# **B.TECH. BIOINFORMATICS**

# **COURSE STRUCTURE**

# Department of Biotechnology & Bioinformatics

#### PROGRAM OUTCOMES

Engineering Graduates will be able to:

- PO1: Engineering knowledge: Apply the knowledge of mathematics, science, engineering fundamentals, and an engineering specialization to the solution of complex engineering problems.
- PO2: Problem analysis: Identify, formulate, review research literature, and analyze complex engineering problems reaching substantiated conclusions using first principles of mathematics, natural sciences, and engineering sciences.
- PO3: Design/development of solutions: Design solutions for complex engineering problems and design system components or processes that meet the specified needs with appropriate consideration for the public health and safety, and the cultural, societal, and environmental considerations.
- PO4: Conduct investigations of complex problems: Use research-based knowledge and research methods including design of experiments, analysis and interpretation of data, and synthesis of the information to provide valid conclusions.
- PO5: Modern tool usage: Create, select, and apply appropriate techniques, resources, and modern engineering and IT tools including prediction and modeling to complex engineering activities with an understanding of the limitations.
- PO6: The engineer and society: Apply reasoning informed by the contextual knowledge to assess societal, health, safety, legal and cultural issues and the consequent responsibilities relevant to the professional engineering practice.
- PO7: Environment and sustainability: Understand the impact of the professional engineering solutions in societal and environmental contexts, and demonstrate the knowledge of, and need for sustainable development.
- PO8: Ethics: Apply ethical principles and commit to professional ethics and responsibilities and norms of the engineering practice.
- PO9: Individual and team work: Function effectively as an individual, and as a member or leader in diverse teams, and in multidisciplinary settings.
- PO10: Communication: Communicate effectively on complex engineering activities with the engineering community and with society at large, such as being able to comprehend and write effective reports and design documentation, make effective presentations, and give and receive clear instructions.
- PO11: Project management and finance: Demonstrate knowledge and understanding of the engineering and management principles and apply these to one's own work, as a member and leader in a team, to manage projects and in multidisciplinary environments.
- PO12: Life-long learning: Recognize the need for, and have the preparation and ability to engage in independent and life-long learning in the broadest context of technological change.

	JAYPEI	E UNIVE	RSITY OF INFORMATION T	ECHN	IOLO	GY, SC	LAN	
	COURSE C	URRICUL	UM OF BT & BI DEPARTMEN	T- 201	8 batcl	n (160 C	REDIT	S)
			B. TECH (BIOINFORMATI	$CS) 1^{st} S$	SEMES	ΓER		
S. No.	Category Code	Subject Code	Name of the Subjects	0	Course H	ours	Credits	Total Hours
				L	Т	Р		
1	HSS	18B11HS111	English and Technical Communication	2	0	0	2	2
2	HSS	18B17HS171	English and Technical Communication	0	0	2	1	2
3		18B11MA112	Basic Mathematics -1 OR	3	1	0	4	4
4	Basic Sciences	18B11BT111	Fundamental Biology	3	0	0	3	3
5		18B17BT171	Fundamental Biology lab	0	0	2	1	2
6	Basic Sciences	18B11PH112	Basic Engineering Physics-I	3	1	0	4	4
7	Engg Science	19B11CI111	Programming for Problem Solving-II	2	0	0	2	2
8	Engg Science	18B17GE173	Engineering Graphics	0	0	3	1.5	3
9	Basic Sciences	18B17PH172	Basic Engineering Physics Lab-I	0	0	2	1	2
10	Engg Science	19B17CI171	Programming for Problem Solving Lab-II	0	0 4		2	4
						Total	17.5	23
			B. TECH (BIOINFORMATIO	$(S) 2^{nd}$	SEMES	TER		
S.No.	Category Code	Subject Code	Name of the Subjects		Course H		Credits	Total Hours
				L	Т	Р		
1	Basic Sciences	18B11MA212	Basic Mathematics-II	3	1	0	4	4
2	Basic Sciences	18B11PH212	Bioinstrumentation Techniques	3	1	0	4	4
3	Engg Science	18B11EC212	Basic Electrical Sciences	3	1	0	4	4
4	Engg Science	18B17EC272	Basic Electrical Sciences lab	0	0	2	1	2
5	Engg Science	18B11CI211	Data Structure & Algorithms	3	1	0	4	4
J	<b>F a</b> ·	18B17CI271	Data Structure & Algorithms Lab	0	0	4	2	4
6	Engg Science	1001/012/1						
-	Engg Science	18BI7GE171	Workshop Practices	0	0	3	1.5	3

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# COURSE CURRICULUM OF BT & BI DEPARTMENT- 2018 batch (160 CREDITS)

			B. TECH (BIOINFORMATIC	$CS) 3^{rd} S$	SEMES	ΓER		
S. No.	Category Code	Subject Code	Name of the Subjects	Cc	ourse Hou	ırs	Credits	Total Hours
				L	Т	Р		
1	HSS	18B11HS311	Interpersonal Dynamics Values and Ethics	3	0	0	3	3
2	Basic Sciences	18B11BI311	Cell and Molecular Biology	3	0	0	3	3
3	Engg Science	20B11BI311	Bioinformatics Data Management	3	1	0	4	4
4	Professional Core	18B11BI312	Microbiology & Immune System	3	1	0	4	4
5	Professional Core	18B11BI313	Biological Computation	3	1	0	4	4
6	Engg Science	20b17BI371	Bioinformatics Data Management Lab	0	0	2	1	2
7	Basic Sciences	18B17BI371	Cell and Molecular Biology Lab	0	0	2	1	2
8	Professional Core	18B17BI372	Microbiology & Immune System Lab	0	0	2	1	2
9	Professional Core	18B17BI373	Biological Computation Lab	0	0	2	1	2
10	Professional Core	18B17BI374	Linux Lab	0	0	2	1	2
						Total	23	28
	•					•		
			B. TECH (BIOINFORMATIC	$CS) 4^{th} S$	SEMES	ΓER		
C M		Subject			л II			Total
S.No.	Category Code	Code	Name of the Subjects		Course H		Credits	Hours
				L	Т	Р		
1	HSS	18B11HS411	Finance and Accounts	3	0	0	3	3
2	Basic Sciences	18B11MA411	Bio-Statistics	3	0	0	3	3
3	Professional Core		Genetic Engineering and Genomics	3	0	0	3	3
4	Engg Science		Object Oriented Programming	3	1	0	4	4
5	Professional Core	18B11BI413	Structural Biology	3	0	0	3	3
6	Professional Core	18B11BI414	Programming Languages for Bioinformatics	3	0	0	3	3
7	Engg Science	18B11CI474	Object Oriented Programming Lab	0	0	2	1	2
8	Basic Sciences	18B11MA412	Bio-Statistics Lab	0	0	2	1	2
9	Professional Core	18B17BI472	Genetic Engineering and Genomics Lab	0	0	2	1	2
10	Professional Core		Structural Biology Lab	0	0	2	1	2
11	Professional Core	18B17BI474	Programming Languages for Bioinformatics Lab	0	0	2	1	2
12	Mandatory Course	18B11GE411	Environmental Studies	2	0	0	Audit	2

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	COURSE CU	RRICULU	M OF BT & BI DEPARTMENT	- 201	8 bate	h (160 C	CREDIT	S)
			B. TECH (BIOINFORMATICS	$(5) 5^{\text{th}} S$	SEMES'	TER		
S. No.	Category Code	Subject Code	Name of the Subjects	Course Hours		Credits	Tota Hours	
	1100	10011110511		L	Т	Р		
1	HSS	18B11HS511	Project Management and Entrepreneurship	3	0	0	3	3
2	Professional Core	18B11BI511	Design and Analysis of Algorithms	3	0	0	3	3
3	Professional Core	18B11BT511	Bioprocess Engineering	3	1	0	4	4
4	Professional Core	18B11BI512	Scripting Languages for Bioinformatics	3	0	0	3	3
5	Professional Core	18B17BI571	Design and Analysis of Algorithms Lab	0	0	2	1	2
6	Professional Core	18B17BT571	Bioprocess Engineering Lab	0	0	2	1	2
7	Professional Core	18B17BI572	Scripting Languages for Bioinformatics Lab	0	0	2	1	2
8	Professional Core	18B17BI573	Structural Bioinformatics Lab	0	0	2	1	2
9	Professional Elective		Departmental Elective-I	3	0	0	3	3
10	Open Elective		Open Elective-I	3	0	0	3	3
10	Project	18B19BI591	Minor Project Part-I	0	0 2		1	2
						Total	24	29
			B. TECH (BIOINFORMATICS	$(5) 6^{\text{th}} $	SEMES	TER		
S.No.	Category Code	Subject Code	Name of the Subjects	Course Hours			Credits	Tota Hours
				L	Т	Р		
1	Professional Core	18B11BI611	Machine Learning for Bioinformatics	3	0	0	3	3
2	Professional Core	18B11BI612	Computer Aided Drug Design	3	0	0	3	3
3	Professional Core	18B17BI671	Machine Learning for Bioinformatics lab	0	0	2	1	2
4	Professional Core	18B17BI672	Computer Aided Drug Design Lab	0	0	2	1	2
5	Professional Core	18B17BI673	Advanced Algorithms for Bioinformatics Lab	0	0	2	1	2
6	Professional Core	18B17BI674	R Language Lab	0	0	2	1	2
7	Professional Elective		Departmental Elective-II	3	0	0	3	3
8	Professional Elective		Departmental Elective-III	3	0	0	3	3
° 9	Open Elective		Open Elective-II (HSS)	3	0	0	3	3
10	Project	18B19BI691	Minor Project Part-II	0	0	4	2	4
11	Mandatory Course		Industrial Training				Audit	
					1	Total	21	27

160 credit scheme (BT & BI Deptt)

	JAYPE	E UNIVE	RSITY OF INFORMATION	N TECH	NOLC	OGY, S	OLAN	
	COURSE C	URRICUL	UM OF BT & BI DEPARTM	ENT- 20	18 bat	ch (160	CREDIT	TS)
			B. TECH (BIOINFORMA	ATICS) 7 <sup>th</sup>	SEME	STER		
S. No.	Category Code	Subject Code			Credits	Total Hours		
1	Professiona 1 Elective		Departmental Elective- IV	L 3	Т 0	P 0	3	3
2	Open Elective		Open Elective - III	3	0	0	3	3
3	Open Elective		Open Elective - IV	3	0	0	3	3
4	Project	18B19BI791	Major Project Part I	0	0	10	5	10
5	HSS		Indian Constitution	1	0	0	Audit	1
						Total	14	20
				th				
	1	1	B. TECH (BIOINFORMA	$\Delta TICS) 8^{m}$	SEME	STER	1 1	
S.No.	Category Code	Subject Code	Name of the Subjects		Course Hours			Total Hours
				L	Т	Р		
1	Professional Elective		Departmental Elective- V	3	0	0	3	3
2	Professional Elective		Departmental Elective- VI	3	0	0	3	3
3	Open Elective		Open Elective-V	3	0	0	3	3
4	Project	18B19BI891	Major Project Part II	0	0	14	7	14
						Total	16	23
			TOTAL CREDITS				160	
			TOTAL HOURS				205	
			HSS				12	
			Basic Science				25	
			Engg Science				28	
			Professional CORE				47	
			Professional Elective				18	
			OE				15	
			PROJECT				15	
			TOTAL CREDITS				160	
								160

#### 160 credit scheme (BT & BI Deptt) JAYPEE UNIVERSITY OF INFORMATION TECHNOLOGY, SOLAN COURSE CURRICULUM OF BT & BI DEPARTMENT- 2018 batch (160 CREDITS) **B. TECH (BIOINFORMATICS) PROFESSIONAL ELECTIVE-I** Subject Total Course Hours Credits S. No. Category Code Name of the Subjects Code Hours L Т Р Professional 18B1WBI531 Structural Bioinformatics 1 3 0 0 3 3 Elective Professional 18B1WBT532 2 Comparative & Functional Genomics 3 0 3 0 3 Elective Total 3 3 **PROFESSIONAL ELECTIVE-II** Subject Total Category Code Course Hours Credits S.No. Name of the Subjects Code Hours L Т Р Professional 18B1WBI631 Advanced Algorithms for Bioinformatics 1 3 0 0 3 3 Elective Professional 18B1WBT632 2 Infectious Diseases 3 0 0 3 3 Elective Total 3 3 **PROFESSIONAL ELECTIVE-III** Subject Course Hours S.No. Category Code Name of the Subjects Credits Total Code Hours Т Р L Professional Dataware housing and Mining for 18B1WBI632 1 3 0 0 3 3 Elective **Bioinformatics** Professional 18B1WBT634 **Bioenergy & Biofuels** 3 0 0 3 3 Elective Total 3 3 PROFESSIONAL ELECTIVE-IV Subject Course Hours S.No. Category Code Name of the Subjects Credits Total Code Hours L Т Ρ Professional 1 18B1WBI731 Computational Systems Biology 3 0 0 3 3 Elective Intellectual Property Rights & Professional

3

0

0

Total

3

3

3

3

18B1WBT734

Elective

Commercialization

160 credit scheme (BT & BI Deptt)

S. No.	Category Code	Subject Code	PROFESSIONAL ELECTIV Name of the Subjects		Course H	Iours	Credits	Total Hours
				L	Т	Р		
1	Professional Elective	18B1WBT831	Genetic Counselling	3	0	0	3	3
2	Professional Elective	18B1WBI831	Computational Molecular Evolution	3	0	0	3	3
						Total	3	3
			PROFESSIONAL ELECTIV	E-VI				
S.No.	Category Code	Subject Code	Name of the Subjects		Course H	Iours	Credits	Total Hours
				L	Т	Р		
1	Professional Elective	18B1WBT833	Diagnostics & Vaccine Manufacture	3	0	0	3	3
2	Professional Elective	18B1WBI834	NGS Data Analysis & Applications	3	0	0	3	3
						Total	3	3

# **B.TECH. BIOINFORMATICS**

# SYLLABUS

# **Fundamental Biology**

# **Pre-requisite:**X<sup>th</sup> Class Biology

# **Course Objectives:**

- 1. This is basic foundation biology course for the students having mathematics background.
- 2. The objectives are to familiarize students with basics of biology.
- 3. Learn about various living organism.
- 4. Learn about different biological at molecular or celluar level.

# **Course Outcomes:**

S. No.	Course Outcomes	Level of Attainment
CO-1	Overview of living system, different life forms and Maintenance of Life.	Familiarity
CO-2	Fundamental understanding of Bio-molecules: Building blocks of living system	Assessment
CO-3	Understanding of structure and function of cell: Prokaryotic and Eukaryotic cells system.	Assessment
CO-4	Understanding the Basic of cellular transport system and cellular inheritance.	Assessment
CO-5	Flow of information in biological system- Central Dogma, DNA replication, Transcription, and Translation	Usage

#### **Course Contents:**

Unit	Contents	Lecture s
		require d
1	<b>General Biology:</b> The nature of life, Characteristics of living organisms, Concept and use of a classification system, brief of five Kingdome and three domain classification system.Concepts of species and hierarchical taxa, biological nomenclature, classical and quantititative methods of taxonomy of plants, animals and microorganisms.	
2	<b>Introduction to bio-molecule: Structure and function relationship:</b> Structure, chemical reactions and biological functions of carbohydrate, lipid, protein and nucleotides.Stablizing interactions (Van der Waals, electrostatic, hydrogen bonding, hydrophobic interaction, etc	8
3	<b>Cell: Basic structure and functions:</b> Unicellular, colonial and multicellular forms; levels of organization of tissues systems; comparative anatomy Structural and biochemical organization of cell. Prokaryotic and Eukaryotic cells. Cell organelles, their molecular composition, structure and functions.	

4	Basic of cellular transport system: Diffusion, Osmosis, Active transport	4
5	Cellular inheritance: Cell division, cell cycle, Mitosis, Meiosis and Inheritance	6
6	<b>Flow of genetic information:</b> The DNA, Search for Genetic Material, RNA World, Genetic Code, Central Dogma, replication, transcription and translation, (initiation, elongation and termination).	8
7	Maintenance of Life: Adjustment and control: Homeostasis, thermoregulation, and osmoregulation, Speciation and selection.	5
	Total Lectures	42

#### Suggested Text Book(s):

- 1. Stryer, Lubert (2002). Biochemistry; Fifth edition. W. H. Freeman and Company.
- 2. Principles of Biochemistry [5th edition], Lehninger.
- 3. NCERT -XII class Biology

#### **Suggested Reference Book(s):**

- 1. Neill, Campbell (1996). Biology; Fourth edition. The Benjamin/Cummings Publishing Company. p. 309,310. ISBN 0-8053-1940-9.
- 2. A. W. Haupt, Fundamental of Biology, 3rd ed. McGRAW-HILL

#### Other useful resource(s):

- 1. https://nptel.ac.in/courses/122103039/
- 2. https://nptel.ac.in/syllabus/122103039/

# **Evaluation Scheme:**

S. No.	Exam	Marks	Duration	Coverage/Scope of
				Examination
1	T-1	15	1 Hour	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3	T-3	35	2 Hours	Entire Syllabus
4	Teaching Assessment	25	Entire	Quiz, Assignment, Attendance, etc.
			Semester	

# Course Outcomes (COs) contribution to the programme Outcomes (POs):

Course outcomes (Fundamental Biology)	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	6-04	PO-10	PO-11	PO-12	PSO-1	PSO-2	Average
CO-1	3	2	2	2	1	3	3	1	1	2	1	2	3	2	1.8
CO-2	2	2	3	2	1	3	3	-	1	2	1	3	3	2	1.8
CO-3	2	3	3	2	2	2	2	-	-	2	2	3	3	2	1.8
CO-4	3	2	2	2	3	2	2	1	-	1	2	3	3	2	2.0
CO-5	3	3	3	3	3	3	2	1	2	2	3	3	2	3	2.5
Average	2.6	2.4	2.6	2.2	2.0	2.6	2.4	1.8	1.6	1.8	1.8	2.8	2.8	2.2	

# **Fundamental Biology Lab**

COURSE CODE: 18B17BT171 COURSE CREDITS: 1 CORE/ELECTIVE: CORE L-T-P:0-0-2

Pre-requisite: X<sup>th</sup> class biology

# **Course Objectives:**

- 1. The objective of this course is to familiarize the students with basic biology laboratory techniques specifically used in modern biotechnology area.
- 2. Learn handling of microorganism
- 3. To learn about safe laboratory practices
- 4. To learn ethics, team work and discipline

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO1	Introduction to basic laboratory practices, microscopy, Bio- safety cabinet and sterilization.	Familiarity
CO2	Fundamental understanding of Biological buffers preparation and application.	Familiarity
CO3	Introduction to microscopic examination of different biological system.	Assessment
CO4	Introduction to analytical technique and application in macromolecular estimation.	Assessment
CO5	Able to understand ethics, team work and discipline.	Usage

#### List of Experiments

S. No.	Description	Hours
	Laboratory safety and basic laboratory Instrumentation Basic	2
	laboratory operation: safety procedure, general safety practice and	
	awareness. (personal safety, eye safety, handling of biologically	
	hazardous material, handling of needles, sharps and chemicals)	
1	To study the different parts and application of simple and compound	2
	microscope	
	To study the fundamental component and application of the Bio-safety	2
	cabinet (BSL) in biotechnology.	
	To study the fundamental of different sterilization method in laboratory	2
	practices (Autoclave, Radiation sterilization)	
	Biological buffers: (Preparation and application)Hands on training on	
	different buffer preparation, purification and pH measurement.	2
2	Application of purified buffer in different biotechnology experiment.	
		2
	Collect water from two different water bodies around you and study them	
	for pH, clarity and presence of any living organism.	2

	<b>Microscopic Analysis of biological sample</b> To perform simple and differential staining of given microorganism and classify them (gram staining)	2
3	Isolation and identification of microbe from given sample: Microscopic examination and motility test.	2
	To perform microscopic examination of unicellular eukaryote organism: identification and characterization	2
4	Analytical estimation of bio-molecule Estimation of Different macromolecules by visible spectrophotometer.	2
4	To study the basic of standard curve preparations and application in biotechnology experiments.	2
Total I	Lab hours	24

# Suggested/Resources:

- 1 Lab manual
- 2 Laboratory exercises in Microbiology - Harley Prescott
- Biotechnology Lab Course: Jeffery M.Becker, Guy A. Caldwell, Eve Ann Zachgo Biology 6<sup>th</sup>edition : Raven Johnson 3
- 4

# EvaluationScheme:

1	Mid Sem. Evaluation	20 Marks
2	End Sem. Evaluation	20 Marks
3	Lab Assessment	60 Marks
	Total	100 marks

# **Course Outcomes (COs) contribution to the Programme Outcomes (POs)**

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	Average
CO1	2	2	3	2	1	3	3	1	2	2	3	3	2.3
CO2	2	2	2	3	2	2	3	2	2	-	1	1	1.9
CO3	3	3	2	3	2	2	2	2	1	2	1	2	2.1
CO4	2	3	2	3	3	2	2	2	2	2	2	2	2.3
CO5	1	1	1	2	1	1	-	3	3	2	3	3	1.8
Average	2.0	2.2	2.0	2.6	1.8	2.0	2.0	2.0	2.0	1.6	2.0	2.2	

# **Cell and Molecular Biology**

COURSE CODE:18B11BI311 COURSE CREDITS: 3 CORE/ELECTIVE: CORE : 3-0-0

# **Pre-requisite:** Knowledge of Biology (10+2)

# **Course Objectives:**

1. The objective of the course is to equip students with a detailed knowledge of molecular structure and components of the cell and to understand howmolecules interact within the cell to promote proper growth, division, and development

#### **Course Outcomes:**

S.N 0.	Course Outcomes	Level of Attainment
CO-1	Able to describe the chemical components of the macromolecules of life and their functions and the structural differences between prokaryotic and eukaryotic cells or between plant and animal cells.	Familiarity
CO-2	Understand how molecular machines within the cell are constructed and regulated so that they can accurately copy, repair, and interpret genomic information.	Assessment
CO-3	Write, discuss or critique about emerging biology-related topics individually or in groups.	Application

#### **Course Contents:**

Unit	Contents	Lectures required							
1	<b>Biological Classification:</b> Introduction to the organization of eukaryotic and prokaryotic cells.								
2	<b>Biomolecules:</b> Carbohydrates: Chemical structures, nature, properties, Classification and Importance in Biological Systems.Lipids: Structure, Classification, Properties and Function.	2+3							
3	Amino acids: Classification, properties, structure, nature. Proteins: Classification, Structure and Function. Enzymes: Classification, Characteristics	3+2							
4	Molecular details of the Cell (Cells and Genomes, Cell Chemistry and Biosynthesis, Proteins)	2+2							
5	Structural functional significance of sub-cellular organelles (Mitochondria, chloroplast, Endoplasmic reticulum, Golgi apparatus, etc.).	3							

6	Nucleic acids:Bases, nucleosides and nucleotides, DNA & RNA structure, rRNA, tRNA and mRNA. Structural organization of DNA and Chromosomes, DNA Replication and Repair.	6			
7	Protein trafficking, Protein synthesis; Protein sorting, transport and secretion.	5			
8	Flow of information in biology (Central Dogma). DNA replication, DNA polymerases, Transcription	6			
9	Cellular transport across membranes; Exocytosis, Endocytosis and Receptor mediated endocytosis.	5			
Total lectures					

#### Suggested Text Book(s):

- 1. De Robertis, E. D. P. and De Robertis, Jr. E. N. F. "Cell and Molecular Biology". Lea and Febiger, New York
- 2. Karp, J. "Cell and Molecular Biology, Concepts and Experiments" Jhon Wiley and Sons Inc. USA
- 3. Stryer, Lubert (2002). Biochemistry; Fifth edition. W. H. Freeman and Company.
- 4. Lehninger "Principles of Biochemistry"

# Suggested Reference Book(s):

- 1. Lodish H, Berk A, Zipursky LS, Matsudaira P, Baltimore D, Darnell J (2000). *Molecular Cell Biology*. W. H. Freeman and Company
- 2. Molecular Biology of the Cell Alberts, B., et al. 4th edition (2002) Garland Science
- 3. Essential Cell Biology Alberts, B., et al; 3rd edition (2010) Garland Science
- 4. Molecular Cell Biology Damell Jr. J., Lodish, H and Baltimore, D. Scientific American Inc., New York
- 5. Neill, Campbell (1996). Biology; Fourth edition. The Benjamin/Cummings Publishing Company. p. 309,310. ISBN 0-8053-1940-9.

