# **Learning Outcomes based Curriculum Framework**

(LOCF) - Curriculum and Syllabus

For

#### M.Sc., MICROBIOLOGY

#### Vision of the University

To provide quality education to reach the un-reached

#### **Mission of the University**

- To conduct research, teaching and outreach programmes to improve conditions of human living.
- To create an academic environment that honours women and men of all races, caste, creed, cultures, and an atmosphere that values intellectual curiosity, pursuit of knowledge, academic freedom and integrity.
- To offer a wide variety of off-campus educational and training programs, including the use of information technology, to individuals and groups.
- To develop partnership with industries and government so as to improve the quality of the workplace and to serve as catalyst for economic and cultural development.
- To provide quality / inclusive education, especially for the rural and un-reached segments of economically downtrodden students including women, socially oppressed and differently abled.

#### **PREAMBLE**

Under the advice and supervision of the University Grants Commission, the Choice Based Credit System (CBCS) curriculum for Microbiology at the postgraduate level has now been transformed into a new system called Learning Outcome Curriculum Framework (LOCF) (UGC). The LOCF approach considers the programme learning outcomes of the M.Sc. degree in Microbiology, as well as the learning outcomes of the courses taught as part of this programme, while keeping in mind the program's graduate

qualities. The curriculum was then created to match the learning objectives. It is expected that students who complete this programme would possess the necessary knowledge, abilities, temperament, and ethics in the field of microbiology. Aside from the curriculum's content, the teaching and learning methods have also been designed to attain these qualities. The curriculum includes a wide range of learning assessment tasks. These exercises would not only aid to assess the students' knowledge/skills, but they would also help to enrich the teaching learning processes.

The study of microorganisms or microbes such as bacteria, viruses, fungus, algae, protozoa, and infectious proteins such as prions is known as microbiology. Microbes are extremely important because their diverse activities range from the transmission of diseases to the production of highly useful products such as antibiotics, vitamins, enzymes, alcohol, fermented foods, and the recycling of nutrients from dead and decaying organic matter in the environment. Microbiology, as a science, plays a vital role in health, agriculture, the environment, and industry. Several breakthroughs in the previous two to three decades that have had a considerable impact on these fields of human endeavour have pushed Microbiology to the forefront of global teaching, research, and development.

The completion of a microbiology graduating course merely establishes a foundation for fundamental knowledge of the field. Microbiology has been transformed and enriched thanks to inventions, advances, and technology. The demand for professional labour necessitates a deep understanding of the subject. It also necessitates the incorporation of cutting-edge information and new technology in order to meet society's evolving needs. Experienced labour is preferred by the public and private sectors. In light of this, the M.Sc. Microbiology LOCF-2022 course is designed to provide comprehensive and up-to-date knowledge of the field, allowing students to enter the public and private sectors with ease. The course is unique in that it requires 6 months of research projects. Students have the option to work at nationally and internationally renowned research institutes and enterprises throughout this time. As a result, professional human resources are produced in accordance with societal needs. Other research parts of the course include scientific writing, research proposal writing, publication preparation, and research poster preparation for conferences, and the entire process also produces new minds to work as scientists.

#### Programme Learning Outcomes of M.Sc., Microbiology

A candidate who receives a PG degree in Microbiology, i.e., M.Sc. degree, must have acquired/developed the competences listed in the Program Employability section. During the course of the study, results and programme specific outcomes will be compared to course outcomes.

#### Programme Employability Outcomes of M.Sc., Microbiology

- 1. Acquire knowledge and comprehension of microbiology ideas as they apply to various fields such as medicine, industry, the environment, genetics, agriculture, and food.
- 2. Demonstrate key practical skills/competencies in dealing with microorganisms for research and use in the lab and land, including the application of appropriate microbiological techniques.
- 3. Capable of applying microbiological knowledge and abilities to analyse microbe-related problems, communicate them to peers/team members/other stakeholders, and implement remedial measures/studies, among other things.
- 4. Gain a wider understanding of the science of microbiology, allowing them to identify difficult societal issues and plan their professional career to discover novel answers.

# **Programme Outcome**

# 1. PO1 Disciplinary Knowledge

This programme offers knowledge on aspects and concepts of microbiology, life-cycle, their physiology and metabolism, diagnostic procedures, imparting knowledge on gene manipulation, bioinformatics, food (agriculture), environment, industrial, medical and pharmaceuticals which enable them to apply solve the problems in microbiology.

#### 2. PO2 Communication Skills

Enable the students to communicate the scientific findings and analysis through Research paper writing, project writing, by giving periodic assignments, seminars, presentation of research findings in conferences and seminars, and presentation of project results in viva-voce. All these aspects are based on different disciplines of microbiology.

# 3. PO3 Critical Thinking

The students get critical thinking on various avenues of standardization of microbiological techniques, disease diagnosis and management.

### 4. PO4 Problem Solving

The understanding of advanced technologies in microbiology enables the students to identify novel medicines from microbes, survey and analyse the problems in disease diagnosis and solving them.

#### 5. PO5 Analytical Reasoning

The practical related to isolation, identification and interpretation of microbiology based research and arriving valid conclusions.

#### 6. PO6 Research Related Skills

One semester (Final semester) Microbiology research based project involves ability to define problem, formulate the hypothesis, draw conclusions and report the results are included in the students' curriculum.

#### 7. PO7 Co-operation / Team Work

Develop an independent thinker and researches effectively and be a part of a team member or lead a team and take part in multidisciplinary research. Team works among the students to establish the role of microorganisms in food, medical and environmental microbiology included in the curriculum create co-operation within student community to solve the problems.

#### 8. PO8 Scientific Reasoning

The programme offers analysis of microbiology based data, its critical evaluation and reasoning.

### 9. PO9 Reflective Thinking

Develop reflective thinking in Microbiology, drug from microbes, resource utilization and management for the benefit of the society.

#### 10. PO10 Information / Digital Library

Almost all the course in Microbiology teaching is based on knowledge dissemination involving ICT. For the project data analysis appropriate analysis is recommended.

#### 11. PO11 Self-directed Learning

Microbiology project works, Practical's and group works make a self-directed approach among the student community.

#### 12. PO12 Multicultural Competence

Microbiology based industrial visits, visit to national organization, and participation of international webinars create the multi-cultural competence among students.

#### 13. PO13 Moral & Ethical Awareness / Reasoning

Create moral and ethical practices in project oriented data collection, presentation of results, entrepreneurship.

# 14. PO14 Leadership Readiness / Qualities

Create start-up/entrepreneurship based grouping of teams and inculcating of leadership qualities among students.

#### 15. PO15 Life-long Learning

Establishment of research forum, invitation of alumni and exchange of knowledge among students creates life-long learning among students.

#### PROGRAM SPECIFIC OUTCOMES (PSOs)

#### PSO1:

Analyse the fundamental ideas and biodiversity of microorganisms (bacteria, fungus, actinomycetes, viruses, algae) in order to develop critical thinking skills in various domains of microbiology. Understand the genetic systems of prokaryotes as well as the physiology, metabolism, and biochemistry of microbes. The structure and functions of the cell, as well as the molecular biology of prokaryotic cellular structure, will be covered. Learn basic Microbiology laboratory skills, techniques, and competence in the use of research tools, clinical approaches, and observation analysis.

#### PSO2:

Demonstrate the importance of pathogen immunity, pathogenesis, cultivation, diagnosis, and control in various health and pharmaceutical sectors using treatments and prophylaxis.

#### PSO3:

Assess and evaluate the demands, potentials, and impacts of microorganisms that are relevant to food, soil, and agriculture, in order to ensure environmental protection and food quality.

#### **PSO4**:

Design appropriate bioprocessing and fermentation techniques, with a focus on being familiar with microbe applications for industrial biomass production and the synthesis of valuable compounds through fermentation.

#### **PSO5**:

Understanding the fundamentals of recombinant DNA technology (RDT) and investigating the use of genetic engineering to generate GMOs, gene therapy, and other applications. Apply bioinformatics, biochemistry, and genetics concepts to the betterment of society through analytical and molecular approaches.

#### **PSO6**:

Analyze the importance of research using statistical methods and present the results in research forums. Ensure biosafety and bioethics in order to promote social responsibility and environmental awareness, as well as obtain Intellectual Property Rights (IPR) for varied research findings. Employability and lifelong learning need the use of computing, communication, and entrepreneurial abilities.

#### Structure of M.Sc., (Microbiology) LOCF degree program:

The overall structure of the course to be implemented from the academic year 2022-2023 onwards is as follows.

- A. Master of Science in Microbiology programme will last for two years. The course will be divided into four semesters, with two semesters in each year.
- B. Four categories of courses will be offered: Core Compulsory Courses: Includes theoretical as well as practical courses, Core Elective Courses: Include theoretical courses and its department specific courses. The students must opt for three core elective courses for the semesters 1, 2 and 3 out of six courses offered by the department, two courses for each semester. Open Elective: For the 1<sup>st</sup>,

2<sup>nd</sup>, 3<sup>rd</sup> and 4<sup>th</sup> semester, students may opt for any one open elective offered through swayam portal (e-PG pathasala-INFLIBNET) and MOOCs. Skill Enhancement Elective Course: A separate project training-based course that leads to a dissertation worth nine credits is also one of the core courses.

C. Candidates must appear for four core compulsory theory courses, one core elective course, and one open elective course from e-PG pathasala-INFLIBNET in semester I. The candidate must finish two practical courses as specified in the syllabus for semester I and the practical examinations for Practical Courses 1 and 2 will be held at the end of the semester. Total credits for the semester I is 26. In semester 2, the candidate must take four core compulsory courses, one core elective course, and one open elective course from MOOCs. Two practical courses will be held in semester II, as specified in the curriculum, and practical examinations for practical courses 3 and 4 will be held at the end of the semester. Totally 26 credits is allotted for the II semester.

D. There will be four core compulsory courses, two practical courses, one core elective course, and one open elective course from MOOCs in Semester III. The practical examinations for practical courses 5 and 6 will be held at the end of the semester III. The total credits for the semester III is 26. Semester IV is devoted exclusively to project/dissertation work with a credit of 9, as well as one open elective paper from e-PG pathasala-INFLIBNET with 3 credits. The elective paper will have a theory examination at the end of semester IV. The total credits for the IV semester is 12.

E. Students will be required to participate in an internship programme for three weeks (21 days) during the second semester summer vacation to learn about research approaches and work presentation in industries or research institutes. The internship will be awarded 100 points and 3 credits in total. Students will work on their internships, complete the experimental/internship work, and submit the internship report within the time frame specified, *i.e.*, before the start of the third semester, i.e., the second year of their study..

F. The entire M.Sc. in Microbiology course will be covered in 19 theory papers: 12 core compulsory courses, 3 core elective courses, 4 open elective courses - 2 from MOOC's and 2 from e-PG pathasala-INFLIBNET, 6 core compulsory practical courses, and an internship (major project with Dissertation). Each core theory course will be addressed in four one-hour weekly lectures, and the

core elective courses and open elective courses will be addressed in three one-hour weekly lectures. Each practical course will be covered in two four-hour practical turns per week. As a result, the students will work for each practical on two days of the week. The total credits for the entire programme is 93.

A complete syllabus for each paper is included, as well as a list of suggested reading that can be augmented with other books/papers. While older editions of books are recommended for certain topics, the books that are commonly prescribed are the most recent editions.

Eligibility for admission: Candidates who have completed Bachelor's degree in science in Microbiology / Biochemistry / Biotechnology / Zoology with Botany ancillary / Botany with Zoology ancillary / Plant Science and Biotechnology / Advanced Zoology and Animal Biotechnology / Biology / Life Science / Nutrition and Dietetics / Medical Lab Technology / Nursing / Genetics / Agriculture / Industrial Microbiology / Immunology / Molecular Biology / Industrial Biotechnology / Environmental Science / Virology / Bioinformatics / B.S.M.S / B.A.M.S / B.U.M.S. / B.F.Sc./ B.E. or B.Tech in Biotechnology or any other degree that may be considered as equivalent top by the Manonmaniam Sundaranar University are eligible for admission

Subject Type	Code	Subject Title	Hrs. /wee k	L	T	P	С	Maximum Marks		ks	
				Hrs/ wee k	Hrs/ wee k	Hrs/ wee k	Credi ts	Exam hours	Inter nal Asses sment	Exter nal Asses sment	Total
SEMESTER	I										
Core Compu	llsory The	ory Courses									
Core -1		Biochemistry & & Biochemical Techniques	4	4	-	-	4	3	25	75	100
Core-2		Cell & Molecular Biology	4	4	-	-	4	3	25	75	100
Core-3		General Microbiology	4	4	-	-	4	3	25	75	100

Core-4	Microbial Physiology &	4	4	T_	_	4	3	25	75	100
	Metabolism	•								
Core Compulsory	Practical Courses	1				L				
Major	Practical I: Core -1 & Core-	4	-	-	4	2	3	50	50	100
Practical -1	2									
Major	Practical II: Core -3 & Core-	4	-	-	4	2	3	50	50	100
Practical –2	4									
<b>Core Elective The</b>	ory – Any one from the following									
Elective -1	Biostatistics & Computer	3	3	-	-	3	3	25	75	100
	Application									
	Aquatic Microbiology									
Open elective										
ePG-	To be selected latter	3	3	-	-	3	3	25	75	100
Pathsala										
	Sub-Total	30	22	-	8	26		200	600	800
SEMESTER II		•	•	•	•	1		•		•
<b>Core Compulsory</b>	Theory Courses									
Core -5	Bacteriology & Virology	4	4	-	-	4	3	25	75	100
Core-6	Mycology & Phycology	4	4	-	-	4	3	25	75	100
Core-7	Immunology	4	4	-	-	4	3	25	75	100
Core-8	Microbial Genetics	4	4	-	-	4	3	25	75	100
Core Compulsory	Practical Courses					I				
Major	Practical 3: Core-5 & Core-	4	-	-	4	2	3	50	50	100
Practical -3	6									
Major	Practical 4: Core-7 & Core-	4	_	-	4	2	3	50	50	100
Practical – 4	8									
<b>Core Elective The</b>	ory Courses – Any one from the f	ollowi	ng							
Elective 2	Food Microbiology	3	3	-	-	3	3	25	75	100
	Marine Microbial									
	Technology									
<b>Open Elective Cou</b>	ITCA	·		·			·	·	·	·

MOOCs on		3	3	-	-	3	3	25	75	100
line course										
	Sub-Total	30	22	-	8	26		200	600	800
Internship						3				100
SEMESTER III				•						
<b>Core Compulsory</b>	Theory Courses									
Core -9	Recombinant DNA	4	4	-	-	4	3	25	75	100
	Technology									
Core-10	Bioprocess Technology	4	4	-	-	4	3	25	75	100
Core-11	Medical Microbiology	4	4	-	-	4	3	25	75	100
Core-12	Bioremediation	4	4	-	-	4	3	25	75	100
<b>Core Compulsory</b>	<b>Practical Courses</b>			•						
Practical 5	Practical 5: Core-9 & Core-10	4	-	-	4	2	3	50	50	100
Practical 6	Practical 6: Core-11 & Core-12	4	-	-	4	2	3	50	50	100
<b>Core Elective The</b>	ory Courses – Any one from the f	ollowi	ng	I.	<u>'</u>		1	1	1	<b>!</b>
Elective 3	Biosafey, Bioethics & IPR	3	3	-	-	3	3	25	75	100
	Bioinformatics	1								
<b>Open Elective Cou</b>	ırse	·		I.	<u>'</u>		1	1	1	<b>!</b>
MOOCs on-		3	3	-	_	3	3	25	75	100
line course										
	Sub-Total	30	22	-	8	26		200	600	800
SEMESTER IV	•		•		•	•	,		1	•
<b>Open Elective Cou</b>	ırse									
ePG-		3	3	-	-	3	3	25	75	100
Pathsala										
Project	Project & Viva voce	-	-	-	-	9		50	50	100
_	Sub-Total					12		75	125	200
•		93	69	-	24	93		675	1925	2700

#### ASSESSMENT FOR STUDENTS

#### **Internal Assessment (Theory Courses - 25 marks):**

Each course has three internal assessments, each worth for 15 marks. The average of the two best results from each of the three tests would be considered for 15 marks. For each course, the student should submit one 5-mark assignment and present a 5-seminar seminar.

# **External Assessment (Theory Courses - 75 marks):**

Question Paper Pattern at end of each semester - University Examination Pattern

Sub. Code:

#### Title of the course

Semester: Max. Marks : 75

Time : 3 hrs

Section-A (10 X 1 mark = 10 marks)

Answer all the questions in one or two sentences

Section – B (5  $\times$  5 marks = 25 marks)

Answer all the questions (minimum 200 words) by selecting either 'a' or 'b'

Section – C (5 X 8 marks = 40 marks)

Answer all the questions (minimum 500 words) by selecting either 'a' or 'b'

Semester	I Semester
Course Type	Core Compulsory Paper - 1
<b>Title of the Course</b>	BIOCHEMISTRY & BIOCHEMICAL TECHNIQUES

<b>Course Code</b>								
Teaching Hou	rs	60 Hours/ Semester : 4 Hours	/ week					
	BIOC	HEMISTRY & HEMICAL TECHNIQUES				ernal: 25, External		
Course Prereq The student sho		e a basic knowledge on chemistr	y and bio	molecul	es			
CODE:	BI	OCHEMISTRY & BIOCHEM	ICAL	L	T	P	C	
TECHNIQUES				4	-	-	4	
<ul> <li>To understand the basic principles of chemistry &amp; physics related to biology</li> <li>To understand the structures and functions of bio-molecules</li> <li>To provide an depth knowledge of metabolic pathways in the living systems</li> <li>To provide an in-depth knowledge on enzymes and bioenergetics</li> <li>To provide an advanced understanding of the core principles and applications of various techniques used in biology/biotechnology</li> </ul>								
Module 1	Bioc	hemistry in General				8 h	ours	
reaction kinetics	, therm	ecules and chemical bonds. Princodynamics, colligative propertien bonding, hydrophobic inte	s). Stabi	lizing in	teraction	is (Vand	er Waals,	
Module 2	Class	ification and Functions of Bion	nolecules	5		12 h	ours	
· ·		, physiological and biochemical acids, vitamins, minerals and hor		ns of ca	rbohydra	ates, am	ino acids,	
Module 3		Bioenergetics, Carbohydrate, Lipid & Aminoacid Metabolism 16 hours						
		getics-basic principles; Equilibr		-				
		Phosphoryl group transfers and eduction reactions and its import		_	_		-	

Metabolism: General Introduction, types. Carbohydrate metabolism - Glycolytic pathway; Gluconeogenesis; Reciprocal regulation of Glycolysis and gluconeogenesis, Pentose phosphate pathway; Pyruvate Dehydrogenase Complex, its mechanism of action and regulation, Kreb's cycle; Electron Carriers, Electron transport and Oxidative phosphorylation; Photophosphorylation; Synthesis of glycogen, glycogenolysis, metabolic diseases related to carbohydrate metabolism. Lipid metabolism - Mobilization and transport of lipids, Oxidation of lipids: beta, alpha & omega oxidation, oxidation of saturated fatty acids, Oxidation of unsaturated fatty acids, and oxidation of odd chain fatty acids; Formation and oxidation of Ketone bodies; Biosynthesis of saturated fatty acids; synthesis of odd chain and unsaturated fatty acids, regulation of fatty acid biosynthesis; Biosynthesis of triglycerols; Biosynthesis of cholesterol and its regulation, metabolic diseases related to lipid metabolism. Amino acids metabolism - Amino acid degradation: deamination and transamination reactions; alpha ketoglutorate, succinate, fumarate and oxaloacetate pathways of amino acid oxidation; Metabolic fate of amino groups: role of glutamate and glutamine, urea cycle: reactions and regulation, Biosynthesis of amino acids, Regulation of amino acid biosynthesis, genetic defects in amino acid metabolism, metabolic diseases related to amino acid metabolism.

