# 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 F	Progressive 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:	11.	$\overline{A}$	<u> </u>		Progr	am Ou	itcome	s (PO	)					rogra	
CLR-1:	describe the basic principle	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 Sto	pichiometry <mark>equations a</mark> nd material balances	ge	edge edge successions ance													
CLR-3:	demonstrate the behavior	of fluids a <mark>nd fluid flo</mark> w phenomena	owledge	S	nent	ation	Usage	ъ		L.			Finance	Вu			
CLR-4:	1.4: describe the principles of filtration working of filtration equipment's and concept of aritation																
CLR-5:	R-5:   illustrate the basic concepts and <mark>laws of t</mark> hermodynamics																
			<u> </u>	Problem	lgin/	onduc	Modern	engety	iron <mark>tain</mark>	S	Individual	J III	Project	흐	7	)-2	53
Course C	Outcomes (CO):	At the end of this course, learners will be able to:	Eng	Po	Des	g g	₩ W	The en	Env Sus	Ethics	Indi	Sol	Proj	Life	PSO	PSO.	PSO
CO-1:	perform unit conversions a	nd <mark>stoichio</mark> metric calculations	2	3	100	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	-	-	-	-	f -		-	-	-	-	-	-	-
CO-4:	formulate the concepts of fil <mark>tration a</mark> nd agitation in processes				€	الخز	-	-	-	- 1	-	-	-	-	-	-	-
CO-5:	comprehend the basic concepts and laws of thermodynamics for different processes				1-54	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	Learning Rationale (CLR): The purpose of learning this course is to:	C.H	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	11 12 Specific Outcomes			
CLR-2:	CLR-2: explain and analyze the basic concepts of convection as applied to various flows and geometry				of		ciety			~						
CLR-3:	illustrate principles of mass transfer, Diffusion phenomena, and calculate mass transfer rates	ring		nt of	ions	ons	socie			Work		inance				
CLR-4:	alunidate the principles of drying different types of drier and calculate drying time for different drying		alysis	velopment	estigations blems	ol Usage	er and	st ≻		Team	tion	8 8	arning			
CLR-5:	demonstrate the concept of distillation, extraction and adsorption	eering	em An	3n/deve	luct inv	ern Too	engineer	ironment tainability	S	vidual &	ommunication	ct Mgt.	ong Le	<del>-</del>	-5	.3
Course C	Outcomes (CO): At the end of this course, learners will be able to:	Engine	Prob	Designation	Cond	Mode	The	Envir	Ethic	Indiv	Com	Project	Life L	PSO.	PS0-2	PSO-
CO-1:	analyze steady state heat co <mark>nduction</mark> and calculate the rate of heat transfer	2	2	18/5/	47-	-	1	-	-1	-	-	-	-	-	-	-
CO-2:	apply the basic concepts of convection and calculate the heat transfer coefficient	32/	3	3	-	-			-	-	-	-	-	-	-	-
CO-3:	interpret mass transfer princ <mark>iples and</mark> solve diffusion problems	- F11-E	2	- 3	-	-	-	-	-	-	-	-	-	-	-	-
CO-4:	calculate drying time for different periods of drying	4.0	2	2	42	-		-	- :	-	-	-	-	-	-	-
CO-5:	comprehend the various types of distillation, extraction and adsorption for different processes		2	- 2	1	_	-	_	_ 1	_	-	-	_	-	_	-

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			A Profession Line	The Contract of the Contract o			
			(	Continuous Learning	Cum	mativa		
	Blo <mark>om's</mark> Level of <mark>Thinkin</mark> g	CLA	Formative 1-1 Average of u (45%)		CL	Learning 4-2 %)	Final Ex	mative amination eightage)
		Theory	100	Practice	Theory	Practice	Theory	Practice
Level 1	Remember	20%	7 A T	122 - 42 - Nº	20%		20%	-
Level 2	Understand	20%		-	20%		20%	-
Level 3	Apply	30%			30%		30%	-
Level 4	Analyze	30%		- 1017	30%	4	30%	-
Level 5	Evaluate	-/4 Y-		- 1	-		-	-
Level 6	Create			- 11911	-	-V /	-	-
	Total	1 2 1	100 %	15/10	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

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

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	Dat	a Book / Codes / Standards		Nil

Course L	earning Rationale (CLR): The purpose of learning this course is to:	7 4			_ "1	rogra	am Ou	itcome	es (PC	<b>)</b> )					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		74	1	of		ety			~						
CLR-3:	determine the types of fatty acids, and use a variety of tests and reagents	edge		nt of	ions	a)	society			Work		Finance				
CLR-4:	become familiar with chromatographic methods and use them to isolate and characterize various biologisubstances	존	alysis	evelopment	investigations problems	Usage	and	t &		Team	tion	∞ర	earning			
CLR-5:	recognize the fundamentals of various reagents and how they interact with biomolecules for measurement	ering	٦	deve		Tool	engineer	Environment Sustainabilit		रू ज	Sommunication	Mgt.				
		e	.oplem	/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:	Eng.	Prof	Des	Som	Moo	ТЪе	Sus	Ethics	ibu	Son	Proj	Ę	PSO	-0Sd	PSO-3
CO-1:	perform basic professional skills related to solutions, pH, and buffer preparation, as well as nume calculations, focusing on the laboratory	ical 3	3	3	} -	1		_		-	1	-	-	-	3	-
CO-2:	identify the various ways in which different types of carbohydrates respond to chemical tests	X 17	3	3	-		-	-	1	-	-	-	-	-	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		3	7	-	-	1	-	-	-	-	-	3	-
CO-5:	describe the measurement of biomolecules in clinical and dietary samples	X 3	3	-	3	1	-	_	- 1	-	-	-	-	-	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	ment			ROLL STATE	21.					
			- Albert		g Assessment (CLA)		4	Sumi	native	
	Bloom's Level of Thinking	exper	ge of first cycle riments 0%)	exper	of second cycle iments 0%)		Examination veightage)	Final Examination (0% weightage)		
	*	Theory	Practice	Theory	Practice	Theory	Practice Practice	Theory	Practice	
Level 1	Remember		15%	- 300 F 10 F	15%	(4) V S	15%	-	-	
Level 2	Understand		20%	100	20%	A . A-	20%	-	-	
Level 3	Apply		25%	3 A 1 2 2 2	25%	12 275	25%	-	-	
Level 4	Analyze	-	25%	The second of	25%	J 445 - 1	25%	=	-	
Level 5	Evaluate		10%		10%	-	10%	-	-	
Level 6	Create		5%	- \	5%	- / (	5 <mark>%</mark>	-	-	
	Total	10	0 %	100	0 %	10	00 %		-	

Course Designers	1514	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	Practice Practice	Theory	Practice		
Level 1	Remember	15%	7.4. 4.6.	15%		15%	-		
Level 2	Understand	25%	A COMPANY	20%	- A-	25%	-		
Level 3	Apply	30%	47.54.5 (6)	25%		30%	-		
Level 4	Analyze	30%	A. J. S. 1777	25%		30%	-		
Level 5	Evaluate			10%	- C - C	4	_		
Level 6	Create		Carried Andrew Pr	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. Muth <mark>ukumar,</mark> 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.		ork		Э			
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		
C 0	viscource (CO)	Engineering	roblem	sign/	Conduct of comple	Modern	ie eng ciety	Environ Sustain	Ethics	ndividual	ommunication	Project	Ę	-So-1	PSO-2 PSO-3
Course O	utcomes (CO):  At the end of this course, learners will be able to:	ш	٩	2 8	ಕಟ	ĭ	투 8	교 <mark> S</mark>	E	<u>=</u>	ပိ	P	Life	PS	8 8
CO-1:	distinguish between prokaryo <mark>tic and e</mark> ukaryotic cells using microscopic analysis		. 3	3	-	-	1	-		-	-	-	-	-	3 -
CO-2:	gain proficiency in identifying the cellular structures	-	7	3	3	- /		-	1	-	-	-	-	-	- 3
CO-3:	acquire skills to isolate cells and cell organelles and relate with cell division	35	3	3	34	- (		-	ė	-	-	-	-	-	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				3	- 7		-	7	-	-	-	-	-	- 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 2771	Continuous Learnin	g Assessment (CLA)		Cumr	native				
	Bloom's Level of Thinki <mark>ng</mark>	expe	experiments exp		of second cycle iments 0%)		Examination eightage)	Final Examination (0% weightage)				
		Theory	Practice	Theory	Practice	Theory	Practice Practice	Theory	Practice			
Level 1	Remember		15%	77.5	15%	A	15%	-	-			
Level 2	Understand		20%	70.75.20	20%	- Table 1	20%	-	-			
Level 3	Apply	-	25%	The second of	25%	1111	25%	-	-			
Level 4	Analyze		25%		25%		25%	-	-			
Level 5	Evaluate		10%		10%	- / /	10%	-	-			
Level 6	Create		5%	- 1777	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	Department	Biotechnology	Data Book / Codes / Standards		Nil

Course L	earning Rationale (CLR):	The purpose of learning this course is to:	1	4		1	ı	rogra	am Ou	tcome	s (PO	)					ogra	
CLR-1:	CLR-1: describe the basics of the fermentation process				1	3	4	5	6	7	8	9	10	11	12		pecifi tcom	
CLR-2:	explain the process of medi	ia formulatio <mark>n and sterili</mark> zation kinetics	0	D D	4		S	1	-	<b>I</b>		ork		9				
CLR-3:	study the basics of reactor of	design an <mark>d its contro</mark> l systems		NIOWIEGIGE Significance	, too	veiopineri	nvestigations k problems	Usage	ъ			am W		inance	р			
CLR-4:	analyze the metabolic stoic	hiometry and energetics of the biochemical process			الرام	do	estig orobl	l Us	r and	∞ <sub>&gt;</sub>		Teal	tion	⊗ ∃	arning			
CLR-5:	CLR-5: illuminate the various types of reactors for suspension and immobilized cell systems			<u> </u>	L 0		t inv olex p	$\vdash$	engineer ety	nability	h.	<u>8</u>	Communication	Project Mgt.	ong Le			
					2 2	utions	duc	dern	erić ety	rol	S	ję.	Ę	ect	Lo	7	7.5	<u>~</u>
Course C	outcomes (CO):	At the end of this course, learners will be able to:		Problem		solu	Con of co	Мос	The	Env	E E	Individual	Col	Proj	Life	PSO-1	PSO	PSO-3
CO-1:	understand the basics of the	e f <mark>ermenta</mark> tion process		1 .	-1	2	ź	T	1	-	-	-	-	-	-	2	2	2
CO-2:	comprehend the process of	m <mark>edia fo</mark> rmulation and sterilization kinetics	1 3	2 2	?	2	2	2		-	=	-	-	-	-	-	2	1
CO-3:	acquire the basics of reactor	o <mark>r design</mark> and its control systems	1 / 20	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		-	3		2	2		_	-	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 Learning Assess  Formative CLA-1 Average of unit test (50%)				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		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:		14	4			Progr	am Oı	itcome	s (PO	))					rogra	
CLR-1:	describe the basics of the	he fermentation pro <mark>cess</mark>		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:					S	Jent	ation	Usage	ъ	. 1				Finan	ning			
CLR-4:			Knowledge	Analysis	velopment	restigations problems		r and	∞ >		Team	ioi	& Fi	ā				
CLR-5:			ering		deve	.≦ ×	T00	engineer stv	ment ability		<u>∞</u>	ommunication	Mgt.	g Le				
	<u>'</u>		7	9	roblem	ign/	comple	Modern		rironme stainab	SS	Individual	חשר	roject	Long	$\Xi$	7-5	5
Course C	Outcomes (CO):	At the end of this course, learners will be able to:	1	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	of the f <mark>ermenta</mark> tion process	187	1 -	13	2		-	7	-		-	-	-	-	2	2	2
CO-2:	comprehend the process of media formulation and sterilization kinetics		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	17.50	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		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%	774 - Cont.	- V					
			mail have	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%	<ul> <li>355 557</li> </ul>	15%	19 1 E	15%	-	-		
Level 2	Understand		20%	100	20%	7A , 4	20%	-	-		
Level 3	Apply	-	25%	7 3 FL 2	25%	111	25%	-	-		
Level 4	Analyze	-	25%	Commence of the Commence of th	25%	COLUMN -	25%	-	-		
Level 5	Evaluate		10%		10%	-	10%	-	-		
Level 6	Create		5%		5%	-	5%	-	-		
	Total		0 %	10	0 %	10	00 %		-		