# Other useful resource(s):

1.Link to NPTEL course contents: <u>https://nptel.ac.in/courses/102104056/</u>

# **EvaluationScheme:**

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3.	T-3	35	2 Hours	Entire Syllabus
4.	Teaching Assessment	25	Entire	Assignment, Quizzes&Attendance
			Semester	

# **Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)**

Course outcomes (Cell and Molecular Biology)	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10	PO-11	PO-12	Average
CO-1	2	2	2	2	2	2	2	2	2	2	1	2	1.9
CO-2	2	2	2	2	2	1	1	2	2	2	1	2	1.8
CO-3	2	2	2	2	2	1	1	1	2	-	2	2	1.5
Average	2.0	2.0	2	2	2	1.3	1.3	1.6	2.0	1.3	1.33	2.0	

# Cell and Molecular Biology lab

COURSE CODE: 18B17BI3731

**COURSE CREDITS: 1** 

#### CORE/ELECTIVE: CORE

#### : 0-0-2

# **Pre-requisite:** Knowledge of Biology (10+2)

#### **Course Objectives:**

- 1. The objective of this course is to familiarize the students with laboratory techniques specifically in microbiology and molecular biology
- 2. At the end of the course, the student will be able to identify and analyze various applications in the field of microbiology and biotechnology.

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO1	Students will learn the basic laboratory practices	Introductory
CO2	Students will learn use and handling microscope	Technical
CO3	The students will be able demonstrate isolation of single bacterial colony through serial dilution	Technical
CO4	Students will able to prepare buffer and extraction of genomic DNA from <i>E. coli</i>	Technical/ Assessment

#### List of Experiments

S.No	Description	Hours
1	laboratory practices	2
2	Introduction to Microscope	2
3	To prepare slides of prokaryotic and eukaryotic cell to observe under microscope.	2
4	Calculation of Molarity, Normality	2
5	Carbohydrate estimation	2
6	To study the biosafety cabinet used in microbiology lab	2
7	Preparation of isolated single bacterial colony through serial dilution.	2
8	To observe difference in cultured plate prepared in laminar air flow and open air	2
9	General Instrumentations for lab. Practices; 1. pH meter	2
10	2. Spectrophotometer	2
11	Introduction to agarose gel electrophoresis	2
12	Preparation of buffer for genomic DNA extraction and	2
13	Isolation of genomic DNA	2
14	Isolation of genomic DNA (continued)	2
Total I	Lab hours	28

# **Suggested Books/Resources:**

- 1. Harley Prescott Laboratory exercises in Microbiology
- 2. Biotechnology Lab course : Jeffery M.Becker, Guy A. Caldwell, Eve Ann Zachgo
- 3. Biology 6<sup>th</sup>edition : Raven Johnson
- 4. Campbell --- Biology 7<sup>th</sup> edition
- 5. NPTL
- 6. Laboratory Manuals

# **EvaluationScheme:**

1	Mid Sem. Evaluation	20 Marks
2	End Sem. Evaluation	20 Marks
3	Lab Assessment	60 Marks
	Total	100 marks

# Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	Average
CO1	2	2	2	2	2	2	2	2	2	2	1	2	1.9
CO2	2	2	2	2	2	1	1	2	2	2	1	2	1.75
CO3	2	2	2	2	2	1	1	1	2	-	2	2	1.72
CO4	2	2	2	2	2	1	1	1	2	2	-	2	1.72
Average	2.0	2.0	2	2	2	1.25	1.25	1.5	2.0	2	1.33	2.0	

# **Biological Computation**

COURSE CODE:18B11BI313 COURSE CREDITS: 4 CORE/ELECTIVE: CORE : 3-1-0

# Pre-requisite: Introduction to Bioinformatics

# **Course Objectives:**

1. To use & develop tools to curate (compare & analyze) biological data.

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	Basic algorithms used in Pairwise and Multiple alignments.	Usage
CO-2	Understanding the methodologies used for database searching, and determining the accuracies of database search.	Usage
CO-3	Application of probabilistic model to determine important patterns.	Familiarity
CO-4	Prediction of structure from sequence and subsequently testing the accuracy of predicted structures.	Usage
CO-5	Determine the protein function from sequence through analyzing data.	Usage
CO-6	Analysis and development of models for better interpretation of biological data to extract knowledge.	Assessment

# **Course Contents:**

Unit	Contents	Lectures required
1	<b>Introduction</b> Meaning of sequence, sequence similarity, homology, meaning of alignment.	1
2	<b>Pairwise Sequence Alignment</b> Different scoring models, Substitution matrices (PAM and BLOSUM), Pairwise Alignment: Concept of Global and Local Alignment, Dot matrix method, Dynamic programming (Needleman-Wunsch algorithm, Smith-Waterman algorithm, Choosing of best scoring matrix, gap penalties, Significance of score, EVD, FASTA and BLAST algorithms, Information theory and Shanon Entropy.	13
3	<b>Multiple Sequence alignment</b> Multiple Sequence Alignment methods (MSA), Scoring of a MSA, Progressive (CLUSTALW and PILEUP), Iterative (Genetic) and Hidden Markov Model (HMM) methods of MSA, Local MSA (Profile and BLOCK analysis, and Pattern searching, and Expectation Maximization (EM) Algorithm (MEME) and Gibbs Sampler.	6
4	Structural Alignment Tools and Protein Tertiary Structure Prediction Structure alignment algorithms & Homology modeling.	3

Total lectu	ures	42
7	<b>RNA Structure Analysis</b> Terminology of RNA secondary structure, inferring structure by comparative sequence analysis, RNA secondary structure prediction, Nussinov folding algorithm, energy minimization and Zuker folding algorithm.	5
6	<b>Phylogenetic Analysis</b> Phylogenetic tree and terminology, different methods of Phylogenetic tree prediction: maximum parsimony, distance (UPGMA, NJ), maximum likelihood methods, bootstrapping, Jacknifing and Phylogenetic analysis by using Bayesian Network.	7
5	Markov Chains and HMM Frequent words in DNA, Consensus word analysis,Transaction and emission matrix, Development of training set, CpG islandprediction using HMM, Application of HMM in genefinding, and Multiple sequence alignment by HMM method.	7

# Suggested Text Book(s):

- 1. D.W. Mount *Bioinformatics: Genome and Sequence Analysis*: (2001) Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York.
- 2. Ian Korf, Mark & Josaph: BLAST, Oreilly Publisher, 2003
- 3. R. Durbin, S. Eddy, A. Krogh and G. Mitchison, *Biological Sequence Analysis: Probabilistic Models of Proteins and Nucleic Acids.* Cambridge University Press.
- 4. A.D. Baxevanis& B.F.F. Oulette*Bioinformatics A practical guide to the Analysis of Genes and Proteins*,2002, Willey International publishers.
- 5. M.J. Bishop and C.J. Rawlings (editors), *DNA and Protein Sequence Analysis---A Practical Approach* IRL Press at Oxford University Press, ISBN 0 19 963464 7 (Pbk)
- 6. J. Pevsner (2002) Bioinformatics and Functional Genomics; Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York

# Suggested Reference Book(s):

- 6. J. Setubal and J. Meidanis(1997) *Introduction to Computational Molecular Biology*, PWS Publishing Co.
- 7. J. Pevsner (2002) Bioinformatics and Functional Genomics; Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York.

#### **Other useful resource(s):**

1. <u>https://onlinecourses.nptel.ac.in/noc19\_bt01/preview</u>

#### **EvaluationScheme:**

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3.	T-3	35	2 Hours	Entire Syllabus

4.	Teaching Assessment	25	Entire	Assignment, Quizzes&Attendance
			Semester	

# Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)

Course outcomes (Biological Computation)	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	P0-7	PO-8	PO-9	PO-10	PO-11	PO-12	Average
CO-1	3	3	3	3	3	1	1	-	1	1	1	3	1.9
CO-2	3	3	3	3	3	1	1	-	1	1	1	3	1.9
CO-3	3	3	3	3	3	1	1	-	1	1	1	3	1.9
CO-4	3	3	3	3	3	1	1	-	1	1	1	3	1.9
CO-5	3	3	3	3	3	1	1	-	1	1	1	3	1.9
CO-6	3	3	3	3	3	1	1	-	1	1	1	3	1.9
Average	3	3	3	3	3	1	1	-	1	1	1	3	

# **Biological Computation Lab**

COURSE CODE: 18B11BI373

COURSE CREDITS: 1

CORE/ELECTIVE: CORE

L-T-P: 0-0-2

Pre-requisite: Basic Programming Skills

# **Course Objectives:**

3. To use and develop bioinformatics programs for comparing &analyzing biological sequence data to identify probable function.

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO1	Basic algorithms used in Pairwise and Multiple alignments.	Assessment
CO2	Understanding the methodologies used for database searching, and determining the accuracies of database search.	Assessment
CO3	Application of probabilistic model to determine important patterns.	Assessment
CO4	Prediction of structure from sequence and subsequently testing the accuracy of predicted structures.	Assessment
CO5	Determine the protein function from sequence through analysis of data.	Assessment
CO6	Analysis and development of models for better interpretation of biological data to extract knowledge.	Assessment

# List of Experiments

S.No	Description	Hours
1.	Overview of Practical classes conducted in the IInd Semester course of "Structural Biology" on RCSB, Visualization softwares, and tools related to secondary and tertiary structure predictions.	4
2.	Different sequence formats such as FASTA, PIR, EMBL, PDB, etc. Different sequence databases, retrieval of sequences from those databases and different ways to store the sequences. Calculation of the score of a pairwise alignment by using a scoring pattern (Take Home).	4
3.	Select a protein family for your mini-project and find out its superfamily. Also select another protein family which belongs to above superfamily and closer to your protein family.	4
4.	Find out structural and functional information about above protein families and superfamily.	4
5.	Write a program to align two sequences using Needleman-Wunsch algorithm?	4
6.	Use EBI (European Bioinformatics Institute) Needle sequence alignment tool to align above two sequences and compare your result with that of Needle tool	4
7.	Use of BLAST on line server to retrieve sequences from a database Develop a program based on BLAST algorithm to carry out database search?	4
8.	Use Clustaw software or on line server to align sequences from a family.	4

9.	Develop a Multiple Sequence Alignment (MSA) program based on ClustalW algorithm.	4
10.	Develop a program to identify motif from a set of sequences. Use on-line motif identification tools to predict motif in a set of sequences.	4
11	Use of Phylip package to infer phylogenetic tree in distance, maximum parsimony (MP) and maximum likelihood (ML) methods.	4
12	Use of Phylip package to determine robustness of inferred tree determined by each method.	4
Total	Lab hours	48

# **Suggested Books/Resources:**

- 1. http://hmmer.org/.
- 2. https://blast.ncbi.nlm.nih.gov/Blast.cgi
- 3. <u>https://www.genome.jp/tools-bin/clustalw</u>
- 4. <u>http://meme-suite.org/</u>
- 5. <u>http://evolution.genetics.washington.edu/phylip.html</u>
- 6. <u>https://www.rcsb.org/</u>
- 7. Mount D.W. : Bioinformatics: Genome and Sequence Analysis: (2001), Cold Spring Harbor Laboratory Press, New York.
- 8. Korf Ian & JosaphMark : BLAST, Oreilly Publisher, 2003
- 9. Durbin R., Eddy S., Krogh A. and G. Mitchison : Biological Sequence Analysis: Probabilistic Models of Proteins and Nucleic Acids. Cambridge University Press
- 10. Pevsner J. : Bioinformatics and Functional Genomics; Cold Spring Harbor Laboratory Press, New York.
- 11. Baxevanis AD &OuletteBFF : Bioinformatics A practical guide to the Analysis of Genes and Proteins, Willey International publishers

# **EvaluationScheme:**

1	Mid Sem. Evaluation	20 Marks
2	End Sem. Evaluation	20 Marks
3	Lab Assessment	60 Marks
	Total	100 marks

# Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	Average
C01	3	3	3	3	3	1	1	-	2	1	1	3	2.00
CO2	3	3	3	3	3	1	1	-	2	1	1	3	2.00
CO3	3	3	3	3	3	1	1	-	2	1	1	3	2.00
CO4	3	3	3	3	3	1	1	-	2	1	1	3	2.00
CO5	3	3	3	3	3	1	1	-	2	1	1	3	2.00
CO6	3	3	3	3	3	1	1	-	2	1	1	3	2.00
Average	3	3	3	3	3	1	1	-	2	1	1	3	

# **Microbiology and Immune System**

COURSE CODE:18B11BI312

#### **COURSE CREDITS: 4**

#### CORE/ELECTIVE: CORE

#### L-T-P: 3-1-0 **Pre-requisite:** None

#### **Course Objectives:**

- 1. The objective of the course is to develop an understanding of basic microbiological and immunological principles and be able to understand different classes of disease causing microorganisms and how they activate and counter-act the host immune system.
- 2. To provide an understanding of the principles of microbiology and immunology and techniques that can serve as a platform for other courses built on biological concepts.

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	Usage of scientific terminologies to describe & express fundamental concepts in Microbiology and Immunology.	Familiarity
CO-2	Able to apply basic principles to understand host-microbe relationship in different Infectious diseases.	Assessment
CO-3	Able to connect and integrate the knowledge obtained for applications related to Microbes, their tools and database.	Usage
CO-4	Able to connect and integrate the knowledge obtained for and applications related to Immunology, Vaccines related informatics.	Usage
CO-5	Able to connect and integrate the knowledge of microbiology and immunology from the perspective of a bioinformatician with special emphasis on microbe-immune interface.	Assessment

#### **Course Contents:**

Unit	Contents	Lectures required
1	<b>History and Introduction to Microbial World</b> : Brief history, contributions of important microbiologists, immunologists. Origin of life and the microbial world, different classes of microbes, good and the bad microbes.	3
2	Forms of microorganisms:	3
	Prokaryotes: Archaea & Bacteria, Cyanobacteria Eukaryotes: Fungi,	
	Algae, Protozoa	
	Viruses – structure, classification, Viral Replication (Lytic and Lysogenis cycle)	
3	Morphology and cell structure of microorganisms:	2
	Morphological features and characteristics of microorganisms, Gram positive and Gram negative bacteria.	

l lect	ures	42
13	<b>Failure of Host Defence Mechanisms:</b> Bacterial Persistence and survival strategies, Autoimmunity, Hypersensitivity	1
10	Immunoinformatics.	
	Resources and tools for Human Microbiota and Infectious Agents,	
12	Bioinformatics-Immunology and Infectious diseases: Bioinformatics	2
	radial immunodiffussion, immunoelectrophoresis, agglutination, ABO blood typing.	
11	Antigen- antibody interactions: Concept, precipitation – double diffusion,	4
	epitopes, haptens, Antigen Recognition by immune system: recognition of antigens by T and B Cells: Antigen processing and presentation, MHCs, role of MHC molecules in antigen presentation and co-stimulatory signals.	т
10	antibody, types of vaccines, active and passive immunization         Antigens:         Immunogenicity,         antipoly	4
У	Antibody, Antigens and Immune receptors: Immunoglobins: structure and function, immunoglobin classes and functions, monoclonal and polyclonal	4
9	Cells of the immune system, cytokines, complement system	Л
	antimicrobial peptide effectors.	
	immunity, Humoral and cell-mediated immune responsePhagocytes and	
0	Specificity, memory, discrimination of self from non- self, Innate and Acquired	т
8	Introduction to Fundamental Concepts in Immunology: Immunology-	4
7	<b>Genetics and Resistance</b> –Plasmids, Bacterial Conjugation, Transformation, Transduction, and Mutation	4
6	<b>Microbial Control:</b> Theory and practice of sterilization, Antibiotics and Concept of Resistance, Physical and chemical control methods in practice.	4
	enumeration techniques, effect of environmental conditions on growth, extremophiles, preservation techniques	
5	Growth of microorganisms: Growth curve of microbes, binary fission,	3
4	<b>Methods in microbiology</b> : Pure culture techniques, Principles of microbial nutrition, culture media and types (simple, complex, enriched, enrichment, selective & differential), replica plating techniques	4

# Suggested Text Book(s):

- Madigan, M.T., Martinko, J.M., Parker, J: Brock Biology of Microorganisms. 10th Edition.: Publisher: Prentice Hall 2003
- 2. Prescott, Harley and Klein: Microbiology, 6th Edition, McGraw Hill 2005.
- 3. Pelczar, Chan and Krieg: Microbiology by; Tata McGraw Hill.
- 4. Roger Y. Stanier,: General Microbiology
- 5. R. Ananthanarayan and CK JayaramPaniker: Textbook of Microbiology
- Kindt, T.J., Goldsby, R.A. and Osborne, B.A. (2007). KubyImmunology .W.H. Freeman and Co., New York, 7<sup>th</sup> Ed.

- 7. Roit, I. (2012). Essential Immunology. Blackwell Scientific Publications, Oxford, 12<sup>th</sup> Ed.
- 8. Pathogenomics: Genome analysis of pathogenic microbes by Hacker J and Dorbindt U. ed. Wiley-VCH

# **EvaluationScheme:**

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3.	Т-3	35	2 Hours	Entire Syllabus
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes&Attendance

# Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)

Course outcomes (Microbiology and Immune System)	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10	P0-11	PO-12	Average
CO-1	2	2	1	2	1	3	3	-	1	-	2	2	1.9
CO-2	2	2	1	2	1	3	3	-	2	1	2	2	1.90
CO-3	2	2	1	2	1	2	2	-	2	-	2	2	1.8
CO-4	3	3	1	2	1	2	2	-	-	-	2	2	2
CO-5	3	3	3	2	3	3	3	-	2	2	2	2	2.54
Average	2.4	2.4	1.4	2.0	1.4	2.6	2.6	0.0	1.8	1.5	2.0	2.0	

# **Microbiology and Immune System Lab**

COURSE CODE: 18B17BI372 COURSE CREDITS: 1 CORE/ELECTIVE: CORE L-T-P: 0-0-2

#### Pre-requisite: Good Laboratory Practices

#### **Course Objectives:**

- 1. The objective of this course is to demonstrate basic microbiological and immunological principles, approaches that enable study of microbe-host immune- interface and enable students to translate the theoretical foundation in the subject into practical understanding.
- 2. Techniques and methods to study different classes of microbes, immune system related molecules will be performed.

S.No.	Course Outcomes	Level of Attainment
CO1	Students will be able to understand and apply basic microbiological techniques and correlate them with their fundamental concepts in the subject.	Introductory
CO2	Students will be able to understand and apply basic immunological techniques and correlate them with their fundamental concepts in the subject.	Introductory
CO3	At the end of the course, students are expected to gain a broad appreciation of the basic methods and their application in the field of microbiology, handle microbial cultures independently, to study applied aspects of microbiology.	Technical
CO4	At the end of the course, students are expected to gain a broad appreciation of the basic methods and their application in the field of immunology along with applied aspects of immunology.	Technical
CO5	At the end of the course, students are expected to gain a broad appreciation of the basic methods and their application in the field of microbiology and immunology along with handle microbial cultures independently, to study applied aspects of microbiology.	Usage

# Course Outcomes:

#### List of Experiments

r		
S.No	Description	Hours

1.	Culture Techniques for Microorganisms:	
	General microbiology procedures and Equipments - use and Safety Considerations, GLP.	
	<b>a.</b> Preparation of culture media for different classes of microbes (bacteria and fungi)	10
	<b>b.</b> Bacterial growth curve: Spectrophotometry	
	<b>c.</b> Culture and Isolation of microorganisms – soil, air, water	
	d. Quantification, Purification of microorganisms	
2.	Microscopy and Staining:	
	Handling, microscopic examination of different classes of	
	microorganisms: Bacteria, fungi	
	a. Simple and differential staining of different shapes and sizes of bacteria –	4
	acid fast, gram staining	
	<b>b.</b> Microscopic examination of specific fungi using Lactophenol cotton blue staining	
3.	Identification/Characterization Techniques for Microorganisms:	
	a. Preservation techniques	
	<b>b.</b> Biochemical characterization	6
	c. Antimicrobial Susceptibility (Disk-diffusion) Test	
4.	Antigen – Antibody Interactions:	
	a. Double diffusion	
	<b>b.</b> Radial Immunodiffusion	8
	c. Rocket Immunoelectrophoresis	
	d. ABO Blood typing	
Total L	ab hours	28

# **Suggested Books/Resources:**

- 1. Benson, Harold J: Microbiological Applications: Laboratory Manual in General Microbiology, McGraw-Hill Higher Education, 2007.
- 2. Cappuccino, James G.: Microbiology: A Laboratory Manual, Pearson Education Sherman, Natalie Asia, 2004.
- 3. Harley, John P.: Laboratory Exercises in Microbiology, Tata McGraw Hill, 2003.
- 4. Dubey, R.C., Maheshwari, D.K.: Practical microbiology, S. Chand and Company Ltd, New Delhi, 2003

#### **EvaluationScheme:**

1	Mid Sem. Evaluation	20 Marks
2	End Sem. Evaluation	20 Marks
3	Lab Assessment	60 Marks
	Total	100 marks

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	Average
CO1	2	2	2	1	-	2	3	3	3	-	1	3	2.2
CO2	3	2	2	2	2	2	1	2	3	-	2	3	2.18
CO3	2	3	-	2	2	2	2	3	3	2	1	2	2.18
CO4	3	3	3	2	1	2	3	-	2	2	2	3	2.36
CO5	2	2	3	3	3	2	2	3	-	3	3	3	2.63
Average	2.4	2.4	2.5	2	2	2	2.2	2.75	2.75	2.33	1.8	2.8	

# Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)

# Linux Lab

#### COURSE CODE: XXXXXXXX

#### COURSE CREDITS: 1

CORE/ELECTIVE: CORE

L-T-P: 0-0-2

# Pre-requisite:None

# **Course Objectives:**

6. To Understand and master Linux and UNIX based OS environment and understanding to various Linux flavors.

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO1	To understand Unix environment	Familiarity
CO2	Familiarize with Unix and Linux commands.	Familiarity
CO3	To learn and master Bash and Shell scripting	Usage
CO4	To learn automating script-based job scheduling in Unix	Usage
CO5	To learn and master administrating and managing superuser-based managing accounts.	Usage
CO6	To run command line scripts of Perl and Python	Usage

# List of Experiments

S.No	Description	Hours	
1.	Introduction to Unix and Shell	2	
2.	Installing Fedora	2	
3.	Unix directories and pathnames and Rules for entering unix commands	2	
4.	Configuring your Linux environment	2	
5.	Working with Hard drives, Listing and Finding Directories and Files	2	
6.	Manipulating Files in Unix	2	
7.	Comparing, Sorting, Modifying, Combining, and Splitting Files, Searching for Lines in a File or Pipeline	2	
8.	Replacing or Removing Text From a File or Pipeline	2	
9.	Using vi to Edit a Text File and Command-Line Editing in the Korn Shell	2	
10.	Writing Bourne Shell Scripts and awk scripts	2	
11.	Additional commands	2	
12.	Network commands	2	

#### **Suggested Books/Resources:**

- 1. Practical Linux, Drew Streib et al Que, Indianapolis, 2000
- 2. Practical Unix, Steve Moritsugu et al, Que, Indianapolis, 2000
- 3. Linux: a practical approach , B. Mohamed Ibrahim 2006

#### **EvaluationScheme:**

1	Mid Sem. Evaluation	20 Marks
2	End Sem. Evaluation	20 Marks
3	Lab Assessment	60 Marks
	Total	100 marks

# **Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)**

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	Average
CO1	3	3	3	3	2	2	1	1	1	1	1	1	1.83
CO2	3	3	3	3	3	1	1	1	1	1	1	3	2.00
CO3	3	3	2	3	2	3	2	1	1	1	2	1	2.00
CO4	3	3	3	2	3	2	1	1	1	1	1	1	1.83
CO5	2	2	3	3	3	3	1	1	1	1	1	1	1.83
CO6	2	3	3	3	2	2	2	2	2	2	2	2	2.25
Average	2.67	2.83	2.80	2.80	2.60	2.20	1.20	1.00	1.00	1.00	1.20	1.40	

# **Genetic Engineering and Genomics**

COURSE CODE:18B11BI412 COURSE CREDITS: 3 CORE/ELECTIVE: CORE L-T-P: 3-0-0

# Pre-requisite: Molecular Genetics

# **Course Objectives:**

- 4. Familiarize the students with the basic concepts in genetic engineering.
- 5. Acquaint the students to versatile tools and techniques employed in genetic engineering and recombinant DNA technology.
- 6. To make the students familiar with basic concepts of technologies.