# Module 4 Nucleic Acid Metabolism & Biocatalysts

15 hours

Nucleic acid metabolism - De novo synthesis of purine and pyrimidine nucleotides; Catabolism of purine and pyrimidine; Disorders of purine and pyrimidine metabolism; Integration of metabolism. Biocatalysts - general principles of catalysis; Enzyme characteristics and classifications; monomeric and oligomeric enzymes; Specificity of enzymes; Active sites and binding site of enzymes, energy considerations, enzyme activity and its measurements, factors affecting enzyme activities; Enzyme kinetics: methods for investigating enzyme kinetics, Michaelis-Menton equation; regulatory enzymes, allosteric enzymes and their regulation; Mechanisms of enzyme catalysis: acid-base catalysis and covalent catalysis; Enzyme inhibition and its types, Enzyme activation; Reaction mechanism of enzymes: chymotrypsin, lysozyme, Structure function relationship of enzymes

# Module 5 Biochemical Techniques

9 hours

Principle, working and applications of centrifugation, filtration; chromatography – Paper, TLC, ion exchange, size exclusion, affinity, adsorption, GLC, HPLC; Electrophoresis – Agarose, PAGE – SDS; Spectrophotometric techniques- UV, Visible, IR, NMR and MASS

# ReferenceBooks

- 1. Cooper. T.G., 2011, The Tools of Biochemistry, Wiley India Pvt. Ltd.
- 2. Donald Voet, Judith G. Voet, 2011, Biochemistry, 4<sup>th</sup> Edition, Willey Science.
- 3. Donald Voet, Judith G. Voet, 2018, Biochemistry, Willey Science.

4.	Jain, J.L., Sunjay Jain, Nitin Jain, 2016, Fundamentals of Biochemistry,	ì
	7 <sup>th</sup> Edition, S. Chand & Company Pvt Ltd.	ı

- 5. Keith Wilson, John Walker, 2012, Principles and Techniques of Biochemistry and Molecular Biology, 7<sup>th</sup> edition, Cambridge University Press.
- 6. Nelson, D.L., and Cox, M.M., 2017, Lehninger Principles of Biochemistry, 7<sup>th</sup> Edition, MacMillan International Edition.
- 7. Reginald H. Garrett, Charles M. Grishm, 2013, Biochemistry, 4<sup>th</sup> Edition, Saunders College Publishers.
- 8. Rodwell, V., Bender, D., Anthony Weil, P., Kennelly, P., Botham, K., 2015, Harpers Illustrated Biochemistry, 30<sup>th</sup> Edition, LANGE.
- 9. Rodwell, V., Bender, D., Anthony Weil, P., Kennelly, P., Botham, K., 2018, Harpers Illustrated Biochemistry, 31<sup>st</sup> Edition, LANGE.
- 10. Satyanarayana, U., 2017, Biochemistry, 5<sup>th</sup> Edition, Books and Allied Pvt. Ltd., Kolkata.
- 11. Trevor Palmer, 2008, Enzymes: Biochemistry, Biotechnology and Clinical Chemistry, 5<sup>th</sup> Edition, Horwood Publishing Limited.

After the completion of the course, the student will be able to

Course Outcomes	Cognitive Level
CO1: Understand the fundamental concepts of chemistry, biology, physics and basic concepts about biochemistry at the atomic level	K1, K2
CO2: Explain the classification, and function of biomolesules like carbohydrates, lipids, proteins, nucleic acids etc.	K1, K2
CO3: Identify the different classes of polymeric biomolecules and their monomeric building blocks, analyze and study the chemical and biochemical properties of biomolecules and also understand	K1, K2, K3, K4, K5

the relationships between biological molecules	
CO4: Understand the concepts of enzyme kinetics in living system	K1, K2, K3, K4
CO5: Critically analyze and interpret the results obtained from biological experiments and understanding of solving biological problems using various techniques	K1, K2, K3, K4

# K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create Mapping of CO with PO

	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	Н	Н	M	M	M	L	L	M	L	Н	L	L	L	L	M
CO2	Н	Н	M	M	M	L	L	M	L	Н	L	L	L	L	M
CO3	Н	Н	M	M	M	L	L	M	L	Н	L	L	L	L	M
CO4	Н	Н	M	M	Н	L	L	M	L	Н	L	L	L	L	M
CO5	Н	Н	M	M	Н	Н	M	M	M	Н	L	M	M	L	Н

H- High, M – Medium, L - Low Mapping of CO with PSO

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6
CO1	Н	M	L	L	Н	L
CO2	Н	M	L	L	Н	L
CO3	Н	M	L	L	Н	L

CO4	Н	M	L	L	Н	L
CO5	Н	M	L	L	Н	L

H - High, M - Medium, L - Low

Semester		I Semes	ter					
<b>Course Type</b>		Core Compulsory Paper-2						
Title of the Co	urse	CELL & MOLECULAR BIOLOGY						
<b>Course Code</b>								
Teaching Hou	ırs	60 Hours/ Semester: 4 Hours/ week						
	CELL BIOLO	& GY	MOLECULAR	Credits: 4	Max. Marks: 100 (Internal 25; External: 75)			
Course Prereq	uisites:			<u> </u>				

The student should possess basic knowledge on general aspects about cell and its various organelles. They may have the brief knowledge on the functions of the cell in the body like

transcription, tra	nslation, genetic codes etc.									
CODE:	CELL AND MOLECULAR RIOLOGY	L	Т	P	C					
	CELL AND MOLECULAR BIOLOGY	4	-	-	4					
Course Objectives	<ul> <li>To understand the molecular compon arrangements and role in making the ce</li> <li>To give a clear knowledge on the tracells by various mechanisms.</li> <li>To impart knowledge on the cytoskelet in cell cycle, cell adhesion and commu</li> <li>To give a knowledge on chromoson expression and regulation.</li> </ul>	ell live.  nsport of  ton of cell  nication	nutries	nts in an their con n cells.	d out of					
Module 1	Introduction to Plasma Membrane									
and uniports, sy	uences - Transport across cell membrane- Diffumports and antiport - Membrane potential sport across epithelia.									
Module 2	Cytoskeleton			12 hours						
movements-intrac Cilia and flagella	and microtubules-structure and dynamics - Melular transport, role and kinesin and dynein, - Cell-cell signaling - Cell surface receptors - Signaling from plasma membrane to nucleus.	signal to	ransduc	tion med	chanisms					
Module 3 Cell-cell Adhesion and Communication 12 ho										
Gap junctions ar	omophilic cell-cell adhesion - Ca <sup>++</sup> independent connexons - cell matrix adhesion – integrated cycle - mitosis and meiosis - cyclins and cycletivity.	grins – c	ollagen	- Non-	collagen					
Module 4	Genome Organization			12 ho						

Hierarchy in organization - chromosomal organization of coding and non-coding DNA – regulation of gene expression - mobile DNA - morphological and functional elements of eukaryotic chromosomes - Genetic analysis in Cell Biology

	somes ceneure ununjus in cen ziology	
Module 5	Intracellular Protein Traffic	12 hours
sorting, post-trans mechanisms - bio	on free and bound polysomes - uptake into ER - membra lational modifications - biogenesis of mitochondria, and blogy of cancer - biology of aging - apoptosis-definition	nuclei - trafficking
significance.		
ReferenceBooks	<ol> <li>Aberts, B., Bray, D., Lewis, J, Ratf, M., Roberts, K 2002, Molecular Biology of the Cell, Garland Pt York.</li> <li>Ajoy Paul, 2011, Text Book of Cell and Molecular Allied Pvt. Ltd., Kolkata.</li> <li>De Robertis, E.D.P. and De Robertis, E.M.F., 1995, Biology, 8<sup>th</sup> Edition, B.I. Waverly Pvt. Ltd.</li> <li>Karp, G., 1996, Cell and Molecular biology: Concept John Wiley &amp; Sons.</li> <li>Lewin, B., 2004, Genes, 8<sup>th</sup> Edition, Pearson Prentices.</li> <li>Lodish, H., Berk, A., Matsudaira, P., Kaiser, C.A., M.P., Zipursky, S.L., and. Darnell, J., 2004, Molecular Edition, W.H. Freeman and Company.</li> <li>Watson, J.D., Hopkins, N.H., Roberts, J.W., Steitt A.M., 1987, Molecular Biology of the Gene, 4<sup>th</sup> Cummings.</li> </ol>	Biology, Books and Cell and Molecular ots and Experiments, Hall. Krieger, M., Scott, lar Cell Biology, 5 <sup>th</sup> z J.A., and Weiner,

After the completion of the course, the student will be able to –

Course Outcome	CognitiveLevel
<b>CO1:</b> Understand and analyze the molecular components of plasma membrane of a cell and their arrangement, their role in transport of micro and	K1, K2. K4
macromolecules between its internal and external environment.	

CO2: Know about microtubules and microfilaments in making the cytoskeleton of a cell and how they help in the transport of molecules needed for cell cycle. Also, know about the cell surface receptors and how they work in cell signaling through signal transduction pathway.	K1, K2
<b>CO3:</b> Acquire knowledge on adhesion molecules and structures that make contact between two adjacent cells; know about the fact lying behind on the growth of cells by mitosis and meiosis and the factors involved in cell cycle regulation.	K3, K4, K5
<b>CO4:</b> Know the facts behind the gene organization of chromosomes and how the genes get expressed and regulated. Know about the use of genes in genetic analyses.	K2, K3, K5
<b>CO5:</b> Understand the process of translation and how the translated protein is getting segregated and reach their respective organelles for their suitable function.	K1, K2

K1- Remember; K2- Understand; K3- Apply; K4- Analyze; K5- Evaluate; K6- Create

# **Mapping of CO with PO**

	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	Н	M	M	M	L	M	M	L	L	M	M	M	M	L	L
CO2	Н	M	M	M	L	M	M	L	L	M	M	M	M	L	L
CO3	Н	M	M	M	L	M	M	L	M	M	M	M	M	L	L
CO4	Н	M	M	M	L	M	M	L	M	M	M	M	M	L	L
CO5	Н	M	M	M	L	M	M	L	M	M	M	M	M	L	L

H - High, M – Medium, L - Low

Mapping of C O with PSO

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6
CO1	Н	L	M	L	M	M
CO2	Н	L	M	L	M	M
CO3	Н	L	M	L	M	L
CO4	Н	L	M	L	M	L
CO5	Н	L	M	L	M	L

H - High, M - Medium, L - Low

Semester		I Semester						
<b>Course Type</b>		Core Compulsory Paper-3						
Title of the Co	urse	GENERAL MICROBIOLOG	Y					
<b>Course Code</b>								
Teaching Hou	rs	60 Hours/ Semester : 4 Hours						
						Max. Marks: 100 nternal: 25, External 75)		
Course Prereq	uisites:							
CODE:		GENERAL MICROBIOLOG	·Y	L	T	P	C	
				4	-	-	4	
Course Objectives	under	ecome familiar with the foundarstand the structure and functions stand the microbial diversity in the stand the microbial diversity in the standard	of a typ	ical prok				
Module 1	Histo	ry of Microbiology	of Microbiology 12 hours					

History and Scope of Microbiology – Generation theory – Contribution of Leuwenhoek, Louis Pasteur, Robert Koch, Edward Jenner, Joseph Lister, Winogradsky, Waksman and John Tyndall. Classification of microorganisms - Haeckel's three kingdom concept, Whittaker's five kingdom concept, Carl Woes three domain system, bacterial classification (outline) according to Bergey's manual of systemic Bacteriology.

# Module 2 Morphology of bacteria

12 hours

Morphological types, cell wall of Gram negative, Gram positive bacteria and halophiles, cell wall synthesis, capsule composition and function, cell membranes in eubacteria, archaebacteria and cyanobacteria, cell membrane-functions, periplasmic space, structure and function of flagella, cilia and pili, gas vesicles, chlorosomes, carboxysomes, magnetosomes and phycobilisomes, reserve food materials — polyhydroxybutyrate, polyphosphates, cyanophycin and sulphur inclusions, general account on mycolpasma.

Module 3 Algae 12 hours

General characteristics, classification, structure and reproduction of algae: Chlorophyta (Green algae), Diatoms, Rhodophyta (Red algae); fungi: cell wall – chemical composition and functions, membranes and their functions, nutritional strategies of fungi, structure and life cycle of fungi Ascomycetes (Aspergillus), Zygomycetes (Mucor), Basidiomycetes (Agaricus) and Protozoa.

# Module 4 Virus 12 hours

Discovery, distinctive properties, morphology and ultra-structure of virus, classification, cultivation and purification assay of virus. Bacteriophages - structural organization and life cycle - lytic, lysogenic. Viral related agents - viroid and prion.

# Module 5 Microscopes 12 hours

Principle and application of bright field, dark field, phase contrast, fluorescence, electron microscope- TEM and SEM, polarized microscope and confocal microscopy.

# ReferenceBooks

- 1. Atlas, R.A. and Bartha, R., 2000, Microbial Ecology, Fundamentals and Application, Benjamin Cummings, New York.
- 2. Dubey, R.C., and Maheswari, D.K., 2013, A text book of Microbiology, Revised Edition, S. Chand and Company Ltd, NewDelhi.
- 3. Madigan, M.T., Martinka, M., Parker, J. and Brock, T.D., 2000, 12<sup>th</sup> Edition, Biology Microorganisms, Prentice Hall, New Jerry.
- 4. Mark Wheelis, 2010, Principles of Modern Microbiology, Jones & Bartlett

India Pvt.	Ltd.,	New	Delhi.
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- 5. Pelczar, M.J., Schan, E.C. and Kreig, N.R., 2010, Microbiology An Application Based Approach, 5<sup>th</sup> Edition, Tata McGraw Hill Publishing Company Limited, New Delhi.
- 6. Prescott, L.M., Harley, J.P. and Helin, D.A., 2008, Microbiology, 5<sup>th</sup> Edition, McGraw Hill, New York.
- 7. Schlegal, H.G., 1995, General Microbiology, 7<sup>th</sup> Edition, Cambridge University Press.
- 8. Stanier, R., Lingraham, Y., Wheelis, M.L. and Painter, R.P., 1986, General Microbiology, 5<sup>th</sup> Edition, Macmillan, London.
- 9. Stryer, L., 2010, Biochemistry, Seventh Edition, W.H. Freeman and Company, New York.
- 10. Tortora G.J., Funke, B.R.and Case, C.L., 2009, Microbiology, Ninth Edition, Dorling Kindersely (India) Pvt. Ltd., Noida.

On the successful completion of the course, students will

Course Outcomes	Cognitive Level
CO1: Understand the history and scope of Microbiology, Contribution of different scientist for the development of microbiology and	
different system of classification of microbes	
CO2: Acquire a thorough knowledge on the structure and function of cell wall and cell membrane of gram positive, gram negative and halophilic bacteria	
<b>CO3:</b> Understand and remember structure and reproduction of Algae and Fungi.	K1, K2
<b>CO4:</b> Develop knowledge on virus and bacteriophages. The difference between the lytic and lysogenic life cycle of virus will be analyzed by the students.	
<b>CO5:</b> Understand the principles and function of different types of microscopes and selection of particular microscope for observing different objects	

# K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create

# **Mapping of CO with PO**

	PO1	PO2	PPO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	Н	M	M	M	M	L	M	L	M	M	M	L	M	L	L
CO2	L	M	Н	M	Н	L	M	L	Н	M	M	L	M	L	L
CO3	M	M	Н	M	L	L	L	L	Н	M	M	L	M	M	L
CO4	L	M	M	M	M	L	M	L	M	M	M	L	L	L	L
CO5	M	M	M	M	Н	L	M	L	M	M	M	L	L	M	L

H - High, M - Medium, L - Low

**Mapping of CO with PSO** 

		PSO2	PSO3	PSO4	PSO5	PSO6
CO1	Н	M	M	L	M	L
CO2	Н	M	M	L	M	L
CO3	Н	M	M	L	M	L
CO4	Н	M	M	L	M	L
CO5	Н	M	M	L	M	L

H - High, M - Medium, L - Low

Semester		I Semester							
Course Type		Core Compulsory Pape	er - 4						
Title of the Cou	ırse	MICROBIAL PHYSIC	LOGY	Y AND N	<b>МЕТАВ</b>	OLISM			
<b>Course Code</b>									
Teaching Hou	rs	60 Hours/ Semester: 4	Hours	/ week					
	Micro Metab	abolism (Inte					Max. Marks: 100 ernal: 25, External 75)		
Course Prerequ	uisites:	The student should have	a basic	knowled	ı		I		
CODE:	N	IICROBIAL PHYSIOL		ND	L	T	P	C	
		METABOLISN	1		4	-	-	4	
Course Objectives	the hi	bbiology. This course cov storical perspective to str he environment and their	ucture,	composi	ition of r				
Module 1	Grow	th of Bacteria		12 hours					
induction of sy physiological ad	nchrony aptation	orth kinetics - batch cultury, factors affecting grown to extreme environment of carbon, energy and elections.	wth - ntal co	nutrition onditions	ı, aeratio	on, tem	perature	and pH,	
Module 2	Bacte	erial Photosynthesis		12 h	ours				
structure of ph	otosyn tosynth	general types of micro thetic pigments – chlo etic bacteria - green sulph	orophyl ur and j	ls, bact purple; n	eriochloi nechanis	rophyll, m of pho	caroten otosynthe	oids and esis - non-	
		on transport, photophosp roxyl propionate cycle.	nioi yiai	ion; car	bon assii	milation	- Calvi	n, reverse	

Nitrogen cycle - ammonification, nitrification, denitrification and nitrogen fixation; nitrogenase enzyme, physiology of nitrogen fixation in symbiotic and free living bacteria, genetics of nitrogen fixation, acetylene reduction assay, transamination and deamination.

