# S. P. Mandali's Ramnarain Ruia Autonomous College

(Affiliated to University of Mumbai)



Syllabus for F.Y, S.Y, T.Y

**Program: BSc** 

**Program Code: Microbiology (RUSMIC)** 

(Credit Based Semester and Grading System for academic year 2020–2021)



# **PROGRAM OUTCOMES**

РО	PO Description
	A student completing Bachelor's Degree in Science program will be
	able to:
PO 1	Recall and explain acquired scientific knowledge in a comprehensive
	manner and apply the skills acquired in their chosen discipline.
	Interpret scientific ideas and relate its interconnectedness to various
	fields in science.
PO 2	Evaluate scientific ideas critically, analyse problems, explore options
	for practical demonstrations, illustrate work plans and execute them,
	organise data and draw inferences.
PO 3	Explore and evaluate digital information and use it for knowledge
	upgradation. Apply relevant information so gathered for analysis and
	communication using appropriate digital tools.
PO 4	Ask relevant questions, understand scientific relevance, hypothesize
	a scientific problem, construct and execute a project plan and
	analyse results.
PO 5	Take complex challenges; work responsibly and independently, as
	well as in cohesion with a team for completion of a task.
	Communicate effectively, convincingly and in an articulate manner.
PO 6	Apply scientific information with sensitivity to values of different
. 0	cultural groups. Disseminate scientific knowledge effectively for
167	upliftment of the society.
PO 7	Follow ethical practices at work place and be unbiased and critical in
24	interpretation of scientific data. Understand the environmental issues
	and explore sustainable solutions for it.
PO 8	Keep abreast with current scientific developments in the specific
	discipline and adapt to technological advancements for better
	application of scientific knowledge as a lifelong learner



# **PROGRAM SPECIFIC OUTCOMES**

PSO	Description		
	A student completing Bachelor's Degree in Science program in the subject of Microbiology will be able to:		
PSO 1	Recall, explain and summarize basic concepts related to cytology, biochemistry, physiology, genetics and reproduction of prokaryotes and compare it with eukaryotes.		
PSO 2	Appreciate and exemplify the diversity in the microbial world and evaluate their ecological role as well as state their significance to humankind.		
PSO 3	Understand the basic concepts associated with growth and control of microorganisms and apply it in pure culture and preservation techniques.		
PSO 4	Differentiate, classify and characterize microorganisms on the basis of their morphological, cultural, biochemical, and molecular properties.		
PSO 5	Explore, compare and evaluate the role of microorganisms in different natural environments as well as plants, animals and humans, and evaluate and exemplify their interrelationships.		
PSO 6	Apply the understanding of microbial processes to diverse science areas such as medical, industrial, agricultural and food and evaluate their potential for human well-being, for tackling environmental issues and exploring sustainable solutions		
PSO 7	Recall and explain the nature of biomolecules and metabolic processes; the role and kinetics of enzymes as well as the thermodynamic laws that drive these reactions.		
PSO 8	Recall the basic working principles of various bioanalytical techniques and tools and apply them to detect, estimate and structurally evaluate biomolecules present in the microbial cells.		
PSO 9	Understand and explain the nature of genetic material and elaborate the molecular mechanisms underlying various genetic processes like replication, transcription, translation, gene transfer and recombination in bacteria; and explain basic concepts in virology.		



PSO 10	Apply the basics of genetics and molecular biology to understand and
	evaluate techniques in genetic engineering and also for the use of
	bioinformatic tools for presentation and processing of data.
PSO 11	Recognize and explain the role of microorganisms in different diseases, attribute pathogenesis mechanisms to their properties and extrapolate it to
	disease diagnosis, treatment and prevention. Outline and recall concepts in
	epidemiology of diseases. Classify and evaluate different chemotherapeutic
	agents.
PSO 12	Recall, classify and summarize mechanisms of defense in humans, detail
	out the functioning of our immune system, correlate it to disease and its
	prevention and outline its association to health.
PSO 13	Understand and outline different biochemical mechanisms and their
	regulation; retrieve and construct biochemical pathways in microbial
	metabolism of major macromolecules and, recall and integrate the
	bioenergetics of metabolic reactions.
PSO 14	Evaluate, exemplify and outline the role of microorganisms in different
	industrial fermentations, summarize technological aspects of bioprocesses,
	recall knowledge about patents, copyright and regulatory practices and
	Quality Assurance.
PSO 15	Demonstrate key practical skills/competencies in working with microbes for
	their study and use in the laboratory as well as outside, including the use of
	good microbiological practices. Analyze problems involving microbes,
16	articulate them and devise innovative and creative solutions.
PSO 16	Hypothesize, design experiments, construct experimental plans, execute
25	them and analyze data with a basic understanding of statistics. Demonstrate
	an ability to be unbiased and critical in interpretation of scientific data
PSO 17	Communicate effectively to express scientific ideas and/or their
	experimental data in an effective, precise and concise manner.



# **PROGRAM OUTLINE**

YEAR	SEM	COURSE	COURSE TITLE	CREDITS
	I	RUSMIC 101	Fundamentals of Microbiology	02
FY		RUSMIC 102	Microorganisms – in the lab and in nature	02
		RUSMICP101	Practicals based on above two courses	02
	II	RUSMIC 201	Microbial world: types and inter-relations	02
		RUSMIC 202	Techniques in Microbiology	02
		RUSMICP201	Practicals based on above two courses	02
	III	RUSMIC 301	Microbial taxonomy and Introduction to Genetics and Molecular Biology	02
		RUSMIC 302	Introduction to Experimental Microbial Biochemistry	02
		RUSMIC 303	Environmental Microbiology	02
SY	RI	RUSMICP301	Practicals based on above three courses	03
	IV	RUSMIC 401	Microbe interactions and host responses	02
		RUSMIC 402	Introduction to Metabolic Pathways and Enzymology	02
		RUSMIC 403	Applied Microbiology	02
		RUSMICP401	Practicals based on above three courses	03



	V	RUSMIC 501	Microbial Genetics	2.5
		RUSMIC 502	Medical Microbiology	2.5
		RUSMICP501	Practical Based on Above Two Courses	3
		RUSMIC 503	Microbial Biochemistry: Part-I	2.5
		RUSMIC 504	Bioprocess Technology	2.5
		RUSMICP502	Practical Based on Above Two Courses	3
TY	VI	RUSMIC 601	Genetics, Bioinformatics & Virology	2.5
		RUSMIC 602	Immunology	2.5
		RUSMICP601	Practical Based on Above Two Courses	3
		RUSMIC 603	Microbial Biochemistry Part II	2.5
		RUSMIC 604	Industrial Microbiology	2.5
		RUSMICP602	Practical Based on Above Two Courses	3
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# **Course Code: RUSMIC 101**

# **Course Title: Fundamentals of Microbiology**

# Academic year 2020-21

COURSE OUTCOME	DESCRIPTION
CO 1	Understand and explain the process of formation of earth and
	evolution of microorganisms on earth.
CO 2	Summarize the key events in the history of microbiology
CO 3	Recognize the scope and relevance of microbiology
CO 4	Recall and explain the nature, correlate function of components that
	make up a prokaryotic cell and identify them microscopically
CO 5	Compare and contrast between structural features of prokaryotic
	and eukaryotic cell
CO 6	Recall the characteristics and structures of biomolecules and
	classify and detect them in various samples
PAMMAR	



Course Code/ Unit	Unit	Course/ Unit Title	Credits/ Lectures
RUSMIC		FUNDAMENTALS OF MICROBIOLOGY	2/45
101			
I		Evolution of Microbes, History and Future of Microbiology	15
	1.1	The Evolution of Microorganisms	07
		<ul> <li>a) Formation and Early History of Earth</li> <li>b) Origin of Cellular life.</li> <li>c) RNA world hypothesis and protein synthesis</li> <li>d) Microbial Diversification</li> <li>e) Endosymbiotic origin of prokaryotes</li> <li>f) Microbial Evolution - Process</li> </ul>	
	1.2	History, Branches and Scope of Microbiology	06
		a) Discovery of microorganisms	
		b) Conflict over spontaneous generation	
		<ul> <li>c) Golden Age of Microbiology-Koch Postulate Medical Microbiology, Immunology</li> <li>d) Development of industrial microbiology an</li> </ul>	
		microbial ecology  e) Scope and relevance of microbiology	
	1.3	Future of Microbiology and unification with other	02
		sciences	
	Q.P	a) Molecular and genomic methods to stud microorganisms     b) Emerging diseases	У
		<ul><li>b) Emerging diseases</li><li>c) Search for extra-terrestrial life</li></ul>	
		d) Bio-based economies	
JI.		Prokaryotic and Eukaryotic Cell Structure	15
2.1	2.1	Prokaryotic Cell Structure and functions	10
		<ul> <li>a) Overview of prokaryotic cell structure</li> <li>b) Cell wall</li> <li>c) Cell membrane</li> <li>d) Components external to cell wall-Capsule, Slim layer, Flagella, Pili, Fimbriae</li> </ul>	е



		<ul> <li>b) Monosaccharides, (Chair and boat conformation) oligosaccharides (maltose,</li> </ul>	
		<ul><li>a) Definition, Classification, Biological role.</li><li>b) Monosaccharides, (Chair and boat</li></ul>	
	3.3	Carbohydrates and glycobiology	04
281	3.2	Water- Structure, properties in brief	01
		Electrovalence, covalent, ester, phosphodiester thioester, peptide, glycosidic.	'
2	M,	f) Types of bonds and their importance	
	A	stereoisomerism in biology.	
		e) Types of Stereoisomers and importance o	f
		and suitable examples only.	
		cells. d) Configuration and Conformation with definitions	
		c) Macromolecules as the major constituents o	f
		b) Universal set of small molecules.	
		variety of functional groups.	
	J. I	a) Biomolecules as compounds of carbon with a	
	3.1	Chemical foundations	02
III		Chemical basis of life	15
		i) Mitosis & meiosis	
		h) Comparison of Prokaryotic and Eukaryotic Cells	
		g) Nucleus –Nuclear Structure	
		f) Chloroplasts	
		<ul><li>d) Eukaryotic ribosomes</li><li>e) Mitochondria</li></ul>	
		Proteasome	
		Golgi apparatus. Lysosome, Autophagy	
		endocytic pathways –Endoplasmic reticulum 8	
		c) Organelles of the Biosynthetic-secretory and	
		filaments, and microtubules, Cilia and Flagella	
		<ul><li>a) Overview of Eukaryotic cell structure</li><li>b) Cytoplasmic matrix, microfilaments, intermediate</li></ul>	(3)
	2.2	Eukaryotic Cell Structure	05
		g) Bacterial endospores and their formation	
		f) Nucleoid, Plasmids	
		magnetosomes, ribosomes, gas vesicles	



a) Fatty acids as basic component of lipids b) Classification, nomenclature, storage lipids and structural lipids. c) Types of lipids with general structure of each and mention examples.  Amino acids & proteins a) General structure and features of amino acids (emphasis on amphoteric nature) b) Classification by R-group, Uncommon amino	02
<ul> <li>b) Classification, nomenclature, storage lipids and structural lipids.</li> <li>c) Types of lipids with general structure of each and mention examples.</li> </ul> Amino acids & proteins <ul> <li>a) General structure and features of amino acids (emphasis on amphoteric nature)</li> </ul>	03
structural lipids. c) Types of lipids with general structure of each and mention examples.  Amino acids & proteins a) General structure and features of amino acids (emphasis on amphoteric nature)	03
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a) General structure and features of amino acids (emphasis on amphoteric nature)	03
a) General structure and features of amino acids (emphasis on amphoteric nature)	
h) Classification by R-group Uncommon amino	
b) Classification by K-group, Officialition affilia	
acids and their functions Peptides and proteins-	
Definition and general features and examples	
with biological role.	
c) Primary, secondary, tertiary, quaternary	
structures of proteins- Brief outline.	
	03
f) Basic structure of RNA and DNA.	
	with biological role. c) Primary, secondary, tertiary, quaternary structures of proteins- Brief outline.  Nucleic acids



- a) Willey, Sherwood and Woolverton, Prescott's Microbiology, 9th edition, 2013, International edition, McGraw Hill.
- b) Michael T. Madigan & J.M. Martin, Brock's Biology of Microorganisms 13th Ed. International edition 2012, Pearson Prentice Hall.
- c) https://www.space.com/search-for-life
- d) https://www.hort.purdue.edu/newcrop/ncnu02/v5-011.html
- e) https://www.weforum.org/agenda/2018/04/can-a-nature-based-economy-help-us-drive-green-growth
- Michael J. Pelczar Jr., E.C.S. Chan, Noel R. Krieg, Microbiology 5th Edition, 1986, Tata McGraw Hill Publishing Company
- g) Conn P. Stumpf, G. Bruening and R. Doi, Outlines of Biochemistry 5/E, 1995, John Wiley & Sons. New York
- h) D. Nelson and M. Cox, Lehninger's Principles of Biochemistry, 4th Edition, 2005, W.H. Freeman and Company
- i) Laurence A. Moran, H. Robert Horton, K. Gray Scrimgeour, Marc D. Perry, Principles of Biochemistry, 5th Edition, 2012, Pearson



# Course Code: RUSMIC 102 Course Title: Microorganisms- in the Lab & in Nature Academic year 2020-21

COURSE	DESCRIPTION
OUTCOME	
CO 1	Understand and explain the principle, construction & functionality
	differences of various microscopes.
CO 2	Classify the microorganisms on the basis of their growth
	requirements & explain the methods of cultivation of different
	microorganisms.
CO 3	Summarize the method & principle of the techniques used for
	visualization of microorganisms.
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CO 4	Infer the significance of different preservation techniques &
	emphasize the role of Culture collection centers.
CO 5	Recall & explain the role of microorganisms in biogeochemical
	cycles & in maintaining balance of the ecosystem
CO6	Illustrate the different types of microbial interactions & explain the
	significance of extremophiles.
60.7	Community begins at a in in a good and to wing a tank minute and to at any in a big
CO 7	Carry out basic staining and culturing techniques and test microbial
	activities using aseptic techniques



Course Code/ Unit	Unit	Course/ Unit Title	Credits/ Lectures
RUSMIC 102		MICROORGANISMS-IN THE LAB & IN NATURE	2/45
I		Cultivating & Visualizing Bacteria	15
	1.1	Microscopy	08
		<ul> <li>a) History of microscopy, Optical spectrum, Lenses and mirrors</li> </ul>	
		b) Simple and compound light microscope	
		c) Dark field Microscopy	
		d) Phase contrast Microscopy	
		e) Electron Microscopy	
	1.2	Nutrition and Cultivation of Microorganisms:	07
		a) Nutritional requirements - Carbon, Oxygen,	
		Hydrogen, Nitrogen, Phosphorus, Sulfur and	
		growth factors.	
		b) Nutritional classification on the basis of source	
		of energy, electron and carbon	
		c) Modes of nutrition: Endocytosis, Phagocytosis,	
		movement of solutes across membranes	
		d) Media Design and composition	
		e) Types of Culture media with examples	
		f) VBNC & oligotrophs	
		g) Anaerobic cultivation	
			4.5
II	02	Pure Culture techniques, Characterization &	15
7	2.1	Preservation of Bacteria Pure Culture Techniques	02
. ~	2.1	r are dutare recrimques	02
		a) Streak plate method	
0/2		b) Pour plate method	
27	2.2	Characterization of Bacteria:	11
		a) Morphological characteristics	
		b) Staining procedures	
		i. Dyes and stains: Types, Physicochemical	
		basis, Fixatives, Mordants, Decolorizers	
		ii. Simple and differential staining	
		iii. Special staining (Cell wall, Capsule, Lipid	



		granules, Spores, Metachromatic granules &	
		Flagella)	
		c) Physicochemical characterization: Influence of	
		environmental factors on growth- oxygen, pH,	
		temperature, osmotic pressure.	
	2.3	Preservation of microorganisms	02
		a) Methods for maintenance and Preservation of	
		Bacteria	
		b) Culture Collection Centers	
III		Microbes in Natural Environments	15
•••	3.1	Microorganisms in Nature	03
		a) Microenvironments	
		b) Introduction to microbial biofilms	
		c) Mixed populations and microbial consortia	
		d) Introduction to Quorum Sensing	
	3.2	Role of microbes in Biogeochemical cycles	06
		a) C- cycle, N- cycle, S- cycle, Iron cycle	
		b) Interaction between elemental cycles	
		XO	
	3.3	Microbial competition and cooperation	04
		a) Types of Microbial Interactions: Mutualism,	
		Cooperation, Commensalism, Predation,	
		Parasitism, Amensalism, Competition with examples	
		b) Functions of symbiosis	
		c) Establishment of symbiosis	
		C) Lotabilorition of Symbiosis	

- a) A.J.Salle, Fundamental Principles of Bacteriology, 1984, McGraw Hill publications
- b) Michael J.Pelczar Jr., E.C.S. Chan ,Noel R , Microbiology TMH 5th Edition
- c) Stanier, Ingraham et al, General Microbiology, 5th Ed. 1987, Macmillan Education Ltd.
- d) Tortora, Funke and Case, Microbiology: An Introduction, 6th Edition.1998, Pearson.
- e) Michael T. Madigan & J.M. Martin, Brock's Biology of Microorganisms 13th Ed. International edition 2012, Pearson Prentice Hall.
- f) Willey, Sherwood and Woolverton, Prescott's Microbiology, 7th edition, 2011, International edition, McGraw Hill.



