energy, Intensive and Extensive properties, State and path functions, processes and its type, equilibrium, enthalpy. Heat capacity- derivation for constant volume and constant pressure processes. First Law of Thermodynamics-Mathematical statement, sign convention, problems, Limitations of First Law of Thermodynamics, Energy balance for closed system. statement of second law of thermodynamics, concept of entropy, Third law of thermodynamics

Learning Resources
Resources

- 1. Himmelblau D.H. and James B. Riggs, Basic Principles and Calculations in Chemical Engineering, 8th Edition, Prentice Hall, 2012
- 2. Bhatt, B.I. and Thakore S.M., Stoichiometry, 5th Edition, Tata McGraw-Hill Publishing Company Ltd., New Delhi, 2010
- 3. Warren L. McCabe, Julian C. Smith and Peter Harriott, "Unit Operations of Chemical Engineering", 7th Edn., McGraw Hill Education (India) Edition, 2022
- Noel de Nevers, Fluid Mechanics for Chemical Engineers, 2nd ed., McGraw Hill International Editions. 1991
- 5. Smith, J.M., Van Ness, H.C., Abbott, M.M., Introduction to Chemical Engineering Thermodynamics, 8 th ed., McGraw Hill International Edition, 2018

	Bloom's Level of Thinking	CLA-1 Aver	Continuous Learnin mative rage of unit test 50%)	CL	Learning A-2 %)	Summative Final Examination (40% weightage)			
	/ 3 /	Theory	Practice	Theory	Practice	Theory	Practice		
Level 1	Remember	20%	FR 3-0, 37/2	20%		20%	-		
Level 2	Understand	20%		20%	6.4	20%	-		
Level 3	Apply	30%	42, 74, 92,251	30%		30%	-		
Level 4	Analyze	30%	to the second	30%		30%	-		
Level 5	Evaluate	-	A 10 1 1 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	10	. 1.7	9 -	-		
Level 6	Create		102 (10 June 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	St. 1 32. 72		-	-		
	Total —	10.11	00 %	100	0 %	10	00 %		

Course Designers		
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
1. Mr. A. Subramaniam, PESCO Beam Environmental Solutions Pvt. Ltd	1. Dr. Lima Rose Miranda, Anna University	1. Dr. S. Kiru <mark>thika, SR</mark> MIST
2. Mr. S. Stalin, Course Director, Chem Skill Development Centre	2. Dr. N. Anantharaman, Former Professor, NIT Trichy	2. Dr. E. Poo <mark>nguzhali</mark> , SRMIST

Course	210402521	Course	CHEMICAL ENGINEERING PRINCIPLES	Course	c	ENCINEEDING SCIENCE	L	Т	Р	С
Code	210032323	Name	CHEMICAL ENGINEERING PRINCIPLES	Category	0	ENGINEERING SCIENCE	3	0	2	4

Pre-requisite Courses	Ni	Co- requisite Courses	Nil	Progressive Courses	Nil	
Course Offeri	ng Department	Chemical Engineering	Data Book / Codes / Star	ndards	Nil	

Course L	earning Rationale (CLR): The purpose of learning this course is to:	di	4			Progr	am Oı	utcome	s (PO))					rograr	
CLR-1:	describe the various modes of heat transfe <mark>r and evaluat</mark> e the rate of steady state heat transfer	1	2	3	4	5	6	7	8	9	10	11	12	Specific Outcomes		
CLR-2:	CLR-2: explain and analyze the basic concepts of convection as applied to various flows and geometry				ciety			~								
CLR-3:	illustrate principles of mass transfer, Diffusion phenomena, and calculate mass transfer rates	edge		nt of	ions	ous	socie			Work		inance				
CLR-4:	elucidate the principles of drying, different types of drier and calculate drying time for different dry periods	줃	alysis	velopment	estigations blems	ol Usage	er and	st ×		Team	tion	8 8	arning			
CLR-5:	demonstrate the concept of distillation, extraction and adsorption	leering	em An	Jn/deve	uct invi	ern Too	engineer	ironment tainability	S	vidual &	ommunication	ct Mgt.	ong Le	<u> </u>	5	.3
Course C	Outcomes (CO): At the end of this course, learners will be able to:	Engine	Prob	Designation	Sono	Mode	The 6	Envir	Ethic	ndiv	Som	Project	Life L	PSO.	PS0-2	-0Sd
CO-1:	analyze steady state heat co <mark>nduction</mark> and calculate the rate of heat transfer	2	2	18/5/	47-		1	-		-	-	-	-	-	-	-
CO-2:	apply the basic concepts of convection and calculate the heat transfer coefficient	3/-	3	3	~	-			-	-	-	-	-	-	-	-
CO-3:	interpret mass transfer princ <mark>iples an</mark> d solve diffusion problems	1004	2	- 3	-	-	7	-	-	-	-	-	-	-	-	-
CO-4:	calculate drying time for different periods of drying	4.04	2	2	25	-	_	-		-	-	-	-	-	-	-
CO-5:	comprehend the various types of distillation, extraction and adsorption for different processes		2	- 2	1	_	-	_	_ 6	_	-	-	-	_	_	-

Unit-1 - Conduction 15 Hour

Introduction to various modes of heat transfer, Concept of rate of heat transfer, heat flux, conduction, Fourier's law of heat conduction, Thermal conductivity, Steady state heat conduction through plane wall, composite wall, hollow cylinder, coaxial cylinders

Unit-2 - Convection and Heat Exchangers

15 Hour

Concept of heat transfer by convection, Newton's law of cooling, Natural and forced convection- Dimensional analysis- Empirical correlations, Heat exchange equipment, Parallel and counter flow, LMTD, heat transfer area

Unit-3 - Mass Transfer and Diffusion 15 Hour

Introduction to Mass Transfer, Diffusion, Types, Fick's law of Diffusion, Molecular diffusion in gases: steady state diffusion of A through non-diffusing B, Gas phase equimolal counter diffusion, Diffusion in Multicomponent gas mixtures, Molecular diffusion in liquids: steady state diffusion of A through non-diffusing B, Liquid phase equimolal counter diffusion, Effect of temperature and pressure on diffusivity

Unit-4 - Drying

15 Hour

Drying - Importance of drying in processes, principles of drying, wet Basis, dry basis, Free moisture, equilibrium moisture, bound and unbound moisture, Mechanism of drying, drying curve, Calculation of drying time under constant drying conditions: constant rate and falling rate period, Total drying time, Classification of dryers, solids handling in dryers, tray, rotary, spray and fluidized bed drier

Unit-5 - Distillation, Leaching and Adsorption

15 Hour

Introduction to Distillation, principle, Raoult's law, relative volatility, Types of distillation, batch distillation - Rayleigh's equation, flash and steam distillation, General principles of extraction, choice of solvent, mixer-settler, Introduction to leaching, adsorption – isotherm

Practice

Practice 1: Heat transfer through composite wall

Practice 2: Heat Transfer through composite lagged pipe

Practice 3: Heat transfer by natural convection

Practice 4: Heat transfer by forced convection

Practice 5: Stefan-Boltzmann apparatus

Practice 6: Double pipe heat exchanger

Practice 7: Shell and tube heat exchanger

Practice 8: Estimation of Diffusivity

Practice 9: Drying characteristics

Practice 10: Batch distillation

Practice 11: Steam distillation

Practice 12: Single stage leaching

Practice 13: Multi stage leaching

Practice 14: Soxhlet Extractor

Practice 15: Adsorption

Learning Resources

- Edition, 2012.
- Engineering", 7th Edn, McGraw Hill Education (India) Edition, 2022. 3. Christie John Geankoplis, "Transport Processes and Separation Process Principles (Includes Unit Operations)", 4thEdn, Pearson India Education Services Pvt. Ltd., 2015.
- 1. Robert E. Treybal, "Mass-Transfer Operations", 3rd Edn., McGraw Hill Education (India) 4. Binay K. Dutta, "Principles of Mass transfer and Separation Processes", Prentice- Hall of India, New Delhi, 2016.
- 2. Warren L. McCabe, Julian C. Smith and Peter Harriott, "Unit Operations of Chemical 5. N. Anantharaman and K. M. Meera Sheriffa Begum, "Mass Transfer Theory and Practice", Prentice Hall of India Pvt. Ltd., New Delhi, 2017.

Learning Assessme	ent		1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	GALL	The Contract of the Contract o							
			Contin	uous Learning	g Assessment (CLA)	7	Cum	mativa				
	Blo <mark>om's</mark> Level of <mark>Thinkin</mark> g	CLA	Formative -1 Average of unit tes (45%)	st	CL	Learning A-2 (%)	Summative Final Examination (40% weightage)					
		Theory	Pra	octice	Theory	Practice	Theory	Practice				
Level 1	Remember	20%	ASSET 11 11 11 11 11 11 11 11 11 11 11 11 11	- <u>11</u> × 7	20%		20%	=				
Level 2	Understand	20%			20%		20%	-				
Level 3	Apply	30%		- 10	30%	-	30%	=				
Level 4	Analyze	30%		 AVV. 	30%	4	30%	-				
Level 5	Evaluate	ela le		- / /	-		-	-				
Level 6	Create			-	-	- - V /	-	-				
	Total	1 7 7	100 %	1900	100) %	10	0 %				

Course Designers		
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
1. Mr. A. Subramaniam, PESCO Beam Environmental Solutions Pvt. Ltd	1. Dr. Lima Rose Miranda, Anna University	1. Dr.S. Kiruthika, SRMIST
2. Mr. S. Stalin, Course Director, Chem Skill Development Centre	2. Dr. N. Anantharaman, Former Professor, NIT Trichy	2. Dr. E. Poonguzhali, SRMIST

ACADEMIC CURRICULA

Professional Core Courses

Regulations 2021



SRM INSTITUTE OF SCIENCE AND TECHNOLOGY

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Kattankulathur, Chengalpattu District 603203, Tamil Nadu, India

Course	21BTC201L	Course	BIOCHEMISTRY LABORATORY	Course		PROFESSIONAL CORE	L	Т	Р	С
Code	ZIBICZUIL	Name	BIOCHEMISTRY LABORATORY	Category	C	PROFESSIONAL CORE	0	0	4	2

Pre-requisite Courses	N		 Nil	Progressive Courses	Nil
Course Offeri	ng Department	Biotechnology	 Data Book / Codes / Standards		Nil

Course L	earning Rationale (CLR): The purpose of learning this course is to:	71			. "	Progra	am Ou	itcome	es (PC))					ogra	
CLR-1:	understand the preparation of laboratory reagents with competence and proficiency	1	2	3	4	5	6	7	8	9	10	11	12		pecifi tcom	
CLR-2:	analyze the different forms of carbohydrates in samples qualitatively using different chemical tests	0		1	of		ety			~						
CLR-3:	determine the types of fatty acids, and use a variety of tests and reagents	edge)	nt of	ions	Φ	society			Work		Finance				
CLR-4:	become familiar with chromatographic methods and use them to isolate and characterize various biologisubstances	조		evelopment	investigations	Š	and	t &		Team	tion	∞ర	earning			
CLR-5:	recognize the fundamentals of various reagents and how they interact with biomolecules for measurement	ering	, A				engineer	Environment Sustainabilit	l.	<u>8</u>	Sommunication	Mgt.				
		e	<u>a</u>	/ugi	duct	Modern		iron	છ	ndividual	שנ	Project	Long	-1)-2	-3
Course C	Outcomes (CO): At the end of this course, learners will be able to:	Engi	, l g	Des		Moo	T _e	Env	Ethics	İpu	Con	Proj	Life	PSO	PSO-	PSO-3
CO-1:	perform basic professional skills related to solutions, pH, and buffer preparation, as well as nume calculations, focusing on the laboratory	rical 3	3	3	} -	7	_	-	-	-	1	-	-	-	3	-
CO-2:	identify the various ways in which different types of carbohydrates respond to chemical tests	No of	3	3	-		-	-		-	-	-	-	-	3	-
CO-3:	explain how various chemicals interact with fatty acids to determine the distinct types	- 3	3		3	-	-	-	-	-	-	-	-	-	-	3
CO-4:	develop methods for separat <mark>ing and</mark> detecting amino acids	- 3	3	44.5	3	1	-	-	1	-	-	-	-	-	3	-
CO-5:	describe the measurement of biomolecules in clinical and dietary samples	25.3	3	-	3	74	-	_	-	-	-	-	-	-	3	-

Unit-1 - Basics of Analytical Biochemistry 12 Hour

Practice:

- 1. Stoichiometric calculations Molecular weight calculation, Molarity, Normality, Molality, % solution, w/w, v/w, v/v, etc.
- 2. Verifying the influence of H+ and OH- ions in the test solutions by pH meter.
- 3. Preparation of biological buffers.

Unit-2 - Qualitative Analysis of Biomolecules - Carbohydrates

12 Hour

Practice:

- 1. Differentiate between aldose and ketose sugars with standards and natural food samples.
- 2. Identify whether the given sugar is pentose/reducing sugar or not with standards and food samples.

3. Distinguishes between mono or disaccharides also to check to reduce or non-reducing disaccharides with standards and food samples such as milk, malted sugars, and sugarcane juice/Jaggery.

Unit-3 - Qualitative Analysis of Biomolecules- Carbohydrates, Fatty Acids /Lipids

12 Hour

Practice:

- 1. Verifying the given carbohydrate is starch polysaccharide.
- 2. Qualitative analysis of fatty acids and cooking oils/fish oils.

Unit-4 - Separation of Biomolecules and Quantitative Analysis of Biomolecules

12 Hour

Practice:

- 1. Separation of amino acids from the mixture and boiled legumes as test samples by TLC and detection by using ninhydrin solution.
- 2. Estimation of reducing sugar-glucose from the blood by 3, 5-Dinitrosalicylic acid (DNS) method.

Unit-5 - Quantitative Analysis of Biomolecules

12 Hour

Practice:

- 1. Estimation of protein from food samples by Lowry's method.
- 2. Quantification of cholesterol from egg yolk by Zak's method.

Learning
Resources

- Biochemistry Practical Manual 2023.
 Varley's Practical Clinical Biochemistry by Gowenlock A.H., 6th Fo
- Varley's Practical Clinical Biochemistry by Gowenlock A.H., 6th Edition, 2022 (8th Reprint), ISBN: 9788123904276, CBS Publishers & Distributors.
- Principles and Techniques of Practical Biochemistry (5th Ed.). Wilson, K., Walker, J. (eds.); Cambridge University Press, Cambridge, 2000, 784 pp., ISBN 0-521-65873-X.
- 4. An Introduction to practical biochemistry (2nd edition): By David T. Plummer. Pp 362 McGraw-Hill Book Company (U.K.) Ltd., London 1978. https://doi.org/10.1016/0307-4412(78)90089-4

Learning Assessr	nent			ROLL STATE	21.						
		_	Continuous Learning Assessment (CLA)								
	Bloom's Level of Thinking	exper	ge of first cycle riments 0%)	exper	of second cycle iments 0%)		Examination reightage)	Summative Final Examination (0% weightage)			
		Theory	Practice	Theory	Practice	Theory	Practice Practice	Theory	Practice		
Level 1	Remember		15%	- 30E 97	15%	M (15)	15%	=	-		
Level 2	Understand		20%	100	20%	4 , 4-	20%	=	-		
Level 3	Apply		25%	3 A 1 2 2 2	25%	75 375	25%	-	-		
Level 4	Analyze	-	25%	The second of	25%	- 14 C	25%	=	-		
Level 5	Evaluate	-	10%		10%		10%	-	-		
Level 6	Create	1	5%	- \	5%	- / (5%	-	-		
	Total	10	0 %	100	0 %	10	00 %		-		

Course Designers	17.11	A 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
Experts from Industry	Experts from Higher Technical Institutions	Internal Ex <mark>perts</mark>
1. Dr. G. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai,	1. Prof. K Subramaniam, IITM, Chennai,	1. Dr. <mark>Pachiapp</mark> an, SRMIST
ramchand@saksinlife.com	suubu@iitm.ac.in	b /60
2. Dr. Karthik Periyasamy, Scientist, Biocon,	2. Prof. R. B. Narayanan, Anna University, Chennai	2. <mark>Dr. S Sub</mark> ashini, SRMIST
karthik.periyasamy@biocon.com	arbeen09@gmail.com	

Course	21BTC202T Course	MICROBIOLOGY	Course		PROFESSIONAL CORE	L	Τ	Р	С
Code	Name	WIICKOBIOLOGT	Category	U	PROFESSIONAL CORE	3	0	0	3

Pre-requisite Courses	N	Co- requisite Courses	Nil Progre	Nil	
Course Offeri	ng Department	Biotechnology	Data Book / Codes / Standards	Nil	

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Course L	earning Rationale (CLR): The purpose of learning this course is to:					Progr	am Ou	tcome	s (PO)					gram
CLR-1:	introduce the concept of Microbiology and <mark>Microorganism</mark> s	1.1	2	3	4	5	6	7	8	9	10	11	12		ecific comes
CLR-2:	understand the growth, metabolism and adaptation of bacteria	dge		Jo	s of					ork		8			
CLR-3:	illustrate the structure and life cycle o <mark>f eukaryot</mark> es	wled			ation	ge	-			≥		inanc	g		
CLR-4:	illustrate the structure and life cycle of viruses	Knowlec	nalysis	velopment	vestigations oblems	Usage	r and	∞ _		Feam	.u	ĕ E	aming		
CLR-5:	analyze the applications of Micro <mark>biology in</mark> various fields		⋖	le ve	inve	100 100	engineer ety	nability	l.	- ∞ - ∞	Communication	Mgt.	Le		
		Engineering	Problem	Design/dev	nduct	dern		nvironr ustaina	SS	ndividual	JI JI	ect N	Long		3
Course C	Outcomes (CO): At the end of this course, learners will be able to:	Eng	Pro	Des		Mod	The	Envi	Ethics	Indi	Con	Project	Life	PSO	PSO-2 PSO-3
CO-1:	illustrate the structure of prok <mark>aryotes</mark>	2	2	2	-	1	7-	-	-	-	-	-	-	2	- -
CO-2:	understanding the growth of prokaryotes	2	2	2	-	2		-		-	-	-	-	2	- -
CO-3:	explain the growth and life c <mark>ycle of m</mark> icrobial eukaryotes	3	2	2	2	- 4		-	-	-	-	-	-	3	- -
CO-4:	discuss the life cycle and pathogenicity of viruses	-3	2	3	-		_	-	-	-	-	-	-	3	- -
CO-5:	discuss the role of microbes and microbial products in various fields	3	2	2	-	3		-	-	-	-	-	-	3	- -

Unit-1 - Microscopy and Structure of Prokaryotes

9 Hour

Introduction to Microbiology, Characterization, Classification and Identification of microbes, Microscopy - Light, Electron and Advanced Microscopy, Structure of prokaryotes - Bacteria, Mycoplasma, Morphology, Structure, Cultivation, Reproduction and Pathogenicity of Actinomycetes

Unit-2 - Metabolism and Adaptation of Prokaryotes

9 Hour

Metabolism of Prokaryotes: Bacteria - Growth curve and kinetics. Quantification of bacterial growth, Microbial metabolism: Non-biosynthetic and biosynthetic pathway. Adaptation mechanism of Halophiles, Alkaliphiles, Psychrophiles, Piezophiles, Xerophiles. Bacterial Recombination: Transformation, Transduction, Conjugation

Unit-3 - Eukaryotes Structure and Methods of Microbial Control

9 Hour

Structure of eukaryotes: Fungi, Algae and Protozoa - Characteristics, Morphology, Reproduction, Physiology and Pathogenicity. Control of Microorganisms: Physical Control and Chemical Control. Antibiotics Unit-4 - Structure of Virus

9 Hour

Virus: Morphology, Structure, Classification and Pathogenicity. Bacteriophages: Lytic and Lysogenic life cycle of bacteriophages. Animal viruses, Plant viruses and Oncoviruses. Plaque assay.

Unit-5 - Applications of Microbiology

9 Hour

Applications of Microbiology: Soil Microbiology - Microbiology - Microbiology - Microbiology - Microbiology - Biofertilizers. Environmental Microbiology - Bioremediation, Bioplastics, Biopolymers. Industrial Microbiology - Microbial metabolites. Medical Microbiology - Antibiotics and Vaccines

Learning Resources
Resources

- 1. Pelczar MJ, Chan ECS and Krein NR: Microbiology, Mc Graw Hill, 10 th Edition, 2016.
- 2. Michael T. Madigan, Kelly S. Bender, Daniel H. Buckley, W. Matthew Sattley and David A. Stahl: Brock Biology of Microorganisms, Pearson. 15 th Edition, 2017.
- 3. Joanne M. Willey, Linda M. Sherwood, Christopher J. Woolverton: Prescott, Harley and Klein's Microbiology, Mc Graw Hill, International Edition, 10 th Edition, 2016.
- 4. Jawetz, MA Brooks, GF Butel JS and Morse SA: Medical Microbiology, Mc Graw Hill, 26 th Edition, 2012

-		100	Continuous Learning Assessment (CLA)						
	Bloom's Level of Thinking	Form CLA-1 Averaç (50	ge of unit test	CL	g L <mark>earning</mark> .A-2 0%)	Summative Final Examination (40% weightage)			
		Theory	Practice	Theory	Prac <mark>tice</mark>	Theory	Practice		
Level 1	Remember	15%		15%		15%	-		
Level 2	Understand	25%		20%	- A- V	25%	-		
Level 3	Apply	30%	42.54.5 6.6	25%		30%	-		
Level 4	Analyze	30%	1 No. 20 17 17 17 17 17 17 17 17 17 17 17 17 17	25%		30%	-		
Level 5	Evaluate			10%	- C - C	4 -	-		
Level 6	Create		State and the result	5%		-	-		
	Total	100)%	10	0 %	10	0 %		

Course Designers		
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
1. Dr. G. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai,	1. Prof. K Subramaniam, IITM, Chennai,	1. Dr. J. Lava <mark>nya, SRM</mark> IST.
ramchand@saksinlife.com	suubu@iitm.ac.in	
2. Dr. Karthik Periyasamy, Scientist, Biocon,	2. Prof. R. B. Narayanan, Anna University, Chennai	2. Dr. R. Muthukumar, SRMIST.
karthik.periyasamy@biocon.com	arbeen09@gmail.com	

Course	21BTC203L Cours	CELL AND MICROBIOLOGY LABORATORY	Course	PROFESSIONAL CORE	L	Т	Р	С	,
Code	21BTC203L Nam	CELL AND WICKOBIOLOGY LABORATORY	Category	PROFESSIONAL CORE	0	0	4	2	

Pre-requisite Courses	Ni	Co- requisite Courses	Nil	Progressive Courses	Nil
Course Offerin	g Department	Biotechnology	Data Book / Codes / Standards		Nil

Course Lo	earning Rationale (CLR): The purpose of learning this course is to:		4	1	٦, ١	rogra	am Ou	tcome	s (PO))					rogram
CLR-1:	provide basic differences between prokaryo <mark>tic and euka</mark> ryotic organisms	1.	2	3	4	5	6	7	8	9	10	11	12		pecific itcomes
CLR-2:	understand the different strategies of organization of cellular structures	ge		of	SL			N		^o rk		8			
CLR-3:	provide hands on training in isolation of cells and cell organelles	Knowledge	S	velopment	investigations ex problems	Usage	ъ	. \		am W		Finance	Вu		
CLR-4:	focus on the cellular response to stimulus		Analysis	lopr	estig	l Us	er and	y t &		Tea	ţi	∞ŏ	arning		
CLR-5:	comprehend the mechanism of bacterial pathogenesis	ering		deve	t inv	Tool	engineer sty	ronment tainability	N	<u>a</u>	nica	Mgt.	ng Le		
Course	utcomes (CO): At the end of this course, learners will be able to:	Engineering	roblem	esign/	Conduct of comple	Modern	ne en	Environ Sustain	Ethics	ndividual	ommunication	Project	ife Lor	-SO-1	PSO-2 PSO-3
-		Ü	- G	٩	9,0	Σ	<u></u>	ш <u>х</u>	Ш	<u>=</u>	Ö	<u>P</u>	ت	ď	
CO-1:	distinguish between prokaryo <mark>tic and e</mark> ukaryotic cells using microscopic analysis		. 3	3	-	-	-	-		-	-	-	-	-	3 -
CO-2:	gain proficiency in identifying the cellular structures	1	-	3	3	- 1	生	-	-	-	-	-	-	-	- 3
CO-3:	acquire skills to isolate cells and cell organelles and relate with cell division	182	3	3	-	- 1	_	-		-	-		-	-	3 -
CO-4:	critique the cell's response to stimuli thereby correlating cell signaling			3	3	- 1	-	-	-	-	-	-	-	-	3 -
CO-5:	integrate cell biology & microbiology to understand the bacterial pathogenesis in host	-		1	3	l - ;	_	-	-	-	-	-	-	-	- 3

Unit-1 - Distinguish Between Prokaryotic and Eukaryotic Cells

Practice:

1. Microscopic observation of cells: Simple staining & Cross section of plant & animal tissues

- 2. Biochemical characterization of bacteria IMVIC tests
- 3. Specific enzyme assays and substrate hydrolysis for bacterial identification

Unit-2 - Visualization of Cellular Structures Using Differential Staining

Practice:

- 1. Cell wall staining Gram staining/ Lactophenol cotton blue staining of fungi
- 2. Nuclear staining of cells using Giemsa
- 3. Bacterial Spore staining.