### Module 4 Microbial Stress Responses

12 hours

Osmotic stress and osmoregulation; aerobic to anaerobic transitions; oxidative stress; pH stress and acid tolerance; thermal stress and heat shock response; nutrient stress and starvation stress. Fermentative pathways in specific group of microbes: alcoholic, lactic acid, formic, mixed, propionic, butyric, butanol, butanediol fermentation. Anaerobic respiration.

#### Module 5 Bioenergetics

12 hours

Principles and laws of thermodynamics, coupling of chemical reactions - TCA cycle, electron transport chain, and chemiosmotic theory of Mitchell. Biomembranes: Fluid mosaic model, transport across membrane - diffusion, osmosis, active transport and group translocation.

- 1. Deb, A.C., 2006, Fundamentals of Biochemistry, New Central Book Agency Pvt. Ltd., Kolkata.
- 2. Donald Voet and Judith G. Voet, 2011, Biochemistry, 3<sup>rd</sup> Edition, John Wiley and Sons, Inc. New York.
- 3. Madigan, M.T., Martinka, M., Parker, J. and Brock, T.D., 2000, Biology Microorganisms, 12<sup>th</sup> edition, Prentice Hall, New Jerry.
- 4. Moat, A.G. and Foster, W., 2002, Microbial Physiology, 4<sup>th</sup> Edition, John Wiley and Sons, New York.

# 5. Nelson, D.L. and Cox, M.M., 2012, Lehingers's Principles of Biochemistry, Sixth Edition, Mac Millan worth Publishers, New Delhi.

- 6. Postgate, J., 1998, Nitrogen Fixation, 3<sup>rd</sup> Edition, Cambridge University Press.
- 7. Salisbury, F.W. and W. Ross, 1992, Plant Physiology, 4<sup>th</sup> Edition, Wardsworth Publishing Company, California.
- 8. Satyanarayana, U. and Chakrapani, U., 2013, Biochemistry, 4<sup>th</sup> Edition Book and Allied Pvt. Ltd., Kolkata.
- 9. Srivastava, M.L., 2008. Microbial Biochemistry, Narosa Publishing House, New Delhi.
- 10. Stryer, L., 2010, Biochemistry, 7<sup>th</sup> Edition, W.H. Freeman and Company, New York.

#### ReferenceBooks

After the completion of the course, the student will be able to

Course Outcomes	Cognitive Level
CO1: Understand the scope and importance of Microbiology, classification schemes, cultivation, preservation and maintenance of microbial cultures. microbial structure and function and study the comparative characteristics of prokaryotes and eukaryotes and also understand the structural similarities and differences among various physiological groups of eubacteria/archaea. : Master skills in aseptic techniques as well comprehend the importance of cleaning and decontamination.	K1, K2, K5, K6
CO2: Understand the microbial transport systems and the modes and mechanisms of energy conservation in microbial metabolism — Autotrophy and heterotrophy Know the various Physical and Chemical growth requirements of bacteria and get equipped with various methods of bacterial growth measurement	K1, K2
CO3: Understand the phases of growth, requirements and factors affecting growth; adaptation to extreme environmental conditions. Nutritional types and metabolic diversity	K1, K2
CO4: Understanding History and types of photosynthesis, state the difference between oxygenic and anoxygenic photophosphorylation. Knowledge on different mechanism of photosynthesis	K1, K2

CO5: Design a mechanism that would allow a bacterium to protect its nitrogenase from oxygen. Analyze the symbiotic relationship that some N<sub>M</sub>-fixing bacteria have with plants. Identify what the bacteria contribute and what the plant contributes. Describe the process of methanogenesis in terms of electron transport and energy generation

# K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create

Mapping of CO with PO

	PO1	PO2	PPO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	Н	M	M	M	Н	M	M	L	M	M	M	M	M	M	L
CO2	Н	M	M	M	Н	M	M	L	M	M	M	M	L	M	L
CO3	Н	M	M	M	Н	M	M	L	M	M	M	M	L	M	M
CO4	Н	M	M	M	Н	M	M	L	M	M	M	M	L	M	M
CO5	Н	M	M	M	Н	M	M	L	M	M	M	M	L	M	M

# H - High, M – Medium, L - Low Mapping of CO with PSO

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6								
CO1	Н	Н	Н	M	Н	M								
CO <sub>2</sub>	Н	M	M	Н	M	M								
CO3	Н	M	L	M	M	L								
CO4	M	Н	M	M	Н	L								
CO5	Н	M	M	M	Н	M								

H - High, M - Medium, L - Low

Semester		I Semester											
Course Type		Core Comp	ulsory Course Pra	ctical - 1									
Title of the Cou	ırse		ISTRY & BIOCH		L TECH	<b>INIQUE</b>	ES AND	CELL					
		& MOLECU	ULAR BIOLOGY	·									
<b>Course Code</b>													
<b>Teaching Hour</b>			mester: 4 hours/w			T							
		HEMISTRY		Credi	ts: 2		Max. Marks: 100						
		HEMICAL	`										
			MOLECULAR			E	xternal:	50)					
	BIOL	OGY											
Course Prerequ						_							
			cs of biochemistry	/ & knov	vledge o	on the c	ell struc	ture and					
functions of var								1					
CODE:		CHEMISTRY			L	T	P	C					
		HNIQUES	AND CEL	L &	_	_	4	2					
	+	ECULAR BI		C	• ,		1	<u> </u>					
	•	• To inculcate/impart skills to perform various tests/assays and experiments.											
	•	To provide qualitative & quantitative analysis of the macromolecules in											
		the given sample and analyze the results.											
	•	• To provide students with a deep insight of the various biochemical reactions and cellular processes through quantitative and qualitative											
Course			the samples provide		ugn qu	anmanv	e and c	quantative					
Objectives	•	•	nd different types o		d their s	tructure							
	•		the biochemical par										
	•	•	e karyotyping of o				d the m	itosis and					
		meiosis proc		01 <b>5</b>	, 00110			100010 4110					
1. Qualitativ	e analy		carbohydrate, lipio	d and nuc	leic acid	ls							
		total carbohy											
			y ninhydrin metho	d									
			owry's method										
5. Quantific	ation of	RNA by Orci	inol method										

- 6. Quantification of DNA by diphenyl amine method
- 7. Analysis of DNA by Agarose gel electrophoresis (Demonstration)
- 8. Separation of any biomolecule by Thin Layer Chromatography
- 9. To study different phases of mitosis in Onion root tip.
- 10. To study meiosis in Grasshopper testis.
- 11. Human buccal smear to show squamous epithelial cells
- 12. To study the Barr body from the smear of buccal epithelial cells (FEMALE)
- 13. Preparation of polytene chromosomes of Chironomous larva/Drosophila.

After the completion of the course, the student will be able to

Course Outcomes	Cognitive Level
CO1: Improved skills to perform various tests/assays and experiments.	K1, K2, K3, K4, K5
CO2: Design and analyze the experiments related with the different molecules and use of the various techniques the kinetics and rationale behind each phenomenon	K1, K2, K3, K4, K5, K6
CO3: Use the techniques, skills, and modern tools necessary for imbalances in various life processes, design a research project, collect and analyze data, and interpret results	K3, K4, K5
<b>CO4:</b> Analyze the mechanism of mitosis, meiosis and enumerate the structure of polytene chromosome of chironomous larvae.	K4, K3

# K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create

Mapping of CO with PO

PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	<b>PO12</b>	PO13	<b>PO14</b>	PO15

CO1	Н	Н	M	L	Н	Н	Н	L	M	M	Н	Н	Н	Н	M
CO2	Н	Н	M	L	Н	Н	Н	L	M	M	Н	Н	Н	Н	M
CO3	Н	Н	M	L	Н	Н	Н	L	M	M	Н	Н	Н	Н	M
CO4	Н	Н	M	L	Н	Н	Н	L	M	M	Н	Н	Н	Н	M
CO5	Н	Н	M	L	Н	Н	Н	L	M	M	Н	Н	Н	Н	M

H - High, M – Medium, L - Low

**Mapping of CO with PSO** 

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6
CO1	Н	M	L	L	Н	L
CO2	Н	M	L	L	Н	L
CO3	Н	M	L	L	Н	L
CO4	Н	M	L	L	Н	L
CO5	Н	M	L	L	Н	L

H - High, M - Medium, L - Low

Semester		I Seme	I Semester							
<b>Course Type</b>		Core Compulsory Course Practical - 2								
Title of the Co	urse	GENERAL MICROBIOLOGY AND MICROBIAL PHYSIOLOGY								
		& METABOLISM								
<b>Course Code</b>										
<b>Teaching Hour</b>	:S	60 hours/semester: 4 hours/week								
	GENE	RAL MICROBIOLOGY Credits: 2 Max. Marks: 100								

	AND MICROBIAL PHYSIOLOGY & METABOLISM		(Internal: 50; External: 50)										
Course Prerequisites:													
The students should know the basics skills on microbiology practicals													
6, F-11-11-11-11-11-11-11-11-11-11-11-11-11													
CODE:	GENERAL MICROBIOLOGY AN	ND L		T	P	C							
CODE:	MICDODIAL DUVCIOLOGY	0											
	MICROBIAL PHYSIOLOGY	&											
	METABOLISM	-		-	4	2							
		-	quire	- adeq	-	2 1 required							

and their growth patterns in different media. The student will also get a thorough

input to analyse end evaluate the difference between different microorganisms.

- 1. Gram's staining.
- 2. Negative staining.
- 3. Simple staining.
- 4. Capsule (Spore) staining.
- 5. AF staining.

Course

**Objectives** 

- 6. Motility of bacteria by Hanging drop method.
- 7. Biochemical test
- 8. Bacterial growth determination by spectrophotometer method.
- 9. Isolation of bacteria from soil and sea water.
- 10. Isolation of fungi from soil and Marine products
- 11. Principle and methods of sterilization
- 12. Preparation of media: nutrient broth, nutrient agar plate, soft agar
- 13. Pure culture techniques: streak plate, spread plate and pour plate
- 14. Motility determination Hanging drop method
- 15. Isolation and enumeration of bacteria from different environmental samples
- 16. Enumeration of bacteria viable count (plate count) and total count (Haemocytometer count) Direct microscopic observation of fungal spores and mycelium
- 17. Staining method: simple, negative, Gram's staining and spore staining
- 18. Fungal slide culture
- 19. Measurement of growth rate and generation time by turbidometry method

After the completion of the course, the student will be able to

Course Outcomes	Cognitive Level
CO1: Able to use different staining methods to differentiate bacteria, bacterial spores and capsules.  Apply different biochemical test for identifying bacteria up to genus level.	K1, K2, K3, K4, K5
CO2: Able to isolate microalgae and fungi from natural environment and preparation of different culture media for growing bacteria, algae and fungi.	K1, K2, K3, K4, K5
CO3: Understand various physical and chemical means of sterilization -Sterilization techniques. Competently prepare different types of media	K1, K2, K3, K5
CO4: Understanding the skill to perform different methods of isolation, enumeration, maintenance and preservation of microorganisms, and skill for viable counting	K1, K2, K3, K5
CO5: Understanding, the knowledge and skill to analyse and differentiate between different types of Microorganisms based on their staining, motility characteristics	K1, K2, K3

# K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create Mapping of CO with PO

	upp	<u> </u>	00 1110													
		PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
C	01	Н	Н	Н	Н	Н	M	Н	Н	Н	M	Н	M	Н	L	M

CO2	Н	Н	Н	Н	Н	M	Н	Н	Н	M	Н	M	Н	L	M
CO3	Н	Н	Н	Н	Н	M	Н	Н	M	M	Н	M	Н	L	M
CO4	Н	Н	Н	Н	Н	M	Н	Н	M	M	Н	M	Н	L	M
CO5	Н	Н	Н	Н	Н	M	Н	Н	M	M	Н	M	Н	L	M

H- High, M – Medium, L - Low Mapping of CO with PSO

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6
CO1	Н	Н	Н	M	Н	L
CO2	Н	Н	Н	M	Н	L
CO3	Н	Н	Н	M	Н	L
CO4	Н	Н	Н	M	Н	L
CO5	Н	Н	Н	M	Н	L

H - High, M - Medium, L - Low

Semester		I Semester							
<b>Course Type</b>		Core Elective Paper 1 A							
Title of the Co	urse	BIOSTATISTICS & COMPUTER APPLICATION							
<b>Course Code</b>									
Teaching Hou	rs	45 Hours/ Semester: 3 Hours/ week							
	<b>BIOS</b>	TATISTICS & COMPUTER	Credits: 3	Max. Marks: 100					
	APPL	ICATION		(Internal: 25, External					
				75)					

**Course Prerequisites:** The student should possess basic knowledge on general aspects about biostatistics, statistical data representations, statistical measures and analysis of biological data, as well as to know the available statistical tools and interpretation of statistical data through computer software.

computer softw	rare.									
CODE:	BIOSTATISTICS AND COMPUTER	T	P C							
	APPLICATION	-	3							
<ul> <li>To know the basic concepts of biostatistics like history &amp; growth of statistics and statistical methods</li> <li>To understand about data, their types, and methods involved in collection of data and presentation of data through various modalities.</li> <li>Enable to understand the statistical measures through determination of averages, deviations, test of significance, hypothetical analysis correlation &amp; Regression analysis, etc.</li> <li>To understand about basic computer application in relation with analysis of biological data by using various computer software technique.</li> </ul>										
Module 1	Statistical Introduction		9 hours							
	atistics, History and growth of statistics, Statist, Samples from populations and Random samplin		nods, T	ypes of	biological					
Module 2	Data Interval and Representation			9 ho	ours					
_	nization and tabulation of data, Diagrammatic aphical representation of data.	represen	itation	of data,	Types of					
Module 3		9 hours								
	ntral Tendency- mean, median and mode, Measu		-		-					
	on measured with quartiles, mean deviation,	variance	and st	tandard	deviation,					
•	neans and variances.  Statistical Hypothesis and Analysis			0.7						
Module 4		<u>9</u> h(	ours							

Proportion of data – Examples of proportion data, Statistical treatment of proportion of data, Chi square test and goodness of fit, Application of Chi square test, Sampling and Hypothesis: One and Two sample hypothesis, Test of Significance, Analysis of Variance: One way and Two way ANOVA classification, Regression and Correlation analysis.

ANOVA classification, Regression and Correlation analysis.												
Module 5	Introduction to Computer	9 hours										
Basic Computer A	Applications: Purpose of computer, types of computer, Hard	ware and Software,										
Programming lang	guage, Commercial software: Windows, MS Word, Excel, Pov	ver point, Statistical										
packages: Sigma s	packages: Sigma stat, SPSS-Intra and Internet, Email, Website Creation, Database in Biology: Pul											
Med, Sequence Analysis, Genome and Protein database genome research.												
	1. Bhose, S.B., 2011, Text book of Computer Application and Biostatist											
	Trinity publishing House, India.											
	2. Bliss, C.I.K., 1967, Statistics in Biology, Vol. I, Mc Gr	aw Hill, New York.										
	3. Campbell, R.C., 1974, Statistics for Biologists, Car	mbridge University										
	Press, New York.											
	4. David Baskeen, 2008, Introduction to Computer Application	cation and Concept,										
	Cengage Learning Public, UK.											
ReferenceBooks	5. Gupta, S.P., 2010, Practical Statistics, S. Chand at	nd Company, New										
	Delhi.											
6. Hand Book of Experimental Immunology, Blackwell Publica												
Oxford. 7. Jerrold H. Zar, 2009, Biostatistical Analysis, 5 <sup>th</sup> Edition,												
										Publications, India.		
	8. Lutz, W., 1967. Statistics Methods as Applied to Immu	nological data, app.										
	In: D.M. Weir (ED).											