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	Students will become aware of concept of genetic engineering and genomics and its applications.	Familiarity and Basics
CO-2	Students will have knowledge of tools and strategies used in genetic engineering.	Technical and strategies
CO-3	Student will acquire knowledge about gene libraries and isolation of genes.	Technical and application
CO-4	Student will develop understanding of DNA and genome sequencing technologies.	Technical and strategies
CO-5	Student will be able to explore domains of genomic technologies.	Application

#### **Course Contents:**

Unit	Contents	Lectures required
1	<b>Introduction:</b> Genetic engineering, Recombinant DNA technology: gene cloning - concept and basic steps - rDNA Glossary, Genomics Concept scope and applications	3
2	<b>DNA modifying enzymes and cloning techniques:</b> Restriction Endonucleases, DNA Ligation Enzymes and, DNA, Gene cloning methods and strategies: Cloning of PCR products, TA cloning, DNA Modifying Enzymes: Nucleases, Kinases, phosphatases, Reverse transcriptase	
3	<b>Cloning and Expression Vectors:</b> Plasmid Vectors, Vectors based on Lambda Bacteriophage, Cosmids, M13 Vectors, Vectors for Cloning Large DNA Molecules, Expression Vectors	6
4	<b>Construction &amp; Screening of genomic libraries:</b> Genomic library, cDNA library, Growing& Storing Libraries, cDNA Cloning (5'&3' RACE)	4

5	Gene transfer Methods: Gene Transfer methods plants and animal cells, Transgenic plants and animals and their applications	4
6	<b>Structural genomics:</b> Genome Analysis, Genomics: Organization and structure of Genomes, genome complexity Sequencing genes and short stretches of DNA: Basic DNA Sequencing, Next generation sequencing technologies	7
7	Mapping and sequencing genomes: Introduction, Molecular Markers Genetic         and Physical Mapping of Genomes, Sequencing of whole genomes, Sequence         analysis of genomic DNA for identification of genes and other features data and         molecular         phylogenetics	6
8	Functional Genomics: RNA expression analysis Comparative genomics	4
9	Application domains of genome technologies: Genomics and Medicine, Genomics and Agriculture	2
otal lec	tures	42

# Suggested Text Book(s):

- 1. Principles of Gene Manipulation and Genomics SEVENTH EDITION S.B. Primrose and R.M. Twyman.
- 2. Recombinant DNA: A Short Course by JD Watson, J. Tooze and DT Kurtz.
- 3. Discovering Genomics, proteomics & bioinformatics. Second edition by A Malcolm Campbell, Davidson College; Laurie J. Heyer Davidson College ; With Foreword by Francis S. Collins

# Suggested Reference Book(s):

- 1. From Genes to Genomes: Concepts and Applications of DNA Technology by JW Dale and M Schantz
- 2. Molecular Biotechnology: Principles & Applications of Recombinant DNA Glick BR and Pasternak JJ
- 3. Genetic Engineering : Amita Rastogi and Neelam Pathak

# **EvaluationScheme:**

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3.	T-3	35	2 Hours	Entire Syllabus
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes&Attendance

# **Course Outcomes (Cos) contribution to the ProgrammeOutcomes(Pos)**

Course outcomes (Genetic Engineering and Genomics)	PO-1	PO-2	PO-3	PO-4	PO-5	9-0d	7-04	8-04	6-0d	PO-10	PO-11	PO-12	Average	
---	------	------	------	------	------	------	------	------	------	-------	-------	-------	---------	--

CO-1	3	3	3	3	3	2	2	2	3	3	2	3	2.67
CO-2	3	3	3	3	3	2	2	2	3	3	2	3	2.67
CO-3	3	3	3	3	3	2	2	2	3	3	2	3	2.67
CO-4	3	3	3	3	3	2	2	2	3	3	2	3	2.67
CO-5	3	3	3	3	3	2	2	2	3	3	2	3	2.67
Average	3.00	3.00	3.00	3.00	3.00	2.00	2.00	2.00	3.00	3.00	2.00	3.00	

# **Genetic Engineering and Genomics Lab**

COURSE CODE: 18B17BI472

#### **COURSE CREDITS: 1**

CORE/ELECTIVE: CORE

#### L-T-P: 0-0-2

#### Pre-requisite: None

# **Course Objectives:**

7. The objective of the course is to give practical exposure to student about basic tools and techniques employed in recombinant DNA technology and genomics.

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment		
CO1	Students will be able to isolate and analyze plasmid vectors.	Technical		
CO2	Students will be cut and ligate DNA fragments/vectors with help of restriction enzymes and ligase.	Technical		
CO3	The students will be able to prepare competent cells	Technical		
CO4	The students will be able demonstrate bacterial transformation with given vectors	Technical		
CO5	Students will be able to perform genome annotations, gene and molecular marker prediction	Technical		

# List of Experiments

S.No	Description	Hours
1	Introduction to rDNA laboratory, w.r.t. working bench, types of instruments and their handling, lab. Preparation of stock solutions of buffers for use in gel running, gel loading, their autoclaving; preparation of working buffers, antibiotic stocks, and storage of buffers required in rDNA practicals with detailed methodology	4
2	Plasmid DNA Preparation: Preparation of LB medium with and without antibiotics for the growth of bacterial cultures, Growth of <i>E. coli</i> , Isolation of Plasmid DNA, Electrtrophoresis of Plasmid DNA and Interpretation of results	4
3	Restriction of given plasmid or $\lambda$ DNA with the restriction enzyme <i>Eco</i> RI and HindIII or any other Restriction Enzymes	4
4	<ul> <li>To perform ligation of <i>λ/Eco</i>R I digest using T4 DNA Ligase Electrophoresis of the uncut and digested DNA and Interpretation of the results</li> <li>Electrophoresis of ligated samples by agarose gel electrophoresis, Interpretation of the results</li> </ul>	4
5	Preparation of competent cells of <i>E. coli</i> transformation	4

6	Transformation of E.coli. DH5 α cells with Empty puc/ pcambia1301/and Confirmation of transformed cells by scoring the expression of LacZ gene.	4					
7	DNASTAR MODULES	4					
8	PRIMER Designing	4					
9	Unknown Gene Prediction Tools and Packages	4					
10	Molecular Markers Prediction (SSR and SNP)	4					
11	Overview of Genomic Resources: Data retrieval and analysis	4					
12	Tools for expression data analysis	4					
Total I	Fotal Lab hours						

# **Suggested Books/Resources:**

- 1. Lab Manual.
- Discovering Genomics, proteomics & bioinformatics. Second edition by A Malcolm Campbell, Davidson College; Laurie J. Heyer Davidson College ; With Foreword by Francis S. Collins
- 3. Virtual Lab

# **EvaluationScheme:**

1	Mid Sem. Evaluation	20 Marks
2	End Sem. Evaluation	20 Marks
3	Lab Assessment	60 Marks
	Total	100 marks

# **Course Outcomes (Cos) contribution to the ProgrammeOutcomes(Pos)**

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	<b>PO7</b>	PO8	PO9	PO10	PO11	PO12	Average
CO1	3	3	3	3	3	2	2	2	3	3	2	3	2.67
CO2	3	3	3	3	3	2	2	3	3	3	2	3	2.75
CO3	3	3	3	3	3	2	2	3	3	3	2	3	2.75
CO4	3	3	3	3	3	2	2	3	3	3	2	3	2.75
CO5	3	3	3	3	3	2	2	3	3	3	2	3	2.75
Average	3.00	3.00	3.00	3.00	3.00	2.00	2.00	2.80	3.00	3.00	2.00	3.00	

# **Structural Biology**

COURSE CODE:18B11BI413

**COURSE CREDITS: 3** 

CORE/ELECTIVE: CORE

L-T-P: 3-0-0

### Pre-requisite: None

#### **Course Objectives:**

7. To visualize, analyze and compare structures of proteins and nucleic acids (DNA), and their subunits. To identify and understand similar structural units (folds and domains) in proteins those have different functions.

### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	Understand the relationship between protein structure and its function.	Familiarity
CO-2	Understand the methods of characterizing protein's structure using X-ray and NMR methods	Familiarity
CO-3	Implementation of bioinformatics tools in understanding protein structures. Understanding the classification of protein databases.	Usage
CO-4	Introduction to protein engineering	Familiarity
CO-5	Understand the structural diversity in nucleic acids.	Familiarity

#### **Course Contents:**

Unit	Contents	Lecture s
		require d
1	<b>Introduction:</b> Structural biology and its significance, Overview of amino acids and their various groups. Unnatural amino acids	1
2	Protein structure: Primary and Secondary Structure	2
3	Motifs &Supersecondary Structure, Tertiary Structure & Fold Types: Different types of secondary structures, Super-secondary structure and their classes, structural and functional domain, tertiary structures of proteins and their classes and sub-classes, Quaternary structures and cooperativity	2
4	<b>Covalent and Non-covalent Forces:</b> H-bonding, base stacking & hydrophobic interaction, paired interaction, torsion angle, solvent interaction in Protein, Role of free energy in Random and Natural states of polypeptide chain	4
5	Mechanisms of Protein Folding: Characterization of Folding Pathways and Mutagenesis Studies	2
6	Interactions of small molecule: Protein-Protein, Protein-DNA and Protein-RNA Interactions	2
7	Types of protein:Membrane Proteins, Fibrous Proteins, Metalloproteins,Carbohydrate Binding Proteins and Metalloenzymes:Structure and Function.	4

8	Protein Structure Determination by X-ray diffraction: Isolation, purification	
	& crystallization of proteins, Basic principles of X-ray diffraction studies, Phase	
	determination, calculation of Electron Density Map, Interpretation of Electron	4
	Density Map, Refinement of	
	the Structures	
9	Techniques: Circular Dichroism and Optical Rotation, Fluctuation	2
	Spectroscopy, Mass Spectrometry	2
10	NMR Techniques for protein structure determination: 1D NMR, 2D NMR	
	(COSY & NOSY) Basic NMR Principles and Parameters, Vector & Product	
	Operator Formalisms, Heteronuclear Correlation Experiments, Resonance	2
	Assignment Strategies, Protein Structure	
	Determination	
11	Protein secondary structure prediction: Principles of secondary structure	
	prediction, Various secondary structure prediction tools (Chou-Fasman, GOR-	
	IV, Neural network), Comparisons of various secondary structure prediction	4
	tools	
12	Structural Classification of Proteins: Principle of protein structure	
	classification (VAST, DALI, SSP), Protein structure classification	3
	Database (SCOP, CATH, DSSP), Profiles and Protein Families	
13	Protein Design: Structural Scaffolds and Enzymatic Function.	3
	Introduction to protein engineering, examples and applications of industrially	
	important enzymes	
14	Nucleic Acid Structures: DNA Tertiary structure(A- and B- DNA, Major and	3
	Minor Grooves of DNA, Z-DNA, Mechanism of specific base sequence	
	recognization in B-DNA, Triple helix DNA, Tetraplex DNA, Introduction to	
	RNA secondary structure	
tal lectu	ires	42

- 1. Introduction to Protein Structure, Carl Branden and John Tooze, Garland Publishing Inc., New York
- 2. Bioinformatics: sequence and Genome Analysis, DW Mount, Cold Spring Harbor Laboratory Press, 200
- 3. Creighton T.E. ed. Protein structure. A practical approach. (2004) Oxford University Press

### **EvaluationScheme:**

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3.	T-3	35	2 Hours	Entire Syllabus
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes & Attendance

Course Outcomes (	COs)	contribution to	the Program	meOutcomes(POs)
Course oures (	~~~,			

Course outcomes (Structural Biology)	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10	PO-11	PO-12	Average
CO-1	2	2	2	2	2	1	1	1	2	2	2	2	1.75
CO-2	2	3	3	3	3	1	1	1	2	2	1	2	2
CO-3	2	2	2	2	3	1	1	1	2	2	1	2	1.75
CO-4	2	3	3	3	2	1	1	1	2	3	2	2	2
CO-5	2	3	3	3	2	1	1	1	2	3	2	2	2
Average	2	2.6	2.6	2.6	2.6	1	1	1	2	2.4	0.6	2	

# **Structural Biology Lab**

COURSE CODE: 18B17BI473 COURSE CREDITS: 1 CORE/ELECTIVE: CORE L-T-P: 0-0-2

Pre-requisite: Thermodynamics and Chemical Processes, Microbiology

#### **Course Objectives:**

- 1. To visualize, analyze and compare structures of proteins and nucleic acids (DNA), and their subunits.
- 2. To develop the ability to design, predict, analyze and compare the protein structures.

### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO1	Understand the relationship between protein structure and its function.	Familiarity
CO2	Understand the methods of characterizing protein's structure using X-ray and NMR methods	Familiarity
CO3	Implementation of bioinformatics tools in understanding protein structures. Understanding the classification of protein databases.	Usage
CO4	Introduction to protein engineering	Familiarity
CO5	Understand the structural diversity in nucleic acids.	Familiarity

### List of Experiments

S.No	Description	Hours
1	Understanding Protein structures and Visualization	2
2	Drawing helical wheel for alpha helix	2
3	Using Rasmol and PyMOL for 3-D visualization	2
4	Analysis of protein-protein interaction and protein-DNA interaction	2
5	Advanced PyMOL usage	2
6	Use of PDBsum for structural analysis	2
7	Protein-Ligand interactions: LIGPLOT	2
8	Secondary structure prediction methods	2
9	PROSITE - Protein signature patterns	2
10	Understanding Ramachandran plots and X-Ray Crystallography	2
11	RNA secondary structure visualization	2
Total La	ıb hours	22

#### **Suggested Books/Resources:**

- 1. Introduction to Protein Structure, Carl Branden and John Tooze, Garland Publishing Inc., New York
- 2. Bioinformatics: sequence and Genome Analysis, DW Mount, Cold Spring Harbor Laboratory Press,

### 200

3. Creighton T.E. ed. Protein structure. A practical approach. (2004) Oxford University Press

### **EvaluationScheme:**

1	Mid Sem. Evaluation	20 Marks
2	End Sem. Evaluation	20 Marks
3	Lab Assessment	60 Marks
	Total	100 marks

Course outcomes (Structural Biology)	POI	P02	P03	P04	P05	P06	P07	PO8	P09	P010	P011	P012	Average
CO1	2	2	2	2	2	1	1	1	2	2	2	2	1.75
CO2	2	3	3	3	3	1	1	1	2	2	1	2	2
CO3	2	2	2	2	3	1	1	1	2	2	1	2	1.75
CO4	2	3	3	3	2	1	1	1	2	3	2	2	2
CO5	2	3	3	3	2	1	1	1	2	3	2	2	2
Average	2	2.6	2.6	2.6	2.6	1	1	1	2	2.4	0.6	2	

# **Programming Languages for Bioinformatics**

COURSE CODE:18B11BI414

COURSE CREDITS: 3

CORE/ELECTIVE: CORE

L-T-P: 3-0-0

### Pre-requisite: None

### **Course Objectives:**

1. To familiarize and master the programming skills in Perl and Python.

### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	Write and execute a script in Perl.	Usage
CO-2	Enable routine and module calls and their implementation using Bioperl.	Familiarity
CO-3	Able to formulate stepwise implementation of a Perl script (from developing a pseudo-code to execute a successful bug-free code) for a given problem in Bioinformatics.	Usage
CO-4	Write and execute a script in Python.	Usage
CO-5	Enable routine and module calls and their implementation using Biopython.	Familiarity
CO-6	Able to formulate stepwise implementation of a Python script (from developing a pseudo-code to execute a successful bug-free code) for a given problem in Bioinformatics.	Usage

### **Course Contents:**

Unit	Contents	Lectures required
1	Crash course in C	2
2	Programming basics	1
3	Sequences and Strings: Storing a DNA sequence, Concatenation, Transcription, Translation	2
4	Arrays and Scalar list, Strings to Array, Operations on Strings	2
5	Subroutines and Command line arguments	3
6	Modules, Calling modules	2
7	Hashes, Data Structures in Perl	4
8	Reading files and writing output formats	3
9	Regular expressions and Perl Operations	3
10	Parsing genbank, PDB, BLAST, and other file formats	3
11	Object-oriented programming, Complex Data Structures, Relational Databases	4

12	BioPerl	3					
13	Introduction to Python	3					
14	BioPython	4					
15							
Total lectu	otal lectures						

- a. Beginning Perl for Bioinformatics By James Tisdall, O'Reilly Media (2001)
- b. Mastering Perl for Bioinformatics By James Tisdall, O'Reilly Media (2003)
- c. Python For Bioinformatics By Sebastian Bassi, Chapman and Hall (2010)

### **EvaluationScheme:**

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3.	T-3	35	2 Hours	Entire Syllabus
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes&Attendance

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	<b>PO7</b>	PO8	PO9	PO10	PO11	PO12	Average
CO1	3	3	3	3	2	2	1	1	1	1	1	1	1.83
CO2	3	3	3	3	3	1	1	1	1	1	1	3	2.00
CO3	3	3	2	3	2	3	2	1	1	1	2	1	2.00
CO4	3	3	3	2	3	2	1	1	1	1	1	1	1.83
CO5	2	2	3	3	3	3	1	1	1	1	1	1	1.83
CO6	2	3	3	3	2	2	2	2	2	2	2	2	2.25
Average	2.67	2.83	2.80	2.80	2.60	2.20	1.20	1.00	1.00	1.00	1.20	1.40	

# **Programming Languages for Bioinformatics Lab**

COURSE CODE: 18B17BI474 COURSE CREDITS: 1 CORE/ELECTIVE: CORE L-T-P: 0-0-2

Pre-requisite: C, Object-oriented data structures.

### **Course Objectives:**

1. To master programming skills in Perl and Python and implement those skills using BioPerl and BioPython.

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO1	Write and execute a script in Perl.	Usage
CO2	Enable routine and module calls and their implementation using Bioperl.	Familiarity
	Able to formulate stepwise implementation of a Perl script (from	Usage
CO3	developing a pseudo-code to execute a successful bug- free code) for a	
	given problem in Bioinformatics	
CO4	Write and execute a script in Python	Usage
CO5	Enable routine and module calls and their implementation using	Familiarity
	Biopython.	
	Able to formulate stepwise implementation of a Python script (from	Usage
CO6	developing a pseudo-code to execute a successful bug- free code) for a	
	given problem in Bioinformatics	

### List of Experiments

S.No	Description	Hours
1.	Installing and learning how to run Perl, Good programming practices	2
2.	File Handling	2
3.	Understanding Sequences and Strings, Operation on Strings - Motif finding	2
4.	Writing a subroutine and calling, Translation of DNA to Protein Sequences	2
5.	Mutating a DNA sequence and generating a random DNA sequence	2
6.	Reading from various file formats	2
7.	Installing Bioperl, Translation of DNA to Protein Sequences using Bioperl	2
8.	Reading and Parsing PDB files using Bioperl	2
9.	Automating BLAST and Parsing BLAST Output using Bioperl	2
10.	Python File handling	2
11	Gene Expression Analysis with Python	2
12	Using BioPython – Part I	2
13	Using BioPython – Part II	2

#### **Suggested Books/Resources:**

- 1. Pauline M. Doran, "Bioprocess Engineering Principles" 8th ed., Academic press, New York, 2003.
- 2. M.L. Shuler and F. Kargi, "Bioprocess Engineering--basic Concepts", 2nd Edn. Prentice-hall Of India Pvt Ltd (2008).
- **3.** Peter F. Stanbury, Stephen J. Hall & A. Whitaker, "Principles of Fermentation Technology", Â Elsevier India Pvt Ltd. (2007).

#### **EvaluationScheme:**

1	Mid Sem. Evaluation	20 Marks
2	End Sem. Evaluation	20 Marks
3	Lab Assessment	60 Marks
	Total	100 marks

course outcomes (COS) contribution to the ringramme outcomes(103)														
CO/P	0   P(	<b>D1</b>	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	Average
CO1	. 3	3	3	3	3	2	2	1	1	1	1	1	1	1.83
CO2	: 3	3	3	3	3	3	1	1	1	1	1	1	3	2.00
CO3	3	3	3	2	3	2	3	2	1	1	1	2	1	2.00
CO4	3	3	3	3	2	3	2	1	1	1	1	1	1	1.83
CO5	; 2	2	2	3	3	3	3	1	1	1	1	1	1	1.83
COG	2	2	3	3	3	2	2	2	2	2	2	2	2	2.25
Avera	ge 2.0	67	2.83	2.80	2.80	2.60	2.20	1.20	1.00	1.00	1.00	1.20	1.40	

## **Environmental Studies**

COURSE CODE:18B11GE411

### **COURSE CREDITS: 2**

### CORE/ELECTIVE: Audit

#### -L-T-P: 2-0-0

### Pre-requisite: None

### **Course Objectives:**

- 1. Identify environmental problems arising due to engineering and technological activities and the science behind those problems.
- 2. Estimate the population- economic growth, energy requirement and demand.
- 3. Analyze material balance for different environmental systems
- 4. Realize the importance of ecosystem and biodiversity for maintaining ecological balance.
- 5. Identify the major pollutants and abatement devices for environmental management and sustainable development.
- 6. Recognizing the major concepts of environmental studies, developing problem solving ability, forecasting the global climate change

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	Introducing basic concept of environmental studies, interdisciplinary nature and scope of the subject	Familiarity
CO-2	Understanding ecosystem services and its functioning as well as equitable use of natural resources.	Assessment
CO-3	Understanding Pollution, A threat to the environment and finding its solutions, Pollutant sampling and monitoring of samples.	Assessment
CO-4	Correlating the concept of Biodiversity and its importance to human mankind	
		Usage
CO-5	Understanding social issues and their impact on environment.	Usage
CO-6	Role of Information Technology in environment and human health	Usage

#### **Course Contents:**

U nit	Contents	Lectures required
1	<b>Unit 1: Multidisciplinary nature of environmental studies:</b> The Multidisciplinary nature of environmental studies: Definition, scope and importance, Need for public awareness, Types of Ecosystems, World Biomes, Ecosystem functioning, Biogeochemical cycles.	
2	Unit 2: Natural resources, their consumption & Protection: Natural resources, their consumption & Protection: Water, Land Energy (Renewable, non-renewable, wind, solar, hydro, Biomass), Mineral, Forest, & Food resources, Role of an individual in conservation of natural resources, Equitable use of resources.	

3	<b>Unit 3: Pollution- a threat to environment:</b> Pollution- a threat to environment: Air, Water & Land pollution, sources & causes, Space pollution, causes & effects, toxicity limits of pollutants. Critical issues concerning global Environment (Urbanization, population growth, global warming, climate change, acid rain, ozone depletion etc.) and the Roots in: Cultural, Social, Political, Commercial, industrial, territorial domains	4
4	<b>Unit 4: Environmental standards &amp; Quality:</b> Environmental standards & Quality: Air, Water & Soil Quality, Pollutant sampling, pollution control systems. Green Chemistry and its applications	3
5	<b>Unit 5: Biodiversity and its conservation:</b> Biodiversity loss: Diversity of flora and fauna, species and wild life diversity, Biodiversity hotspots, threats to biodiversity	4
6	<b>Unit 6: Social Issues and the Environment:</b> Waste land reclamation, consumerism and waste products, eco-consumerism, dematerialization, green technologies, eco-tourism. Water conservation, rain water harvesting, watershed management. Environment protection act, Air (prevention and control of population) act; Water (prevention and control of pollution) act, Wildlife protection act, Forest conservation act, Issues involved in enforcement of environmental legislation National Environmental Policy; Function of pollution control boards (SPCB and CPCB), their roles and responsibilities.	4
7	<b>Unit 7: Human Population and the environment:</b> Population growth, variation among nations. Population explosion—Family Welfare Programme. Environment and human health. Humanrights. Value education.HIV/AIDS. Women and Child Welfare. Role of Information Technology in environment and humanhealth. Case Studies.	4
8	Unit 8: Field work: Field Work: Explore the surrounding flora & fauna (Study of common plants, insects, birds document environmental assets), documentation of industries in local region and their possible effects, measure of water, air and land quality, Visit to a local polluted site- Urban/Rural /Industrial / Agricultural, Study of simple ecosystems-pond, river, hill slopes etc	4
Total l		30

- 1. Environmental Studies By: M. P. Poonia and S.C. Sharma, Khanna Publishers
- 2. Textbook of Environmental Studies for UG Courses ErachBharucha, University Press
- 3. Joseph, B., 2005, Environmental Studies, Tata McGraw Hill, India.