Unit-3 - Isolation of Cells/Cell Organelles and Cell Division

Practice:

- 1. Isolation of bacteria by pour plate/spread plate and culturing techniques (Streak, Slant & Deep).
- 2. Isolation of Chloroplast from leaves and determination of chlorophyll content
- 3. Mitosis cell division in vegetative cells

12 Hour

12 Hour

12 Hour

Unit-4 - Response of Cell to Stimuli

12 Hour

Practice:

- 1. Stomatal movement in response to stimulus
- 2. Bacterial motility using hanging drop technique
- 3. Determination of cell viability using tryphan blue

Unit-5 - Understand the Mechanism of Bacterial Pathogenesis

12 Hour

Practice:

- 1. Bacterial Growth curve
- 2. Antibiotic sensitivity tests using Kirby Bauer assay
- 3. Adherence of Enteropathogenic E.coli on host cells.

Learning
Resources

- 1. Lab manual
- Chaitanya, k. V. Cell and molecular biology: A Lab Manual. India, PHI Learning, 2013.
- 3. Lorrence H. Green, Emanuel Goldman. Practical Handbook of Microbiology: Fourth Edition, CRC Press. Taylor and Francis; 2021.
- 4. Julio E.Cellis. Cell Biology: A Laboratory Handbook. (2008). United Kingdom: Academic Press

Learning Assessm	nent		- 10-2		THE PARTY IN		4		
			7 277	Continuous Learnin	g Assessment (CLA)		Cume	native	
	Bloom's Level of Thinki <mark>ng</mark>	expe	nge of first cycle riments 30%)	exper	of second cycle iments 0%)		Examination reightage)	Final Exa	amination ightage)
		Theory	Practice	Theory	Practice	Theory	Practice Practice	Theory	Practice
Level 1	Remember		15%	77.5	15%	A	15%	-	-
Level 2	Understand		20%	70.75 200	20%	-12	20%	-	-
Level 3	Apply	-	25%	44.	25%	100	25%	-	-
Level 4	Analyze		25%		25%	-	25%	-	-
Level 5	Evaluate		10%		10%	- / /	10%	-	-
Level 6	Create		5%	- 7777	5%	-/ "	5 <mark>%</mark>	-	-
	Total	1	00 %	10	0 %	10	00 %		-

Course Designers		
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
1. Dr. G. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai,	1. Prof. K Subramaniam, IITM, Chennai, suubu@iitm.ac.in	1. D <mark>r.S.Sujath</mark> a, SRMIST
ramchand@saksinlife.com	VITEARN FAD round	
2. Dr. Karthik Periyasamy, Scientist, Biocon,	2. Prof. R. B. Narayanan, Anna University, Chennai	2. Dr.J.Lavanya, SRMIST
karthik.periyasamy@biocon.com	arbeen09@gmail.com	

Course	21BTC204T	Course	DIODDOCECC DDINICIDI EC	Course	_	PROFESSIONAL CORE	L	Т	Р	С	
Code	210102041	Name	DIOPROCESS PRINCIPLES	Category	C	PROFESSIONAL CORE	3	0	0	3	

Pre-requisite Courses	Nil	Co- requisite Courses	Nil	Progressive Courses	Nil
Course Offering Depart	tment	Biotechnology	Data Book / Codes / Standards		Nil

Course L	earning Rationale (CLR):	The purpose of le <mark>arning this co</mark> urse is to:		-			_ 1	rogra	am Ou	tcome	s (PO)					ogra	
CLR-1: describe the basics of the fermentation process				1	2	3	4	5	6	7	8	9	10	11	12		pecifi tcom	
CLR-2:	explain the process of media	formulatio <mark>n and sterili</mark> zation kinetics		a fir		of	SL			I		ork		9				
CLR-3:	study the basics of reactor de	esign an <mark>d its contro</mark> l systems		Nilowiedge Significant of the significant of the si	,	velopment	vestigations x problems	Usage	ъ			am W		inance	рu			
CLR-4:	analyze the metabolic stoichi	ometr <mark>y and ene</mark> rgetics of the biochemical process			Allalysis	lopn	estig orobl	- Os	r and	∞ _{>}		Teal	tion	⊗ ⊢	arning			
CLR-5:	illuminate the various types o	f rea <mark>ctors for</mark> suspension and immobilized cell systems	T 1 1 1		٠ ا		t inv	\vdash	engineer ety	nability		<u>a</u>	Communication	Project Mgt.	ong Le			ł
		The state of the s	275			sign/de utions	duc	dern	er et	ron	S	Jġ.	<u>ا</u>	ect	Lor	7	7-5	5
Course C	outcomes (CO):	A <mark>t the en</mark> d of this course, learners will be able to:	3,444			Des solu	Con	Moc	The	Env	E E	Individual	Col	Proj	Life	PS0-1	PSO	PSO-3
CO-1:	understand the basics of the	f <mark>ermenta</mark> tion process		1		2	-	T	7	-	-	-	-	-	-	2	2	2
CO-2:	comprehend the process of n	nedia formulation and sterilization kinetics	1 3	2	2	2	2	2 -		-	=	-	-	-	-	-	2	1
CO-3:	acquire the basics of reactor	design and its control systems	C / X	2	-4	2	- 1	2		-	-	-	-	-	-	2	2	1
CO-4:	evaluate the metabolic stoichiometry and energetics of the biochemical process		1.34	2 .	3	1	2	-	-	-		-	-	-	-	2	-	-
CO-5:	explore the various types of reactors for suspension and immobilized cell systems		ماوال	3		2	2	- 1	_	-	1	-	-	-	-	2	2	2

Unit-1 - Microbial Cell Factories

9 Hour

Cellular systems as molecular factories and its industrial importance, Isolation and improvement of industrially important organisms, Types of fermentation, Upstream and downstream bioprocess, Process flow sheets of primary and secondary metabolites production- eg. ethanol, lactic acid, lysine, poly-L-lactic acid, lipase, rhamnolipid, streptomycin, insulin, Interferon, monoclonal antibody, tumour necrosis factor inhibitor, Pneumococcal conjugate vaccine.

Unit-2 - Design and Preparation of Media for Bioprocess

9 Hour

Bioreaction theory, Kinetics of biological systems, Growth patterns and kinetics of cells, Quantifying cell growth kinetic parameters, Optimization of cell growth environment, Types of media and classes of medium components. Media formulation and optimization of medium for the industrially important cultures - Microbial, plant and animal cells, Sterilization, Types of sterilization - batch, continuous and air sterilization

Unit-3 - Bioprocess Design - Instrumentation and Control Systems

9 Hour

Fermentation facility, equipment and space requirements - Fermenter design and its configuration, Body construction, Agitators, Stirrer glands and bearings, Spargers and valves, Aseptic operation and containment, Bioinstrumentation and its control - Methods of measuring process variables, Online analysis of chemical factors, Control systems, Combination of methods of the controller, Troubleshooting in a fermentation plant.

Unit-4 - Fundamentals of Biological Engineering

9 Hour

Material and energy balances for reactive and non-reactive systems; Stoichiometry of growth and product formation; Degree of reduction, electron balance and theoretical oxygen demand, Determination of stoichiometric coefficients, Theoretical prediction of yield coefficients, Conductive and convective heat transfer; Overall heat transfer coefficient, Bio-thermodynamics.

Unit-5 - Bioreactors for Suspension and Immobilized Cultures

9 Hour

Strategies for choosing a bioreactor, Microbial and immobilized cell system, Active and passive immobilization of Cells, novel reactors - Airlift Bioreactor, Fluidized Bed Bioreactor, Membrane Bioreactor, Photobioreactor, Biofilm reactor, Single-use bioreactors, Various modes of operation in Bioreactors, Performance equation of a batch, fed-batch and continuous reactors, Stability analysis of bioreactor.

Learning Resources	1. 2.	Pauline M. Doran "Bioprocess Engineering Principles", 2nd Edition, Academic Press, 2012. Michael L. Shuler, Fikret Kargi, Matthew DeLisa "Bioprocess Engineering: Basic Concepts", 3rd Edition, Prentice-Hall, 2017.	Hall, Stephen J., Stanbury, Peter F., Whitaker, Allan, "Principles of Fermentation Technology", 3rd Edition, Butterworth– Heinemann, 2017.
			I i .

Learning Assessm	ent	/ .00									
	Bloom's Level of Thinking Continuous Learn Formative CLA-1 Average of unit test (50%)			CL	Learning A-2 %)	Summative Final Examination (40% weightage)					
		Theory	Practice	Theory	Practice Practice	Theory	Practice				
Level 1	Remember	15%	-	15%		15%	-				
Level 2	Understand	25%		20%		25%	-				
Level 3	Apply	30%	Activities	25%	1/2	30%	-				
Level 4	Analyze	30%	47.5	25%	400	30%	-				
Level 5	Evaluate	/~ ·	A Section 2787	10%		-	-				
Level 6	Create			5%			-				
	Tota <mark>l</mark>	100	0%	100) %	10	0 %				

Course Designers	ままず (**) ** ** ** ** ** ** ** ** ** ** ** **	3 7.
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
1. Dr. S. Sam Gunasekar, Orchid Chemicals and Pharmaceuticals	1. Prof. K Subramaniam, IITM, Chennai,	1. Dr. V. Vinoth Kumar, SRMIST
Ltd., Chennai., sam@orchidpharma <mark>.com</mark>	suubu@iitm.ac.in	
2. Dr. G. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai,	2. Prof. R. B. Narayanan, Anna University, Chennai	2. Dr. P. Radha, SRMIST
ramchand@saksinlife.com	arbeen09@gmail.com	

Course	21BTC205I	Course	BIOPROCESS PRINCIPLES LABORATORY	Course	_	PROFESSIONAL CORE	L	Т	Р	C	
Code	21010200L	Name	DIOPROCESS PRINCIPLES LABORATORY	Category		PROFESSIONAL CORE	0	0	4	2	

Pre-requisite Courses	Ni	Co- requisite Courses	Nil	Progressive Courses	Nil
Course Offering	g Department	Biotechnology	Data Book / Codes / Standards		Nil
			THE NAME OF THE OWNER,		

Course L	earning Rationale (CLR):	The purpose of learning this course is to:	**	17	4			Progr	am Oı	itcome	s (PO))					rogra	
CLR-1:	describe the basics of the	ne fermentation pro <mark>cess</mark>		1-1	2	3	4	5	6	7	8	9	10	11	12		pecifi utcom	
CLR-2:	CLR-2: explain the process of media formulation and sterilization kinetics					of	SI	1	-			Work		9				
CLR-3:					W	Jent	ation	Usage	ъ	. 1				Finan	ning			
CLR-4:			Knowledge	Analysis	velopment	restigations problems		r and	∞ >		Team	ion	& Fi	ā				
CLR-5:			ering		deve	.≦ ×	T00	engineer stv	ment ability		<u>∞</u>	ommunication	Mgt.	g Le				
	<u>'</u>		741	9	roblem	ign/	comple	Modern		rironme stainab	SS	Individual	nuı	roject	Long	7	7-5	5
Course C	Outcomes (CO):	At the end of this course, learners will be able to:	Salah Salah	Engi	Pro	Des	o d	Mod	The	Envi S <mark>us</mark>	Ethics	lpdi	Con	Proj	Life	PSO	PSO.	PSO
CO-1:	understand the basics of	f the f <mark>ermenta</mark> tion process	413	1	13	2		-	7	-		-	-	-	-	2	2	2
CO-2:	comprehend the proces	s of <mark>media fo</mark> rmulation and sterilization kinetics	B. 1	2	2	2	2	2	4-	-	-	-	-	-	-	-	2	1
CO-3:	acquire the basics of re-	acto <mark>r design</mark> and its control systems		2	4/22	2	1	2		-	-	-	-	-	-	2	2	1
CO-4:	evaluate the metabolic stoichiometry and energetics of the biochemical process		3	3	-1	2	-	-	-		-	-	-	-	2	-	-	
CO-5:	explore the various types of reactors for suspension and immobilized cell systems		TE 113	3	12	- 2	2	3		-	-	-	_	-	-	2	2	2

Unit-1 - Microbial Cell Factories

Practice:

- 1. Estimation of glucose by DNS assay method
- 2. Production of enzymes by solid state fermentation
- 3. Production of enzymes by submerged fermentation
- 4. Effect of pH and temperature on enzyme activity

Unit-2 - Design and Preparation of Media for Bioprocess

Practice:

- 1. Batch sterilization kinetics
- 2. Measurements of Cell Biomass Concentration
- 3. Medium optimization by Plackett Burman design

Unit-3 - Bioprocess Design - Instrumentation and Control Systems

Practice:

- 1. Fermenter operation Demonstration/Explanation
- 2. Methods of measuring process variables during yeast fermentation in fermenter

12 Hour

Unit-4 - Fundamentals of Biological Engineering

12 Hour

Practice:

- 1. Microbial growth kinetics to determine the doubling time
- 2. Microbial growth kinetics to determine the yield coefficient
- 3. Enzyme kinetics Michaelis Menten Kinetics and Lineweaver Burk Plot

Unit-5 - Bioreactors for Suspension and Immobilized Cultures

12 Hour

Practice:

- 1. Preparation of immobilized cells/ enzyme
- 2. Enzyme immobilization kinetics
- 3. Production of ethanol by yeast

Learning
Resources

1. Debabrata Das, Debayan Das," Biochemical Engineering- A Laboratory Manual" Jenny Stanford Publishing, 2021.

Learning Asses	ssment) <u>/</u>	- 50 L 50%	Pit of Cont.						
	_		end hour	Continuous Learnin)	4	Sumi	mative		
	Bloom's Level of Thinking	expe	ge of first cycle riments 0%)	exper	of second cycle iments 0%)		Examination reightage)	Final Examination (0% weightage)			
		Theory	Practice	Theory	Practice	Theory	Practi <mark>ce</mark>	Theory	Practice		
Level 1	Remember		15%	 355 557 	15%	74 - 1 S	15%	-	-		
Level 2	Understand		20%	100	20%	A . 4	20%	-	-		
Level 3	Apply	-	25%	7 3 FL 2	25%	112 22	25%	-	-		
Level 4	Analyze	-	25%	Commence of the Commence of th	25%	Crant -	25%	-	-		
Level 5	Evaluate	-	10%		10%		10%	-	-		
Level 6	Create		5%		5%	-	5%	-	-		
	Total		0 %	10	0 %	10	00 %		-		

Course Designers		4 ¹
Experts from Industry	Experts from Higher Technical Institutions	Internal E <mark>xperts</mark>
1. Dr. S. Sam Gunasekar, Orchid Chemicals and Pharmaceuticals	1. Prof. K Subramaniam, IITM, Chennai,	1. Dr. <mark>M.Venkate</mark> sh Prabhu, SRMIST
Ltd., Chennai.sam@orchidpharma.com	suubu@iitm.ac.in	5 /6/
2. Dr. G. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai,	2. Prof. R. B. Narayanan, Anna University, Chennai	2. Dr. Vinoth kumar, SRMIST
ramchand@saksinlife.com	arbeen09@gmail.com	

Course	21BTC206T Course	CENETICS AND CYTOGENETICS	Course	PROFESSIONAL CORE	L	Т	Р	C	;
Code	Name	GENETICS AND CYTOGENETICS	Category	PROFESSIONAL CORE	3	0	0	3	j

Pre-requisite Courses	N	Co- requisite Courses	Nil Progre	Nil
Course Offeri	ng Department	Biotechnology	Data Book / Codes / Standards	 Nil

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Course L	earning Rationale (CLR):	The purpose of learning this course is to:		4		. "	Progr	am Ou	itcome	s (PO))					rograi	
CLR-1:	describe the fundamental	Laws of Genetics and interaction of genes	1	2	3	4	5	6	7	8	9	10	11	12		pecifi Itcom	
CLR-2:	explain the concepts and	explain the concepts and experiments i <mark>n the prepa</mark> ration of linkage map				ટા			1		Work		8				
CLR-3:	describe the elements of Genetic Counseling				velopment	vestigations problems	Usage	ъ					Finance	ning			
CLR-4:	analyze gene transfer and its role in mapping in bacteria				udo	estig	l Us	r and	∞ >		Team	tion	& ∃	arni			
CLR-5:	differentiate factors that lead to genetic variation in a population		neering	Analysis	deve		Tool	gineer	ment ability	h.	<u>8</u>	Communication	Mgt.	g Le			
			inee	roblem	lign/	nduct in complex	Modern	er er	ironm tainal	S	ndividual	nwu	roject	Lon	7-1	7-7	
Course C	outcomes (CO):	A <mark>t the en</mark> d of this course, learners will be able to:	Engi	Pro	Des	5 6	Moc	The	Sus	Ethic	<u>la</u>	S	Proj	Life	PSO	PSO.	PSO-
CO-1:	analyze the pattern of inheritance of genes and its interaction				2	2	Ŧ	1	-	- 1	-	-	-	-	3	-	-
CO-2:	construct linkage maps from inheritance pattern of different genes		3	3	3	2			-	<u> </u>	-	-	-	-	3	-	-
CO-3:	illustrate the role of Gene	tic Counselor and techniques in genetic testing	3	2	2	3	- (_	-	i-	-	-	-	-	3	-	-
CO-4:	illustrate gene mapping based on the type of recombination in Bacteria			3	3	2	-	-	-	-	-	-	-	-	2	-	-
CO-5:	analyze genetic variations in a population				Tarre	2	- 5		-	-	-	-	-	-	2	-	-

Unit-1 - Pattern of Inheritance and Gene Interaction

9 Hour

Mendel's Experiments - Law of segregation, Law of independent assortment - Problems in Mendelian inheritance; Allelic interaction - Lethal genes, Non-allelic interaction - Epistasis, Duplicate genes, Complementary and inhibitory genes; Multiple allelism - ABO, Rh factor in Humans; Cytoplasmic inheritance; Mechanisms of sex determination and sex linked inheritance; Epigenetics - histone modification, methylation - x-inactivation, dosage compensation, Lyon hypothesis

Unit-2 - Linkage and Chromosome Mapping

9 Hour

Chromosome structure, Chromosome organization, Giant chromosomes - polytene chromosome, Lampbrush chromosome; Linkage - Arrangement and types of linkage; Crossing over - Frequency of recombination, Cytological basis of crossing over - Stern's experiment; Chromosome mapping - Mapping by two factor cross, Mapping by three factor cross, Interference and Coincidence, Solving Problems, Combining of map segments, Preparation of linkage map; Somatic cell hybridization - HAT selection procedure

Unit-3 - Basic Human Genetics

9 Hour

Mutation - classification, structural chromosomal aberration - deletion, duplication-tandem and dispersed repeats, inversion, translocation; Numerical aberration; Genetic counseling - History and pedigree construction - Autosomal and X-linked, Diagnosis - Human karyotype preparation, FACS, FISH, Counseling, Follow-up - Prenatal diagnosis - amniocentesis, chorionic villus sampling; Multifactorial inheritance - congenital malformation, diabetes, comparative genome hybridization

Unit-4 - Bacterial Genetics

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Bacterial genetics, Mechanisms of recombination, Transformation in bacteria - Mapping by transformation, Recombination by generalized transduction - Mapping by generalized transduction, Specialized transduction by lambda phage - Mapping by specialized transduction; Recombination by conjugation - Mapping by Interrupted mating analysis, Preparation of linkage map in bacteria, Fine structure mapping by Merozygote analysis

Unit-5 - Population Genetics 9 Hour

Population genetics, Allele frequency - Calculation of allele frequency in a population, Calculation of genotype frequency - Hardy-Weinberg equilibrium, Applications of Hardy Weinberg equilibrium; Changes in allele frequency - Changes in allele frequency by mutation, changes in allele frequency by migration - migration dynamics, changes in allele frequency by selection - selection dynamics, Random genetic drift - Loss of heterozygotes, Genetic equilibrium

Learning	1.	Gardner, Simmons, Sunstad, "Principles of Genetics," 8 th edition	- John Wiley and Sons,	2.	Monroe W. Strickberger, "Genetics," 3 rd edition – Phi Learning, 2015
Resources		Inc., 2006.	COLUMN TO A STREET	3.	Peter Sunstad and Michael Simmons "Principles of Genetics" 7th edition, Wiley, 2015

Learning Assessm	ent		3	~ A A						
			Sumi	mative						
	Bloom's Level of Thinking	CLA-1 Avera	native ge of unit test 0%)	Life Long CLA (10	1-2	Final Examination (40% weightage)				
	/ 2 /	Theory	Practice	Theory	Practice	Theory	Practice			
Level 1	Remember	15%	S. J. S. W.	15%		15%	-			
Level 2	Understand	25%		20%	V-2	25%	-			
Level 3	Apply	30%	Marine Medicine in	25%		30%	-			
Level 4	Analyze	30%	Charles " All a mar	25%		30%	-			
Level 5	Evaluate			10%	- 1		-			
Level 6	Create		Service Committee and the	5%			-			
	T <mark>otal —</mark>	100	0 %	100	%	10	0 %			

Course Designers		
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
1. Dr. C. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai,	1. Prof. K Subramaniam, IITM, Chennai,	1. Dr. S. Barathi, SRMIST
ramchand@saksinlife.com	subbu@iitm.ac.in	
2. Dr. Karthik Periyasamy, Scientist, Biocon,	2. Prof. R. B. Narayanan, Anna University, Chennai	2. Dr. K.T. Ramya Devi, SRMIST
karthik.periyasamy@biocon.com	arbeen09@gmail.com	

Course	21BTC207T Course	MOLECUL AR RIOLOGY	Course	`	PROFESSIONAL CORE	L	Т	Р	С	
Code	Name	MOLECULAR BIOLOGY	Category	,	PROFESSIONAL CORE	3	0	0	3	

Pre-requisite Courses	N	Co- requisite Courses	Nil	Progressive Courses	Nil
Course Offering	g Department	Biotechnology	Data Book / Codes / Standards		Nil

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Course L	Learning Rationale (CLR): The purpose of learning this course is to:		4		. "1	Progr	am Ou	itcome	s (PO)					gram	1
CLR-1:	know the structures of nucleic acids and th <mark>eir role as he</mark> reditary materials	-1	2	3	4	5	6	7	8	9	10	11	12		ecific comes	
CLR-2:	adopt the structure of nucleic acids for their expression and regulation	dae	,	Jo	SI					ork		8				
CLR-3:	explain the basis and mechanism of protein synthesis and activity			nent	gations ems	Usage	ъ	N.		am W		inan	ρ			
CLR-4:	understand the regulatory role of nucleic acids in cell functioning	Knowle		elopment	estiga	US	r and	∞ × >		Теаг	ion	& Fi	arning			
CLR-5:	scrutinize the controlling events of gene expression under anabolic and catabolic conditions	ering	, A	deve	.≦ ≼	Tool	ngineer 'y	nment		al &	mmunication	Mgt.	ng Le			
		_ e	<u> </u>	lign/	ompl	dern	et ei	io,	S	Individua	JIII	Project	Long	7	2 2	
Course C	Outcomes (CO): At the end of this course, learners will be able to:	Eng.	, E	Des	ည်	Moc	The	Sus	Ethics	Indi	Son	Proj	Life	PSO	PSO PSO	
CO-1:	reminisce the structure of nuc <mark>leic acid</mark> s at the DNA and RNA levels	-	3		-	-	7-	-		-	-	-	-	-	2 3	
CO-2:	comprehend the analysis of functioning of nucleic acids	1.5	2	2		-4		-	<u> </u>	-	-	-	-	-	2 2	
CO-3:	relate the expression of DNA at the different levels	3	West.	1	-		-	-	-	-	-	-	-	-	3 3	
CO-4:	assess the mechanisms of protein synthesis with the genetic code	- 3	2	3	-	-	-	-	-	-	-	-	-	-	3 3	
CO-5:	invoke the various regulatory elements and mechanisms controlling gene expression	3	2	2	1	- 1	-	-	-	-	-	-	-	-	3 3	1

Unit-1 - Structure and Composition of Nucleic Acids

9 Hour

Genetic information and its perpetuation; Development of molecular biology; History of nucleic acids; Landmark experiments of DNA as the genetic material; Modes of DNA replication; DNA constituents; DNA structure and its stability; DNA models; A-, B- and Z-DNA forms; Central dogma; DNA topology