Course Designers		4 <sup>1</sup>
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	Ρ	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	tcome	s (PO	)					rogra	
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 ıtcom	
CLR-2:	explain the concepts and	explain the concepts and experiments i <mark>n the prepa</mark> ration of linkage map				SC	1				Work		8				
CLR-3:	describe the elements of Genetic Cou <mark>nseling</mark>				velopment	vestigations problems	Usage	ъ					Finance	ning			
CLR-4:	analyze gene transfer and its role in mapping in bacteria				udo	estig	- S	r and	ج × ح ×		Team	tion	& ∃	arni			
CLR-5:	i: differentiate factors that lead to genetic variation in a population		neering	Analysis	(i)		T00	gineer	ment	h.	al &	Sommunication	Mgt.	g Le			
			jed	roblem	/ugi	nduct in complex	Modern	er er	ironn tainal	S	ndividual	nwu	roject	Lon	7-1	)-2	<u>-</u> 3
Course C	outcomes (CO):	A <mark>t the en</mark> d of this course, learners will be able to:	- L		Des	305	Mo	The	Env	Ethic	Indi	Cor	Proj	Life	PSO	PSO	PSO-
CO-1:	analyze the pattern of inheritance of genes and its interaction				2	2	-	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	2	-	-	-	-	-	-	-	-	2	-	-
CO-5:	analyze genetic variations in a population				200	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

у пои

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						
				Assessment (CLA)	Laborina	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
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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	21BTC207T 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	٦
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	dge	,	of	SL			L		ork		99				
CLR-3:	explain the basis and mechanism of protein synthesis and activity	Knowlec	w	nent	gations ems	Usage	ъ	N.		am W		inan	ρ			
CLR-4:	understand the regulatory role of nucleic acids in cell functioning			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.	ong Le			
		_ e	<u> </u>	lign/	ompl	dern	et ei	io,	S	Individua	JIII	Project	힏	7	2 2	
Course C	Outcomes (CO):  At the end of this course, learners will be able to:	En iii	,   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		2	2		-4		-	<u> </u>	-	-	-	-	-	2 2	
CO-3:	relate the expression of DNA at the different levels	3	Variation of the state of the s	1	-			-	-	-	-	-	-	-	3 3	1
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	-	-	-	-	-	-	-	-	-	3 3	

#### 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		ourse	С			F	PROFE	SSIOI	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 /	Standards			T+.				I	Vil							
Course Los	arning Rationale	(CLD): The	purpose of learning this	course is to:	V					roara	m Out	como	· (DO)				I	Dro	ogran	n
CLR-1:			s DNA in p <mark>rokaryotes</mark>	Course is to.			2	- 3	4	5	6	7	8	9	10	11	12	Sp	ecific	•
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CLR-4:			le <mark>ment an</mark> d gene transcri <sub>l</sub>	ots	477	g Kr	naly	/elop	vest	O lo	er a	nt &		ž Te	atior	t. &	Learning			
CLR-5:	know DNA dama	ge in prokaryote	S		277	erin	m A	Jn/dev ons	ct in	P L	engineer ety	nme	l.	ual {	unic	t Mg	ng L			
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Unit-1 - Ge	nomic DNA Isola	tion and An <mark>alys</mark>	sis	William Town	- 61		. 164.0			7									12 F	lour
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	re Analysis or Ge re Analysis Genon									1										
Unit-2 - Pla	smid DNA Isolati		is		7.5														12 F	lour
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	or Plasmid DINA ir ive Analysis of Pla			LEARN L	EAP	<ul> <li>I</li> </ul>	E7	M	-											
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	tal RNA Isolation	and Analysis							4										12 F	lour
Practice:	of Total RNA from	F coli																		
2. Quantitat	ive Analysis of To	tal RNA																		
3. Qualitativ	e Analysis of Tota	al RNA																		

# 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)	( PS)			
	Bloom's Level of Thinking	expe	ge of first cycle riments 0%)	experi	of second cycle iments 0%)		eightage)		amination ightage)
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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:	M.	11	4			Progr	<mark>am O</mark> ı	tcome	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	пg			
CLR-4:	LR-4: describe the structured and unstructured models of microbial system				Analysis	udoli	vestigations problems	l Us	r and	y k		Team	ig.	∞ర	arni			
CLR-5:	CLR-5: discuss modern tools in Bioprocess Engineering			ering	_	n/development	t inv	Modern Tool	engineer sty	ronment tainability		al &	Sommunication	Mgt.	ng Le			
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Course C	Outcomes (CO):	At the end of this course, learners will be able to:	Table 1	Engi	Po	Des	Con	₩ W	Soci	Env S <mark>us</mark>	Ethics	ng.	Sol	Proj	Life	PSO.	PSO-2	PSO-3
CO-1:	understand the ideal and i	non <mark>-ideal sy</mark> stems in bioprocess engineering	15 TX	3	- 3	3	-			-	-	-	-	-	-	1	-	-
CO-2:	gain knowledge on fluid flow and its mixing property			3	2	1	2	-	4	-		-	-	-	-	2	2	-
CO-3:	acquire knowledge in transp <mark>ort phen</mark> omena and scale up studies			3	2	1	1	-	-	7 -	- 6	-	-	-	-	2	2	2
CO-4:	-4: understand structured and Unstructured models			2	1	. 3	1	-	-	-		-	-	-	-	2	-	-
CO-5:	apply modern tools in model <mark>ling of b</mark> ioprocess 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	1.	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 press
		2012

- 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 Learning native ge of unit test 9%)	CL	g Learning .A-2 0%)	Final Ex	mative amination eightage)
		Theory	Practice	Theory	Practice	Theory	Practice
Level 1	Remember	15%	-	15%		15%	-
Level 2	Understand	25%	2.4	20%		25%	-
Level 3	Apply	30%	20 E 10 E 10	25%	A 1-3	30%	-
Level 4	Analyze	30%	N. J. M. 1777	25%		30%	-
Level 5	Evaluate			10%	1-2	4 -	-
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	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	[4] [4] 전 전 등 전 경기를 위해 있는 [4] [4] [4] [4] [4]	
2. Dr. G. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai,	2. Dr.N.Selvaraj, IITG	2. Dr.P.Radh <mark>a, SRMIS</mark> T
ramchand@saksinlife.com		

Course Code 21BTC210L Course Name BIOPROCESS ENGINEERING LABORATORY Category	C PROFESSIONAL CORE	3 C 4 2
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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:	U =	Program Outcomes (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	egp		o	SL		-			ork		99				
CLR-3:	analyze the oxygen mass transfer coefficient and deactivation kinetics	N N		velopment	stigations	Usage	ъ			Μ		Finan	Б			
CLR-4:	evaluate the model parameters in microbial growth	중	alysis	lop	estig		er and	× ×		Team	tion	∞ర	arning			
CLR-5:	discuss the modern tool of programming microbial cultures	ring	. Ana	/deve	t in K	<u>8</u>	engineer stv	meniabilit		<u>8</u>	ommunication	Project Mgt.	g Le			
		inee.	roblem	/ugi	duc	Jern	et et	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	İndi	Co	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				-		-	-		-	-	-	-	2	2	-
CO-5:	learn the modern tool for programming the microbial cultures	1	2	3	-	3	<u> </u>	-		-	-	-	-	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.	Ψ.	3. Shijie Liu,"Bioprocess Engineering Kinetics, Sustainability, and
Resources	<ol><li>Arvind Kumar Bhatt, "Basic Biotechniques for Bioprocess and Bioentrepreneurship" Academic Press, Elsevier, 2023</li></ol>	-	Reactor Design"Elsevier, 2020.

earning Asses	ssment			18 July 1998	124 CT 16				
			well had	Continuous Learning	g Assessment (CLA)				
	Bloom's Level of Thinking	experi	e of first cycle ments %)	exper	of second cycle iments 0%)		eightage)		amination pightage)
	*	Theory	Practice	Theory	Practice	Theory	Practi <mark>ce</mark>	Theory	Practice
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Level 2	Understand		20%	1000	20%	A , 4	20%	-	-
Level 3	Apply	-	25%	73.5-1.2	25%	112 23	25%	-	-
Level 4	Analyze	-	25%	the same of the control of the contr	25%	C1245 -	25%	-	-
Level 5	Evaluate		10%		10%	- 1	10%	-	-
Level 6	Create		5%	).l.	5%	- / /	5%	-	-
	Total	100	) %	10	0 %	10	0 %		-

Course Designers	1594	
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 CARDY 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 MANIPUL 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
		OIENOR.	