Practicals	2 Credits
PRACTICAL-1	
Fundamentals of Microbiology	C.V
<ol> <li>Demonstration of Pasteur's experiment to refute Spontaneous Generation theory.</li> <li>Demonstration of microbes in air, cough, on table surface, finger tips, fomites etc.</li> </ol>	
<ol> <li>Study of prokaryotic subcellular structures by special staining: Cell wall, capsule, endospore, flagella, lipid, metachromatic granules.</li> <li>Study of Motility (Hanging Drop Preparation)</li> <li>Wet mount of Hay infusion</li> <li>Permanent slides of eukaryotic microorganisms</li> </ol>	
Qualitative detection     a. Carbohydrates- Benedicts, Molisch's test.     b. Proteins, amino acids- Biuret, Ninhydrin.     c. Nucleic acid detection by DPA and Orcinol	
PRACTICAL-2	
Microorganisms – In the Lab and in Nature	
<ol> <li>Parts of a microscope</li> <li>Micrometry</li> <li>Dark field and Phase Contrast Microscopy:         (Demonstration)</li> <li>Monochrome staining</li> <li>Gram staining</li> <li>Negative Staining</li> <li>Nutritional requirements- Designing media using food material</li> <li>Preparation of standard laboratory Culture Media:         <ul> <li>Liquid medium (Nutrient Broth)</li> <li>Solid Media (Nutrient agar, Sabouraud's agar)</li> <li>Preparation of slant, butts&amp; plates</li> </ul> </li> <li>Inoculation techniques and Study of Growth:         <ul> <li>Inoculation of Liquid Medium</li> </ul> </li> </ol>	
	PRACTICAL-1  Fundamentals of Microbiology  1. Demonstration of Pasteur's experiment to refute Spontaneous Generation theory. 2. Demonstration of microbes in air, cough, on table surface, finger tips, fomites etc.  1. Study of prokaryotic subcellular structures by special staining: Cell wall, capsule, endospore, flagella, lipid, metachromatic granules. 2. Study of Motility (Hanging Drop Preparation) 3. Wet mount of Hay infusion 4. Permanent slides of eukaryotic microorganisms  1. Qualitative detection a. Carbohydrates- Benedicts, Molisch's test. b. Proteins, amino acids- Biuret, Ninhydrin. c. Nucleic acid detection by DPA and Orcinol  PRACTICAL-2  Microorganisms – In the Lab and in Nature  1. Parts of a microscope 2. Micrometry 3. Dark field and Phase Contrast Microscopy: (Demonstration) 4. Monochrome staining 5. Gram staining 6. Negative Staining 7. Nutritional requirements- Designing media using food material 8. Preparation of standard laboratory Culture Media: a. Liquid medium (Nutrient Broth) b. Solid Media (Nutrient agar, Sabouraud's agar) c. Preparation of slant, butts& plates 9. Inoculation techniques and Study of Growth:



Unit-II	<ol> <li>Pure culture techniques- Streak plate method</li> <li>Study of Colony Characteristics of bacteria.</li> <li>Use of Differential &amp; Selective Media         <ul> <li>(MacConkey&amp; Salt Mannitol Agar), Enriched (Blood Agar) &amp; enrichment (Ashby's Mannitol broth)</li> </ul> </li> <li>Effect of environment on growth         <ul> <li>Temperature</li> <li>pH</li> <li>Osmotic pressure</li> </ul> </li> <li>Demonstration of anaerobic jar</li> <li>Methods of Preservation of culture- Soil stock, oil overlay and preparation of glycerol stocks, lyophilization (demo)</li> </ol>
Unit-III	Dip slide technique to demonstrate microbial biofilms     Crowded plate technique for demonstration of antibiosis     Demonstration of bacteroid forms of <i>Rhizobia</i>
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### **Modality of Assessment**

#### **Theory Examination Pattern:**

#### A. Internal Assessment- 40%- 40 Marks per paper

Sr No	Evaluation type	Marks
1	One Assignment/Case study/Project/ Presentation	15
2	One class Test (multiple choice questions / objective)	20
3	Active participation in routine class instructional deliveries	05
	TOTAL	40

#### B. External Examination- 60%- 60 Marks per paper

#### **Semester End Theory Examination:**

- 1. Duration These examinations shall be of **two hours** duration.
- 2. Theory question paper pattern:
  - a. There shall be three questions each of 20 marks one on each unit.
  - b. All questions shall be compulsory with internal choice within the questions.

#### **Paper Pattern:**

Question	Options	Marks	Questions Based on
Q.1) A)	Any 3 out of 5	15	
Q.1) B)	Any 1 set out of 2 (i & ii or i & ii)	03 &02	Unit I
Q.2) A)	Any 3 out of 5	15	l loit II
Q.2) B)	Any 1 set out of 2 (i & ii or i & ii)	03 &02	Unit II
Q.3) A)	Any 3 out of 5	15	11-2-111
Q.3) B)	Any 1 set out of 2 (i & ii or i & ii)	03 &02	Unit III
	TOTAL	60	



#### **Practical Examination Pattern:**

#### A. Internal Examination: 40%- 40 Marks

Particulars	Paper I	Paper II
Journal	05	05
Experimental tasks	10	10
Participation	05	05
Total	20	20

#### B. External Examination: 60%- 60 Marks

#### **Semester End Practical Examination:**

Particulars	Paper I	Paper II
Laboratory work	25	25
Spots/Quiz/Viva	05	05
Total	30	30

#### PRACTICAL BOOK/JOURNAL

The students are required to present a duly certified journal for appearing at the practical examination, failing which they will not be allowed to appear for the examination.

In case of loss of Journal and/ or Report, a Lost Certificate should be obtained from Head/ Coordinator / In charge of the department; failing which the student will not be allowed to appear for the practical examination.

# Overall Examination & Marks Distribution Pattern

#### Semester I

Course	101			1	02		Grand Total
	Internal	External	Total	Internal	External	Total	
Theory	40	60	100	40	60	100	200
Practicals	20	30	50	20	30	50	100



**Course Code: RUSMIC 201** 

**Course Title: Microbial World: types and inter-relations** 

Academic year 2020-21

COURSE OUTCOME	DESCRIPTION
CO 1	Understand the structure, cultivation and significance of viruses
CO 2	Explain and compare the features of Rickettsia, Chlamydia and
	Mycoplasma
CO 3	Summarize the characteristics and infer significance of
	Actinomycetes and Archaebacteria
CO 4	Categorize microorganisms like Protozoa, Algae and Fungi into
	different groups based on their characteristics
CO 5	Infer the medical and industrial significance of Protozoa, Algae and
	Fungi
CO 6	Explain the types and role of normal flora on human body and infer
	its significance
CO 7	Organizing the events of development of infection in human system
	and summarize the factors affecting host immune system



Course Code/ Unit	Unit	Course/ Unit Title	Credits/ Lectures
RUSMIC		MICROBIAL WORLD: TYPES AND	2/45
201		INTER-RELATIONS	
I		Microbial, world (Viruses Rickettsia, Actinomycetes and Archaea)	15
	1.1	Viruses	07
		<ul> <li>a) Historical highlights, General properties of viruses, prions, viroids</li> <li>b) Structure of viruses-capsids, envelopes, genomes–TMV, Influenza, and T4 as representatives</li> <li>c) Cultivation of viruses- overview</li> </ul>	
	1.2	Rickettsia, Chlamydia, Mycoplasma	02
		General features and medical significance	
	1.3	Actinomycetes	02
		a) General features     b) Examples- Nocardia and Streptomyces     c) Importance: ecological, commercial and medical	
	1.4	Archaea	02
		a) Introduction- Major Archaeal physiological groups,     b) Archaeal cell wall, lipids and membranes     c) Ecological importance	
	1.5	Cyanobacteria& Myxobacteria	02
II		Microbial World (algae, fungi, yeasts, slime molds, protozoa)	15
	2.1	Protozoa	04
Self		a) General characteristics b) Major categories of Protozoa based on motility, reproduction c) Medically important Protozoa d) Life cycle of Entamoeba	
	2.2	Algae	05
		a) Characteristics of algae: morphology, Pigments, reproduction	



	1		
		b) Cultivation of algae	
		c) Major groups of Algae –an overview	
		d) Biological, Medical and economic importance	
		e) Differences between Algae and Cyanobacteria	
		,	
		f) Medical, ecological &Commercial application	
	2.3	Fungi and Yeast	05
	2.4	a) Characteristics: structure, Reproduction	
		b) Cultivation of fungi and yeasts	, ( ^ ) ·
		c) Major fungal divisions- overview	
		d) Life cycle of yeast	
		e) Biological and economical importance	<b>V</b>
	2.5	Slime molds and Myxomycetes	01
III		Microbe- Human interactions	15
	3.1	Normal flora of the human body	04
		a) Skin, Nose &Nasopharynx, Oropharynx,	
		Respiratory tract, Eye, External ear	
		b) Mouth, Stomach, Small intestine, Large intestine	
		c) Genitourinary tract	
		d) Gnotobiotic animals	
		e) Introduction to the concept of microbiome	
		e) introduction to the concept of microbiome	
	3.2	Development of infection	07
		a) Portal of entry and infectious dose	
		b) Attaching to host	
		c) Surviving defenses	
		d) Virulence factors	
		e) Process of infection	
		f) Portal of exit	
		g) Patterns of an infection- localized, systemic,	
		focal, mixed, primary, secondary, acute and	
		chronic infections	
	$\mathcal{O}$ $Y$	h) Signs and symptoms of disease	
	Ma.	11) Signs and symptoms of disease	
	3.3	Host defense against infection: Overview	04
		a) Factors affecting host defense: Species	
M.		resistance, racial resistance and Individual	
		resistance	
2 7		b) Introduction to innate and adaptive defences,	
		Barriers at portal of entry: Physical barriers,	
		Chemical defenses, genetic resistance.	



- a) Willey, Sherwood and Woolverton, Prescott's Microbiology, 9th edition, 2013, International edition, McGraw Hill.
- b) Michael T. Madigan & J.M. Martin, Brock's Biology of Microorganisms 13th Ed. International edition 2012, Pearson Prentice Hall.
- c) Tortora, Funke and Case, Microbiology: An Introduction, 10th Edition, 2010, Pearson.
- d) Kathleen Park Talaro & Arthur Talaro Foundations in Microbiology International edition 2002, McGraw Hill.
- e) Jacquelyn Black, Laura Black, Microbiology, Principles and Explorations, 9th Ed, 2015, Wiley
- f) Michael J. Pelczar Jr., E.C.S. Chan, Noel R. Krieg, Microbiology 5th Edition, 1986, Tata



# Course Code: RUSMIC 202 Course Title: Techniques in Microbiology Academic year 2020-21

COURSE OUTCOME	DESCRIPTION
CO 1	Understand and explain the growth pattern with the phases of growth for bacteria.
CO 2	Summarize the physical, chemical &cultivation-based methods for enumeration of microorganisms.
CO 3	Recall & exemplify the mechanisms of physical & chemical antimicrobial agents.
CO 4	Infer the significance of different preservation techniques & emphasize the role of Culture collection centers.
CO 5	Understand & explain the concept and need of biosafety levels.
CO 6	Summarize the modern microscopic techniques & explain the molecular methods for detection of microorganisms.
CO 7	Execute & perform the techniques used for enumeration of microorganisms & evaluate the microbicidal action of physical & chemical agents.



Course Code/ Unit	Unit	Course/ Unit Title	Credits/ Lectures
RUSMIC 202		TECHNIQUES IN MICROBIOLOGY	2/45
I		Microbial Growth	15
	1.1	Growth Curve & Mathematical Expression of Growth Curve	05
		a) Definition of Growth, Growth phases	
		b) Determining growth constant & growth rate	
	1.2	a) Direct microscopic count i) Breed's count, ii) Petroff-Hausser counting chamber iii) Haemocytometer b) Viable count using Spread plate and Pour plate technique c) Measurements of cell constituents. d) Turbidity measurements—Brown's opacity tubes and spectrophotometer techniques e) Factors affecting growth pattern	10
II		Control of Microorganisms	15
	2.1	Definition of terms	01
	2.2	Physical agents for control of microorganisms (mode of action, advantages, disadvantages and applications)	06
		a) High temperature-moist heat and dry heat	
	•	b) Low temperatures	
W.		c) Radiation	
		d) Osmotic pressure	
		e) Desiccation	
		f) Physical removal of microorganisms using bacteriological filters	



	2.3	Chemical agents for control of microorganisms (mode of action, advantages, disadvantages and applications of all major groups of antimicrobial agents)	04
	2.4	Evaluation of Chemical disinfectants	01
	2.5	Chemotherapeutic & antimicrobial agents- types & examples (tabular form)	01
	2.6	Biosafety in Microbiology	02
		a) Biosafety general principles and terminology with equipment     b) Biological containment and laboratory safety levels	
III		Modern techniques in Microbiology	15
	3.1	Modern Microscopy	03
		a) Fluorescence microscopy     b) Confocal Microscopy	
	3.2	Molecular methods of microbe detection	10
		<ul> <li>a) Identification and quantification using nucleic acid probes and labeled antibodies (Eg: ELISA &amp; its Types, FISH)</li> <li>b) Microbial activity measurements using radioisotopes and microelectrodes</li> <li>c) PCR, Electrophoretic techniques, Hybridization techniques, Blotting techniques</li> </ul>	
	3.3	Introduction to Metagenomics, community DNA analysis	02

- a) Microbiology TMH 5th Edition by Michael J.Pelczar Jr., E.C.S. Chan ,Noel R. Krieg
- b) A.J.Salle, Fundamental Principles of Bacteriology, 1984, McGraw Hill Book Company Inc.
- c) Prescott, Hurley Klein-Microbiology, 5th edition, International edition 2002, McGraw Hill.
- d) Prescott's Microbiology, 7th Edition; Joanne M. Willey, Linda M. Sherwood, Christopher J.Woolverton, 2011, McGraw Hill International
- e) Michael T.Madigan & J.M. Martin, Brock, Biology of Microorganisms 11th Ed. International edition, 2006, Pearson Prentice Hall.
- f) Principles and Techniques of Biochemistry and Molecular Biology by Keith Wilson and John Walker, 7th edition, 2010, Cambridge University Press.



Course code	PRACTICALS	2 Credits
RUSMICP	SECTION-1	
201	MICROBIAL WORLD: TYPES AND INTER-RELATIONS	C
Unit-I	<ol> <li>Demonstration of Bacteriophages in sewage</li> <li>Isolation of Actinomycetes from soil and Slide Culture technique for Actinomycetes</li> <li>Biogas production using methanogens</li> <li>Cultivation of algae</li> </ol>	
Unit-II	<ol> <li>Isolation of yeast, and other fungi</li> <li>Fungal Wet mounts &amp; Study of Morphological Characteristics Mucor, Rhizopus, Aspergillus, Penicillium</li> <li>Slide culture of fungi</li> <li>Cultivation of fungi- static and shaker conditions</li> <li>Permanent slides of Algae, Protozoa</li> <li>Demonstration of protozoa in hay infusion</li> </ol>	
Unit-III	<ol> <li>Normal flora of the skin, oral cavity and intestine.</li> <li>Role of fomites</li> <li>Cough plate technique</li> </ol>	
RUSMICP	SECTION-2	
202	TECHNIQUES IN MICROBIOLOGY	
Unit-I	Study of growth curve of bacteria     Enumeration of microorganisms using Haemocytometer & Breed's Count     Enumeration of microorganisms Brown's opacity tubes     Viable count: Spread plate and pour plate	
Unit-II	<ol> <li>Demonstration of efficiency of autoclave</li> <li>Effect of UV Light on bacteria</li> <li>Effect of surface tension on bacterial growth</li> <li>Study of Oligodynamic action</li> <li>Effect of dyes, phenolic compounds and chemotherapeutic agents on bacteria- disc diffusion method</li> <li>Demonstration of MIC of an antibacterial agent</li> </ol>	
Unit-III	<ol> <li>Introduction to laboratory equipment for electrophoresis, PCR</li> <li>Assignment on any modern method used in microbial detection</li> </ol>	



# **Modality of Assessment**

#### **Theory Examination Pattern:**

#### A. Internal Assessment- 40%- 40 Marks per paper

Sr No	Evaluation type	Marks
1	One Assignment/Case study/Project/ Presentation	15
2	One class Test (multiple choice questions / objective)	20
3	Active participation in routine class instructional deliveries	05
	TOTAL	40

# B. External Examination- 60%- 60 Marks per paper

#### **Semester End Theory Examination:**

- 1. Duration These examinations shall be of **two hours** duration.
- 2. Theory question paper pattern:
  - a. There shall be three questions each of 20 marks one on each unit.
  - b. All questions shall be compulsory with internal choice within the questions.