Unit-2 - Replication and Repair of DNA

9 Hour

Basic rules for replication; Chemistry of DNA synthesis; Types and the mechanisms of DNA replication; Replication enzymes; DNA polymerases in prokaryotic and eukaryotic replications; Proof reading activity of DNA polymerase; Topoisomerases; Events in the replication fork; Models of DNA replication; DNA repair mechanism

Unit-3 - Transcription and Post Transcription

9 Hour

Basic features of RNA synthesis; RNA polymerases; Types and function of RNA polymerases; DNA promoters- structure and function; Epigenetics Fundamentals; RNA transcription; Transcription of mRNA, rRNA, and tRNA genes; RNA processing; Posttranscriptional modifications of mRNAs; RNA editing-RNAi and miRNAs

Unit-4 - Translation and Post Translation

9 Hour

Coding of genetic information; Outline of translation; Translation in prokaryotes and eukaryotes; Polyribosome; Posttranslational modifications; Protein folding and sorting; Protein targeting into mitochondria and nucleus;

Unit-5 - Gene Regulation

9 hour

General aspects of Regulation; Gene regulators; Silencers and Enhancers; Operons; Positive and negative gene regulations; The operon models; Lac, Trp, Ara and Gal operons and their regulations

Learning Resources

1. Robert Weaver, Molecular Biology, McGraw-Hill, 2011

2. James D Watson, Molecular Biology of Gene, Pearson Publisher, 2017

			Continuous Learning	Assessment (CLA)		C	mative				
	Bloom's Level of Thinking	CLA-1 A <mark>vera</mark>	native ge of unit test 0%)	CL	g Learning _A-2 <mark>0%)</mark>	Final Examination (40% weightage)					
		Theory	Practice	Theory	Practice	Theory	Practice				
Level 1	Remember	15%	OTTEN	15%		15%	-				
Level 2	Understand	25%		20%		25%	-				
Level 3	Apply	30%	5	25%		30%	-				
Level 4	Analyze	30%	-	25%		30%	-				
Level 5	Evaluate		-	10%		-	-				
Level 6	Create		- A - A A	5%	2	<u>-</u>	-				
	Total	10	0 %	10	00 %	10	0 %				

Course Designers	A SAME SWILL ST.	
Experts from Industry	Experts from Higher Technical Institutions	Internal Exp <mark>erts</mark>
1. Dr. S. Sam Gunasekar, Orchid Chemicals and Pharmaceuticals	1. Dr. Aravind Rengan, Indian Institute of Technology Hyderabad.	1. Dr. N. Selvamurugan, SRMIST
Ltd., sam@orchidpharma.com	aravind@bme.iith.ac.in	
2. Dr. D. Gunaseelan, BIOCON Ltd.,	2. Dr. K. Subramanian, Indian Institute of Technology Madras.	2. Dr. S. Barathi, SRMIST
guna.sachin@gmail.com	subbu@iitm.ac.in	

Course Code	21BTC208L	Course Name	MOLECULAR	BIOLOGY LABORATORY		ourse egory	С			F	PROFE	SSIO	NAL C	ORE			L 0	T 0	P 4	C 2
Pre-requis		Nil	Co- requisite Courses	Nil		Progra		е						Nil						
Course C	Offering Departme	ent	Biotechnology	Data Book / Codes / S	Standards		٠.,	٠.,				I	Nil							
Course Lo	arning Rationale	(CLD): The	purpose of learning this	course is to:	V				-	roara	m Out	oomo	c (DO)					Dro	gram	n
CLR-1:			s DNA in p <mark>rokaryotes</mark>	course is to.		1	2	- 3	4	5	6	7	8	9	10	11	12	Sp	ecific	С
	Ů			W.		-1.1		3	4	5	0	1	0		10		12	Out	come	25
CLR-2:	evaluation of the					edge		t of	Sus			l.		Nork		& Finance				
CLR-3:			il el <mark>ement and</mark> gene transc		A	owle	.S	mer	gatio	sage	and			am V	_	-ina	ing			
CLR-4:	dissection of extra	achromosomal e	le <mark>ment and</mark> gene transcrip	ts		N K	alys	elop	/esti) ()	era	nt &		Te	ation	∞ .:	Learning			
CLR-5:	know DNA dama	ge in prokaryote	s	Siz 37	277	Engineering Knowledge	Problem Analysis	Design/development of solutions	Conduct investigations of complex problems	Modern Tool Usage	engineer ety	Environment 8 Sustainability		ndividual & Team Work	Communication	Project Mgt. 8	Jg L			
						gine	ppler	sign	ng lug	derr	The en society	viror stair	Ethics	ividı	mm	ject	Life Long l	PS0-1	PS0-2	PSO-3
	tcomes (CO):		<mark>ne en</mark> d of this course, lea	rners will be able to:	1 4 12	굡		Designation	ರ ಕ	Mo	The	E S	击	pu	ပိ	Prc	Life	PS		
CO-1:			<mark>cellu</mark> lar organisms		N 783 1	- 1	. 3			+	/-	-		-	-	-	-	-	2	3
CO-2:	comprehend the i	isolation an <mark>d ch</mark> a	<mark>arac</mark> terization of genetic m	aterials	Back.	3	2	2	-	-4		-	-	-	-	-	-	-	2	2
CO-3:	retrospect the ger	netic mater <mark>ials a</mark>	<mark>t di</mark> fferent levels			3	25	1	- E			-	-	-	-	-	-	-	3	3
CO-4:	relate the co-exis	tence of th <mark>ese n</mark>	n <mark>ate</mark> rials		1 4 1/19	-3	3	3	-	-	-	-	-	-	-	-	-	-	3	3
CO-5:	invoke the geneti	c defect ca <mark>usin</mark> g	cell death		Or or the	3	3	3	-	- 5	-	-	-	-	-	-	-	-	3	3
						35	34			- 1										
Unit-1 - Ge Practice:	nomic DNA Isola	tion and An <mark>alys</mark>	sis	Year and the second	, 4					-0		-							12 H	lour
	of Genomic DNA t	from E.coli								_	7									
	ive Analysis of Ge			. 10						7										
	e Analysis Genon		<u> </u>	A)																
Unit-2 - Pla Practice:	smid DNA Isolati	on and Analysi	's						- <		+	" /							12 H	Iour
	Practice: 1. Isolation of Plasmid DNA from E.coli 2. Quantitative Analysis of Plasmid DNA																			
	ive Analysis of Pla			Ammer, to	WAL.	· 1.	Εď	۱IJ												
	re Analysis of Plas																		12 H	Ja
Practice:	tal RNA Isolation	anu Anaiysis							4										12 П	iour
1. Isolation	of Total RNA from																			
	ive Analysis of To																			
3. Qualitati	e Analysis of Tota	I KNA																		

Unit-4 - DNA Cloning Enzymes

12 Hour

Practice:

- 1. Restriction Enzyme Digestion of DNA
- 2. Ligation of DNA Fragment into Plasmid
- 3. E.coli Transformation

Unit-5 - DNA Damage

12 Hour

Practice:

1. Effect of UV radiation on Bacterial Growth

Learning Resources 1. Molecular Cloning, A Laboratory Manual by M. R. Green and J. Sambrook, 2012, Cold Spring Harbor Laboratory Press

2. Molecular Biology Techniques, A Classroom Laboratory Manual, 2019, Elsevier Press

	and the second second			Continuous Learning	g Assessment (CLA)	(40)			
	Bloom's Level of Thinking	expe	ge of first cycle riments 0%)	experi	of second cycle iments 0%)		eightage)		amination ightage)
		Theory	Practice	Theory	Practice	Theory	Prac <mark>tice </mark>	Theory	Practice
Level 1	Remember	-	15%	A 11 4 A 11 A 11	15%	Lain -	15%	-	-
Level 2	Understand	GA I	20%	(10 map) 10 /	20%	7 4 7 7	20%	-	-
Level 3	Apply	1-2	25%	 A) 1/2 (1) (1) 	25%	# VE	25%	-	-
Level 4	Analyze		25%	100	25%	ALL ALL	25%	-	-
Level 5	Evaluate		10%	The same of a	10%	100	10%	-	-
Level 6	Create	-	5%	to the state of	5%	- 24	5%	-	-
	Total	10	0 %	100	0 %	10	0 %		-

Course Designers			
Experts from Industry	Experts from Higher Technical Institutions	Internal Exp <mark>erts</mark>	
1. Dr. S. Sam Gunasekar, Orchid Chemicals and Pharmaceuticals	1. Dr. K. Subramanian, Indian Institute of Technology Madras.	1. Dr. N. Selvamurugan, SRMIST	
Ltd., sam@orchidpharma.com	subbu@iitm.ac.in		
2. Dr. D. Gunaseelan, BIOCON Ltd.,	2. Dr. Sudha Warrier, Professor and Dean, Manipal University,	2. Dr. S. Barathi, SRMIST	
guna.sachin@gmail.com	sudha.warrier@mannipal.edu		

Course	21RTC200T	Course	BIOPROCESS ENGINEERING	Course	_	PROFESSIONAL CORE	L	Т	Р	С
Code	21B1C2091	Name	BIOFROCESS ENGINEERING	Category)	PROFESSIONAL CORE	3	0	0	3

Pre-requ	Λ	Co- requisite	Progressive	Nii
Course	s	" Courses	Courses	IVII
Course	Offering Department	Biotechnology	Data Book / Codes / Standards	Nil

Course L	earning Rationale (CLR):	The purpose of learning this course is to:	W	11	4			Progr	<mark>am O</mark> ı	itcome	s (PO)					ogran	
CLR-1:	enumerate the Ideal and I	Ion- Ideal Reacto <mark>rs</mark>		1	2	3	4	5	6	7	8	9	10	11	12		pecific tcome	
CLR-2:	LR-2: discuss the fluid flow and its mixing in the reactor			dge		of	SL		L T			Work		9				
CLR-3:	CLR-3: explain the mass and heat transfer in the reactor, and scale up in Bioreactor			Knowledge	S	nent	ation	Usage	ъ			N N		Finance	ng			
CLR-4:					Analysis	ldoli	vestigations problems	l Us	r and	∞ ×		Team	figur	∞ర	arni			
CLR-5:	.R-5: discuss modern tools in Bioprocess Engineering			ering	_	n/development	t inv	Modern Tool	engineer sty	ronment tainability		<u>8</u>	Sommunication	Mgt.	ig Le			
			- 7/4	9	roblem	.ಠ.≌	onduct in f complex	Jern	enç ety	io <mark>tai</mark>	cs	ndividual	l E	Project	Long	7	7-2	2
Course C	Outcomes (CO):	At the end of this course, learners will be able to:	Table Sec.	Engi	Po	Des	Con	₩ W	Soci	Sus	Ethics	ng.	S	Proj	Life	PSO.	PSO-2	PSO-3
CO-1:	understand the ideal and i	non <mark>-ideal sy</mark> stems in bioprocess engineering	64 - 13.	3	- 3	3	-			-	-	-	-	-	-	1	-	-
CO-2:	gain knowledge on fluid flo	w <mark>and its m</mark> ixing property	Mar.	3	2	1	2	-	4	-	- 1	-	-	-	-	2	2	-
CO-3:	acquire knowledge in trans	sp <mark>ort phen</mark> omena and scale up studies		3	2	1	1	-	-	7 -	- 6	-	-	-	-	2	2	2
CO-4:	CO-4: understand structured and Unstructured models		1, 1	2	1	. 3	1	-	-	-		-	-	-	-	2	-	-
CO-5:	apply modern tools in modelling of bioprocess system			1	_ 1-	3	3	3	-	-		-	-	-	-	2	2	-

Unit-1 - Ideal and Non-Ideal Bioreactors

9 Hour

Ideal Batch, Fed-Batch, Continuous, Enzymatic catalyzed reaction in CSTR, CSTR with Recycle, Ideal Plug flow reactor. Reactors with Nonideal mixing-mixing times in RTD, Models for Non-ideal reactors-Tanks in Series Model- Dispersion models.

Unit-2 - Fluid Flow and Mixing in Bioreactors

9 Hour

Classification in fluids, Reynolds Number, Viscosity, Momentum Transfer, Non-Newtonian fluid, Rheological Properties of Fermentation Broths, Factors Affecting Broth Viscosity, Mixing- Power Requirements for Mixing- Scale-Up of Mixing Systems- Improving Mixing in Fermenters- Effect of Rheological Properties on Mixing- Role of Shear in Stirred Fermenters

Unit-3 - Transport Phenomena and Scaleup in Bioreactors

9 Hour

Gas liquid mass transfer in cellular systems, Determination of Oxygen Transfer Rates, Forced Convection mass transfer, Correlation for Mass Transfer Coefficients, and Interfacial areas. Heat Transfer correlations. Scale up concerns in Microbial, Mammalian and plant cell Process-Scale up criteria-Selection of scaleup criteria-scaleup of genetically engineered cell culture fermentation.

Unit-4 - Models in Bioprocess

9 Hour

Model classification- Model Formulation- Unstructured Mo<mark>dels- Phase</mark>s of batch growth cycles-Monod Models-Multiple substrate models and model Inhibition, Models of growth and non-growth product inhibition, Models for the growth of fungi, Plant cell and Animal cells, Structured models- Models of metabolites and growth-compartmental Models-Models of product formation.

Unit-5 - Modelling and Simulation in Bioprocessing

9 Hour

Introduction to modelling and Simulation. Modelling and simulation of Batch, Fed-Batch and Continuous system using MATLAB. Artificial Intelligence and Machine Learning in bioprocessing. Introduction of object-oriented modelling in bioprocess using Python.

Learning	 James E.Bailey, David F.Ollis "Biochemical Engineering Fundamentals", 2nd Edition Mc Graw Hill. 1986.
Resources	2. Pauline M. Doran "Bioprocess Engineering Principles", 2nd Edition, Academic pres

- S.N.Mukhopadhyay "Process Biotechnology Fundamentals", 2nd Edition, 2004.
 Michael L. Shuler, Fikret Kargi, Matthew De Lisa "Bioprocess Engineering: Basic Concepts", 3rd Edition, Prentice-Hall, 2017.
- 5. Ravindra Pogaku, "Horizons in Bioprocess Engineering" Springer, 2019

	Bloom's Level of Thinking	CLA-1 Avera	Continuous Leaming native ge of unit test 0%)	CL	g Learning LA-2 0%)	Summative Final Examination (40% weightage)					
		Theory	Practice	Theory	Practice	Theory	Practice				
Level 1	Remember	15%		15%	2 -	15%	-				
Level 2	Understand	25%	ALC: U.S.	20%	- V	25%	-				
Level 3	Apply	30%	27 (77) (4)	25%	100 m	30%	-				
Level 4	Analyze	30%	Sec. 277	25%		30%	-				
Level 5	Evaluate			10%		-	-				
Level 6	Create		A 1981 WAREHOUSE	5%		-	-				
	Tot <mark>al</mark>	100	0%	- 10	00 %	10	0 %				

Course Designers		
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
1. Dr. S. Sam Gunasekar, Orchid Chemicals and Pharmaceuticals	1. Dr.S.Senthil Kumar, IITG	1. Dr.M.Venkatesh Prabhu, SRMIST
Ltd., Chennai,sam@orchidpharma.com		
2. Dr. G. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai,	2. Dr.N.Selvaraj, IITG	2. Dr.P.Radh <mark>a, SRMI</mark> ST
ramchand@saksinlife.com		

	Course Code	21BTC210L	Course Name	BIOPROCESS ENGINEERING LABORATORY	Course Category	С	PROFESSIONAL CORE	L T P C
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Pre-requisite Courses	Co- requisite Courses	Nil Progressive Courses	Nil
Course Offering Department	Biotechnology	Data Book / Codes / Standards	Nil

Course Le	earning Rationale (CLR): The purpose of learning this course is to:		7			Progra	<mark>am</mark> Ou	tcome	es (PO))					ogran	
CLR-1:	explain the Residence Time Distribution in Stirred tank and Plug flow reactor	1	2	3	4	5	6	7	8	9	10	11	12		oecific tcome	
CLR-2:	describe the rheological and mixing behavior of fermented fluid	dge		o	SL					ork		99				
CLR-3:	analyze the oxygen mass transfer coefficient and deactivation kinetics	N N		velopment	stigations	Usage	ъ			ΜW		Finan	Б			
CLR-4:	evaluate the model parameters in microbial growth	穒	alysis	lop	estig		er and	× ×		Team	tion	∞ర	arning			
CLR-5:	discuss the modern tool of programming microbial cultures	iring	. Ana	/deve	t in K	<u>8</u>	engineer sty	meniabilit		रू ज	ommunication	Project Mgt.	g Le			
		inee in	roblem	/ugi	duc	Jern	et e	iron	S	ndividual	nwu	ect	Long	7	2-0Sc	~
Course O	utcomes (CO): At the end of this course, learners will be able to:	Engir	Pro	Des	Sob	Mo	The	Env Sus	Ethi	lndi	Con	Proj	Life	PSO-1	PS(PSO-3
CO-1:	explore the Residence Time Distribution studies in Stirred tank and Plug flow reactor	3	3	2	-	+	7-	-	-	-	-	-	-	2	-	-
CO-2:	understand the rheological and mixing behavior of fermented fluid	. 3	3	1		- 1		-		-	-	-	-	2	2	-
CO-3:	measure the oxygen mass transfer coefficient and deactivation kinetics parameters	3	3	2		- 5		-	-	-	-	-	-	2	2	-
CO-4:	estimate the model parame <mark>ters in m</mark> icrobial growth	- 3	3	1	-		-	-		-	-	-	-	2	2	-
CO-5:	learn the modern tool for programming the microbial cultures	1	2	3	-	3		-		-	-	-	-	2	2	-

Unit-1 - Non-Ideal Reactors 12 Hour

Practice:

- 1. RTD studies in Stirred tank reactor
- 2. RTD studies in Plug flow reactor

Unit-2 - Fluid Flow and Mixing in Bioreactors

Practice:

- 1. Rheological study of fermented fluids
- 2. Regime analysis of a stirred tank reactor
- 3. Determination of mixing time in a stirred tank reactor

Unit-3 - Transport Phenomena and Scale-up in Bioreactors

Practice:

- 1. Determination of KLa by power correlation method
- 2. Determination of KLa by dynamic gassing out method
- 3. Deactivation kinetics of enzymatic reaction
- 4. Deactivation kinetics of microbial growth

12 Hour

12 Hour

Unit-4 - Models in Bioprocess 12 Hour

Practice:

- 1. Estimation of unstructured model parameters of bacterial culture
- 2. Estimation of unstructured model parameters of yeast culture

Unit-5 - Modelling and Simulation in Bioprocessing

Practice:

- 1. Modelling and simulation of Batch culture using MATLAB
- 2. Modelling and simulation of continuous culture using MATLAB
- 3. Modelling and simulation of Fed culture using MATLAB
- 4. Modelling of batch reactor using Python

Learning	1. Hans-Peter Schmauder,"Methods in Biotechnology" Taylor and Francis Ltd, 2003.	Ŧ	3. Shijie Liu, "Bioprocess Engineering Kinetics, Sustainability, and
Resources	Arvind Kumar Bhatt, "Basic Biotechniques for Bioprocess and Bioentrepreneurship" Academic Press, Elsevier, 2023		Reactor Design"Elsevier, 2020.

earning Asse	ssment			18 July 1998	124 CT 16							
			well had	Continuous Learning	g Assessment (CLA)							
	Bloom's Level of Thinking	experi	CLA-1 Average of first cycle experiments (30%)		of second cycle iments 0%)		eightage)	Final Examination (0% weightage)				
	*	Theory	Practice	Theory	Practice	Theory	Practi <mark>ce</mark>	Theory	Practice			
Level 1	Remember		15%	 355 557 	15%	JAN JE	15%	-	-			
Level 2	Understand		20%	1000	20%	A , A	20%	-	-			
Level 3	Apply	-	25%	73.5	25%	172 237	25%	-	-			
Level 4	Analyze	-	25%	the same of the control of the contr	25%	Charles -	25%	-	-			
Level 5	Evaluate		10%		10%	- 1	10%	-	-			
Level 6	Create		5%) (5%	- / /	5%	-	-			
	Total	100) %	10	0 %	10	0 %		-			

Course Designers	1,1,14	
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
1. Dr. S. Sam Gunasekar, Orchid Chemicals and Pharmaceuticals	1. Dr.S.Senthil Kumar, IITG, senthilkumar@iitg.ac.in	1. Dr.M.Venkatesh Prabhu, SRMIST
Ltd., Chennai.sam@orchidpharma.com	DI CARN IN	
2. Dr. G. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai,	2. Dr.N.Selvaraj, IITG, selva@iitg.ac.in	2. Dr.P.Radha, SRMIST
ramchand@saksinlife.com		

Course	21BTC301J	Course	GENE MANIDUL ATION AND GENOMICS	Course	C	PROFESSIONAL CORE	L T P	С
Code	210103013	Name	GENE MANIPULATION AND GENOMICS	Category	C	PROFESSIONAL CORE	3 0 2	4

Pre-requisite Nil	Co- requisite Courses	Nil Progressive Courses	Nil
Course Offering Department	Biotechnology	Data Book / Codes / Standards	Nil
		OID NO.	

Course L	earning Rationale (CLR): The purpose of learning this course is to:	1 1	7		- I	rogra	m Ou	tcome	es (PO)				Pı	rograr	n
CLR-1:	assess the basic concepts and principles of utilization of different expression vectors for cloning from the perspective of engineers	1	2	3	4	5	6	7	8	9	10	11	12		pecifi itcom	
CLR-2:	demonstrate the different strategies of gene cloning and construction of genomic and cDNA libraries	lge		of	SI	1				ork		се				
CLR-3:	analyze the concepts of structural and functional genomics with advanced cutting-edge technologies	owledge	w	Jent	stigations	sage	ъ			Μ		Finan	рu			1
CLR-4:	assess the applications of recombinant DNA technology in animals, plants, and microbial organisms	X S	alysi	elopment	estig	l Us	r and	∞ >		Teal	tion	& Fi	arnii			1
CLR-5:	develop and apply the strategies on altering gene expression in vitro and in vivo	eering	em Ana	n/deve	luct inve	m Too	ngineer v	nment		dual &	Communication	st Mgt.	ong Le	_	2	3
Course O	utcomes (CO): At the end of this course, learners will be able to:	Engin	Proble	Desig	Cond	Mode	The e	Enviro Susta	Ethics	Individual	Comn	Project	Life L	PSO-1	PSO-2	PSO-3
CO-1:	describe the foundations of modern biotechnology	-		3	- 1			-		-	-	-	-	-	2	-
CO-2:	design and conduct experim <mark>ents inv</mark> olving genetic manipulation		54	2	- 1		2	-	1	-	-	-	-	-	-	3
CO-3:	illustrate the steps involved in the production of biopharmaceuticals in microbial and mammalian cell system	s 2	27.	4	-	- 1	=	-	2	-	-	-	-	-	-	3
CO-4:	apply modern biotechnology in the different areas like medicine, microbes, environment, and agriculture	3	-	-4	-		3	-		-	-	-	-	-	-	3
CO-5:	discuss the cutting-edge techniques and their applications such as plant transformation, protein expression and genomic DNA library construction etc.	n 3		2	-	-5		-	2	-	-	-	-	-	-	3

Unit-1 - Overview of Cloning and Vectors

15 Hour

Introduction to genomics and gene regulation; Fundamental requirement for DNA cloning; Prokaryotic and eukaryotic vectors; Phage vectors; Strategies for gene cloning; Enzymes in genetic engineering

Practice: 1. Genomic DNA isolation

2. Double digestion of Genomic DNA

Unit-2 - Preparation and Screening of DNA library

15 Hour

DNA Library; Preparation of DNA Libraries; Genomic DNA library; Overlapping and non-overlapping DNA fragments; Choice of vectors; Evaluation of genomic DNA library; cDNA library; Purification and separation of mRNA; cDNA synthesis; cDNA library construction; Evaluation of cDNA library; Screening libraries; Polymerase chain reaction (PCR) and its applications

Practice: 1. Double digestion of Vector

- 2. Preparation of recombinant vector
- 3. E. coli Transformation

Unit-3 - DNA Sequencing and Genomics

15 Hour

DNA sequencing strategies; Principles of DNA sequencing; Sanger's Dideoxy sequencing method; Automated DNA sequencing; Next generation sequencing; Genome sequencing; Next generation sequencing and its applications; Methods of nucleic acid detection; Random priming; Nick translation and End labeling; RNA labeling; Non-isotopic labeling; Structural genomics; comparative genomics; Microarray

Practice: 1. Colony PCR

2. Functional Assay

Unit-4 - Analysis and Manipulation of Gene Expression and Function

15 Hour

Regulation of gene expression at different levels; Factors influencing gene expression; Epigenetic regulation; Protein expression in prokaryotic and eukaryotic cells; Alteration of gene expression by mutagenesis; Methods for site directed mutagenesis