### Suggested Reference Book(s):

- 1. Nebel, B.J. & Wright, R.T., 1993, Environmental Science, 8th Edition, Prentice Hall, USA.
- 2. Chiras D D.(Ed.). 2001. Environmental Science Creating a sustainable future. 6th ed. Jones &Barlett Publishers.
- 3. David Laurance. 2003. Environment Impact assessment, Wiley publications.
- 4. Chhokar KB, Pandya M & Raghunathan M. 2004. Understanding Environment. Sage publications, NewDelhi.

### **Other useful resource(s):**

- 1. Issues of the journal: Down to Earth, published by Centre for Science and Environment.
- 2. Audio visuals from: Discovery, National Geographic etc.

- https://nptel.ac.in/courses/120108002/
   https://nptel.ac.in/courses/120108005

# EvaluationScheme:

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Syllabus covered up to T-1
2	T-2	25	1.5 Hours	Syllabus covered up to T-2
3.	Т-3	35	2 Hours	Entire Syllabus
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes, Attendance.

Course outcomes (EVS)	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	9-04	PO-10	PO-11	PO-12	Average
CO-1	2	2	2	2	2	1	1	1	2	2	2	2	1.75
CO-2	2	3	3	3	3	1	1	1	2	2	1	2	2
CO-3	2	2	2	2	3	1	1	1	2	2	1	2	1.75
CO-4	2	3	3	3	2	1	1	1	2	3	2	2	2.08
CO-5	2	3	3	2	2	1	1	1	1	1	3	2	1.83
CO-6	2	2	2	2	1	1	1	2	2	2	2	2	1.75
Average	2	2.5	2.5	2.33	2.16	1	1	1.1	1.5	2	1.8	2	

# **Bioprocess Engineering**

COURSE CODE:18B11BT511

COURSE CREDITS: 4

CORE/ELECTIVE: CORE

: 3-1-0

Pre-requisite: Thermodynamics and Chemical Processes, Microbiology, Biochemistry

### **Course Objectives:**

- 1. Learn various bioprocess related terms and principles
- 2. Learn about microbial growth kinetics in various mode of fermentation
- 3. Learn about the principles and application of Mass transfer and Sterilization
- 4. Develop an understanding of important concepts and design aspects of bioreactors
- 5. Learn about the functioning of various bioreactors
- 6. Learn about the principle of scaling up and scaling down of bioprocesses

### Course Outcomes:

S.No.	Course Outcomes	Level of Attainment
CO-1	Able to use correct biological terms to describe & analyze phenomena/ problems in bioprocesses	Familiarity
CO-2	Able to apply engineering principles to address issues in various bioprocesses	Assessment
CO-3	Able to analyze bacterial growth kinetics (homogeneous reaction) in batch /continuous/ Fed-batch reactor and sterilization	Assessment
CO-4	Able to understand and to solve problems related to bioprocess phenomena including mixing, Mass transfer and sterilization	Assessment
CO-5	To develop a strong foundation about bioreactor designs and their applications	Usage
CO-6	Able to understand the basis of bioprocess scale up and the related basic design calculations	Usage

#### **Course Contents:**

Unit	Contents	Lectures required
1	Introduction:Role of bioprocess engineer, Microbial process development, Quality control management, Fermentation Economics	3
2	Kinetics of Microbial growth: Batch culture, Kinetic implications of endogenous and maintenance metabolism. Continuous culture, Modifying continuous reactors: Chemostat with recycle and multistage Chemostat Systems. Modifying batch reactors: Fed- batch operation, Perfusion systems.	7
3	<b>Sterilization:</b> Design of batch and continuous sterilization processes, kinetics of thermal death of cells and spores.	2
4	<b>Mixing:</b> Mixingequipments, flow patterns in reactors, mixing mechanism, power consumption and shear properties of sparged and agitated vessels and various mixing agitators.	4

9	Scale up and Scale down:Scale up of bioprocesses and its difficulties. Scale up criteria for bioreactors based on oxygen transfer, power consumption and impeller tip speed. Scale down.	5
8	<b>Bioreactor design and analysis:</b> Bioreactor configurations and its utilities, Analysis of ideal and non-ideal reactors. Multiphase reactors: packed-bed reactors, bubble-column bioreactors, fluidized bed bioreactors, trickle-bed reactors. Practical considerations for bioreactor construction, Bioreactors instrumentation and control. Bioprocess Considerations: Animal cell cultures & plant cell cultures	6
7	<b>Immobilized Cell Systems (ICS):</b> Immobilization and its limitations, Active and passive immobilization, applications of immobilized cell biocatalysts. Diffusional limitations in ICS. Bioreactor considerations.	3
6	<b>Strain Improvement and Media Formulation:</b> Strain improvement of industrially important microorganisms, Media formulation industrial fermentations.	5
5	<b>Mass Transfer:</b> Role of diffusion in bioprocessing, film theory, convective mass transfer, oxygen uptake in cell cultures. Oxygen transfer in fermenters: measuring dissolved-oxygen concentration, estimating oxygen solubility, mass transfer correlation, measurement of $k_La$ , oxygen transfer in large vessels.	7

- 1. Pauline M. Doran, "Bioprocess Engineering Principles", 8th ed., Academic press, New York, 2003.
- 2. M.L. Shuler and F. Kargi, "Bioprocess Engineering--basic Concepts", 2nd Edn. Prentice-hall Of India Pvt Ltd (2008).
- 3. Peter F. Stanbury, Stephen J. Hall & A. Whitaker, "Principles of Fermentation Technology", Â Elsevier India Pvt Ltd. (2007).

#### Suggested Reference (s):

- KlaasVan't Riet, Johannes Tramper, "Basic Bioreactor Design", 2nd ed., Marcel Dekker, Inc., New York, 1991.
- 2. Bailey and Ollis, "Biochemical Engineering Fundamentals", 2nd ed., McGraw-Hill Book Company, New York, 1986.
- 3. MccabeL.Warren, Smith C. Julian and Peter Harriott, "Unit Operations of Chemical Engineering", 6th ed., McGraw Hill International Edition, New York, 2001.

#### 4. Abhilasha S. Mathuriya, "Industrial Biochnology" 1<sup>st</sup>ed., Ane Books Pvt. Ltd., New Delhi, 2009.

#### **Other useful resource(s):**

- 1. NPTEL Course Content:
  - i) Bioreactors by Prof. Suraish Kumar, IIT Madras https://nptel.ac.in/courses/102106053/
  - ii) Industrial Biotechnology by Prof. Debabrata Das, IIT Kharagpur.... https://nptel.ac.in/courses/102105058/
  - iii) Aspects of Biochemical Engineering by Prof. Debabrata Das, IIT Kharagpur https://nptel.ac.in/courses/102105064/
- 2. Link to topics related tocourse:
  - i) Mass Transfer by Prof. Bishnupada Mandal, IIT Guwahati https://nptel.ac.in/courses/103103034/13#

### **EvaluationScheme:**

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Unit 1-2
2	T-2	25	1.5 Hours	Unit 1-5
3.	T-3	35	2 Hours	Whole Syllabus
4.	Teaching Assessment	25	Entire Semester	Inform class time to time (Quizzes, Presentation, Assignments)

Course outcomes (Bioprocess Engineering)	P0-1	P0-2	PO-3	PO-4	PO-5	PO-6	P0-7	PO-8	PO-9	PO-10	PO-11	PO-12	Average
CO-1	-	1	1	1	-	-	2	1	3	2	-	3	1.75
CO-2	3	2	2	1	-	-	-	1		1	-	3	1.86
CO-3	3	3	3	3	2	-	2	1	2	1	-	2	2.20
CO-4	3	3	3	1	-	-	-	1	2	1	-	1	1.88
CO-5	3	1	2	1	2	2	-	1	-	2	-	1	1.67
CO-6	3	3	3	3	-	2	2	1	3	1	2	2	2.27
Average	3.00	2.17	2.33	1.67	2.00	2.00	2.00	1.00	2.50	1.33	2.00	2.00	

# **Bioprocess Engineering Lab**

COURSE CODE:18B17BT571 COURSE CREDITS: 1

CORE/ELECTIVE: CORE

: 0-0-2

### Pre-requisite: Microbiology Lab, Biochemistry Lab

#### **Course Objectives:**

- 1. Provide exposure to the students with hands on experience on various practices in Bioprocess Engineering.
- 2. Enable students to link the theoretical knowledge of bioprocess engineering with the experiments.
- 3. Learn how to operate bench scale fermentor
- 4. Learn how to determine various Monod's Kinetics parameter

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO1	Able to apply practical knowledge to understand the various important process engineering aspects involved in biotechnology industries	Familiarity
CO2	Able to design experiments and analyze various data related to various practices in bioprocess engineering	Assessment
CO3	Ability to apply theoretical concepts for data analysis and interpretation and their documentation	Assessment and Usage
CO4	Able to run fermenter and also to analyze their results	Usage
CO5	Able to understand and determine various growth kinetics parameters in a batch culture	Assessment and Usage
CO6	Able to work in a team to accomplish the experiments and to document the experiments properly in lab note books	Assessment

#### List of Experiments

S.No.	Description	Hours
1	Introduction of Lab and lab safety	1
2	Describe the various parts of the bench-top fermenter (bioreactor) along with their functions.	1
3	To determine the thermal death point of a microbial culture.	2
4	To determine the thermal death time of a microbial culture.	2
5	To estimate the reducing sugar concentration in a given sample using DNS method.	2
6	To estimate the sugar concentration in fresh and spent media using DNS method.	2
7	To establish the correlation between OD and dry cell weight.	2

8	To study the different phase of microbial growth.	2
9	To study growth kinetics parameters of E. coli.	4
	a) Specific growth rate ( $\mu$ ) h <sup>-1</sup>	
	b) Maximum specific growth rate $(\mu_m) h^{-1}$	
	c) Saturation constant $(K_s)$ gm/l	
	d) Growth yield coefficient $(Y_{x/s})$ gm cell/gm substrate.	
	e) Productivity of biomass gm cell/litre/h.	
10	To study the effect of varying carbon substrate on specific growth rate	2
11	Determination of Volumetric mass transfer coefficient (K <sub>L</sub> a) using dynamic gassing out method (Virtual Lab)	2
12	Preparation of Immobilized yeast cells in calcium alginate beads	2
Total Lab	o hours	24

### Suggested/Resources:

- 1. M.L. Shuler and F. Kargi, "Bioprocess Engineering--basic Concepts", 2nd Edn. Prentice-hall Of India Pvt Ltd (2008).
- 2. Lab Manual
- 3. Pauline M. Doran, "Bioprocess Engineering Principles", 8th ed., Academic press, New York, 2003.
- 4. Peter F. Stanbury, Stephen J. Hall & A. Whitaker, "Principles of Fermentation Technology", Â Elsevier India Pvt Ltd. (2007).
- 5. <u>http://iitd.vlab.co.in/?sub=63</u>

#### **EvaluationScheme:**

1	Mid Sem. Evaluation	20 Marks
2	End Sem. Evaluation	20 Marks
3	Lab Assessment	60 Marks
	Total	100 marks

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	Average
CO1	3	3	3	3	1	3	1	2	1	2	1	3	2.17
CO2	3	3	3	3	2	2	1	2	1	2	-	3	2.27
CO3	3	3	3	3	2	2	1	2	1	2	-	3	2.27
CO4	3	3	3	3	1	2	2	3	2	2	2	3	2.42
CO5	3	3	3	3	1	2	3	3	3	2	2	3	2.58
CO6	-	-	-	-	-	-	-	-	3	3	1	3	2.5
Average	3.00	3.00	3.00	3.00	1.40	2.20	1.60	2.40	1.83	2.17	1.50	3.00	

# **Scripting Language for Bioinformatics**

COURSE CODE: 18B11BI512

**COURSE CREDITS: 3** 

CORE/ELECTIVE: CORE

: 3-0-0

Pre-requisite: Basic programming principal

### **Course Objectives:**

1. To apply, and develop scripting languages codes and implement them towards the analysis of biological data. Additionally to Develop web based applications for the problems in biology. All students will be able to develop their own websites.

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	Able to apply design principles to develop web based applications specially for biological data analysis	Familiarity
CO-2	To understand working on world wide web through implementations	Familiarity and Assessment
CO-3	Use various methods from computational biology to implement their programmatic versions	Assessment
CO-4	Able to design new web pages and web sites	Assessment and Usage
CO-5	Able to developed programs to describe and analyze problems in biology	Assessment and Usage

### **Course Contents:**

Unit	Contents	Lecture Hours
1	Introduction to Internet and World Wide Web. An overview of scripting languages, with applications towards biological data and sequence analysis. Complexity of DNA problems and their computational implications and applications. Introduction to HTML, DHTML, XML. accessing different objects of the HTML page, Dynamic page generation, Cascading Style Sheets (CSS).	12
2	JAVASCRIPT: Document object model, Elements of the document object model, basic principles of JS, object based programming using JavaScript; data types and structures, array and string handling, function implementations, XML: DTD, XML schemas, XML document structure, retrieving data from database in XML format; various bio based versions of XML.	10
3	PHP: PHP beginning to advanced level, data types, array and string handling, mathematical expressions and functions in PHP, PHP programming (implementation of object model), Database connectivity using PHP.	12

4	Hands-on practice on above mentioned programming languages. Implementation of programming skills for solving problems in biology. Development of bioinformatics based small applications and web based applications.	
Total Lectu	ires	42

- 1. HTML the complete reference, 2004, TMH.
- 2. Beginning PHP and Professional PHP, 2009, Wrox, Wiley Dreamtech.
- 3. JavaScript: The complete Reference, 2004, TMH.

### **Suggested Reference Book(s):**

- 1. Biological Sequence Analysis: Probabistic models of proteins and nucleic acids (1998) Durbin R., et
  - al, Cambridge University press.

2. Many other reference and text books on these scripting languages available in the library.

### **EvaluationScheme:**

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3.	T-3	35	2 Hours	Entire Syllabus
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes & Attendance

### **Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)**

Course outcomes (Scripting Languages for Bioinformatics)	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	P0-7	PO-8	P0-9	PO-10	PO-11	PO-12	Average
CO-1	2	2	1	3	2	2	2	1	3	2	3	2	2.08
CO-2	3	2	2	2	-	-	-	1	-	1	2	2	1.88
CO-3	3	3	3	3	2	2	1	1	2	1	-	2	2.09
CO-4	3	3	3	1	-	-	-	-	2	2	2	1	2.13
CO-5	3	2	2	2	2	2	-	-	-	2	-	1	2.00
Average	2.8	2.4	2.2	2.2	2	2	1.5	1	2.33	1.6	2.33	1.6	

# Scripting Language for Bioinformatics Lab

COURSE CODE:18B17BI572 COURSE CREDITS: 1

CORE/ELECTIVE: CORE

L-T-P: 0-0-2

### Pre-requisite: Basic concepts in programming

### **Course Objectives:**

- 1. To apply, and develop scripting languages codes and implement them towards the analysis of biological data.
- 2. Additionally to develop web based applications.

### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO1	To understand working on world wide web through implementations for client and server side programming	Familiarity
CO2	Able to developed programs to describe and analyze problems in biology.	Assessment
CO3	Able to design new web pages and web sites.	Assessment
CO4	To understand coordination of HTML, Java Script and PHP.	Assessment
CO5	Able to develop web based applications especially for biological data analysis.	Assessment and Usage

### List of Experiments

S.No	Description	Hours
1	Introduction to HTML, DHTML, XML and accessing different objects of the HTML page and dynamic page generation.	2
2	HTML page and dynamic page generation.         HTML code for basic understanding of the syntax including the use of nesting of lists.	2
3	HTML code for creating a webpage including hyperlinks and images.	2
4	Construction of DTD schema,a sample xml document to represent evolutionary tree.	2
5	Construction of XML schema, a sample xml document to represent a pathway.	2
6	Implementation of Session, request, report objects in an ASP application.	2
7	Create a MySql/ MSacessesdatabase tables and execute all SQL queries.	2
8	Development of a PHP program to take set of sequences and find out conserved sequences.	2
9	Write a PHP program to construct a pathway.	2
10	Write a PHP program to connect mysql database and execute all SQL commands.	2
11	Construct a PHP interface for a given ER model.	2
12	Write a PHP program to find out ORFs existing in a given genomic sequence.	2

13	Write a PHP program to find out annotation and sequence from a fasta file	2
Total Lab	hours	26

#### Suggested/Resources:

- a. A P Thomas: HTML the complete reference, 2nd Edition, TMH, 2004.
- b. A P Thomas, <u>F Schneider</u>: JavaScript: The complete Reference, 2nd Edition, TMH, 2004.
- c. D W Mercer et al: Beginning PHP and Professional PHP, Wrox, Wiley Dreamtech, 2009.
- d. R Durbin *et al*: Biological Sequence Analysis: Probabistic models of proteins and nucleic acids, Cambridge University press, 1998.

### **Evaluation Scheme:**

1	Mid Sem. Evaluation	20 Marks
2	End Sem. Evaluation	20 Marks
3	Lab Assesment	60 Marks
	Total	100 marks

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	Average
CO1	2	2	1	3	2	2	2	1	3	2	3	2	2.08
CO2	3	2	3	2	1	-	-	1	-	1	2	3	2.00
CO3	2	3	2	3	3	2	1	1	2	2	-	2	2.09
CO4	3	3	3	1	-	1	-	-	2	2	2	2	2.11
CO5	3	2	2	2	2	2	-	-	-	2	-	1	2.00
Average	2.6	2.4	2.2	2.2	2	1.75	1.5	1	2.33	1.8	2.33	2	

# **Structural Bioinformatics Lab**

COURSE CODE:18B17BI573 COURSE CREDITS: 1 CORE/ELECTIVE: CORE

: 0-0-2

### Pre-requisite: Structural Biology

### **Course Objectives:**

To develop the ability to design, predict, analyze and compare the protein structures as well as predict the function of target proteins.

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
	Understanding the fundamental concepts of structural biology (chemical	Familiarity
CO1	building blocks, structure, superstructure, folding, etc.)	
CO2	To Understand and use structural databases and software for structure	Familiarity
	visualization	
CO3	To understand the algorithms used in Structure determination and quality	Assessment
	assessment	
	To perform protein structure comparison and the hierarchical nature of	Usage
CO4	biomacromolecular structure classification	
CO5	To understand the methodology of protein structure prediction and	Assessment
	assessment	
CO6	To understand the methodology of sequence- and structure-based	Assessment
	functional site prediction	

## List of Experiments:

S.No	Description	Hours
1	Homology modeling using Swiss-Modeller and Modeler standalone software	2
2	Searching and analyzing SCOP and CATH database; analysis of protein structure (protein ligand and protein-protein complexes) using pdbsum, ligplot and dimplot	2
3	Prediction protein secondary structure using above methods and comparison, calculating Q3 value and SOV score.	2
4	Prediction of tertiary structure of protein sequences (using Modeller and GenThreader)	2
5	Error estimation and precison of predicted protein structures (Procheck, What IF, Errat, Verify3D, etc.)	2
6	Comparing protein structures (using CE, DALI, Comparer, SARF2, SSAP, VAST) and statistical analysis of results	2
7	Secondary structure assignment of protein structure using DSSP, STRIDE, DEFINE and P-Curve and statistical analysis of results	2

8	Predicting structural domains (using PRODOM) and binding sites (acSite, SiteMatch, SiteFinder, etc.)	2
9	Ab initio prediction of various energy components of protein structure and validation of structure	2
10	Prediction of PBSA and GBSA energy components using molecular simulation technique	2
11	Filling of gaps in protein structure, energy minization and validation of protein structure	2
12	Setting of MD simulation job of protein structure, interpretation of results and refinement of structure.	2
Total L	ab hours	24

### **Suggested/Resources:**

- 1. Structural Bioinformatics (2nd Edition), Jenny Gu (Editor), Philip E. Bourne (Editor)
- 2. D.W. Mount Bioinformatics: Genome and Sequence Analysis: (2001) Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York.
- 3. Molecular Modeling: Principles & Applications, Andrew R. Leach, Prentice Hall

#### **EvaluationScheme:**

1	Mid Sem. Evaluation	20 Marks
2	End Sem. Evaluation	20 Marks
3	Lab Assessment	60 Marks
	Total	100 marks

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	Average
C01	3	3	3	3	2	2	1	1	1	1	1	1	1.83
CO2	3	3	3	3	3	1	1	1	1	1	1	3	2.00
CO3	3	3	2	3	2	3	2	1	1	1	2	1	2.00
CO4	3	3	3	2	3	2	1	1	1	1	1	1	1.83
CO5	2	2	3	3	3	3	1	1	1	1	1	1	1.83
CO6	2	3	3	3	2	2	2	2	2	2	2	2	2.25
Average	2.67	2.83	2.80	2.80	2.60	2.20	1.20	1.00	1.00	1.00	1.20	1.40	

# **Machine learning for Bioinformatics**

COURSE CODE:18B11BI611

COURSE CREDITS: 3

CORE/ELECTIVE: CORE

L-T-P: 3-0-0

Pre-requisite: Molecular biology, Python

### **Course Objectives:**

- 1. Learn what is machine learning
- 2. Learn algorithms used in machine learning.
- 3. Learn how to implement machine learning for biological problems.
- 4. Apply machine learning to practical projects.
- 5. Use machine learning and data mining in one project.