Course L	earning Rationale (CLR): The purpose of learning this course is to:	1 1	7		- I	rogra	<mark>m</mark> Ou	tcome	es (PO	)				Pı	ograr	n
CLR-1:	perspective of engineers		2	3	4	5	6	7	8	9	10	11	12		pecifi tcom	
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	. 1		Μ		Finan	ри			
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	ion	& Fi	arnii			
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	nabilit		dual &	Communication	ct Mgt.	ong Le	_	2	e
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	-		Z-	-		-	-	-	-	-	2	-
CO-2:	design and conduct experiments involving genetic manipulation		5.4	2	- 1		2	-	1	-	-	-	-	-	-	3
CO-3:	illustrate the steps involved in the production of biopharmaceuticals in microbial and mammalian cell system	s 2	美石	4	-	- 1	H.	-	2	-	-	-	1	-	-	3
CO-4:	apply modern biotechnology in the different areas like medicine, microbes, environment, and agriculture	3	-	-	-		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
Resources
. 100001.000

- 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			of the Park Inc.	A Warning					
	Blo <mark>om's</mark> Level of <mark>Thinkin</mark> g	VIS7	CLA-1 Averag	Continuous Learnir ative ge of unit test %)	Summative Final Examination (40% weightage)					
	-		Theory	Practice	Theory	Practice	Theory	Practice		
Level 1	Remember		15%	The first section of	P 12 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	15%	15%	-		
Level 2	Understand		25%		1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	20%	25%	-		
Level 3	Apply		30%		A service and	25%	30%	-		
Level 4	Analyze		30%	. III	-	25%	30%	-		
Level 5	Evaluate	1	-	- 11.7	-	10%	-	-		
Level 6	Create			- 1	-	5%		-		
	Total	1	100	)%	100	0%	100	0 %		

Course Designers	The state of the 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	A LILITED A	
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	N	Co- requisite Courses	Nil Progress Course		Nil
Course Offe	ring 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	)				_	ograr	
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 of immune system functioning, ways to strengthen immune system and how human body is designed and protected to fight against various pathogens</mark>	ering	m Analysis	g g	act inves	Tool	engineer	Environment 8		∞ ~	Sommunication	Project Mgt. &	ong Lear		0.1	က
Course (	Outcomes (CO):  At the end of this course, learners will be able to:	Engine	Problem	Design/ solution	Condi	Modern	The el	Enviro	Ethics	Individual	Comn	Projec	Life Lo	PSO-	PS0-2	PSO-(
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	-	-	-	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	<b>C</b>	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 15%)	CLA-2-I	Practice %)	Summative Final Examination (40% weightage)				
		Theory	Practice	Theory	Practice	Theory	Practice			
Level 1	Remember	15%		//	15%	15%	-			
Level 2	Understand	25%	11 - JUST	-	20%	25%	-			
Level 3	Apply	30%	- 1	-	25%	30%	-			
Level 4	Analyze	30%	- 1/1/1/	-	25%	30%	-			
Level 5	Evaluate	1 7 -	- //4///	-	10%	-	-			
Level 6	Create	·			<i>5</i> %	-	-			
	Total	- 10	00 %	100	)%	10	0 %			

Course Designers	VALUE OF PURPLE SALES	
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
1. Dr. C. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai,	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	mative							
	Bloom's Level of Thinking	CLA-1 Avera	native ge of unit test 0%)	CL	g Learning .A-2 0%)	Final Examination (40% weightage)					
		Theory	Practice	Theory	Practice	Theory	Practice				
Level 1	Remember	15%	A 420 A 12	15%	- A-	15%	-				
Level 2	Understand	25%	F-10 (1) (4)	20%	( ) ( )	25%	-				
Level 3	Apply	30%	Sec. 27.0	25%		30%	-				
Level 4	Analyze	30%		25%	- C- J	30%	-				
Level 5	Evaluate		A THE WATER A TO	10%		-	-				
Level 6	Create	-	Charles Mary and	5%			-				
	To <mark>tal</mark>	10	0 %	10	0 %	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, subbu@iitm.ac.in	1. Dr. Priya Swaminathan, SRMIST
ramchand@saksinlife.com	Martin Bridge Back To The Alleran	
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	, C
Code	210103041	Name	ANIMAL 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

															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	Knowledge		of	SL	1				ork		9				
CLR-3:	CLR-3: inculcate the understanding of cell culture technique and production of valuable products from them					stigations	age	ъ			N W		Finan	Б			
CLR-4:					elopment	estig	l Us	er and	± >		Team	ţion	∞ర	arning			
CLR-5:					deve	t inv	18 18	enginee	ironment tainability		ual &	ommunication	Project Mgt.	ig Le			
			ineering	oblem	ign/	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:	CO-2: impart an understanding about Embryo transfer, fertilization methods and animal production				3	-			-		-	-	-	-	-	3	3
CO-3:	CO-3: provide knowledge about different culture techniques, Characterization of cell lines and in vitro testing of drugs			3	2	-	-	-	-		-	-	-	-	3	3	-
CO-4:	CO-4: provide knowledge about improvement of animals to increase the yield and quality of animal products				-5	3			-	-	-	-	-	-	3	3	-
CO-5:	0-5: 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

э поиг

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%	200	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
ramchand@saksinlife.com	Mark the Park of the Control of	
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:	$I^{\dagger}$	4 5		Ī	Progr	<mark>am</mark> Ou	tcome	s (PO	)					ogran	
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		ent of	SI					ork		9				
CLR-3:					stigations	age	ъ			Μ		Finan	б			
CLR-4:	distinguish between cell viability and cell cytotoxicity	Knowledge	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 aty	onment sinability		<u>∞</u>	ommunication	Project Mgt.	g Le			
		Enginee	roblem	sign/d	duc	Jern	et e	viron staina	S	ndividual	nuu	ect	Lon		2-0Sc	<u>ب</u>
Course (	urse Outcomes (CO):  At the end of this course, learners will be able to:				55	Mod	The	Env Sus	Ethics	İpu	Sol	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 8	-	3	2	- 1		-		-	-	-	-	-	-	3
CO-3:	acquire skills to perform fluore <mark>scent st</mark> aining procedures to visualize cellular content	15 NF	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		3 /	- 50 L 50%	175 × 6 m.						
			mel had	Continuous Learnin		)	4				
	Bloom's Level of Thinking	Level of Thinking (30%)			of second cycle iments 0%)		Examination reightage)	Final Examination (0% weightage)			
	9	Theory	Practice	Theory	Practice	Theory	Practi <mark>ce</mark>	Theory	Practice		
Level 1	Remember		15%	<ul> <li>355 557</li> </ul>	15%	79 V S	15%	-	-		
Level 2	Understand		20%	100	20%	A , 4	20%	-	-		
Level 3	Apply	_	25%	7 3 FL 2	25%	100	25%	-	-		
Level 4	Analyze	-	25%	Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the Commence of the 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	17,74	
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
Dr. G. N. Ramchand, Saksin Life sciences Pvt Ltd, Chennai, ramchand@saksinlife.com	1. 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 DIOTECHNOLOGY	Course	C	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:		7			ı I	rogra	am Ou	tcome	s (PO	))					ograi	
CLR-1:	understand the genome org	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		of	SL		- i			ork		9				
CLR-3:	employ different methods for	or the de <mark>velopment</mark> of transgenic plants		wlec	S	velopment	stigations	age	pu			M ≪		Finan	Б			
CLR-4:	interpret the mechanisms for	or the p <mark>lant to co</mark> pe with biotic and abiotic stresses		Kno	Analysis	lopi	estig probl	l Us	er an	۲ × ×	A.	Team	ţio	∞ F	arning			
CLR-5:	apply the classical and mod	dern <mark>plant bre</mark> eding techniques for crop improvements		ring		(1)	t inv	T 90	inginee ty	ment		<u>रू</u>	Communication	Project Mgt.	g Le			l
	•			inee	roblem	sign/de	duc	lern	et c	ron	g	Individual	nur	ect	Long	7	)-2	
Course C	Outcomes (CO):	At the end of this course, learners will be able to:	12.72	Engir	Prof	Des	Cou	Moo	The	Envi	Ethic	lndi	Sol	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	-	3	-		7	3	-	-	-	-	-	-	2	-
CO-2:	demonstrate the mechanish	m and role of plant tissue culture for mass multiplications	/	3	2	3	1			-	-	-	-	-	-	-	3	-
CO-3:	establish the various metho	o <mark>ds of ge</mark> netic manipulation in plants	1.2.8	3	2	-74		3		-		-	-	-	-	-	3	-
CO-4:	discuss the molecular aspe	cts of plant adaptability to various stresses		-3		2	-	-	_	3		-	-	-	-	-	-	3
CO-5:	apply the significance of pla	a <mark>nt breed</mark> ing and genetic manipulations of plants for economic importance		3		11	-	3		3	-	-	-	-	-	-	3	-

#### 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 9%)	CLA	g Learnin <mark>g</mark> A-2 – 0%)	Final Examination (40% weightage)			
	/ /	Theory	Practice	Theory	Practice	Theory	Practice		
Level 1	Remember	15%	100	15%	-	15%	-		
Level 2	Understand	25%	20 E 10 E 10	20%		25%	-		
Level 3	Apply	30%	\$ 3.50 TW	25%		30%	-		
Level 4	Analyze	30%		25%	2.5	30%	-		
Level 5	Evaluate		8, CH (842) C + C	10%		-	-		
Level 6	Create	-	Called States and	5%			-		
	Total -	100	0 %	10	0 %	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	1. 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 21DTC4011	Course	PLANT BIOTECHNOLOGY LABORATORY	Course	C	PROFESSIONAL CORE	L T	Р	C	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	ng Department	Biotechnology	Data Book / Codes / Standards	Nil

Course L	earning Rationale (CLR):	The purpose of learning this course is to:	- 4	<u> </u>			Progr	<mark>am</mark> Ou	itcome	s (PC	))				_	rograr	
CLR-1:	relate the growth and develo	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	K	Analysis	lopi	estigal probler		er and	∞ >	l.	Teal	tion	∞ర	arning			
CLR-5:	apply the classical techniqu	es fo <mark>r crop im</mark> provement	Engineering	Ang	sign/development	ě i	_	enginee etv	ment		<u>8</u>	ommunication	Mgt.	g Le			
	<u> </u>		inee	Problem	)ugi	. o =	Modern	erc et<		જ	ndividua	nur	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	Col	Proj	Life	PS0-1	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 technic	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	ĘΖ	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							
			5.47.5	Continuous Learning		20 20 21 14 1					
	Bloom's Level of Thinking	exper	ge of first cycle riments 0%)	CLA-2 Average experi (30	ments		Examination veightage)	Final Examination (0% weightage)			
		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	A - A	25%	- 1/ /	25%	-/ A	2 <mark>5%</mark>	-	-		
Level 5	Evaluate	-	10%	- 1730	10%	7 ,	10%	-	-		
Level 6	Create	- t	5%	- 1/2//	5%	1-1	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

	earning Rationale (CLR): The purpose of learning this course is to:						_				_						D.	'oaro	
Course L	earning Rationale (CLR):	The purpose of learn	ing this course is to:					1	rogra	am Oı	utcome	es (PC	))					rograi 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	duct from so <mark>lid –liquid</mark> pha	se		dge	1	of	SL			N.		ork		9				
CLR-3:				owlec	S	elopment	stigations	Usage	ъ			N W		Finan	ning			ł	
CLR-4:	learn the methods of purific	ation o <mark>f produc</mark> ts			조	Analysis	ndol	estig	-	r and	م × «		Team	ţio	& ∃	earni			
CLR-5:	learn the methods of polish	ing a <mark>nd formu</mark> lation of pro	ducts for packaging	4	ering		deve	t inv	T00	engineer	ment		<u>ھ</u>	ommunication	Project Mgt.	Ľ			
					(I)	roblem	ign/	ag di	ern	eng 4		S	ndividual	חוו	ect	Long	7	)-2	<u>ښ</u>
Course O	outcomes (CO):	At the end of this co	rse, learners will be able to:	, rait	Engin	Prof	Des	o So	Moc	The S	Env	Ethic	lpd	Con	Proj	Life	PS0-1	PSO-2	PSO-3
CO-1:	categories the products into	o <mark>various </mark> sectors	5 2 2 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		1	2	1	-	-	7	-	-	-	-	-	-	1	2	1
CO-2:	identify the unit operation for	o <mark>r se</mark> paration		7	2	3	1	-	1		-		-	-	-	-	2	2	1
CO-3:	adapt the best methods of	i <mark>solation</mark> of products		H.S	2	2	2	<i>3</i> -	- 1		-		-	-	-	-	2	2	1
CO-4:	identify the sophisticated ed	q <mark>uipmen</mark> t for purification		7	-2	3	2	-	-	-	-	-	-	-	-	-	2	2	2
CO-5:	know the polishing and forr	m <mark>ulation o</mark> f the products			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