Practice: 1. RNA isolation

2. cDNA synthesis

Unit-5 - Applications of Cloning

3. Semi-quantitative PCR

15 Hour

Medical applications; Human and genetic diseases; DNA vaccines; Gene therapy; Study of gene function in vivo; Embryonic stem cells; Applications in Embryonic stem cells; Transgenics; Methods of producing transgenic mice; Over-expression; Gene knock-in; Gene <mark>knock-out; C</mark>onditional knock-out; Genome editing; CRISPER-Cas9; Guide RNA; Gene inactivation

Practice: 1. Quantitative PCR

2. Fold and Relative Gene Expression

Learning
Learning
Resources
. 10000.000

- 1. Jeremy W. Dale and Malcolm von Schantz, "From Genes to Genomes," John Willey and 3. S. B. Primrose and R. M. Twyman, "Principles of Gene Manipulation and Genomics" 7th Sons Publications, 2002
- 2. Old. R.W and Primrose. S.B, "Principles of Gene Manipulation, An Introduction to Genetic Engineering," Blackwell Scientific Publications, 2014
- Edition, Wiley-Blackwell, 2006
- 4. T A Brown Gene Cloning and DNA Analysis: An Introduction 8th Edition, Wiley Blackwell Publisher 2020

Learning Assessm	ent			A Property of	A William					
	Blo <mark>om's</mark> Level of <mark>Thinkin</mark> g	VS	CLA-1 Averag	Continuous Learnin ative ge of unit test %)	Summative Final Examination (40% weightage)					
	-		Theory	Practice	Theory	Practice	Theory	Practice		
Level 1	Remember		15%	The fact that the N	F 12 - 12 - 12 - 12	15%	15%	-		
Level 2	Understand		25%			20%	25%	-		
Level 3	Apply	-	30%	1		25%	30%	-		
Level 4	Analyze		30%	- 1134	-	25%	30%	-		
Level 5	Evaluate	-	-	- 1.9	-	10%	-	-		
Level 6	Create			- 1/ 1/	-	5%		-		
	Total	1	100) %	100	0%	100	0 %		

Course Designers	S. C. S.	
Experts from Industry	Experts from Higher Technical Institutions	Inter <mark>nal Exper</mark> ts
1. Dr. C. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai,	1. Prof. K Subramaniam, IITM, Chennai, subbu@iitm.ac.in	1. Dr. N. Selvamurugan, SRMIST
ramchand@saksinlife.com	DI LILII LIV	
2. Dr. Karthik Periyasamy, Scientist, Biocon,	2. Prof. R. B. Narayanan, Anna University, Chennai	2. Dr. S. Barathi, SRMIST
karthik.periyasamy@biocon.com	arbeen09@gmail.com	. * /

Course	21BTC302J	Course	IMMUNOLOGY	Course	C	PROFESSIONAL CORE	L	Т	Р	С
Code	210103023	Name	IIVIIVIONOLOGT	Category	C	PROFESSIONAL CORE	3	0	2	4

Pre-requisite Courses	Ni	Co- requisite Courses	Nil	ressive urses	Nil
Course Offeri	ng Department	Biotechnology	Data Book / Codes / Standards		Nil

Course L	earning Rationale (CLR): The purpose of learning this course is to:	_A			- I	rogra	m Ou	tcome	s (PO)				_	ogran	
CLR-1:	introduce the science of immunology and a detailed study of various types of immune cells	1	2	3	4	5	6	7	8	9	10	11	12	_ '	pecific tcome	
CLR-2:	provide knowledge about immune systems produced molecules and their classification, structure, and function		· .					£								
CLR-3:	provide students with experience in methods used in immunology, particularly the use of specific antibody in biomolecular applications	dge		oę	ns of	A.	society	tainability		Work		ce				
CLR-4:	provide knowledge about major histocompatibility complex and acquired immune system, their cells and its interaction and how they fight against infectious diseases	Knowledge	/sis	velopment	stigations lems	Usage	and so	& Susta	N.	eam W	C.	Finance	ning			
CLR-5:	provide knowledge about dysreg <mark>ulation o</mark> f immune system functioning, ways to strengthen immune system and how human body is designed and protected to fight against various pathogens	ering	m Analysis	g g	ct inves	Tool	engineer	Environment 8		∞ ~	Sommunication	Project Mgt. &	ong Lear			က
Course (Outcomes (CO): At the end of this course, learners will be able to:	Engine	Problem	Design/ solution	Condu	Modern	The el	Enviro	Ethics	Individual	Comm	Projec	Life Lo	PSO-	PS0-2	PSO-3
CO-1:	describe the immune system, their structure, classification and function	12	- 14	2	- 1	- /		-		-	-	-	-	1	-	1
CO-2:	summarize genetic control of antibody diversity, monoclonal antibodies and cellular immunology	251	문진	2	2	2	_	-		-	-	-	-	1	-	2
CO-3:	determine various methods to assess immune function, their application and interpretation of the results	L.	-	-4	2	3	-	-		-	-	-	-	3	-	3
CO-4:	outline major histocompatibility complex, types, function and the role of acquired immune cells signalling and its function	1	4	2	3	-5		-	-	-	1	-	1	2	-	2
CO-5:	categorize hypersensitive immu <mark>ne reac</mark> tion, autoimmunity, vaccination and cancer immunology and Illustrate the processes function to protect hyman body against infective agents	-	-	-	3	4	1	-	- 1	-	-	-	-	1	-	2

Unit-1 - Immune System for Health

15 Hour

Overview of the immune system; Development and differentiation of the hematopoietic stem cells, Myeloid and Lymphoid lineage; Lymphatic system; Lymphoid organs – types; Innate lymphoid cells; Rhesus group types; incompatible blood transfusion and hemolytic disease; Receptors of Innate Immune system; Types of Immune cells, Innate Immunity; Anatomical and Physiological barriers; Acquired Immunity, Clonal selection theory; Comparative immunity - Plant Immune system, Vertebrate and Invertebrate Immune system; Immunogens, Antigens and Haptens; Requirements for immunogenicity; major classes of antigens; antigen recognition by T and B lymphocytes

Practice: 1: Laboratory safety principles and Blood grouping; Agglutination principle, blood group types, 2: Total Leukocyte count; Types of blood cells - Leukocyte counting, 3: Differential Leukocyte count

Unit-2 - Immunity of Secretory Proteins

15 Hour

Immunoglobulin structure, types and function; Antibodies biological and functional properties - Proteolytic digestion of antibodies; Monoclonal antibodies production and applications; B Cell differentiation -B cell receptor structure and B cell signal transduction; Antibody diversity - Light chain synthesis; Heavy chain synthesis;; Cytokine types and function; Cytokine receptor structure; Role of cytokines in diseases; Complement system - Regulation of complement pathway; Role of complement proteins in diseases

Practice: 1. Antigen – Antibody reaction I – Widal test- slide method, 2. Antigen – Antibody reaction II -rapid plasma reagin (RPR) test, 3. Single radial immunodiffusion (SRID) - titer value, zone of equivalence

Unit-3 - Methods to Assess Immune Status 15 Hour

Isolation of immune cells from Human and animals; Antigen- antibody interaction; antibody affinity and avidity; Hemaagglutination reaction - Coombs test – direct and indirect; precipitation reaction; Quantitative Immuno assays; passive Immunodiffusion; Precipitation reaction; Active Immunodiffusion – Rocket immunoelectrophoresis, SDS-PAGE and Western blot; Quantitative Immuno assays - Radio-immunoassay, Immunoprecipitation; Immunofluorescence – Direct and indirect; Immunohistochemistry; flow cytometry, ELISA and types; Cell culture and experimental models, analysis of gene expression

Practice: 1. Ouchterlony gel diffusion - Antigen-Antibody specificity, 2. Active Immunodiffusion I - Rocket Immunoelectrophoresis, 3. Active immunodiffusion – II – Counter Current Immunoelectrophoresis

Unit-4 - T Cell Signalling and Major Histocompatibility Complex

15 Hour

Major histo-compatibility Complex(MHC) – types and function; antigen processing and presentations – Endogenous and Exogenous; Diversity of MHC molecules;; Antigen – Antibody interaction Standard and test antigen; Rocket Immunoelectrophoresis; Biology of T lymphocyte - T cell receptors and interaction with MHC; T-cell maturation - T-cell activation and differentiation; Thymic selection – Positive and negative selection; T-cell activation and cytokine secretion; Cytokine control of TH1 and TH2 CD4+; Function of CD8+ T cells, T Regulatory cells; T-cell and B-cell cooperation, Pathways of Activation

Practice: 1. Enzyme linked Immunosorbent assay (ELISA) – Qualitative, 2. Enzyme linked Immunosorbent assay (ELISA) – Quantitative, 3. Immunoprecipitation

Unit-5 - Immunity of Infection, Autoimmune Disorder and Cancer

15 Hour

Hypersensitive reactions - Type I, Type II, Type III and Type IV reaction; Immune responses to infectious diseases introduction; Viral disease-HIV infection; Bacterial disease-Tuberculosis; Parasitic disease - Malaria; Evading Mechanisms of pathogens; Vaccine history and principle; Active and passive Immunization; DNA vaccine, Edible vaccine and Adjuvants; Cancer Immunology introduction; Evidence for cancer Immunity; cancer Immuno therapy; Autoimmunity introduction; Genetic Basis of Autoimmunity; Classification of auto-immunity

Practice: 1. SDS-PAGE, 2. Western blotting – Demo, 3. Flow cytometry - Demo

Learning	1. Sudha Gangal, Shu <mark>bha</mark> ngi Sontakke	, Textbook of basic and clinical immunology,	2. Jenni Punt, Sharon Stranford, Patricia Jones, Judith A Owen, Kuby Immunology, 8th	ed., W. H.
Resources	Universities Press, 2 <mark>013</mark>		Freeman and Company, 2018	

Learning Assessm	nent	2	Continuous Learnin	g Assessment (CLA)		0				
	Blo <mark>om's</mark> Level of <mark>Thinkin</mark> g	CLA-1 Avera	mative age of unit test 5%)	CLA-2-I	Practice %)	Summative Final Examination (40% weightage)				
		Theory	Practice	Theory	Practice	Theory	Practice			
Level 1	Remember	15%	12.	//	15%	15%	-			
Level 2	Understand	25%	- INT.	-	20%	25%	-			
Level 3	Apply	30%	- 1	-	25%	30%	-			
Level 4	Analyze	30%	- 11111	-	25%	30%	-			
Level 5	Evaluate	1 7 -	- /(\$76)	-	10%	-	-			
Level 6	Create				<i>5</i> %	-	=			
	Total	10	00 %	100)%	10	0 %			

Course Designers	VALUE OF STATE OF SALES	
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
1. Dr. C. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai,	1. Dr. Joe Varghese, CMC Vellore, joevarghese@cmcvellore.ac.in	1. Dr.S.Nageswaran, SRMIST
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2. Dr. Karthik Periyasamy, Scientist, Biocon,	2. Prof. K Subramaniam, IITM, Chennai, subbu@iitm.ac.in	2. Dr.S.Rupachandra, SRMIST
karthik.periyasamy@biocon.com		

Course	21BTC303T	Course	PROTEIN ENGINEERING	Course	C	PROFESSIONAL CORE	L	Τ	Р	()
Code	210103031	Name	PROTEIN ENGINEERING	Category		PROFESSIONAL CORE	3	0	0	:	3

Pre-requisite Courses	N	Co- requisite Courses	Nil Progressive Courses	Nil
Course Offerin	ng Department	Biotechnology	Data Book / Codes / Standards	Nil
			CALLY NO.	

Course L	earning Rationale (CLR):	The purpose of learning this course is to:	. //		<u> </u>			Progr	am Ou	itcome	s (PC))				Pi	rograr	n
CLR-1:	distinguish the organization			1	2	3	4	5	6	7	8	9	10	11	12	_	pecifi itcom	
CLR-2:	appraise the structure-funct	ion correlat <mark>ion in sele</mark> cted proteins		ge		of	ည			l.		å		8				
CLR-3:	understand Mutagenesis ba	sed prot <mark>ein design</mark>		Knowledge	(0	nent	stigations	age	70			, m		Financ	Б			
CLR-4:	construct 3D structure of pr	otein fr <mark>om amin</mark> o acid sequence		Κno	Analysis	lopi	estiga	ool Usage	r and	∞ × >		Tear	ţio	∞ర	arni			
CLR-5:	discuss on the experimenta	I tec <mark>hniques a</mark> vailable for protein structure characterization		Engineering	Ang	ign/development	t inve	-	jineer	ment	N	<u>∞</u>	ommunication	Mgt.	g Le			
		1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		inee	Problem	ign/e	duct	Modern	engine etv	ro Lai		ndividual	JML	Project I	Long	7)-2	53
Course C	Outcomes (CO):	At the end of this course, learners will be able to:	- 42	Eng	Prof	Des	o do	Moc	The		Ethics	Indi	S	Proj	Life	PSO-1	PSO-2	PSO-3
CO-1:	outline proteins and its prop	pe <mark>rties at t</mark> he elemental, molecular and structural levels		- (2			1	7	-	-	-	-	-	-	-	3	-
CO-2:	group the proteins based or	n super secondary structure of protein with its function		Æ,	2	1	-	3	-	-	-	-	-	-	-	-	3	-
CO-3:	integrate protein biochemist	try to design efficient protein structures	40.5	Kir y	2	- 1	<i>-</i>	3	-	-		-	-	-	-	-	3	-
CO-4:	scoring and validating the n	<mark>nethods</mark> of obtain protein structural data	7	4		1	2	3	_	-	4	-	-	-	-	-	-	3
CO-5:	mutagenesis experiments to	o test protein stability and/or function	-	2	-	Les	2	3	-	-	-	-	-	-	-	-	-	3

Unit-1 - Characteristics of Proteins 9 Hour

Structure of amino acids- Properties of amino acids- Role of Glycine and Proline in structure determination- Ramachandran plot and its significance- Interactions that stabilize secondary -Structures, Structural features of alpha helices- Parallel beta-strand structure-Anti-parallel beta-strand structure- Beta turns- loops and other secondary structures- Super- Secondary structures- Difference between motifs & domains- Types of motifs, Types of domains, Monomeric and polymeric proteins- hydrophobic collapse & theories of folding- Levinthal paradox- Role of chaperones- and heat shock proteins

Unit-2 - Structural features of Different Classes of Proteins

9 Hour

Role of Transcription factors in gene - Nature of interaction between p53 and DNA- effect of mutations in the DNA binding domain of p53- Effects of mutations in the oligomerization and Nuclear localization region-Structural elucidation of leucine zipper- Interaction of leucine zipper and DNA- - Structural elucidation of GPCR- Types of GPCR- Mechanism of activation of GPCR- Structural features of serine proteases

Unit-3 - Experimental Protein Structure and Functional Analysis

9 Hour

Methods of generating crystals- (ITC) Principle- Instrumentation of ITC- Determination enthalpy- entropy and free energy- Prediction of binding energy and multiple binding sites by ITC- Prediction of 3D structure from amino acid sequence, Homology modelling and threading

Unit-4 - Increasing Efficacy of Proteins

9 Hour

Protein Engineering in Basic and Applied Biotechnology- engineering new protein function- Engineering enzymes- Specificity- stability- antibodies- Denovo designs Fusion proteins- Protein engineering in Vaccine development- Protein engineering in biosensors- Case Study: Enhancing binding affinity of T4 lysozyme- Enhancing stability in T4 lysozyme

Unit-5 - Protein Expression Purification and Characterization

9 Hour

The isolation and characterization of proteins, Recombinant DNA technology and protein expression- Protein Digestion Techniques- Chemical and Enzymatic- Mass spectrometry - Tandem LC MS-/MS- Tools for mass spectrum analysis

Learning	
Resources	

- 1. Whitford, David. Proteins: Structure and Function. Wiley, 2013.
- 2. Tooze, John, and Branden, Carl Ivar. Introduction to Protein Structure. United States, CRC Press, 2012.
- 3. Ben-Tal, Nir. Kessel, Amit. Introduction to Proteins: Structure, Function, and Motion. United Kingdom: CRC Press, Taylor & Francis Group, 2018.
- 4. Buxbaum, Engelbert. Fundamentals of Protein Structure and Function. Germany: Springer International Publishing, 2015
- Lilia Alberghina, Protein Engineering For Industrial Biotechnology, Taylor & Francis, 2003.
 Chatwal. G. R, "Instrumental methods of Chemical Analysis", Himalaya Publishing House, 5th Edition, 2011.

			Cum	mativa						
	Bloom's Level of Thinking	CLA-1 Avera	native ge of unit test 9%)	CL	g Learning .A-2 0%)	Summative Final Examination (40% weightage)				
		Theory	Practice	Theory	Practice	Theory	Practice			
Level 1	Remember	15%	44, 144	15%	- A- V	15%	-			
Level 2	Understand	25%	10 E 10 E 10 E	20%	() () () ()	25%	-			
Level 3	Apply	30%	10 July 1777	25%		30%	-			
Level 4	Analyze	30%		25%		30%	-			
Level 5	Evaluate		8, THE WAY SEE A 12	10%		-	-			
Level 6	Create	-	Carlotte March 1980	- 5%			-			
	Total	100	0 %	10	0 %	100 %				

Course Designers		
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
1. Dr. C. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai,	1. Prof. K Subramaniam, IITM, Chennai, subbu@iitm.ac.in	1. Dr. Priya Swaminathan, SRMIST
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2. Dr. Karthik Periyasamy, Scientist, Biocon,	2. Prof. R. B. Narayanan, Anna University, Chennai	2. Dr. Vasantharekha R, SRMIST
karthik.periyasamy@biocon.com	arbeen09@gmail.com	

Course	21BTC304T	Course	ANIMAL BIOTECHNOLOGY	Course	C	PROFESSIONAL CORE	L T P	С
Code	210103041	Name	ANIMAL BIOTECTINOLOGI	Category	C	PROFESSIONAL CORE	3 0 0	3

Pre-requisite Courses	N	Co- requisite Courses	Nil	ogressive Courses	Nil
Course Offer	ring Department	Biotechnology	Data Book / Codes / Standards		Nil

															D.		
Course L	earning Rationale (CLR):	The purpose of learning this course is to:		1 .			Progr	am Oı	utcome	es (PC))					ogran pecific	
CLR-1:	provide a basic understanding	of animal bre <mark>eding and</mark> animal health	1	2	3	4	5	6	7	8	9	10	11	12		tcome	
CLR-2:	develop an understanding on r	aising ani <mark>mals using</mark> assisted reproductive techniques	ge		of	SL	1				ork		9				
CLR-3:	CLR-3: inculcate the understanding of cell culture technique and production of valuable products from them						age	ъ			N W		Finan	Б			
CLR-4:	provide an understanding of al	terati <mark>on of ani</mark> mal body biological system	Knowled	Analysis	elopment	estigations problems	l Us	er and	× × ∞		Team	ţion	∞ర	arning			
CLR-5:	CLR-5: give emphasis to transgenesis thereby improving livestock production						18 18	enginee	ironment tainability		ual &	ommunication	Project Mgt.	ig Le			
			ineering	oblem	ign/d	용병	ern	eu €	a Series	S	/idu	חת	ect	Long	7)-2	<u>ښ</u>
Course C	Outcomes (CO):	At the end of this course, learners will be able to:	Eng	Prok	Desi	Col	Mod	The	Sust	EF	Individu	Con	Proj	Life	PSO-1	PS0-2	PSO-3
CO-1:	familiarize the students about using vaccines	breeding, biological markers for genetic diseases and managing animal hea	Ith -	3	3	-		Z	-		-	-	-	-	-	3	3
CO-2:	0-2: impart an understanding about Embryo transfer, fertilization methods and animal production				3	-			-		-	-	-	-	-	3	3
CO-3:	0-3: provide knowledge about different culture techniques, Characterization of cell lines and in vitro testing of drugs			3	2	-	-	-	-		-	-	-	-	3	3	-
CO-4:	O-4: provide knowledge about improvement of animals to increase the yield and quality of animal products				-5	3			-	-	-	-	-	-	3	3	-
CO-5:	1-5: familiarize the students about livestock improvement using molecular pharming			445	-	2	- 5		-	-	-	-	-	-	-	3	3

Unit-1 - Animal Improvement for Desired Traits and Animal Health

9 Hour

Breeding, different types of breeding; Marker assisted Selection - Gene mapping and identification of genes of economic importance in farm animals; Animal Health: Common viral, bacterial and parasitic diseases affecting animals; Vaccines for animal health; Developing diagnostic kits for animal diseases

Unit-2 - Embryo Transfer and Animal Propagation

9 Hour

Assisted reproductive techniques in animals: Artificial insemination; In vitro fertilization- Superovulation, MOET, Embryo transfer, — Pregnancy diagnosis — Sexing of embryos, Embryo splitting; Cryopreservation of embryo; Cloning for conservation of endangered species; Stem cell technology & its applications

Unit-3 - Animal Cell Culture

9 Hour

Principles of sterile techniques and cell propagation — Primary cell culture, secondary cell culture, continuous cell lines, suspension cultures; Chemically defined and serum free media for cell culture; Preservation and characterization of animal cells; Scaling up of animal cell culture; organ culture; 3D printing; Application of animal cell culture in vitro testing of drugs; Cell culture as source of therapeutic protein production

Unit-4 - Biotechnology in Livestock Production

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Manipulation of Growth hormone – somatotropic hormone – Thyroid hormone; Probiotics as growth promoters, Mode of action & uses of probiotics; Manipulation of lactation – Lactogenesis – galactopoiesis; Manipulation of rumen microbial digestive system; Manipulation of wool growth

Unit-5 - Transgenesis and Molecular Pharming

9 Hour

Trangenesis, Gene editing using CRISPR Cas9, Transgenic animals, Methods of producing transgenic animals, knockin, knock out, mutation models; Transgenic animals as models for human diseases; Transgenic animals in livestock improvement- Therapeutic protein expression using transgenic animals, Animal as bioreactors; Ethical issues in animal biotechnology, 3R's and alternative for animal models - In vitro testing & insilico modeling

	1.	Animal Biotechnology: Recent concepts and developments - P.Ramadas, MJ.	Р
Learning		Publications, 2015.	4
Resources	2.	Animal Breeding and Genetics; Aggrey, S.E.; Rekaya, R. Spangler, M.L., Ed.;	
		Springer: New York, NY, USA, 2022.	

- 3. Animal Biotechnology M.M.Ranga, 3rd edition, 2007.
- 4. Culture of Animal cells; a manual of basic technique R.lan Freshney, 4th edition, Wiley publications, 2006.
- 5. Textbook of Animal Biotechnology P.Ramadas & S.Meerarani, 2nd edition, 2002.

Learning Assessm	ent				*, ``					
	Bloom's Level of Thinking	Form CLA-1 Averag	ge of unit test	Life Long	g Learning _A-2 0%)	Summative Final Examination (40% weightage)				
		Theory	Practice	Theory	Practice Practice	Theory	Practice			
Level 1	Remember	15%		15%	2 - 1	15%	-			
Level 2	Understand	25%	4.5	20%	1/2-	25%	-			
Level 3	Apply	30%	47-17-18-18	25%	(P)	30%	-			
Level 4	Analyze	30%	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	25%		30%	-			
Level 5	Evaluate			10%	1-2	-	-			
Level 6	Create			5%		-	-			
	Tot <mark>al</mark>	100)%	- 10	00 %	100	0 %			
			A COLUMN TO A STATE OF		(a)					

Course Designers		
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
1. Dr. G. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai,	Prof. K Subramaniam, IITM, Chennai, subbu@iitm.ac.in	1. Dr.S.Sujath <mark>a, SRMI</mark> ST
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2. Dr. Karthik Periyasamy, Scientist, Biocon,	2. Prof. R. B. Narayanan, Anna University, Chennai	2. Dr.K.Venkatesan, SRMIST
karthik.periyasamy@biocon.com	arbeen09@gmail.com	

Course	210102051	Course	ANIMAL BIOTECHNOLOGY LABORATORY	Course	C	PROFESSIONAL CORE	L T	Р	(;
Code	21D1C303L	Name	ANIMAL BIOTECHNOLOGY LABORATORY	Category	C	PROFESSIONAL CORE	0 0	4	1	.]