After the completion of the course, the student will be able to

Course Outcomes	Cognitive Level
CO1: Understand about the basic concepts of Biostatistics, its history and development, meaning of data, different types of statistical data, etc.	K1, K2

CO2: Know about how to collect biological data and its presentation by following various methodologies like tabulation, diagrammatic and graphical representations.	K2, K3
CO3: Interpret and determine the biological data through some basic statistical measures like measures of Central tendency, Dispersion and variation analysis.	K3, K4, K5
CO4: Test and analyse the biological data through hypothetical assumption or creation by following some statistical treatments with Chi square test, students 't' test, ANOVA test, Correlation and Regression analysis	K4, K5, K6
CO5: Know about computer application in statistical data analysis, study about available basic software's & hardware's and statistical packages related to data base in biology.	K3, K4, K6

K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create

# **Mapping of CO with PO**

	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	L	Н	M	L	M	Н	L	M	L	Н	M	L	Н	L	M
CO2	L	Н	M	L	M	Н	L	M	L	Н	M	L	Н	L	M
CO3	L	Н	M	L	M	Н	L	M	L	Н	M	L	Н	L	M
CO4	L	Н	M	L	M	Н	L	M	L	Н	M	L	Н	L	M
CO5	L	Н	M	L	M	Н	L	M	L	Н	M	L	Н	L	M

H – High, M – Medium, L - Low Mapping of CO with PSO

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6
CO1	L	L	L	L	L	Н
CO2	L	L	L	L	L	Н
CO3	L	L	L	L	L	Н
CO4	L	L	L	L	M	Н
CO5	L	L	L	L	M	Н

H - High, M - Medium, L - Low

Semester		I Semester									
<b>Course Type</b>		Core Elective Paper – 1B									
Title of the Co	urse	AQUATIC MICROBIOLOGY									
Course Code											
Teaching Hou	rs	45 Hours/ Semester: 3 Hours/ we	eek								
	AQUA	ATIC MICROBIOLOGY	Credi	its: 3	Max. Marks: 100 (Internal: 25, External 75)						
Course Prerequisites: Have a basic knowledge on microbiology											
CODE:		A OLIA TRICIA MICROPIOLOGNI		L	T	P	C				
		AQUATIC MICROBIOLOGY		3	-	-	3				

		1								
Course Objectives	<ul> <li>To know about the composition of waste water, the biological and chemical</li> <li>To get an idea about the water disinfection, wasted drinking water treatment and biotechnological appropriate treatment</li> </ul>	te water treatment,								
Module 1	Characteristics of Water & its Composition	9 hours								
water, composition	eristics of water, microorganisms in sewage and seawater, saper of domestic waste water, COD, BOD, total organic carbon cases, indicator microorganisms.	1 0								
Module 2 Water and waste water disinfection 9 hours										
Water and waste water disinfection – factors influencing disinfection, type of disinfectants, their mode of action, toxicology.										
Module 3	Waste water treatment	9 hours								
foaming, biofilm dewatering, condit	process – biology of sludge, nutrient removal, pathogen reactors, waste stabilization ponds. Sludge processing – scritions, stabilization, composting, anaerobic digestion of waste and bio-odours from waste water treatment plants and their control of the stabilization of the stabili	reening, thickening, e water and sludge,								
	Drinking water treatment	9 hours								
Drinking water tactivated carbon to formation, problem	reatment – storage, prechlorination, coagulation, water s reatment, biological treatment, disinfection. Water distributions caused by biofilms. Other biological problems associated wome devices for water treatment.	on system – biofilm								
Module 5	Biotechnological applications of waste treatment	9 hours								
	Biotechnological application in waste treatment – bioagumentation, use of enzyme, use of immobilized cells, biosensors, application of recombinant DNA technology for waste water reuse.									
1. Gabriel Britton (Ed.), 1994, Waste water Microbiology, Wiley – Liss Publications.  2. Purohit, S.S., 2000, Microbiology Fundamental and Applications, 6 <sup>th</sup>										

Edition, Agrobios Publishers.

- 3. Chatwall, G.R., (Ed), 2003, Encyclopedia of Environmental Water Pollution, Vol. 1 3, Chatwall, Annol Publications Pvt. Ltd., New Delhi.
- 4. Gerhard Rheinheimer (Ed.), 1986, Aquatic Microbiology Methods, 4<sup>th</sup> Edition, John Wiley & Sons.
- 5. Ian T. Paulsen and Andrews J. Holmes (Eds.), 2013, Environmental Microbiology-Methods and Protocols, 2<sup>nd</sup> Edition, Humana Press.

After the completion of the course, the student will be able to

Course Outcomes	Cognitive Level
<b>CO1:</b> Understand the characteristics of water and waste water and the biological and chemical indicators	K1, K2, K3, K4
CO2: Factors affecting water & waste water treatment, their types and mode of action	K1, K2, K3
CO3: Understand how to treat the waste water, its methods and its implications	K2, K3, K4, K5
<b>CO4:</b> Understand how to treat the drinking water, its methods and its implications	K2, K3, K4, K5
CO5: Biotechnological applications in water treatment	K3, K4, K5

## K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create

Mapping of CO with PO

		PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
C	01	Н	M	M	M	L	M	M	M	L	L	L	L	L	L	M

CO2	Н	M	M	M	L	M	M	M	L	L	L	L	L	L	M
CO3	Н	M	M	M	L	M	M	M	L	L	L	L	L	L	M
CO4	Н	M	M	M	L	M	M	M	L	L	L	L	L	L	M
CO5	Н	M	M	M	L	M	M	M	L	L	L	L	L	L	M

H – High, M – Medium, L - Low Mapping of CO with PSO

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6
CO1	Н	M	Н	M	L	L
CO2	Н	M	Н	M	L	L
CO3	Н	M	Н	M	L	L
CO4	Н	M	Н	M	L	L
CO5	Н	M	Н	M	L	L

H - High, M - Medium, L - Low

Semester		II Semester						
<b>Course Type</b>		Core Compulsory Paper - 5						
Title of the Course BACTERIOLOGY & VIROLOGY								
<b>Course Code</b>								
<b>Teaching Hou</b>	rs	60 Hours/ Semester : 4 Hours/ week						
	BACT	TERIOLOGY & VIROLOGY	Credits: 4	Max. Marks: 100				
				(Internal: 25, External				
				75)				

Course Prerequ	isites: The students should have a basic knowled	dge on m	icrobiol	ogy									
CODE:		L	Т	P	С								
	BACTERIOLOGY & VIROLOGY	ACTERIOLOGY & VIROLOGY  4											
Course Objectives	<ul> <li>To introduce knowledge on the structure of bacteria, viruses,</li> <li>Provide knowledge on fundamentals of bacterial &amp; viral classification</li> <li>Develop understanding of infection processes, host range, cultivation and purification assays using molecular techniques</li> </ul>												
Module 1	Morphology and Structure of Bacteria			12 h	ours								
cilia – pili – gas v <b>Module 2</b>	nbranes – structure – composition, properties. sesicles – chromosomes  Classification of Microorganisms  eckel's three kingdom concept – Whitaker's fiv			12 h	ours								
concept of Carl	Woese, basis of microbial classification – clang to the Bergey's Manual of Determinat	ssificatio	n and s	salient f	eatures of								
Module 3	History, Morphology and Structure of Virus			12 h									
origin and evolusedimentation, elerange, transmission viruses: Criteria bacteria, plants ar plant and animal embryonated eggs cell cultures, cell s		sical- motic nature fomenclate ICTV classay and cultures	orphologory of virus and assificated mainter.  Expense	gy and ses, biological classification of vinance of rimental on and r	structure, ogical host ication of viruses of bacterial, animals, monolayer								
Module 4	Purification of viruses			12 h	ours								

Need for virus purification. Extraction of viruses from tissues, clarification, concentration of viruses in clarified extracts by physical and chemical methods, further purification of viruses by rate zonal / equilibrium density gradient centrifugation, criteria of virus purity, quantitation and preservation of purified virus preparations. Quantitation of viruses: Infectivity assay methods (plaque, pock, end point, local / systemic assay of plant viruses), physical (EM), serological (HA, HI, immunofluorescence, ELISA) and chemical (viral protein and nucleic acid based) approaches. Major characteristics of the following virus families / genera / groups: Adenoviridae, Bromoviridae, Bunyaviridae, Caulimoviridae, Flaviviridae, Geminiviridae, Hepadnaviridae, Herpesviridae, Orthomyxoviridae, Paramyxoviridae, Parvoviridae, Picornaviridae, Potyviridae, Poxviridae, Reoviridae, Retroviridae, Rhabdoviridae, Tobamovirus, Insect Viruses: Biology of major RNA and DNA viruses of insects and their applications.

### Module 5 Bacteriophages

12 hours

Biology of major RNA (MS2, Q $\beta$ , Ø $\delta$ ) and DNA (T-even and T-odd, lambda, Mu, Øx174, M13) bacteriophages, biology of cyanophages. Algal and fungal viruses: Biology of viruses of Phycodnaviridae, Partitiviridae and Totiviridae. Biology of sub-viral agents: Satellite viruses, sat-RNAs, viroids virusoids and prions, Concept of molecular parasitism, importance of viruses in human welfare with suitable examples.

- 1. A Practical Guide to Clinical Virology, 2002, 2<sup>nd</sup> edition, Haahcim, Pattison & Whitley-Wiley
- 2. Bergey's Manual of Systemic Bacteriology
- 3. Corat, H.F., Kimball, P.C., and Levy, J.A., 1994, Virology, 3<sup>rd</sup> Edition, Blackwell Scientific Publications, Oxford.
- 4. Dimmock, N.J., and Primrose, S.B., 1994, Introduction of Modern Virology, 4<sup>th</sup> Edition, Blackwell Scientific Publications, Oxford.
- 5. John B Carter, 2013, Virology: Principles and Applications: A Review, John Wiley & Sons.
- 6. Vinod Singh, 2010, Text of Bacteriology, International Book Distributing Co.
- 7. Lennetter, E.H., 1984, Diagnostic Procedures for Viral and Rickettsial Diseases, American Public Health Association, NY.
- 8. Mathews, R.E., 1992, Functional of Plant Virology, Academic Press, San Diego.
- 9. Morg, C.and Timbury, M.C., 1994, Medical Virology, 10<sup>th</sup> Edition,

## ReferenceBooks

Churchil Livingston, London.

- 10. Topley and Wilson, 1995, Text book on Principles of Bacteriology, Virology and Immunology, Edward Arnold, London.
- 11. Williams Hayes, 1985, The Genetics of Bacteria and Their Viruses, Blackwell Scientific Publishers, London.

After the completion of the course, the student will be able to

Course Outcomes	Cognitive Level
CO1: Knowledge on how to compare and contrast the structure of	K1
cell membranes and cell walls in Bacteria and Archaea. To	
differentiate between Gram positive and Gram negative bacteria;	
Explain how specialized structures (e.g., pili/fimbriae, capsules,	
lipopolysaccharides, spores, or flagella) enable a microbe to	
survive in a given environment	
CO2: Gives understanding on the classification of microbes	K1
CO3: Understanding on nomenclature and classification of	K1, K3
viruses-bacterial viruses, plant viruses; knowledge about viruses	
and the chemical nature of viruses, different types of viruses	
infecting animals, plants and bacteria; Deep knowledge and	
understanding and skill on the principal purposes of cultivating	
viruses. Describe three ways in which viruses are cultivated	
CO4: Understand the knowledge of skills on methods of	K1, K3, K6
purification of viruses; the possible causative agents, modes of	
transmission, virulence factors, diagnostic techniques and	
prevention/treatment for different viral diseases of human, plant	

K2, K1

K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create Mapping of CO with PO

	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	Н	M	M	M	M	M	M	Н	Н	M	M	M	L	L	M
CO2	Н	M	M	M	M	M	M	Н	Н	M	M	M	L	L	M
CO3	Н	M	M	M	M	M	M	Н	Н	M	M	M	L	L	M
CO4	Н	M	Н	M	M	M	M	Н	Н	M	M	M	L	L	M
CO5	Н	M	M	M	M	M	M	Н	Н	M	M	M	L	L	M

H – High, M – Medium, L - Low Mapping of CO with PSO

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6
CO1	Н	Н	M	L	L	L
CO2	Н	Н	M	L	L	L
CO3	Н	Н	M	L	L	L
CO4	Н	Н	M	L	L	L
CO5	Н	Н	M	L	L	L

# $\boldsymbol{H}$ - High, $\boldsymbol{M}-Medium,\,\boldsymbol{L}-Low$

Semester		II Semester									
<b>Course Type</b>		Core Compulsory Paper - 6									
Title of the Co	ırse	MYCOLOGY & PHYCOLO	MYCOLOGY & PHYCOLOGY								
<b>Course Code</b>											
Teaching Hou	rs	60 Hours/ Semester : 4 Hours/ week									
	MYC	OLOGY & PHYCOLOGY		Iax. Marks: 100 ernal: 25, External 75)							
Course Prereq	uisites:	The students should have a gene	eral know	ledge on	microb	oiology					
CODE:		Mygol ogy a byygol og	787	L	T	P	C				
		MYCOLOGY & PHYCOLOG	OGY & PHYCOLOGY 4 -								
Course Objectives		udy the distribution, life cycle as enrich the learners on isolation					and algae				
Module 1	Morp	ohology & Life Cycle				12 hours					
Acrasiomycetes,	Hydron ycetes,	gy, classification, structure and comyxomycetes, Myxomycetes, P Oomycetes, Zygomycotina – ii.	lasmo-di	ophorom	ycetes,	Chytridi	omycetes,				
Module 2	Class	ification of Fungi				12 h	ours				
Ascomycotina – Hemiascomycetes, Plectomycetes, Pyrenomycetes, Discomycetes, Laboulbenomycetes, Loculoascomycetes, Basidiomycotina, Teliomycetes, Hymenomycetes. Deuteromycotina – Hypomycetes, Coelomycetes, Blastomycetes.											
Module 3	Fung	al Diseases				12 h	ours				
Heterothalism, s	ex horr	mones in fungi; physiological s	pecializa	tion phy	logeny	of fungi;	Lichens-				

classification of lichens, habit and habitat and economic importance. Mycorrhizaecto mycorrhia, endomycorrhiza, vesicular arbuscular mycorrhiza. fungi as insect symbiont, fungal diseases systemic and subcutaneous, mycoses, candidiasis, pneumocystis, blastomycoses, deterophytosis. Module 4 Microalgae 12 hours Isolation of microalgae, commercially important phytoplankton, commercial production of microalgae, photobioreactor, harvesting technology, phytoplankton in aquaculture industry, oil industry and carotenoid industry. 12 hours Module 5 Algae Distribution of algae, classification of algae, algal nutrition, algal structure and reproduction. Chlorophyceae, Pheophyceae, Rhodophyceae, Diatoms, Euglenoids, algal ecology and algal biotechnology. 1. Alexopoulos, C.J., Mims, C.W., and Blackwell, M., 1996, Introductory Mycology, 4<sup>th</sup> Edition, John and Sons, Inc. 2. Alexopoulos, C.J. and C.W. Mims, 1979, Introduction to Mycology, 3<sup>rd</sup> Edition, Wiley Eastern Ltd., New Delhi. 3. Gopal Bhattacharya, 2013, Text Book of Mycology, Neha Publishers & Distributors. 4. Kumar, H.D., 1990, Introductory Phycology, 2<sup>nd</sup> Edition, Affiliated East Western Press. Mehrotra, R.S. and K.R. Anexia, 1990, An Introduction to ReferenceBooks Mycology, New Age International Publishers. 5. Sreekumar, S., 2015, Microbiology, Phycology, Mycology, Lichenology and Plant Pathology, Medtech Publishers. 6. Vashishta, B.R. and Sinha, A.K., 2008, Fungi, S Chand and Company Ltd. New Delhi. 7. Vashishta, B.R., 2005, Algae, 3<sup>rd</sup> Edition, S Chand and Company Limited. New Delhi.

After the completion of the course, the student will be able to

Course Outcomes	<b>Cognitive Level</b>

CO1: Acquire knowledge on morphology and life cycle of	K1, K2
different divisions of fungi.	
CO2: Develop ideas on the division of fungi such as	K1, K2
Ascomycotina, Basidomycotina and Deuteromycotina	
CO3: Obtain knowledge on Lichens, Mycorrhiza and Fungal	K1, K2, K3, K4
diseases such as cutaneous, subcutaneous and systemic.	
CO4: Acquire knowledge on isolation, culture, harvest and	K1, K2
commercial importance of algae.	
CO5: Understand about the distribution, classification, structure	K1, K2, K4
and reproduction of algae.	

K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create

Mapping of CO with PO

	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	M	L	Н	M	L	L	L	L	L	M	M	M	L	L	L
CO2	M	L	Н	M	L	L	M	L	L	M	M	M	L	L	L
CO3	M	L	Н	Н	Н	L	M	M	Н	M	M	M	L	L	L
CO4	M	L	Н	M	Н	L	M	M	Н	M	M	M	L	Н	L
CO5	M	L	Н	L	M	L	M	M	Н	M	M	M	L	Н	L

**Mapping of CO with PSO** 

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6
CO1	Н	M	M	L	L	L
CO2	Н	M	M	L	L	L
CO3	Н	M	M	L	L	L
CO4	Н	M	Н	Н	L	L
CO5	Н	M	M	L	L	L

H - High, M - Medium, L - Low

Semester		II Semester								
Course Type		Core Paper - 7								
Title of the Cou	ırse	IMMUNOLOGY								
<b>Course Code</b>										
Teaching Hou	rs	60 Hours/ Semester : 4 Hours	/ week							
	IMMU	UNOLOGY	Cred	x. Marks: 100 Internal: 25, External: 75)						
Course Prerequ	uisites:									
CODE:				L	T	P	С			
		IMMUNOLOGY		4	-	-	4			
Course Objectives										
Module 1 Immunology- Fundamental Concepts and Anatomy 12 hour							ours			

Components of innate and acquired immunity; phagocytosis; complement and inflammatory responses; organs and cells of the immune system - primary and secondary lymphoid organs and tissues; haematopoiesis- immune cells; antigens - immunogens, haptens and super antigens.

## Module 2 Immune Components and Vaccinology

12 hours

Lymphokines and cytokines; immunoglobulins-structure, classes and biological functions; genetic organization of immunoglobulin genes and generation of antibody diversity; applications and monoclonal antibodies; major histocompatibility complex - mhc genes, structure and functions; immunization practices: active and passive immunization; vaccines.

### Module 3 Immune Effector Responses

12 hours

Humoral immune responses - B cell maturation, activation and differentiation and humoral immune response; cell mediated immune response- T cell development and activation, mechanism of cell mediated immune response; T-cell and B-cell receptors; antigen processing and presentation; antigen recognition; interaction of T-cell and B-cell; immunological memory and immunotolerance.

## Module 4 Antigen-Antibody Interactions

12 hours

Affinity, avidity, lattice theory; immuno precipitation and agglutination techniques, immunoelectrophoresis, radio-immunoassay, ELISA, immunoblotting, immunofluorescence, flow cytometry; monoclonal antibodies and hybridoma technology; CMI techniques- lymphoproliferation assay, mixed lymphocyte reaction, cell cytotoxicity assays, apoptosis.