#### Paper Pattern:

Question	Options	Marks	Questions Based on
Q.1) A)	Any 3 out of 5	15	
Q.1) B)	Any 1 set out of 2 (i & ii or i & ii)	03 &02	Unit I
Q.2) A)	Any 3 out of 5	15	l leste II
Q.2) B)	Any 1 set out of 2 (i & ii or i & ii)	03 &02	Unit II
Q.3) A)	Any 3 out of 5	15	11
Q.3) B)	Any 1 set out of 2 (i & ii or i & ii)	03 &02	Unit III
	TOTAL	60	



#### **Practical Examination Pattern:**

#### A. Internal Examination: 40%- 40 Marks

Particulars	Paper I	Paper II
Journal	05	05
Experimental tasks	10	10
Participation	05	05
Total	20	20

#### B. External Examination: 60%- 60 Marks

#### **Semester End Practical Examination:**

Particulars	Paper I	Paper II
Laboratory work	25	25
Spots/Quiz/Viva	05	05
Total	30	30

#### PRACTICAL BOOK/JOURNAL

The students are required to present a duly certified journal for appearing at the practical examination, failing which they will not be allowed to appear for the examination.

In case of loss of Journal and/ or Report, a Lost Certificate should be obtained from Head/ Coordinator / Incharge of the department; failing which the student will not be allowed to appear for the practical examination.

#### **Overall Examination & Marks Distribution Pattern**

#### Semester II

Course	201			2	02		Grand Total
	Internal	External	Total	Internal	External	Total	
Theory	40	60	100	40	60	100	200
Practicals	20	30	50	20	30	50	100



#### **Course Code: RUSMIC 301**

# Course Title: MICROBIAL TAXONOMYAND INTRODUCTION TO GENETICS AND MOLECULAR BIOLOGY

COURSE	DESCRIPTION
OUTCOME	
CO 1	Differentiate between vast pool of microbes on the basis of
	morphological, cultural, biochemical and genetic characteristics
CO 2	Understand, apply and evaluate techniques in microbial taxonomy
CO 3	Construct phylogenetic trees using simple computational tools
CO 4	Recall Mendelian genetics and critique the deviations from Mendelian
	genetics
CO 5	Discriminate the structure of DNA and RNA focusing on the different
	forms of DNA
CO6	Understand the central dogma of molecular genetics
C07	Explain prokaryotic transcription and translation process and interpret
	the significance of the important events from initiation to the
0	termination of the process
CO8	Extrapolate the role of omics in molecular biology studies



Course Code/ Unit	Unit	Course/ Unit Title	Credits/ Lectures
RUSMIC		MICROBIAL TAXONOMY AND	2 / 45
301		INTRODUCTION TO GENETICS AND	700
301			
		MOLECULAR BIOLOGY	
I		Techniques in Microbial Taxonomy	15
	1.1	Introduction to microbial Taxonomy and Taxonomic	01
		ranks	
	1.2	Techniques for studying Microbial Taxonomy	08
		a) Microscopic & macroscopic morphology and	
		biochemical characteristics,	
		b) Chemical Analysis	
		c) Serological analysis	
		d) Genetic & molecular analysis: i. Nucleic acid	
		sequencing and finger printing ii. G+C content iii.	
		Nucleic acid hybridization iv. Amino acid	
		sequencing	
		e) Community DNA analysis	
		,,	
	1.3	Introduction to Microbial Phylogeny	05
		a) Phylogenetic Trees	
		i. Types	
		ii. Construction (an overview)	
	(24)	b) Numerical Taxonomy	
		by Trainerieal Fazieriening	
	1.4	Bergey's Manual of Systematic Bacteriology	01
		a) Understanding classification and identification	
2 Alla		schemes for bacteria using Bergey's manual	
II		Classical Genetics (Mendelian & Neomendelian)	15
		& Nucleic acid structure	
	2.1	Mendelian genetics:	04
		a) Genotype and Phenotype	
		b) Mendel's Experiments design	



		<ul> <li>c) Monohybrid cross and dihybrid cross, Mendelian Laws of inheritance</li> <li>d) Trihybrid Cross</li> </ul>	
	2.2	Neomendelian genetics	05
		<ul> <li>a) Multiple alleles</li> <li>b) Modification of dominance relationships</li> <li>c) Incomplete dominance</li> <li>d) Codominance (both with their molecular explanations)</li> <li>e) Essential and lethal genes</li> <li>f) Gene expression and effect of environment</li> <li>g) Maternal effect</li> <li>h) Gene interactions and modified Mendelian ratios</li> </ul>	Y.G.
	2.3	Structure of DNA:	03
		Different 3D forms and unusual structures DNA methylation	
	2.4	Structure of chromosomes	01
	2.5	Structure of RNA	02
III		Gene Expression in Bacteria	15
	3.1	Central dogma of Molecular Biology	01
	3.2	Transcription in prokaryotes	06
	28	a) RNA biosynthesis b) Prokaryotic transcription i. Prokaryotic promoters ii. Initiation, elongation and termination	
	3.3	Translation	06
PAMI		a) Components of protein synthesis apparatus: Genetic code, mRNA, Ribosomes b) Degeneracy of genetic code c) Protein synthesis	
	3.4	Comparison of eukaryotic & prokaryotic transcription & translation	01
	3.5	Introduction to the concept of Omics: Genomics and Proteomics	01



- a) Prescott's Microbiology, Joanne M. Willey, Linda M. Sherwood, Christopher J.Woolverton, Edition, 7th Edition, 2011, McGraw Hill International
- b) Madigan, Martinko, Dunlap and Clark, Brock Biology of Microorganisms, 12thedition, 2009, Pearson Education
- c) Peter J. Russell, "iGenetics A molecular approach", 3rd edition, 2010, Benjamin Cummings.7
- d) Stanier R.Y. And Other, MacMillan General Microbiology, 5th edition,1987, MACMILLAN PRESS LTD
- e) D. Nelson & M. Cox, Lehninger"s Principles Of Biochemistry,4th Edition ,2005, (W.H.Freeman& Co., (LPE)
- f) James Watson, Molecular Biology of Gene, 5th edition,2004, Pearson Benjamin Cummings CSHL Press.
- g) Benjamin A Pierce, Genetics: A conceptual approach ,2002, W.H. Freeman



#### **Course Code: RUSMIC 302**

# Course Title: INTRODUCTION TO EXPERIMENTAL MICROBIAL BIOCHEMISTRY

# Academic year 2020-21

COURSE OUTCOME	DESCRIPTION
CO 1	Understand the process of designing experiments & analyze the
	experimental data statistically.
CO 2	Implement the use of web directories & databases for
	biochemical studies
CO 3	Recall & compare the different cell disintegration methods &
	elaborate the working principles of centrifugation,
	electrophoretic & chromatographic techniques used for studying
	cell analytes.
CO 4	Illustrate the principles of protein separation & purification.
CO 5	Compare the utility & perform the techniques for the estimation
	of biomolecules.
CO 6	Understand the principle, instrumentation & application of
"Hy	different laboratory instruments used in biochemical studies.
CO 7	Design an experiment for extraction, purification & estimation of
	biomolecules, & evaluate the statistical relevance of the data
	generated.



Course Code	Unit	Course/ Unit Title	Credits/ Lectures
RUSMIC 302		INTRODUCTION TO EXPERIMENTAL MICROBIAL BIOCHEMISTRY	2/45
I		Designing and Analysis of experimental data, General laboratory techniques:  Electrochemical sensors	15
	1.1	Designing experiments:	02
		a) Aims of laboratory experiments     b) Outline of Scientific method     c) Experimental design     d) Analytical considerations and experimental error	
	1.2	Analysis of experimental data:	07
	RA	<ul> <li>a) Data presentation: Dot diagram, Bar diagram, Histogram, Frequency curve, Calibration methods: Linear regression, Internal standards</li> <li>b) Assessment of precision -Mean, Median, Mode, Standard deviation, coefficient of variation and variance</li> <li>c) Assessment of performance of an analytical technique -performance indicators</li> <li>d) Poisson and Normal distribution</li> <li>e) Assessment of accuracy&amp; Validation of analytical data population statistics, confidence limit and confidence interval; Students t factor, Q test, F test, ANOVA</li> </ul>	
	1.3	Using computers in biochemistry	02
AMI		Using web directories, biological databases and tools (eg. NCBI, EMBL)	
	1.4	General and routine laboratory procedures:	04
		Theoretical and practical aspects of:  a) Preparation and use of buffers b) Electrochemical sensors: pH meter c) Oxygen electrode d) Biosensors	



II		Fractionation of microbial cells and separation techniques	15
	2.1	Disintegration of cells	02
		a) Physical methods	_
		b) Chemical methods	
	2.2	Separation Techniques	03
		a) Centrifugation techniques: i. Basic principles of sedimentation ii. Types of centrifuges and their use: preparative & analytical, ultracentrifuges iii. Density Gradient & isopycnic centrifugation	
		b) Electrophoretic techniques: i.General Principles ii.Factors affecting electrophoresis iii.Support media- Agarose gels and PAGE	03
		c) Chromatographic Techniques: i. General principles ii. Types and applications- Partition, adsorption, ion exchange, affinity and size exclusion iii. Modes- Paper, TLC, HPLC, GC, Reverse Phase	07
III		Purification & Estimation of biomolecules	15
	3.1	Separation and purification of proteins	03
2 ANN	ARA	<ul> <li>a) Criteria for purity</li> <li>b) Methods of separation/ concentration of proteins based on: <ol> <li>i. Size and mass</li> <li>ii. Polarity</li> <li>iii. Solubility</li> <li>iv. Specific binding sites</li> <li>v. Concentration of proteins - Dialysis, Ultrafiltration</li> <li>c) Choice of methods</li> </ol> </li> </ul>	



3.2	Estimation of Biomolecules	12
	a) Visible and UV spectrophotometry	03
	i. Principles	
	ii. Instrumentation	
	iii. Applications	_
	b) Preparation of bacteria for analysis	01
	<ul> <li>c) Methods for chemical analysis (Basic principles of all methods to be covered)</li> </ul>	08
	<ul> <li>i. Methods of elemental analysis: Carbon by Slyke's method, Nitrogen by Microkjelhdahl method, Phosphorus by</li> </ul>	
	Fiske-Subbarow method  ii. Estimation of Carbohydrates by	
	Phenol and Anthrone Method	
	iii. Estimation of Reducing Sugars	
	iv. Estimation of Proteins	
	v. Estimation of Amino acids	
	vi. Extraction of Lipids and estimation of total lipid	
	vii. Estimation of Nucleic acids	

- a) Norris & Ribbon, Methods in Microbiology, Vol.5B, Edition, 1971, Academic Press
- b) J. Jayaraman, Laboratory Manual in Biochemistry, 2003, New Age International Publishers
- c) D. Nelson & M. Cox, Lehninger's Principles Of Biochemistry,4th Edition, 2005, W.H.Freeman & Co., (LPE)
- d) B.K. Mahajan. Jaypee brothers, Methods in biostatistics for medical & research workers. 6thedition, Medical Publishers (P) ltd.
- e) Rodney Boyer, Modern experimental biochemistry by 3rd Edition ,2000, Benjamin Cummings
- f) I.H. Segel, Biochemical calculations, 2nd Edition 2004, Wiley India
- g) Wilson and Walker, Principles and Techniques of Biochemistry and Molecular Biology 7th Ed, 2010. Cambridge University Press
- h) Stanier R.Y. And Other, General Microbiology, 5th edition, 1989 MacMillan Press.
- i) Plummer David, An Introduction to Practical Biochemistry, 1979, TMH
- j) Wayne Daniel, Biostatistics: A Foundation for Analysis in Health Sciences, 10th edition, 2013, Wiley.



# Course Code: RUSMIC 303 Course Title: ENVIRONMENTAL MICROBIOLOGY

## Academic year 2020-21

CO 1 Understand the distribution and characterization of microbes in various habitats/ecosystems  CO 2 Explain role of air as a medium of microbial dispersion  CO 3 Differentiate between microbial flora of marine and fresh environments  CO 4 Execute microbiological techniques for studying microbiota of air, and and terrestrial environments  CO 5 Implement routine bacteriological analysis techniques for assessing with quality and attribute the results to sources of contamination  CO 6 Recall steps in sewage treatment and check effectivity of treat processes
CO 3  Differentiate between microbial flora of marine and freshven environments  Execute microbiological techniques for studying microbiota of air, and terrestrial environments  CO 5  Implement routine bacteriological analysis techniques for assessing valuality and attribute the results to sources of contamination  Recall steps in sewage treatment and check effectivity of treat
co 4  Execute microbiological techniques for studying microbiota of air, aquant terrestrial environments  Co 5  Implement routine bacteriological analysis techniques for assessing was quality and attribute the results to sources of contamination  Recall steps in sewage treatment and check effectivity of treat
and terrestrial environments  CO 5  Implement routine bacteriological analysis techniques for assessing water quality and attribute the results to sources of contamination  Recall steps in sewage treatment and check effectivity of treat
quality and attribute the results to sources of contamination  Recall steps in sewage treatment and check effectivity of treat
(40 %)
CO 7 Implement microbiological analysis of a soil ecosystem with an understanding of the most appropriate technique
Apply basic principles of environmental microbiology for understar and solving environmental problems –bioremediation



Course Code	Unit	Course/ Unit Title	Credits/ Lectures				
RUSMI C 303		ENVIRONMENTAL MICROBIOLOGY					
I		Air & Fresh Water Microbiology	15				
	1.1	Air Microbiology	05				
		a) Origin, distribution, number and kinds of microorganisms in air, Factors affecting microbial survival in air					
		<ul> <li>b) Enumeration of microorganisms in air: Impingement in liquids, Impaction on solids, Filtration, Sedimentation, Centrifugation, Electrostatic Precipitation.</li> </ul>					
		c) Air borne pathogens and diseases, droplets and droplet nuclei					
		d) Air sanitation- methods and application					
	1.2	Fresh water microbiology	10				
		a) General: Groups of natural waters, factors affecting kinds of microorganisms found in aquatic environments and nutrient cycles in aquatic environments					
		<ul> <li>b) Fresh Water environments and microorganisms found in Lakes, ponds, rivers, marshes, bogs and springs</li> </ul>					
	R	c) Potable water: Definition, water purification and pathogens transmitted through water.					
	71,	d) Microorganisms as indicators of water quality					
2AM		e) Bacteriological examination of water-sampling, routine analysis, SPC, membrane filter technique, Standards for water quality					
II		Marine and Sewage Microbiology	15				
	2.1	Marine Microbiology	05				
		a) Characteristics of marine environments					
		b) Diversity& characteristics of marine					



		microorganisms and their importance	
		<ul> <li>c) Ecosystems of Deep-sea Hydrothermal vents and Subterranean Water</li> </ul>	
	2.2	Sewage Microbiology	10
		a) Types of waste water	
		b) Characteristics of waste water	C
		<ul> <li>c) Modern waste water treatment: Primary, Secondary and tertiary treatment (oxidation ponds, activated sludge, trickling filters, anaerobic digestor).</li> </ul>	
		d) Removal of pathogens by sewage treatment Processes	
		e) Sludge Processing	
		<ul> <li>f) Disposal of Solid Waste, Modern Sanitary Landfills, Composting</li> </ul>	
III		Soil & Geo Microbiology	15
	3.1	Soil Microbiology	03
		<ul> <li>a) Soil – Definition, composition, function, Textural Triangle</li> </ul>	
		b) Types of Soil microorganisms & their activities	
	3.2	Methods of studying soil microorganisms	05
		a) Sampling	
		b) Cultural methods	
		c) Physiological methods	
	2	<ul><li>d) Immunological methods (Tabulation of the immunological methods)</li><li>e) NA based method</li></ul>	
		f) Radioisotope technique	
	3.3	Geo Microbiology	03
	5.5	a) Carbon cycle	- 03
Q1		b) Nitrogen cycle	
		c) Sulphur cycle	
		d) Phosphorus cycle	



3.4	Biodegradation and Bioremediation	4
	a) Microbial leaching	
	b) Metal transformations	
	c) Petroleum degradation	
	d) Degradation of xenobiotics	
		/.