rning Assessme		1777	Continuous Learnin	g Assessment (CLA)		Cum	mativa
	Blo <mark>om's</mark> Level of <mark>Thinkin</mark> g	Form CLA-1 Averag (45	ge of unit test		-Practice 5%)	Final Ex	mative amination eightage)
		Theory	Practice	Theory	Practice	Theory	Practice
Level 1	Remember	15%	MY - N/17/	-	15%	15%	-
Level 2	Understand	25%	- 1	-	20%	25%	-
Level 3	Apply	30%	- 1/1/1	-	25%	30%	-
Level 4	Analyze	30%	- 1/2%	<u>-</u>	25%	30%	-
Level 5	Evaluate				10%	-	-
Level 6	Create	<-		7-	5%	-	-
	Total	/ 100	1%	4 13 10	00 %	10	0 %

Course Designers	V Links	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	1. 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	O
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			0.	licon	25
CLR-3:			age that could be caused to the environment		ledc		ant 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	<del>560  </del> 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	-	"	-	-	-	-	-	-	<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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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>	Summative Final Examination (40% weightage)				
		Theory	Practice	Theory	Practice	Theory	Practice			
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Level 2	Understand	25%		20%		25%	-			
Level 3	Apply	30%	3	25%		30%	-			
Level 4	Analyze	30%	-	25%		30%	-			
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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, subbu@iitm.ac.in	1. Dr. DVL Sara <mark>dha, SRM</mark> IST
ramchand@saksinlife.com		
2. Dr. Karthik Periyasamy, Scientist, Biocon,	2. Prof. R. B Narayanan Anna University, Chennai, arbeen 09@gmail.com	2. Dr Lilly M Sale <mark>ena. SR</mark> MIST
karthik.periyasamy@biocon.com		

# ACADEMIC CURRICULA

# UNDERGRADUATE/ INTEGRATED POST GRADUATE DEGREE PROGRAMMES

(With exit option of Diploma)

(Choice Based Flexible Credit System)

Regulations 2021

Volume - 8D
(Syllabi for Biotechnology (Computational Biology)
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 Code	21BTE208T	Course Name	SEQUENCE ANALYSIS AND MOLECU	LAR PHYLOGENY Course Category	PROFESSIONAL ELECTIVE	L T	P 0	3	
Pre-requisi	ite		Co- requisite	Progres	sive				
Courses		Nil	Courses	Nil Cours		Nil			

Data Book / Codes / Standards

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Course L	earning Rationale (CLR)	: The purpose of learning this course is to:		14	7	72	Progi	ram O	utcom	es (PC	<b>D</b> )					rogra	
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CLR-2:	Analyze sequence patte	erns and <mark>profiles.</mark>	ge		of	SL	1				Work		8				
CLR-3: Apply the concept of the evolutionary tree.		Knowledge	S	velopment	stigations	Usage	ъ			≽		inan	Б				
CLR-4:	Understand the various	methods of phylogenetic tree construction.	Α̈́	nalysis	ldol	estig		r and	∞ - >		Team	ig.	∞ ∃	aming			
CLR-5:	Interpret the application	of <mark>Biomole</mark> cular sequence analysis and tools.	Engineering	roblem Ana	ydeve	. .≦ x	ŏ	engineer	ironment tainability		ual &	mmunication	t Mgt.	ong Le			
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CO-2:	Infer the different kinds	of sequence patterns and profiles	2	3	生元		3	- 4		-		-	-	-	2	-	2
CO-3:	Explore the phylogeneti	<mark>c relatio</mark> nship between species	3	3	1-0	2	-		-	-	-	-	-	2	2	-	2
CO-4:	Formulate an approach	to analyze phylogenetic inference and methods	2	3	dia-	2	-	-,7	-	-		-	-	-	-	2	2
CO-5:	Understand and implem	n <mark>ent biom</mark> olecular sequence analysis and tools.	2		اينا	2	3		-	_	<b>.</b>	-	-	2	-	2	2

Unit-I: Biological Databases 9 hour

Biological databases - Classification: Primary and secondary databases - NCBI, EMBL and DDBJ - PIR: UniProtKB - Structure database: PDB, SCOP and CATH - Specialized database- KEGG, TCGA, PUBCHEM, Bibliographic database- File formats.

#### Unit-II: Sequence Alignment, Patterns and Profiles

9 hours

Nil

Sequence alignment classification: Pairwise alignment and Multiple sequence alignment (MSA) -Scoring matrices Illustration — PAM and BLOSUM - Needleman Wunsch algorithm - Smith-Waterman algorithm — Database searching algorithm- BLAST - Methods and Tools for MSA - Progressive alignment - Domain & Motif - Family — Superfamily-Profiles-Patterns.

#### Unit-III: Concept of Evolutionary Tree

**Course Offering Department** 

Biotechnology 8 1

9 hours

Terminology of phylogenetics, definition of Dendrogram, Cladograms, & Phylograms, Types of phylogenetic trees (rooted vs. unrooted trees), gene tree & Species tree, Newick format of tree representation, True tree & Inferred tree. Phylogenetic Inference, Distance methods – Fitch Margoiasch, UPGMA, WPGMA, Neighbor-Joining method, Character-based method - Maximum Parsimony methods, Maximum Likelihood method.

#### Unit-IV: Phylogenetic Inference – Evaluation methods

9 hours

Bootstrapping, Randomized and jack-knifing methods- Molecular Clock- Rate heterogeneity- LBA problem- Phylogenomics- Supertree & Super matrix method-Phylogenetic analysis software: PHYLIP, MEGA, PhyML, and MrBayes.

#### Unit-V: Biomolecular Sequence Analysis and Tools

9 hour

Gene prediction in prokaryotes and eukaryotes-ORF finder- primer design- secondary structure prediction of RNA and proteins - Chou Fasman - GOR - TMHMM, Promoter and regulatory element prediction - RNA structure prediction.

		1.	David W. Mount Bioinformatics: Sequence and Genome Analysis Cold Spring	3.	Molecular Evolution and Phylogenetics Masatoshi Nei and Sudhir Kumar, Oxford University
Lea	rning		Harbor Laboratory Press, 2004	-	Press, 2000.
Res	ources	2.	Lesk, Arthur. Introduction to bioinformatics. Oxford University Press, 2019.	4.	Dandekar, Thomas, and Meik Kunz. Bioinformatics: An Introductory Textbook. Springer
				3.3	Nature, 2023.

earning Assessi	ment		Continuous Learnin	g Assessment (CLA)	$\theta$	0			
	Bloom's Level of T <mark>hinking</mark>	CLA-1 Avera	native ge of unit test 0%)	Life-Long CL/ (10		Summative Final Examination (40% weightage)			
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Level 2	Understand	25%		25%		25%	-		
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Co	ourse Designers	ساعيها	1.07		-
Ex	perts from Industry	Ex	perts from Higher Technical Institutions	Inte	r <mark>nal Exp</mark> erts
1.	P. V. Prasad, Founder, Helix Bios <mark>ciences</mark>	1.	Dr.Sanjeev Kumar Singh, Department of Bioinformatics, Alagappa	1.	Dr. S. K. M. Habeeb, SRMIST
			University, Tamil Nadu, India. sanjeevslab@gmail.com		
2.	Mr. Sudheendra Rao, Director, DataLore Labs, Bengaluru	2.	Dr. K. Sekar, Department of Computational and Data Sciences, Indian	2.	Dr. S. Shobana, SRMIST
			Institute of Science, Bangalore, India. Sekar@iisc.ac.in		

Course Code 21	BTE319T Course Name	ADVANCED GEI	NOMICS	Course E	PROFESSIONAL ELECTIVE	L T P C 3 0 0 3
Pre-requisite Courses	Nil	Co- requisite Courses	Nil	Progressive Courses	Nil	
Course Offerin	ng Department	Riotech <mark>nology</mark>	Data Book / Codes / Standa	rds T	Nil	

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Course L	earning Rationale (CLR): The pur	pos <mark>e of learnin</mark> g this course is to:		-41	$\mathcal{I}$	/ [	Progr	am Oı	utcome	es (PC	D)					rogram	
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CLR-3:	Create an interest about approach	es of studying genomics	Jowled	w	Jent	estigations	sage	ъ		• 1			inan	Бu			
CLR-4:	Demonstrate the methods used for	r performing genome annotation	Αĥ	nalysis	velopment of	estig		er and	∞ 		Team	io	& Fi	amii			
CLR-5:	Apply genomics in various <mark>fields</mark>		ining	Αng	/deve	ex p	T00	nginee	ment ability		<u>ळ</u>	mmunication	Mgt.	ıg Le			
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CO-1:	Illustrate knowledge about genome	e structure and human genome project	. J-	2	2	7	2			-	12-	-	-	-	-	-	2
CO-2:	Discuss about different DNA seque	encing methods	3	3	1	-	2			-	-	-	-	-	-	-	2
CO-3:	Outline the different approaches in	nvolved in studying genomics and transcriptomics	111-2	2	20	2	-	3	-	-	15-	-	-	-	2	-	-
CO-4:	Demonstrate applicability of metho	ods involved in gene prediction and genome annotation	2	, -	120	-51	2	2	-	-	12	-	-	-	-	-	3
CO-5:	Explain various applications of ger	nomics in different fields	2	2	1 1-1	-	2		-	-	-	-	-	-	-	-	3

#### Unit-I: Genome Organization and Structure

9 Hour

Organization of nuclear DNA in eukaryotes; Genome structure – Genome sizes – Genome sequence complexity – Human genome project - Goals and outcomes of human genome project - Human genome - significance of other genome projects - Genome assembly

#### Unit-II: DNA Sequencing Strategies

9 Hour

History and Evolution of DNA sequencing strategies; Principles, methods, advantages, disadvantages and comparison of Sanger sequencing, Illumina sequencing, Ion Torrent sequencing, PacBio sequencing, Nanopore sequencing

#### Unit-III: Genomic Approaches

9 Hour

Principles, methods, advantages, disadvantages and applications - Whole genome sequencing – whole transcriptome sequencing – Small RNA sequencing – ChipSeq – Exome Sequencing – Amplicon sequencing – Metagenomics – Single Cell Genomics - Microarray

#### Unit-IV: Genome Annotation

9 Hour

Bioinformatics tools for genome annotation - Detecting open reading frames – principles, methods and tools for gene prediction; Homology based approach to find genes – Non-coding RNA, Extra coding DNA, function identification of new gene, fusion genes, gene ontology, non-homology-based approach to find genes- genome annotation tools - Introduction to Bioconductor

#### Unit-V: Applications of Genomics

9 Hour

Case studies depicting applications in Pharmacogenomics and Personalized Medicine – GWAS, MWAS and mGWAS Studies in agriculture – Single-Cell RNA sequencing application - Application of NGS in Human diseases: Cancer and neurodegenerative diseases.