Module 5 12 hours

Immunity to infection: Bacteria, viral, protozoan and parasitic infections; hypersensitivity- Type I-IV- detection methods; autoimmunity- mechanism, types of autoimmune diseases; Transplantation immunology, immunological basis of graft rejection —tissue matching and immuno suppression; tumour immunology, cancer and the immune system; Immunodeficiency diseases.

#### ReferenceBooks

- Abas, Lichtman and ShivPillai, 2021, Cellular and Molecular Immunology, 10<sup>th</sup> Edition, Elsevier
- Jeffrey K. Actor, 2011, Elsevier Integrated Review Immunology and Microbiology, 2<sup>nd</sup> Edition.
- 3. Joseph A. Bellanti, 2016, Immunology IV: Clinical Applications in Health and Disease, Georgetown University School of Medicine, Washington, DC.
- 4. Kuby, J., 1999, Immunology, W.H. Freeman and Company, New

York.
5. Murphy and Weaver, 2016, Janeway's Immunobiology 9th Edition,
W.W. Norton & Company.
6. Paul, 2012, Fundamental Immunology, 7 <sup>th</sup> Edition, Lippencott
Williams & Dilkins, Kluwer.
7. Punt, Stranford, Jones and Owen, 2019, Kuby Immunology, 8 <sup>th</sup>
Edition, W.H. Freeman and Company, New York.
8. Rich, Fleisher, Shearer, Schroeder, Frew and Weyand, 2018, Clinical
Immunology: Principles and Practice, 5 <sup>th</sup> Edition, Elsevier.
9. Roitt, 2000, Essential Immunology, Blackwell Scientific Publishers.
10. Williams, 2011, Immunology: Mucosal and Body Surface Defences,

At the end of the course the students can expect to learn the following

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Course Outcomes	Cognitive Level
CO1: Understand the fundamental concepts of immunity and Gain an in-depth knowledge of characteristics and functions of the organs and cells of the immune system. Assimilate knowledge on the characteristics that make the molecules to act as antigens.	K1, K2
CO2: Knowledge on the molecular components of the immune system like lymphokines, cytokines, immunoglobulins and understand the organisation of genes and antibody repertoire. understand the cell surface proteins essential to differentiate self and non-self molecules and the pathways for antigen processing and presentation.	K1, K2
CO3: Demonstrate an understanding of the overlapping roles of innate and adaptive immunity and able to compare and contrast	K1, K2, K4

the unique roles of humoral and cell-mediated immunity in response to pathogens. Gained knowledge on Prophylactic measures and different types of vaccines.	
CO4: The understanding of the antigen antibody reactions will enable the students to develop the skill to decide on the selection of specific assays according to the samples and need. Gain potential for research projects/ novel diagnostic kits development based on disease diagnosis and prophylaxis.	K3, K4, K5, K6
CO5: Comprehend the overreaction by our immune system; Apply the knowledge of molecular and cellular basis of immunity to understand the role of immunity in infections, transplantation, autoimmune disease, immune deficiency and cancer. Gain potential for development of research projects based on immunological changes and its advancements.	K2, K3, K6

K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create

**Mapping of CO with PO** 

	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	M	L	L	L	L	L	L	L	L	M	L	L	L	L	L
CO2	M	L	M	L	L	L	L	L	L	L	M	L	L	L	L
CO3	M	L	M	L	L	M	L	L	L	L	M	L	M	L	M
CO4	Н	Н	Н	Н	Н	Н	M	Н	M	L	M	M	M	Н	Н
CO5	Н	Н	Н	Н	M	Н	M	M	M	M	M	M	M	Н	Н

# $H-High,\,M-Medium,\,L\text{ - }Low$

**Mapping of CO with PSO** 

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6
CO1	L	L	L	L	L	L
CO2	L	M	L	M	L	L
CO3	L	L	L	L	L	L
CO4	Н	M	L	L	L	M
CO5	Н	Н	L	L	L	M

H - High, M - Medium, L - Low

Semester		II Semester							
<b>Course Type</b>		Core Paper - 8							
Title of the Co	urse	MICROBIAL GENETICS							
<b>Course Code</b>									
Teaching Hou	ırs	60 Hours/ Semester : 4 Hours/	week						
	MICI	ROBIAL GENETICS	its: 4	Max. Marks: 100 (Internal: 25, External: 75)					
Course Duomas	nicitec								
Course Prereq	uisites								
-	•	re a basic knowledge on genetic m	aterial o	f prokary	votes				
-	•		aterial o	f prokary L	votes T	P	C		

Module 1	Basics of Microbial Genetics	12 ho	urs
Course Objectives	<ul> <li>To provide an understanding of the genetic constitue special emphasis on various approaches</li> <li>To impart thorough knowledge on gene regular mechanisms.</li> <li>To understand the different modes of gene regulation and the importance of bacterial transposition and its ap</li> </ul>	nts of bacteration and in microor	transfer
	To provide a comprehensive detail on microbial genom	nes	

Gene, chromosome – prokaryotic & eukaryotic chromosomes, genome, organisation in prokaryotes and eukaryotes – DNA content law of DNA constancy & C value paradox – cot curve. Prokaryotic transposable elements – Insertion Sequences, composite and non-composite transposons, Replicative and Non replicative transposition, Mu transposon. Eukaryotic transposable elements - Yeast (Ty retrotransposon), Drosophila (P elements), Maize (Ac/Ds). Uses of transposons and transposition. Mutations and mutagenesis: Definition and types of Mutations; Physical and chemical mutagens; Molecular basis of mutations; Functional mutants (loss and gain of function mutants); Uses of mutations. Reversion and suppression: True revertants; Intra- and inter-genic suppression; Ames test; Mutator genes.

# Module 2 Bacterial Genetics 12 hours

Bacterial chromosomes, E. coli chromosome structure, circular genetic map, plasmids - structure, classification, copy control, incompatibility, F-factor, col and R plasmids, Genetic mapping of E. coli, genetics of quorum sensing in bacteria. Gene concept - regulation of bacterial gene expression. Lactose system - coordinate regulation, Lac components, positive and negative regulation, catabolite repression. Tryptophan operon - attenuation. Arabinose operon and its regulation.

#### Module 3 Gene Transfer & Recombination 12 hours

Gene Transfer Genetics & Mechanisms: Transformation - Discovery, mechanism of natural competence. Conjugation - Discovery, mechanism, Hfr and F' strains, Interrupted mating technique and time of entry mapping. Transduction - Generalized transduction, specialized transduction, LFT & HFT lysates, Mapping by recombination and co-transduction of markers; Recombination: types homologous or general, site specific and random recombination, general recombination between homologous DNA- Holliday model, double strand model of general recombination, enzymes involved in recombination, rec - proteins. Chromosomes & DNA replication in archaea, gene

transfer in archaea, deletion mapping, complementation, intragenic complementation, heteroduplex mapping, foot printing, chromosome walking and chromosome jumping.

## Module 4 DNA Damage & Repair

12 hours

DNA damages - Types, hit theory, UV radiation. DNA repair: post irradiation effects on survival levels - photo reactivation, liquid holding recovery. Biochemistry repair mechanism: excision, recombination and SOS repair.

## Module 5 Fungal, Bacterial & Archael Genetics

12 hours

Yeast as a model organism for eukaryotic genetic research – chromatin structure, genetic recombination and gene regulation. Meiotic & mitotic mapping, gene conversion, heterothallism & mating type, tetrad analysis & linkage detection in Neurospora. Structure, genome organisation and replication of bacteriophages – T7, T4 – rII locus,  $\infty X174$  & M13 phages, genetics of Agrobacterium.

- 1. Das, S, and Dash, H.R., 2018, Microbial Diversity in the Genomic Era, 1<sup>s</sup> Edition, Academic Press.
- Freifelder, D., 2008, Microbial Genetics, 18<sup>th</sup> Edition, Narosa Publishing House, New Delhi.
- 3. Gardner, Simmons and Snustad, 2010, Principles of Genetics, 8<sup>th</sup> Edition, Wiley India Pvt. Ltd.
- 4. Jeyanthi, G.P., 2009, Molecular Biology, MJP Publishers, Chennai.
- Johnson, 1994, Molecular Genetics of Yeast A Practical Approach, Oxford University Press.
- Kalia, V.C., Shouche, Y., Purohit, H.J. and Rahi, P., 2017, Mining of Microbial Wealth and MetaGenomics 1<sup>st</sup> Edition, Springer Nature Singapore Pvt. Ltd.
- 7. Klug, Cummings and Spencer, 2016, Concepts of Genetics, 10<sup>th</sup> Edition, Pearson Education, India.
- 8. Krebs, J.E., Elliott, S and Goldstein, 2017, Lewin's GENES XII, 12<sup>th</sup> Edition, Jones and Bartlett Publishers.
- 9. Lewin B., 2017, Gene XII, Oxford University Press Oxford.
- 10. Moore and Frazer, 2002, Essential Fungal Genetics, Springer.
- Primrose and Twyman, 2003, Principles of Genome Analysis and Genomics, Blackwell Publishing Company.

#### ReferenceBooks

- 12. Snyder, L, Peters, Henkin and Champness, 2013, Molecular Genetics of Bacteria 4<sup>th</sup> Edition, ASM Press.
- 13. Stanley R. Maloy, John E.C. and Freifelder, D., 2008, Microbial Genetics, Narosa Publishing House, New Delhi.
- Streips and Yasbin, 2002, Modern Microbial Genetics, 2<sup>nd</sup> Edition, Wiley Publications.

At the end of the course the students can expect to learn the following

Course Outcomes	Cognitive
<b>CO1</b> : Understanding the structure and functions of genomes of different microbial groups and microbial genetics.	K1, K2, K4
CO2: Understanding the processes behind mutations and other genetic changes.	K1, K2, K3, K4
CO3: Identifying and distinguishing genetic regulatory mechanisms at different levels.	K2, K3, K4, K5
<b>CO4:</b> Able to understand the different recombination mechanisms in microorganisms, and the basic genetics behind the DNA damage and repair mechanisms.	K2, K3, K4
CO5: Able to plan basic experiments in microbial genetics concerned with clarifying phenotypes and their relationship with the genotype using common methods in microbial genetics	K2, K3, K4, K5

# K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create Mapping of CO with PO

	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	Н	M	M	L	M	L	M	L	M	M	M	M	L	L	Н

CO2	Н	M	M	L	M	L	M	L	M	M	M	M	L	L	M
CO3	Н	M	M	L	M	L	M	L	M	M	M	M	L	L	Н
CO4	Н	M	M	M	M	M	M	L	M	M	M	M	L	L	M
CO5	Н	M	M	M	M	M	M	L	M	M	M	M	L	L	Н

H – High, M – Medium, L - Low

**Mapping of CO with PSO** 

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6
CO1	Н	M	M	L	Н	L
CO2	Н	M	M	L	Н	L
CO3	Н	M	M	L	Н	L
CO4	Н	M	M	L	Н	M
CO5	Н	M	M	L	Н	M

H - High, M – Medium, L – Low

Semester		II Semester						
<b>Course Type</b>		Core Compulsory Course Pra	ectical - 3					
Title of the Co	urse	BACTERIOLOGY & VIROLOGY AND MYCOLOGY &						
		PHYCOLOGY						
<b>Course Code</b>								
<b>Teaching Hour</b>	rs	60 hours/semester: 4 hours/week						
	BACT	TERIOLOGY & VIROLOGY	Credits: 2	Max. Marks: 100				
	AND	MYCOLOGY &		(Internal: 50;				

	PHYCOLOGY		E	xternal:	50)
Course Prere	quisites:				
The student sl	hould have a basic practical knowledge on micro	biology a	nd espec	ially on	isolation
of microorgan	isms				
CODE:	BACTERIOLOGY & VIROLOGY AND	L	T	P	С
0022	MYCOLOGY & PHYCOLOGY	-	-	4	2
Course Objectives	Understand the architecture of viruses, Know the methods used for isolating, cultivate To understand the isolation, identification, con To know about the techniques for culture and	unting and	harvest	of micro	C
1 Table!	n of hostoriophages from savage water	iuciiiiica	11011 01 11	angai spo	nes

- 1. Isolation of bacteriophages from sewage water
- 2. Cultivation of viruses in embryonated Eggs: different routes of inoculation.
- 3. Virus inclusion bodies (slides)
- 4. Determination of stability of plant virus in cell sap- TIP, DEP, LIV.
- 5. Determination of chlorophylls in healthy and virus diseased leaves.
- 6. Purification of viruses by different chemical and physical methods
- 7. Isolation and identification of microalgae from seawater.
- 8. Harvest of microalgal cells by different methods.
- 9. Enumeration of algal cells by counting chamber method
- 10. Isolation of marine fungi from marine environment.
- 11. Mushroom cultivation technique.
- 12. Identification of fungal spore by staining.
- 13. Identification of fungal diseases in selected plants and animals (Etiology).

After the completion of the course, the student will be able to

Course Outcomes	Cognitive Level
CO1: Acquire hands on experiences on isolation methods, identification, different methods of culture, media preparation, enumeration, and harvesting of microalgae	

CO2: Acquire hands on experiences on isolation methods,	K1, K2, K3, K4, K5
identification, of fungal disease diagnosis	
<b>CO3:</b> Understanding skill on different methods of isolation and	K1, K2, K3
cultivation of viruses	
CO4: Understanding on virus inclusion bodies	K1, K2
<b>CO5:</b> Understanding and knowledge on purification of viruses by	K1, K2, K3
chemical and physical methods	

# K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create

**Mapping of CO with PO** 

	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	Н	M	Н	M	M	M	M	M	M	M	Н	M	Н	M	L
CO2	Н	M	Н	M	M	M	M	M	M	M	Н	M	Н	M	L
CO3	Н	M	Н	M	M	M	M	M	M	M	Н	M	Н	M	L
CO4	Н	M	Н	M	M	M	M	M	M	M	Н	M	Н	M	L
CO5	Н	M	Н	M	M	M	M	M	M	M	Н	M	Н	M	L

## H – High, M – Medium, L – Low Mapping of CO with PSO

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6
CO1	Н	Н	M	L	L	L
CO2	Н	Н	M	L	L	L
CO3	Н	Н	M	L	L	L
CO4	Н	Н	M	L	L	L

CO5	Н	Н	M	L	L	L
~ ~ ~				_	_	

H – High, M – Medium, L - Low

Semester		II Semester										
<b>Course Type</b>		Core Compulsory Course Practical - 4										
Title of the Co	ourse	IMMUNOLOGY AND MICE	ROBIAL	GENET	ICS							
<b>Course Code</b>												
Teaching Hou	ırs	0 hours/semester: 4 hours/week										
		UNOLOGY AND	Cred	its: 2	Max	x. Mark	s: 100					
	MICE	ROBIAL GENETICS			-	nternal:	,					
					E	xternal:	50)					
Course Prered	_											
		ve a basic practical knowledge o	n microb	oiology, e	speciall	y the iso	lation of					
microorganism	is and in	nmunological techniques		Π	_	Π	I					
CODE:	IN	MMUNOLOGY AND MICRO	BIAL	${f L}$	T	P	C					
		GENETICS		-	-	4	2					
Course Objectives	•	To get an idea about blood gro To get an idea about the antige To inculcate/impart skills to pe To provide an complete idea al To provide students with a especially transformation in many	n-antiboderform vaccout the real deep in the content of the conten	rious test nutation s insight g	s/assays studies i	in micro	organisms					
1.		Identification of lymphoid orga	ans and c	ells.								
2.		Preparation of serum, plasma a	_									
3.		Direct agglutination to determi	ne ABO	blood gro	ouping.							
4.		Determination of differential le	eukocyte	count.								
5.	5. Isolation and enumeration of RBC from human blood.											
6.	Antigen- antibody reaction – Precipitation - ODD/SRID/CID											
7.												
agglutin	ation											

8. Cell viability/cytotoxicity assay

9. Enzyme Linked Immune Sorbant Assay – Demonstration

10. Isolation of nucleic acid and characterization by gel Electrophoresis

11. Inactivation of microorganisms by different mutagens.

12. Production, isolation and characterization of mutants.

13. Determination of mutation rate – natural and induced

14. Isolation, characterization and curing of plasmids.

15. Preparation of competent cells and transformation of E.coli using plasmid DNA

After the completion of the course, the student will be able to

Course Outcomes	Cognitive Level
CO1: Understand the immune system of mammal and fish	K2
CO2: Learn the working principle of Antigen antibody reactions and apply it for disease diagnosis (understand/apply)	K2
CO3: Learn experiments of hematology and identify the defects of immune cells	K3, K5
CO4: Able to develop projects related to development of immunostimulants and its application	K5
CO5: Understand practically how gene transfer occurs in microorganisms, and able to understand the mutation studies in microorganisms	K2, K4, K5

# K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create Mapping of CO with PO

| PO1 | PO2 | PO3 | PO4 | PO5 | PO6 | PO7 | PO8 | PO9 | PO10 | PO11 | PO12 | PO13 | PO14 | PO15 |

CO1	Н	M	Н	L	L	L	Н	L	M	M	Н	M	L	L	L
CO2	Н	M	Н	L	L	L	Н	L	M	M	Н	M	L	L	L
CO3	Н	M	Н	L	L	L	Н	L	M	M	Н	M	L	L	L
CO4	Н	M	M	L	L	L	M	L	M	M	Н	M	L	L	L
CO5	Н	M	M	L	L	L	M	L	L	M	Н	M	L	L	L

H – High, M – Medium, L - Low Mapping of CO with PSO

		PSO2	PSO3	PSO4	PSO5	PSO6
CO1	Н	Н	L	L	M	L
CO2	Н	Н	L	L	M	L
CO3	Н	Н	L	L	M	L
CO4	Н	M	L	L	Н	L
CO5	Н	M	L	L	Н	L

H - High, M – Medium, L – Low

Semester		II Semester						
<b>Course Type</b>		Core Elective Course – 2A						
Title of the Co	urse	FOOD MICROBIOLOGY	FOOD MICROBIOLOGY					
<b>Course Code</b>								
<b>Teaching Hou</b>	rs	45 Hours/ Semester: 3 Hours	/ week					
	FOOI	O MICROBIOLOGY	Credits: 3	Max. Marks: 100				
				(Internal: 25, External				