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	Different types of machine learning and its utility in bioinformatics	Familiarity
CO-2	Application of Hidden Markov Model and Artificial neural networks to different types of bioinformatics data	Assessment
CO-3	Determination of Bayesian Network (BN) from expression data.	Assessment
CO-4	Application of symbolic machine learning (SML) methods to predict cleavage site of HIV- protease from training data of positive and negative cases.	Assessment
CO-5	Optimization of weights in a supervised and unsupervised neural network, and application of supervised learning to predict sub- cellular localization of a protein.	Assessment
CO-6	Application of stochastic context-free grammar (SCFG) to predict RNA secondary structure.	Assessment

#### **Course Contents:**

Unit	Contents	Lectures required					
1	1 Introduction: Overview of intelligent systems and machine learning						
2	Hidden Markov Model (HMM): Viterbi algorithm, Forward algorithm,	10					
	Backward algorithm, Profile-HMM, Baum-Welch algorithm to optimize HMM- profile, Multiple alignment and database searching using profile- HMM						
3	<b>Symbolic Machine Learning:</b> Nearest neighbour approach to predict secondary structure, Decision tree methods, Identification tree methods	6					
4	<b>Bayesian Network (BN):</b> Calculation of statistical significance by using Bayesian methods, Factorization and Markov blanket rule, d-separation, Equivalence classes, Learning of Bayesian network, Learning of Gaussian network	9					
5	Artificial Intelligence (AI):Search strategies, logic, deduction, and pathways comparison	4					

6	Artificial Neural Network (ANN):Basics and introduction to terminologies, Supervised and non-supervised learning, Feed forward back propagation error method, Application of ANN methods: Protein sub-cellular localization and secondary structure prediction	7
7	<b>Stochastic Context Free Grammar (SCFG):</b> Transformational grammar, Parsing, Chomsky hierarchy (regular, context-free, context-sensitive, and unrestricted grammar), Automata, Context-free grammar, Application of SCFG for prediction of secondary structure of RNA	5
Total	Lectures	42

- a. R. Durbin, S. Eddy, A. Krogh, and G. Mitchison (1998), Biological Sequence Analysis: Probabilistic Models of Proteins and Nucleic Acids. Cambridge University Press
- b. Edward Keedwell and Ajit Narayanan (2005), Intelligent Bioinformatics: The Application of Artificial Intelligence Techniques to Bioinformatics Problems, Wiley
- c. P Baldiand S Brunak, BIOINFORMATICS: The Machine Learning Approach

### Suggested Reference Book(s):

- 1. Olson et al., 2018. Data-driven advice for applying machine learning to bioinformatics problems
- 2. Husmeier D, Dybowski R, and Roberts S (2005), Probabilistic Modeling in Bioinformatics and Medical Informatics, Springer
- 3. Nat Cell Biol. 2001 Aug;3(8):E190-5. Review. PubMed PMID: 11483980
- 4. Kim JB, Porreca GJ, Song L, Greenway SC, Gorham JM, Church GM, Seidman CE, Seidman JG. Polony multiplex analysis of gene expression (PMAGE) in mouse hypertrophic cardiomyopathy. Science. 2007 Jun 8;316(5830):1481-4. PubMed PMID: 17556586
- 5. MacBeath G, Schreiber SL. Printing proteins as microarrays for high-throughput function determination. Science. 2000 Sep 8;289(5485):1760-3. PubMed PMID: 10976071.
- Shankar J, Wu TD, Clemons KV, Monteiro JP, Mirels LF, et al. (2011) Influence of 17b-Estradiol on Gene Expression of Paracoccidioides during Mycelia-to- Yeast Transition. PLoS ONE 6(12): e28402. doi:10.1371/journal.pone.0028402
- 7. Mary V. Relling, William E. Evans Nature. Author manuscript; available in PMC 2016 Jan 13.
- 8. Published in final edited form as: Nature. 2015 Oct 15; 526(7573): 343–350. doi: 10.1038/nature15817

#### **Other useful resource(s):**

1. Link to NPTEL course contents: https://nptel.ac.in/courses/106104019/

2.Link to topics related tocourse:

- i. <u>https://www.advancedsciencenews.com/machine-learning-for-bioinformatics-and-neuroimaging/</u>
- ii. https://www.tutorialspoint.com/artificial intelligence/artificial intelligence neural networks.htm

iii. https://www.analyticsvidhya.com/blog/2017/09/understaing-support-vector-machine-example-code/ EvaluationScheme:

S. No	Exam	am Marks		Coverage / Scope of Examination				
1	T-1	15	1 Hour	Syllabus covered upto T-1				
2	T-2	25	1.5 Hours	Syllabus covered upto T-2				
3.	T-3	35	2 Hours	Entire Syllabus				
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes&Attendance				

Course outcomes (Machine learning for Bioinformatics)	P0-1	PO-2	PO-3	PO-4	PO-5	PO-6	P0-7	PO-8	PO-9	PO-10	PO-11	PO-12	Average
CO-1	2	2	2	2	2	2	2	2	2	2	1	2	1.92
CO-2	2	2	2	2	2	1	1	2	2	2	1	2	1.75
CO-3	2	2	2	2	2	1	1	1	2	-	2	2	1.73
CO-4	2	2	2	2	2	1	1	1	2	2	-	2	1.73
CO-5	2	2	2	2	2	1	1	1	2	-	-	2	1.7
CO-6	2	2	2	2	2	1	1	1	-	-	-	2	1.67
Average	2.0	2.0	2.0	2.0	2.0	1.16	1.16	1.33	2.0	2.0	1.33	2.0	

# Machine learning for Bioinformatics Lab

COURSE CODE:18B17BI671 COURSE CREDITS: 1 CORE/ELECTIVE: CORE

L-T-P: 0-0-2

### Pre-requisite: None

#### **Course Objectives:**

- 1. Develop an understanding of important concepts and their implementation in machine learning in the context of biological problems.
- 2. Implementation in machine learning in the context of biological problems

Course	Outcomes:

S.No.	Course Outcomes	Level of Attainme	ent		
CO1	Implementation of KNN using Perl/Python	Assessme	ent		
CO2	Implementation of ANN using Perl/Python		Assessme	ent	
CO2	Application of Hidden Markov Model for CpG island prediction	Assessme	Assessment		
CO3	Application of HMMER package and Pfam database	Assessment			
CO4	Application of Transformational Grammars in bioinformatics	Assessment			
CO5	Application of SVM in bioinformatics		Assessme	ent	
List of Ex	periments				
S.No	Description	Hou	rs		
1	Calculation of sensitivity, specificity, accuracy for a given classifier	2			
2	Implementation of crisp KNN for a microarray file	2			
3	Implementation of fuzzy KNN for a microarray file 2				
4	Identification tree construction using See5 and Weka	2			

2

2

2

2

2

2

2

24

1. http://hmmer.org/.

**Total Lab hours** 

5

7

8

9

10

11

12

6

2. https://www.cs.waikato.ac.nz/ml/weka/https://nptel.ac.in/courses/106104019/26

Implementation of perceptron on LOGIC GATES

Calculation of pseudo amino acid composition

Implementation of ANN using SNNS software

HMM using HMMER package

Stochastic context free grammar

Calculation of AAC and DPC for SVM and ANN input files

Implemenation of SVM using SVM-light, LIBSVM and Weka

Implementation of HMM for prediction of CpG islands

3. https://www.rulequest.com/download.html

## **EvaluationScheme:**

1	Mid Sem. Evaluation	20 Marks
2	End Sem. Evaluation	20 Marks
3	Lab Assessment	60 Marks
	Total	100 marks

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	<b>PO7</b>	PO8	PO9	PO10	PO11	PO12	Average
C01	3	3	3	3	2	2	1	1	1	1	1	1	1.83
CO2	3	3	3	3	3	1	1	1	1	1	1	3	2.00
CO3	3	3	2	3	2	3	2	1	1	1	2	1	2.00
CO4	3	3	3	2	3	2	1	1	1	1	1	1	1.83
CO5	2	2	3	3	3	3	1	1	1	1	1	1	1.83
Average	2.67	2.83	2.80	2.80	2.60	2.20	1.20	1.00	1.00	1.00	1.20	1.40	

# **Computer Aided Drug Design**

### COURSE CODE: 18B11BI612

### COURSE CREDITS: 3

### CORE/ELECTIVE: CORE

: 3-0-0

### Pre-requisite: None

### **Course Objectives:**

1. To design potential lead molecules against any disease that may be explored further as a potential candidate for the drug development.

### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	Feasibility study of a drug development project	Familiarity
CO-2	Design and optimize lead molecules against drug target, and using ligand- basedapproach	Usage
со-3	Determination of pharmacophore from lead molecules and active sites and use of pharmacophore for lead discovery	Usage
CO-4	Development of potential drug molecule and pharmacophore databases for virtualscreening	Assessment
CO-5	Use of molecular fragments for lead discovery and implementation of statisticalapproaches for lead molecule discovery	Usage
CO-6	Bioavailability prediction of a drug and working capability in drug designing softwarelike, Discovery Studio and molecular dynamics software like AMBER 8.0, On-line tools, etc.	Assessment

### **Course Contents:**

Unit	Contents	Lectures required
1	<b>Introduction:</b> Drug design: A billion dollar baby and drug design techniquesteam work, Economic factors involved in drug design, Irrational vs.	4
	rational approaches, Drug target identification & computer-aided drug design processes, Case study related to drug target identification (Viral Targets -HIV Example)	
2	<b>Computational Approaches to Drug Design:</b> Structure-based (receptor fitting)and ligand-based (receptor mapping) molecule design, lead molecules design in a research environment (crossing the barriers), tools used in both environments	3
3	<b>Receptor Fitting (Lead discovery&amp; refinement)</b> : Utility, Binding-site predictions: Stereoelectronic factors, receptor flexibility, tight binding; Docking: Introduction, search algorithms, scoring (MM, Grid, etc.), validation of results, comparisons of search and scoring methods; Docking processes and analysis of results; <i>De novo</i> design, database searching & high throughput virtual screening(HTVS); and applications. Introductions to docking & molecular modeling packages (DS Studio; Schrodinger Inc, Ligbuilder, etc.)	9
4	<b>Receptor Fitting (Lead optimization)</b> : Molecular simulation methods used for binding and free energy calculation, Calculation on Free Energy Perturbation(FEP) of 1. Thermolysin with 2 ligands, Molecular mechanics Poisson-Boltzmann surface area methods: molecular basis of HIV protease drug	4

	resistance	
5	<b>Receptor Mapping (Pharmacophore)</b> : The pharmacophore concept, Determination of pharmacophore from a set of active molecules, Design of pharmacophore using various algorithms, Creating pharmacophore model from active site, Practical utility (searching compound databases) and A case study of new lead design	5
6	<b>Chemoinformatics:</b> Introduction, representing 2D & 3D structures, 2D chemical database applications & molecular descriptors and their classifications, database searching and applications in CADD	4
7	<b>Receptor Mapping (Quantitative structure activity relationship</b> (QSAR)): QSAR methodology, biological and physicochemical parameters, feature selection(PLS, PCA, MLR, etc.), model building and validation, QSAR applications in drug design, Quantitative structure-property relationships (QSPR), CoMFA, 3D and nD-QSAR methods	6
8	<b>Fragment-based Lead Discovery</b> : Fragment and substructure discovery and evaluation, virtual fragment scanning (trends, applications and web-based tools) & capture methods for fragment-based discovery	4
9	<b>ADMET</b> : Oral bioavailability, drug half-life in the bloodstream, BBB permeability, toxicity, Lipinski rule of five, The impact of physiochemical properties on the control of drug-like properties.	4
Tota	al Lectures	42

- 1. David C Young : Computational Drug Design (A guide for computational and medicinal chemists) Wiley & Sons, Inc., New Jersey, USA
- 2. Holtje H.-D, Sippl W., Rognan D. and FolkersG. : Molecular Modeling, Basic Principles and Applications Wiley-VCHGmbH& Co. KGaA
- 3. Leach AR : Molecular Modeling: Principles and Applications: Prentice Hall, Edinburg UK.
- 4. Zartler ER & Shapiro MJ : Fragment-based Drug Discovery (A practical approach), Wiley & Sons, Inc., West Sussex, UK Flower DR : Drug design: cutting edge approaches, RSC publication, Cambrige, UK

### Suggested Reference Book(s):

- 1. Merz KM, D Ringe, : Drug Design: Structure and Ligand-based Approaches. Reynolds CH Cambridge University Press
- 2. Opera TI :Chemoinformatics in Drug Discovery, Wiley-VCH, GMBH
- 3. Hubbard RE : Structure-based drug discovery (An overview), RSC publication, Cambride, UK

#### **EvaluationScheme:**

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3.	T-3	35	2 Hours	Entire Syllabus
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes & Attendance

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	Average
C01	3	3	3	3	2	2	1	1	1	1	1	1	1.83
CO2	3	3	3	3	3	1	1	1	1	1	1	3	2.00
CO3	3	3	2	3	2	3	2	1	1	1	2	1	2.00
CO4	3	3	3	2	3	2	1	1	1	1	1	1	1.83
C05	2	2	3	3	3	3	1	1	1	1	1	1	1.83
C06	2	3	3	3	2	2	2	2	2	2	2	2	2.25
Average	2.67	2.83	2.80	2.80	2.60	2.20	1.20	1.00	1.00	1.00	1.20	1.40	

## **Computer Aided Drug Design Lab**

COURSE CODE: 18B17BI672 COURSE CREDITS: 1 CORE/ELECTIVE: CORE L-T-P: 0-0-2

# Pre-requisite: None

### **Course Objectives:**

1. To design potential lead molecules against any disease that may be explored further as a potential candidate for the drug development.

### Course Outcomes:

S.No.	Course Outcomes	Level of Attainment
CO1	Feasibility study of a drug development project	Familiarity
CO2	Design and optimize lead molecules against drug target, and using ligand- basedapproach	Usage
CO3	Determination of pharmacophore from lead molecules and active sites and use of pharmacophore for lead discovery	Usage
CO4	Development of potential drug molecule and pharmacophore databases for virtualscreening	Assessment
CO5	Use of molecular fragments for lead discovery and implementation of statistical approaches for lead molecule discovery	Usage
CO6	Bioavailability prediction of a drug and working capability in drug designing softwarelike, Discovery Studio and molecular dynamics software like AMBER 8.0, On-line tools, etc.	Assessment

#### **List of Experiments**

S.No	Description	Hours				
1	Installation of various drug design software and assignment 'Project'	2				
2	Generation of 3D optimized structure of a "Ligand" molecule					
3	Preparation of target and ligand molecules for docking					
4	"Virtual library Preparation" of lead molecules					
5	Docking of ligands into a receptor (active site)	2				
6	Flexible docking of ligand with target	2				
7	Fragment docking using ' <i>De Novo</i> ' Receptor and ' <i>De Novo</i> ' Links (LUDI algorithm)	2				
8	Pharmacophore modeling of ligands	2				
9	Pharmacophore-based database searching and <i>de novo</i> design of ligand against an active site	2				
10	Development of 3D QSAR model by using "Discovery Studio"	2				
11	ADME property and toxicity predictions of lead molecule (using TOPKAT)	2				
12	Energy minimization and molecular dynamics (MD) target molecule by using "Simulation" module of "Discovery Studio"	2				
13	Estimates binding free energy of ligands and receptor using	2				
	CHARMmimplicitsolvation models					
Total La	b hours	26				

### **Suggested Books/Resources:**

1. David C Young : Computational Drug Design (A guide for computational and medicinal chemists) Wiley & Sons, Inc., New Jersey, USA

- 2. Holtje H.-D, Sippl W., Rognan D. and FolkersG. : Molecular Modeling, Basic Principles and Applications Wiley-VCHGmbH& Co. KGaA
- 3. Leach AR : Molecular Modeling: Principles and Applications: Prentice Hall, Edinburg UK.
- 4. Accelrys: User Manuals Discovery Studio.
- 5. AMBER : AMBER 11 Users' Manual, Scripps Research Institute, USA
- 6. GROMACS:Gromacs User Manual

#### **EvaluationScheme:**

1	Mid Sem. Evaluation	20 Marks
2	End Sem. Evaluation	20 Marks
3	Lab Assessment	60 Marks
	Total	100 marks

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	<b>PO7</b>	PO8	PO9	PO10	PO11	PO12	Average
CO1	3	3	3	3	2	2	1	1	1	1	1	1	1.83
CO2	3	3	3	3	3	1	1	1	1	1	1	3	2.00
CO3	3	3	2	3	2	3	2	1	1	1	2	1	2.00
CO4	3	3	3	2	3	2	1	1	1	1	1	1	1.83
CO5	2	2	3	3	3	3	1	1	1	1	1	1	1.83
CO6	2	3	3	3	2	2	2	2	2	2	2	2	2.25
Average	2.67	2.83	2.80	2.80	2.60	2.20	1.20	1.00	1.00	1.00	1.20	1.40	

# **Advanced Algorithms for Bioinformatics Lab**

COURSE CODE:18B17BI673

COURSE CREDITS: 1

CORE/ELECTIVE: CORE

L-T-P: 0-0-2

**Pre-requisite:** Basics of algorithms and programming, data structures. Some knowledge of objectoriented technology and is also desirable

### **Course Objectives:**

- 1. Develop the ability to design, implement and manipulate algorithms.
- 2. Develop computer programs for Bioinformatics solutions to life and health science problems.
- 3. Apply programming concepts to various biological examples and real life applications.

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO1	Able to understand algorithmic principles	Familiarity
CO2	To write programs for specific computational biology problems	Assessment
CO3	Analyze problems in biology and able to design new protocols and algorithms for biological data analysis	Assessment
CO4	Able to analyze biological data through programs	Assessment
CO5	Implement algorithms for bioinformatics problems and their assessments	Assessment and Usage

### List of Experiments

S.No	Description	Hours
1	Program to solve the US change problem.	2
2	Program to deal with Tower of Hanoi problem.	2
3	Program to generate Fibonacci series using recursive algorithm and few other programs.	2
4	Program to generate distinct sub-strings in a given DNA sequence using combinatorial and other methods.	2
5	Program to generate palindrome of a string and for a nucleotide sequence, translation and reverse translation, find out the GC content in a sequence.	2
6	Program to implement dynamic programming to solve local, semi-global, and global alignment of biological sequences.	2
7	Program to generate redundant nucleotide sequences from given amino acid sequence using standard genetic code system and ambiguous character codes.	2
8	Implementation of fragment assembly algorithms to make contigs.	2
9	Program to predict genes using statistical approaches.	2
10	Program to predict genes using similarity based approaches.	2
11	Program to generate restriction map of DNA sequence using Brute force algorithm.	2

12	Program to generate restriction map of DNA sequence using PDP (Partial Digest Problem) algorithm.	2
13	Motif finding algorithms implementations in DNA and Protein sequences.	2
14	RNA structure algorithms and their implementations.	2
Total L	ab hours	28

### Suggested/Resources:

- a. P A Pevzner: Computational Molecular Biology: An algorithmic approach, PHI, 2004.
- b. N C Jones and P A Pevzner: An Introduction to Bioinformatics Algorithms, Ane Books, 2004.
- c. G. Benson and R. Page: Algorithms in Bioinformatics, Springer Verlag, 2004.
- d. C J Date: An Introduction to Database Systems, Addison-Wesley Longman Publishing Co., USA, 1990.
- e. I IMandoiu and A Zelikovsky: Bioinformatics Algorithms: Techniques and Applications, Wiley Interscience Press, 2008.
- f. R Durbin*et al*: Biological Sequence Analysis: Probabistic models of proteins and nucleic acids, Cambridge University press, 1998.

#### **EvaluationScheme:**

1.	Mid Sem. Evaluation	20 Marks
2.	End Sem. Evaluation	20 Marks
3.	Lab Assessment	60 Marks
	Total	100 marks

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	<b>PO7</b>	PO8	PO9	PO10	PO11	PO12	Average
C01	2	2	1	3	2	2	2	1	3	2	3	3	2.17
CO2	3	2	3	2	1	-	-	-	1	1	2	3	2.00
CO3	2	2	2	2	3	2	-	1	2	3	-	1	2.00
CO4	2	3	3	2	1	1	1	-	2	2	2	2	1.91
CO5	3	2	2	2	2	2	-	-	-	2	2	1	2.00
Average	2.4	2.2	2.2	2.2	1.8	1.75	1.5	1	2	2	2.25	2	

# R Language Lab

COURSE CODE:18B17BI674

COURSE CREDITS: 1

CORE/ELECTIVE: CORE

L-T-P: 0-0-2

Pre-requisite: Basic concepts in programming

#### **Course Objectives:**

- 1. Students will learn how to program in R and how to use R for effective data analysis.
- 2. Students will learn how to install and configure software/packages necessary for a statistical programming environment; discuss generic programming language concepts as they are implemented in a high-level statistical language.
- 3. The course covers practical issues in statistical computing which includes programming in R, reading data into R, accessing R packages, writing R functions, debugging, and organizing and commenting R code.
- 4. Topics in statistical data analysis and optimization will provide working biological and real life examples.

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment		
CO 1	Able to apply various approaches in R Software and tools to understand concept of Statistics	Familiarity		
	Use various programming techniques in R to implement algorithmic	Familiarity and		
CO 2	methods from computational biology to describe and analyze problems in biology	Assessment		
CO 3	Implement various packages of R for their application in bioinformatics	Assessment		
CO 4	To understand Bioconductor and its association with Bioinformatics	Assessment		
CO 5	To apply R with universal features through available packages	Usage		

#### List of Experiments

S.No	Description           Introduction of R and R-studio, overview and History of R.			
1				
2	R-console input and evaluation, Data Types, Basic Arithmetic operation.			
3	Reading Tabular data and large tables: File handling.			
4	Textual data formats, Connection: interface to outside world.	2		
5	Sub setting- Basics, lists, Matrices, Partial matching, Removing missing values	2		
6	Vectorized operations.	2		
7	Control structure- introduction, if-else, for loops, while loops, Repeat, Next, Break.			
8	R-functions, Loop function and debugging.	2		
9	Scoping Rules- Symbol binding, rules and optimizing examples, Coding standardDates and times.	2		
10	Bioconductors: Introduction.	2		
11	Simulation in R- str function, Generate random numbers, Simulating linearModels, Random sampling.	2		
12	R-profiler- collect detailed information on how your R functions are running and to identify bottlenecks that can be addressed. The profiler is a key tool in helpingYou optimize your programs.	2		
13	Bioconductors package usage for analysis of genomic data - I	2		
14	Bioconductors package usage for analysis of genomic data - II	2		
Total Lab hours				

## **Suggested/Resources:**

- 1. An Introduction to R
- 2. Quick-R
- 3. <u>R for Beginners</u>
- 4. Kim Seefeld's <u>R-introduction for Biostatistics</u>
- 5. P Dalgaard:<u>Introductory Statistics with R</u>, 2<sup>nd</sup> Edition, Springer Nature, 2008.
- 6. W P Krijnen: Applied Statistics for Bioinformatics using R, 2009.
- 7. R Gentlemen: R programming for Bioinformatics, CRC Press, 2008.

## **EvaluationScheme:**

1	Mid Sem. Evaluation	20 Marks
2	End Sem. Evaluation	20 Marks
3	Lab Assessment	60 Marks
	Total	100 marks

## **Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)**

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	<b>PO7</b>	PO8	PO9	PO10	PO11	PO12	Average
C01	3	2	2	3	2	2	2	1	3	2	3	3	2.33
CO2	3	3	3	2	1	1	-	-	1	1	2	3	2.00
CO3	2	2	2	3	3	2	1	1	3	3	1	2	2.08
CO4	3	3	3	2	2	1	1	-	2	3	3	2	2.27
CO5	3	2	2	2	2	2	-	1	1	2	2	3	2.00
Average	2.8	2.4	2.4	2.4	2	1.6	1.33	1	2	2.2	2.2	2.6	

## Phytopharmaceuticals and Biologicals

COURSE CODE: 18B1WBT531

## COURSE CREDITS: 3

CORE/ELECTIVE: ELECTIVE

L-T-P: 3-0-0

## Pre-requisite: Basic understanding of biology

## **Course Objectives:**

a. The objective of the course is to develop an understanding and basic knowledge on Indian medicinal herbs, its commercial value, quality control and industrial standards for commercialisation of phytopharmaceuticals and biologicals.