Learning	1.	Sandy B. Primrose and Richard Twyman, Principles of Genome Analysis and
Resources		Genomics, 3rd Edition, Wiley, 2009.
	2.	Stuart M. Brown, Next-Generation DNA Sequencing Informatics, 2nd Edition, Cold
		0 1 11 1 1 1 1 0 1 1 0045

- Shademan B, Biray Avci C, Nikanfar M, Nourazarian A. Application of Next-Generation Sequencing in Neurodegenerative Diseases: Opportunities and Challenges. Neuromolecular Med. 2021 Jun;23(2):225-235. Wang Y, et al. GWAS, MWAS and mGWAS provide insights into precision
- Spring Harbor Laboratory Press, New York. 2015 3. David W. Mount., "Bioinformatics: Sequence and Genome Analysis", 2nd Edition, agriculture based on genotype-dependent microbial effects in foxtail millet. Nat Commun. 2022 Oct 7;13(1):5913. Cold Spring Harbor Laboratory Press, New York. 2004

Learning Assessr	ment								
	Bloo <mark>m's</mark> Level of <mark>Thinking</mark>	CLA-1 Ave	Continuous Learnin rmative rage of unit test 50%)	CL	Learning A-2 0%)	Summative Final Examination (40% weightage)			
		Theory	Practice	Theory	Practice	Theory	Practice		
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Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
Dr. V.L.Ramprasad, MedGenome Labs Ltd, Bengaluru, ramprasadv@medgenome.com	<ol> <li>Dr. Ranjith Kumavath, Pondicherry University, Pondicherry, rnkumavath@gmail.com</li> </ol>	1. Dr <mark>.S. K. M.</mark> Habeeb, SRMIST
Dr. Sanjay Singh, Eurofins Genomics India Pvt Ltd.,     Bengaluru, sanjay.singh@xoin.eurofinasia.com	Dr. S. Mahalingam, Indian Institute of Technology Madras,     Chennai, mahalingam@iitm.ac.in	2. Dr. R. Satish, SRMIST

Course Code	21BTE320T	Course Name	ALGORITHMS IN COMPU	FATIONAL BIOLOGY	Course E	PROFESSIONAL ELECTIVE	3	T 0	P 0	3
Pre-requis		Nil	Co- requisite Courses	Nil	Progressive Courses	Nil				
Course O	ffering Departme	ent	Biotechnology Biotechnology	Data Book / Codes / Stand	ards	Nil				

Course Le	earning Rationale (CLR): The purpose of learning this course is to:			$\cup A$	7 34	Progr	am O	utcome	es (PC	D)					ogra	
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CO-2:	implement various sequ <mark>ence ali</mark> gnment algorithms	3	3	3			- "		-	-	-	-	-	2	-	2
CO-3:	perform genome assembly and regulatory motifs	3	3	2	1	-	-	-	-	-	-	-	-	2	-	2
CO-4:	analyze high throughpu <mark>t data an</mark> alysis	1	3	3	2	-	-	-	-	12	-	-	-	2	-	2
CO-5:	integrate biological interaction with network analysis		3	3	1	3-	-,4	-	-	-	-	-	-	-	2	2

#### Unit-I: Introduction to Algorithms

9 Hour

Background - algorithms - complexity - brute force, recursive techniques - molecular biology primer - different data types - anecdotes of algorithms and data structures - breakthroughs - Probability - Descriptive statistics - Hypothesis testing - Heuristics algorithm

#### Unit-II: Sequence Alignment Algorithms

9 Hour

Dynamic programming -Sequence alignment methods - sequence database search - suffix trees - space efficient alignment using divide & conquer - addressing challenges with pseudoknots, Principal component analysis- Poisson distribution - Regression analysis- Hidden Markov Model - Neural networks.

#### Unit-III: Genome Assembly and Regulatory Motif

9 Hour

Genome assembly - Overlap layout consensus - Burrow wheeler and De Bruijn graph assembly - greedy algorithm types - for shortest common superstring - Regulatory motif discovery - brute force - greedy algorithms - the Pattern branching algorithm - Genetic Algorithm - Bayesian statistics.

#### Unit-IV: High-Throughput Data Analysis

9 Hour

Analysis of bulk and single cell expression - data normalization - dimension reduction - K-means clustering - bi-clustering - BiMax, a divide and conquer algorithm - differential expression analysis - pseudo time analysis using minimum spanning tree - Bootstrapping - Survival analysis.

#### Unit-V: Network Analysis Algorithm

9 Hour

Network analysis: clique finding - regulatory networks - co-expression networks - Scale free networks - Small world network - Density based clustering - Tools for network analysis - Mcode algorithm - shortest path algorithm.

Learning Resources	1. 2.	Pevzner, P., Compeau, P. Bioinformatics Algorithms: An Active Learning Approach. United States: Active Learning Publishers, 2018. Gusfield, D. Algorithms on Strings, Trees, and Sequences: Computer Science and Computational Biology. United States: Cambridge University Press, 1997	3.Baum, J. O., Zvelebil, M., Zvelebil, M. J. Understanding Bioinformatics. United Kingdom: Garland Science, 2008
		CUENC	

Learning Assessn	ment		POSTER A	ALL ALL				
			<ul> <li>Continuous Learnin</li> </ul>	Summative				
	Bloom's Level of Think <mark>ing</mark>	CLA-1 Avera	ative ge of unit test %)	CL	Learning A-2 0%)	Final Examination (40% weightage)		
	/67	Theory	Practice	Theory	Practice	Theory	Practice	
Level 1	Remember	15%		15%	V-S	15%	-	
Level 2	Understand	25%	A 10 Com 17	25%		25%	-	
Level 3	Apply	30%		30%		30%	-	
Level 4	Analyze	20%		20%		20%	-	
Level 5	Evaluate	10%	<b>自己的人人</b> (1)	10%		10%	-	
Level 6	Create		Start Brown Block	St. 1 1 1 1 1 1 1 1 1 1	17 July 19 19 19 19 19 19 19 19 19 19 19 19 19		=	
	<b>Total</b>	100	) %	100	0 %	100	0 %	

Course Designers		
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
Dr. V.L.Ramprasad, MedGenome Labs Ltd, Bengaluru. ramprasadv@medgenome.com	Dr Shandar Ahamad, Jawaharlal Nehru University, New Delhi.     shandar@jnu.ac.in	1. Dr. TAnand, SRMIST
Dr. N. Mathan, Allianz Biosciences (P) Ltd, Puducherry.     nm@abpl.co.in	2. Dr. Michael Gromiha,Indian Institute of Technology Madras, Chennai. gromiha@iitm.ac.in	2. Dr. V. Sivaram <mark>akrishnan</mark> , SRMIST

Course Code	21BTE321T	Course Name	PYTHON FOR BIOIN	IFORMATICS	Course E	PROFESSIONAL ELECTIVE	L T P 3 0 0	3
Pre-requis		Nil	Co- requisite Courses	Nil	Progressive Courses	Nil		
Course O	ffering Departme	ent	Biotech <mark>nology</mark>	Data Book / Codes / Sta	andards	Nil		
				CALLEY V	N. 18 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1			

Course L	earning Rationale (CLR): The purpos <mark>e of learnin</mark> g this c	ourse is to:	Program Outcomes (PO)					Program Specific									
CLR-1:	Understand the basic of python coding	U'	1	2	3	4	5	6	7	8	9	10	11	12		tcom	
CLR-2:	Summarize about Python libraries for data analysis	with middle	work Work														
CLR-3:	CLR-2: Summarize about Python libraries for data analysis  CLR-3: Exercise biopython modules to analzye the biological sequence  CLR-4: Analyze sequence annotation objects				nent	ation	sage	ъ					Finance	ĝ			
CLR-4:	R-3: Exercise biopython modules to analyze the biological sequence  R-4: Analyze sequence annotation objects  R-5: Implement outhor in molecular highers  R-6: Implement outhor in molecular highers  R-7: Summanze about Python in molecular highers  R-7: Summanze about Python in molecular highers  R-7: Summanze about Python in molecular highers  R-7: Summanze about Python in molecular highers  R-7: Summanze about Python in molecular highers  R-7: Analyze sequence annotation objects  R-8: Implement outhor in molecular highers  R-8: Implement outhor																
CLR-5:	CLR-5: Implement python in molecular biology		eering	⋖	Jn/deve	anduct inve	ρ	engineer stv	ironment tainability	S	ndividual &	nmunication	Project Mgt.	ong Le	<u>-</u>	-2	က္
Course C	Outcomes (CO):  At the end of this course, lear	ners will be able to:	Engine	Problem	Designation	Sonc	Modern	The e	Envir Susta	Ethics	ndiv	Com	Proje	∏e [	PSO-1	PS0-2	PSO
CO-1:	Learn the basics of Python coding necessary for biological	problem analysis	2	2		7-	-		=-	-	-	-	-	2	-	-	2
CO-2:			2	40	2	2	-	- "		-	-	-	-	-	-	-	2
CO-3:			2	2	2	2	-	-	-	-	-	-	-	-	-	-	2
CO-4:				2	2	2	-	-	-	-	12	-	-	-	-	-	2
CO-5:	Application of Bio-python coding practices for biological sc	ences —	3	3	3	3	-	-,-	-	-	-	-	-	-	-	-	3

Unit-I: Basics of Python Programming	9 Hour
Introduction to python- Keywords and identifiers - Datatypes - Operators - Python statements - if - Elif and Else For and while loop - Python functions: Built in and user defined as a contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the contraction of the	ned Functions - Lists and Tuple.
Unit-II: Python for Data Analysis	9 Hour
Introduction to Python library for data analysis - Numpy-Arrays - Indexing and selection - Operation - Exercise - Pandas - Series - DataFrames - Conditional filtering	
Unit-III: Python for Sequence Objects	9 Hour
Biopython - working with Python for Genomic data analysis - String operators - Indexing and slicing biological sequences - Sequences as objects: Seq object into strings Unknown sequence contents, partially defined sequence contents-Mutableseq Objects	s, -Comparing sequence objects:
Unit-IV: Python for Sequence Annotation Objects	9 Hour
Sequence annotation objects: Sequence record objects from FASTA and Genbank-Sequence feature objects-Parsing FASTA Sequence format - Parsing Genbank sequence	ce file format. Parsing or reading
sequence alignments: Multiple and pairwise sequence alignment	
Unit-V: Application of Python Modules in Molecular Biology	9 Hour
Introduction to Biopython library-Python code-Identify the GC Content-Counting nucleotides-coding nucleotide sequences-reverse complements-Transcription-Translation	-Translation table