C				<b>75</b> )	
related to food	isites: The students should have a basic know	ledge on f	cood and	d microo	rganisms
CODE:	EOOD MICROPIOLOGY	L	T	P	C
	FOOD MICROBIOLOGY	3	-	-	3
Course Objectives					
Module 1	Introduction to Food Microbiology			9 ho	ours
micro-organisms of characteristics — cl	orne bacteria, airborne fungi, micro-organisms of plants, micro-organisms of animal origin, n				
influencing micro	lassification and importance. Principles of food (anaerobic conditions, high temperature, low bial growth in food: extrinsic and intrinsic faming, processing for heat treatment-D and F value.)	w temper factors- cl	ature &	drying	removal of g), factors
influencing micro	lassification and importance. Principles of food (anaerobic conditions, high temperature, low bial growth in food: extrinsic and intrinsic for	w temper factors- cl	ature &	drying	emoval of g), factors atives and
influencing micro food additives- can  Module 2  Contamination, p vegetables & frui	lassification and importance. Principles of food anaerobic conditions, high temperature, low bial growth in food: extrinsic and intrinsic finning, processing for heat treatment-D and F value Contamination and Spoilage  Treservation and spoilage of cereals & its its and their products, meat and its product eir products; poultry and its products—spoil	factors- chalues  products, s, milk a	sugar	greserva 9 ho and its products,	emoval of g), factors atives and purs products, fish and
influencing micro food additives- can  Module 2  Contamination, p vegetables & frui seafood's and the spoilage and chara	lassification and importance. Principles of food anaerobic conditions, high temperature, low bial growth in food: extrinsic and intrinsic finning, processing for heat treatment-D and F value Contamination and Spoilage  Treservation and spoilage of cereals & its its and their products, meat and its product eir products; poultry and its products—spoil	factors- chalues  products, s, milk a	sugar	greserva 9 ho and its products,	emoval of g), factors atives and purs products, fish and tection of
influencing micro food additives- can Module 2  Contamination, p vegetables & frui seafood's and the spoilage and chara Module 3  Bacterial and not Clostridium, E.cot algae, fungi and	lassification and importance. Principles of food anaerobic conditions, high temperature, low bial growth in food: extrinsic and intrinsic finning, processing for heat treatment-D and F value and their products, meat and its product eir products; poultry and its products—spoil acterization  Food Borne Infection and Intoxications  n-bacterial with examples of infective and li, Salmonella, Shigella, Staphylococcus, Vibra viruses. Food borne outbreaks—laborate	products, s, milk a age of ca toxic typrio, Yersin ory testin	sugar nd its j anned f	9 ho and its products, coods-Des	products, fish and tection of tection of tection of tection, protozoa, prevention
influencing micro food additives- car  Module 2  Contamination, p vegetables & frui seafood's and the spoilage and chara Module 3  Bacterial and not Clostridium, E.co. algae, fungi and measures – food of	lassification and importance. Principles of food anaerobic conditions, high temperature, low bial growth in food: extrinsic and intrinsic finning, processing for heat treatment-D and F value and their products, meat and its product eir products; poultry and its products—spoil acterization  Food Borne Infection and Intoxications In-bacterial with examples of infective and li, Salmonella, Shigella, Staphylococcus, Vibra	products, s, milk a age of ca toxic typrio, Yersin ory testin	sugar nd its j anned f	9 ho and its products, coods-Des	products, fish and tection of burs  Bacillus, Protozoa, prevention

Cheese, yoghurt, butter milk, sour cream Fermented vegetables; sauerkraut, pickles, olives and soy sauce. Fermented meat, fermented Indian foods – leavening of bread. Food spoilage: Spoilage of fruit and vegetables, spoilage of cereal and cereal products – cereal grains, and bread, spoilage of meat and meat products – Bacon and Ham, spoilage of milk and milk products – butter and frozen desserts. Food borne diseases – indicators of pathogens & food poisoning.

# Module 5 Food Produced by Microbes Fermented foods microbial cells as food (Single cell protein)

Fermented foods, microbial cells as food (Single cell protein), mushroom cultivation. Bioconversions – production of alcohol – fermented beverages (beer and wine), industrial enzymes production (amylases, proteases, and cellulases); amino acid production (glutamic acid and lysine).

1. Adams, M.R. and Moss, M.O., 2008, Food Microbiology, RSC Publishing, Cambridge, UK.

9 hours

- 2. Benwart, G.J., 1987, Basic Food Microbiology, CBS Publishers & Distributors, New Delhi.
- 3. Blackburn C. de W., 2006, Food Spoilage Microorganisms, Woodhead Publishing, Cambridge, UK
- 4. Deak, T. and Beuchat, L.R., 1996, Hand Book of Food Spoilage Yeasts, CRC Press, New York.
- 5. Frazier, W.C., and Westhoff, D.C., 1988, Food Microbiology (Reprint 1995), Tata McGraw Hill Publishing Ltd., New Delhi.
- 6. Garbutt, J., 1997, Essentials of Food Microbiology, Arnold International Students Edition, London.
- 7. Jay J.M., 2000, Modern Food Microbiology, 6th Edition, Chapman & Hall, New York.
- 8. Prescott, L.M., Harley, J.P. and Helin, D.A., 2008, Microbiology, 5th Edition, McGraw Hill, New York.
- 9. Ray, B., 2000, Fundamental Food Microbiology, 2nd Edition. CRCPress. New York.
- 10. Robinson R.K. (ed.), 2002, Dairy Microbiology Handbook, 3rd Edition, Wiley Interscience.

After the completion of the course, the student will be able to

Co	ourse Outcomes	Cognitive Level

### ReferenceBooks

CO1: Gain knowledge about food as a substrate for various	K1, K2, K6
microbes, the role of factors and its importance, understand about	
the principles and application of different types of food	
preservation technique, understand the basis of food safety	
regulations and use of standard methods and procedures for the	
microbiological analysis of food; Acquire, discover, and apply the	
theories and principles of food microbiology in practical, real-	
world situations and problems.	
CO2: Understanding the knowledge on various food	K1, K2, K3
<b>CO2:</b> Understanding the knowledge on various food contamination and spoilage	K1, K2, K3
Contamination and sponage	
CO3: Acquire a thorough understanding of food borne diseases,	K1, K2, K3
testing methods, and preventive technique.	
COA. I same shout the various formanted maduate and its various	V1 V2 V2
<b>CO4:</b> Learn about the various fermented products and its various	K1, K2, K3
stage spoilage.	
CO5: Understand the basic knowledge about the fermentation	K1, K2
process and the requirements, designing of fermentation process.	
Acquire the knowledge about the production of antibiotic and	
enzymes	

K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create Mapping of CO with PO

	PO1	PO2	PPO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	Н	M	Н	M	M	L	M	M	M	M	M	M	Н	Н	M
CO2	Н	Н	Н	M	M	M	M	Н	M	M	M	L	L	Н	M

CO3	Н	Н	M	M	L	L	M	M	M	Н	M	M	L	Н	M
CO4	Н	Н	Н	M	M	L	M	M	M	M	M	M	L	Н	M
CO5	Н	Н	Н	M	M	Н	M	M	M	M	M	M	M	L	M

H – High, M – Medium, L - Low Mapping of CO with PSO

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6
CO1	M	Н	M	M	M	M
CO2	Н	M	M	M	M	M
CO3	Н	M	L	M	L	L
CO4	Н	Н	M	M	Н	L
CO5	Н	M	M	M	Н	M

H - High, M - Medium, L - Low

Semester		II Semester						
<b>Course Type</b>		Core Elective Course – 2B						
Title of the Co	urse	MARINE MICROBIAL TECHNOLOGY						
<b>Course Code</b>								
Teaching Hou	ırs	45 Hours/ Semester : 3 Hours/ week						
	MAR	INE MICROBIAL	Credits: 4	Max. Marks: 100				
	TECH	INOLOGY		(Internal: 25, External				
				75)				

Course Prerequisites: The students should have a basic knowledge on marine microbiology and techniques used in microbiology

CODE:		L	T	P	C						
	MARINE MICROBIAL TECHNOLOGY	3	-	-	3						
Course Objectives	<ul> <li>To get an idea about the vaccination on disease control, role of microbe in industry</li> <li>Acquire knowledge on microbial probiotics</li> <li>To get an idea about the microalgal and halobacterial technology</li> </ul>										
Module 1	Gene Technology for Disease Control			9 ho	urs						
proteins in aquati shrimp by gene t	nt type of vaccines in microbial control; anti- ic important pathogens. Transgenic fish: de ransfer technology. RNA interference: princi- c in microbial control.	velopme	nt of tr	ansgenic	fish and						
Module 2	Marine Microbes for Industry			12 h	ours						
– A general revie	ation and maintenance of industrially important in the word primary and secondary metabolites production of alcohol, organic solvents, anti- es.	duction -	- SCP I	Productio	n – basic						
Module 3	Microalgal Production Technology			12 h	ours						
Culture system-p	hotobioreacter, harvesting of biomass, Bio-	fuel pro	oduction	ı- fats,	oils, and						
hydrocarbons: ca Genetic engineerii	rotenoid production, Microalgae for aquacult ng of microalgae.	ure and	waste	water ti	eatments,						
Module 4	Probiotics			12 h	ours						
Definition-characteristics of probiotics: Probiotic microbes-production and characterization of probotics-mode of action of probiotics in fish nutrution larval rearing. Immune modulation and disease management											
Module 5	Halobacterial Technology			12 h	ours						
	cation – Ecology – structure and functions of co										
	potentials of Halobacterium - important pr	oducts a	and their	ir uses -	- role of						
halobacteria in sal	t purification.										

	1. Chandra Bhusan Singh, 2011,
	Introduction to Microbiology and Biotechnology, Neha Publishers and
	Distributors.
	2. Jonatha M. Gott, 2004, RNA
	Interface, Editing and Modification: Methods and Protocol (Methods in
	Molecular Biology), Humana Publisher.
ReferenceBooks	3. Michael A. Borowitzka and Laglay J. Borowitzka (Eds.), Micro algal
	Biotechnology, University of Cambridge Press, New York.
	4. Patel, A.H., 2015, Text book of
	Industrial Microbiology, 2 <sup>nd</sup> Edition, McMillan, India.
	5. Peter Marian, M., John, J.A.C., Immanuel G., and Michael Babu, M.,
	2002, A Text Book of Marine Natural Products, M.S. University,
	Tirunelveli.

After the completion of the course, the student will be able to

Course Outcomes	Cognitive Level
<b>CO1:</b> Vaccination, techniques used in rDNA technology for the development of transgenic fish	K1, K2, K3
CO2: Understand the role of microbes in industrial production of products such as SCP, alcohol, organic acids, antibiotics etc	K1, K2, K3
CO3: Know about the role of microalgae on biofuel production, aquaculture and waste water treatment	K2, K3, K4
CO4: Role of probiotic microbes in fish nutrition, characterization, mode of action	K2, K3, K4
CO5: Role of halobacteria on salt purification, their structure, potential	K2, K3, K4

## K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create

## **Mapping of CO with PO**

	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	Н	M	M	Н	M	M	M	L	M	M	M	M	M	L	L
CO2	Н	M	M	M	M	M	M	L	M	M	M	Н	M	L	L
CO3	Н	M	M	M	M	M	M	L	M	M	M	Н	M	L	L
CO4	Н	M	M	M	M	M	M	L	Н	M	M	Н	M	L	L
CO5	Н	M	M	M	M	M	M	L	M	M	M	Н	M	L	L

H – High, M – Medium, L - Low

**Mapping of CO with PSO** 

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6
CO1	Н	Н	Н	L	Н	L
CO2	Н	Н	Н	L	Н	L
CO3	Н	Н	Н	L	M	L
CO4	Н	Н	Н	L	L	L
CO5	Н	Н	Н	L	L	L

H - High, M - Medium, L - Low

Semester		III Semester						
Course Type Core Compulsory Paper - 9 Title of the Course RECOMBINANT DNA TECHNOLOGY								
Title of the Co								
<b>Course Code</b>								
Teaching Hou	rs	60 Hours/ Semester : 4 Hours	/ week					
	RECOMBINANT DNA TECHNOLOGY			its: 4		fax. Marks: 100 rnal: 25, External 75)		
Course Prereq biotechnologica		The student should have a basi	c knowle	edge abo	ut the b	iotechno	logy and	
CODE:	D. D. C.		T 0 0 T 7	L	T	P	C	
	REC	COMBINANT DNA TECHNO	LOGY	4	-	-	4	
Course Objectives	•	To understand the principle be recombinant DNA technology To gain knowledge on gene methods and its importanc biotechnology. To teach students about intelle understand the process of pater To impart knowledge of safety for transgenic research	e manipu e in p ctual pro nt filing	lation ulant, and	sing ge imal ar	enetic end nd envir	ngineering ronmental nake them	
Module 1	Basic	s of rDNA Research				12 h	ours	
Cloning vectors vector –YAC v	– plasm ectors, m – La	engineering; DNA modifying enids, cosmids, phasmids, phagement viral vectors — SV 40 and ad mbda, PL / PR Promoter, T7 pro	nids, exp eno viru	ression v s; Prom	ectors, otors: L	integrati ac Z pr	ng shuttle omoter –	
Module 2 Cloning Methodologies 12 hours						ours		

Steps in cloning: sticky and blunt end cloning. Gene Transfer methods; Cloning from mRNA – synthesis of cDNA, cloning cDNA– cDNA library. Cloning from genomic DNA – genomic library. Metagenomic library; Shot gun cloning. Screening of recombinants – phenotypic expression of characters--α complementation, – Blotting techniques – Western, Northern and Southern. Mapping of human genes – Human genome project.

### Module 3 Techniques

12 hours

PCR – gene amplification, primer designing, optimization, variation in the PCR (RAPD, RFLP, RACE, RT-PCR). DNA sequencing – Sanger's method, Maxam Gilbert's method, automated sequencing and micro array, Next generation sequencing; Expression system: Prokaryotic and eukaryotic expression systems and their application in E. coli, Streptomyces, Yeast, Baculovirus and animal hosts.

#### Module 4 Application

12 hours

Cloning of human insulin, interferon in E.coli. Recombinant vaccine development – HBs Ag in yeast. Cloning for commercial production of antibiotics (Penicillin), bio steroid transformation, production of biopolymers – Xanthumgum, melanin biosynthesis in E.coli, adhesive biopolymer in yeast; Gene Therapy; Plant genetic engineering: Ti plasmid, CaMV vector, direct DNA delivery methods – micro projectile bombardment, microinjection and electroporation.

#### Module 5 Advances in Transgenic Technology and IPR

12 hours

Gene silencing; Types and mechanism of gene silencing, genetic factors of silencing, formation of antisense mRNA, inhibition of gene expression by antisense RNA, gene silencing in crop plants: tomato, Si RNA and disease control. Transgenic and GM organisms- guidelines for rDNA and transgenic research; Safety issues and IPR.

## ReferenceBooks

- 1. Brown, T.A., 2021, Gene Cloning and DNA Analysis: An Introduction, 8<sup>th</sup> Edition, Wiley- Blackwell Publishing, U.K.
- 2. Glick, B.R. and Patten, C.L., 2017, Molecular Biotechnology: Principles and Applications of Recombinant DNA, 5<sup>th</sup> Edition, ASM Press, USA.
- 3. Glick, B.R., Pasternak, J.J. and Patten, C.L., 2009, Molecular Biotechnology, 4<sup>th</sup> Edition, ASM Press, USA.
- 4. Green, M. and J. Sambrook, J., 2012, Molecular Cloning: A Laboratory Manual, 4<sup>th</sup> Edition, Cold Spring Harbour Laboratory Press, USA.
- 5. Nicholl, D.S.T., 2008, An Introduction to Genetic Engineering, 3<sup>rd</sup> Edition, Cambridge University Press.
- 6. Primrose, S.B. and Twyman, R.M., 2016, Principles of Gene Manipulation

- 7. Willey, J. M., Sandman, K. and Wood, D. 2019, Prescott's Microbiology, 11<sup>th</sup> Edition, McGraw Hill Higher Education, USA.
- 8. Winnacker, E.L., 1986, From Genes to Clones, Reprinted by Panima Publishing Corporation, New Delhi.

After the completion of the course, the student will be able to

Course Outcomes	Cognitive Level
CO1: Acquire in-depth understanding of the exploitation of restriction and DNA-modifying enzymes, in recombinant DNA technology, along with the use of linkers and adapters. Gained detailed knowledge of the use of different cloning vectors and different types of expression vectors. Expertise in selection and evaluation of proper tools for rDNA research	K1, K2, K5, K6
CO2: Understand the various steps involved in construction of chimeric molecules and construction of genomic, metagenomic and cDNA libraries, and whole genome sequencing. Gained expertise in development of projects in engineering of genes	K1, K2, K5, K6
CO3: Learn and apply various techniques like PCR and Next generation sequencing technologies for development of research projects. Acquire knowledge on different types of expression vectors used to express heterologous proteins in bacteria, yeast, insect cells and mammalian cells. Evaluate the use of different expression systems.	K1, K2, K4, K6
CO4: Apply the acquired knowledge of genetic engineering for	K3, K4, K5, K6

development of products of human therapeutic interest. Understand gene therapy and plant Genetic engineering to create novel GE crops.	
CO5: Understand and apply transgenic techniques for research and development. Appreciate the potential for safety associated with rDNA research and evaluate and ensure the preventive measures/guidelines to be undertaken. Become aware of the ethics involved in biotechnology research.	K2, K3, K5, K6

## K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create

# **Mapping of CO with PO**

	PO1	PO2	PPO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	Н	L	L	L	L	M	L	L	L	M	L	L	L	L	L
CO2	Н	L	M	L	L	M	L	Н	L	L	M	L	L	L	L
CO3	Н	Н	M	L	L	M	Н	Н	L	M	M	Н	M	L	M
CO4	Н	M	M	Н	Н	Н	Н	Н	M	L	M	M	M	Н	Н
CO5	Н	M	Н	Н	M	Н	Н	M	M	M	M	Н	Н	Н	Н

# H-High, M-Medium, L-Low

**Mapping of CO with PSO** 

Mapp	apping of CO with 100												
	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6							
CO1	Н	M	M	Н	Н	L							

CO2	Н	M	M	Н	Н	L
CO3	Н	M	M	Н	Н	Н
CO4	Н	M	M	Н	Н	Н
CO5	Н	Н	M	Н	Н	Н

H - High, M - Medium, L - Low

Semester		III Semester					
<b>Course Type</b>		Core Compulsory Paper - 10					
Title of the Co	urse						
<b>Course Code</b>		BIOPROCESS TECHNOLOGY					
Teaching Hou	rs	60 Hours/ Semester : 4 Hours/ week					
BIOP		ROCESS TECHNOLOGY	Credits: 4	Max. Marks: 100			
				(Internal: 25; External			
				75)			

**Course Prerequisites:** The student should acquire knowledge on general aspects about fermentation technology, which includes microbes responsible for production of bioactive metabolites in confined system. The produced metabolites will be processed by employing various techniques and the industrial and pharmaceutical applications of the identified metabolites will be studied.