## **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	Able to understand pharmacopoeial standards of ayurvedic products	Familiarity
CO-2	Able to understand the phytopharmaceuticals, monographs and quality control of medicinal herbs	Usage
CO-3	Able to understand the extraction and evaluation process of phytopharmaceuticals	Technical
CO-4	Able to understand pharmacopoeial standards of Indian Pharmacopoeia	Familiarity
CO-5	Able to understand the monographs, specifications and quality control of biologicals	Usage

## **Course Contents:**

Unit	Contents	Lectures required
1	Intoduction to phytopharmaceuticals – medicinal herbs & its importance and biologicals. Over view of Ayurvedic and Indian Pharmacopoeia	4
2	Pharmacopoeial standards of Ayurvedic (Used by manufacturers, regulators and other stakeholders for quality control of medicinal herbs and finished products against internationally recommended specifications).	7
3	Monographs of medicinal herbs, specifications and standards for identification, evaluation, processes and clinical applications	8
4	Evaluation of physiochemical parameters of herbal drugs, extraction, Identification & Assay of Herbals Drugs	6
5	Pharmacopoeial standards of Indian Pharmacopoeia (Used by manufacturers, regulators and other stakeholders for quality control of active pharmaceutical ingredients (APIs) and finished products against internationally recommended specifications).	3
	Monographs on Blood and Blood related products, Monographs on Human Vaccines (The specifications cover the various tests for critical quality parameters Approved in Academic Council held on 25.10.2018	

6	of the vaccine, procedures and acceptance criteria)	7
7	Monographs of Erythropoietin Injection, Interferon Injection, streptokinase solution, Human Insulin, etc. Bacterial Endotoxin Test, Sterility Test, Test for Microbial Contamination, etc.	7
Total lect	ures	4 2

- 1. The Ayurvedic Pharmacopoeia Of India, First Edition, Published By Pharmacopoeia Commission For Indian Medicine & Homoeopathy Ghaziabad (2016)
- 2. Indian Pharmacopoeia published by the Indian Pharmacopoeia Commission (IPC)

## **EvaluationScheme:**

S. No	Exam	Marks	Duration	Coverage / Scope of Examination			
1	T-1	15	1 Hour.	Syllabus covered upto T-1			
2	T-2	25	1.5 Hours	Syllabus covered upto T-2			
3.	T-3	35	2 Hours	Entire Syllabus			
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes&Attendance			

## **Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)**

Course outcomes (Phytopharmaceuti cals and Biologicals)	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10	PO-11	PO-12	Average
CO-1	2	2	2	2	2	2	2	2	2	2	1	2	1.9
CO-2	2	2	2	2	2	1	1	2	2	2	1	2	1.75
CO-3	2	2	2	2	2	1	1	1	2	2	2	2	1.75
CO-4	2	2	2	2	2	1	1	1	2	2	1	2	1.6
CO-5	2	2	2	2	2	1	1	2	2	2	1	2	1.75
Average	2.0	2.0	2	2	2	1.25	1.25	1.5	2.0	2	1.25	2.0	

## **Bioenergy and Biofuels**

COURSE CODE: 18B1WBT634

## COURSE CREDITS: 3

## CORE/ELECTIVE: ELECTIVE

## L-T-P: 3-0-0

Pre-requisite: Microbiology, Bioprocess Engineering, Downstream Processing, Bioresource Technology

## **Course Objectives:**

The shortage of fossil fuels and its environmental consequences, Bioenery and Biofuel technology seems to be a alternative for generation of energy and fuels. This sector facing various technical, process and social problems for implementation. Based on these aspects the objectives of the course are framed as

- 1. Introduction of existing and possible Bioenergy and Biofuels technoloies
- 2. Discussion of technical, process and economic issues related to first, second and third generation biofuels along with Physico chemical techniques

## **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	Advantages and disadvantages of Bioenergy and Biofuels over fossil fuels	Familiarity
CO-2	Technical barriers in Bioenergy and Biofuel Technology	Assessment
CO-3	Whole biorefinery approaches for economical implementation into the market	Usage
CO-4	Conversion technologies of waste to Biofuels, Bioproducts, and Bioenergy	Usage
CO-5	Conversion of waste and Mixed feedstock to Biofuels, Bioenergy and Bioproducts	Usage

## **Course Contents:**

Unit	Contents	Lectures required
1	<b>Introduction to Biofuels and Bioenergy:</b> Definition, Global Energy Outlook, Sustainability, Biomass Feedstocks, Processes and Technologies, Environment and Ecology	4
2	<b>Crop Oils, Biodiesel, and Algae Fuels:</b> Vegetable Oils, Algae Oil Extraction of Straight Vegetable Oil, Manufacture of Biodiesel	12
3	<b>Ethanol from Corn and Lignocellulosics:</b> Fuel Ethanol from Corn, Corn Ethanol as Oxygenated Fuel, Chemistry of Ethanol Fermentation, Corn-to-Ethanol Process Technology, By- Products/Coproducts of Corn Ethanol, Ethanol as Oxygenated and Renewable Fuel, Ethanol Vehicles, Lignocellulose and Its Utilization, Lignocellulose Conversion, Agricultural Lignocellulosic Feedstock, Cellulosic Ethanol Technology; Energy Balance for Ethanol Production from Biomass, Process Economics and Strategic Direction.	12
4	<b>Fast Pyrolysis and Gasification of Biomass:</b> Biomass and Its Utilization, Analysis and Composition of Biomass, Chemistry of Biomass Gasification, Fast	7

	Pyrolysis of Biomass, Biomass Gasification Processes, Utilization of Biomass Synthesis Gas	
5	<b>Conversion of Waste to Biofuels, Bioproducts, and Bioenergy &amp; Mixed Feedstock:</b> Types of Waste and Their Distributions, Strategies for Waste Management, Waste Preparation and Pretreatment for Conversion, Technologies for Conversion of Waste to Energy and Products, Economic and Environmental Issues Related to Waste Conversion, Future of the Waste Industry, Advantages and Disadvantages of Mixed Feedstock, Transportation, Storage, and Pretreatment, Gasification Technologies, Liquefaction Technologies, Future of Mixed Feedstock.	7
Total Le	ectures	42

- 1. Biofuels and Bioenergy: Processes and Technologies by Sunggyu Lee and Y. T. Shah, CRC Press
- 2. Bioenergy and Biofuel from Biowastes and Biomass by Samir K. Khanal, Rao Y. Surampalli, Tian C. Zhang, Buddhi P. Lamsal, R. D. Tyagi and C.M. Kao, ASCE Publishers .

## **Suggested Reference Book(s):**

1. Review and research articles from Science Direct, Springer, Wiley and PubMed Publishers

## **EvaluationScheme:**

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3.	T-3	35	2 Hours	Entire Syllabus
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes&Attendance

## **Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)**

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	Average
CO1	3	3	3	3	1	3	1	2	1	2	1	3	2.17
CO2	3	3	3	3	2	2	1	2	1	2	-	3	2.27
CO3	3	3	3	3	2	2	1	2	1	2	-	3	2.27
CO4	3	3	3	3	1	2	2	3	2	2	2	3	2.42
CO5	3	3	3	3	1	2	3	3	3	2	2	3	2.58
Average	3.00	3.00	3.00	3.00	1.40	2.20	1.60	2.40	1.60	2.00	1.00	3.00	

## **Intellectual Property Rights and Commercialization**

COURSE CODE: 18B1WBT734

COURSE CREDITS: 3

CORE/ELECTIVE: ELECTIVE

L-T-P: 3-0-0

## **Pre-requisite:** None

### **Course Objectives:**

- b. To provide an insight and understanding about different aspects of protection of inventions and research developments
- c. Learn about procedures for filling protection through Intellectual Property Rights.
- d. To provide scopes of protection of diverse intellectual properties and its commercialization for socio-economic improvement.

## **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	To enable students with basic concepts and knowledge of intellectual property rights.	Awarness
CO-2	To apply and execute different types of IP protection in research and academics.	Assessment and technical skills
CO-3	Able to understand about the mechanisms of different IP protections, registrations and applications	Technical
CO-4	To be capable of tackling issues related to IP and its commercialization	Assessment
CO-5	Able to learn the strategies for effective IP management and commercialization	Analytical skills
CO-6	To apply the knowledge of IPR for the benefit generation and for mass utilization	Usage

## **Course Contents:**

Unit	Contents	Lectures required
1	Introduction: Introduction of Intellectual properties and rights conferred	4
	. Integration of Intellectual Property, Bioethics and Biosafety for	Т
	biological and applied sciences in research and academia.	
2	<b>Types of IP tools</b> :Different types of IPR( Patents, copyrights and related rights, Trademark, Tradesecret, Integrated circuit layout, Geographical indications, Traditional language and the patents of	10
	indications, Traditional knowledge, Industrial designs and PBR) Drafting Patent Application and Documentation	
	Revocation of Patent, Litigation and Infringement	
	Rationale of different IPR, their mechanism of protection and provisions in Law	
3	<b>International Agreements</b> and Treaties:International IP treaties (Madrid Agreement, Trademark law treaty, Patent Law treaty etc.)	8
	WIPO, EPC, WTO, and TRIPS. International agreements relevant to biotechnology-associated IP	v

4		
	<b>Commercialization</b> : Methods of commercialization,Impact of commercialization. Financing	6
5	<b>IP Management for value addition:</b> Strategies for IP Management and commercialization. IP audit, IP insurance	
	Bioentreprenuership management	4
6	<b>Licensing/Assignment</b> : Types of licensing and modes to carry out, Assignments and its benefits, Compulsory Licensing	8
	Commercialization for social and economic prosperity with case studies	~
Total le	ctures	42

## Methodology

The course will be covered through lectures, presentations and vedios. Apart from discussions on topics covered through lectures and assignments, students have to carry out research paper analysis.

## Suggested Text Book(s):

- i. Intellectual Property Rights & Copyright By Bouchoux.
- ii. Intellectual Property Licensing Strategies by Thompson Reuters

**Suggested Reference Book(s):** 

- 1. Intellectual Property Rights, the WTO and Developing Countries: The TRIPS ...Book by Carlos María Correa
- 2. Perspectives on Commercializing Innovation by F. Scott Kieff (Editor), Troy A. Paredes (Editor

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3.	T-3	35	2 Hours	Entire Syllabus
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes & Attendance

## Course Outcomes (COs) contribution to the Programme Outcomes(POs)

Course outcomes (Intellectual Property Rights & Commercializat ion)	PO-1	PO-2	PO-3	PO-4	PO-5	9-0-6	PO-7	PO-8	PO-9	PO-10	PO-11	PO-12	Average
CO-1	2	2	2	2	2	1	1	1	1	1	2	2	15

CO-2	2	2	1	2	2	2	1	1	1	1	1	2	1.5
CO-3	2	2	2	2	3	1	1	1	2	2	1	2	1.7
CO-4	2	2	3	3	2	1	1	1	2	2	2	2	1.9
CO-5	2	2	2	2	2	1	2	1	1	2	1	2	1.6
CO-6	2	2	2	2	2	2	1	1	2	2	2	2	1.8
Average	3.4	3.4	2	2.1	2.1	1.3	1.1	1	1.5	1.6	1.5	2	

## **Peptide Therapeutics**

COURSE CODE: 18B1WBT631

## COURSE CREDITS: 3

## CORE/ELECTIVE: ELECTIVE

## L-T-P: 3-0-0

## Pre-requisite: General Chemistry

## **Course Objectives:**

- 1. To develop an understanding of important concepts and design aspects of peptides
- 2. To learn various therapeutic applications of peptides.
- 3. Apply basic knowledge to design peptides for various therapeutic purposes

## **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	Able to understand various peptide design consideration and other important structural aspects of peptides	Familiarity
CO-2	Able to understand and learn the concept of solid phase peptides synthesis	Familiarity
CO-3	To develop methods of peptides and proteins for their quality control and apply them in handling Therapeutic peptides and proteins	Assessment
CO-4	To understand the mechanism of action of Antibiotic, Anticancer, Antihypertensive and Opioid peptides	Usage
CO-5	To develop a strong foundation therapeutic peptide design and their applications	Usage

## **Course Contents:**

Unit	Contents	Lectures required
1	Peptides, synthetic peptides & their classification based on structure, engineering bioactive peptide based therapeutic molecules,	7
2	Principle and practice of solid phase peptide synthesis, solid           support, protection scheme, peptide acid and amide, Purification of peptides, quality control of peptides	7
3	Antimicrobial host defense peptides, Anticancer peptides, Opioid Peptides, Antihypertensive Peptides, Peptides in clinical trial ,chemical biology of Oxytocin, valinomycin and enkephalins	18
4	Preformulationstudies, Formulation development, Aggregation in protein formulation, novel formulation approaches, Lyophilization, Pharmaceutical Processing, and Handling of Therapeutic Peptides and Proteins	6
5	Circular dichroism, UV, IR, Mass and fluorescence spectroscopy of peptides	4
	Total lecture	42

- 1. Ajay K Banga, "Therapeutic peptides and protein: formulation processing and delivery system, Second edition, Taylor and Francis.
- 2. Lehninger Principles of Biochemistry Cox, M.M. and Nelson, D.L. and Lehninger A. L. 4th edition.
- 3. Biochemistry- J.M. Berg, J.L.Tymoczko, and LubertStryer; 5<sup>th</sup> edition W.H. Freeman and Company, New York, USA.

## **Suggested Reference Book(s):**

1. Gregory A. Grant, "Synthetic peptides A Users Guide" 2<sup>nd</sup> ed. W. H. Freeman and Company

## **EvaluationScheme:**

Assessment	Max. marks	Duration	Course Covered
T1 Test	15	1 hr.	Syllabus covered upto T-1
T2 Test	25	1.5 hrs.	Syllabus covered upto T-2
End Term Test	35	2 hrs.	Whole Syllabus
TeacherAssessment(Based on Assignments, quizzes etc.)	25	Whole Sem	Inform class time to time
Total	100		

## **Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)**

Peptide Therapeutics	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10	PO-11	PO-12	Average
CO-1	2	3	2	2	2	1	-	-	2	3	3	3	1.92
CO-2	2	3	3	3	2	2	-	-	2	3	3	3	2.17
CO-3	3	3	3	3	3	2	-	-	2	3	3	3	2.33
CO-4	3	3	2	3	3	2	-	-	2	3	3	3	2.25
CO-5	3	3	2	3	2	2	-	-	2	3	3	3	2.17
Average	2.60	3.00	2.40	2.80	2.40	1.80	0.00	0.00	2.00	3.00	3.00	3.00	

## Nano-Biotechnology

COURSE CODE: 18B1WBT633

COURSE CREDITS: 3

CORE/ELECTIVE: ELECTIVE

L-T-P: 3-0-0

## Pre-requisite: Basic of Physics, Chemistry and Biology

## **Course Objectives:**

- 1. Introduction to Nanomaterial and various material used for obtaining nano-materials
- 2. Learn various approaches or methods used for nanomaterial synthesis.
- 3. To learn various analytical techniques used for nanomaterial characterization.
- 4. Learn various applications of nanomaterial in health care, agriculture and environmental monitoring

## **Course Outcomes**

S. No.	Course Outcomes	Level of Attainment
CO-1	Introduction to Nano (Basics to Nanoscience and Nanotechnology)	Familiarity
CO-2	Introduction to the two approaches (bottom up and top down) followed for the synthesis of nanomaterial and fundamental properties of Nano-materials(Nano-effect)	Assessment & Technical
CO-3	Introduction to various technique used for the characterization of nanostructures and nanomaterial.	Assessment & Technical
CO-4	Fundamental understanding of nanomaterial/nano- biotechnological application in health and disease.	Usage
CO-5	Fundamental understanding of nanomaterial/nano- biotechnological application in Environment and food - detection and mitigation	Usage

**Course Contents:** 

Unit	Contents	Lectures required
1	<b>Introduction, History &amp; Applications:</b> Various definitions and Concept of Nano- biotechnology & Historical background. Fundamental sciences and broad areas of Nano-biotechnology. Various applications of Nano- biotechnology, Cell – Nanostructure interactions	6
2	<b>Synthetic methodologies:</b> Introduction to the two approaches (bottom up and top down) followed for the synthesis of nanomaterials: Lithography method, Electrochemical method, Mechanical Method, Chemical Synthesis, Chemical vapour deposition, Molecular self-assembly, Laser Induced assembly.	10
3	<b>Techniques used for the characterization of nanoparticles:</b> Principles of microscopy-light, electron, fluorescent confocal, scanning and transmission microscopes, different fixation and staining techniques for	13
	EM. Principles of spectroscopy-UV, visible, CD, FTIR, NMR, and ESR spectroscopy, structure determination using X-ray diffraction, analysis using light scattering.	
4	Nano-biotechnologicalapplications in health and disease:Properties of different types of nanoparticles normally used in health and disease. Diagnostics and theranostics application of nanomaterials in health sciences.	6
	Nanobiotechnological applications in Environment andfood - detection and mitigation:Properties of different types of nanoparticles normally used in	

5	environmental and food sciences. Detection and removal of toxic metal ion from polluted sample and detection and removal of pathogen form food sample.	7
Tota	l Lectures	42

- 1. C. A. Mirkin and C. M. Niemeyer. Nanobiotechnology II more concepts and applications. (2007) Wiley VCH.
- 2. P. Boisseau, P. Houdy, M. Lahmani, Nanoscience: Nanobiotechnology and Nanobiology

## Suggested Reference Book(s):

- 1. A. Nouailhat, An Introduction to Nanoscience and Nanotechnology, Wiley
- 2. D.A Phoenix, W. Ahmed, Nanobiotechnology, One Central Press Ltd, UK
- 3. L. Filipponi, D. Sutherland, Nanotechnologies: Principles, Applications, Implications and Hands-on Activities.Directorate- European commission

#### Other useful resource(s):Link to NPTEL course contents

- https://nptel.ac.in/courses/118107015/
- https://onlinecourses.nptel.ac.in/noc17\_bt17/preview
- http://videos.gitam.edu/nptel/nano.html

## **Evaluation Scheme:**

S. No.	Exam	Marks	Duration	Coverage/Scope of
				Examination
1	T-1	15	1 Hour	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3	T-3	35	2 Hours	Entire Syllabus
4	Teaching Assessment	25	Entire	Quiz, Assignment, Attendance,
			Semester	etc.

Course Outcomes (COs) contribution to the programme Outcomes (POs):

Course outcomes (NanoBiotech.)	P0-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10	PO-11	PO-12	Average
CO-1	2	2	1	1	2	3	2	1	1	1	2	2	1.7
CO-2	2	3	2	2	3	2	2	2	1	1	3	2	2.1
CO-3	3	3	2	2	3	2	2	2	1	1	3	2	2.2
CO-4	2	2	3	3	2	3	2	2	2	2	3	2	2.3
CO-5	2	2	3	3	2	3	3	2	2	1	3	2	2.3
Average	2.2	2.4	2.2	2.2	2.4	2.6	2.2	1.8	1.4	1.2	2.8	2.0	

## COURSE CODE: 18B1WBT632

## **COURSE CREDITS: 3**

## CORE/ELECTIVE: ELECTIVE

L-T-P: 3-0-0

## Pre-requisite: Microbiology, Immunology

## **Course Objectives:**

- 1. Learn to define the basic concepts related to infectious diseases, immunology and epidemiology.
- 2. Able to understand the basic forms, functions, behaviour, and diversity of infectious agents and their interactions with the host
- 3. Able to analyse the underlying principles of mode of action and resistance towards the agents used to treat infectious diseases

S.No.	Course Outcomes	Level of Attainment
CO-1	The students would have knowledge of infectious diseases for practical use in medicine and biotechnology.	Familiarity
CO-2	The students would have in-depth knowledge of basic concepts related to infectious diseases, immunology and epidemiology.	Assessment
CO-3	The students would develop knowledge and understanding of the basic form, function, behavior, and diversity of infectious agents and their interaction with the host.	Assessment
CO-4	The students would develop knowledge and skill about important techniques used to study host –pathogen interactions.	Assessment/Usage
CO-5	The students would have sound knowledge of mode of action and resistance towards the agents used to treat infectious diseases.	Assessment/Usage

## **Course Outcomes:**

## **Course Contents:**

Unit	Contents	Lectures required
1	Introduction to infectious diseases Infectious and non- infectious diseases, Epidemiology of infectious diseases. Infectious agents, socio economic	
	impact of infectious diseases	6
2	Host response to infections Bacterial, mycobacterial, viral, helminth, fungal	3
3	Biology of infectious agents Morphology, classification, life cycle, pathogenecity, mechanism of replication in bacteria, viruses, protozoa and fungal pathogens	5
4	Pathology of Infectious diseases Pathogenesis, Clinical pathology, Gross pathology, Microscopic pathology.	4
5	Biology of major Infectious diseases HIV/AIDS, Tuberculosis, malaria, dengue, West Nile virus, chikungunya virus, diarrheal diseases, sexually transmitted infections, influenza, viral hepatitis, Ebola	8
6	Diagnostic techniques for infectious diseases Immunohistology, Immunohistochemistry/In situ hybridization, Polymerase chain reaction based methods, Flow Cytometry.	5
7	Antimicrobials against infectious agents Antimicrobial agents, mechanism of action, Antibiotic resistance, various mechanisms of antibiotic resistance	5
8	Emerging infectious diseases and their Social Impact Emergenceof SARS, Zika virus, Ebola and other newly reported diseases along with their Social	

	and Scientific Impact.	4
9	Biological Weapons Introduction, Concept and examples	2
Total Lectures	42	

- Evolution of infectious disease. Ewald PW. Oxford University Press, New York.1994. ISBN 0-19-511139-7.
- Emerging Infections 1. Scheld WM, Armstrong D and Hughes JM, Editors. ASM Press, Washinton, DC. 1998. ISBN 1-55581-123-3
- 3. Emerging Infections 2. Scheld WM, Craig WA and Hughes JM, Editors. ASM Press, Washington, DC. 1998. ISBN 1-55581-141-8.
- Emerging Viruses. Morse SS, Editor. Oxford University Press, New York. 1993. ISBN 0-19-510-484-6.
- 5. Modern Infectious Disease Epidemiology: Concepts, Methods, Mathematical Models, and Public Health (Statistics for Biology and Health) Kramer; 2010

## **EvaluationScheme:**

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3.	T-3	35	2 Hours	Entire Syllabus
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes&Attendance

## **Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)**

Course outcomes (Infectious diseases)	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	P0-7	PO-8	6-04	PO-10	PO-11	PO-12	Average
CO-1	2	3	3	3	3	3	3	3	2	2	1	3	2.6
CO-2	3	3	3	3	3	3	3	3	3	2	-	3	2.9
CO-3	1	3	3	3	1	1	3	3	3	3	-	3	2.5
CO-4	3	1	3	3	3	3	2	1	-	-	3	2	2.4
CO-5	1	2	3	3	-	-	3	3	2	2	2	2	2.3
Average	2	2.4	3	3	2.5	2.5	2.8	2.6	2.5	2.25	2	2.6	

## **Genetic Counselling**

COURSE CODE: 18B1WBT831

COURSE CREDITS: 3

CORE/ELECTIVE: ELECTIVE

L-T-P: 3-0-0

Pre-requisite: Molecular Biology, Genetics

## **Course Objectives:**

- 1. To provide an understanding of the basis of genetic counseling, diagnostic testing and management for a variety of types of disorders and also the ethical and legal considerations.
- 2. The students will understand the nature of the non-directive counseling process and the need to educate the patient and family to make informed decisions relating to complex genetic situations.

## **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	To understand basics of genetic counselling, the nature of the directive and non-directive genetic counselling process and to counsel the patients with genetic diseases and help them in decision making.	Familiarity
CO-2	To understandgenetic basis of various diseases (Chromosomal, monogenic and oligogenic disorder).	Assessment
CO-3	To understand gene therapy, its role in genetic disorders and recent developments in gene therapy.	Assessment
CO-4	To understand risk assessment in genetic counseling and ethical issues in genetic counselling.	Usage

## **Course Contents:**

Unit	Contents	Lectures required
1	Genetic Counselling an overview: Counseling, diagnostic	2
1	testing and management of genetic disorders.	
2	Human chromosomal disorders: mutations, types of mutations, chromosomal aberrations, quad screen test, Amniocentesis, karyotyping, Down's syndrome, Patau's syndrome, Edward's syndrome, Turner's syndrome, X- chromosome related syndromes.	6
3	Human allelic diseases (monogenic and oligogenic): Cystic fybrosis. Glucose-6-phospho dehydrogenase deficiency Bradet- Biedl syndrome and some important autosomal recessive and dominant disorders	6
4	<b>Muscle disorders:</b> Duchenne muscular dystrophy, Becker's muscular dystrophy, limb-girdle muscular dystrophies and cardiac muscle disorders.	4
5	Neurological disorder: Alzheimer, Hutington, Parkinson, Lewy body dementia and Schizophrenia	4
6	<b>Genetic basis of neoplastic disorders:</b> Retinoblastoma, Wilms tumor, Colectral cancer, and Blooms syndrome	6
7	Prenatal Genetic counseling	2

8	Gene therapy: Principles of molecular genetic-based therapies and treatment with recombinant proteins or genetically engineered vaccines The technology of classical gene therapy Gene therapy for inherited disorders Gene therapy for neoplastic disorders The ethics of human gene therapy	7
9	Genetic counseling risk assessment	3
1 0	Genetic counseling and ethical issues	2
Total Leo	ctures	42

- 1. Strachan T and Read AP (2010) Human Molecular Genetics -4, Garland Science, 4<sup>th</sup> Ed.
- 2. Pasternak JJ (2005) An introduction to Human Molecular Genetics: Mechanisms of Inherited Diseases. *Hoboken (New Jersey): John Wiley & Sons*, 2<sup>nd</sup> Ed.
- 3. Evans C (2006) Genetic Counselling A psychological approach. New York, NY, US: Cambridge, 1<sup>st</sup> Ed.