Learning	Jitendra Kumar Verma, Prashant Johri, Sudip Paul. Application of	3. Online Sources: https://wiki.python.org/moin/BeginnersGuide/Programmers.
Resources	Machine learning., Springer Singapore, 2020	4. http://biopython.org/DIST/docs/tutorial/Tutorial-1.48.pdf
	2. Dheepak Khemani. A First course in Artificial Intelligence., McGraw Hill	
	Education Pvt Ltd., 2014	

		S. Tree	SCIEN	$CE_{A\lambda}$					
Learning Assessm	nent								
	Bloom's Level of T <mark>hinking</mark>	CLA-1 Avera	Continuous Learning ative ge of unit test %)	Life-Long CL	g Learning .A-2 0%)	Final Exa	Summative Final Examination (40% weightage)		
		Theory	Practice	Theory	Practice	Theory	Practice		
Level 1	Remember	15%	等级 经销售工业	15%		15%	-		
Level 2	Understand	25%		20%		25%	-		
Level 3	Apply	30%	1.20 1.30	25%		30%	-		
Level 4	Analyze	20%	Start Branch	25%	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	20%	-		
Level 5	Evaluate	10%		10%		10%	-		
Level 6	Create	1.77 (6) (1.10)	51 797 E.S.	5%			-		
	<b>Total</b>	100	)%	10	0 %	100	) %		

Course Designers		
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
Dr. C. Ramakrishnan, Principal Scientist, Molecular Design Division Aroniter Co., Seoul, South Korea ramakrishnan@arontier.co	1. Dr Shandar Ahamad, Jawaharlal Nehru University, New Delhi. shandar@jnu.ac.in	1. Dr. Thirumurthy Madhavan, SRMIST
2. Mr. Sudheendra Rao, Director, DataLore Labs, Bengaluru	Dr. Balachandran Manavalan, Department of Integrative Biotechnology, College of Biotechnology and Bioengineering, Sungkyunkwan University, Suwon 16419, Gyeonggi-do, Republic of Korea, bala2022@skku.edu	2. Dr. Habeeb. S. K. M, SRMIST

Course Code	21BTE322T	Course Name	DATA SCIENCES I	N BIOLOGY	Course E E	PROFESSIONAL ELECTIVE	L T P C 3 0 0 3
Pre-requisite Courses	е	Nil	Co- requisite Courses	Nil	Progressive Courses	Nil	
Course Offe	ering Departm	ent	Biotechnology	Data Book / Codes / S	tandards	Nil	
				a beautiful and the first of	5 S. A. E. L		

Course L	earning Rationale (CLR): The purpos <mark>e of learnin</mark> g this course is to:	Program Outcomes (PO)					Program Specific									
CLR-1:	summarize the biological file formats	1	1 2 3 4 5 6 7 8 9 10 11 12					12	Outcomes							
CLR-2:	acquire knowledge about different data analysis tools	ge	edge edge edge													
CLR-3:	perform data analysis using mathematical and statistical approaches	wed					Б									
CLR-4:	analyzing the graphical data	Αno	nalysis	elopment	estig probl		r and	م <sub>&gt;</sub>		Team	ţį	∞ర	ami			
CLR-5:	integration of data science in biology	aring	. ⋖	deve	tiny	70 20	jineer	ment ability		ual &	ommunication	Mgt.	g Le			
			oblem	/ugi	op de	ern	eng eng	ron	S	vidu	E	oject	Long	0-1	0-2	5
Course C	outcomes (CO):  At the end of this course, learners will be able to:	Engi	Pg	Desi	Con	Mod	The	Envi	Ethi	Indi	Com	Proje	_ie	PSC	PSC	5
CO-1:	identify the need for biological data formats	3	2		15.5	-		_	-	-	-	-	-	2		-
CO-2:	outline the various data analysis tools in biological sciences	3	3	1.0		-	- "		-	-	-	-	-	2		- ]
CO-3:	apply the big data analysis approach to biological data	77	70-1	2	1	3	- 1	-	-	-	-	-	-	2		-
CO-4:	analyze biological data using graph theory	-	-	2	3	-	-	-	-		-	-	-	-	- 1	2
CO-5:	application of data science in biology	3	-	1 -	2	-	- 3	-	-		-	-	-	-	- 2	2

#### Unit-I: Data Formats in Biological Science

9 hours

Databases and file formats: GenBank, VCF, BED, SRA, FASTA, and Fastq - Mapping data formats: SAM - BAM and CRAM - General Feature Format/General Transfer Format for Genes, DNA, RNA, and Protein sequences - Binary Base Call format - Protein structure file formats: PDB, PSF and mmCIF - small molecule file formats: SMILES, MOL2, SDF and MultiMOL2.

#### Unit II: Data Analysis Tools for Biologists

9 hours

Gene Database tools: Ensemble, Gene Cards, and UCSC Genome browser - Protein Analysis Tools: UniProt, Expasy, InterProscan, Motif prediction and ProFunc - Protein Visualization Tools: Rasmol, and Chimera - Structure Prediction Tools: SwissPDB, Alpha fold, iTasser, and PDBeMotif - Pathway Database: KEGG-GeneMania-STRING-Biogrid.

#### Unit-III: Overview of Big Data

9 hours

Big Data and its relevance in biological science – Its Characteristics: OMICS and Cancer Research - Sources and generation of Big Data in biological science: High throughput techniques - Challenges and opportunities in analyzing Big Data.

#### Unit-IV: Data Mining and Statistical Analysis

9 hours

Introduction to data mining and visualization in biological science - Data preprocessing and cleaning techniques for biological data: KNIME, GALAXY, Open Refine - Statistical analysis of biological data: Regression, Clustering, Classification, and Resampling analysis - Machine learning algorithms for analyzing OMICS data: SVM, RF, ANN and Deep Learning

#### Unit-V: Data visualization and Applications

9 hour

Data visualization - techniques and tools: MiBiOmics and Graph Bio – Application: Machine learning principles for Biomarker determination – Machine learning approaches for Drug discovery-Predictive modeling for disease diagnosis - Immune engineering: Epitope prediction and Network analysis for understanding signaling pathways involved in immunology – Neurosciences: Image processing analysis using machine learning approaches.

Data Mining and Analysis:
lge University Press. 2014.

earning Assessr	ment		Continuous Loarnin	g Assessment (CLA)			
	Bloom's Level <mark>of Think</mark> ing	Formati CLA-1 Average (50%	ve of unit test	Life-Long CL (10	Summative Final Examination (40% weightage)		
		Theory	Practice	Theory	Practice	<u>The</u> ory	Practice
Level 1	Remember	15%	The state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the state of the s	15%		15%	-
Level 2	Understan <mark>d</mark>	25%	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	20%		<b>25</b> %	-
Level 3	Apply	30%	100 m 1 m	25%	-	30%	-
Level 4	Analyze	20%	The same of the same of the	25%		20%	-
Level 5	Evaluate	10%		10%		10%	-
Level 6	Create	A Lording	- 1	5%	- (	-	-
	Total	100 %	ó	100	0 %	10	0 %

Course Designers		•
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
Dr. C. Ramakrishnan, Principal Scientist, Molecular     Design Division Aroniter Co., Seoul, South Korea ramakrishnan@arontier.co	Dr. Sanjeev Kumar Singh, Department of Bioinformatics, Alagappa University, sanjeevslab@gmail.com	1. Dr. V. Sivaramakrishnan, SRM IST
2. Mr. Sudheendra Rao, Director, DataLore Labs, Bengaluru	2. Dr. P. Gautam, Department of Biotechnology, Anna University, Tamil Nadu, India. pgautam@annauniv.edu	2. Dr. S.K.M. Habeeb, SRM IST

Course Code	21BTE425T	Course Name	R PROGRAMMING FOR BIOLOGISTS	Course Category	Е	PROFESSIONAL ELECTIVE	L 3	T 0	P 0	<u>C</u>

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

Course L	earning Rationale (CLR): The purpos <mark>e of learning</mark> this course is to:		4	M	12	Prog	ram O	utcom	es (PC	<b>D</b> )					rogram	
CLR-1:	generate knowledge about the fundamentals of R programming	1	2	3	4	5	6	7	8	9	10	11	12		pecific itcome	
CLR-2:	provide a detailed understandin <mark>g of statist</mark> ical approaches in R	dge		of	SL	1		\	. 1	ork		9				
CLR-3:	,		S	velopment	restigations problems	age	ъ	1		N N		inan	Бu			
CLR-4:				udol	estig probl	l Usa	rand	∞ >		Teal	ion	& F	ami			
CLR-5:	understand the importance of transcriptomics and pathway based packages	aring	Toc   An Initing		Mgt.	ng Le										
		inee	roblem	ign/	gric	ern	eng	ai g	S	vidu	nuı	oject	Lon	0-1	0-5	က္
Course Outcomes (CO): At the end of this course, learners will be able to:		Eng	Po	Des	Col	Moo	The	Sus	ΕĒ	Indi	Col	Proj	Life	PSC	PSC	PSC
CO-1:	illustrate knowledge about the fundamentals of R programming	J-	2	2	1	2			-	-	-	-	-	2		2
CO-2: discuss about the importance of R in statistics and biological sciences		3	3	100		2	- (		-	-	-		-	-	2	2
CO-3: outline the significance of bioconductor packages for whole genome analysis		T	2	17.0	2	3	-	_	-	-	-	-	2	-	-	-
CO-4:	CO-4: demonstrate the applicability of bioconductor packages to do exome research		3	100	- 55	3	- [	-	-	12	-	-	-	-	3	3
CO-5:			3	16	1	3	- 7		-	-	-	-	-	-	2	2

Unit-1 : Fundamentals of R 9 Hours

Introduction to R – Installation and R-Studio - Datatypes & Data structures - scalars, vectors, matrices, data frames, lists, array, factors and tables; Loops and Conditionals; Operators - arithmetic, relational, logical; Strings, Input/Output (I/O) - Introduction to basic statistical functions and packages

Unit-2: Statistics in R

Statistics in R

Challenge of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statistics of the statist

Statistics in R - mean, median, mode, standard deviation, functions and strings - concepts of statistical significance - F-test, t-test, Anova - one- and two-way, Chi square test - Linear algebra, correlation, regression. PCA, Data Visualization - charts & types - ggplot2 - heatmaps, correlogram

#### Unit-3: Bioconductor and Genome Analysis

9 Hours

Introduction to Bioconductor – Installation – CRAN vs Bioconductor – BiocManager – Discovering and Installing packages - string manipulation, BioStrings - Genome annotation - features - databases, bioconductor packages - read alignment: - Genome Annotation: BSGenome – Genomic Features – RSAM Tools – Genomic Alignments - Rsubread, Orthology. eg.db – Gene Ontology database