Pre-requisite Courses	N	Co- requisite Courses	Nil	ogressive Courses	Nil
Course Offer	ring Department	Biotechnology	Data Book / Codes / Standards		Nil

Course I	Learning Rationale (CLR): The purpose of learning this course is to:		4 5		Ī	Progr	<mark>am</mark> Ou	tcome	s (PO)					ogram	
CLR-1:	provide the basics of cell culture media and primary cell culture	1	2	3	4	5	6	7	8	9	10	11	12		pecific tcome	
CLR-2:	understand the rationale of sub culturing of cells and maintaining it	ge		of	SI					ork		9				
CLR-3:	analyzing the cellular content using specific staining methods	Knowledge	W	nent	stigations	age	ъ			Μ		Finan	б			
CLR-4:	distinguish between cell viability and cell cytotoxicity		Analysis	velopm	estig	l Us	r and	۲ ×		Теаг	tion	≪ E	arning			
CLR-5:	comprehend the applications of an <mark>imal cell</mark> culture	ering		Ó	t inv	P 8	engineer sty	onment sinability		<u>∞</u>	ommunication	Project Mgt.	g Le			
		<u>e</u>	roblem	sign/d	duc	Jern	et etv	viron staina	S	ndividual	nuu	ect	Lon		2-05	<u>ب</u>
Course (Outcomes (CO): At the end of this course, learners will be able to:	Engir	Pro	Des	55	Mod	The	Env Sus	Ethics	İpu	Son	Proj	Life	PSO.	PS(PSO-3
CO-1:	develop hands on training in pr <mark>imary ce</mark> ll culture techniques	=	2	3		1	7-	-	-	-	-	-	-	-	3	-
CO-2:	gain proficiency in culturing an <mark>d maint</mark> aining cell lines	1.5	-	3	2	- 1		-		-	-	-	-	-	-	3
CO-3:	acquire skills to perform fluorescent staining procedures to visualize cellular content	. s 82-5	3	3		- 5		-		-	-	-	-	-	3	-
CO-4:	critique the toxicity of drugs in <mark>vitro</mark>	2 3		3	3	-	-	-		-	-	-	-	2	3	-
CO-5:	utilize cell culture techniques in emerging fields of animal biotechnology	-	1 -	14	3	2		-		-	-	-	-	-	-	3

Unit-1 - Media Preparation and Primary Cell Culture

Practice:

- 1. Preparation & Sterilization of media for animal cell culture
- 2. Isolation of Hepatocytes and checking its viability
- 3. Isolation and culturing fibroblasts from chick embryo

Unit-2 - Cell Culture and Maintenance

Practice:

- 1. Cell passaging
- 2. Cryopreservation of cells
- 3. Revival of Cryopreserved cells.

Unit-3 - Rapid Staining Procedures for Analysis of Cellular Content using Specific Fluorochromes

Practice:

- 1. Mitochondrial & Nuclear staining using fluorochromes
- 2. Detection of apoptosis using Annexin V
- 3. Detection of mycoplasmal contamination by Hoechst staining

12 Hour

12 Hour

Unit-4 - Cell Viability and Cell Cytotoxicity Assays

12 Hour

Practice:

- 1. Determination of Cell viability by MTT assay
- 2. Assessment of Cytotoxicity by LDH assay
- 3. Clonogenic assay

Unit-5 - Applications of Cell Culture

12 Hour

Practice:

- 1. Determination of glucose uptake by the cells using 2NBDG method
- 2. Demonstration on sorting of cells by flow cytometry
- 3. Mammalian cell transfection using lipofectamine

Learning Resources 1. Capes-Davis & Ian Freshney " Freshney's Culture of Animal Cells: A Manual of Basic Technique 2. ATCC Animal Cell culture guide and Specialized Applications", 8th Edition, ISBN: 978-1-119-51304-9, 2021 Wiley-Blackwell

Learning Asses	ssment		<i>)</i>	St. 10. 35%	771 - 611						
			met had	Continuous Learnin			4 1 9 1				
	Bloom's Level of Thinking	el of Thinking (30%)			of second cycle iments 0%)		Examination reightage)	Final Examination (0% weightage)			
	9	Theory	Practice	Theory	Practice	Theory	Practice Practice	Theory	Practice		
Level 1	Remember		15%		15%	79 - V E	15%	-	-		
Level 2	Understand		20%	100	20%	5 A , A=	20%	-	-		
Level 3	Apply	_	25%	7 3 FL 2	25%	100	25%	-	-		
Level 4	Analyze	-	25%	Commence of the Commence of th	25%	London -	25%	-	-		
Level 5	Evaluate	-	10%		10%	. ·	10%	-	-		
Level 6	Create	-	5%		5%	-	5%	-	-		
	Total	10	00 %	10	0 %	10	00 %		-		

Course Designers	1,5,94	
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
Dr. G. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai, ramchand@saksinlife.com	Prof. K Subramaniam, IITM, Chennai, subbu@iitm.ac.in	1. Dr.S.Sujatha, SRMIST
2. Dr. Karthik Periyasamy, Scientist, Biocon,	2. Prof. R. B. Narayanan, Anna University, Chennai	2. Dr.K.Venkatesan, SRMIST
karthik.periyasamy@biocon.com	arbeen09@gmail.com	

Course	21BTC306T	Course	DI ANT DIOTECHNOLOCY	Course	_	PROFESSIONAL CORE	L	T !	Р	С
Code	210103001	Name	PLANT BIOTECHNOLOGY	Category	C	PROFESSIONAL CORE	3	0	0	3

Pre-requisite Courses	N	Co- requisite Courses	Nil	ogressive Courses	Nil
Course Offer	ring Department	Biotechnology	Data Book / Codes / Standards		Nil

Course L	earning Rationale (CLR):	The purpose of learning this course is to:	11	\overline{A}	1		F	rogra	<mark>am</mark> Ou	tcome	s (PO))					ogran	
CLR-1:	understand the genome or	ganization and <mark>gene express</mark> ion in plants		1	2	3	4	5	6	7	8	9	10	11	12	_	pecifi tcom	
CLR-2:	exercise the plants as prod	uction syste <mark>ms by alte</mark> ring the plant hormones for growth and development		dge	1	of	SL		- i			ork		9				
CLR-3:	employ different methods f	or the de <mark>velopment</mark> of transgenic plants		wlec	S	velopment	stigations	age	pu			ΜW		Finan	Б			
CLR-4:	interpret the mechanisms f	or the p <mark>lant to co</mark> pe with biotic and abiotic stresses	7	S S	Analysis	lopi	estig	l Us	er an	۲ × ۷ ×	A.	Team	tion	∞ F	arning			ł
CLR-5:	apply the classical and mo	dern <mark>plant bre</mark> eding techniques for crop improvements	Jir	ring		(D)	t inv	T 90	inginee ty	ment		<u>∞</u>	Communication	Project Mgt.	g Le			ł
				inee	roblem	sign/de	duc	lern	et c	ron	g	Individual	nur	ect	Long	Ξ)-2	-3
Course C	outcomes (CO):	At the end of this course, learners will be able to:	A 19	Engi	Prof	Des	of S	Moo	The	Envi	Ethic	Indi	Col	Proj	Life	PSO-1	PSO-2	PSO-3
CO-1:	discuss the structure, organ	ni <mark>zation of</mark> plant genomes and gene regulation		3	1	3	-	7	1	3	-	-	-	-	-	-	2	-
CO-2:	demonstrate the mechanis	m <mark> and rol</mark> e of plant tissue culture for mass multiplications	/ -	3	2	3				-	-	-	-	-	-	-	3	-
CO-3:	establish the various metho	o <mark>ds of ge</mark> netic manipulation in plants	. 4 8	3	- 2	- 1		3		-		-	-	-	-	-	3	-
CO-4:	discuss the molecular aspe	cts of plant adaptability to various stresses		3		2	-	-	_	3		-	-	-	-	-	1	3
CO-5:	apply the significance of pl	a <mark>nt breed</mark> ing and genetic manipulations of plants for economic importance		3		1-1	-	3		3	-	-	-	-	-	-	3	- 1

Unit-1 - Plant Genomes: the Organization and Expression of Genes

9 Hour

Plant DNA, chromatin, chromosome structure. Nuclear genome, genome size, and organization. Chloroplast and mitochondrial - Genome structure, evolution, expression, and gene regulations. Eukaryotic gene expressions and its regulation - Transcription and translation levels: Organellar self-splicing, introns, and horizontal DNA transfer, RNA modification, post-transcriptional gene silencing (PTGS), Micro RNA - Production and interfering with the gene for silencing, DNA instability, Transposable elements in plants.

Unit-2 -Techniques for in Vitro Propagation of Plants

9 нои

Introduction to plant tissue culture. Plasticity and totipotency of plant cells. The culture environment - physical and chemical factors. Plant growth hormones - classes and their roles. Stages of plant tissue culture. Culture types. Cybrids production, haploid production. Production of secondary metabolites.

Unit-3 - Tools and Techniques for Transgenic Plant Development

9 Hour

Introduction to Agrobacterium-mediated gene transfe<mark>r and Biolo</mark>gy. Ti-plasmid-process of T-DNA transfer and integration, transformation in the plant. Direct gene transfer methods - advantages and disadvantages. Basic features of vectors, optimization, and binary vectors. Alternative markers and reporter genes. The genetic manipulation of pest resistance crop plants, and Clean gene technology.

Unit-4 - Biotic and Abiotic Stresses of Plants

9 Hour

Plant stresses - Biotic stress: Plant-pathogen interactions, prokaryotes, fungi, and viruses. Disease resistance, natural disease resistance in plants. Biotechnological approach - Overexpression of PR-proteins. Herbs as biotic stress factors. Abiotic stresses: Natural and plant responses - The nature of water deficit stress. Various approaches for tolerance - salt, cold, and heat stress - Molecular mechanisms.

Unit-5 - Genetic Improvements in Agriculture

9 Hour

Introduction to crop improvement, crop plant domestication, and beyond. Breeding technologies: Advances in breeding technologies - Modern molecular plant breeding - Transgenic plants. Emerging technologies circumvent some concerns about transgenics. Applications of breeding. The second green revolution. Metabolic engineering: Molecular farming of carbohydrates, lipids, and protein. Producing fine chemicals, Plant-derived compounds as drugs. Current demand - the plants as alternative fuels

Learning Resources
Resources

- Slater. A, Scott.N.W and Fowler,M.R, "Plant Biotechnology The genetic manipulation of plants", Oxford University Press 2008
- Agnès Ricroch, Surinder Chopra, Marcel Kuntz. Plant Biotechnology (2021). Springer Nature Switzerland AG 2021 Publisher. ISBN: 978-3-030-68344-3. Published: 31 August 2021. https://doi.org/10.1007/978-3-030-68345-0. 2nd Edition.
- 3. C Neil Stewart Jr. "Plant Biotechnology and Genetics: Principles, Techniques, and Applications (2016)"- John Wiley & Sons, Inc., New Jersey ISBN: 978-1-118-82012. 2nd Edition.
- 4. Malik Zainul Abdin, Usha Kiran, Kamaluddin, Athar Ali. Plant Biotechnology: Principles and Applications (2017). Springer Publisher, Singapore. ISBN: 978-981-10-2959-2 Published: 17 March 2017. https://doi.org/10.1007/978-981-10-2961-5.

			Continuous Learning	Assessment (CLA)		Cum	mative		
	Bloom's Level of Thinking	CLA-1 Avera	native ge of unit test 0%)	CLA	g Learnin <mark>g</mark> A-2 – 0%)	Final Examination (40% weightage)			
	/ /	Theory	Practice	Theory	Practice	Theory	Practice		
Level 1	Remember	15%		15%	4	15%	-		
Level 2	Understand	25%	20 E 10 E 10	20%		25%	-		
Level 3	Apply	30%	S. J. S. 1777	25%		30%	-		
Level 4	Analyze	30%		25%	7	30%	-		
Level 5	Evaluate		A CHARLES A C	10%		-	-		
Level 6	Create	-	Charles March 1981	5%			-		
	Total -	10	0 %	10	00 %	10	0 %		

Course Designers		
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
Dr. G. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai, ramchand@saksinlife.com	Prof. K Subramaniam, IITM, Chennai, suubu@iitm.ac.in	1. R. Pachaiappan, SRMIST
Dr. Karthik Periyasamy, Scientist, Biocon, karthik.periyasamy@biocon.com	2. Prof. R. B. Narayanan, Anna University, Chennai arbeen09@gmail.com	2. S. Rupach <mark>andra, S</mark> RMIST

Course 21DTC4041	Course	PLANT BIOTECHNOLOGY LABORATORY	Course	C	PROFESSIONAL CORE	L T	Р	(1
Code	Name	PLANT BIOTECHNOLOGY LABORATORY	Category	C	PROFESSIONAL CORE	0 0	4	2	

Pre-requisite Courses	N	Co- requisite Courses	Nil Progressive Courses	Nil
Course Offeri	ing Department	Biotechnology	Data Book / Codes / Standards	Nil

Course L	earning Rationale (CLR):	The purpose of learning this course is to:		<u> </u>			Progr	<mark>am</mark> Ou	tcome	s (PC))				_	rograr	
CLR-1:	relate the growth and devel	opment of nat <mark>ural and in vit</mark> ro growth of plants for production systems	1	2	3	4	5	6	7	8	9	10	11	12	_	pecific otcom	
CLR-2:	comprehend the methods o	f nucleic ac <mark>ids isolati</mark> on from plants	dge		of	SL					ork		8				
CLR-3:	apply various gene transfer	methods in plants	Knowledge	S	Jent	stigations	ge	ъ			۸ ×		Finan	Б			
CLR-4:	employ different steps for th	e prod <mark>uction of</mark> plant secondary metabolites	Kno	Analysis	lopi	estigal probler		er and	م × ×	l.	Teal	tion	∞ర	arning			
CLR-5:	apply the classical techniqu	es fo <mark>r crop im</mark> provement	=ngineering	Ang	sign/development	ě i	_	enginee etv	ment		<u>8</u>	ommunication	Mgt.	g Le			
			inee	Problem)ugi	. o =	Modern	erc et<		જ	ndividua	nuc	Project	Long	7	7.5	53
Course O	outcomes (CO):	At the end of this course, learners will be able to:	Eng	Prof	Des		Moc	The en	Envirol Sustair	Ethics	İndi	Sol	Proj	Life	PSO-1	PSO-2	PSO-3
CO-1:	develop in vitro plants for m	a <mark>ss multi</mark> plication	3	2	3	-	5-	7	-	-	-	-	-	-	3	-	-
CO-2:	contrast the different techne expression	iques for the isolation of nucleic acids for cloning and quantification of gene	2	3	2	Ž	- 1		-		-	-	-	-	-	3	-
CO-3:	demonstrate the different st	eps for gene transfer methods and verify the transgene in plants	3	E7	1,1-1	3	2	-	-		-	-	-	-	3	-	-
CO-4:	establish the cells for the pr detection	oduction of bioactive plant secondary metabolites and methods for isolation and	3	2	Li.	-	- }		-		-	-	-	-	2	3	-
CO-5:	design the methods for the	p <mark>roductio</mark> n of best traits and apply the plant pathology for crime investigation	3	2	-	3	-		-	1	-	-	-	-	_	3	_

Unit-1 - Techniques for in Vitro Propagation of Plants

Practice:

1. Preparation of plant tissue culture media - Murashige and Skoog's (MS) medium

2. Plant tissue culture - Direct and Indirect Organogenesis

Unit-2 - Plant Genomic DNA and RNA Isolation Techniques

Practice:

- 1. Isolation of plant genomic DNA Salk line & CTAB methods Qualitative and quantitative analysis of DNA
- 2. Extraction of total RNA from plant tissues using Trizol reagent Qualitative and quantitative analysis of RNA

Unit-3 - Techniques for Transgenic Plant Development

12 Hour

12 Hour

12 Hour

Practice:

- 1. Transform the binary vector (pCAMBIA 1301) to Agrobacterium tumefaciens
- 2. Screening of Agrobacterium colonies for confirming transformation of pCAMBIA 1301 by colony PCR and Agrobacterium Mediated gene transformation by Co-cultivation of plant leaf discs
- 3. Screening of transgenic plant tissues GUS Reporter assay

Unit-4 - Plant Secondary Metabolites - Production, Isolation and Detection

12 Hour

Practice:

- 1. Development of Cell suspension culture for the production of secondary metabolites
- 2. Extraction and detection of plant secondary metabolites extract Flavonoid quercetin from onion dried peels and alkaloid caffeine from Camellia sinensis Tea / Detection by TLC and HPLC

Unit-5 - Applications of in Vitro Propagation & Plant Pathology

12 Hour

Practice:

- 1. Cybrids production through protoplast fusion
- 2. Somatic embryogenesis through endosperm culture
- 3. Crime scene investigation

Learning Resources

- 1. Plant Biotechnology Practical Manual 2023.
- 2. C Neil Stewart Jr. "Plant Biotechnology and Genetics: Principles, Techniques, and Applications (2016)"- John Wiley & Sons, Inc., New Jersey ISBN: 978-1-118-82012. 2nd Edition
- 3. Maheshwari, S.C. (1990). Tissue Culture, Molecular Biology and Plant Biotechnology A Historical Overview. In: Sangwan, R.S., Sangwan-Norreel, B.S. (eds) The Impact of Biotechnology on Agriculture.. Current Plant Science and Biotechnology in Agriculture, vol 8. Springer, Dordrecht. https://doi.org/10.1007/978-94-009-0587-0_1
- Çelik, Ö. (2018). Introductory Chapter: New Age Molecular Techniques in Plant Science. In (Ed.), New Visions in Plant Science. IntechOpen. https://doi.org/10.5772/intechopen.79360.
- Methods in Plant Molecular Biology and Biotechnology by Bernard R. Glick. Published November 29, 2017, by CRC Press. ISBN 9780367412128

earning Asses	ssment			201 1 4 1 10							
			\$74° 5	Continuous Learning		re popular in	/				
	Bloom's Level of Thinking	expe	ge of first cycle riments 0%)	CLA-2 Average experi (30	ments		Examination veightage)	Final Examination (0% weightage)			
	1	Theory	Practice	Theory	Practice	Theory	Practice Practice	Theory	Practice		
Level 1	Remember	-	15%		15%		15%	=	-		
Level 2	Understand	-	20%	- 11	20%	-	20%	-	-		
Level 3	Apply		25%	- 1011	25%	-	25%	-	-		
Level 4	Analyze		25%	- 1/ /	25%	-/ h	2 <mark>5%</mark>	-	-		
Level 5	Evaluate		10%	- 17 111	10%	7 ,	10%	-	-		
Level 6	Create	- t	5%	- 1/2//	5%	11	5%	-	-		
	Total	10	0 %	100) %	/ 10	00 %		-		

Course Designers	/ II EAKN · LEAD rein b	
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
Dr. G. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai, ramchand@saksinlife.com	Prof. K Subramaniam, IITM, Chennai, suubu@iitm.ac.in	1. R. Pachaiappan, SRMIST
Dr. Karthik Periyasamy, Scientist, Biocon, karthik.periyasamy@biocon.com	2. Prof. R. B. Narayanan, Anna University, Chennai arbeen09@gmail.com	2. S. Rupachandra, SRMIST

Course	21BTC402J	Course	BIO SEPARATION TECHNOLOGY	Course	C	PROFESSIONAL CORE	L T P	С
Code	210104023	Name	BIO SEPARATION TECHNOLOGY	Category	U	PROFESSIONAL CORE	3 0 2	4

Pre-requisite Courses	Co- requisite Courses	Nil Progressive Courses	Nil
Course Offering Department	Biotechnology	Data Book / Codes / Standards	Nil

C	samina Dationala (CLD):	The mount of leave	in with in any way in the		_					O-	.4	(DC					Pi	rograi	<u> </u>
Course L	earning Rationale (CLR):	The purpose of learn	ng this course is to:	X - X - X - X - X - X - X - X - X	1	-			rogra	am Ol	utcome	es (PC	")	1				pecifi	
CLR-1:	know the importance of bio	separation and its recove	ry economically		1	2	3	4	5	6	7	8	9	10	11	12		tcom	
CLR-2:	learn the separation of prod	luct from so <mark>lid –liquid</mark> pha	se		dge		of	SL			l N		ork		9				
CLR-3:	know the techniques of isol	ation of b <mark>io-produc</mark> ts	(65	an William	owlec	S	nent	stigations roblems	Usage	ъ			M		Finan	ning			
CLR-4:	learn the methods of purific	ation o <mark>f produc</mark> ts	N 434	3 3 4 5	조	Analysis	elopment	estig probl	-	r and	∞ >		Team	ţio	& ∃	earni			
CLR-5:	learn the methods of polish	ing a <mark>nd formu</mark> lation of pro	ducts for packaging	22W25	ering		deve	t inv	Т00	engineer stv	ment		<u>ھ</u>	ommunication	Project Mgt.	Ľ			
	<u>, </u>			77.5	(I)	roblem	ign/	duc	ern	enç et >		S	ndividual	nul	ect	Long	7	-5	က္
Course O	outcomes (CO):	At the end of this cou	ırse, learners will be able to:	rational garage	Engin	Prok	Desi	Con	Мод	The	Envi Sust	Ethic	lndj	Son	Proj	Life	PSO-1	PS0-2	PSO-3
CO-1:	categories the products into	v <mark>arious s</mark> ectors		A district of	1	2	1	-	7	7-	-	-	-	-	-	-	1	2	1
CO-2:	identify the unit operation for	o <mark>r se</mark> paration	1000	O William I	2	3	1	-			-	1	-	-	-	-	2	2	1
CO-3:	adapt the best methods of i	solation of products	THE STATE OF THE S	Water Mary	2	2	2	<i>3</i> -	-		-		-	-	-	-	2	2	1
CO-4:	identify the sophisticated ed	<mark>quipmen</mark> t for purification		77 4 77	- 2	3	2	-	-	-	-	-	-	-	-	-	2	2	2
CO-5:	know the polishing and form	n <mark>ulation of the products</mark>		1 1	2	2	2	-	- 7	-	-	-	-	-	-	-	2	2	2

Unit-1 - Bioproducts Classification and Disruption Techniques

Classification of Bioproducts, Engineering Analysis, Analytical methods, Cell disruption Methods- Physical, Chemical, Mechanical and Biological methods.

Practice:

Cell disruption Techniques

1. Cell disruption by Sonication, 2. Cell disruption by High Pressure Homogenisation, 3. Chemical and Enzymatic method of cell disruption

Unit-2 - Separation of Insolubles

15 Hour

15 Hour

Electrical Double layers, Schulze–Hardy Rule, Flocculation Rate, Polymeric Flocculants, Sedimentation-Principles, Methods and Coefficients, Filtration Principles and Theory, Conventional Filtration-Filtration Equipments and Media, Scaleup and Design of Filtration Systems, Cross flow filtration-Microfiltration, Centrifuges, Scaleup of Centrifugations.

Practice:

Recovery Methods

1. Cell separation by Flocculation, 2. Cell separation by Batch filtration, 3. Cell separation by Microfiltration, 4. Cell separation by Centrifugation

Unit-3 - Concentration of Solubles

15 Hour

Extraction-Batch, Staged, Differential Extraction, Aqueous two phase Extraction, Supercritical Extraction, Batch Adsorption, Adsorption in CSTR and Fixed Bed, Precipitation-Different methods of precipitation, Ultrafiltration, Dialysis and Electrodialysis.

Practice:

Protein Concentration Methods

1. Protein concentration by Precipitation methods, 2. Protein concentration by Ultrafiltration, 3. Protein Concentration by Aqueous two-phase extraction

Unit-4 - Protein Purification 15 Hour

Chromatography Column Dynamics, Plate Models, Chromatography Column Mass Balance with Negligible Dispersion, Dispersion Effects in Chromatography, Gradients and Modifiers, Adsorbent Types, Particle Size and Pressure Drop in Fixed Beds, Equipment, Scaleup.

Practice:

Purification of Protein

- 1. Protein purification by gel column chromatography
- 2. Protein purification by ion exchange chromatography

Unit-5 - Polishina

15 Hour

Crystallization Principles, Batch Crystallizers, Process Crystallization of Proteins, Crystallizer Scaleup and Design, Drying Principles, Dryer Description and Operation, Scaleup and Design of Drying Systems, Case studies.

Practice:

Polishing of Biomaterial

- 1. Crystallization Techniques
- 2. Freeze drying of biomaterials

Learning
Resources

- Harrison. R.G., Todd. P., Rudge S.R, Petrides. D.P, "Bioseparation Science and Engineering" Oxford University press, 2003.
- 2. Belter. P.A., Cussler, E., "Bioseparations", Wiley, 1985.