- a) Raina M. Maier, Ian L. Pepper, Charles P. Gerba, Environmental Microbiology, 2nd Edition, 2010, Academic Press
- b) A.J. Salle, Fundamental Principles of Bacteriology, 7th Editon,1974, Tata McGraw Hill Publishing Company
- c) Air Quality Standards NAAQS Manual, Volume I, 2011
- d) Joanne M. Willey, Linda M. Sherwood, Christopher J.Woolverton Prescott's Microbiology, 8th Edition, 2011, McGraw Hill International Edition
- e) Frobisher, Hinsdill, Crabtree, Goodheart, Fundamentals of Microbiology, 9th Edition, 1974, Saunders College Publishing
- f) Barbara Kolwzan, Waldemar Adamiak (E Book) Oficyna Wydawnicza Politechniki Wroclawskiej, Wroclaw, 2006
- g) N.S Subba Rao, Introduction to Environmental Microbiology –Soil Microbiology -4th Edition ,2000, Oxford and IBH Publishing Co. Pvt Ltd
- h) Michael T. Madigan & J.M. Martin, Brock's Biology of Microorganisms 13th Ed. International edition 2012, Pearson Prentice Hall.



Course code	PRACTICALS	3 CREDITS
RUSMICP301	PRACTICAL-1	CKEDIIS
	Microbial Taxonomy and Introduction to Genetics	
	and Molecular Biology.	
	Isolation and identification of a bacterial isolate	<del>, (,)</del>
	Problems on Mendelian genetics	
	Extraction of DNA from onion and <i>E. coli</i>	
	Problems on genetic code	r
	Construction of phylogenetic tree.	
RUSMICP302	PRACTICAL -2	
	Introduction to Experimental Microbial	
	Biochemistry	
	Introduction to experimental design	
	2. Lab common sense workshop	
	3. Biostatistics problems	
	4. Study of pH meter and preparation of buffers	
	5. Density gradient centrifugation	
	6. Demonstration of agarose gel electrophoresis	
	7. Demonstration of PAGE	
	8. Separation of amino acids using paper	
	chromatography	
- 0	9. Separation of carbohydrates using TLC	
67	10. Demonstration of column chromatography	
	11. Demonstration of HPLC, HPTLC and GC	
	12. Determination of λmax	
	13. Verification of Beer's law and determination of	
X	extinction coefficient	
	14. Large scale cultivation of bacteria /yeast/ fungi	
	15. Determination of Dry and wet Weight	
	16. Disintegration of cells using physical & chemical	
	methods and separation of biomolecules	



	17.	Estimation of Amino acids by Ninhydrin method	
	18.	Estimation of Proteins by Biuret method	
	19.	Bradford's Method for protein estimation	
	20.	Estimation of Reducing Sugars by DNSA method	
	21.	Estimation of RNA by orcinol method	/,
	22.	Estimation of DNA by diphenylamine method	
	Not	e: All the above methods will also be analyzed using	
		statistical methods covered in theory	
RUSMICP303		PRACTICAL-3	
		Environmental Microbiology	
	1.	Enumeration of microorganisms in air and study its	
		load after fumigation	
	2.	Determination of microbial load using air impinger	
	3.	Study of halophilic and haloduric bacteria from	
		marine samples	
	4.	Routine analysis of water	
	5.	Use of membrane filter technique for bacteriological	
		analysis of water	
	6.	Rapid detection of E.coli by MUG technique-Demo	
	7.	Visit to Sewage treatment plant	
	8.	BOD of untreated and treated sewage	
	9.	Buried slide technique to study soil flora	
	10	. Mapping of soil flora- building phylogenetic trees	
	11	. Enrichment and isolation of Cellulose degraders,	
		Sulphate reducers and Phosphate solubilizers from	
		soil	
	12	. Setting up Winogradsky's Column	
	13	. Developing compost pits	
PII.			



# **Modality of Assessment**

## **Theory Examination Pattern:**

## A) Internal Assessment- 40%- 40 Marks per paper

Sr No	Evaluation type	Marks
1	One Assignment/Case study/Project/ Presentation	15
2	One class Test (multiple choice questions / objective)	20
3	Active participation in routine class instructional deliveries	05
	TOTAL	40

## B) External Examination- 60%- 60 Marks per paper

## **Semester End Theory Examination:**

- 1. Duration These examinations shall be of **two hours** duration.
- 2. Theory question paper pattern:
  - a. There shall be three questions each of 20 marks one on each unit.
  - b. All questions shall be compulsory with internal choice within the questions.

#### Paper Pattern:

Question	Options	Marks	Questions Based on
Q.1) A)	Any 3 out of 5	15	11.21
Q.1) B)	Any 1 set out of 2 (i & ii or i & ii)	03 &02	Unit I
Q.2) A)	Any 3 out of 5	15	I Ind II
Q.2) B)	Any 1 set out of 2 (i & ii or i & ii)	03 &02	Unit II
Q.3) A)	Any 3 out of 5	15	1154111
Q.3) B)	Any 1 set out of 2 (i & ii or i & ii)	03 &02	Unit III
	TOTAL	60	



#### **Practical Examination Pattern:**

A) Internal Examination: 40%- 60 Marks

Particulars	Paper I	Paper II	Paper III
Journal	05	05	05
Experimental tasks	10	10	10
Participation	05	05	05
Total	20	20	20

## B) External Examination: 60%- 90 Marks

## **Semester End Practical Examination:**

Particulars	Paper I	Paper II	Paper III
Laboratory work	25	25	25
Spots/Quiz/Viva	05	05	05
Total	30	30	30

## PRACTICAL BOOK/JOURNAL

The students are required to present a duly certified journal for appearing at the practical examination, failing which they will not be allowed to appear for the examination.

In case of loss of Journal and/ or Report, a Lost Certificate should be obtained from Head/ Co-ordinator / Incharge of the department; failing which the student will not be allowed to appear for the practical examination.

## **Overall Examination & Marks Distribution Pattern**

#### Semester III

Course	3	301		3	302		3	03		Grand Total
	Internal	External	Total	Internal	External	Total	Internal	External	Total	
Theory	40	60	100	40	60	100	40	60	100	300
Practicals	20	30	50	20	30	50	20	30	50	150



# **Course Title: Microbe Interactions and Host Responses**

CO 2 Evaluate the ecological, medical and evolutionary significance microbial interactions with plants, animals and other micro organism  CO 3 Outline the strategies through which pathogens develop infection and demonstrate presence of some virulence factors in known isolates  CO 4 Understand the concepts and terminologies used in epidemiology and correlate disease transmission to disease control  CO 5 Apply the understanding of epidemiology studies in solving publications.	COURSE	DESCRIPTION
CO 2 Evaluate the ecological, medical and evolutionary significance microbial interactions with plants, animals and other micro organism  CO 3 Outline the strategies through which pathogens develop infection and demonstrate presence of some virulence factors in known isolates  CO 4 Understand the concepts and terminologies used in epidemiology and correlate disease transmission to disease control  CO 5 Apply the understanding of epidemiology studies in solving publications.	OUTCOME	
CO 2 Evaluate the ecological, medical and evolutionary significance microbial interactions with plants, animals and other micro organism  CO 3 Outline the strategies through which pathogens develop infection and demonstrate presence of some virulence factors in known isolates  CO 4 Understand the concepts and terminologies used in epidemiology at correlate disease transmission to disease control  CO 5 Apply the understanding of epidemiology studies in solving publications.	CO 1	Exemplify microbial interactions with plants, animals and other
CO 3  Outline the strategies through which pathogens develop infection and demonstrate presence of some virulence factors in known isolates  CO 4  Understand the concepts and terminologies used in epidemiology and correlate disease transmission to disease control  Apply the understanding of epidemiology studies in solving put health concerns		microorganisms
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and demonstrate presence of some virulence factors in known isolates  CO 4  Understand the concepts and terminologies used in epidemiology at correlate disease transmission to disease control  CO 5  Apply the understanding of epidemiology studies in solving publications.		microbial interactions with plants, animals and other micro organisms
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correlate disease transmission to disease control  CO 5  Apply the understanding of epidemiology studies in solving publication health concerns		isolates
correlate disease transmission to disease control  CO 5  Apply the understanding of epidemiology studies in solving publication health concerns	_	
CO 5 Apply the understanding of epidemiology studies in solving publication health concerns	CO 4	Understand the concepts and terminologies used in epidemiology and
health concerns		correlate disease transmission to disease control
	CO 5	Apply the understanding of epidemiology studies in solving public
		health concerns
CO6 Understand the key components of innate and acquired immu	CO6	Understand the key components of innate and acquired immune
system and summarize their role in overcoming disease		system and summarize their role in overcoming disease
CO7 Compare the different types of immunoglobulins and understand the	CO7	Compare the different types of immunoglobulins and understand their
function in protection		function in protection



Course Code/ Unit	Unit	Course/ Unit Title	Credits/ Lectures
RUSMIC		MICROBE INTERACTIONS AND HOST	2/45
401		RESPONSES	, C) <sup>v</sup>
I		Microbial interactions with plants, animals and	15
		other microbes	
	1.1	Microbial associations with plants	08
1.2		<ul> <li>a) Phyllosphere</li> <li>b) Rhizosphere &amp; Rhizoplane</li> <li>c) Mycorrhizae</li> <li>d) Nitrogen fixation: Biochemistry of nitrogen fixation, nodulation in <i>Rhizobia</i>, <i>Azolla-Anabena</i> symbiosis, Actinorhizae, Stem nodulating <i>Rhizobia</i></li> <li>e) Fungal &amp; Bacterial endophytes</li> <li>f) Plant pathogens -Fungal, bacterial and viral diseases</li> <li>Microbial interactions with animals</li> <li>a) Zoo xanthallae, Zoo chlorellae- invertebrates</li> <li>b) Bacterial flora in the Rumen</li> <li>c) Microbe- insect interactions</li> <li>d) Introduction to Zoonotic diseases</li> </ul>	05
	1.3	Microbe -Microbe interactions	02
	R	<ul><li>a) Lichen</li><li>b) Endosymbionts of Protozoa</li><li>c) Parasitism in microbes</li></ul>	
II		Microbial invasion in Human hosts	15
	2.1	Development of an infection	08
Q.P.		<ul> <li>a) Bacterial virulence factors <ol> <li>Adherence factors</li> <li>Invasion of host cells and tissues</li> <li>Toxins- Exotoxins and Endotoxins</li> <li>Enzymes</li> <li>Evading host defense- Antigenic variation, Antiphagocytic factors and Intracellular pathogenicity </li> </ol></li></ul>	



		vi. Iron sequestration	
		vii. The role of Biofilms	
		<ul> <li>b) Measuring bacterial virulence: Infective dose &amp; Lethal dose, limulus amoebocyte assay</li> </ul>	
	c) Pathogenic properties of viruses, fungi and		
		protozoa	
	2.2	Introduction to epidemiological concepts	07
		a) Reservoirs of infection	
		<ul><li>b) Modes of disease transmission</li><li>c) Epidemiological terminology: epidemic, endemic,</li></ul>	
		pandemic, sporadic, incidence rate, prevalence	<b>Y</b>
		rate, mortality, morbidity	
		d) Controlling epidemics: Controlling reservoirs,	
		controlling transmission- Immunization strategies-	
		passive and active, Surveillance	
III		Host responses to infection	15
	3.1	Cells, Tissues and Organs of the Immune System	04
		a) Cells of the immune system- Lymphoid and Myeloid	
		cells, NK cells	
		b) Organs of the immune system- Introduction to	
		primary and secondary lymphoid organs and their roles	
	3.2	Immune responses- Innate defense mechanisms	04
		a) Phagocytosis – Recognition, Destruction,	
		b) Inflammation- Acute and Chronic	
		c) Fever	
	2 2	d) Molecular defenses- IFN, complement, ACP	07
	3.3	Immune responses- Acquired Defense  a) Outline and characteristics of Adaptive Immune	U/
		response	
16		b) Immunoglobulins – basic and fine structure	
		c) Immunoglobulin classes and biological activities	
		d) Antigenic determinants on immunoglobulins –	
(A)		isotypes, allotypes, idiotypes	
*		<ul> <li>e) Protective functions of antibodies- Opsonization,</li> <li>Complement mediated lysis, viral neutralization and</li> </ul>	
		toxin neutralization	
		f) Introduction to Cell mediated immunity	
		•	



- a) Willey, Sherwood and Woolverton, Prescott's Microbiology, 9th edition, 2013, International edition, McGraw Hill.
- b) Michael T. Madigan & J.M. Martin, Brock's Biology of Microorganisms 13th Ed. International edition 2012, Pearson Prentice Hall.
- c) Stanier, General microbiology 5th edition ,1987, Macmillan publication
- d) Tortora, Funke and Case, Microbiology: An Introduction, 10th Edition, 2010, Pearson.
- e) Kathleen Park Talaro & Arthur Talaro Foundations in Microbiology International edition 2002, McGraw Hill.
- f) Jacquelyn Black, Laura Black, Microbiology, Principles and Explorations, 9th Ed, 2015,
   Wiley
- g) Brooks, Carroll, et al, Jawetz, Melnick & Adelberg's Medical Microbiology, 26th Ed McGraw Hill Lange 2013
- h) https://www.eurofins.com.au/biopharma-services/testing-solutions/sterile-products-testing/endotoxin-or-lal-test/
- i) Ingraham and Ingraham, Introduction to Microbiology, by 2nd Ed ,2000, Brooks/Cole
- j) Thomas J. Kindt, Barbara A. Osborne, Richard A. Goldsby, Kuby Immunology, 6th ed, W. H. Freeman & Company 2005



# Course Title: INTRODUCTION TO METABOLIC PATHWAYS AND ENZYMOLOGY

# Academic year 2020-21

COURSE	DESCRIPTION
OUTCOME	
CO 1	Understand the concepts and types of metabolism. Compare the
	metabolic strategies & recall the role of Omics in biochemical studies
CO 2	Explain the regulatory junctions of metabolic pathways.
CO 3	Recall the properties & classes of enzymes. Illustrate Enzyme-
	substrate interaction models & recognize the significance of cofactors
	& coenzymes.
CO 4	Evaluate enzyme kinetics & the change in activity in presence of
	variables.
CO 5	Explain the principles of Bioenergetics & attribute the role of energy
	currency molecule
CO 6	Understand & apply the laws of thermodynamics to microbial
	metabolism.
CO 7	Implement experimental procedures for enzyme purification and
	enzyme kinetics studies



Course Code	Unit	Course/ Unit Title	Credits/ Lectures	
RUSMI		INTRODUCTION TO METABOLIC	2/45	
C 402		PATHWAYS AND ENZYMOLOGY		
		Introduction to Metabolism	15	
	1.1	Introduction to biochemical reactions:	04	
		a) Central role of chemical reactions in life		
		b) Characteristics of biochemical reactions		
	1.2	Introduction to metabolism:	06	
		a) Metabolism- Catabolism & Anabolism		
		b) Types of Metabolic pathways		
		c) Metabolic networks, use of different software		
		d) Primary and secondary metabolism		
		e) C- skeleton, Energy and reducing power		
		requirements		
	1.3	Metabolic strategies: Managing metabolic network	04	
		a) Role of enzymes, enzyme clustering &		
		multienzyme complexes		
		b) Functional coupling		
		c) Compartmentalization in cells		
	1.4	Introduction to omics: Metabolome & Metabolomics	01	
II		Enzymology	15	
	2.1	Introduction to enzymes:	06	
		a) General properties of enzymes		
	07	b) How do enzymes accelerate reactions?		
		c) Classification of enzymes		
		d) Enzyme kinetics: Rate law for a simple catalyzed		
		reaction, Michaelis-Menten equation and its		
		derivation, other plots to determine velocity of		
01		reactions		
	2.2	Modifying enzyme catalysis rates	05	
		a) Effect of temperature and pH		
		b) Effect of Inhibitors- Reversible and irreversible,		
		competitive, Non-competitive and uncompetitive		
		inhibitors		



a) Different types and reactions catalyzed by coenzymes (in tabular form) b) Water soluble coenzymes (NAD, Nicotinic acid) c) Fat soluble vitamins and their examples. d) Inorganic cofactors	04
coenzymes (in tabular form) b) Water soluble coenzymes (NAD, Nicotinic acid) c) Fat soluble vitamins and their examples.	
<ul><li>b) Water soluble coenzymes (NAD, Nicotinic acid)</li><li>c) Fat soluble vitamins and their examples.</li></ul>	
c) Fat soluble vitamins and their examples.	
d) Inorganic cofactors	
Principles of Bioenergetics	1:
Bioenergetics & thermodynamics:	0(
a) Energy transformations	
b) Thermodynamic quantities, standard –free energy	
c) Difference between ΔG & ΔGo"	
2 ATP and it's role	0
a) Structure of ATP, phosphoryl group transfer and	
ATP	
donor	
Biological oxidation-reduction reactions	04
	<ul> <li>a) Energy transformations</li> <li>b) Thermodynamic quantities, standard –free energy</li> <li>c) Difference between ΔG &amp; ΔGo"</li> </ul> 2.2 ATP and it's role <ul> <li>a) Structure of ATP,phosphoryl group transfer and ATP</li> <li>b) Types of energy –rich compounds</li> <li>c) Multi-roles of ATP inorganic phosphoryl group donor</li> </ul>



- a) Principles of Biochemistry by Geoffery Zubay (1988) 4th Edition Wm.C. Brown Publishers
- b) Outlines Of Biochemistry,5/E,Conn P.Stumpf,G.Bruening & R.Doi,John Wiley & Sons,New York 1995
- c) Fundamentals of Enzymology: Cell and Molecular Biology of Catalytic Proteins 3rd Edition Nicholas Price and Lewis Stevens
- d) Lehninger: Principles Of Biochemistry,4th Ed., D. Nelson & M. Cox, W.H.Freeman & Co., (LPE)
- e) A biologist"s Physical Chemistry by John Gareth Morris.
- f) Rodney Boyer
- g) Stanier, General microbiology 5th edition ,1987, Macmillan publication
- h) Principles of Biochemistry by Robert Horton (2011) 5<sup>th</sup> Edition Pearson Publishers.