Unit-4 : Exome Analysis 9 Hours

Exome - significance of exome data analysis - SNP - SNV - Packages - Gwascat, VariantAnnotation, SeqVarTools, StructuralVariantAnnotation, RAREsim, XNAString, VariantFiltering - heatmap - pharmacogenomics: PharmacoGx, drugTargetInteractions

#### Unit-5: Transcriptome and Pathway Analysis

9 Hours

Transcriptomics - features - tools: dada2 - easyRNASeq, Differential expression: Limma, edgeR, RNASeqR, deseq2, DNAfusion, pathwayPCA, Co-expression profile: GeneGeneInteR, gCrisprTools, Enrichment analysis: EGSEA - DEScan2 - GSVA - pathview

Learning Resources	<ol> <li>Dylan Z. Childs &amp; Andrew P Beckerman. Second Edition, OUP Oxford. 2017</li> <li>Dan MacLean. R Bioinformatics Cookbook. First Edition, Packt Publishing Ltd. 2019</li> </ol>	<ol> <li>Matloff N. "The art of R programming: A tour of statistical software design". No Starch Press; 2011</li> <li>https://bioconductor.github.io/BiocWorkshops/r-and-bioconductor-for-everyone-an-introduction.html</li> </ol>
	- CNENC	

earning Assessn	ment		Continuous Loarnin	Assessment (CLA)				
	Bloom's Level of Thi <mark>nking</mark>	CLA-1 Avera	native ge of unit test 0%)	Life-Long CL (10	4-2	Summative Final Examination (40% weightage)		
		Theory	Practice	Theory	Practice	Theory	Practice	
Level 1	Remember	15%	A 10 July 200	15%		15%	-	
Level 2	Understand	25%	<b>表达的过程的</b>	20%		25%	-	
Level 3	Apply	25%	Mark personal process	25%		25%	-	
Level 4	Analyze	25%	A SA PARA HAL	30%		25%	-	
Level 5	Evaluate	10%	13. P 13.0 (	10%	957 3 - Za	10%	-	
Level 6	Create		Real Property Services	20 July 18 18 18 18 18 18 18 18 18 18 18 18 18		÷ -	-	
	<b>Total</b>	10	0 %	100	) %	100	) %	

Course Designers		
Experts from Industry	Experts from Higher Technical Institutions	Internal Expe <mark>rts</mark>
Dr. C. Ramakrishnan, Principal Scientist, Molecular     Design Division, Aroniter Co., Seoul, South Korea, ramakrishnan@arontier.co	<ol> <li>Dr Shandar Ahamad, Jawaharlal Nehru University, New Delhi.shandar@jnu.ac.in</li> </ol>	1. Dr. S. K. M. Habeeb, SRMIST
Mr. Sudheendra Rao, Director, DataLore Labs,     Bengaluru	Dr. Michael Gromiha, Indian Institute of Technology     Madrasgromiha@iitm.ac.in	2. Dr. T. Anand, SRMIST

Course Code	21BTE426T	Course Name	SYSTEMS BIOLOGY - M	ODEL <mark>ING AND SIMULATION</mark>	Course Category	PROFESSIONAL ELE	CTIVE	L 3	T 0	P 0	C 3
Pre-requis		Nil	Co- requisite Courses	Nil	Progressive Courses	***	Nil				
	offering Departm	ent	Biotechnology	Data Book / Codes / Stand		Nil					-

Course Le	earning Rationale (CLR):	: The purp <mark>ose of learn</mark> ing th	is course is to:					1	Progi	am Oı	utcom	es (PC	<b>)</b>					rogram	
CLR-1:	understand importance of	of model <mark>ling biologi</mark> cal system			1	2	3	4	5	6	7	8	9	10	11	12	_	pecific Itcome	
CLR-2:	acquire knowledge on fu	ındam <mark>enta</mark> l <mark>prin</mark> ciples that gov	ern biological system modelling	18.17	dge		of	SL		Α.		Se ork							
CLR-3:					wlec	alysis	pment	stigations oblems		ъ			N N		Finance	Вu			
CLR-4:							elopr	/estig probl	$\supseteq$	ar and	t &		Tea	ation	∞	arni			
CLR-5:								ot inv	10 10	ingineer tv	ment ability		ळ	ပ	roject Mgt.	ng Le			
	appy design principles in synthetic bloogy					roblem	ign/	duc		ety	rol	S	/idu	mmuni	ect	Lon	-1	7.5	53
Course O	ourse Outcomes (CO):  At the end of this course, learners will be able to:		N. 188.	Eng	Prot	Des	Cod	Moo	The	Env	Eŧ	Indi	Sol	Proj	Life	PSO-1	PSO.	PSC	
CO-1:	review the basic principle	<mark>les invol</mark> ved in modelling biolog	ical systems	11 17	3	2	1	-,1	-	- /	-3	-	-	-	-	-	-	-	2
CO-2: discuss knowledge on basic biological process that can be modelled and simulated				The second	3	3	- 3	4	-	-	-	-	-	-	-	-	-	-	2
CO-3:					3	_ 3	2	- 1	-		-	-	1	-	-	-	-	-	2
CO-4:						3	3	2	-	- 2	-	-	-	-	-	-	-	-	2
CO-5:					F.	3	3	3	-	- %		-	-	-	-	-	-	-	2

#### Unit-I: Systems Biology Introduction

9 Hour

Principles of Systems Biology, Reductionist Vs Holism approach - Challenges in modelling biological systems - The practice of modelling - Building the model - Simulating the biological model - Troubleshooting biological models - Graph theory and biological networks

#### Unit-II: Standard Models and approaches

9 Hour

Kinetic laws for modelling Biochemical reactions - Mass-action kinetic models - Modelling enzyme kinetics - Co-operativity: Hill kinetics - Biochemical systems theory - Discrete dynamic models:

Boolean networks - Flux balance analysis approach - Metabolic control analysis - Perturbation of metabolic networks - Metabolic engineering - Drug target identification - Scope of genome scale metabolic models.

#### Unit-III: Simulation of Signaling Pathways

9 Hour

Introduction, structure, and functions of signalling mechanisms - Signalling paradigm – Receptor Ligand Interaction - Selected biological process: Glycolysis as a model - G protein model - MAP kinase cascade - Signalling: Simple motifs - Feedback mechanisms.

#### Unit-IV: Modelling Biological Process

9 Hour

Biological oscillation - Glycolytic oscillation: Coupling of oscillator - Cell cycle modelling - Models of budding yeast cell cycle - Ageing model - Evolution and self-organization - Boolean network models - Prediction of biological systems from optimality principles - Modelling tools.

#### Unit-V: Synthetic Biology

9 Hour

Designing Biological circuits - The biobricks - Classic circuit design experiment - Designing an oscillator: the repressilator - Systems-theoretic approaches - Automating circuit design - Toggle switch - Designing modules - Exploring the design space - Automating circuit design - Concepts of redundancy, modularity, exaptation, robustness - Adelman's classic experiment - DNA data storage - The E-cell project.

		CIENCE
Learning	1.	Raman K. An Introduction to Computational Systems Biology: Systems-Level 3.Marchetti, L., Thanh, V. H., Priami, C. Simulation Algorithms for Computational
Resources		Modelling of Cellular Networks. Chapman and Hall/CRC; 2021. Systems Biology. Germany: Springer International Publishing., Germany, 2017
	2.	Wierling, C., Kowald, A., Liebermeister, W., Klipp, E. Systems Biology: A
		Textbook., Wile <mark>y, Germa</mark> ny, 2016

Learning Assessr	ment	4	(A. J. Style 1999)				
		The second second	Continuous Learnin	g Assessment (CLA)		Cum	mativa
	Bloom's Level <mark>of Think</mark> ing	Formative CLA-1 Average of (50%)			Learning A-2 1%)	Final Ex	mative amination eightage)
		Theory	Practice	Theory	Practice	<u>The</u> ory	Practice
Level 1	Remember	15%		15%		15%	-
Level 2	Understan <mark>d                                    </mark>	25%	37 Sec. 37	20%		<mark>25</mark> %	-
Level 3	Apply	30%	1 11 - 10 To a	25%	4	30%	-
Level 4	Analyze	30%	The second second	25%		30%	-
Level 5	Evaluate	3.00		10%			-
Level 6	Create	و المناصلات ا	- 1	5%			-
	T <mark>otal</mark>	100 %		100	) %	10	0 %

Course Designers	- 4		
Experts from Industry		Experts from Higher Technical Institutions	Internal Experts
Dr. V.L.Ramprasad, MedGenome Labs Ltd,     Bengaluru ramprasadv@medgenome.com	V	Dr Shandar Ahamad, Jawaharlal Nehru University, New Delhi.shandar@jnu.ac.in	1. Dr. TAnand, SRM IST
2. Dr. N. Mathan, Allianz Biosciences (P) Ltd,	- e_	Dr. Michael Gromiha, Indian Institute of Technology Madras,	2. Dr. Sivaramakrishnan, SRMIST
Puducherry nm@abpl.co.in	<u> </u>	Chennai. gromiha@iitm.ac.in	

Course	01DTE/107T	Course	COMPLITATIONAL APPROACHES IN DRUG DISCOVERY	Course	Е	PROFESSIONAL ELECTIVE	L	T	Р	С
Code	21B1E42/1	Name	COMPUTATIONAL APPROACHES IN DRUG DISCOVERT	Category		PROFESSIONAL ELECTIVE	3	0	0	3

Pre-requisite	Nil	Co- requisite	Nil	Progressive Progressive	Nil
Courses	, ,,,,	Courses	Courses		
Course Offeri	ng Department	Biotec <mark>hnology</mark>	Data Book / Codes / Stand	dards	Nil
			V TITLE AL		

Course Lo	earning Rationale (CLR)	: The purpose of learni	ng this course is to:		Program Outcomes (PO)												ogran		
CLR-1:	CLR-1: acquire the knowledge to manipulate the chemical structures of 2D and 3D			1	2	3	4	5	6	7	8	9	10	11	12		pecific tcome		
CLR-2: gain knowledge in basics of protein chemistry and functions		-the Miss	dge		of	ટા	J)		, I	١,١	ork		9						
CLR-3: understand the drug discovery process and computational approaches		nowlec	ဟ	elopment	stigations	age	ъ			M M		Finan	Ð						
CLR-4: obtaining to information to generate various computational models		Kno	nalysis	lopr	estig	l Us	er and	t &		Tea	ation	∞ర	earni						
CLR-5:	CLR-5: applications of 3D-QSAR in Drug design		gring	⋖	deve	t in	700	gineer	men		<u>ळ</u>	ပ	Project Mgt.	ng Le					
			1000		nee	roblem	ign/	ag E	ern	encet	ron Tain	SS	/idu	mmuni	ect	Ę	0-1	0-5	က္
Course O	utcomes (CO):	At the end of this cou	rse, learners will be able to:	A Section	Engil	Po	Des	o co	Moo	The	Env	EF	Indiv	Con	Proj	Life	PSC	PSC	PSC
CO-1:	apply the computational	l approaches to generate	2D and 3D chemical structures	Salary Million	1	1-1	1	152	1			-	-	-	-	-	-	-	1
CO-2: learn the basics of Protein world to design drug molecules		2	wit o	2		-	- "		-	-	-	-	2	-	-	2			
CO-3:			2	2	2		-	- 1	-	-	-	-	-	-	-	-	2		
CO-4: apply knowledge on computational models and learn best approaches for drug design		3	2	100	-55		- [	-	-	-	-	-	-	-	-	2			
CO-5:	apply 3D-QSAR in Drug	<mark>rdesign</mark>	W/2 - 2 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 -	12 X T	2	2		1.4	2	- 7	-	-	-	-	-	-	-	-	2