CODE:		${f L}$	T	P	C
	BIOPROCESS TECHNOLOGY	4	-	-	4
Course Objectives	<ul> <li>To understand the basic concepts fermentation and its optimized condition parts &amp; functions</li> <li>To know about microbes responsible isolation, preservation and maintenance</li> </ul>	ons, types for produ	s of bio	reactors of metabo	and their olites, its

media required including different nutrients necessary for their growth Enable to understand the methods of downstream processing of metabolites. Immobilization techniques involved in preservation of metabolites To understand the elementary idea on production and preservation of industrially important food products. Module 1 General Account on Fermentation 12 hours Types of fermentation – surface, submerged solid state, adhesive, batch, continuous and fed batch principle of chemostat and thermostat, Bioreactors – types, parts and their functions – optimization conditions, aeration, agitation, foam control process control equipment's. Isolation, Preservation & Maintenance of Industrially Module 2 12 hours **Important Microbes** Isolation, preservation and maintenance of industrial important microorganisms-Microbial growth kinetics and microbial death kinetics- media for industrial fermentation: Source of nutrients, types of media: synthetic and crude media for industrial fermentation: Source of nutrients, types of media; synthetic and crude media; Advantage and disadvantages. **Downstream Processing** 12 hours Module 3 Introduction, removal of microbial cells and solid matter, foam separation, precipitation, filtration, centrifugation, cell disruption, liquid extraction, chromatography, membrane process. Drying and crystallization, effluent treatment -BOD, COD and disposal of effluents. 12 hours Module 4 **Immobilization Technique** Whole cell immobilization and their industrial applications, production of chemicals: alcohol (ethanol); acids (citric, acetic and gluconic acids), solvents (glycerol, acetone and butanol), antibiotics (penicillin, streptomycin and tetracycline); amino acids (lysine and glutamic acid); Single Cell Protein. Module 5 **Introduction to Food Technology** 12 hours Elementary idea of canning and packing – sterilization and pasteurization of food products technology of typical food and food products (bread, cheese & idli) – basic food preservation techniques. 1. Alba, S., Humphrey, A.E. and Millis N.F., 1985, Biochemical ReferenceBooks

Engineering, Univ. of Tokyo press, Tokyo.

2.	Anton Moser.	1988, Bioprocess	Technology.	Springer-Verl	ag, Austria.
∠.	rinton moser,	1700, Diopiocess	1 cermonogy,	opinizer ven	us, musum.

- 3. Baily, J.E. and Ollis, D.F., 2008, Biochemical Engineering Fundamentals, Mc Graw Hill book Co., New York.
- 4. Berenjian, A., 2019, Essentials in Fermentation Technology, Springer.
- 5. Kartan, P., 2017, Advances in Bioprocess Technology, Delve Publications.
- 6. Pogaku Ravindra, 2015, Advances in Bioprocess Technology, Springer-Verlag.
- 7. Rao, D.G., 2010, Introduction to Biochemical Engineering, 2<sup>nd</sup> Edition, Tata McGraw Hill education ltd., New Delhi.
- 8. Svenska, P., 2000, Bioprocess Technology, Fundamentals and Applications, Royal Institute of Technology, Stockholm.

Course Outcomes	Cognitive Level
CO1: Understand about the basic concepts of fermentation technology and its types, bioreactors and their types and conditions responsible for fermentation.	K1, K2
<b>CO2:</b> Know about how to isolate industrially important microbes, their preservation, nutrition and media and their types required for the growth of microbes.	K2, K3
<b>CO3:</b> Find out the suitable downstream processing technique to process the metabolites obtained after fermentation for industrial usage.	K3, K4
<b>CO4:</b> Determine the industrial and pharmaceutical applications of produced products	K5, K6

CO5: Make elementary idea on preservation of produced products in large scale level	K4, K5, K6

# K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create

# Mapping of CO with PO

	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	Н	M	L	L	L	M	M	L	L	M	M	M	M	L	L
CO2	Н	M	L	L	L	M	M	L	L	M	M	M	M	L	L
CO3	Н	M	L	L	L	M	M	L	L	M	M	M	M	L	L
CO4	Н	M	L	L	L	M	M	L	L	M	M	M	M	L	L
CO5	Н	M	L	L	L	M	M	L	L	M	M	M	M	L	L

# H – High, M – Medium, L - Low Mapping of CO with PSO

Mapping of CO with 150										
	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6				
CO1	M	L	L	Н	L	L				
CO2	M	L	L	Н	L	L				
CO3	M	L	L	Н	L	L				
CO4	M	L	L	Н	L	L				
CO5	M	L	L	Н	L	L				

H - High, M - Medium, L - Low

Semester		III Se	emester								
Course Type		Core Compulsory Paper - 11									
Title of the Co	urse	se MEDICAL MICROBIOLOGY									
<b>Course Code</b>											
Teaching Hou	rs	60 H	ours/ Sen	nester :	4 Hours	/ week					
	MED	CAL I	MICROE	BIOLOG	GY	Cred	its: 4	Max	x. Mark	s: 100	
								(Intern	nal: 25, 1	External	
								75)			
Course Prereq	uisites:	The st	udents sh	ould hav	ve a basio	c knowle	dge on m	icrobiol	ogy		
							Γ		1	1	
CODE:		MED	TOAT NA	ICDOD		<b>T</b> 7	${f L}$	T	P	C	
		MEDICAL MICROBIOLOG			Y	4	-	-	4		
	It co	vers al	ll biology	y of ba	acteria, v	viruses a	nd other	patho	gens rel	ated with	
	infect	ious di	iseases in	human	s. The c	ourse wil	ll provide	e the co	nceptual	basis for	
	under	understanding pathogenic microorganisms and particularly address the									
Course	funda	fundamental mechanisms of their pathogenicity. It will also provide opportunities									
Objectives	for a student to develop diagnostic skills in microbiology, Gain knowledge of										
Objectives		morphology, cultural characteristics, epidemiology, laboratory diagnosis of									
		emerging and reemerging infectious diseases and the role of the Biosafety									
	profe	ssional	s in Biom	edical R	Research	Laborato	ries				
Module 1		rical obiolog	<b>Events</b>	and	Develo	pment	in Mo	edical	12 h	ours	
Medically impor	tant mi	crobes	<ul> <li>bacteria</li> </ul>	a, fungi,	, algae, v	rirus and	parasites	. Labor	atory ma	nagement	

Medically important microbes – bacteria, fungi, algae, virus and parasites. Laboratory management – safety in containment laboratory, collection and transport of clinical samples, microbiological examination of urine, blood, faeces, cerebrospinal fluid, throat swabs, sputum, pus and wound exudates, normal flora of human systems – skin, respiratory tract, gastrointestinal tract and genitourinary tract, nosocomial infections, common types of hospital infections and their diagnosis and control.

Module 2	Pathogenic Bacteria	12 hours
Module 2	Pathogenic Bacteria	12 hours

Establishment, spreading, tissue damage and anti phagocytic factors; mechanism of bacterial adhesion, colonization and invasion of mucus membranes of respiratory, enteric and urogenital tracts. Role of aggressions, depolymerising enzymes, organotrophism, variation and virulence. Classification of pathogenic bacteria: *Staphylococcus*, *Streptococcus*, *Pneumococcus*, *Neisseria*, *Corynebacterium*, *Bacillus*, *Clostridium*, non-sporing anaerobes, organisms belonging to Enterobacteriaceae. Vibrios, Non-fermenting Gram negative Bacilli, *Yersinia*; *Haemophilus*; *Bordetella*, *Brucella*; *Mycobacterium*, *Spirochaetes*, *Actinomycetes*; *Rickettsiae*, *Chlamdiae*. Sexually transmitted diseases – Syphilis.

Module 3 Yeast 12 hours

General characteristics, pathogenesis and laboratory diagnosis and control measures of: Yeast–Cryptococcus neoformans. Yeast like fungus – *Candida* spp. Filamentous fungi – *Aspergillus* and *Penicillium*. Dimorphic fungus – *Blastomyces dermatidis*. Morphology and life cycle: Intracellular parasites– *Cryptosporidium* and *Plasmodium*. Intralumen parasites–*Entameoba histolytica* and *Ascaris lumbricoides*. Parasitic zoonosis– *Toxoplasma* and *Taenia*.

## Module 4 DNA viruses 12 hours

Morphology, pathogenesis and laboratory diagnosis and control measures of: DNA viruses – Hepatitis B virus. RNA viruses – Flavi virus (dengue), Retrovirus – HIV, viral zoonosis –rabies, classification of antibiotics based on mode of action: antibacterial (penicillin), antiviral (amantidine), antifungal (amphotericin) antiparasitic drugs (quinine and metraindazole). Emerging and reemerging infections (Chickungunya, Zika virus, H1N1 and Ebola). National programmes in prevention of infectious diseases

## Module 5 Control of Diseases 12 hours

Laboratory control of antimicrobial therapy; various methods of drug susceptibility testing, antibiotic assay in body fluids. Brief account on available vaccines and schedules; Passive prophylactic measures; medically important parasites – disease diagnosis, control & prevention, protozoan disease, nematode diseases.

### ReferenceBooks

- 1. Anathanarayan, R., and Jeyaram Panikers, C.K., 2013, Text Book of Microbiology, 9<sup>th</sup> Edition, Jain Book Depot, New Delhi.
- 2. Arora, D.R., Brij Bala Arora, 2015, Textbook of Microbiology, CBS, Chennai.
- 3. Awetz Melnick and Adelberg's Medical Microbiology, 21<sup>st</sup> Century, 2010.

Appleton & Lange.
4. Bhattacharjee, R.N., 2015, Introduction to Microbiology, 1 <sup>st</sup> Edition
Kalyani Publishers, New Delhi.
5. Connie R Mahon, 2010, Textbook of Diagnostic Microbiology, 3 <sup>rd</sup>
Edition, Pearson Publishers.
6. David Greenwood, Richard Slack, John Peutherer, 2012, Medical
Microbiology, Churchill Livingstone.
7. Myra Wilkinson, 2011, Medical Microbiology, Scion Publishing Ltd.
8. Patrick Murray, Ken Rosenthal and Michael Pfalle, 2015, Medical
Microbiology, 8th Edition, Academic Press, New York.
9. Patrick R. Murray, 2015, Medical Microbiology, Elsevier.

Course Outcomes	Cognitive Level
CO1: Gain information about the concepts of medical microbiology and gain knowledge on medically important microorganisms Understanding, knowledge, practical and communication skill on collection of different clinical samples, their transport, culture and examination by microscopy, staining and biochemical methods for the diagnosis of bacterial, fungal and protozoan diseases.	K1, K2, K5, K6
CO2: Gain knowledge of morphology, cultural characteristics, biochemical tests, epidemiology, laboratory diagnosis etc of bacterial pathogens	K2, K7
CO3: Knowledge about the life cycle, pathogenesis, diagnosis and treatment of yeast, fungal and protozoan diseases	K1
<b>CO4:</b> Understanding and knowledge on RNA,DNA viral disease, Mode of action of antibiotics Emerging reemerging infections;	K1

vaccine production and treatment	
CO5: Gain knowledge on various chemotherapeutic agents and	K1
their mode of action including alternatives of antibiotics and	
Alternative and Complimentary medicine	
·	

K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create

# Mapping of CO with PO

	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	Н	M	Н	M	M	Н	M	Н	M	Н	M	M	Н	Н	M
CO2	Н	Н	Н	M	Н	M	M	Н	Н	Н	M	L	L	Н	M
CO3	Н	Н	M	M	Н	M	M	M	M	Н	M	M	M	Н	M
CO4	Н	M	Н	M	Н	L	M	M	M	Н	M	M	L	Н	M
CO5	Н	M	Н	M	Н	Н	M	M	M	Н	M	M	M	L	M

H – High, M – Medium, L - Low Mapping of CO with PSO

mapping of CO with 150								
	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6		
CO1	Н	Н	Н	M	Н	M		
CO2	Н	M	M	Н	M	M		
CO3	Н	M	L	M	M	L		

CO4	Н	Н	M	M	Н	L
CO5	Н	M	M	M	Н	M

H - High, M - Medium, L - Low

Semester		III Semester							
<b>Course Type</b>		<b>Core Compulsory Paper - 12</b>							
Title of the Co	ourse	BIOREMEDIATION							
<b>Course Code</b>									
Teaching Ho	urs	60 Hours/ Semester : 4 Hours	/ week						
		EMEDIATION	Cred	its: 4	Ma	x. Mark	s: 100		
					(Intern	nal: 25, 1 75)	External		
Course Prerecagents	quisites:	The students should have a bas	ic knowl	edge on	pollutio	n, their	causative		
CODE:				L	T	P	С		
		BIOREMEDIATION		4	-	-	4		
Course Objectives	•	To get an idea about the bioren Acquire knowledge on treatm bacteria on waste water treatme To get a knowledge on bioreme	ent of i	industrial ludge pro	waste				
Module 1	Biore	emediation				12 h	ours		
In situ bioremediation, intrinsic bioremediation, engineered in situ remediation and ex situ bioremediation, solid phase and slurry phase systems and factors affecting slurry phase system.									
Module 2 Industrial Waste Water Sources and Treatment Strategies 12 hours									
Introduction and	d targets	, wastewater flow fractions from	industri	al plant,	waste w	ater froi	n sanitar		

and employee facilities, wastewater from in-plant water preparation, kinds and impacts of wastewater components, hazardous substances, corrosion inducing substances, typical treatment sequence in a wastewater treatment, plant and wastewater composition and treatment strategies in the food processing industry.

#### Module 3 **Bacterial Metabolism in Waste Water Treatment Systems** 12 hours

Decomposition of organic carbon compounds in natural and man-made ecosystems, basic biology, mass and energy balance of aerobic biopolymer degradation, mass and energy balance for aerobic and anaerobic glucose respiration and sewage sludge stabilization. General considerations for the choice of aerobic and anaerobic wastewater treatment systems, anaerobic degradation of carbohydrates in wastewater, protein, neutral fats and lipids.

#### Module 4 **Activated Sludge Process**

12 hours

Historical development, single and two stage processes, single sludge carbon, nitrogen and phosphorus removal, waste water characteristics, removal of organic carbon, nitrification, denitrification, phosphorus removal, environmental factors, carbon and nitrogen removal process and post denitrification with external organic carbon.

#### Module 5 **Bioremediation of Heavy Metals**

12 hours

Xenobiotics, microbial degradation of xenobiotics, microbial leaching – microorganisms used in leaching, chemistry of leaching, direct, indirect leaching, leaching process, examples of bioleaching, genetically modified microorganisms (GMO) in bioremediation and environmental concern.

- 1. Agarwal, S.K., 2009, Environmental Microbiology, APH Publishing Corporation, New Delhi.
- 2. Arvind Kumar, 2004, Environmental Biotechnology.
- 3. Chatterji, A.K., 2011, Introduction to Environmental biotechnology, PHI Learning Private Limited, New Delhi.
- 4. Dubey, R.C., 2001, A text book of Biotechnology.
- - 5. Garg, K.L., and Mukherji, K.G., 1993, Recent Advances in Bioremediation and biodegradation, Vol. I & II.
  - 6. Jordening, H.J., and Winter, J., 2005, Environmental Biotechnology.
  - 7. Maier, R.M., Pepper, I.L., and Gerba, C.P., 2000, Environmental Microbiology, Academic Press.
  - 8. Pelczer, M.J., and Chan, E.C.S. 1993, Microbiology, McGraw Hill Education Private Limited, New Delhi.

## ReferenceBooks

Course Outcomes	Cognitive Level
CO1: Different types of bioremediation	K1, K2
CO2: Industrial waste water from different sources and their treatment	K1, K2, K3
CO3: Basic biology of aerobic and anaerobic water treatment	K2, K3, K4
CO4: Factors affecting sludge, their treatment	K2, K3, K4
CO5: Role of microorganisms on bioremediation of heavy metals, their treatment	K2, K3, K4, K5

K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create

# **Mapping of CO with PO**

	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	Н	M	L	L	L	M	M	L	M	M	M	M	M	M	L
CO2	Н	M	M	L	L	M	M	L	M	M	Н	Н	M	M	L
CO3	Н	M	M	L	M	M	M	L	M	M	M	M	M	M	M
CO4	Н	M	L	L	M	M	M	L	M	M	M	Н	M	M	M
CO5	Н	M	Н	L	Н	M	M	M	M	M	Н	Н	M	M	Н

H – High, M – Medium, L - Low Mapping of CO with PSO

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6
CO1	M	L	Н	L	M	L
CO2	M	L	Н	L	M	L
CO3	M	L	Н	L	M	L
CO4	M	L	Н	L	M	L
CO5	M	L	Н	L	M	L

# H - High, M – Medium, L – Low

Semester	III Semester					
Course Type	Core Compulsory Course Practical - 5					
Title of the Course	RECOMBINANT DNA TEC	CHNOLOGY AN	D BIOPROCESS			
	TECHNOLOGY					
Course Code						
<b>Teaching Hours</b>	60 hours/semester: 4 hours/w	eek				
REC	COMBINANT DNA	Credits: 2	Max. Marks: 100			
TEC	CHNOLOGY AND		(Internal: 50;			
BIO	PROCESS TECHNOLOGY		External: 50)			

**Course Prerequisites:** The students should know the basics of microbes which are having industrial and pharmaceutical importance and also to produce different metabolites (products) through Fermentation technology

CODE:	RECOMBINANT DNA TECHNOLOGY	L	T	P	C
	AND BIOPROCESS TECHNOLOGY	-	-	4	2
Course Objectives	<ul> <li>To isolate industrially important microb</li> <li>To determine and optimize the culture c</li> <li>To utilize the substrates for production commercially important products by the</li> </ul>	onditions of ethar	s for mic nol, citr	crobial g ic acid a	rowth.