**Course Title: APPLIED MICROBIOLOGY** 

# Academic year 2020-21

COURSE OUTCOME	DESCRIPTION		
CO 1	Understand and explain the significance of microbes in fermentation industry and compare the techniques used for their screening		
CO 2	Compare different types of fermentations and fermentation processes used for industrial productions		
CO 3	Exemplify components used in industrial fermentation media with an understanding of its role in the process		
CO 4	Summarize the general principles of food spoilage by microorganisms and compare methods used for food preservation		
CO 5	Execute experimental procedures for detection of microbes in food and dairy products and comment on its quality		
CO 6	Recall the sources of microorganisms in milk and explain the significance of pasteurization techniques		
CO7	Outline and analyze the manufacturing processes of different fermented dairy products		
CO 8	Apply knowledge of contamination, preservation, and quality control in food and dairy product manufacturing industries		



Course Code	Unit	Course/ Unit Title	Credits/ Lectures 2/45	
RUSMIC		APPLIED MICROBIOLOGY		
403				
I		Industrial Microbiology	15	
	1.1	Strains of industrially important microorganisms	04	
		a) Desirable characteristics of an industrial strain     b) Principles and methods of primary and secondary screening		
	1.2	Types of fermentations:	02	
		a) Aerobic b) Anaerobic c) Solid state fermentation		
	1.3	Types of fermentation processes:	02	
		a) Surface and Submerged     b) Batch, continuous, fed-batch fermentation process		
	1.4	Media for industrial fermentations	05	
		a) Production and Inoculum media b) Media components: - Carbon source, nitrogen source, amino acids and vitamins, minerals, water, buffers, antifoam agents, precursors, inhibitors and inducers		
,	1.5	Inoculum development	02	
II		Food Microbiology	15	
	2.1	Introduction:	01	
SVIII		Significance, food as a substrate and sources of microorganisms in food		
	2.2	Intrinsic and extrinsic factors affecting the microbial growth in food	02	
	2.3	General Principles of spoilage	04	
		Spoilage of fresh foods: fruits and vegetables, eggs, meat, poultry and seafood		



	2.4	General principles of food preservation	04
		(principle of each method and example of foods only)	
		High temperature, low temperature, drying, radiations	
		and food additives and preservatives (tabular	
		representation), Asepsis, introduction to HACCP,	
		Regulation	
	2.5	Food borne diseases	1)
	2.6	Methods of detection of microorganisms in food:	3
		Overview of cultural, microscopic, physical, chemical	
		and bioassay methods	
III		Dairy Microbiology	15
	3.1	Milk- Definition, composition, sources of	2
		contamination of milk	_
	3.2	Pasteurization of milk	3
		LTHT, HTST, UHT	
	3.3	Milk products: production and spoilage of:	7
		a) Yoghurt	
		b) Butter	
		c) Cheese-Cheddar and Cottage cheese	
		d) Fermented milks	
	3.4	Quality control of milk	3
		a) Rapid platform tests	
		b) Microbiological analysis of milk : SPC, Coliform	
		count, LPC, Psychrophiles, Thermophilic count,	
		DRT	
		1	

- a) Fundamental Food Microbiology by Bibek Ray, Arun Bhunia (2007), 4th edition CRC Press
- b) Food Microbiology by Frazier 5th ed (1971), McGraw-Hill Education.
- c) Modern Food Microbiology by James Jay 6th ed(2000), Springer US.
- d) Applied Dairy Microbiology by Marth & Steele(2001), CRC Press
- e) BIS standards, FSSAI



- f) Casida L. E., "Industrial Microbiology" 2009 Reprint, New Age International (P) Ltd, Publishers, New Delhi
- g) Stanbury P. F., Whitaker A. & Hall--S. J., 1997, "Principles of Fermentation, Technology", 2nd Edition, Aditya Books Pvt. Ltd, New Delhi.
- h) Prescott and Dunn's "Industrial Microbiology" 1982 4th Edition, McMillan Publishers
- , india.

  RANNARAIN RUIA ARITO NON PROPERTIES COLLIES i) H. A. Modi, 2009. ""Fermentation Technology"" Vol 2, Pointer Publications, India.



Course code	PRACTICALS	3 Credits
RUSMIC	SECTION-1	
P401	MICROBE INTERACTIONS AND HOST RESPONSES	
	1. Isolation of Rhizobium from root nodules	60
	2. Demonstration of fungi and algae in lichens	
	3. Isolation of Xanthomonas from spoilt citrus fruit	
	Study of virulence factors – Enzymes – Streptokinase, Coagulase, Hemolysin, Lecithinase	
	5. Demonstration of biofilm formation by pathogens on catheters	
	6. Assignment on classical stages, signs and symptoms of any one	
	microbial disease	
	7. Staining of blood film to demonstrate different types of leucocytes	
	8. Phagocytosis (Demonstration)	
	9. Study of plant microbe interactions: Screening for Auxin production (PGP from Rhizosphere)	
	10. Case studies and problems on Epidemiology	
	11. How to develop epidemiological surveys	
RUSMIC	SECTION-2	
P402	INTRODUCTION TO METABOLIC PATHWAYS AND ENZYMOLOGY	
	1. Using KEGG, Ecocyc, metacyc, biocyc and Brenda for	
	understanding metabolic networking	
67	2. Qualitative detection of	
	a. Amylase	
	b. Lipase	
	c. Protease	
	d. DNase	



		T
	e. Catalase	
	f. Oxidase	
	g. Carbohydrate fermentation	
	h. Dehydrogenase	
	3. Production and purification of an enzyme	4
	4. Assay of an enzyme and determination of enzyme units	1,0,
	5. Determination of km and Vmax of an enzyme	
	6. Effect of environment on enzyme activity:	
	a. Effect of temperature	
	b. Effect of pH	
	c. Effect of enzyme concentration	
	7. Effect of inhibitors	
RUSMIC	SECTION-3	
P403	APPLIED MICROBIOLOGY	
	1. Isolation of antibiotic producers from soil- Wilkin's overlay method.	
	2. Determination of microbial counts in food using dip slide technique	
	(demonstration)	
	3. Isolation of food spoilage agent	
	4. Determination of TDT and TDP	
	5. Determination of Salt and sugar tolerance	
	6. Determination of MIC of a preservative	
	7. Visit to Food/Dairy industry	
	8. Rapid platform tests of raw and pasteurized milk.	
67	9.Microbiological analysis of raw and pasteurized Milk.	
	10. Microbiological analysis of Butter, Cheese.	
	11. Surface and submerged fermentation.	
	12. Testing a packaged meat product for its microbial load.	



# **Modality of Assessment**

## **Theory Examination Pattern:**

## A) Internal Assessment- 40%- 40 Marks per paper

Sr No	Evaluation type	Marks
1	One Assignment/Case study/Project/ Presentation	15
2	One class Test (multiple choice questions / objective)	20
3	Active participation in routine class instructional deliveries	05
	TOTAL	40

## B) External Examination- 60%- 60 Marks per paper

## **Semester End Theory Examination:**

- 1. Duration These examinations shall be of **two hours** duration.
- 2. Theory question paper pattern:
  - a. There shall be three questions each of 20 marks one on each unit.
  - All questions shall be compulsory with internal choice within the questions.

#### Paper Pattern:

Question	Options	Marks	Questions Based on	
Q.1) A)	Any 3 out of 5	15	11.21	
Q.1) B)	Any 1 set out of 2 (i & ii or i & ii)	03 &02	Unit I	
Q.2) A)	Any 3 out of 5	15		
Q.2) B)	Any 1 set out of 2 (i & ii or i & ii)	03 &02	Unit II	
Q.3) A)	Any 3 out of 5	15	11	
Q.3) B)	Any 1 set out of 2 (i & ii or i & ii)	03 &02	Unit III	
	TOTAL	60		



#### **Practical Examination Pattern:**

A) Internal Examination: 40%- 60 Marks

Particulars	Paper I	Paper II	Paper III
Journal	05	05	05
Experimental tasks	10	10	10
Participation	05	05	05
Total	20	20	20

## B) External Examination: 60%- 90 Marks

## **Semester End Practical Examination:**

Particulars	Paper I	Paper II	Paper III
Laboratory work	25	25	25
Spots/Quiz/Viva	05	05	05
Total	30	30	30

## PRACTICAL BOOK/JOURNAL

The students are required to present a duly certified journal for appearing at the practical examination, failing which they will not be allowed to appear for the examination.

In case of loss of Journal and/ or Report, a Lost Certificate should be obtained from Head/ Coordinator / Incharge of the department; failing which the student will not be allowed to appear for the practical examination.

## **Overall Examination & Marks Distribution Pattern**

#### **Semester IV**

Course	ourse 401		Course 401 402			403			Grand Total	
	Internal	External	Total	Internal	External	Total	Internal	External	Total	
Theory	40	60	100	40	60	100	40	60	100	300
Practicals	20	30	50	20	30	50	20	30	50	150



**Course Title: Microbial Genetics** 

# Academic year 2020-21

COURSE	DESCRIPTION
OUTCOME	
CO 1	Understand and differentiate between population and
	quantitative genetics and compare model organisms used in genetic
	studies.
CO 2	Summarize different natural plasmids and transposons
	present in prokaryotes and be able to compare and contrast
	between different plasmids.
CO 3	Understand the coherence of the molecular mechanisms
	involved in DNA replication and outline different enzymes and
	proteins associated with both prokaryotic and eukaryotic DNA
	replication
CO 4	Identify, interpret and classify mutations in DNA followed by
	mechanism of DNA repair
CO 5	Test the effect of mutagens on bacteria and identify mutants
CO 6	Solve and interpret problems based on mapping of bacterial
	genes using transformation, transduction and conjugation
CO 7	Retrieving basic concepts of homologous recombination and genetic
	exchange among prokaryotes



Course Code	Unit	Course/ Unit Title	Credits/ Lectures
RUSMIC		MICROBIAL GENETICS	2.5/60
501			
I		Branches of Genetics, Plasmids, Transposons	15
	1.1	Overview of branches of Genetics	04
		a) Transmission, Molecular,	
		b) Population Genetics: Hardy-Weinberg Law-	
		principle and violation of assumptions (Mutation,	
		Migration, Genetic Drift, Natural Selection)	
		c) Quantitative Genetics: Characteristics, concept of	
		Heritability, QTLs, Response to selection	
	1.2	Model Organisms	03
	1.2	a) Characteristics of a model organism	03
		b) Examples of select model organisms used in	
		study: <i>E.coli</i> , Yeast, Mouse, <i>Caenorhabditis</i>	
		elegans, Arabidopsis thaliana	
	1.3	Plasmids	04
		a) Physical nature	
		b) Detection and isolation of plasmids	
		c) Plasmid incompatibility and Plasmid curing	
		d) Cell to cell transfer of plasmids	
		e) Types of plasmids	
		i. Resistance Plasmids	
	7	ii. Plasmids encoding Toxins and other Virulence characteristics	
	0	iii. col factor	
		iv. Degradative plasmids	
		iv. Degradative plasmids	
	1.4	Transposable elements in Prokaryotes	04
		a) Insertion sequences	
$Q_{ij}$		b) Transposons	
		i. Types	
		ii. Structure and properties	
		iii. Mechanism of transposition	
		iv. Transposon mutagenesis	
		v. Integrons	



II	DNA Replication	15
2.1	Historical perspective	04
	a) Conservative	
	b) Dispersive	
	c) Semi-conservative	
	d) Bidirectional	
	e) Semi-discontinuous DNA replication	
2.2	Prokaryotic DNA replication	`04
	Details of molecular mechanism involved in Initiation,	
	Elongation and Termination	
2.3	Enzymes and proteins associated with DNA replication	04
	<ul> <li>a) Primase</li> <li>b) Helicase</li> <li>c) Topoisomerase</li> <li>d) SSB</li> <li>e) DNA polymerases</li> <li>f) Ligases</li> </ul>	
	g) Ter and Tus proteins	
2.4	Eukaryotic DNA replication	02
	<ul><li>a) Molecular details of DNA synthesis</li><li>b) Replicating the ends of the chromosomes</li></ul>	
2.5	Rolling circle mode of replication	01
III	Mutation and Repair	15
3.1	Mutation	10
RAMINAR!	<ul> <li>a) Terminology: alleles, homozygous, heterozygous, genotype, phenotype, Somatic mutation, Germline mutation, Gene mutation, Chromosome mutation, phenotypic lag, hotspots and mutator genes</li> <li>b) Fluctuation test.</li> <li>c) Types of mutations: Point mutation, reverse mutation, suppressor mutation, frameshift mutation, conditional lethal mutation, base pair substitution, transition, transversion, missense mutation, nonsense mutation, silent mutation, neutral mutation, pleiotropic mutations.</li> <li>d) Causes of mutation: Natural/spontaneous mutationreplication error, depurination, deamination. Induced mutation: principle and mechanism with illustrative diagrams for – <ol> <li>i. Chemical mutagens- base analogues, nitrous acid, hydroxyl amine, intercalating agents and alkylating agents.</li> <li>ii. Physical mutagen</li> </ol> </li> </ul>	



		iii. Biological mutagen (only examples)	
		e) Ames test	
		f) Detection of mutants	
	3.2	DNA Repair	05
		a) Mismatch repair	
		b) Light repair	_
		c) Repair of alkylation damage	
		d) Base excision repair	$C_{\sim}$
		e) Nucleotide excision repair f) SOS repair	
IV		f) SOS repair  Genetic Exchange	15
	4.1	Gene transfer mechanisms in bacteria & homologous	10
		recombination	
		a) Transformation	04
		i. Introduction and History	
		ii. Types of transformation in prokaryotes—Natural	
		transformation in Streptococcus pneumoniae,	
		Hemophilus influenzae and Bacillus subtilis	
		iii. Mapping of bacterial genes using transformation	
		iv. Problems based on transformation.	
		b) Conjugation	05
		i. Discovery of conjugation in bacteria	03
		ii. Properties of F plasmid/Sex factor	
		iii. The conjugation machinery	
		iv. Hfr strains, their formation and mechanism of	
		conjugation	
		v. F' factor, origin and behavior of F' strains,	
		Sexduction.	
		vi. Mapping of bacterial genes using conjugation	
		(Wolman and Jacob experiment).	
		vii. Problems based on conjugation c) Transduction	03
		i. Introduction and discovery	03
	7	ii. Generalized transduction	
		iii. Use of Generalized transduction for mapping	
	K	genes	
		iv. Specialized transduction	
		v. Problems based on transduction	
	4.0	December 1 and a land of	00
	4.2	Recombination in bacteria	03
(L)		a) General/Homologous recombination	
		i. Molecular mechanism	
		ii. Holliday model of recombination	
		b) Site –specific recombination	



- a) Peter J. Russell, "Genetics-A molecular approach", 2nd edition, 2006.
- b) Benjamin A. Pierce, "Genetics a conceptual approachl", 3rdedition, 2008, W. H. Freeman and company.
- c) R. H. Tamarin, "Principles of genetics",2004, Tata McGraw Hill.
- d) D, Nelson and M. Cox, "Lehninger's Principles of biochemistry",4thedition,2005, Macmillan worth Publishers.
- e) M.Madigan, J. Martinko, J.Parkar, "Brock Biology of microorganisms", 12th edition, 2009, Pearson Education International.
- f) Fairbanks and Anderson, "Genetics", 1999, Wadsworth Publishing Company.
- g) Willey, Sherwood and Woolverton, Prescott's Microbiology, 7th edition, 2013, International edition, McGraw Hill.
- h) Robert Weaver, "Molecular biology", 3rd edition, McGraw Hill international edition.
- i) Nancy Trun and Janine Trempy, "Fundamental bacterial genetics", 2004, Blackwell Publishing.
- j) Snustad, Simmons, "Principles of genetics", 3rd edition, John Wiley & sons, Inc.
- k) Stanier, Ingraham, "General Microbiology",5th edition, Macmillan
- I) Benjamin Lewin, "Genes IX", Jones and Bartlett publishers.
- m) JD Watson, Bake, Bell, Gann, Levine, Losick, "Molecular biology of the gene", 5th edition, Person