#### Unit-I: Representation and Manipulation of Chemical Structures

9 Hour

2D Chemical Structures- Chemical notations, Graph theory, Connection table, Morgan algorithms Structure and substructure searching, 3D Chemical Structures- Experimental 3D Databases-Conformational search and analysis: Systematic and random search analysis -Molecular descriptors- Calculation of physical and Chemical Data: Empirical approaches to the calculation properties-Molecular mechanics: DFT-Molecular Dynamics Simulation

#### Unit-II: Protein Modeling

9 Hour

Amino acids, Protein structure and confo<mark>rmational</mark> properties, Determination and validation of protein 3D structures-Ramachandran Plot, Protein structure database, Enzyme mechanisms, protein secondary and tertiary structure prediction methods: Homology and ab initio methods

#### Unit-III: Drug Design and Development

9 Hour

Drug Discovery Process-Target Identification-Lead Identification-Lead optimization-Identification of active part-Pharmacophore-functional group modification-Introduction to drug discovery process and computational approaches-Drug-Likeness and Compound Filters

#### **Unit-IV: Computational Models**

9 Hour

Virtual Screening-Structure based virtual screening-Protein\_Ligand Docking: Searching algorithms, Scoring function, applications-Pharmacophore concept of Bioisosterism:3D-Pharmacophore, Model generation, Lead hoping- Ligand based pharmacophore search, structure based pharmacophore search

#### Unit-V: 3D-QSAR in Drug Design

y Hour

Introduction of Deriving a Quantitative Structure Activity Relationship (QSAR) Equation: Simple and Multiple Linear Regression-Designing a QSAR Experiment: Selection of Descriptors, Experimental Design, Indicator variables, Molecular field analysis-Partial Least Squares-cross validation-Evaluation of QSAR models

Learning	1. Andrew Leach. Molecular Modelling: Principles & Applications, Second edition,	3. John Gasteiger & Thomas Engel. Chemoinformatics, Wiley-VCH, 2003
Resources	Pearson, 2013	
	2. Hugo Kubinyi. 3D-QSAR in Drug design, Volume-3, Kluwer Academic Publishers,	- */VA
	UK, 1998	(V).

	8	Artist		
_earning Assessm	ent		(S) 14 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	
		Continuous Learn	ing Assessment (CLA)	Summative
	Blo <mark>om's</mark> Level <mark>of Thinki</mark> ng	Formative CLA-1 Average of unit test (50%)	Life-Long Learning CLA-2 (10%)	Final Examination (40% weightage)
		Theory Practice	Theory Practice	Theory Practice
Level 1	Remember	15% -	15%	- 15% -
Level 2	Understan <mark>d                                    </mark>	25%	20%	25% -
Level 3	Apply	25% -	25%	25% -
Level 4	Analyze	25%	30%	25% -
Level 5	Evaluate	10%	10%	10% -
Level 6	Create	1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1		
	Total	100 %	100 %	100 %

Course Designers			A-3
Experts from Industry	$\leq$	Experts from Higher Technical Institutions	Interna <mark>l Experts</mark>
Dr. C. Ramakrishnan, Principal Scientist, Molecular     Design Division Aroniter Co., Seoul, South Korea.     ramakrishnan@arontier.co	Į	Dr Shandar Ahamad, Jawaharlal Nehru University, New Delhi. shandar@jnu.ac.in	1. Dr. Thirumurthy Madhavan, SRMIST
Mr. Sudheendra Rao, Director, DataLore Labs, Bengaluru.		2. Dr. Balachandran Manavalan, Assistant Professor, Department of Integrative Biotechnology, Sungkyunkwan University, Suwon 16419, Gyeonggi-do, Republic of Korea bala2022@skku.edu	2. Dr. Habeeb. S. K. M, SRMIST

Course Code	21BTE428T	Course Name	MACHINE LEARNING IN	BIOLOGICAL SCIENCES	Course Category	Е	PROFESSIONAL ELECTIVE	L T P C 3 0 0 3
Pre-requisir Courses		Nil	Co- requisite Courses	NiT	Progres		Nil	
Course Off	fering Departme	ent	Biot <mark>echnology</mark>	Data Book / Codes / Stand	ards		Nil	
				V	7/1		× 100	

Course Le	earning Rationale	The purpose of learning this course is to:	Program <mark>Outcomes</mark> (PO)							Program Specific								
CLR-1:	R-1: Understand various ML enviro <mark>nment</mark>						4	5	6	7	8	9	10	11	12	Outcomes		
CLR-2: acquire the basis of python coding						of	SI	7	17	À		Work		æ				
CLR-3:	CLR-3: provide the knowledge about ML libraries			wiedge	S	nent	vestigations problems	age	ъ			am W		Finance	Б		1	
CLR-4:	R-4: manipulate Data of various ML algorithms			5	alysis	elopment	estig probl	Us	r and	± ₹		Tea	tion	⊗ E	arni			
CLR-5:	CLR-5: learn ML algorithms and their application in biological dataset			ering	An	deve s	.⊑ ॲ	T00	ingineer tv	men		<u>∞</u>	ommunication	roject Mgt.	ng Le		1	
		5 2 4 . W		dine	oblem	ign/ tion	onduct compl	lem	a a	iron	S	Individua	nur	ect	틸	SO-1	SO-2	5-3
Course O	utcomes (CO):	At the end of this course, learners will be able to:	4 27 4 8 4	Eng	Prol	Des	Conc of co	Mod	The	Env	Ethics	İbdi	Sol	Proj	Life	PSC	PSC	PSO.
CO-1:	implement ML environm	<mark>ent usin</mark> g python librarires		2	( - Tr	2	- 4	2	- 1		-	-	-	-	-	-	-	2
CO-2:	learn the basics of Pyth <mark>on codin</mark> g necessary for ML					2	7 - 1	2	- 1	-	-		-	-	-	-	-	2
CO-3:	discuss the application of Numpy, Pandas and Matplotlib				2	-		2		-	-		-	-	-	-	- 1	2
CO-4:	learn how to use Scikit-learn to apply powerful machine learning algorithms			2			2	2		7	-		-	-	-	-	-	2
CO-5:	apply knowledge on ML	pply knowledge on ML models and learn best practices for biological dataset			-	-	2	2	- 1		-	-	-	-	-	-	-	2

#### Unit-I: Introduction of basic concepts and various ML environment

9 Hour

Machine learning pathway overview-Types of ML Algorithms: Supervised and Unsupervised learning - classification-regression-clustering-Dimensionality reduction-Model selection-Preprocessing-Feature engineering and data Preparation: Dealing with outliers, missing data

Unit-II: Python for ML 9 Hour

Introduction to Python: Installation of python software-Google Collab tool-Jupyter notebook-Basics of python programming, Variables -Identifies, Reserved word list, Arithmetic operators, Objects, Comparison operators, Logical operators-Functions-arguments-modules-String operators-List

Unit-III: Numpy, Pandas and Matplot for ML

9 Hour

Numpy: Arrays, Indexing and selection, Operation, Exercise-Pandas-Series-Data Frames: Conditional filtering, Exercise-Matplotlib: basics, figure parameters, Exercise

Unit-IV: ML Algorithms 9 Hour

Supervised learning algorithm: Support Vector Machine (SVM): Theory and intuition, SVM with Scikit, SVM Model generation-Random Forest (RF) algorithms: Theory and intuition, RF with Scikit, RF model generation -Unsupervised learning algorithm: K means clustering overview, theory, and coding - Cross-validation

### Unit-V: Application of ML Algorithms and Model Implementation in Biological Dataset

9 Hour

Case study: Introduction to various Biological Dataset construction-Data preparation, data preprocessing, dataset distributions, cross validation Test and Training split, implementation of cross validation Leave One Out (LOO), 10 fold cross validation, 5 fold cross validation, ML model performance-External validation

Learning Resources	I. Inamuddin, Jorddy N. Cruz, Moamen Salah El-Deen Refat, Tariq A. Altalhi. Drug design using Machine learning, Wiley, 2022     Alexander Heifetz. Artificial Intelligence in Drug design, Springer USA, 2021	3. Jitendra Kumar Verma, Prashant Johri, Sudip Paul. Application of Machine learning, Springer Singapore, 2020
	2. Alexander Helletz. Artificial Intelligence in Drug design, Springer USA, 2021	

Learning Assess	ment	3.17	Continuous Learnin	g Assessment (CLA)		<u> </u>			
	Bloom's Level <mark>of Think</mark> ing	Formative CLA-1 Average of (50%)		Life-Long CL	Learning A-2 9%)	Summative Final Examination (40% weightage)			
		Theory	Practice	Theory	Practice	<u>The</u> ory	Practice		
Level 1	Remember	15%	7.1	15%		<u>15</u> %	-		
Level 2	Understan <mark>d                                    </mark>	25%	30 to 30 to 30 to 30 to 30 to 30 to 30 to 30 to 30 to 30 to 30 to 30 to 30 to 30 to 30 to 30 to 30 to 30 to 30	20%		25%	-		
Level 3	Apply	30%	1 11 - 10 To a	25%	- L	30%	-		
Level 4	Analyze	30%	and the second second	25%		30%	-		
Level 5	Evaluate	5.00		10%			-		
Level 6	Create	to the letter	- 1/2/2	-	7 - 53	-	-		
	Total	100 %		100	0 %	10	0 %		

Course Designers		
Experts from Industry	Experts from Higher Technical Institutions	Internal Experts
Dr. C. Ramakrishnan, Principal Scientist, Molecular     Design Division Aroniter Co., Seoul, South Korea.     ramakrishnan@arontier.co	Dr Shandar Ahamad, Jawaharlal Nehru University, New Delhi.     shandar@jnu.ac.in	1. Dr. Thirumurthy Madhavan, SRMIST
Mr. Sudheendra Rao, Director, DataLore Labs, Bengaluru.	<ol> <li>Dr. Balachandran Manavalan, Assistant Professor, Department of Integrative Biotechnology, Sungkyunkwan University, Suwon 16419, Gyeonggi-do, Republic of Korea bala2022@skku.edu</li> </ol>	2. Dr. Habeeb. S. K. M, SRMIST



## SRM INSTITUTE OF SCIENCE AND TECHNOLOGY

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

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