## **EvaluationScheme:**

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3.	T-3	35	2 Hours	Entire Syllabus
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes&Attendance

## **Course Outcomes (COs) contribution to the Programme Outcomes(POs)**

Course outcomes (Genetic Counseling)	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10	PO-11	PO-12	Average
CO-1	2	3	3	2	2	2	1	3	2	2	1	3	2.1
CO-2	2	2	3	2	2	1	-	3	2	2	1	3	2
CO-3	2	3	2	2	2	1	-	3	2	3	1	2	2
CO-4	2	3	2	2	2	2	-	3	2	3	1	2	2
Average	2.0	2.75	2.5	2	2	1.5	1	3	2	2.5	1	2.5	

## **Comparative and Functional Genomics**

COURSE CODE: 18B1WBT532

## COURSE CREDITS: 3

#### CORE/ELECTIVE: ELECTIVE

L-T-P: 3-0-0

Pre-requisite: Molecular Biology, Biochemistry

## **Course Objectives:**

- 1. The course is intended to provide thorough understanding of the genomics i.e. modern technologies in whole genome sequencing, genome mining, comparative genomics, global gene function technologies, protein structure & function technologies at the genome level, etc.
- 2. The course will explore that how technological innovations fostered by the Human Genome Project, will lead to significant advances in our understanding of diseases that have a genetic basis and, more importantly, how health care will be delivered from this point forward

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	Students will have a thorough understanding of various genomic technologies such as whole genome mapping & sequencing, genome annotation, global gene cloning and gene expression technologies, comparative genomics, introduction to pharmacogenomics	Familiarity
CO-2	The students will know the vast amount of genome information in publically available databases and how to access and best utilize for practical purposes.	Assessment
CO-3	Able to analyze the gene expression data sets to derive the biologically meaning information	Assessment
CO-4	Able to apply the knowledge of function genomics in public health	Usage

## **Course Contents:**

Unit	Contents	Lectures required						
1	1 Introduction to genomics: Genome organization of Model organism- E. coli, Yeast, Mice, A. thaliana, Human etc. Genome statistics							
2	<b>First and 2<sup>nd</sup> generation sequencing:</b> Sanger sequencing and next generation sequencing; Reverse termination sequencing, Single cell RNA sequencing or single cell RNA sequencing and Applications	8						
3	Comparative genomics:Genome Annotation i.e. Mining Genomic Sequence Data, gene prediction methods, Physical mapping,	8						
	Metagenomics, evolutionary relationship, Genome Analysis, Functional maps (Transcriptome, proteome, metabolome) Metabolic network maps							
4	<b>Functional genomics tools:</b> Hybridization and sequencing based approaches. Serial Analysis of Gene Expression-SAGE, DNA- Microarray, Application of DNA Microarray, cDNA-PCR, etc.	8						
	SNP:SNP Technologies: Platforms & Analysis	7						
5	Haplotyping: Concepts and Applications and relevance in cancer Biology							
6	<b>Regulation of gene expression:</b> Gene Function Technologies (Gene Targeting, Gene Silencing (RNAi), micro RNA-human and Drosophila	4						
7	Biomarkers Pharmacogenomics: Concepts and Applications in Healthcare Role of genotype in drug metabolism Identification & Utilisation of cancer bio-marker	4						
'otal Le	ectures	42						

- 1. Discovering Genomics, proteomics & bioinformatics. Second edition by A Malcolm Campbell, Davidson College; Laurie J. Heyer Davidson College ; With Foreword by Francis S. Collins
- 2. Molecular Biology of the Gene (1987) Watson J. D., Hopking N., Robast J. and Steiz, J.
- 3. BIOINFORMATICS: A Practical Guide to the Analysis of Genes and Proteins (Third edition) Andreas D. Baxevanis& B. F. Francis Ouellette

## Suggested Reference Book(s):

- Ronaghi M. Pyrosequencing sheds light on DNA sequencing. Genome Res. 2001 10. Jan;11(1):3-11. Review. PubMed PMID: 11156611
- 2. Schulze A, Downward J. Navigating gene expression using microarrays—a technology review. Nat Cell Biol. 2001 Aug;3(8):E190-5. Review. PubMed PMID: 11483980
- Kim JB, Porreca GJ, Song L, Greenway SC, Gorham JM, Church GM, Seidman CE, Seidman JG. Polony multiplex analysis of gene expression (PMAGE) in mouse hypertrophic cardiomyopathy. Science. 2007 Jun 8;316(5830):1481-4. PubMed PMID: 17556586
- 4. MacBeath G, Schreiber SL. Printing proteins as microarrays for high-throughput function determination. Science. 2000 Sep 8;289(5485):1760-3. PubMed PMID: 10976071.
- Shankar J, Wu TD, Clemons KV, Monteiro JP, Mirels LF, et al. (2011) Influence of 17b-Estradiol on Gene Expression of Paracoccidioides during Mycelia-to- Yeast Transition. PLoS ONE 6(12): e28402. doi:10.1371/journal.pone.0028402
- 6. Mary V. Relling, William E. Evans Nature. Author manuscript; available in PMC 2016 Jan 13.
- 7. Published in final edited form as: Nature. 2015 Oct 15; 526(7573): 343–350. doi: 10.1038/nature15817

## Other useful resource(s):

1. Link to NPTEL course contents: https://nptel.ac.in/courses/102104056/

## **EvaluationScheme:**

uluulo				
S. No	Exam	Marks	Duration	<b>Coverage / Scope of Examination</b>
1	T-1	15	1 Hour.	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3.	T-3	35	2 Hours	Entire Syllabus
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes&Attendance

## **Course Outcomes (COs) contribution to the Programme Outcomes(POs)**

Course outcomes	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	6-04	PO-10	PO-11	PO-12	Average
CO-1	2	2	2	2	2	2	2	2	2	2	1	2	1.9
CO-2	2	2	2	2	2	1	1	2	2	2	1	2	1.75
CO-3	2	2	2	2	2	1	1	1	2	-	2	2	1.7
CO-4	2	2	2	2	2	1	1	1	2	2	-	2	1.7
Average	2.0	2.0	2	2	2	1.25	1.25	1.5	2.0	2	1.33	2.0	

## **Computational Molecular Evolution**

COURSE CODE: 18B1WBI831

**COURSE CREDITS: 3** 

CORE/ELECTIVE: ELECTIVE

L-T-P: 3-0-0

Pre-requisite: Basic knowledge of computational biology, evolutionary biology, and functional genomics

## **Course Objectives:**

1. The objective of the course is to develop functional and evolutionary genomic understanding of biological entities for various kinds of lineages for biological data types such as DNA, RNA, Genes, and Proteins etc.

## **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	Able to understand the holistic approaches of molecular evolution	Familiarity
CO-2	Combining acquisition, integration and management of experimental evolutionary data with computer aided analysis	Assessment
CO-3	Use various methods from computational genomics and proteomics to learn their functional aspects of controlling biological processes by incorporating evolutionary information	Assessment
CO-4	Able to analyze various kind of biological sequence data and identify their limiting factors to propose new design principles for the analysis of biological data	Assessment and Usage
CO-5	Applications of evolutionary analysis through various available approaches	Usage

## **Course Contents:**

Unit	Contents	Lectures required
1	<b>Introduction:</b> Introduction to molecular evolution and its role in the regulation of various biological processes. Introduction to evolutionary biology, functional genomics, basics of data types for biological sequences their importance and applications in further analysis.	6
2	<b>Biomolecules:</b> To understand various entities and their role in evolution: Genes, operons, regulons, stimulons, genomes, proteins, proteomes etc.	6
3	<b>Codon usage and patterns:</b> Patterns of base composition and codon usage, gene duplication, pseudogenes, orphan genes, gene gain and gene loss processes, overlapping and nested genes, exonization, intron and RNA editing, functional convergence, hypothetical proteins and their annotations.	6
4	<b>Mutation and selection:</b> Various kinds of mutations and selection pressure and related theories.	6
5	<b>New data and evolution:</b> Description of models, methods and algorithms that are most useful for analysing the ever-increasing supply of molecular sequence data, with a view to furthering our understanding of the evolution of genes, proteins, and genomes.	10
	Models of evolution: Models of nucleotide substitution, models of amino acid and codon substitution, phylogeny reconstruction: Overview, Maximum Likelihood	

	6	methods, Bayesian methods, comparison of methods and tests on trees, molecular clock and estimation of species divergence times, neutral and adaptive protein evolution, simulating molecular evolution	•
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42

#### **Total Lectures**

#### Suggested Text Book(s):

- 1. Molecular Evolution by Dan Graur and Wen-Hsiung Li, Sinauer Associated Inc. Pub., USA .
- 2. Computational Molecular Evolution by Ziheng Yang, Oxford Series in Ecology and Evolution.
- 3. Molecular Evolution: A phylogenetic approach Rodric DM Page and Edward C Holmes, Blackwell Science Ltd.

## **Suggested Reference Book(s):**

- 1. Inferring Phylogenies J. Felsenstein, . Sinauer Associated Inc. Pub., USA.
- 2. Bioinformatics and Molecular Evolution by Paul G Higgs, Blackwell Publishing.
- 3. Molecular Evolution and Phylogenetics Masatoshi Nei and Sudhir Kumar, Oxford University Press.

#### **EvaluationScheme:**

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3.	T-3	35	2 Hours	Entire Syllabus
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes & Attendance

## **Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)**

Course outcomes (Computational Molecular Evolution)	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10	PO-11	PO-12	Average
CO-1	2	2	2	3	2	3	1	1	3	2	3	2	2.17
CO-2	2	2	3	2	2	-	_	-	1	1	2	2	1.89
CO-3	3	3	2	2	2	2	1	1	3	2	1	3	2.08
CO-4	2	2	3	1	1	-	-	-	2	2	2	1	1.78
CO-5	3	2	3	2	2	3	-	-	-	2	-	1	2.25
Average	2.4	2.2	2.6	2	1.8	2.67	1	1	2.25	1.8	2	1.8	

## **Diagnostics & Vaccine Manufacturing**

COURSE CODE: 18B1WBT833 COURSE CREDITS: 3 CORE/ELECTIVE: ELECTIVE L-T-P: 3-0-0

### **Pre-requisite:**Immunology

## **Course Objectives:**

- 1. To familiarize the students with the principles & applications of the latest state-of-the-art bio-molecular diagnostic techniques/technology used in laboratories the world over.
- 2. The safety aspects, quality control, quality assurance and validation of PCR based diagnostics and laboratory safety.
- 3. Knowledge of various technologies employed in vaccine production and examine their use in developing vaccines against human and animal pathogens. The safety aspects, quality control, quality assurance and validation of vaccine production and will also be covered.

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	The students would be able to identify and analyze what DNA based approach and methodology should be used for diagnostic purpose in different settings, their comparative advantages and limitations.	
CO-2	The students would be able to identify and analyze what antigen - antibody based approach and methodology should be used for diagnostic purpose in different settings, their comparative advantages and limitations.	
CO-3	The students would have in-depth knowledge of various types of vaccines and approaches used for their production.	Familiarity
CO-4	The students would have in-depth knowledge of quality control and assurance considerations used in the industry for diagnostics.	Assessment & Technical
CO-5	The students would have in-depth knowledge of antimicrobial susceptibility and its application in the industry for diagnostics.	Assessment & Technical

## **Course Contents:**

Unit	Contents	Lecture s require d
1	<b>General Introduction</b> : Biotechnology in the diagnosis of infectious diseases and vaccine development, Biotechnology in Vaccine production, Recent developments in vaccine technology.	2
2	Immunodiagnostics: Antigen – Antibody Interaction, Lattice Theory, Precipitin Curve, Simple Immunodiffusion (Radial Immunodiffusion – Qualitative, Quantitative); Double Diffusion (Mechanism of Reaction of Identity, Partial – Identity, and Non- Identity); Immunoelectrophoresis; Rocket Electrophoresis, Western Blot, Immunofluorescence, Agglutination – Antibody titer, Prozone Phenomenon, Direct and Indirect Agglutination, Hemagglutination, ABO Blood	10

	typing, Agglutination Inhibition; Immunofluorescence, Radioimmunossay (including advantages and disadvantages).	
3	<b>ELISA:</b> Theory, Designing an ELISA method, Types – Direct, Indirect, Sandwich, Competitive, Dot ELISA.	2
4	<b>PCR:</b> concept, protocol, strategy. Types of PCR – Strategy and Applications - Nested, Semi-nested, Real time, RT-PCR, Asymmetric PCR, Inverse PCR, Multiplex PCR.	3
5	<b>QC &amp; QA of PCR and Real Time based diagnostics</b> : Theory, Application, and Trouble shooting. Importance of controls. Best Fit Assay, Optimization and Standardization of PCR based diagnostics.	3
6	<b>AST</b> : Concept, KB Method. Laboratory methodologies for bacterial antimicrobial susceptibility testing – concepts, antibiotics –, resistance, mechanism. Disk diffusion, tube dilution, microbroth dilution methods.	4
7	Biosafety and biosecurity in the medical microbiology laboratory and animal facilities.	2
8	Different types of vaccines, i.e., sub-unit vaccines, recombinant vaccines, synthetic vaccines, idiotypic based - vaccines, DNA vaccines, glycoconjugate vaccines, deletion vaccines.	3
9	Examples of different vaccines - Rabies vaccines, PPRV vaccines, Chimeric vaccines - JEV/DENV/Westnile, Meningococcal conjugate & protein based vaccines, Oral B subunit + whole cell cholera vaccine, Multicellular Parasite vaccines, Novel Vaccines against <i>Mycobacterium tuberculosis</i> , Mycoplasma vaccines, Protozoal &rickettsial vaccines.	8
10	Genetic basis of attenuation, vaccine vectors, large-scale production of vaccines and automation. Vaccine delivery system and approaches to enhance immunogenicity - immunomodulators and, immunomodulation adjuvant. Delivery of particulate antigens through liposomes, microspheres etc.	5
Total lectur	es	42

- 1. Burtis, C. A., Ashwood, E. R.,:Tietz textbook of Clinical Chemistry &Bruns, D. E. Molecular Diagnostics, Saunders, 2006
- 2. World Organization for: Manual of Diagnostic Tests and Vaccines for Animal Health Terrestrial Animals, Volumes I & II, 6th Edition, 2010.
- 3. Rao, J. R.:Molecular Diagnostics: current technology and Applications, Horizon Bioscience, U. K., 2006.
- 4. Review and Research Publications available on-line
- 5. Immunology: Kuby

## **EvaluationScheme:**

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3.	T-3	35	2 Hours	Entire Syllabus
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes&Attendance

Course Outcomes	(COs) contribution	n to the Programm	eOutcomes(POs)
Course Outcomes	(005) contribution	i to the i i ogi ammi	

Course outcomes (Diagnostics & Vaccine Manufacturing)	PO-1	PO-2	PO-3	P0-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10	PO-11	PO-12	Average
CO-1	3	1	1	1	1	1	1	1	0	0	0	2	1
CO-2	1	2	3	1	1	1	1	0	0	2	2	2	1.33
CO-3	1	2	2	2	1	2	1	1	1	2	2	1	1.5
CO-4	2	2	2	3	1	1	3	1	1	1	1	3	1.75
CO-5	1	2	3	2	2	2	1	2	2	2	2	3	2
Average	1.6	1.8	2.2	1.8	1.2	1.4	1.4	1	0.8	1.4	1.4	2.2	

## Traditional Bioprocesses & their upscaling

COURSE CODE: 18B1WBT832

#### **COURSE CREDITS: 3**

CORE/ELECTIVE: ELECTIVE

L-T-P: 3-0-0

Pre-requisite: Microbiology, Bioprocess Engineering, Downstream Processing

## **Course Objectives:**

Traditional bioprocess gives the lying fundamentals of the pre-historic bioprocess technologies and its scaleup studies on the industrial sectors. Moreover, these gives insights about the technological developments till the existing industrial bioprocesses. Bioprocessing of biopharmaceuticals and immobilization technology also gives an opportunity to study the recent advancements in this field. By keeping the mentioned points, the course objectives were framed as follows

- 1. Introduction of traditional bioprocesses and its upscaling
- 2. Discussion of bioprocessing of biopharmecuticals, recent advances in Immobilization technology and Fermentation technology

**Course Outcomes:** 

S.No.	Course Outcomes	Level of Attainment
CO-1	Introduction to Traditional Bioprocesses and its production technologies	Familiarity
CO-2	Bioprocessing of Biopharmaceuticals	Assessment
CO-3	Recent trends in Immobilization technology and bioreactor technology	Assessment
CO-4	Upscaling studies of bioprocess products	Usage
CO-5	Scale up considerations of different bioprocess commodities	Usage

## **Course Contents:**

Unit	Contents	Lectures required					
1	Traditional Bioprocesses: Introduction and Technology Advancemnets:Industrial production of organic acids (Citric acid; Glutamic acid; Lactic acid;Kojic acid; Ascorbic acid) ; Industrial production of Antibiotics(Pencillins; Cephalosporins;Tetracyclins); Industrial production of Amino acids (Lysine;Threonine; Aspartate)						
2	Bioprocessing of Biopharmaceuticals: Overview of USP and DSP aspects of Biopharmaceuticals; Upstream processing of Mab production; Downstream Processing of Mab's ; Process optimization for Mab production; Protein therapeutics	6					
3	<b>Recent advances in Reactor technology:</b> Bioreactors for solid state fermentation ; Photobioreactors for microalgal products; Bioreactors for pharmaceuticals	9					
4	Advances of enzyme immobilization techniques:Sol-gel chemistry and immobilization; Immobilization on nano-particles; Cross-linked Enzyme Aggregates; Surface analysis technology of Immobilized Enzymes	5					
5	Scale up considerations of different bioprocess commodities: Bioprocess aspects of sugar alcohols; Production of flavours in food industries; Flavour production in fermented foods; Bioprocess aspects of Nutraceuticals; Probiotics, Prebiotics and Synbiotics;	8					
tal lect		42					

- 1. Microbial Technology: Microbial processes by Henry J. Peppler, D. Perlman
- 2. Microbial Biotechnology by Alexander N. Glazer and Hiroshi Nikaido
- 3. Industrial Biotechnology by Wim Soetaert and Erick J. Vandamme
- 4. Immobilization of Enzymes and Cells by Jose M.Guisan
- 5. Biofilms in Medicine, Industry and Environmental Biotechnology by Piet Lens et al. **Suggested Reference Book(s):**

# 1. Review / Research articles from Science Direct, Springer, Wiley and Pub Med **EvaluationScheme:**

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3.	T-3	35	2 Hours	Entire Syllabus
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes&Attendance

**Course Outcomes (COs) contribution to the Programme Outcomes(POs)** 

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	Average
CO1	3	3	3	3	1	3	1	2	1	2	1	3	2.17
CO2	3	3	3	3	2	2	1	2	1	2	-	3	2.27
CO3	3	3	3	3	2	2	1	2	1	2	-	3	2.27
CO4	3	3	3	3	1	2	2	3	2	2	2	3	2.42
CO5	3	3	3	3	1	2	3	3	3	2	2	3	2.58
Average	3.00	3.00	3.00	3.00	1.40	2.20	1.60	2.40	1.60	2.00	1.00	3.00	

## **Structural Bioinformatics**

COURSE CODE: 18B1WBI531 COURSE CREDITS: 3 CORE/ELECTIVE: ELECTIVE L-T-P: 3-0-0

## Pre-requisite: Structural Biology

## **Course Objectives:**

1. To develop the ability to design, predict, analyze and compare the protein structures as well as predict the function of target proteins.

## **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	Understanding the fundamental concepts of structural biology (chemical building blocks, structure, superstructure, folding, etc.)	Familiarity
CO-2	To Understand and use structural databases and software for structure visualization	Familiarity
CO-3	To understand the algorithms used in Structure determination and quality assessment	Assessment
CO-4	To perform protein structure comparison and the hierarchical nature of biomacromolecular structure classification	Usage
CO-5	To understand the methodology of protein structure prediction and assessment	Assessment
CO-6	To understand the methodology of sequence- and structure-based functional site prediction	Assessment

## **Course Contents:**

Unit	Contents	Lectures required
1	<b>Introduction of protein structure:</b> Overview of syllabus and protein structure (amino acids and peptide bonds; primary, secondary, super-secondary, tertiary and quaternary structure of proteins).	3
2	Fundamental concepts of structural biology:Chemical	6
	building blocks, structure, superstructure, folding, etc.; the physical forces that shape macromolecules; structural databases (protein data bank, SCOP database, CATH database and other structure based databases)	
3	Secondary structure of protein:Computational methods for prediction of secondary structure of protein sequences (Chou- Fasmann, GOR and Neural Networks) and reliability (Q3 value and SOV score)	3
4	<b>Tertiary structure of protein:</b> Prediction of tertiary structures of protein sequences (Homology and Threading methods); structure quality assessment.	6

5	<b>Protein structures comparison and alignment:</b> General approach of alignment and comparison, comparison algorithm & optimization, statistical analysis of results, multiple structural alignment.	3
6	Analysis of 3D structures:Secondary structure assignment, assignment of hydrogen bonds, coulomb hydrogen bond calculation, empirical hydrogen bond calculation, assignment methods of secondary structure (DSSP, STRIDE, DEFINE, P- Curve)	3
7	<b>Identifying structural domains in protein:</b> How structural domains are defined? First and second generation algorithms for domain assignments, domain assignment based on graph theoretical methods, prediction of binding sites and characterization.	3
8	<b>Ab initio protein structure prediction:</b> Empirical force field for biomolecular simulations, Potential Energy Function (bond length potential, bond angle potential, torsional potential, van der Waals potential and coulomb potential), classical representations of electrostatics (Poisson-Boltzmann, Generalized Born and Colombic).	6
9	<b>Energy minimization techniques:</b> Concept of local and global minima, energy minimization protocol, energy minimization algorithms (steepest descent, conjugate gradient, Newton Raphson)	3
10	<b>Molecular Dynamics simulations:</b> Monte Carlo Simulations, Techniques for efficient conformational search: Simulated Annealing, Calculation of Free energy using simulation techniques.	6
Total Le	ectures	42

- 1. Structural Bioinformatics (2nd Edition), Jenny Gu (Editor), Philip E. Bourne (Editor)
- 2. D.W. Mount Bioinformatics: Genome and Sequence Analysis: (2001) Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York.
- 3. Molecular Modeling: Principles & Applications, Andrew R. Leach, Prentice Hall

## **Evaluation Scheme:**

Assessment	Max. marks	Duration	Course Covered
T1 Test	15	1 hr.	Syllabus covered up to Test-1
T2 Test	25	1.5 hrs.	Syllabus covered up to Test-2
End Term Test	35	2 hrs.	Whole Syllabus
Teacher Assessment (Based on Assignments, quizzes etc.)	25	Whole Semester	Inform class time to time
Total	100		

CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	Average
CO1	3	3	3	3	2	2	1	1	1	1	1	1	1.83
CO2	3	3	3	3	3	1	1	1	1	1	1	3	2.00
CO3	3	3	2	3	2	3	2	1	1	1	2	1	2.00
CO4	3	3	3	2	3	2	1	1	1	1	1	1	1.83
CO5	2	2	3	3	3	3	1	1	1	1	1	1	1.83
CO6	2	3	3	3	2	2	2	2	2	2	2	2	2.25
Average	2.67	2.83	2.80	2.80	2.60	2.20	1.20	1.00	1.00	1.00	1.20	1.40	

## Course Outcomes (COs) contribution to the Programme Outcomes (POs)

## **Advanced Algorithms for Bioinformatics**

COURSE CODE: 18B1WBI631

COURSE CREDITS: 3

CORE/ELECTIVE: ELECTIVE

L-T-P: 3-0-0

Pre-requisite: Basics of data structures, algorithms, and basic methods in computational biology

## **Course Objectives:**

- 1. The overall objective of the course is to develop an understanding of algorithms implementation for solving problems in biology.
- 2. To evaluate existing algorithms, possible improvements and for their implementations.