**Course Title: Medical Microbiology** 

COURSE OUTCOME	DESCRIPTION
CO 1	Understand modern alternatives to Koch's postulates
CO 2	Summarize the basic aspects of clinical and diagnostic microbiology and implement bacteriological investigations using good laboratory practices
CO 3	Understand, interpret and explain the coherence between pathogenesis mechanisms of microorganisms, clinical manifestation of disease and prophylactic measures of representative bacterial, fungal and parasitic infections in various organ systems
CO 4	Extrapolate the understanding of representative infections of skin, respiratory system, urinary tract, gastro intestinal tract central nervous system to other infections within the same system
CO 5	Given a few key clinical features, design and execute lab diagnostic procedures for any given pathological specimen and test antibiotic susceptibility of the isolated pathogen
CO6	Differentiate between the different classes of antibiotics on the basis of their mechanism of action
C07	Attribute strategies through which microbes acquire anti-microbial resistance
CO8	Check and evaluate drugs/ antibiotics for their efficacy by demonstrating their action on microorganisms



Course Code	Unit	Course/ Unit Title	Credits/ Lectures
RUSMIC 502		MEDICAL MICROBIOLOGY	2.5/60
I		Study of Infectious diseases-I	15
	1.1	Associating Microbes to disease	02
		a) Koch's Postulate and modern alternatives to it b) Molecular Koch's postulates	
	1.2	Introduction to Clinical and diagnostic Microbiology	05
		a) Phases of diagnostic cycle- Pre analytic, analytic and post analytic     b) Introduction to Molecular and immunological methods	
	1.3	Study of Infectious Diseases-I (with Emphasis on Characteristics of the Aetiological Agent, Pathogenesis & clinical features, Laboratory Diagnosis and Prevention)	08
		Respiratory diseases:  a) Strep throat by <i>S. pyogenes</i> b) Diphtheria c) Common cold d) Tuberculosis e) Pneumonia caused by <i>K. pneumoniae</i>	
II	70	Study of Infectious Diseases II (With emphasis on cultural characteristics of the aetiological agent, pathogenesis, laboratory diagnosis and prevention)	15
	2.1	Study of skin infections	05
apple	P.	<ul> <li>a) Leprosy</li> <li>b) Pyogenic skin infections caused by <i>Pseudomonas</i>,</li> <li>S. pyogenes and S. aureus.</li> <li>c) Fungal infections- Oral Thrush, Dermatophytosis</li> </ul>	
	2.2	Study of gastrointestinal tract infections	08
		<ul> <li>a) Enteric fever- Salmonella</li> <li>b) Shigellosis</li> <li>c) Infections due to pathogenic E. coli strains</li> <li>d) Rotavirus diarrhoea</li> <li>e) Dysentery due to Entamoeba histolytica</li> </ul>	



	2.3	. Study of urinary tract infections	02
		a) Predisposing factors	
		b) List of causative agents	
		c) Pathogenesis and laboratory diagnosis	
III		Study of Infectious Diseases III	15
		(With emphasis on cultural characteristics of the	
		aetiological agent, pathogenesis, laboratory diagnosis and	, ( ^ Y
		prevention)	
	3.1	Study of vector-borne infections	03
		a) Rickettsial diseases (Tabular form), b) Malaria	
	3.2	Study of sexually transmitted infectious diseases	07
		a) Syphilis	
		b) AIDS	
		c) Gonorrhea	
	3.3	Study of central nervous system infectious diseases	05
	3.3	a) Tetanus	US
		b) Polio	
		c) Meningococcal meningitis	
		o, memigeocosii momigi	
IV		Chemotherapy of infectious agents	15
	4.1	Introduction to Chemotherapeutic agents	03
		a) Attributes of an ideal chemotherapeutic agent and	
		related definitions	
		b) Selection and testing of antibiotics for bacterial isolates by Kirby-Bauer method and other assays	
		(E-test & Checker Board Assay)	
		(E test & Officerer Board 7133dy)	
	4.2	Mode of action of antibiotics	08
	4.2	Mode of action of antibiotics  a) Cell wall (Beta-lactams- Penicillin and	08
	4.2		08
	4.2	a) Cell wall (Beta-lactams- Penicillin and Cephalosporins, Carbapenems)     b) Cell Membrane (Polymyxin and Imidazole)	08
	4.2	<ul> <li>a) Cell wall (Beta-lactams- Penicillin and Cephalosporins, Carbapenems)</li> <li>b) Cell Membrane (Polymyxin and Imidazole)</li> <li>c) Protein Synthesis (Aminoglycosides-Streptomycin,</li> </ul>	08
	4.2	a) Cell wall (Beta-lactams- Penicillin and Cephalosporins, Carbapenems)     b) Cell Membrane (Polymyxin and Imidazole)     c) Protein Synthesis (Aminoglycosides-Streptomycin, Macrolide (Tetracycline and Chloramphenicol)	08
	4.2	<ul> <li>a) Cell wall (Beta-lactams- Penicillin and Cephalosporins, Carbapenems)</li> <li>b) Cell Membrane (Polymyxin and Imidazole)</li> <li>c) Protein Synthesis (Aminoglycosides-Streptomycin, Macrolide (Tetracycline and Chloramphenicol)</li> <li>d) Nucleic acid (Quinolones, Nalidixic acid,</li> </ul>	08
	4.2	<ul> <li>a) Cell wall (Beta-lactams- Penicillin and Cephalosporins, Carbapenems)</li> <li>b) Cell Membrane (Polymyxin and Imidazole)</li> <li>c) Protein Synthesis (Aminoglycosides-Streptomycin, Macrolide (Tetracycline and Chloramphenicol)</li> <li>d) Nucleic acid (Quinolones, Nalidixic acid, Rifamycin)</li> </ul>	08
	4.2	<ul> <li>a) Cell wall (Beta-lactams- Penicillin and Cephalosporins, Carbapenems)</li> <li>b) Cell Membrane (Polymyxin and Imidazole)</li> <li>c) Protein Synthesis (Aminoglycosides-Streptomycin, Macrolide (Tetracycline and Chloramphenicol)</li> <li>d) Nucleic acid (Quinolones, Nalidixic acid,</li> </ul>	08
	R	<ul> <li>a) Cell wall (Beta-lactams- Penicillin and Cephalosporins, Carbapenems)</li> <li>b) Cell Membrane (Polymyxin and Imidazole)</li> <li>c) Protein Synthesis (Aminoglycosides-Streptomycin, Macrolide (Tetracycline and Chloramphenicol)</li> <li>d) Nucleic acid (Quinolones, Nalidixic acid, Rifamycin)</li> <li>e) Enzyme inhibitors (Sulfa drugs, Trimethoprim)</li> </ul>	
PANK	4.2	<ul> <li>a) Cell wall (Beta-lactams- Penicillin and Cephalosporins, Carbapenems)</li> <li>b) Cell Membrane (Polymyxin and Imidazole)</li> <li>c) Protein Synthesis (Aminoglycosides-Streptomycin, Macrolide (Tetracycline and Chloramphenicol)</li> <li>d) Nucleic acid (Quinolones, Nalidixic acid, Rifamycin)</li> </ul>	08
	R	a) Cell wall (Beta-lactams- Penicillin and Cephalosporins, Carbapenems) b) Cell Membrane (Polymyxin and Imidazole) c) Protein Synthesis (Aminoglycosides-Streptomycin, Macrolide (Tetracycline and Chloramphenicol) d) Nucleic acid (Quinolones, Nalidixic acid, Rifamycin) e) Enzyme inhibitors (Sulfa drugs, Trimethoprim)  List of common antibiotics	
	R	a) Cell wall (Beta-lactams- Penicillin and Cephalosporins, Carbapenems) b) Cell Membrane (Polymyxin and Imidazole) c) Protein Synthesis (Aminoglycosides-Streptomycin, Macrolide (Tetracycline and Chloramphenicol) d) Nucleic acid (Quinolones, Nalidixic acid, Rifamycin) e) Enzyme inhibitors (Sulfa drugs, Trimethoprim)  List of common antibiotics used for treating viral, fungal and parasitic diseases, New	



- a) Brenda Wilson, Abigail Salyer And Dixie Whitt, Bacterial Pathogenesis A molecular approach 3rdEd ASM press 2011
- b) Gary. W. Procop, Dierdre Church et al, Koneman's Color Atlas and Textbook of Diagnostic Microbiology, Seventh Ed, Walters Kluwer, 2017
- c) Willey, Sherwood and Woolverton, Prescott's Microbiology, 9th edition, 2013, International edition, McGraw Hill.
- d) Brooks, Carroll, et al, Jawetz, Melnick & Adelberg's Medical Microbiology, 26th Ed McGraw Hill Lange 2013
- e) Ananthanarayan and Panicker's, Textbook of Microbiology, 10th edition, Ed by Reba Jology, : Kanugo, Universities Press, 2017
  - f) Goering, Dockerel et al, Mim's Medical microbiology, 5th Ed 2013, Saunders



Course code	PRACTICALS	3 Credits
RUSMIC P 501	Practical Based on 501	
1 301	UV survival curve – determination of exposure time leading to 90% reduction	. (3)
	2. Isolation of mutants using UV mutagenesis	
	<ol> <li>Replica plate technique for selection &amp; characterization of mutants – auxotroph &amp; antibiotic resistant</li> </ol>	
	4. Isolation and detection of plasmid DNA.	
	5. Preparation of competent cells and transformation	
	Demonstration of conjugation.	
RUSMIC P 501	Practical Based on 502	
	<ol> <li>Assignment on sample collection, transport and processing of any one pathological sample</li> </ol>	
	2. Rapid detection of infection in samples from CNS	
	3. Rapid Direct tests for identification of pathogens-	
	a. Acid fast staining of <i>M. tuberculosis/ M.leprae</i> .	
	b. Metachromatic granule staining for <i>C.diphtheriae</i>	
	c. Catalase test	
	d. Bile solubility test	
	e. Slide coagulase test for <i>S.aureus</i>	
	f. Spot indole test	
	g. Oxidase test	
	h. Modern methods for identification of pathogens.	
	4. Identification of isolates obtained from following samples by	
	morphological, cultural and biochemical properties from-	
" M	a. Nasal/ throat swabs (URT infection)	
25	b. Sputum (LRT infection)	
	c. Skin swab/ pus (Skin infection)	
	d. Identification of <i>Candida</i> species using the germ tube test and growth on Chrom agar	
	e. Stool (GI tract infection)	



- f. Urine (UTI infection)
- 5. Demonstration of malarial parasite in blood film
- 6. Selection and testing of antibiotics using the Kirby-Bauer method
- 7. Determination of MIC of an antibiotic by E-test
- 8. Synergistic action of two drugs
- 9. Determination of MBC of an antibiotic.
- 10. Detection of βlactamase in S.aureus.
- 11. Role of plasmids in antibiotic resistance through curing of the PANNAR AIM RUIA AUTONO MONO DE PANNAR AIM RUIA AUTONO DE PANNAR AUTONO DE PANN plasmid



# **Course Title: Microbial Biochemistry Part I**

# Academic year 2020-21

COURSE OUTCOME	DESCRIPTION
OOTOOME	
CO 1	Understand the membrane architecture & critique the modes of solute
	transportation.
CO 2	Compare & contrast the mechanism of ATP synthesis in Prokaryotes
	& Eukaryotes.
CO 3	Summarize & differentiate the catabolic pathways of carbohydrates &
	deconstruct its amphibolic nature.
CO 4	Outline & evaluate the different fermentative pathways in bacteria.
CO 5	Paraphrase the anabolic pathways for carbohydrate synthesis.
CO 6	Organize the tally sheet of energetics for different catabolic substrates
	and solve problems based on these.
CO 7	Execute & evaluate the experimental aspects of metabolic reactions &
	differentiate organisms on the basis of their metabolic differences.



Biological Membranes & Transport   15	Course Code	Unit	Course/ Unit Title	Credits/ Lectures
1.1 Composition and architecture of membrane  a) Lipids b) Integral & peripheral proteins & interactions with lipids c) Permeability and outer membrane- a barrier d) Aquaporins e) Mechanosensitive channels  1.2 Methods of studying solute transport a) Using Whole cells b) Using Liposomes c) Using Proteoliposome  1.3 Solute transport across membrane proteins. b) Transporters grouped into Superfamilies' c) Co transport across plasma membrane (Uniport, Antiport, Symport) d) Active transport & electrochemical gradient e) Ion gradient provides energy for secondary Active transport e) Lactose transport f) ATPases and transport g) ABC transporters e.g. Histidine transport f) ATPases and transport g) ABC transporters e.g. Histidine transport h) Shock sensitive system — Role of binding proteins e.g. Maltose uptake i) Phosphotransferase system j) Schematic representation of various Membrane transport mechanisms in. E. coli  1.4 Other examples of solute transport c) Bacterial protein export c) Bacterial protein export c) Bacterial membrane fusion central to many biological processes			MICROBIAL BIOCHEMISTRY PART I	2.5/60
a) Lipids b) Integral & peripheral proteins & interactions with lipids c) Permeability and outer membrane- a barrier d) Aquaporins e) Mechanosensitive channels  1.2 Methods of studying solute transport a) Using Whole cells b) Using Liposomes c) Using Proteoliposome  1.3 Solute transport across membrane proteins. b) Transporters grouped into Superfamilies' (c) Co transport across plasma membrane (Uniport, Antiport, Symport) d) Active transport & electrochemical gradient e) Ion gradient provides energy for secondary Active transport e.g. Lactose transport f) ATPases and transport g) ABC transporters e.g. Histidine transport h) Shock sensitive system — Role of binding proteins e.g. Maltose uptake i) Phosphotransferase system j) Schematic representation of various Membrane transport mechanisms in. E. coli  1.4 Other examples of solute transport c) Bacterial protein export c) Bacterial protein export c) Bacterial membrane fusion central to many biological processes	ı		Biological Membranes & Transport	15
b) Integral & peripheral proteins & interactions with lipids c) Permeability and outer membrane- a barrier d) Aquaporins e) Mechanosensitive channels  1.2 Methods of studying solute transport a) Using whole cells b) Using Liposomes c) Using Proteoliposome  1.3 Solute transport across membrane a) Passive transport facilitated by membrane proteins. b) Transporters grouped into Superfamilies' ' c) Co transport across plasma membrane (Uniport, Antiport, Symport) d) Active transport & electrochemical gradient e) lon gradient provides energy for secondary Active transport e.g. Lactose transport f) ATPases and transport g) ABC transporters e.g. Histidine transport h) Shock sensitive system – Role of binding proteins e.g. Maltose uptake i) Phosphotransferase system j) Schematic representation of various Membrane transport mechanisms in. E. coli  1.4 Other examples of solute transport c) Bacterial protein export c) Bacterial protein export c) Bacterial membrane fusion central to many biological processes		1.1	Composition and architecture of membrane	02
a) Using whole cells b) Using Liposomes c) Using Proteoliposome  1.3 Solute transport across membrane a) Passive transport facilitated by membrane proteins. b) Transporters grouped into Superfamilies' (c) Co transport across plasma membrane (Uniport, Antiport, Symport) d) Active transport & electrochemical gradient e) lon gradient provides energy for secondary Active transport e.g. Lactose transport g) ABC transporters e.g. Histidine transport h) Shock sensitive system – Role of binding proteins e.g. Maltose uptake i) Phosphotransferase system j) Schematic representation of various Membrane transport mechanisms in. E. coli  1.4 Other examples of solute transport b) Bacterial protein export c) Bacterial membrane fusion central to many biological processes  II Bioenergetics and Bioluminescence			<ul> <li>b) Integral &amp; peripheral proteins &amp; interactions with lipids</li> <li>c) Permeability and outer membrane- a barrier</li> <li>d) Aquaporins</li> </ul>	
a) Using whole cells b) Using Liposomes c) Using Proteoliposome  1.3 Solute transport across membrane a) Passive transport facilitated by membrane proteins. b) Transporters grouped into Superfamilies' ' c) Co transport across plasma membrane (Uniport, Antiport, Symport) d) Active transport & electrochemical gradient e) lon gradient provides energy for secondary Active transport e.g. Lactose transport f) ATPases and transport g) ABC transporters e.g. Histidine transport h) Shock sensitive system — Role of binding proteins e.g. Maltose uptake i) Phosphotransferase system j) Schematic representation of various Membrane transport mechanisms in. E. coli  1.4 Other examples of solute transport b) Bacterial protein export c) Bacterial protein export c) Bacterial membrane fusion central to many biological processes  II Bioenergetics and Bioluminescence		1.2	Methods of studying solute transport	02
a) Passive transport facilitated by membrane proteins. b) Transporters grouped into Superfamilies' ' c) Co transport across plasma membrane (Uniport, Antiport, Symport) d) Active transport & electrochemical gradient e) lon gradient provides energy for secondary Active transport e.g. Lactose transport f) ATPases and transport g) ABC transporters e.g. Histidine transport h) Shock sensitive system – Role of binding proteins e.g. Maltose uptake i) Phosphotransferase system j) Schematic representation of various Membrane transport mechanisms in. E. coli  1.4 Other examples of solute transport b) Bacterial protein export c) Bacterial membrane fusion central to many biological processes  II Bioenergetics and Bioluminescence			a) Using whole cells b) Using Liposomes	
proteins. b) Transporters grouped into Superfamilies' ' c) Co transport across plasma membrane (Uniport, Antiport, Symport) d) Active transport & electrochemical gradient e) Ion gradient provides energy for secondary Active transport e.g. Lactose transport f) ATPases and transport g) ABC transporters e.g. Histidine transport h) Shock sensitive system – Role of binding proteins e.g. Maltose uptake i) Phosphotransferase system j) Schematic representation of various Membrane transport mechanisms in. E. coli  1.4 Other examples of solute transport a) Iron transport: A special problem b) Bacterial protein export c) Bacterial membrane fusion central to many biological processes  II Bioenergetics and Bioluminescence		1.3		80
a) Iron transport: A special problem b) Bacterial protein export c) Bacterial membrane fusion central to many biological processes  Bioenergetics and Bioluminescence 15		RR	proteins. b) Transporters grouped into Superfamilies' (c) Co transport across plasma membrane (Uniport, Antiport, Symport) d) Active transport & electrochemical gradient e) Ion gradient provides energy for secondary Active transport e.g. Lactose transport f) ATPases and transport g) ABC transporters e.g. Histidine transport h) Shock sensitive system – Role of binding proteins e.g. Maltose uptake i) Phosphotransferase system j) Schematic representation of various Membrane transport mechanisms in. <i>E. coli</i>	
b) Bacterial protein export c) Bacterial membrane fusion central to many biological processes    II   Bioenergetics and Bioluminescence   15	01,	1.4		03
<u> </u>	<i>Q</i> -		<ul><li>b) Bacterial protein export</li><li>c) Bacterial membrane fusion central to many</li></ul>	
2.1 Biochemical mechanism of generating ATP 01	II			15
a) Substrate level		2.1	Biochemical mechanism of generating ATP  a) Substrate level	01