- Nooralabettu Krishna Prasad, "Downstream Process Technology: A New Horizon In Biotechnology", PHI Learning Private Limited 2013
- 4. Mihir K Purkait; Randeep Sing, "Membrane Technology in separation science, CRC Press Taylor & Francis Group, 2018

•	ent	2.57 2024	Continuous Learnin	g Assessment (CLA)		Cum	mativa
	Blo <mark>om's</mark> Level of <mark>Thinking</mark>	Form CLA-1 Averaç (45	ge of unit test		-Practice 5%)	Final Ex	mative amination eightage)
		Theory	Practice	Theory	Practice	Theory	Practice
Level 1	Remember	15%	NY - 1/177.	-	15%	15%	-
Level 2	Understand	25%	- 1	-	20%	25%	-
Level 3	Apply	30%	- 11/1/1	-	25%	30%	-
Level 4	Analyze	30%	- 1/2%	<u>-</u>	25%	30%	-
Level 5	Evaluate				10%	-	-
Level 6	Create	<-		7-	5%	-	-
	Total	100)%	4 13 10	00 %	10	0 %

Course Designers	V LIMIT	4.00
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
Dr. G. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai, ramchand@saksinlife.com	Dr.S.Senthil Kumar, IITG, senthilkumar@iitg.ac.in	1. Dr.M.Venkatesh Prabhu, SRMIST
Dr. Karthik Periyasamy, Scientist, Biocon, karthik.periyasamy@biocon.com	2. Dr.N.Selvaraj, IITG, selva@iitg.ac.in	2. Dr.P.Radha, SRMIST

ACADEMIC CURRICULA

Non-Credit Course

Regulations 2021



SRM INSTITUTE OF SCIENCE AND TECHNOLOGY

(Deemed to be University u/s 3 of UGC Act, 1956)

Kattankulathur, Chengalpattu District 603203, Tamil Nadu, India

Course Code	21BTM191T	Course Name	BIOETHICS AND IPR	-	ourse tego:	-	М				NON	CRED	IT			-	L T 1 0	P 0	0 0
Pre-requis	s	Nil	Co- requisite Nil	****		gres ours							Nil	1					-
Course C	Offering Departme	ent	Biotechnology Data Book / Codes /	Standards								Nil							
Course Lea	arning Rationale	(CLR): The	purpose of learning this course is to:	NO	7	-			Progr	ram Oı	utcome	s (PO)				Р	rogra	m
CLR-1:		` '	s in Biotec <mark>hnology Res</mark> earch		1	2	3	4	5	6	7	8	9	10	11	12		pecif	
CLR-2:			with biotechnology Research		<u>o</u>		-							10			01	licon	25
CLR-3:			age that could be caused to the environment		ledc		ent c	tions	e e		. 1		Wo		Finance				l
CLR-4:			Values to be inculcated in ethical decision making		ريا ا	ysis	bmd	stiga oble	Jsac	and	∞ _	N.	eam	5	Ë	Learning			l
CLR-5:			inment of risk group organisms	560-1 3	ng	√nai	selc	nves x pr	00	eer	ent		∞ ∞	catic	gt. &	Lea			l
CLK-3.	Know the require	ITIETIIS TOT COTILA	minent of risk group organisms		eeri	me	n/de	uct i	E	igi >	onm inab	(0	dual	nun.	T W	ong	_	2	က
Course Ou	tcomes (CO):	At t	he end of this course, learners will be able to:		Engineering Knowledge	Problem Analysis	Design/development of solutions	Conduct investigations of complex problems	Modern Tool Usage	The engineer and society	Environment 8 Sustainability	Ethics	ndividual & Team Work	Communication	Project Mgt.	Life Long l	PS0-1	PSO-2	PSO-3
CO-1:	define Principles	of Bioethics and	aspects related to IP protection	E	-	3	3	_	-	- 0	-	7	-	-	-	-	-	-	<u> </u>
CO-2:	•		safety precautions in biotechnology research	4.58. L	- '	3	2	77	-	1			-	-	-	-	-	-	-
CO-3:	explain concepts	pertaining to ex	rercising personal and environmental safety		- 10	2	75	124	3			1-5	-	-	-	_	-	-	-
CO-4:			I decisions in healthcare research		1	2	3		_	_	-		-	-	-	_	-	-	-
CO-5:	discriminate diffe	rent biosafe <mark>ty le</mark>	vels and different forms of IP	Op 19		2			_		-		-	_	-	-	_	-	-
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	sic Principles of											1						3	Hou
			<mark>f anim</mark> als in research and Ethical issues in Clinical Trials, Ethi	ical issues in	Stem	Cell	Resea	arch, E	thical	Issues	in In v <mark>i</mark> i	tro Fer	<mark>ti</mark> lizatio	on					
	bal Health Ethic				,										, ,	- · ·			Hou
	ems and Institutions Is in Bioethics	ons, Synaptoger	<mark>esis and</mark> development of sensory-motor system, Ethical issu	ies in Organ t	ransį	oianta	ation, E	siobani	king, E	tnicai	issues	ın Reg	enera	tive ivie	eaicine,	Religi	ous a	ına Cı	itura
	safety Regulation	ons							7									3	Ηοι
			of various regulatory bodies, Biosafety Rules for GMOs, Biod	liversity and E	nviro	onme	nt cons	servatio	on. CE	BD and	Cartao	ena Pi	rotoco	I					
Unit-4 - For		,	/ ITEARN	FAD			1 1 1	L	- Andrews									3	Ηοι
		graphical indica	tions, N <mark>ovelty and</mark> Utility, Patentable subjects and protection i	in biotechnolo	ogy, E	Biodiv	versity												
Unit-5 - Pat								لسيا		.*									Ηοι
Basic princi	ples and general r	equirements of p	patent law Pa <mark>tents and met</mark> hods of application of patents-Lega	al implications	, Obi	ective	es of th	e pater	nt syst	em, TF	RIPs-GA	ATT-Ini	ternati	ional co	nventi	ons. Pa	atent C	Coope	ratic

2. The Indian Patent Act and Rules, 2015, Gol, India.

1. Singer and Viens (Eds.) Bioethics – Cambridge University Press, Cambridge, 2008

Learning Resources

arning Assessm			Continuous Learning	g Assessment (CLA)		C	
	Bloom's Level of Thinking	CLA-1 Avera	native ge of unit test %)	CI	g Learning LA-2 <mark>0%)</mark>	Final Ex	mative amination eightage)
		Theory	Practice	Theory	Practice	Theory	Practice
Level 1	Remember	15%	ALTERNA	15%		15%	-
Level 2	Understand	25%		20%		25%	-
Level 3	Apply	30%	3	25%		30%	-
Level 4	Analyze	30%	-	25%		30%	-
Level 5	Evaluate		-	10%	7	-	-
Level 6	Create		*-A	5%	7	-	-
	Total	100) %	10	00 %	10	00%

Course Designers		
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
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ACADEMIC CURRICULA

UNDERGRADUATE/ INTEGRATED POST GRADUATE DEGREE PROGRAMMES

(With exit option of Diploma)

(Choice Based Flexible Credit System)

Regulations 2021

Volume - 8B
(Syllabi for Biotechnology w/s Regenarative Medicine
Programme Courses)



SRM INSTITUTE OF SCIENCE AND TECHNOLOGY

(Deemed to be University u/s 3 of UGC Act, 1956)

Kattankulathur, Chengalpattu District 603203, Tamil Nadu, India

ACADEMIC CURRICULA

Professional Elective Courses

Regulations 2021



SRM INSTITUTE OF SCIENCE AND TECHNOLOGY

(Deemed to be University u/s 3 of UGC Act, 1956)

Kattankulathur, Chengalpattu District 603203, Tamil Nadu, India

Course	21BTE206T	Course	MOLECULAR CELL BIOLOGY	Course	Е	PROFESSIONAL ELECTIVE	L	Т	Р	С	
Code	210102001	Name	WOLECULAR CELL BIOLOGY	Category		PROFESSIONAL ELECTIVE	3	0	0	3	

Pre-requisite Courses	N	Co- requisite Courses	Nil Progress Course	Nil
Course Offeri	ing Department	Biotechnology	Data Book / Codes / Standards	 Nil

Course Le	earning Rationale (CLR): The purpose of learning this course is to:	Program Outcomes (PO)										rogran				
CLR-1:	provide basic knowledge of stem cell specific gene expression in lineage-based tissues from the perspective of engineers	1	1 2 3 4 5 6 7 8 9 10 11 12						12		pecific itcome					
CLR-2:	identify the role of epigenetic regulation in stem cell proliferation and differentiation	edge		. <	of	ų.	ciety			k						
CLR-3: deliver the knowledge on signaling molecules and molecular mechanisms that regulate the stem cell proliferation and differentiation				evelopment of	stigations lems	Usage	S		N.	m Work		Finance	lug			
CLR-4:					nvestigat problems	I Us	and and	t &		Team	tion	∞	arning			
CLR-5:	LR-5: utilize the strategies for novel gene editing techniques for tissue engineering					T00	engineer	rironment tainability	1	<u>ल</u>	ommunication	Mgt.	g Le			
		gineering	oblem	ign/d/ tions	duct	lern		iron	છ	ndividual	חתר	roject	Long	-1)-2	-3
Course O	utcomes (CO): At the end of this course, learners will be able to:	Eng	Prof	Des	5 6	Mod	The	Env	Ethics	İpul	Coll	Proj	Life	PSO-1	PSO-2	PSO-
CO-1:	identify gene regulation in stem cells	2	2	2	2	-	1	-		-	-	-	-	3	-	-
CO-2:	analyze gene expression in stem cells and artificial generation of pluripotency				3	-		} -		-	-	-	-	3	-	
CO-3:	identify the applications of g <mark>rowth fac</mark> tor signaling and their receptor molecules			2	2	-	-	-		-	-	-	-	3	-	-
CO-4:	0-4: analyze the regulation of molecules involved in self-renewal of stem cells				3	-		-		-	-	-	-	2	-	-
CO-5:	discuss stem cell death and survival mechanisms	3	2	1,14	2	_	-	-		-	-	-	-	2	-	-

Unit-1 - Molecular Biology of life

9 Hour

The Molecules of Life- Genomes, Cell Architecture, and Cell Function- Chemical Foundations-Covalent Bonds and Noncovalent Interactions -Chemical Building Blocks of Cells -Nucleic acids, genetic material-Primary and secondary structure of DNA - Watson & Crick model -Hogsteen base pairing, Triple helix, Quadruple helix -DNA super-coiling- Linking number- satellite DNA replication - Meselson & Stahl experiment bi— directional DNA replication- Proteomics of DNA replication - Overview of differences in prokaryotic and eukaryotic DNA replication -Role of telomerase in aging and cancer- Mutagens, DNA mutations and their mechanism- Telomere replication in eukaryotes DNA Repair- DNA mismatch, Base-excision- Nucleotide-excision and direct repair DNA recombination- Homologous, site-specific and DNA transposition- Gene regulation and operon.

Unit-2 - Gene Regulation and Transcription

9 Hou

Overview of Central dogma- Characteristics promoter and enhancer sequences - Transcriptional bubble - prokaryotic and eukaryotic RNA polymerase -RNA synthesis- Fidelity of RNA synthesis. Inhibitors of transcription -Differences in prokaryotic and eukaryotic transcription -Regulatory elements- Mechanism of transcription regulation - Transcription of Protein-Coding Genes and Formation of Functional mRNA- Splicing - nuclear export of mRNA - mRNA stability-Role of gene expression in microRNA -LncRNA, snoRNA, piRNA- srRNA, siRNA and shRNA -Genetic code: Elucidation of genetic code- Codon degeneracy, Wobble hypothesis and its importance- Prokaryotic and eukaryotic ribosomes- Chromosomal Organization of Genes and Noncoding DNA- Molecular Mechanisms of Transcription Repression and Activation

Unit-3 - Protein Functions 9 Hour

Membrane Proteins: Structure and Basic Functions- cytoskeletal proteins Extra cellular matrix- cell-cell junctions, various types of transport across cell membrane - Protein sorting and trafficking, cargo proteins-Growth factor signaling, cell-cell communication - Mechanism of action of different class of hormones- Cell cycle – Molecules controlling cell cycle- Carcinogens and Caretaker Genes in Cancer- Recombination to Repair Double-Strand Breaks in DNA-Cell culture and immortalization of cells and its applications- Restriction Enzymes, Cloning and Libraries- DNA Cloning and Characterization - Molecular Analysis Using Cloned Sequences

Unit-4 - Molecular Biology Techniques

9 Hour

Molecular Genetic Techniques- Inactivating the Function of Specific Genes in Eukaryotes - Cloning a Human Gene - Locating and Identifying Human Disease Genes - Inactivating the Function of Specific Genes in Eukaryotes- Molecular basis of Organ culture- Molecular Basis of Pluripotency- Induced pluripotency- Lineage tracing experiments in stem cells- Characterization and maintenance of murine and human embryonic stem cells- Differentiation of embryonic Stem Cells- Embryonic stem cell cloning- Therapeutic cloning of stem cells- Genomic Reprogramming.

Unit-5 - Molecular Diagnosis and Cell Therapy

9 Hour

Proteomics, Metabolomics, Microbiomics and Systems Biology -Newborn screening: Neonatal PKU- Cystic fibrosis and sweat tests- Prenatal diagnosis of diseases, amniotic fluid- Fetal blood examination-Karyotyping, Chromosomal abnormalities by cytogenetics- Restriction fragment length polymorphism (RFLP)- Nuclear injection- stem cell transplantations for sickle-cell anemia, hemophilia- Stem cell transplantation for cancer (leukemia and myeloma- Muscular dystrophy and stem cell therapy- Stem cell therapy, Neurodegenerative disease- and human embryonic stem cells, Stem cell transplantation- Dementia-Neurodegenerative disease- CRIPSR/Cas9 system-gene editing- Applications of CRISPR/Click chemistry techniques.

Learning
Learning Resources
Resources

- 1. Harvey Lodish, Arnolg Berk "Molecular Cell Biology," 9 th edition Mcmillan 2021; ISBN:9781319208523
- 2. Gerald Karp, "Karp's Cell and Molecular Biology," 9th edition Wiley, 2019, ISBN: 978-1-119-59816-9
- Lewin's Genes. Joycelyn E. Krebs, Elliott S. Goldstein, Stephen T. Kilpatrick. 12th Edition, Jones and Bartiett Publishers Inc. ISBN: 978-1284104493

earning Assessm	nent		- 10	Continuous Learnin	g Assessment (CLA)			
	Bloom's Level of <mark>Thinking</mark>	Formative Life Long Learn CLA-1 Average of unit test CLA-2 (50%) (10%)		A-2	Final Ex	mative amination eightage)		
		7	Theory	Practice	Theory	Practice	Theory	Practice
Level 1	Remember	_	15%	7.00	15%		15%	-
Level 2	Understand		25%		20%		25%	-
Level 3	Apply		30%	A STATE OF THE STATE OF	25%		30%	-
Level 4	Analyze		30%		25%		30%	-
Level 5	Evaluate			The second second	10%			-
Level 6	Create			- 4,077	5%	-4		-
	Total		100 %	1.7	10	0 %	10	0 %

Course Designers	300	
Experts from Industry	Experts from Higher Technical Institutions	/ Internal Experts
1. Dr.Ramesh Babu Boga, BogaR Laboratories, Hyderabad,	1. Prof. N. Gopalan, Central University Tamil Nadu, email:	1. Dr.Kanagaraj Palaniyandi, SRMIST
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2. Dr. Archana Khosa Kakkar, IDRS Labs Private Limited,	2. Dr. Ajaikumar B. Kannumakkara , IITG,	2. Dr. N. Selvamurugan, SRMIST
Bangalore, archgana.kk@idrslabs.com	kunnumakkara@iitg.ernet.in	

Course	21BTE311T	Course	CELL COMMUNICATION AND SIGNALING	Course	Е	PROFESSIONAL ELECTIVE	L	Т	Р	С
Code	ZIBIEJIII	Name	CELL COMMUNICATION AND SIGNALING	Category	E	PROFESSIONAL ELECTIVE	3	0	0	3

Pre-requisite Courses	N	Co- requisite Courses	Nil Progressi Courses	Nil
Course Offeri	ng Department	Biotechnology	Data Book / Codes / Standards	Nil

Course L	earning Rationale (CLR): The purpose of learning this course is to:	111	4			Progr	am Oı	itcome	s (PO)					rograi	
CLR-1:	provide basic concepts of gene expression patterns from the perspective of engineers	1	2	3	4	5	6	7	8	9	10	11	12	_	pecifi ıtcom	
CLR-2:	identify the role of epigenetic regulation in adult stem cells			14				ility								
CLR-3: identify the external and internal signaling molecules that regulate the stem cell proliferation and differentiation				int of	stigations of	a	society	ustainability		Work		Finance	_			
CLR-4: analyze the self-renewal and cell death mechanisms in stem cells				sign/development	tigal ems	Usage	and	S		eam	_		aming			i
CLR-5:	clr-5: analyze the molecular mechanism of stemness- signaling pathways and transcription factors and diseases				prob	T00	engineer a	ment &		_ ∞ŏ	ommunication	Mgt. &	မ			
Course C	Outcomes (CO): At the end of this course, learners will be able to:	Engine	Problem,	Design/c	Conduct	Modern	The en	Environment	Ethics	Individual	Comm	Project	Life Long I	PSO-1	PS0-2	PSO-3
CO-1:	apply the basic understandin <mark>g of gen</mark> e regulation in stem cells	2	2	2	2	-	4	-	-1	-	-	-	-	3	-	-
CO-2:	CO-2: manipulate the gene expres <mark>sion in s</mark> tem cells and artificial generation of pluripotency				3	-	-	7 -		-	-	-	-	3		-
CO-3:	CO-3: identify the applications of growth factor signaling and their receptor molecules				2	-	-	-		-	-	-	-	3	-	-
CO-4: apply the regulation of molecules involved in self-renewal of stem cells				3	3	-	-	-	-:	-	-	-	-	2	-	-
CO-5:	discuss the stem cell death a <mark>nd survi</mark> val mechanisms	3	2	100	2	-	*	-	-10	-	-	-	-	2	-	-

Unit-1 - Cell-Cell Communication 9 Hour

Heterotypic and homotypic cell-cell contacts and how these contacts mediate intracellular communication-Cell-matrix communication including mechanisms of cell motility -Pluripotency associated transcription factors-Tissue specific multi potency-Stem cells with no tissue specificity-Transcriptional network controlling pluripotency in ES cells-Alternative splicing in embryonic stem cells- Niche required for inducing stem cell control-Homeostasis and Feed-back regulation in niche- Cytokines and growth factors maintenance of stemness-Modeling for stem cell asymmetry- Pluripotency genes, expression and regulation

Unit-2 - Receptor Mediated Cell Signaling

9 Hour

Cell surface receptor mediated signal transduction-Growth factor and receptors-tyrosine kinases Mediated signaling (RasRaf-MAP-MEK)-Wnt -signaling-Notch signaling pathways-Hedgehog singling-Hippo signaling-JAK-STAT- nuclear signaling-NF-κB signaling pathways-TGFβ -activation/nodal BMP-signaling-FGF signaling pathways-Hemotopoiesis and signaling molecules-Progenitor cell differentiation factors-Colony stimulating factor and its receptor signaling pathways. Glycoprotein and proteoglycan structure and biology including molecular gradients and their involvement in embryology and disease.

Unit-3 - Aging and Senescence Signaling

9 Hour

Stem cell aging and apoptosis-Regulation and significance apoptosis in stem cells-Stem cell necrosis-Intrinsic – extrinsic pathways of apoptosis-Death ligands, cytokines and tumor necrosis factor-Role of apoptosis in hematopoiesis-Apoptosis resistance in stem cells-Anti-apoptotic molecules expression in stem cells-Caspase mediated apoptosis-Apoptosis transcription factors and regulators-Heat shock proteins- The role of ubiquitination in signal transduction and protein degradation-The role of reactive oxygen species as secondary messengers-How cells respond to stress signals in homeostasis and disease (e.g. autophagy and ER stress).

Unit-4 - Neural Cell Differentiation and Signaling

9 Hour

Neural stem cells-Neural progenitors-The heterogeneity of adult neural stem cells-Emerging complexity of neural niche-Neural stem cell signaling-Neural stem cell homeostasis-Galecitin-1 in neural stem cells-Neurotransmitter-induced stem cell differentiation-cholinergic-dopaminergic signals-Nerve cell growth factor-Induced regeneration of neuronal cells-Neurosphere culture-Astrocyte, oligodendrocyte differentiation-Glial cell differentiation-Pathophysiology of neuronal stem cell signaling-Multiple sclerosis, Parkinson's and Alzhimer's -How microRNA/IncRNA regulate cell signaling.

Unit-5 - Methods in Cell Signaling 9 Hour

How different techniques are used to study cell signaling-Regeneration, Stem Cells, and the Evolution of Tumor Suppression- Smads - Polycomb genes- Cellular signaling of Akt/PKB - β-catenin- Induced pluripotency (iPSc)- Epithelial-mesenchymal transition (EMT)- EMT markers- Growth factor induced differentiation of stem cells- Pancreatic stem cells- Beta cell differentiation factors and transplantation- Stem cell therapy for obesity- Leukemia, lymphoma and Myeloma- Bone marrow transplantation- Cytokine and chemokine therapies- Cancer stem cell - cell survival and tumor maintenance- Mechanism of cancer stem cell resistance-Targeting cancer stem cells- Selective killing of cancer stem cells- Nano carrier mediated drug delivery

Learning	 Hancock John T "Cell signaling," 	' <mark>Oxford Univers</mark> ity Press- 2016; ISBN	: 2 Handbook of Cell signaling, Edward A Dennis and Ralph A Bradshaw. Elsevier, 2003 ISBN:
Resources	9780199658480		9780121245467

Learning Assessm	nent								
	Bloom's Level of Think <mark>ing</mark>	CLA-1 Avera	Continuous Learning mative age of unit test 0%)		Learning A-2 0%)	Summative Final Examination (40% weightage)			
		Theory	Practice	Theory	Practice	Theory	Practice		
Level 1	Remember	15%		15%	- C- C	15%	-		
Level 2	Understand	25%		20%		25%	-		
Level 3	Apply	30%	Charles To the Control	25%		30%	-		
Level 4	Analyze	30%		25%	. 1 . 7	30%	=		
Level 5	Evaluate	A - 2.	William Commence	10%		-	-		
Level 6	Create	F 4	All the second of the second o	5%	- 5	-	-		
	Total	10	0 %	100	0 %	100) %		

Course Designers		
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
1. Dr.Ramesh Babu Boga, BogaR Laboratories, Hyderabad,	1. Dr. Ravisankar B, University of Madras, email:	1. Dr.Kanagaraj Palaniyandi, SRMIST
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2. Mr.J.B. Vijayakumar BioArtis Life Sciences Pvt. Ltd. email:	2. Dr. Ajaikumar B. Kannumakkara , IITG,	2. Dr. R. Satish, SRMIST
contact@bioartis.in	kunnumakkara@iitg.ernet.in	La Value and the second

Course	21BTE312T	Course	STEM CELL TECHNIOLOGY	Course	Е	PROFESSIONAL FLECTIVE	L	Т	Р	С
Code	ZIBIEJIZI	Name	STEW CELL TECHNOLOGY	Category	E	PROFESSIONAL ELECTIVE	3	0	0	3

Pre-requisite Nil	Co- requisite Courses	Nil Progressive Courses	Nil	
Course Offering Department	Biotechnology	Data Book / Codes / Standards	Nil	

Course L	earning Rationale (CLR): The purpose of learning this course is to:	111	Program Outcomes (PO)												rograi	
CLR-1:	provide basic knowledge on embryogenesis from the perspective of engineers	1	2	3	4	5	6	7	8	9	10	11	12	_	pecifi ıtcom	
CLR-2:	create an interest to know about the different types of embryonic stem cells, its isolation, and cloning	O)		14	of		ciety			~						
CLR-3:	illustrate awareness about adult and cancer stem cells, IPSCs and importance of stem cell niches	Knowledge		nt of	stigations lems	Θ	socie			Work		ance				i
CLR-4:					tigat	Usage	and		h.	eam	_	Fina	ning			i
CLR-5:	generate interest on applications <mark>and uses</mark> of stem cells and to develop strategies for tissue engineering and create awareness on ethics and regulations of stem cell research	ering K	n Analysis	development	t inves x probl	Tool	engineer a	ment 8 ability	. 1	~ ∞	ommunication	Mgt. &	ıg Lear			Í
		nginee	oblem	ign/	op (e)	dern		ion		ndividual	nwu	roject	Long	0-1)-2	PSO-3
Course C	Outcomes (CO): At the end of this course, learners will be able to:	Eng	Pro	Des	Con	ωМ	The	Env Sus	Ethics	Indi	Cor	Pro	Life	PS(PSO	PS(
CO-1:	recall knowledge about embr <mark>yogenes</mark> is, stem cells and its characteristics		120	18	2	-	1	-	3	-	-	-	-	-	-	2
CO-2:	interpret knowledge on different types of stem cells isolation of ESCs and cloning	V -	11.	19	2	-	-		3	-	-	-	-	-	-	2
CO-3:	interpret about adult and cancer stem cells, IPSCs and stem cell niches	Mr.E.	17.74	7.	.2	-	-	-	3	-	-	-	-	-	-	3
CO-4:	analyze the role of signaling pathways, epigenetics control of stem cells	. 44		177	- 3	-	-	-	3	-	-	-	-	-	-	2
CO-5:	evaluate the application of stem cells for different diseases and reconstruct knowledge on tiss engineering for regenerative medicine	ie -			3		\subseteq	-	3	-	-	-	-	-	3	-