## **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	Able to apply algorithmic principles to address problems in biology	Analytical
CO-2	Use various methods from computational biology to implement their algorithmic versions	Usage
CO-3	Analyze problems in biology and able to design new protocols and algorithms for biological data analysis	Analytical
CO-4	Able to analyze the algorithms in computational biology and identify their limiting factors to propose new design principles	Analytical
CO-5	Assessment of biological complexity through algorithmic principles	Analytical

**Course Contents:** 

Unit	Contents	Lectures required
1	<b>Introduction:</b> An overview of Algorithms, Sequence and String search algorithms with mathematical formulations for similarity and distance scoring systems with their algorithmic implementations.	6
2	Genome Assembly:Complexity of DNA problems and their simulatory solutions.Genomeassemblyalgorithms, their computationalimplications and applications.	7
3	<b>Graph based Algorithms:</b> Graph algorithms in bioinformatics and their applications to fragment assembly, Eulerian and Hamiltonian Cycle Problem, Interval graph algorithm, shortest superstring problem and its mapping with traveling salesman problem.	4
4	<b>Motif and Regulatory element's Algorithms:</b> Algorithms for finding regulatory motifs in genomic sequences through profiles and consensus approaches. Brute Force Motif Search, Median String Search algorithms and their refinements. Algorithms for Sequencing by hybridization (SBH), use of spectrum approach to solve SBH problem.	5
5	<b>Gene prediction:</b> Algorithmic approaches for Contig assembly to super- contigs. Computational challenges for gene prediction, popular algorithms and their implementations for gene prediction. Exon chaining and Spliced Alignment Problems.	7
6	Brute Force and branch and bound algorithms:Brute Force and branch and bound algorithms for Partial Digest Problem, restriction mapping, partial digest	

	and double digest problems and their solutions through multiset and homometric sets.	5
7	<b>MSA advancedments:</b> Progresssive and iterative refinements of MSA algorithms, Barton-Sternberg Iterative Refinement Algorithm, STAR and TREE alignment approaches, Greedy and Entropy approach for MSA.	5
8	<b>Graph based MSA advancements:</b> Partial Order (PO)-MSA, and A- Bruijn Alignment (ABA) algorithm for MSA. Combinatorial dynamic programming approach for MSA.	3
Total Le	ctures	42

- 1. Computational Molecular Biology: An algorithmic approach (2004), P.A. Pevzner, PHI.
- 2. An Introduction to Bioinformatics Algorithms (2004) N.C. Jones and P.A. PevznerAne Books.
- 3. Algorithms in Bioinformatics (2004), G. Benson and R. Page (Eds): Springer Verlag.

## **Suggested Reference Book(s):**

1. Bioinformatics Algorithms: Techniques and Applications, I.I. Mandoiu and A Zelikovsky,

Wiley Interscience Press.

2. Biological Sequence Analysis: Probabistic models of proteins and nucleic acids (1998) Durbin R., et al, Cambridge University press.

## **EvaluationScheme:**

S. No	No Exam		Duration	Coverage / Scope of Examination			
1	T-1	15	1 Hour.	Syllabus covered upto T-1			
2	T-2	25	1.5 Hours	Syllabus covered upto T-2			
3.	T-3	35	2 Hours	Entire Syllabus			
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes & Attendance			

## **Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)**

Course outcomes (Advanced Algorithms for Bioinformatics)	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10	P0-11	PO-12	Average
CO-1	2	3	1	3	2	2	2	1	3	2	3	3	2.25
CO-2	3	2	2	1	-	1	-	-	-	1	2	3	1.88
CO-3	3	3	3	3	2	2	2	1	2	_	-	2	2.30
CO-4	3	3	3	1	-	-	-	-	2	1	2		2.14
CO-5	3	1	2	2	2	2	-	1	-	2	-	1	1.78
Average	2.8	2.4	2.2	2	2	1.75	2	1	2.33	1.5	2.33	2.25	

## Datawarehousing and Mining for Bioinformatics

COURSE

CODE:18B

1WBI632

COURSE

CREDITS:

3

ELECTIVE

L-T-P: 3-0-0

Pre-requisite: Molecular Biology, Biochemistry

## **Course Objectives:**

- 1. Learn to develop and use datawarehouse
- 2. Learn feature selection methods
- 3. Learn methods for data mining.
- 4. Apply data mining techniques in biological datasets.
- 5. Learn and apply cross-validation.

## **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	Students will have a thorough understanding of various datawarehousing components and architecture.	Familiarity
CO-2	Students will understand various types of data models.	Assessmen
CO-3	Students will understand how to perform feature selection and derive association rules	Assessmen
CO-4	Students will understand how to perform various types of data mining, including clustering, neural networks etc.	Usage

#### **Course Contents:**

Unit	Contents	Lectures required
1	Knowledge Discovery Process - Understanding the business intelligence cycle	2
2	Introduction to Data warehousing – Components of data warehouse, Architecture, lifecycle & related core terms.	2
3	<b>Types of Data warehouse design methodologies</b> – Top Down Approach, Bottom Down approach, Hybrid design Approach	3
4	<b>Data Models</b> - Dimensional Data Modeling (Star Schema, Snowflake Schema); Relational Data Modeling; Conceptual, Physical & Logical Data Model.	3
5	Multidimensional Analysis – OLAP & OLTP Approaches	2
6	Building & Maintaining the data warehouse – ETL design & development;	2
7	Introduction to Data Mining - concepts and techniques for the	2
	discovery of patterns hidden in large data sets	

	Grouping of data - Classification and Clustering Methods; Decision Tree, Neural	
8	Network, Nearest Neighbor, Genetic Algorithm	7
9	<b>Feature Selection Methods -</b> Wrapper & Filter Approach, Correlation analysis, PCA	7
10	Association Rule Learning Based Methods - Apriori Algorithm	7
11	Statistical techniques involved in data mining – regression based model development	1
12	Cross Validation Techniques - Jackkniffing, Bootstrapping, Sensitivity, Specifcity, Accuracy	2
	Total Number of Lectures	42

- a. Kimball, R., Margy, R. : The Data Warehouse Toolkit, 2nd Edition: The Complete Guide to Dimensional Modeling, John Wiley &SonsMolecular Biology of the Gene (1987) Watson J. D., Hopking N., Robast J. and Steiz, J.
- b. Inmon, B. :Building the Data Warehouse, John Wiley & Sons.

## Suggested Reference Books(s):

- 1. Pei, Han and Kamber, Data mining: Concepts and techniques third edition, Elsevier, 2011
- 2. Data Mining: Practical Machine Learning Tools and TechniquesKim JB, Porreca GJ, Song L, Greenway SC, Gorham JM, Church GM, Seidman CE,
- 3. Introduction to Data Mining, Tan, Steinbach and Vipin Kumar, Pearson Education, 2016

### **Other useful resource(s):**

- a. Link to NPTEL course contents:https://onlinecourses.nptel.ac.in/noc18\_cs14/preview
- b. Link to topics related tocourse:
- iv. https://nptel.ac.in/courses/102104063
- v. https://nptel.ac.in/courses/102106069/
- vi. https://nptel.ac.in/courses/102106026/

## **EvaluationScheme:**

S. No	Exam	Marks	Duration	Coverage / Scope of Examination
1	T-1	15	1 Hour.	Syllabus covered upto T-1
2	T-2	25	1.5 Hours	Syllabus covered upto T-2
3.	T-3	35	2 Hours	Entire Syllabus
4.	Teaching Assessment	25	Entire Semester	Assignment, Quizzes&Attendance

## **Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)**

Course outcomes (Datawarehousing and Mining for BI)	PO-1	P0-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10	PO-11	PO-12	Average
CO-1	2	2	2	2	2	2	2	2	2	2	1	2	1.9
CO-2	2	2	2	2	2	1	1	2	2	2	1	2	1.75
CO-3	2	2	2	2	2	1	1	1	2	-	2	2	1.7
CO-4	2	2	2	2	2	1	1	1	2	2	-	2	1.7
Average	2.0	2.0	2	2	2	1.25	1.25	1.5	2.0	2	1.33	2.0	

## Department of Biotechnology & Bioinformatics

## Intellectual Property Rights and Commercialization

COURSE CODE: 18B1WBT732

COURSE CREDITS: 3

CORE/ELECTIVE:

ELECTIVE L-T-P: 3-0-0

#### **Pre-requisite:** None

## **Course Objectives:**

- 1. To provide an insight and understanding about different aspects of protection of inventions and research developments
- 2. Learn about procedures for filling protection through Intellectual Property Rights.
- **3.** To provide scopes of protection of diverse intellectual properties and its commercialization for socio-economic improvement.

#### **CourseOutcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	To enable students with basic concepts and knowledge of intellectual property rights.	Awarness
CO-2	To apply and execute different types of IP protection in research and academics.	Assessment and technical skills
CO-3	Able to understand about the mechanisms of different IP protections, registrations and applications	Technical
CO-4	To be capable of tackling issues related to IP and its commercialization	Assessment
CO-5	Able to learn the strategies for effective IP management and commercialization	Analytical skills
CO-6	To apply the knowledge of IPR for the benefit generation and for mass utilization	Usage
	CourseContents:	
Unit	Contents	Lectures required
1	Introduction:Introduction of Intellectual properties and rights conferred . Integration of Intellectual Property, Bioethics and Biosafety for biological and applied sciences in research andacademia.	4
2	Types of IP tools:Different types of IPR( Patents, copyrights and related rights, Trademark, Tradesecret, Integrated circuit layout, Geographical indications, Traditional knowledge, Industrial designs and PBR)Drafting Patent Application and Documentation Revocation of Patent, Litigation and Infringement Rationale of different IPR, their mechanism of protection and provisions in Law	

 

 3
 International Agreements and Treaties:International IP treaties (Madrid Agreement, Trademark law treaty, Patent Law treaty etc.)

 WIPO, EPC, WTO, and TRIPS. International agreements relevant to biotechnology-associated IP
 8

4		
	<b>Commercialization</b> : Methods of commercialization,Impact of commercialization. Financing	6
5	<b>IP Management for value addition:</b> Strategies for IP Management and commercialization. IP audit, IP insurance Bioentreprenuership management	4
6	<b>Licensing/Assignment</b> :Types of licensing and modes to carry out, Assignments and its benefits, Compulsory Licensing Commercialization for social and economic prosperity with case studies	8
Total lec		42

- 1. Intellectual Property Rights & Copyright By Bouchoux.
- 2. Intellectual Property Licensing Strategies by Thompson Reuters

#### **Suggested Reference Book(s):**

- 1. Intellectual Property Rights, the WTO and Developing Countries: The TRIPS ...Book by Carlos María Correa
- 2. Perspectives on Commercializing Innovation by F. Scott Kieff (Editor), Troy A. Paredes (Editor

#### Course Outcomes (COs) contribution to the Programme Outcomes(POs)

Course outcomes (Intellectual Property Rights & Commercializat ion)	PO-1	P0-2	PO-3	P0-4	PO-5	PO-6	P0-7	PO-8	PO-9	PO-10	PO-11	PO-12	Average
CO-1	2	2	2	2	2	1	1	1	1	1	2	2	15
CO-2	2	2	1	2	2	2	1	1	1	1	1	2	1.5
CO-3	2	2	2	2	3	1	1	1	2	2	1	2	1.7
CO-4	2	2	3	3	2	1	1	1	2	2	2	2	1.9
CO-5	2	2	2	2	2	1	2	1	1	2	1	2	1.6
CO-6	2	2	2	2	2	2	1	1	2	2	2	2	1.8
Average	3.4	3.4	2	2.1	2.1	1.3	1.1	1	1.5	1.6	1.5	2	

## **Diagnostics & Vaccine Manufacturing**

COURSE CODE:18B1WBT833 COURSE CREDITS: 3 CORE/ELECTIVE: ELECTIVE L-T-P: 3-0-0

#### Pre-requisite:Immunology

#### **Course Objectives:**

- 1. To familiarize the students with the principles & applications of the latest state-of-the-art biomolecular diagnostic techniques/technology used in laboratories the world over.
- 2. The safety aspects, quality control, quality assurance and validation of PCR based diagnostics and laboratory safety.
- 3. Knowledge of various technologies employed in vaccine production and examine their use in developing vaccines against human and animal pathogens. The safety aspects, quality control, quality assurance and validation of vaccine production and will also be covered.

#### **Course Outcomes:**

S.No.	Course Outcomes	Level of Attainment
CO-1	The students would be able to identify and analyze what DNA based approach and methodology should be used for diagnostic purpose in different settings, their comparative advantages and limitations.	Usage
CO-2	The students would be able to identify and analyze what antigen - antibody based approach and methodology should be used for diagnostic purpose in different settings, their comparative advantages and limitations.	Isage
CO-3	The students would have in-depth knowledge of various types of vaccines and approaches used for their production.	Familiarity
CO-4	The students would have in-depth knowledge of quality control and assurance considerations used in the industry for diagnostics.	Assessment & Technical
CO-5	The students would have in-depth knowledge of antimicrobial susceptibility and its application in the industry for diagnostics.	Assessment & Technical

#### **Course Contents:**

Unit	Contents	Lectures required
1	<b>General Introduction</b> : Biotechnology in the diagnosis of infectious diseases and vaccine development, Biotechnology in Vaccine production, Recent developments in vaccine technology.	2
2	<b>Immunodiagnostics:</b> Antigen – Antibody Interaction, Lattice Theory, Precipitin Curve, Simple Immunodiffusion (Radial Immunodiffusion – Qualitative, Quantitative); Double Diffusion (Mechanism of Reaction of Identity, Partial – Identity, and Non- Identity); Immunoelectrophoresis; Rocket Electrophoresis, Western Blot, Immunofluorescence, Agglutination – Antibody titer, Prozone Phenomenon, Direct and Indirect Agglutination, Hemagglutination,	10
	ABO Blood typing, Agglutination Inhibition; Immunofluorescence, Radioimmunossay	

	(including advantages and disadvantages).	
3	<b>ELISA:</b> Theory, Designing an ELISA method, Types – Direct, Indirect, Sandwich, Competitive, Dot ELISA.	2
4	<b>PCR:</b> concept, protocol, strategy. Types of PCR – Strategy and Applications - Nested, Semi-nested, Real time, RT-PCR, Asymmetric PCR, Inverse PCR, Multiplex PCR.	3
5	<b>QC &amp; QA of PCR and Real Time based diagnostics</b> : Theory, Application, and Trouble shooting. Importance of controls. Best Fit Assay, Optimization and Standardization of PCR based diagnostics.	3
6	<b>AST</b> : Concept, KB Method. Laboratory methodologies for bacterial antimicrobial susceptibility testing – concepts, antibiotics –, resistance, mechanism. Disk diffusion, tube dilution, microbroth dilution methods.	4
7	Biosafety and biosecurity in the medical microbiology laboratory and animal facilities.	2
8	Different types of vaccines, i.e., sub-unit vaccines, recombinant vaccines, synthetic vaccines, idiotypic based - vaccines, DNA vaccines, glycoconjugate vaccines, deletion vaccines.	3
9	Examples of different vaccines - Rabies vaccines, PPRV vaccines, Chimeric vaccines – JEV/DENV/Westnile, Meningococcal conjugate & protein based vaccines, Oral B subunit + whole cell cholera vaccine, Multicellular Parasite vaccines, Novel Vaccines against <i>Mycobacterium tuberculosis</i> , Mycoplasma vaccines, Protozoal &rickettsial vaccines.	8
10	Genetic basis of attenuation, vaccine vectors, large-scale production of vaccines and automation. Vaccine delivery system and approaches to enhance immunogenicity - immunomodulators and, immunomodulation adjuvant. Delivery of particulate antigens through liposomes, microspheres etc.	5
otal lectu	ires	42

- 1. Burtis, C. A., Ashwood, E. R.,:Tietz textbook of Clinical Chemistry &Bruns, D. E. Molecular Diagnostics, Saunders, 2006
- 2. World Organization for: Manual of Diagnostic Tests and Vaccines for Animal Health Terrestrial Animals, Volumes I & II, 6th Edition, 2010.
- Rao, J. R.:Molecular Diagnostics: current technology and Applications, Horizon Bioscience, U. K., 2006.
- 4. Review and Research Publications available on-line
- 5. Immunology: Kuby

**Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)** 

course outcomes (cos) contribution to the Programme outcomes (ros)													
Course outcomes (Diagnostics & Vaccine Manufacturing)	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	6-04	PO-10	PO-11	PO-12	Average
CO-1	3	1	1	1	1	1	1	1	0	0	0	2	1
CO-2	1	2	3	1	1	1	1	0	0	2	2	2	1.33
CO-3	1	2	2	2	1	2	1	1	1	2	2	1	1.5
CO-4	2	2	2	3	1	1	3	1	1	1	1	3	1.75
CO-5	1	2	3	2	2	2	1	2	2	2	2	3	2
Average	1.6	1.8	2.2	1.8	1.2	1.4	1.4	1	0.8	1.4	1.4	2.2	

## **Industrial Enzymes Technologies**

COURSE CODE:18B1WBT731 COURSE

CREDITS: 3 CORE/ELECTIVE: ELECTIVE L-

T-P: 3-0-0

Pre-requisite: Enzyme production purification and applications

## **Course Objectives:**

1. The objective of the course is to develop an understanding of important aspects of production and purification of industrially important enzyme and their application in industry. Course Outcomes:

S.No.	Course Outcomes	Level of Attainment
CO-1	To develop an understanding of basic concepts of enzymes.	Familiarity
CO-2	To understand the basic mechanism of action and working behaviour of enzymes	Assessment
CO-3	To familiarize the students with various applications of enzymes in laboratory as well as Industrial scale.	Assessment
CO-4	To conceptualize about immobilized enzyme technology, and other specific enzymes and their applications.	Usage
CO-5	To familiarize the students with present potential of enzyme in industrial application and improved activity of the enzyme using various molecular biology techniques.	Usage
CO-6	To understand the principle and function of enzyme in various adverse conditions like high temperature and pH(s).	Assessment

#### **Course Contents:**

Unit	Contents	Lectures required
1	<b>Enzymes: Basic concepts:</b> Enzymes as powerful and highly specific catalysts, Classification of enzymes, free energy and enzymes, the formation of the transition state, catalytic strategies. General properties: Enzyme specificity, stability and structure, Factors affecting enzyme activity; effect of pH and Temperature, Substrate and Enzyme concentration.	5
2	<b>Enzyme kinetics</b> : Michaelis-Menten kinetics, evaluation of parameters in the Michaelis-Menten equation, 3-D structure of active site, Kinetics of single and bi-substrate enzyme catalysed reactions, Inhibition & its kinetics.	5
3	<b>Enzyme preparation techniques:</b> Sources of enzymes, production,Recovery and purification of intracellular products: cell disruption, chromatographic techniques. Analytical assays of purity level of enzymes.	3
4	<b>Enzyme preparation and application in industries:</b> Application of enzymes in leather, glucose syrup production, starch and sugar industry, Dairy and food industry, Beverage industry, Textile industry. Hydrolysis of starch and cellulose. Catalytic functions of Cellulase, lipase, esterase laccase amylase, glucose isomerase, protease, xylanase, invertase, peroxidises. Other applications of enzymes in solution: medical applications of enzymes, non-hydrolytic enzymes in current and developing industrial technology.	10
5	<b>Enzyme engineering:</b> Mechanisms and manifestations of protein denaturation. Strategies for enzyme stabilization: Physical and chemical modifications, Selection, directed evolution and Rational design. design and construction of mutant enzymes,	5

6	<b>Immobilized-enzyme technology</b> : Introduction, enzyme immobilization method: Entrapment, carrier-binding and cross-linking method. Medical and analytical applications of immobilized enzymes.	8
7	<b>Specified Enzymes and applications:</b> Thermozymes, Cold adapted enzymes, Ribozymes, Hybrid enzymes, Diagnostic enzymes, Therapeutic enzymes: Characteristics, principles and applications.	6

- 1. Devasena, T., "Enzymology", 1st ed., Oxford University Press, 2010
- 2. Berg, J.M., Tymoczko, J.L. and Stryer, L., "Biochemistry", 5th ed., W.H. Freeman and Company, New York, 2002.
- 3. Nelson D.L., Cox M.M., "Lehninger Principles of Biochemistry", 5th ed., W.H. Freeman and Company, New York, 2008.

## **Suggested Reference Book(s):**

- 1. Pye, E.K. and Wingard, L.B., "Enzyme Engineering II", Plenum Press, 1974.
- 2. Illanes A, "Enzyme Biocatalysis", Springer Science, 2008.

## **Other useful resource(s):**

https://nptel.ac.in/course.php?disciplineId=102

## Course Outcomes (COs) contribution to the ProgrammeOutcomes(POs)

Course outcomes	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	P0-7	PO-8	9-04	PO-10	PO-11	PO-12	Average
CO-1	2	2	3	1	1	2	2	3	2	2	1	2	1.9
CO-2	1	2	2	2	2	1	-	2	2	2	1	3	1.8
CO-3	2	2	2	2	2	2	1	1	2	-	2	2	1.8
CO-4	2	3	2	1	2	1	-	1	3	2	-	2	1.9
CO-5	1	3	1	3	2	2	1	2	1	2	1	3	1.8
CO-6	1	1	-	2	2	2	2	1	2	2	1	3	1.7
Average	1.5	2.1	1.6	1.8	1.8	1.6	1	1.6	2	2	1.2	2.5	

## NGS Data Analysis and Applications

## COURSE CODE: 18B1WBI834 COURSE

CREDITS: 3 CORE/ELECTIVE: ELECTIVE

L-T-P: 3-0-0

## **Pre-requisite:**

#### **Course Objectives:**

- 1. To acquire basic knowledge of Next Generation Sequencing approaches in genomic research.
- 2. To understand computational and statistical principles for analysing genomic data.
- 3. To understand and formulate questions to investigate deep sequencing data.

#### **Course Outcomes**

S.No.	Course Outcomes	Level of Attainment
CO-1	The students will acquire knowledge of Next Generation Sequencing technologies used in genomics and genetics research.	Familiarity
CO-2	The students will learn about the various platforms used in NGS.	Assessment
CO-3	The students will learn about the tools and techniques used in NGS data analysis.	Usage
CO-4	The students willknowtheapplications and scopes of genomics research using the latest genome-wide data centric approaches.	Assessment

**Course Contents:** 

Unit	Contents	Lectures required			
1		4			
	Overview of Sequencing Technologies: Historical perspective of sequencing	7			
	technologies. Overview of modern sequencing technologies. Sanger				
	Sequencing. Next Generation sequencing platforms. Whole genome				
	sequencing.				
2	Next Generation Sequencing Data analysis: Data acquisition and base	10			
	calling. Quality of sequencing data. Read mapping and genome assembly.	10			
	Structural and functional annotation of genomes. Statistics and algorithms				
	used in different steps in data analysis.				
3	Technologies for Transcriptomics and Regulatory Genomics: Chip-				
	seqAnlysis. Peak finding. Motif Discovery. RNA-seq Analysis. Differential				
	gene expression analysis. Exome sequencing and analysis. Exome	9			
	sequencing.				
4	Software and pipelines for NGS Data Analysis: Chip-seq analysis				
	Piplines. RNA-seq analysis pipelines. Software used for assembly and				
	differential gene analysis. Basics of Genome Browsers. Annotation pipelines.	9			
5	Genome Sequencing and Applications in Genetics Studies: Algorithms				
	and application in studying regulation of gene expression. Emerging				
	technologies of single cell gene expression analysis. Metagenomics. Variant	10			
	detection. Time series analysis. Pathway Analysis.	<u>10</u> <b>42</b>			
Total lectures					

#### Suggested Text Book(s):

- 1. Genomic Data Analysis by Megahed Mohammad, LAP Lambert Academic Publishing
- 2. Deep Sequencing Analysis by Noam Shomrom, Springer.

Course outcomes	P0-1	PO-2	PO-3	PO-4	PO-5	9-04	PO-7	PO-8	6-04	PO-10	PO-11	PO-12
CO-1	2	2	1	2	2	2	2	1	2	2	2	2
СО-2	1	2	1	2	2	2	2	2	2	2	1	2
СО-3	2	3	3	3	3	2	3	2	2	1	2	2
CO-4	2	3	3	3	3	2	3	2	2	1	2	2