		b) Oxidative	
		,	
		c) Photo Phosphorylation	
	2.0	Flacture transport shalin	00
	2.2	Electron transport chain	03
		a) Universal Electron acceptors that transfer	
		Electrons to ETC.	
		b) Carriers in ETC	
		<ul> <li>i. Hydrogen carriers – Flavoproteins, Quinones</li> </ul>	
		ii. Electron carriers-Iron sulphur proteins,	, ( ) ·
		Cytochromes	
		c) Mitochondrial ETC	
		i.Biochemical anatomy of mitochondria	
		ii.Complexes in Mitochondrial ETC	
		iii. Schematic representation of Mitochondrial ETC	
		iii. Schematic representation of Mitochondrial ETC	
	2.3	Prokaryotic ETC	03
		a) Organization of electron carriers in bacteria	
		b) Generalised electron transport pathway in bacteria	
		c) Different terminal oxidases	
		d) Branched bacterial ETC	
		e) Pattern of electron flow in <i>E. coli</i> – aerobic an	
		anaerobic	
		f) Pattern of electron flow in Azotobacter vinelandii	
		1) Fattern of electron flow in Azolobacter virielandii	
	0.4	ATD 41 1	0.4
	2.4	ATP synthesis	04
		a) Explanation of terms – Proton motive force, Proton	
		Coupling sites, P: O ratio, Redox potential	
		b) Free energy released during electron transfer from	
		to O <sub>2</sub> .	
		c) Chemiosmotic theory	
		d) Structure & function of Mitochondrial ATP	
		synthase (No Kinetics)	
		, ,	
		e) Mechanism by Rotational catalysis	
	2	f) Structure of bacterial ATP synthase	
		g) Inhibitors of ETC, Inhibitors of ATPase,	
	V-	Uncouplers, Ionophores	
	0.5	Other medes of memoration of electrical and a	00
	2.5	Other modes of generation of electrochemical energy	02
1112		a) ATP hydrolysis	
6/2.		b) Oxalate formate exchange	
		c) Product efflux, Definition- Lactate efflux	
		d) Bacteriorhodopsin - Definition, Significance,	
		Function as proton pump	
	2.6	Bioluminescence	02
		a) Brief survey of bioluminescent systems	
		b) Biochemistry of light emission	
		c) Schematic diagram	
		d) Significance / Application	
L	·	, ,	



III		Methods of Studying Metabolism & Catabolism of Carbohydrates	15
	3.1	Experimental Analysis of metabolism	03
		<ul> <li>a) Goals of the study</li> <li>b) Levels of organization at which metabolism is studied.</li> <li>c) Metabolic probes</li> <li>d) Use of radioisotopes in biochemistry <ul> <li>i. Pulse labelling</li> <li>ii. Assay &amp; study of radio respirometry –to differentiate EMP &amp; ED</li> <li>e) Use of biochemical mutants.</li> <li>f) Sequential induction technique</li> </ul> </li> </ul>	
	3.2	Catabolism of Carbohydrates	12
		<ul> <li>a) Breakdown of polysaccharides – glycogen, starch,</li> <li>b) cellulose.</li> <li>c) Breakdown of oligosaccharides – lactose, maltose, sucrose, cellobiose</li> <li>d) Utilization of monosaccharides – fructose, Galactose.</li> <li>e) Major pathways- <ul> <li>i. Glycolysis (EMP)</li> <li>ii. HMP Pathway &amp; Significance of the pathway</li> <li>iii. ED pathway,</li> <li>iv. TCA cycle &amp; Significance of the cycle</li> <li>v. Anaplerotic reactions</li> <li>vi. Glyoxylate bypass,</li> <li>vii. Incomplete TCA in anaerobic bacteria</li> <li>viii. Amphibolic role of EMP and TCA cycle</li> <li>ix.Energetics of Glycolysis, ED and TCA-Balance sheet and efficiency calculation</li> </ul> </li> </ul>	
IV	7	Fermentative Pathway & Anabolism of Carbohydrates	15
	4.1	Fermentative pathways (With structures and enzymes)	04
PANK		a) Lactic acid fermentation – i. Homofermentors ii. Heterofermentors iii. Bifidobacterium pathway (Schematic) b) Alcohol fermentation i. by ED pathway in bacteria ii. by EMP in yeasts	
	4.2	Other modes of fermentations in microorganisms	05
		a) Mixed acid b) Butanediol c) Butyric acid	



	d) Butanol-acetone e) Propionic acid (Acrylate pathway and succinate propionate pathway)	
4.3	Anabolism of Carbohydrates	06
	<ul> <li>a) General pattern of metabolism leading to synthesis of a cell from Glucose</li> <li>b) Gluconeogenesis</li> <li>c) Biosynthesis of Glycogen</li> <li>d) Biosynthesis of Peptidoglycan</li> <li>e) Role of carriers in synthesis of LPS and capsule</li> </ul>	I.E.G.K.

- a) Stanier R. Y., Ingraham. J. L, Wheelis. M. L, Painter. P. R., General Microbiology, 5<sup>th</sup> edition, 1987, The Macmillan press Ltd.
- b) Conn, E.E., P. K. Stumpf, G.Bruening and R. Y. Doi, Outlines of Biochemistry, 5<sup>th</sup> edition, 1987. John Wiley &Sons. New York.
- c) Gottschalk, G., Bacterial Metabolism, 2<sup>nd</sup> edition, 1985, Springer Verlag.
- d) White, D., The Physiology and Biochemistry of Prokaryotes, 3rd edition, 1995, Oxford University Press.
- e) Nelson, D. L. and M.M. Cox, Lehninger, Principles of biochemistry. 4<sup>th</sup> edition, 2005, W. H. Freeman and Company.
- f) Rose, A.H. Chemical Microbiology, 3<sup>rd</sup> edition, 1976, Butterworth-Heinemann.
- g) Zubay, G. L, Principles of Biochemistry, 4th edition, 1996, Wm. C. Brown publishers
- h) Mathews, C.K., K.E. van Holde, D.R. Appling, S.J. Anthony-Cahill, Biochemistry, 4<sup>th</sup> edition, 2012, Pearson.
- i) Wilson and Walker, Principles & techniques of Biochemistry & Molecular Biology, 7<sup>th</sup> edition, 2010, Cambridge University Press.
- j) Madigan, M.T. and J.M. Martinko, Brock Biology of Microorganisms, 11<sup>th</sup> edition, 2006, Pearson Prentice Hall;
- k) Cohen, G.N., Microbial Biochemistry. 2<sup>nd</sup> edition, 2006, Springer.



**Course Title: Bioprocess Technology** 

Academic year 2020-21

COURSE	DESCRIPTION		
OUTCOME			
CO 1	Understand and execute the process for isolation and strain		
	improvement of industrially important microorganisms		
CO 2	Outline the types and significance of sterilization process in		
	fermentation industry		
CO 3	Design the process of Inoculum development at various levels of		
	scale-up		
CO 4	Understand the assembly and working of typical fermenters and apply		
	the knowledge to operate fermenters in microbiological industries		
CO 5	Understand, attribute and apply methods of recovery and purification		
	of fermentation products		
CO 6	Recall, infer and apply methods in industrial effluent treatment and		
	correlate it to environment protection		
CO 7	Understand and use spectroscopic techniques in Biological analysis		
CO 8	Recognize the significant role of different organizations in genesis of		
1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 -	Intellectual Property Rights, categorize and use different types of		
ON,	intellectual property rights in protection of intangible properties		



Course Code	Unit	Course/ Unit Title	Credits/ Lectures
RUSMIC 504		BIOPROCESS TECHNOLOGY	2.5 /60
		Upstream Processing	15
	1.1	Strains and Strain Improvement of industrial microorganisms	11
		a) Isolation of industrially important microorganisms     b) Improvement of industrial microorganisms     i. Selection of induced mutants for primary metabolites     ii. Isolation of induced mutants for secondary metabolites	
	1.2	Sterilization	04
		<ul> <li>a) Introduction to the concept of media sterilization and Nabla factor</li> <li>b) Design and methods of batch sterilization</li> <li>c) Design and methods of continuous sterilization</li> </ul>	
II	II Fermenter equipment and control		15
	2.1	Design of fermenter	05
		<ul> <li>a) Inoculum development</li> <li>b) Basics of fermenter <ol> <li>Aseptic operation and containment</li> <li>Body construction</li> <li>Aeration and agitation</li> <li>Achievement and maintenance of aseptic condition</li> <li>Valves- function in general and examples</li> <li>Steam Traps- function in general and examples</li> </ol> </li></ul>	
	2.2	Types of fermenters	05
RAM	AR	a) Acetator b) Cavitator c) Tower fermenter d) Cylindro conical fermenters e) Air lift fermenters i. Outer loop fermenters ii. Inner loop fermenters f) Cyclone column g) Packed tower (generator) h) Rotating disc fermenters i) Bubble cap fermenters	
	2.3	Control of Variables	05
		a) Types of variables	



		b) Sensing and control of	
		i. pH	
		ii. Temperature	
		iii. Dissolved oxygen	
		iv. Flow measurement	
		v. Pressure	
		vi. Inlet/ Exit gas analysis	
		vii. Foam sensing	
			. (^^
III		Downstream processing	15
	3.1	Downstream processing	12
		a. Recovery & Purification of fermentation products:	
		i. Introduction	
		ii. Precipitation	
		iii. Filtration - theory, filter-aids, batch filters (Plate	
		and frame filters), continuous filters (Rotary	
		vacuum),	
		iv. Centrifugation: flocculating agent, range of	
		centrifuges - Basket, tubular bowl.	
		b. Cell disruption methods: Physico-chemical.	
		<ul><li>c. Liquid – Liquid extraction, Solvent recovery,</li></ul>	
		d. Chromatography –Ion exchange &Adsorption	
		e. Membrane processes – Ultrafiltration, reverse	
		osmosis, liquid membranes.	
		f. Drying, Crystallization, Whole broth processing	
	3.2	Environmental aspects	3
		a) Modern methods of effluent treatment	
		b) Carbon Credits	
IV		Bioinstrumentation And IPR	15
	4.1	Bioinstrumentation	8
	8	Principles, working and applications of:	
		a) Spectrophotometry (I. R)	
	1	b) Atomic absorption (AAS) & Atomic Emission	
		spectroscopy (Flame photometry)	
		c) Mass Spectroscopy- MALDI ToF, ESI	
	4.2	Intellectual Property Rights	7
		a) Introduction to Intellectual Property	1
		b) Genesis of IPR - GATT, WTO, TRIPS, World	3
		Intellectual Property Organization (WIPO)	
		c) Types of Intellectual Property – Patents, Copyright,	3
		Trademark, Trade secret, Plant varieties protection	
		act, Industrial Designs, Geographical Indications	
L	i	I	



- a) Casida L. E., "Industrial Microbiology" (2009) Reprint, New Age International (P)Ltd, Publishers, New Delhi
- b) Stanbury P. F., Whitaker A. &Hall--S. J., (1997), "Principles of Fermentation Technology", 2nd Edition, Aditya Books Pvt. Ltd, New Delhi.
- c) H. A. Modi, (2009). \_ 'Fermentation Technology "Vols 1 & 2, Pointer Publications, India
- d) Okafor Nduka (2007) \_ 'Modern Industrial Microbiology and Biotechnology ", Science Publications Enfield, NH, USA.
- e) G Y Shitole and Ram Sable (2012) Environmental Degradation Issues and Challenges (Research publication)
- f) Crueger W. and Crueger A. (2000) "Biotechnology -"A Textbook of Industrial Microbiology",2nd Edition, Panima Publishing Corporation, New Delhi.
- g) Principles and Techniques of Biochemistry and Molecular Biology by Wilson/Walker 7th Edition
- h) Brian Mcneil & Linda M. Harvey, Practical Fermentation Technology, John Wiley and Sons. Pvt. Ltd. (2008).
- i) https://www.wipo.int/edocs/pubdocs/en/intproperty/450/wipo\_pub\_450.pdf WIPO Publication No. 450(E) ISBN 978-92-805-1555-0



Course code	PRACTICALS	3 Credits		
RUSMICP502	Practical Based on 503	_		
	Isolation and detection of Mitochondria			
	2. Isolation and study of Bioluminescent organisms	1.0°		
	Study of oxidative and fermentative metabolism			
	Carbohydrate fermentation tests			
	<ol><li>Mixed acid fermentations- Detection of organic acids by TLC</li></ol>			
	Study of Homo and Heterofermentation in Lactic acid     bacteria			
	7. Detection of enzyme phosphatase			
	8. Quantitative assay of Phosphatase			
	9. Anaerobic fermentation			
RUSMICP502	Practical Based on 504			
	Strip Plate Technique			
	2. Streak Plate Technique			
	3. Gradient plate technique for isolation of mutants.			
	Production and detection of vitamin B12 by bioautography.			
	5. Anaerobic digestion of Industrial effluent- Generation and detection of methane			
	Demonstration of IR spectroscopy and analysis of IR spectrum of one compound			
	7. Demonstration of GC-MS/ LC-MS			



## **Modality of Assessment:**

### **Theory Examination Pattern:**

## A. Internal Assessment- 40%- 40 Marks per paper

Sr No	Evaluation type	Marks
1	One Assignment/Case study/Project/ Presentation	15
2	One class Test (multiple choice questions / objective)	20
3	Active participation in routine class instructional deliveries	05
	TOTAL	40

# B. External Examination- 60%- 60 Marks per paper Semester End Theory Examination:

- 1. Duration These examinations shall be of **two hours** duration.
- 2. Theory question paper pattern:
  - a. There shall be four questions each of 15 marks on each unit.
  - b. All questions shall be compulsory with internal choice within the questions.

### Paper Pattern:

Questions	Options	Marks	Total marks	Questions on
Q.1) A)	Any 2 out of 3	10		
Q.1) B)	Any 1 set out of 2 (i & ii or i & ii)	03 & 02	15	Unit I
Q.2) A)	Any 2 out of 3	10		
Q.2) B)	Any 1 set out of 2 (i & ii or i & ii)	03 & 02	15	Unit II
Q.3) A)	Any 2 out of 3	10		
Q.3) B)	Any 1 set out of 2 (i & ii or i & ii)	03 & 02	15	Unit III
Q.4) A)	Any 2 out of 3	10		
Q.4) B)	Any 1 set out of 2 (i & ii or i & ii)	03 & 02	15	Unit IV



#### **Practical Examination Pattern:**

## A. Internal Examination: 40%-80 Marks

Practical	I		II	
Particulars	Paper I	Paper II	Paper III	Paper IV
Journal	05	05	05	05
Experimental tasks	10	10	10	10
Participation	05	05	05	05
Total	20	20	20	20

### B. External Examination: 60%- 120 Marks

### **Semester End Practical Examination:**

Particulars	Practical I	Practical II
Laboratory work	50	50
Spots/Quiz/Viva	10	10
Total	60	60

### PRACTICAL BOOK/JOURNAL

### Semester V:

The students are required to present a duly certified journal for appearing at the practical examination, failing which they will not be allowed to appear for the examination.