- To know the methodology for production of enzymes and antibiotics with the help of microbes
- To learn the immobilization technique for long term storage of microbial enzymes.
- 1. Safety, Reagent preparation and SOP in rDNA lab
- 2. Plasmid isolation and Restriction analysis
- 3. Analysis of recombinants- blue & white colony screening
- 4. PCR analysis of 16s gene
- 5. Analysis of protein by SDS-PAGE.
- 6. Demonstration blotting techniques
- 7. Isolation of industrially important microorganisms for microbial processes.
- 8. Determination of thermal death point (TDP) and thermal death time (TDT) of microorganisms for design of a sterilizer.
- 9. Comparative studies on ethanol production using different substrates.
- 10. Microbial production of citric acid using different substrates.
- 11. Microbial production of antibiotics (Penicillin).
- 12. Production and estimation of alkaline protease.
- 13. Use of alginate for cell immobilization.

Course Outcomes	Cognitive Level
CO1: Understand the techniques of rDNA technology	K2, K3
<b>CO2:</b> Isolate the industrially important microbes from different sources. Determine the culture conditions for microbial growth, especially to know the thermal death point and thermal death time of microbes	K2, K3, K4
CO3: Produce ethanol and citric acid by the identified microbes	K3, K4,K5

by utilizing different substrates	
CO4: Acquire knowledge on microbial production of antibiotics especially Penicillin and enzyme like alkaline protease.	K2, K3, K4, K5
<b>CO5:</b> Apply the immobilization technique for long term storage of microbial enzymes	K2, K3, K4, K5

# $K1-Remember,\,K2-Understand,\,K3-Application,\,K4-Analysis,\,K5-Evaluate,\,K6-Create$

# **Mapping of CO with PO**

	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	Н	M	L	L	Н	M	M	L	L	M	M	M	L	L	L
CO2	Н	M	L	L	Н	M	M	L	L	M	M	M	M	Н	L
CO3	Н	M	L	L	Н	M	M	L	L	M	M	M	M	M	L
CO4	Н	M	L	L	Н	M	M	L	L	M	M	M	M	L	L
CO5	Н	M	L	L	Н	M	M	L	L	M	M	M	M	L	L

# H – High, M – Medium, L - Low

**Mapping of CO with PSO** 

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6
CO <sub>1</sub>	Н	L	L	M	Н	L
CO <sub>2</sub>	Н	L	L	M	Н	L

CO3	Н	L	L	M	Н	L
CO4	Н	L	L	Н	M	L
CO5	Н	L	L	Н	M	L

H - High, M - Medium, L - Low

G		TIT C								
Semester		III Semester		_						
Course Type										
Title of the Co	urse	MEDICAL MICROBIOLOG	Y AND	BIORE	MEDIA	TION				
<b>Course Code</b>										
Teaching Hou	rs	0 hours/semester: 4 hours/week								
	MED	ICAL MICROBIOLOGY	Cred	its: 2	Max	x. Mark	s: 100			
	AND	BIOREMEDIATION			<b>(I</b>	nternal:	50;			
					`	xternal:				
Course Prerec	nuisites	: The students should have a l	basic pra	ctical sk	ill on r	nicrobio	logy a	nd		
chemistry	1		· · · · · ·				. 67			
CODE.	MED	ICAL MICROBIOLOGY AND	D	L	Т	P	С			
CODE:	BIOF	REMEDIATION			<del>-</del>	-				
				-	-	4	2			
	•	To identify microorganisms	s of re	levance	to hea	althcare	and	the		
		pharmaceutical industry and th								
Course	•	To evaluate microbial content			v testing	2				
Objectives	•	To determine the microbial count and Do, COD, BOD in waste water								
- · · <b>y</b> · · · · · · · ·		To determine the role of micro								
1 T 1	1 1 1		organism	15 111 1110 1	regradai	ion or pr	ubiles			

- 1. Isolation and identification of normal flora of skin.
- 2. Isolation and identification of *Streptococci* from teeth.
- 3. Serological test (WIDAL).
- 4. Antimicrobial sensitivity test by Kirby Baur method.
- 5. Diagnostic Bacteriology: Laboratory diagnosis (isolation & identification) i) Pyogenic infections Streptococci  $\alpha$ ,  $\beta$  and  $\gamma$  haemolysis. Staphylococci differentiation coagulase

- test. ii) UTI infection E.coli, Proteus, Pseudomonas
- 6. Dilution sensitivity test MIC
- 7. Isolation of E. coli from sewage water samples with the help of EMB agar
- 8. To determine the Total Dissolved Solids (TDS), Total Suspended Solids (TSS) in given water sample.
- 9. Quantitative estimation of nitrate in given water sample.
- 10. To determine the amount of Dissolved Oxygen (DO) present in given water sample.
- 11. To determine the Biological Oxygen Demand (BOD) of given waste water sample
- 12. Estimation of pollution load of a natural sample (e.g. river water /industrial waste water)
- 13. Low density plastic/bioplastic degradation using bacterial isolates

Course Outcomes	Cognitive Level
CO1: Technique and skill to isolate pathogenic bacteria in human disease with respect to infections of the respiratory tract, gastrointestinal tract, urinary tract, skin and soft tissue	K1, K2, K3
CO2: Knowledge and skill on serological diagnosis of samples, Recognize the biochemical basis for antibiotic resistance and ways of controlling spread of antibiotic resistance.	K1, K2, K3
CO3: Demonstrate practical skills in fundamental microbiological techniques, opportunities to develop informatics and diagnostic skills, including the use and interpretation of laboratory tests in the diagnosis of infectious diseases	K1, K2, K3, K4, K5
<b>CO4:</b> Understand and get the skill for the determination of TDS, TSS, DO, BOD, COD and <i>E.coli</i> in waste water	K3, K4, K5
CO5: Understand the mechanism behind the degradation of	K2, K3, K4

plastics by microorganism	

K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create

Mapping of CO with PO

P	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,														
	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	Н	M	Н	Н	Н	M	M	M	Н	M	Н	M	M	L	M
CO2	Н	M	Н	Н	Н	M	M	M	Н	M	Н	M	M	L	M
CO3	Н	M	Н	Н	Н	M	M	M	Н	M	Н	M	M	L	M
CO4	Н	M	M	M	Н	M	M	M	Н	M	Н	M	M	L	M
CO5	Н	M	M	M	Н	M	M	M	Н	M	Н	M	M	L	M

H – High, M – Medium, L - Low

**Mapping of CO with PSO** 

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6
CO1	Н	Н	Н	L	L	L
CO2	Н	Н	Н	L	L	L
CO3	Н	Н	Н	L	L	L
CO4	Н	M	Н	L	M	L
CO5	Н	M	Н	L	Н	L

H - High, M - Medium, L - Low

Semester		III Semester									
<b>Course Type</b>		Core Elective Paper – 3A									
Title of the Co	urse	<b>BIOSAFEY, BIOETHICS &amp;</b>	IPR								
<b>Course Code</b>											
<b>Teaching Hou</b>	rs	45 Hours/ Semester: 3 Hours	s/ week								
		Max. Marks: 100 ternal: 25, External 75)									
Course Prereq	uisites:	Students should have a knowled	ge on bio	ological 1	regulatio	ns					
CODE:				L	T	P	С				
0022		BIOSAFEY, BIOETHICS & I	PR	3	-	-	3				
Course Objectives											
Module 1	Intro	duction to Bio-safety and Regu	llations			9 h	ours				
Experimental papplications, deg	rotocol gradatio	and definitions, national and in approvals, levels of conta n of pollutants, Bacterial mining ganisms and their release in e	inment, g- vaccin	environ es- Biol	mental	aspects of	of biotech				
Module 2	Bioet	hics				9 h	ours				
studies on huma materials devel	n and ar	ethics- norms in India- Licensin nimal subjects- Ethical clearance -environmental safety and impelated to animal models	, ELSI. I	Bioethics	for co	smetics	and nano				
Module 3	IPR -	- Introduction & Registration				9 h	ours				
		Basic concepts and need for Ins., IPR in India and Abroad –			•		10 0				

WTO to WIPO –TRIPS, Nature of Intellectual Property, Industrial Property, technological Research, Inventions and Innovations – Important examples of IPR. Meaning and practical aspects of registration of Copy Rights, Trademarks, Patents, Geographical Indications, Trade Secrets and Industrial Design registration in India and Abroad.

# Module 4 IPR – Agreements, Legislations, Digital Products and Law 9 hours

International Treaties and Conventions on IPRs, TRIPS Agreement, PCT Agreement, Patent Act of India, Patent Amendment Act, Design Act, Trademark Act, Geographical Indication Act. Digital Innovations and Developments as Knowledge Assets – IP Laws, Cyber Law and Digital Content Protection – Unfair Competition – Meaning and Relationship between Unfair Competition and IP Laws – Case Studies. Infringement of IPRs, Enforcement Measures, Emerging issues – Case Studies.

# Module 5 Enforcement of IPRs and Patents 9 hours

Infringement of IPRs, Enforcement Measures, Emerging issues – Case Studies. Definition of patents- conditions for patent ability: Novelty testing- composition of patent - patenting of biotechnological discoveries - commercialization- Biotech companies - Natures implications-screening and selection of genetic materials for patenting- Public attitude - Genetic counselling.

- 1. Cartagena Protocol on Biosafety, 2006, Ministry of Environment and Forest, Government of India, New Delhi
- 2. Gopalakrishnan, N.S., and Agitha, T.G., 2014, Principles of Intellectual Property, 2<sup>nd</sup> Edition, Eastern Book Company, Lucknow.
- 3. Kshitij Kumar Singh, Springer, 2015, Biotechnology and Intellectual Property Mittal, D.P., 1999, Rights: Legal and Social Implications, Indian Patents Law, as amended by Patents (Amendment) Act 1999, Taxman Publication.
- 4. Laboratory Biosafety Manual, 2004, 3<sup>rd</sup> Edition, World Health Organization, Geneva.
- 5. Narayanan, P., 2021, Intellectual Property Law, 3<sup>rd</sup> Edition, Eastern Law Book House Ltd.
- Sasson, A., 1988, Biotechnologies and Development, UNESCO Publications.
- 7. Sasson, A., 1993, Biotechnologies in Developing Countries: Present and Future, UNESCO Publications.

### ReferenceBooks

- 8. Singh, K., Intellectual Property Rights on Biotechnology, BCIL, New Delhi.
- 9. Wadera, B.L., 2010, Law relating to Patents Trade Marks Copyright Designs and Geographical Indications, Delhi University Law Publishing.

Course Outcomes	Cognitive Level
CO1: Remember the basic knowledge on biosafety, national and international guidelines and regulations for biosafety. Gain knowledge on organization rDNA guidelines, approval of protocols, Analyze the benefits of degradation of pollutants, bacterial mining, vaccines and biological pesticides. Evaluate the use of environmental aspects of biotech applications, and genetically modified organisms and their release in environment	K1, K2, K3
CO2: Remember the definition of ethics, Gain knowledge on ethical clearance, ELSI (Ethical, Legal and Social Implications, Comprehend the application of ethical principles in India, Analyze the norms of bioethics in India, Acquire knowledge on licensing of animal house and norms for conducting studies on human and animal subjects – regarding ethical clearance	K1, K2, K3
CO3: Understand and remember concepts of IPR, copyrights etc	K1, K2, K3
CO4: Remember the IPRs, Treaties, TRIPS agreement and laws	K1, K2, K3
CO5: Acquire knowledge on obtaining Indian patent and international patents	K1, K2, K3

	PO1	PO2	PPO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	M	Н	L	L	L	M	L	L	L	L	L	L	M	L	L
CO2	L	M	L	L	L	1	L	L	L	L	M	L	M	L	L
CO3	M	M	L	L	L	Н	M	M	M	L	L	L	M	L	L
CO4	L	L	M	M	M	Н	L	M	L	L	L	L	M	L	L
CO5	L	L	L	L	L	L	L	L	L	L	M	M	M	M	M

**Mapping of CO with PSO** 

	PSO1	PSO2		PSO4	PSO5	PSO6
	1501	1502	1505	1504	1505	1500
CO1	M	L	Н	L	M	Н
CO2	M	L	M	L	M	Н
CO3	Н	Н	Н	L	L	Н
CO4	M	M	L	M	M	Н
CO5	L	L	M	L	L	Н

H - High, M - Medium, L - Low

Semester	III Semester
<b>Course Type</b>	Core Elective Paper – 3B

Title of the Cou	rse	BIOINFORMATICS						
<b>Course Code</b>								
Teaching Hour	S	45 Hours/ Semester: 3 Hours	/ week					
	BIOI	NFORMATICS	Cred	its: 4		x. Marks mal: 25, l 75)		
Course Prereq bioinformatics	uisites	: The students should have	a basic	knowl	edge o	n compi	uter and	
CODE:		<b>D.O.D. D. C.I. M.</b> GG	BIOINFORMATICS         L         T         P         C           3         -         -         -         -					
		BIOINFORMATICS						
Course Objectives Module 1	• • Intro	To know about the protein, the using the bioinformatics tools. To get an idea about the sequent To get a gene prediction and production of Bioinformatics.	nce simil	arity usii				
EMBnet-, intrane	t and	of bioinformatics, internet at internet packages, basics, WWV quence databases, protein sequen	V, HTMI	L, URLs				
Module 2	Prote	ins				9 ho	ours	
primary, secondar domain database	ry, sup s, Ra ds, pro	er-secondary, tertiary and quate machandran map, prediction of ediction of 3D structures, home	rnary stru of protei	ucture, p in struc	rotein s ture, se	equence :	motif and structure	
Module 3	Seque	ence Similarity				9 ho	ours	
		nce similarity, pair-wise sequences, Scoring matrices, PAM and						

algorithm, multiple sequence alignments (MSA), Importance of MSA, Clustal W and Phylip. Definition and description of phylogenetic trees and various types of trees, methods and programs for phylogenetic tree construction.

#### Module 4 Gene Prediction

9 hours

Prediction of genes, gene prediction in prokaryotes and eukaryotes, promoters, splice sites, regulatory regions, Comparative genomics, functional genomics, DNA microarray, basic concepts on identification of disease genes, OMIM database, Pharmacogenomics, Identification of SNPs, SNPs databases (DbSNP), Metabolic pathways, databases such as KEGG, EMP.

#### Module 5 Proteomics

9 hours

Introduction to proteomics, steps in proteomics research, two-dimensional gel electrophoresis, mass spectrometry, MALDI, ESI, protein identification and characterization strategies, 2D Gel maps, applications of proteomics, proteomics in disease diagnosis, protein arrays – basic principles, drug designing, drug designing approaches, chemoinformatics.

- Andrew Leach, and Valerie J. Gillet, 2003, An Introduction to Chemoinformatics, Kluwer Academic Publishers.
- 2. Arthur, M. Lesk, 2003, Introduction to Bionformatics, Oxford University Press, New Delhi.
- 3. Attwood, T.K., and Parry-Smith, D.J., 2004, Introduction to Bioinformatics, Pearson Education Ltd., New Delhi.
- 4. Baxevanis, A., and Quellette, B.F., 1998, Bioinformatics: A practical guide to the Analysis of Genes and proteins, Wiley-Interscience, Hoboken, NJ.

### ReferenceBooks

- 5. Baxevanis, A.D., Davison, D.B., Page, R.D.M., and Petsko, G.A., 2004, Current Protocols in Bioinformatics, John Wiley & Sons Inc., New York.
- 6. Branden, Carl, Tooze & John, 1991, Introduction to Protein Structure, Garland Publishing.
- 7. Durbin R., Eddy, S., Krogh, A., and Mitchison, G., 1998, Biological Sequence Analysis: Probabilistic Models of Proteins and Nucleic Acids, Cambridge University Press.
- 8. Higgins, D., and Taylor, W., 2000, Bioinformatics Sequences, Structure and Databanks, Oxford University Press, New Delhi.

- 9. Mount and David, 2004, Bioinformatics: Sequence and Genome Analysis, Cold Spring Harbour Laboratory Press, New York.
- 10. Swindell, S.R., Miller, R.R., and Myers, G.S.A., 1996, Internet for the Molecular Biologist, Horizon Scientific Press, Wymondham, UK.

Course Outcomes	Cognitive Level
CO1: Basci bioinformatics and the tools involved	K1, K2, K3
CO2: Structure of protein and their different structural predictions	K2, K3, K4
CO3: Understand the different methods in sequence similarity	K2, K3, K4
CO4: Understand the knowledge on gene prediction in prokaryotes and eukaryotes, basic concepts of disease genes	K2, K3, K4, K5
CO5: Acquire knowledge on proteomics & chemo-informatics, the techniques involved in protein identification and characterization.	K2, K3, K4, K5

K1 – Remember, K2 – Understand, K3 – Application, K4 – Analysis, K5 – Evaluate, K6 – Create

	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PO13	PO14	PO15
CO1	Н	M	M	Н	M	M	M	M	L	Н	Н	M	M	L	M
CO2	Н	M	Н	Н	M	M	M	M	L	Н	Н	M	M	L	M

CO3	Н	M	Н	Н	M	M	M	M	L	Н	Н	M	M	L	M
CO4	Н	M	Н	Н	M	M	M	M	L	Н	Н	M	M	L	M
CO5	Н	M	Н	Н	M	M	M	M	L	Н	Н	M	M	L	M

H – High, L – Low, M - Medium

**Mapping of CO with PSO** 

	PSO1	PSO2	PSO3	PSO4	PSO5	PSO6
CO1	Н	L	L	L	Н	L
CO2	Н	L	L	L	Н	L
CO3	Н	L	L	L	Н	L
CO4	Н	L	L	L	Н	L
CO5	Н	L	L	L	Н	L

H - High, M - Medium, L - Low