Unit-1 - Stem Cells-Characteristics and Types

9 Hour

Overview of Stem cells - "Stemness": Definitions, Criteria - Embryonic and adult stem cells - Types and classification of stem cells based on potency - Types of stem cells - Embryonic stem cells (ESCs), Adult stem cells (ASCs) - Differences between ESCs and ASCs- Similarities between ESCs and ASCs- Identification and characterization of ESCs and ASCs at cellular level and molecular level

Unit-2 - Embryonic Stem Cells- Isolation and Cloning

9 Hour

ESCs –IVF, Primate and Mouse ES cells, Markers - Nuclear transfer technology in ES cells - Human ESCs -Isolation and culturing of hESC's - Differentiation of stem cells - Enzymatic and Mechanical isolation – Immunosurgical Isolation - Stem Cells derived from early mouse embryos-ES, EC, EG, TS cells - Primed Cells - Naïve Cells - Epiblast Stem Cells - ESC markers - Therapeutic cloning using ESCs - Reproductive cloning using ESCs

Unit-3 - Adult Stem Cells - Sources, Types and Niches

9 Hour

Adult stem cells (ASC)-advantages and disadvantages - Sources of ASCs and its properties and its role as specialised cells in differentiation - Transdifferentiation - Fusion experiments - Experiments on transdifferentiation - Induced pluripotent stem cells (iPSCs)-Methodology, Applications - Cancer stem cells- Isolation Characterization, Properties, origin, theories - CSCs and Metastasis - Stem Cell Niche – Drosophila testis and ovary nich - Human intestinal epithelia niche

Unit-4 - Signaling Pathways and Epigenetics in Stem Cells

9 Hour

ESC pluripotency and signaling- JAK-STAT pathway -Activin/Nodal/TGFβ Signaling Pathway - FGF Signaling Pathway - Wnt signaling and Insulin-like growth factors - HSC signaling pathways- Notch, Wnt, TGF,, SMAD signalling-Epigenetic control of stem cells- Effect of TSA on stem cell differentiation - Effects of histone demethylases - Epigenetic control in pluripotent stem cells, somatic cells, germ cells -Epigenetics in iPSCs

Unit-5 - Application of Stem Cells and Ethics in Stem Cell Research

9 Hour

Stem Cells in Tissue Engineering - Therapeutic Applications - Parkinson's disease - Bone defects - Stem Cells for Spinal Cord Injury- Common strategies toward regeneration of the damaged spinal cord- Stem Cell treatment for diabetes- Cardiac tissue engineering using stem cells-Stem cell treatment for burns Transplantable matrices - Ethics of Stem Cell Research- The Ethics of Using Human Embryonic Stem Cells in Research - Regulations governing Stem Cell research-ICMR, Drugs and Cosmetic Act

Learning Resources
Resources

- 1. Robert Lanza, Edited by: Robert Lanza and Anthony Atala, "Essentials of Stem Cell Biology"3rd Edition, Academic Press, Copyright © 2014 Elsevier Inc. 4.
- Huang G, Ye S, Zhou X, Liu D, Ying QL. Molecular basis of embryonic stem cell self-renewal: from signaling pathways to pluripotency network. Cell Mol Life Sci. 2015, May; 72(9):1741-57.
- The Science of Stem Cells by Jonathan M W Slack, John Wiley & Sons, 16-Jan-2018 - Science - 272 pages.
- 4. Stem Cells Biology and Application by Mary Clarke, Jonathan Frampton · 2020 CRC Press.

earning Assessm	nent	- A.V.	Continuous Learnin	g Assessment (CLA)					
	Bloom's Level of Thin <mark>king</mark>	CLA-1 Avera	native ge of unit test 0%)	Life Long CLA (10	1-2	Summative Final Examinatio (40% weightage			
		Theory	Practice	Theory	Practice	Theory	Practice		
Level 1	Remember	15%	2. 74. 9221	15%		15%	-		
Level 2	Understand	25%	Carlot and the same	20%		25%	-		
Level 3	Apply	30%	A CONTRACTOR OF THE PROPERTY O	25%		30%	-		
Level 4	Analyze	30%	188 1 1 T - 1897 917	25%		30%	-		
Level 5	Evaluate	S 100 177 15	NAME OF THE PARTY	10%	- C	-	-		
Level 6	Create	42.7 77.31	13.1	5%		0 -	-		
	Total	10	0%	100	%	100	0 %		

Course Designers		
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
1. Dr. B.R.Desikachari, Medical Director, Westminster Health Care,	1. Prof .Halagowder D, Univ. of Madras, hdrajum@yahoo.com	1. Dr. Devi. <mark>A, SRMI</mark> ST
Chennai, brdesikachari@hotmail.com		
2. Dr. A. Premkumar, Ph.D., GVK Biosciences, Hyderabad	2. Dr. Sudha Warrier, Associate Professor, Manipal University,	2. Dr. N.S <mark>elvamuru</mark> gan, SRMIST
aprem70@yahoo.com	sudha.warrier@mannipal.edu	

Course	21BTF313T	Course	BIOMATERIALS IN TISSUE ENGINEERING	Course	Е	PROFESSIONAL ELECTIVE	L	Т	Р	С
Code	ZIBIEJIJI	Name	BIOWATERIALS IN 1135UE ENGINEERING	Category	E	PROFESSIONAL ELECTIVE	3	0	0	3

Pre-requisite Courses	N	Co- requisite Courses	Nil Progressi Courses	Nil
Course Offeri	ng Department	Biotechnology	Data Book / Codes / Standards	Nil

Course Lo	earning Rationale (CLR):	The purpose of learning this course is to:	Program Outcomes (PO)											rogra			
CLR-1:	R-1: demonstrate the basic knowledge on biomaterials from the perspective of engineers				3	4	5	6	7	8	9	10	11	12	_	pecifi itcom	
CLR-2:	CLR-2: analyze biological tissue engineering problems with biomaterials				of	SL		L "			Work		8				
CLR-3:	demonstrate basic concep	ts regardin <mark>g design o</mark> f drug delivery system using different biomaterials	Knowledge	S	nent	ation	Usage	ъ		L			inan	рu	ļ		
CLR-4: analyze the design of artificial tissues and their medical applications				alysis	elopment	restigations problems		er and	t &		Team	tion	& F	arni			
CLR-5: analyze the regulatory strategies and commercialization of biomaterials				A	deve	.≦ ×	100 100	engineer ety	nment nability	, 1	रू ज	mmunication	Mgt.	ng Le			
			ine	plem	ign/	onduct f comple	Modern		<u>a</u> 5	S	ndividual	nur	əct	Lol	-1-	7.5	53
Course O	utcomes (CO):	At the end of this course, learners will be able to:	Engi	Prof	Des	o o	₩ W	The	Env Sus	Ethics	İndi	Col	Proje	Life	PSO.	PSO-2	PSO.
CO-1:	appraise the fundamental	con <mark>cepts of</mark> biomaterials and their impact with biological response	-		2	-	-		-	2	-	-	-	-	2	3	-
CO-2:	CO-2: comprehend various biomaterials compatibility and their interaction with tissue growth				2	2	-	4	-	2	-	-	-	-	2	3	-
CO-3:	assessment of bio mimetic	es <mark>and dru</mark> g delivery system applications	1. 6	i rije	2	2	-	-	7 -	3	-	-	-	-	2	3	-
CO-4:	dissect the biological probl	le <mark>ms in tis</mark> sue engineering that require engineering expertise to solve them	2	1.20	2	2	-	3	-	3	-	-	-	-	2	3	-
CO-5:	translate highestarials as scatfolds for various clinical applications and assess regulatory controls in globa				- 2	2	_	3	-	3	-	-	-	-	-	3	-

Unit-1 - Introduction to Biomaterials

9 Hour

Properties of biomaterials (chemical, physical, mechanical and thermal), Elements of Biomaterials, preparation and characterization of biomaterials (metal, bio ceramic polymeric materials), Evaluation of biomaterials and biological responses

Unit-2 - Basic Concepts in Tissue Engineering

9 Hour

Fundamentals of tissue engineering, Tissues, Organization of tissues in vertebrate body, Cell sources, Stem cells, Cell lineages, Cell-biomaterial interactions, Cell-biomaterial response, Assessment of biocompatibility of biomaterials, cell viability assays, MTT and cytotoxicity assays, Antibacterial assessment of biomaterials, In vitro and In vivo evaluation of biomaterials

Unit-3 - Bioactive Molecules and Their Delivery in Tissue Engineering

9 Hour

Stimuli responsive in biomaterials, Bio mimetics, Dental and bone, Designing nanoparticles for drug delivery, Targeted delivery, Proteins, peptides, DNA, RNAs, oligos in drug delivery, Surface modifications, Applications in drug delivery, Advantages and limitations of biomaterials in drug delivery

Unit-4 - Biomaterials in Biomedical Applications

9 Hour

Tissue engineering, wound care and suture materials, vascular implants and bio-inspired materials, Biomimetic devices, Organ transplant, Tissue Construction, Bio artificial tissues, Connective tissues, Regeneration of connective tissues, Targeting ligands in drug delivery, Targeting ligands in cancer treatment, Cell growth and repair, Drug discovery, Impact of drug discovery and development

Unit-5 -Biomaterials and Their Marketing in Medicine

9 Hour

Technical considerations of biomaterials, Commercialization of biomaterials, Regulatory strategies for biomaterials, Clinical development with biomaterials, Clinical evaluation of biomaterials, Approval threshold of biomaterials, Supply chain of biomaterials, Strategies of global marketing, Regulatory controls in global marketing, Global authorization of biomaterials, Post-market surveillance approaches for biomaterials, Good manufacturing practice for biomaterials

	1.	Hench L. Larry, and Jones J., (Editors), Biomaterials, Artificial organs and Tissue	
Learning		Engineering, Woodhead Publishing Limited, 2005	L
Resources	2.	Ulrich Meyer, Thomas Meyer, Jörg Handschel, Hans Peter Wiesmann (2009):	١.
		Fundamentals of Tissue Engineering and Regenerative Medicine, Springer	

- cial organs and Tissue 3. Regenerative Medicine and Tissue Engineering, Edited by Jose A. Andrades, ISBN 978-953-51-1108-5, Publisher: InTech,2013
 - 4. S. Amato and B. Ezzell, (Editors), Regulatory Affairs for Biomaterials and Medical Devices, Woodhead Publisher, 2015

	Bloom's Level of Thinking	CLA-1 Avera	Continuous Learning native ge of unit test	Life Long	y Learning A-2	Summative Final Examination (40% weightage)			
		Theory (50	Practice	Theory	0%) Practice	Theory	Practice		
Level 1	Remember	15%		15%		15%	-		
Level 2	Understand	25%	ACT 1.18.71	20%	1/4- 10	25%	-		
Level 3	Apply	30%	77 - 17 1 10	25%	A 200	30%	-		
Level 4	Analyze	30%	N. J. St. 1777	25%		30%	-		
Level 5	Evaluate			10%		4 -	-		
Level 6	Create			5%		-	-		
	Tot <mark>al</mark>	100)%	10	0 %	10	0 %		

Course Designers	
Experts from Industry	Experts from Higher Technical Institutions Internal Experts
1 Dr. Giridharan, Life Cell, giridharan.a@lifecell.in	1 R. Jayakumar, Ph. D, Amrita Medical Center, Kochi 1 Dr. N. Selvamururgan, SRMIST jayakumar77@yahool.com
2 Dr. Gokuladhas Krishnan , Director, Laboratory, World Sterr	Cell 2 N. Srinivasan, Ph. D., Chettinad Health City, Chennai, 2 Dr. K. Venkatesan, SRMIST
Clinic, Chennai, care@worldstemcellclinic.com	srinivasanibms@gmail.com

Course	21DTE214T C	Course	NANOTECHNOLOGY IN REGENERATIVE MEDICINE	Course	П	PROFESSIONAL ELECTIVE	L	Т	Р	С	
Code	21B1E3141	Name	NANOTECHNOLOGY IN REGENERATIVE MEDICINE	Category		PROFESSIONAL ELECTIVE	3	0	0	3	

Pre-requisite Courses	Ni	Co- requisite Courses	Nil	Progressive Courses	Nil
Course Offerin	g Department	Biotechnology	Data Book / Codes / Standards		Nil

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Course L	_earning Rationale (CLR):	The purpose of learning this course is to:			1	- 1	rogr	am Ou	tcome	s (PO))					rogra	
CLR-1:	understanding the basic co	oncepts of nanom <mark>edicine</mark>	1	2	3	4	5	6	7	8	9	10	11	12	_	pecifi ıtcom	
CLR-2:	exploring various types of	nanomaterial <mark>s and thei</mark> r applications	lge		of	SI	1		1		ork		g				
CLR-3:	demonstrate the cutting-ed	dge nanom <mark>edicine tec</mark> hnologies for diagnosis and therapeutic applications	Knowledge	S	nent	ation	age	ъ			Μ		inance	ρ			
CLR-4:			S	Analysis	evelopment	vestigations problems	ol Usage	r and	∞ >		Team	ion	⊗ E	arning			
CLR-5:	understanding the issues i	related <mark>to toxicit</mark> y and environmental impact of nanomaterials	Engineering			ě i	P	engineer etv	ment ability	N	<u>8</u>	ommunication	Project Mgt.	<u> </u>			
			nee	len-	ign/d	onduct	ern	erc etv		တ္သ	ndividual	l III	ect	Long	7)-2	5.
Course (Outcomes (CO):	A <mark>t the en</mark> d of this course, learners will be able to:	Eng	Problem	Desi	o Col	Modern	The en	Envir <mark>on</mark> Sustain	Ethics	lndi	Con	Proj	<u>l</u> e	PSO-1	PSO-2	PSO.
CO-1:	recall the basics of nanobio	tec <mark>hnology</mark>	3	-	-	2		7	-	- I	-	-	-	-	3	-	-
CO-2:	classify the nanomaterials a	ns <mark>vehicles</mark> for drug delivery	-	2	3	2	- /		-	-	-	-	-	-	3	-	-
CO-3:	organize various types of na	an <mark>omedica</mark> l devices and their applications	3	12-4	3	3	-		-		-	-	-	-	3	-	-
CO-4:	CO-4: infer the efficient methods in the development of nanobiosensors and their applications		947	7.	3	2	- 1	-	-	-	-	-	-	-	2	-	-
CO-5:	interpret the toxicity of nano	om <mark>aterials</mark> and its remediation	<u> </u>	3	17	2	- ;	-	-	2	-	-	-	-	2	-	-

Unit-1 - Basics of Nanobiotechnology in Relation to Nanomedicine

9 Hour

Scientific principles of nanomedicine, Nanotools – types & various techniques of detection, Scanning Tunneling microscope, Atomic Force Microscope, Functional biological nanomaterials, nanoengines, Nanomaterials and their Production, Various synthesis methods of Nanomaterials, Nanodevices-Quantum Computing, Spintronic Materials and Devices, Impact of nanotechnology - Scientific and technical Impacts, Environmental Impacts, Grand challenges of nanomedicine, Ethical, Legal, and Social Issues, Government Promotion of Advancements in nanomedicine, Government Evaluation, Policy and Regulation of Nanotechnology.

Unit-2 - Nano Based Drug Delivery Systems

9 Hour

Nanomaterials as vehicles for drug delivery, Types of Nanomaterials, criteria and selection of Nanomaterials, Sources of Nanomaterials, Drug loading and release, biodegradation, Nanopatterning, Electrospinning Technology, nanopolymers, Classification of biopolymers, magnetic nanoparticles – preparation and properties, Applications of Magnetic Nanoparticles, Properties and applications of Nanotubes, Nano immunotherapy, Nanomaterials for vaccine delivery- Types of nanomaterials as vaccine adjuvants, Nanotechnology and Diagnostic Imaging, Nanomaterials as contrast agents in clinical use

Unit-3 - Nanotechnology and its Applications in Medicine

9 Hour

Nanorobots in medicine, nanorobots in nanosurgery, Nanocameras and its applications, Recombinamers, nanochips, nanoimplants, nanomaterials for bone and cartilage applications, nanomaterials for vascular applications and skin disorders, Nanomaterials in 3D Bioprinting, nanoparticle-based therapy for genetic diseases, Cell Delivery of Therapeutic Nanoparticles, nanomaterials for delivery in cells- nerve cell repair, Applications of Nanofibers in Tissue Engineering, nanomaterials for stem cells growth, Stem Cell Tracking with Nanoparticles, Nanomaterials for Stem Cell Imaging, Nanotechnology in the regulation of stem cell behavior

Unit-4 - Nano Biosensors: Properties and Applications

9 Hour

Introduction- nanobiosensors, Biosensing Techniques, unique properties of nanobiosensors, Preparation of nanobiosensors- immobilization strategies, covalent conjugation technique, Self-assembled monolayer nanomaterial, Nano biosensors for protein and DNA detection, Detection methods – optical detection and electronic detection, In vivo Biosensors, Nanowire Biosensors, Cantilever Biosensors, Applications – DNA nanobiosensor, Protein biosensor, whole cell biosensor applications, Nanobiosensor in diagnostics, Biosensors in forensic sciences

Unit-5 - Nanotoxicology 9 Hour

Overview of Nanotoxicology in Humans and the Environment, Physico-chemical characteristics dependent toxicity, Potential Adverse Effects of Engineered Nanomaterial Exposure, Respiratory Response to Pulmonary Exposure, Oral Exposure, Dermal Exposure, Handling, storage and disposal of nanomaterials, Remediation in case of nanomaterials spills, In vitro and in vivo toxicity assessment of nanoparticles, Embryonic Toxicity of Nanoparticles, Mapping Exposure onto Nanoscale, Toxicity Measures, Factors Affecting Nanoparticle Dose–Exposure and Cell Response, Green Synthesis of Nanoparticles – mechanism and Applications, Nanoparticles: Environmental Problems, nanotoxicity regulations, nanomaterials intellectual property perspective

Learning
Resources

- Melba Navarro and Josep A. Planell. Nanotechnology in Regenerative Medicine (2012), Humana Press
- 2 Jamie R. Lead, Shareen H. Doak and Martin J. D. Clift, Nanotoxicology in Humans and the Environment (2021), Springer Press
- Haiyan Xu and Ning Gu, Nanotechnology in Regenerative Medicine and Drug Delivery
 Therapy (2020), Springer Press

earning Assessm.	nent		Continuous Learnin	g Assessment (CLA)			
	Bloom's Level of Thi <mark>nking</mark>	(1/ \lambda 1 \lambda\text{Vorago of unit toot}		Life Long L CLA (10%	-2	Final Exa	mative amination eightage)
	1.2	Theory	Practice	Theory	Practice	Theory	Practice
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Level 2	Understand	25%	A 18 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	20%		25%	-
Level 3	Apply	30%	18 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	25%		30%	-
Level 4	Analyze	30%	Mar 1997 1997	25%		30%	-
Level 5	Evaluate	227 777 317		10%	1	-	-
Level 6	Create		St. 1977 L. P. L.	5%		-	-
	Total	100)%	100	%	10	0 %

Course Designers	P. Land St.	
Experts from Industry	Experts from Higher Technical Institutions	Internal Expe <mark>rts</mark>
1 Dr. S Natarajan, Advisor / Sr. Vice President - R & D; Sami Labs	1 Prof. Sundara Ramaprabhu, Department of Physics	1 Dr. Ram <mark>kumar K</mark> M, SRMIST
Limited,Bangalore. mail@samilabs.com	IIT-Madras. ramp@iitm.ac.in; ramp@physics.iitm.ac.in	. V . / 9 /
2 Dr. Gokuladhas Krishnan, Director, Laboratory,	2 Prof. Ashok M. Raichur, Department of Materials Engineering,	2 Dr. N <mark>. Selvamu</mark> rurgan, SRMIST
World Stem Cell Clinic, Chennai,care@worldstemcellclinic.com	IISc, Bangalore. amr@materials.iisc.ernet.in	

Course		Course	TISSUE ENGINEERING FOR REGENERATIVE MEDICINE	Course	_	PROFESSIONAL ELECTIVE	L	Τ	Р	С	
Code	21B1E4161	Name	11330E ENGINEERING FOR REGENERATIVE WIEDIGINE	Category	Е	PROFESSIONAL ELECTIVE	3	0	0	3	

Pre-requisite Courses	N	Co- requisite Courses	Nil Progress Course	Nil
Course Offeri	ing Department	Biotechnology	Data Book / Codes / Standards	Nil

Course L	Learning Rationale (CLR): The purpose of learning this course is to:	$\Box B$	4	1-		Progr	<mark>am O</mark> ı	itcome	es (PO))					rogra	
CLR-1:	from the viewpoint of engineers, explain the foundations of tissue engineering and tissue restoration	1	2	3	4	5	6	7	8	9	10	11	12		Specifi utcom	
CLR-2:	describe understanding of tissue engineering's clinical applications			14				ility								
CLR-3:	describe understanding of tissue engineering's clinical applications	g		₹	Is of	b.	society	ustainability	l.	Work		g				
CLR-4:	state engineering students to think more on artificially generated tissues for their tissue engineer applications	Š	Sis	velopment	stigations	Jsage	and so	S		eam W	<u>_</u>	Finance	arning			
CLR-5:	talk about the technology of 3D-bioprinting and Explain the methods for cutting-edge bioactive tiss engineering research	ering Knc	n Analysis	n/develo	prof	Tool Usage	engineer	ment &		∞ ~	ommunication	Mgt. &	ong Lear			
Course C	Outcomes (CO): At the end of this course, learners will be able to:	Engine	Problem	Design	Conduct	Modern	The en	Environment	Ethics	Individual	Commu	Project	Life Lor	PS0-1	PSO-2	PSO-3
CO-1:	examine the elements that make up the tissue architecture	3	2	2	11-3	-	1	-	-1	-	-	-	-	-	2	-
CO-2:	show the traits of stem cells <mark>and thei</mark> r importance in medicine	3	3	3	1	-		<i>i</i> -		-	-	-	-	-	3	2
CO-3:	describe knowledge of biomaterials' characteristics and wide range of applications	1	1			_	-	-	-	-	-	-	-	-	2	-
CO-4:	analyze the role of tissue engineering and stem cell therapy in organogenesis	3	3	_ 2	1 K -	-		-		-	-	-	-	-	2	2
CO-5:	create novel biomaterials and emerging techniques for creating effective tissue and organ replacem and examine the in vivo and in vitro testing of biomaterials	ents 3	3	2	2	7-	X	-	-	-	-	-	-	-	3	3

Unit-1 - Basics of Tissue Engineering

9 Hour

Cellular Basis of Regeneration-Molecular Basis of Regeneration; Overview of tissue engineering-Simple terms used in tissue engineering-Present possibility of development in tissue engineering -Therapeutic application of tissue engineering; Components used in tissue engineering-Primary cells, cell lines and immortalization of cells; Evaluation of tissue characteristics, appearance, cellular component -Cell differentiation, cell migration, and processes determining a cell's fate; Extracellular matrix (ECM) constituent and their regulation of cell behavior-Mechanical measurements of the ECM component-Physical properties of the ECM component properties of the ECM component properties of the ECM component properties of the ECM component properties of the ECM component properties of the ECM component properties of the ECM component properties of the ECM component properties of the ECM component properties of the ECM component properties of the ECM component properties of the ECM component properties of the ECM component properties of the ECM component properties of the ECM component properties of the ECM component properties of the ECM component properties of the ECM compo

Unit-2 - Interactions and Culture of Cells and Tissue

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Tissue kinds-Tissue constituents -Tissue healing-Engineering wound healing-Sequence of events of wound healing; Three-Dimensional Cell Culture-Organ Culture-Organ Culture; Basic Wound Healing Overview-The use of growth factors: Function of VEGF/angiogenesis-Various methods of angiogenesis and its significance-Fundamental characteristics of growth factors, Cell-Matrix Interactions-Cell-Cell Interactions; Telomeres and Self-renewal

Unit-3 - Biomaterials and Their Implementations

9 Hour

Basics of the science of biomaterials; Idea of biocompatibility; Categories of biomaterials-Fundamental traits of biomaterials-Disinfection and sterilization of biomaterials; Biomaterials' physico-chemical characteristics: Mechanical (elasticity, yield stress, ductility, toughness, strength, fatigue, hardness, wear resistance)- Tribological (friction, wear, lubricity)-Morphological and texture, Physical (electrical, optical, magnetic, thermal)-Chemical and biological characteristics; Rudiments in contact with the surface of a biomaterial: blood composition, plasma proteins, cells, tissues; Scaffolds' function in tissue engineering-Biopolymers; Modifications of Biomaterials-In vitro testing of biomaterials-In vivo testing of biomaterials

Unit-4 - Stem Cells and Their Applications in Tissue Engineering

9 Hour

Overview of Stem Cells-Various kinds of Stem cells-Hematopoietic differentiation pathway of stem cells-Potency of stem cells-Plasticity of stem cells-Sources of embryonic stem cells-Sources of hematopoietic and mesenchymal stem cells; Stem Cell markers, FACS analysis; Types & sources of stem cell with characteristics: Embryonic stem cells and Adult stem cells-Comparison between- embryonic and adult stem cells; Bone marrow, primordial germ cells; Cancer stem cells; Induced pleuripotent stem cells

Unit-5 -Therapeutic Aspects of Tissue Engineering

9 Hour

Discussion on Stem cell therapy- Therapies for spinal cord injury, muscular dystrophy-Orthopedic applications-Stem cells and Gene therapy; Tissue engineering of bones-Tissue engineering of cartilage-Neural tissue engineering-Skin tissue engineering-Cardiovascular tissue Engineering-Therapeutic applications; Overview on the basic principles for Biofabrication and 3D printing-Methods and materials-Applications of Biofabrication and 3D printing: Lab-on-chip, Organ-on-chip; Innovative bioactive research; Regenerative medicine

Learning Resources

- 1 Clemens Van Blitterswijk, Jan De Boer, "Tissue Engineering", 2nd Edition Academic Press, 2014.
- 2 Robert Lanza, Robert Langer, Joseph Vacanti, "Principles of Tissue Engineering", 4th Edition -Academic Press, 2013.
- 3 John P. Fisher, Antonios G. Mikos, Joseph D. Bronzino, Donald R. Peterson, "Tissue Engineering: Principles and Practices", 1st Edition - CRC Press, 2017
- 4 Buddy D. Ratener, Allan S. Hoffman, Frederick J. Schoen, Jack E. Lemons, "Biomaterial Science: An Introduction to Material in Medicine", 3rd edition – Academic Press. 2013.
- 5 Lijie Grace Zhang, John Fisher, Kam Leong, "3D Bioprinting and Nanotechnology in Tissue Engineering and Regenerative Medicine", 1st Edition Academic Press, 2015.

Learning Assessme	nt			1,862.5			
	Bloom's Level of <mark>Thinking</mark>	Form CLA-1 Avera (50	Summative Final Examination (40% weightage)				
		Theory	Practice	Theory	Practice	Theory	Practice
Level 1	Remember	20%	100 May 100 Ma	15%	-	15%	-
Level 2	Understand	20%	N 727 2	20%	-	25%	-
Level 3	Apply	30%	100	25%		30%	-
Level 4	Analyze	30%		25%	- U	30%	-
Level 5	Evaluate			10%		9 -	-
Level 6	Create		- 4114	5%	- 4	-	-
	Tota <mark>l </mark>	10	0 %	100	0 %	100	0 %

Course Designers		
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
1 Dr. Harikrishna Varma, SCTIMST, Thiruvan <mark>anthapura</mark> m, India; —	1 Dr. Sourabh Ghosh, IIT Delhi, India, sghosh08@textile.iitd.ac.in	1 Dr. Koustav Sarkar, SRMIST
head-bmtw@sctimst.ac.in	VITEARA · LEAD round	
2 Dr. Dipak Datta, CDRI, Lucknow, India; dipak.datta@cdri.res.in	2 Dr. Rathindranath Baral, CNCI, Kolkata., India,	2 Dr. N. Selvamururgan, SRMIST
	baralrathin@hotmail.com	

Course	21BTF417T	Course	BIOREACTORS IN TISSUE ENGINEERING	Course	Е	PROFESSIONAL ELECTIVE	L	Т	Р	С
Code	210164171	Name	BIOREACTORS IN 1135UE ENGINEERING	Category	E	PROFESSIONAL ELECTIVE	3	0	0	3

Pre-requisite Courses	N	Co- requisite Courses	Nil	Progressive Courses	Nil
Course Offer	ing Department	Biotechnology	Data Book / Codes / Standards		Nil

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Course L	Learning Rationale (CLR): The purpose of learning this course is to:	U/	7	4			Progr	am Ou	tcome	s (PO)					rograr	
CLR-1:	provide the basic concepts of tissue engineering and bioreactors from the perspective of engineers		1 1	2	3	4	5	6	7	8	9	10	11	12		pecifi itcom	
CLR-2:	identify the 3D- culture of stem cells an <mark>d organoge</mark> nesis	Ç	e Ge		of o	દા					Work		g				
CLR-3:	identify the role of stem cells in clinical applications of different disease conditions		owledge	S	Jent	vestigations problems	Usage	ъ			\ 		Finance	Бū			
CLR-4:	identify the safety and efficacy of b <mark>ioreactors</mark>	2		Analysis	ndol	/estig probl		r and	× ×		Team	fion	∞ర	arni			
CLR-5:	create the strategies for designing clinically relevant bioreactors	2	= `	4	n/development of ons		T00	engineer ety	Environment Sustainability		<u>∞</u>	mmunication	Mgt.	g Le			
				plem	fign/	nduct in complex	Modern	ne eng ciety	ron	SS	Individual	l III	Project	Long	7)-2	က္
Course C	Outcomes (CO): At the end of this course, learners will be able to:			g S	Desi	Con	Mod	The	Envi Sus <mark>i</mark>	Ethics	lpdi	Con	Proj	Life	PSO.	PSO	PSO
CO-1:	apply the basic understanding of large-scale production stem cells in bioreactors	17 / 2	2 - 2	2	2	2	-	/	-		-	-	-	2	3	-	-
CO-2:	discuss the 3D- culture syste <mark>ms and</mark> artificial organs	i., 13	3	2	2	3	_	4	-	-1	-	-	-	2	3	-	-
CO-3:	identify the bioreactor-based strategies to generate organoids		3	2	2	2	-	_	- 1	-	-	-	-	2	3	-	-
CO-4:	understand the role of bioreactors in the development of drug development and therapy		3 .	3	3	3	-	-	-		-	-	-	2	2	-	-
CO-5:	explain the large scale production of stem cells	- 4	3 - 2	2		2	_		-		_	-	-	-	2	-	_

Unit-1 - Tissue Organization and Bioreactors

9 Hour

Complexity and organization of the organ system-Measurement of tissue characteristics, appearance, tissue types-Types of Bioreactors- Cells for tissue engineering— Perfusion Bioreactors for 3D cultures, Spinner Flask Bioreactor-Rotating Wall Bioreactor, Compression Bioreactor, Strain Bioreactor-static culture, stem cell cultivation in scaffold Bioreactor systems -Hydrostatic pressure Bioreactor, Flow Perfusion Bioreactor, Combined Bioreactor- Clinic grade production of mesenchymal stem cells.

Unit-2 - Scaffold and Functionalized Tissue Engineering

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Functional tissue engineering and role of Biomechanics in a 3D environment—Controlled release strategies in tissue engineering -Tissue fabrication technology, microfabrication -Bioreactors role in tissue engineering of Cartilage-Cardiovascular tissue, Vascular tissue, musculoskeletal tissue and Skin—Bone-microfluidic devices and microbioreactors for stem cell micro environment—Perfusion bioreactors for granulocyte progenitor cell growth; Bioreactor stimulation-Mechanics and Controlled Parameters of Bioreactors—Engineering stem cell niches in bioreactors—Oxygen tension, Scaffold/substrate cues-Decellularized ECMs, Mechanical forces, Electrical stimulation, Flow shear rate, and paracrine and autocrine factors.

Unit-3 - Applications of Bioreactors

9 Hour

Novel approaches in bioreactor systems for stem cell seeding of vascularized bioscaffolds-Bioreactor-based strategies with reconstructive applications- Stem cell cultivation in scaffold-bioreactor systems; Physiological biomimicry-Understanding Mechanical forces on organs and functional aspects-Control and Feedback Control in Mechatronics for mechanical stimulation; Scaffolds and Constructs for Bioreactor Systems – Organoids and organ-on-chip-Boprinting- Applications of growth factors-Role of VEGF. Angiogenesis, Basic properties, Cell-Matrix, Cell-Cell Interactions, Control of cell migration in tissue engineering.

Unit-4 - Biomaterial and Tissue Engineering

9 Hour

Biomaterials: Properties of Biomaterials, Surface, bulk, mechanical and biological properties-Scaffolds & tissue engineering, Types of Biomaterials, biological and synthetic materials-Biopolymers, Applications of biomaterials, Sensing and Automation in bioreactor systems-Bioreactors in drug discovery and implant testing; Bioreactors in clinics-Stem cell cultivation in scaffold-bioreactor systems-Large-scale bioreactor cultivation of pluripotent stem cells-Engineering of functional bone tissue from human stem cells-Miniature bioreactors for precise, systematic studies of stem cell environments.

Unit-5 - Clinical Applications of Bioreactors

9 Hour

Clinical applications - Stem cell therapy, Molecular Therapy-In vitro organogenesis, Neurodegenrative diseases-spinal cord injury, heart disease, diabetes, burns and skin ulcers-muscular dystrophy, orthopedic applications-Stem cells and Gene Therapy-Physiological models, tissue engineering therapies, product characterization-components, safety, efficacy. Preservation – Product and process design toward industrial tissue engineering manufacturing - Patent protection and regulation of tissue engineered products, ethical issues in tissue engineering.

Learning	1	Bioreactors: Design			Novel	Applications,	2016,	Chaudri,	Wiley	2	Tissue engineering and Regeneration, 2022, Heinz Redl, Springer Publications, ISSN: 2731-0558
Resources		pulications; ISBN: 9	78352768336	9				23 [1]			

Learning Assessm	ent		1 J							
			Continuous Learning	Cum	mativa					
	Bloom's Level of Thinking	CLA-1 Aver	mative age of unit test 50%)	Life Long L CLA (10%	-2	Summative Final Examination (40% weightage)				
		Theory	Practice	Theory	Practice	Theory	Practice			
Level 1	Remember	15%	- 18 A. S. W. S. W.	15%		15%	-			
Level 2	Understand	25%		20%		25%	-			
Level 3	Apply	30%	367 Par (455) 17 Par (4	25%		30%	-			
Level 4	Analyze	30%	Street Allen and	25%	7	30%	-			
Level 5	Evaluate		A 1 1 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	10%	. 3 - / .	-	-			
Level 6	Create		PRESENT STATES TO A	5%		-	-			
	Total -	1	00 %	100	%	10	0 %			

Course Designers	EST NOTATION OF THE PARTY OF THE STREET	L_
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
Dr.Ramesh Babu Boga, BogaR Laboratories, Hyderabad, brameshb@msn.com	1 Dr. Ajaikumar B. Kannumakkara , IITG, kunnumakkara@iitg.ernet.in	1 Dr. Kanagaraj Palaniyandi, SRMIST
2 Mr.J.B. Vijayakumar BioArtis Life Sciences Pvt. Ltd. email: contact@bioartis.in	2 Dr. Suttur S Malini, University of Mysore, drssmalini@gmail.com	2 Dr. R. Satish, SRMIST

Course	21BTE418T Cours	DEVELOPMENTAL BIOLOGY IN TISSUE ENGINEERING	Course	_	PROFESSIONAL ELECTIVE	L	Т	Р	С	
Code	Nam	DEVELOPMENTAL BIOLOGY IN TISSUE ENGINEERING	Category	E	PROFESSIONAL ELECTIVE	3	0	0	3	

Pre-requisite Courses	N	Co- requisite Courses	Nil	Progressive Courses	Nil
Course Offeri	ng Department	Biotechnology	Data Book / Codes / Standards		Nil

Course L	earning Rationale (CLR):	The purpose of learning this course is	s to:	H	-4			Progr	am Ou	tcome	s (PO)					rograi	
CLR-1:	describe cell-cell interaction	ons from the context of tissue engineering	1.0	1	2	3	4	5	6	7	8	9	10	11	12	_	pecifi ıtcom	
CLR-2:	illustrate the types of cell	specification a <mark>nd germ la</mark> yers		dge		of	SL					Work		9				
CLR-3:	provide information on ne	urulation an <mark>d types of</mark> mesoderm tissues	Marin Marin	Knowledge	တ	nent	ation	Usage	ъ			×		Finance	рu			
CLR-4:	summarize on heart and g	gut tube d <mark>evelopm</mark> ent	4.17.2.17.4		Analysis	velopment of	vestigations problems		er and	× ×		Team	ig.	∞ర	arni			
CLR-5:	appraise on ageing and ty	rpes of <mark>regenerat</mark> ion		ering		deve	t inve	<u>0</u>	enginee ety	Environment Sustainability		<u>a</u>	mmunication	Mgt.	ng Le			
			Sales State of the Sales	9	roblem	ign/dev tions	induct inv complex	Modern		iron tain	S	Individual	E E	Project	Long	7)-2	5.3
Course C	Outcomes (CO):	At the end of this course, learners wi	II be able to:	Engi	Prof	Des	Con	₩ W	The	Env S <mark>us</mark>	Ethics	lpi	Cou	Proj	Life	PSO.	PS0	PSO
CO-1:	interpret on the basics of	sign <mark>aling me</mark> chanisms	1 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	-		-	2	-	/	-	3	-	-	-	-	-	-	2
CO-2:	recall the concepts in cel	I co <mark>mmitme</mark> nt and morphogen gradients		<i>j</i> - '.	Ğ-	40	2	-	4	-	3	-	-	-	-	-	-	2
CO-3:	describe the genetics of n	eu <mark>ral tube</mark> and kidney development		- X	17.		2	-	-	7 -	3	-	-	-	-	-	-	3
CO-4:	analyze the development	of <mark>heart an</mark> d the digestive organs		134	1.7	1.5	3	-	-	-	3	-	-	-	-	-	-	2
CO-5:	understand the processes	s o <mark>f germ c</mark> ell migration and types of tissue	regeneration		4.0	-	3	-		-	3	-	-	-	-	_	3	-

Unit-1 - Cell Communication in Development

9 Hour

Differential cell affinity – Cadherins and ce<mark>ll adhes</mark>ion – Cell migration – Cell induction – Cell competence – Paracrine factors – Signal transduction cascades – RTK pathway – Jak-STAT pathway – Juxtracrine signaling – Notch pathway – Developmenta<mark>l signals</mark> from ECM – Epithelial-to-mesenchymal transition

Unit-2 - Cell Specification and Germ Layers

9 Hour

Cell commitment – Cell determination - Cell specification – Autonomous, Conditional and Syncytial specifications - Morphogen gradients – Cell fate – Cell lineage – Stem cells in development – Stem cell niches – Regulatory microenvironments – Germ layers – Ectoderm, mesoderm and endoderm

Unit-3 - Neurulation, Somitogenesis and Kidney Development

9 Hour

Neurulation – Primary and secondary neurulation - Neural tube formation – DV axis of the neural tube – Brain organization – Ectodermal placodes – Types of mesoderm – Somite formation – Specification of intermediate mesoderm – Development of mammalian kidney

Unit-4 - Formation of Heart, Gut Tube and Respiratory Tube

9 Hour

Specification of lateral plate mesoderm – Cardiac precursor cells – Determination of cardiac domains – Formation of heart chambers – Specification of gut tissue – Development of liver and pancreas – Origin and development of respiratory tube

Unit-5 - Concepts on Tissue Regeneration

9 Hour

Genetic causes of ageing -Germplasm -Specification of primordial germ cells - Germ cell migration - Regeneration - Epimorphic regeneration - Morphallactic regeneration - Regeneration in mammalian liver

	1	Developmental Biology (2020): Scott F. Gilbert and Michael J.F. Barresi, Twelfth Edition,	3	Principles of Development (2015): Lewis Wolpert, Cheryll Tickle and Alfonso Arias, Fifth Edition,
Learning		Oxford University Press, Inc.		Oxford Publishers, Inc.
Resources	2	Essential Developmental Biology (2012): J.M.W. Slack, Third Edition, Wiley-Blackwell	4	Principles of Developmental Genetics (2014) S.A. Moody (Ed.) Second Edition,
		Publishers	- 1	Academic Press

			Continuous Learning Assessment (CLA)								
	Bloom's Level of Thinking	Form CLA-1 Averaç (50	ge of unit test	CL	g Le <mark>arning</mark> LA-2 0%)	Summative Final Examination (40% weightage)					
		Theory	Practice	Theory	Practice	Theory	Practice				
Level 1	Remember	15%		15%	2 - 1	15%	-				
Level 2	Understand	25%	A	20%	- A- 10	25%	-				
Level 3	Apply	30%	45.50	25%	A 100 TO	30%	-				
Level 4	Analyze	30%	A. J. S. W.	25%		30%	-				
Level 5	Evaluate			10%	1-2	4 -	-				
Level 6	Create		St. 200 (1977) 19 19 19	5%		-	-				
•	Tot <mark>al</mark>	100	1%	- 10	00 %	10	0 %				

Course Designers	STORES CONTRACTOR STORES AND AND AND AND AND AND AND AND AND AND	
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
1 Dr. B.R.Desikachari, Medical Director, Westminster Health Care	, 1 Dr. K. Subramaniam, Indian Institute of Technology Madras,	1 Dr. S. Kiran <mark>kumar,</mark> SRMIST
Chennai, brdesikachari@hotmail.co <mark>m</mark>	Chennai,subbu@iitm.ac.in	
2 Dr. V.L.Ramprasad, MedGenome Labs Ltd, Bengaluru	2 Dr. Sudha Warrier, Associate Professor, Manipal University	2 Dr. N. Selv <mark>amuruga</mark> n, SRMIST
ramprasadv@medgenome.com	sudha.warrier@mannipal.edu	

Course	21BTE419T Cou	ADVANCED IMMUNOLOGY AND VASCULAR ENGINEERING	Course	_	PROFESSIONAL ELECTIVE	L	Τ	Р	С	1
Code	Nar		Category	E	PROFESSIONAL ELECTIVE	3	0	0	3	

Pre-requisite Courses	N	Co- requisite Courses	Nil Progressi Courses	Nil
Course Offeri	ng Department	Biotechnology	Data Book / Codes / Standards	 Nil

Course L	earning Rationale (CLR): The purpose of learning this course is to:	111	4			Progr	<mark>am Ο</mark> ι	itcome	es (PO)					rograi	
CLR-1:	provide the most recent advancement in the field of immunology from the perspective of bioengineers	1	2	3	4	5	6	7	8	9	10	11	12	_	pecifi utcom	
CLR-2:	enrich with knowledge on immunobiology and immune responses related to regeneration and transplants		h.	4	2	<u>.</u>	Ü	oility								
CLR-3:	recognizing the issue of shortage of organ donors as the major limitations in the transplantation and finding solution for the same	vledge		ant of	vestigations or oblems	<u>o</u>	society	stainability	N.	Work		ance	_			
CLR-4:	learning of various treating metho <mark>ds for inj</mark> ury and the significance of vascular engineering	S S	Sis.	bme	tigal	Sag	and	S		eam	_	뜶	ming			
CLR-5:	train and develop skills among the students to explore strategies for Immunotherapy and Stem cells therapy	eering Know	em Analysis	n/development of].⊑ ⊑	m Tool Usage	engineer a	Environment &		~	Sommunication	ct Mgt. &	ong Lear	_	2	က
Course C	Outcomes (CO): At the end of this course, learners will be able to:	Engin	Problem	Desig	Conduct	Modern	The e	Enviro	Ethics	Individual	Comn	Project	Life L	PSO-1	PSO-2	PSO-3
CO-1:	acquire knowledge on the latest tools for diagnosis of diseases	2	1	- 16	-	-	-		-+	-	-	-	-	-	-	2
CO-2:	gain knowledge in molecula <mark>r and im</mark> munological basis of diagnosis	2	2	17-	2	-	-	-		-	-	-	-	-	2	2
CO-3:	able to appreciate the releva <mark>nce of c</mark> linical immunology	2	3	772	- 2	2	-	-	- :	-	-	-	-	-	3	2
CO-4:	acquire knowledge on vascu <mark>lar biolo</mark> gy and vascular tissue engineering	3	3	51	2	2	-	-		-	-	-	-	-	3	2
CO-5:	acquire knowledge on host vs Graft rejection, neovascularization and the significance of immune syste in these processes	n 2	3	-	2	7-	Š	-		-	-	-	-	2	3	2

Unit-1 - Basics of Immunology 9 Hour

Components of innate and acquired immunity: Phagocytosis; Complement and Inflammatory responses; Haematopoesis; Organs and cells of the immune system – primary and secondary lymphoid organs; Lymphatic system; Lymphocyte circulation; Lymphocyte homing; Mucosal and Cutaneous associated Lymphoid tissue. (MALT & CALT); Mucosal Immunity; Antigens – immunogens, haptens; Major Histocompatibility Complex – MHC genes, MHC and immune responsiveness and disease susceptibility, HLA typing; Immunological considerations for stem cell banking (self-study)

Unit-2 - Antigen and Antibodies 9 Hour

Immunoglobulins-basic structure, classes & subclasses of immunoglobulins, antigenic determinants; Mutagenic organization of immunolobulin genes; B-cell receptor; Immunoglobulin superfamily; Principles of cell signaling; Basis of self – non-self-discrimination; Kinetics of immune response, memory; B cell maturation, activation and differentiation; Generation of antibody diversity; T-cell maturation activation and differentiation; and T-cell receptors; Functional T Cell Subsets; Cell-mediated immune responses, ADCC; Cytokines-properties, receptors and therapeutic uses; Antigen processing and presentation – endogenous antigens, exogenous antigens, non-peptide bacterial antigens and super-antigens; Cell-cell co-operation, Hapten-carrier system; Current status of Immunotherapy (self-study)

Unit-3 - Transplantation 9 Hou

Immunologic targeting of cancer stem cell population- tumor- initiating cells and their immuno targeting; CML of Haematopoietic stem cells-allogenic transplantation of HSC- Graft versus leukemia effect, T-cells, B-cells and NK cells as mediators of graft versus leukemia –Malignant progenitors targeting by graft- versus-leukemia; Cells and factors involved in transplant acceptance and rejection; (self-study); Recent advances in transplantation

Unit-4 - Tissue Regeneration 9 Hour

Stem cells in Regenerative Biology, Stem cell therapy for treatment of ulcers, skin burns, neurodegenerative diseases, Immunological considerations and the potential barriers for stem cell therapy; Immuno-suppressive therapies (self-study)- Immunological Aspects of Allogeneic and Autologous Mesenchymal Stem Cell Therapies

Unit-5 - Grafts and Vascularization

9 Hour

Significance of acellular grafts in regeneration; tissue injury and immune responses; potential barriers to engraftment of human pluripotent stem cells; Transplantation – Immunological basis of graft rejection (self-study); mast cells in allograft rejection Mouse models of graft-versus-host disease; Clinical transplantation and immunosuppressive therapy; Importance of vascularization in Tissue Engineering, Signaling pathways of angiogenesis, Vascular Remodeling; Stem cells and scaffolds for vascular regeneration.

	1	Vascularization: Regenerative Medicine and Tissue Engineering, edited by Eric M.	3	The Immunological Barriers to Regenerative Medicine. Editors-Paul J. Fairchild, Humana Press
Learning		Brey, CRC Press 2017		2013
Resources	2	Stem Cell Transplantation, edited by Carlos López-Larrea, Antonio López Vázquez,	4	Immunotherapy in Bioregenerative Medicine, by Dmytro Klokol, Dato Sri Mike K. S. Chan,
		and Beatriz Suárez Álvarez. Springer 2016		Roni Lara Moya · 2022

Learning Assessment								
	Bloom's Level of T <mark>hinking</mark>		Continuous Learnin Formative CLA-1 Average of unit test (50%)		ng Assessment (CLA) Life Long Learning CLA-2 (10%)		Summative Final Examination (40% weightage)	
	•		Theory	Practice	Theory	Practice	Theory	Practice
Level 1	Remember		15%	William Street St.	15%	- 4	15%	-
Level 2	Understand		25%	MARCH 1984 1994	20%		25%	-
Level 3	Apply		30%	NE 1721 74	-25%		30%	-
Level 4	Analyze		30%	W. Marketta 2011	25%	-	30%	-
Level 5	Evaluate		472-	The second section of the second	10%			-
Level 6	Create				5%		-	-
	Total		100) %	100 %	ý –	100) %

Course Designers	1.7	
Experts from Industry	Experts from Higher Technical Institutions	Internal Exp <mark>erts</mark>
1 Dr. Vani, Jeevan Stem Cell Foundation, Chennai,	1 Dr. S. Sittadjody, Research Fellow, Institute for Regenerative	1 Dr. N <mark>. Selvamu</mark> rurgan, SRMIST
stemcell@jeevan.org	Medicine, Winston-Salem, USA. ssdjody@gmail.com	
2 Dr. Satheesh K. Sainathan, Study Director, Phenotypic Services,	2 Dr. Yuvaraj Sambandam, Assistant Professor, Surgery, Transp	plant 2 D <mark>r. R. Satis</mark> h, SRMIST
Eurofins Discovery, St Charles, Missouri, United States	Surgery Division, North western University, USA	



SRM INSTITUTE OF SCIENCE AND TECHNOLOGY

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B.Tech / M.Tech (Integrated) Programmes-Regulations 2021- Volume-8-Biotechnology - Syllabi(Revised August 2024) - Control Copy