In case of loss of Journal and/ or Report, a Lost Certificate should be obtained from Head/ Coordinator / In charge of the department; failing which the student will not be allowed to appear for the practical examination.

#### **Overall Examination and Marks Distribution Pattern**

#### Semester V

Course		501			502		503			504			Grand Total
	In	Ex	Total	In	Ex	Total	In	Ex	Total	In	Ex	Total	
Theory	40	60	100	40	60	100	40	60	100	40	60	100	400
Practical	20	30	50	20	30	50	20	30	50	20	30	50	200



## Course Title: Gene Manipulation, Bioinformatics, & Virology

## Academic year 2020-21

COURSE	DESCRIPTION
OUTCOME	
CO 1	Understand and explain the fundamentals of gene manipulation
CO 2	Implement bioinformatics tools for genetic analysis and
	structure building
CO 3	Correlate structure and function of important cell components
	of prokaryotic and eukaryotic cells
CO 4	Recalling and categorising various genes and proteins involved in
	functioning of prokaryotic and eukaryotic structures
CO 5	Summarizing the structure, classification, enumeration,
	cultivation and life cycle of viruses.
CO 6	Recognise and compare the commonly used terms like
	cancer, prions, viroids and their replication mechanisms
CO 7	Independently illustrate regulation of lytic and lysogenic
	pathway of lambda phage
CO 8	Test the presence of coliphages and execute experiments for
	their enumeration



Course Code	Unit	Course/ Unit Title	Credits/ Lectures
		OFNE MANUBUL ATION DIGINEODMATIOS	
RUSMIC		GENE MANIPULATION, BIOINFORMATICS,	2.5/60
601		&VIROLOGY	C
I		Gene Manipulation And Bioinformatics	15
	1.1	Basic Principles of Gene Manipulation	07
		a) Cutting and joining DNA: Restriction	
		endonucleases, Ligases, Linkers and Adapters	
		b) Cloning vectors: Characteristics of a good vector,	
		Plasmid vectors, Bacteriophage λ, Expression	
		vectors	
		c) Cloning strategies: Genomic libraries, cDNA	
		libraries, PCR	
	1.2	Bioinformatics	06
	1.2	a) Introduction	
		i. Definition, aims, tasks and applications of	
		Bioinformatics.	
		ii. Overview of prominent Databases, tools	
		and their uses	
		iii. Importance, Types and classification of	
		databases	
		iv. Nucleic acid sequence databases- EMBL,	
		GenBank, Ensembl	
		v. Protein sequence databases-PIR, SWISS-PROT, TrEMBL	
		vi. Protein structure databases: PDB, Cn3D.	
		vii. Pathway analysis: KEGG.	
		b) Applications:	
		i. Transcriptome, Metabolomics,	
		Pharmacogenomics,	
		ii. Phylogenetic analysis, Phylogenetic tree,	
		Annotation, SNPs	
(~)		iii. Sequence alignment global v/s local	
		alignment, FASTA file format, BLAST.	
		iv. Genomics- structural, functional and	
		comparative genomics.	
		v. e. Proteomics- structural and functional	
		proteomics.	



	1.3	Emerging techniques in Genome sciences	02
		a) Microarray technologies	
		b) Karyotyping	
		c) CRISPR-based technologies and applications	
		c, crack resident sections and approximent	
II		Cell Biology	15
	2.1	Structure and function of Prokaryotic cell	07
		a) Cell wall	
		b) Capsule	
		c) Flagella	
		d) Endospore	•
	2.2	Cytoskeleton and cell motility in eukaryotes	08
		a) Cytosol, Ergastoplasm and cytoskeleton	
		b) Structure and function: Microtubules,	
		Microfilaments, Intermediate filaments	
		c) Microtubular organelles – Cilia, Flagella and	
		centrioles	
		d) Microfilament structures and role of associated	
		proteins	
		e) Molecular motors: Myosins, Kinesins, Dyenin	
		Pacia Viralemy	4.5
III	3.1	Basic Virology Viral architecture	15 04
	3.1	a) Capsid, viral genome and envelope	04
		b) Structure of TMV, T4, Influenza virus, HIV	
		b) Gradiale of Tiviv, 14, Illiadiza viido, Filv	
	3.2	Viral classification	02
	0.2	VII al Glassification	02
	3.3	The viral replication cycle	04
		a) attachment,	
		b) penetration,	
		c) uncoating,	
		d) types of viral genome and their replication,	
		e) assembly,	
16	7/	f) maturation and release	
	3.4	Cultivation of viruses	05
		a) cell culture techniques,	
		b) embryonated egg,	
		c) laboratory animals,	
		d) Cell culture methods:	
		e) Equipment required for animal cell culture,	
		f) Isolation of animal tissue	
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IV		Advanced Virology	15
	4.1	Life cycle of viruses	05
		a) T4 phage,	
		b) TMV,	
		c) Influenza Virus and	
		d) HIV	
	4.2	Visualization and enumeration of virus particles	03
		a) Measurement of infectious units	
		i. Plaque assay	
		ii. Fluorescent focus assay	
		iii. Infectious centre assay	
		iv. Transformation assay	
		v. Endpoint dilution assay.	
		b) Measurement of virus particles and their	
		components	
		i. Electron microscopy	
		ii. Atomic force microscopy	
		iii. Haemagglutination	
		iii. Measurement of viral enzyme activity.	
	4.3	Regulation of lytic and lysogenic pathway of lambda	03
		phage	
	4.4	Role of viruses in cancer	02
		a) Definitions,	
		b) characteristics of cancer cell,	
		c) cancer multi step process,	
		d) Human DNA tumor viruses-	
		i. EBV,	
		ii. Kaposi's sarcoma virus,	
	7	iii. Hepatitis B and C virus,	
	R-1	iv.Papilloma Virus	
	4.5	Prions and viroids	02



- a) R. H. Tamarin, (2004), "Principles of genetics", Tata McGraw Hill.
- b) M. Madigan, J. Martinko, J. Parkar, (2009), "Brock Biology of microorganisms", 12th ed., Pearson Education International.
- c) Fairbanks and Anderson, (1999), "Genetics", Wadsworth Publishing Company.
- d) Prescott, Harley and Klein, "Microbiology" 7th edition McGraw Hill international edition.
- e) Edward Wagner and Martinez Hewlett, (2005) "Basic Virology", 2nd edition, Blackwell Publishing
- f) Teri Shors, (2009), "Understanding viruses", Jones and Bartlett publishers.
- g) S.Ignacimuthu, (2005), "Basic Bioinformatics", Narosa publishing house.
- h) Robert Weaver, (2008), "Molecular biology", 3rd ed. McGraw Hill international edition.
- i) Primrose and Twyman, (2001), "Principles of gene manipulation and genomics", 6thed, Blackwell Publishing
- j) Arthur Lesk, (2009), "Introduction to Bioinformatics", 3rd Edition, Oxford University Press
- k) Snustad, Simmons, "Principles of genetics", 3rdedn. John Wiley & sons, Inc.
- I) Lodish, Scott." Molecular cell biology,7th edn, Macmillan higher education, International ed.
- m) Flint, Enquist, Racanillo and Skalka, "Principles of virology", (2009)3rdedn. ASM press
- n) T. K. Attwood & D. J. Parry-Smith, (2003), "Introduction to bioinformatics", Pearson education
- o) Benjamin Lewin, (2014) 9th edition, "Genes IX", Jones and Bartlett publishers.
- p) JD Watson, Baker (2004) 5thedn. "Molecular biology of the gene", CSHL Press and Benjamin Cummings
- q) Jonathan Pevsner, Bioinformatics and Functional Genomics, 3rd Edition, 2015, Wiley Blackwell
- r) Jin Xiong, Essential Bioinformatics, 1st Edition, 2006, Cambridge University Press



**Course Title: Immunology** 

COURSE OUTCOME	DESCRIPTION
CO 1	Evaluate molecules for their antigenicity and explain role of haptens in
	elucidating molecular nature of antigens
CO 2	Compare and contrast between different isotypes of antibodies and
	recall their roles in immune mechanisms
CO 3	Outline mechanisms of antigen processing and presentation and the
	molecules involved thereof
CO 4	Retrieve the process of T and B cell maturation, activation and
	proliferation
CO 5	Summarize and compare the effector responses- Humoral Immunity &
	Cell Mediated Immunity
CO 6	Extrapolate the role of immune system in disease: Unregulated
	response- Hypersensitivity; exemplify the different types
CO 7	Understand the mechanism of Antigen-Antibody interaction & illustrate
	and execute immunological techniques for disease diagnosis
CO 8	Apply the concept of immunity for protection from disease by
	development of vaccine



Course Code/ Unit	Unit	Course/ Unit Title	Credits/ Lectures
RUSMIC 602		IMMUNOLOGY	2.5/60
I		Antigens, Antibodies and MHC	15
	1.1	Antigens	05
		<ul> <li>a) Immunogenicity versus antigenicity</li> <li>b) Factors that influence immunogenicity, Contribution of the biological system to immunogenicity</li> <li>c) Epitopes / antigen determinants (only concepts)</li> <li>d) Haptens and antigenicity</li> <li>e) Immunogenicity of some natural substances – native globular proteins, polysaccharides, lipids, nucleic acids Types of antigens – heterophile antigens, isophile antigens, sequestered antigens, super antigens, bacterial and viral antigens.</li> </ul>	
	1.2	Immunoglobulins	07
	25	a) Immunoglobulins – basic and fine structure b) Immunoglobulin classes and biological activities c) Antigenic determinants on immunoglobulins – isotypes, allotypes, idiotypes d) Immunoglobulin Superfamily e) Monoclonal antibodies, Production (Diagrammatically) & applications	
	1.3	MHC complex and MHC molecules	03
2 AM		a) Structure of class I, and class II molecules; class III molecules     b) Peptide – MHC interaction	
II		Antigen presentation and Activation of Immune cells	15
	2.1	Antigen processing and presentation	02
		<ul><li>a) Antigen presentation- professional and nonprofessional cells</li><li>b) Antigen processing and presentation</li></ul>	



	2.2	Receptor Ligand interactions and activation in T cells	05
		<ul> <li>a) TcR, (alpha-beta, gamma-delta TcR), TcR-CD3 complex structure &amp; functions, Accessory molecules.</li> <li>b) T cell activation, T cell differentiation, Subsets of T cells (TH1, TH2, TH17, T reg), Formation of Memory cells</li> </ul>	
	2.3	Receptor Ligand interactions and activation in B cells	05
		<ul> <li>a) B- cell receptors, Receptor associated molecules, receptor clustering. Antigen processing by B cells B cell activation and differentiation –Antigen recognition and presentation by B cells, Formation of germinal centres and memory cells.</li> <li>b) B-cell responses to Thymus dependent and independent antigens</li> </ul>	
	2.4	Cytokines	03
		a) Properties, types and functions     b) Cytokines secreted by Th1 and Th2 cells	
III		Immune Responses and their Detection	15
	3.1	Humoral Response	05
		<ul> <li>a) Introduction of Humoral response, Primary and secondary responses</li> <li>b) Affinity maturation and somatic hyper mutation, Ig diversity, class switching</li> <li>c) Effector functions of antibodies- Neutralization, opsonization, Complement fixation and ADCC</li> </ul>	
	3.2	Cell mediated effector response	03
		a) Generation and target destruction by Cytotoxic T cells. b) Killing mechanism of NK cells.	
Uh.	3.3	Antigen-Antibody reactions	06
5r		a) Precipitation, b) Agglutination, c) Passive agglutination, d) Agglutination inhibition, e) Radioimmunoassay (RIA), f) Enzyme immunoassays (EIA), g) Immunofluorescence, h) western blot technique	



	3.4	Immunodiagnostics	01
		Modern immunology based diagnostic tests	
IV		Vaccines, Immunohematology And Hypersensitivity	15
	4.1	Vaccines	05
		a) Active and passive immunization b) Types of vaccines - Killed and attenuated vaccines, Whole organism vaccines, Purified macromolecules as vaccines, recombinant viral vector vaccines, DNA vaccines c) Use of adjuvants in vaccine d) New vaccine strategies, Ideal vaccine	NECK.
	4.2	Immunohematology	05
		<ul> <li>a) Human blood group systems, ABO, secretors and non-secretors, Bombay Blood group</li> <li>b) Rhesus system and list of other blood group systems.</li> <li>c) Haemolytic disease of new born, Coombs test.</li> </ul>	
	4.3	Hypersensitivity	05
		Coombs and Gell's classification Type I to Type IV hypersensitivity - Mechanism and manifestation.	

- a) Thomas J. Kindt, Barbara A. Osborne, Richard A. Goldsby, Kuby Immunology, 6th ed, W. H. Freeman & Company 2005
- b) Oven, Punt, Stranford, Kuby Immunology,7th ed W.H. Freeman, 2013
- c) Sulabha Pathak, Urmi Palan, Immunology: Essential and Fundamental, 3rd Ed, Anshan Ltd, 2011
- d) Davis, Dulbecco, Eisen and Ginsberg, Microbiology, 4th ed, Lippincott Williams and Wilkins, 1990.



COURSE	PRACTICALS	3
CODE		Credits
RUSMIC	Practical Based on 601	
P601		
	1. Isolation of genomic DNA of E. coli and measurement of its	
	concentration by UVVIS.	1,0
	Restriction digestion of plasmid DNA	
	Demonstration of PCR	
	Bioinformatics practical On Line Practical	
	a. Visiting NCBI and EMBL websites & list services available,	
	software tools available and databases maintained	
	b. Visiting & exploring various databases mentioned in syllabus	
	i. Using BLAST and FASTA for sequence analysis	
	ii. Fish out homologs for given specific sequences (by teacher	
	<ul> <li>decide sequence of some relevance to their syllabus and related to some biological problem e.g. evolution of a</li> </ul>	
	specific protein in bacteria, predicting function of unknown	
	protein from a new organism based on its homology)	
	iii. Six frame translation of given nucleotide sequence	
	iv. Restriction analysis of given nucleotide sequence	
	v. Pair-wise alignment and multiple alignment of a given	
	protein sequences	
	vi. Formation of phylogenetic tree	
	5. Enrichment of coliphages from sewage	
	6. Enumeration of phages- Phage assay (pilot & proper).	
	7. Demonstration of chick embryo inoculation	
RUSMIC	Practical Based on 602	
P601		
	1. Antigen Preparation: 'O'& 'H' antigen preparation of	
	1. Antigen Preparation: 'O'& 'H' antigen preparation of Salmonella. Confirmation by slide agglutination	
	2. Electrophoresis of serum.	
	3. Demonstration of soluble antigens by precipitation reaction.	
	Immunodiagnostics- Dreyer's drop Widal test	
	Diagnosis of syphilis- TRUST antigen kit	
07	Demonstration of ELISA	
	7. Blood grouping – Direct & Reverse typing	
	8. Major and minor compatibility test	
	Determination of Isoagglutinin titre	
	10. Coomb's Direct test	



## **Course Title: Microbial Biochemistry Part II**

## Academic year 2020-21

COURSE	DESCRIPTION
OUTCOME	
CO 1	Categorize lipids into different classes based on their structure
CO 2	Map the steps in the biochemical pathway for metabolism of lipids
CO 3	Outline pathways for biochemical synthesis, degradation and recycling of nucleic acids
CO 4	Explain mechanisms of catabolism of protein and synthesis of amino acid synthesis in the cell
CO 5	Compare and contrast between various levels of metabolic regulation
CO 6	Explain process of prokaryotic photosynthesis and attribute it to photosynthetic pigments, photochemical apparatus and light and dark reactions
CO 7	Compare and contrast metabolism of different inorganic compounds and outline the concept of Lithotrophy
CO 8	Execute and implement enzyme assays and testing of metabolic processes



Course Code	Unit	Course/ Unit Title	Credits/ Lectures
RUSMIC 603		MICROBIAL BIOCHEMISTRY PART II	2.5/60
I		Lipid Metabolism & Catabolism Of Hydrocarbons	15
	1.1	General introduction to Lipids	02
		<ul> <li>a) Lipids and their functions</li> <li>b) Action of lipases on triglycerides /tripalmitate</li> <li>c) Phospholipids and their properties</li> <li>d) Common phosphoglycerides in bacteria</li> </ul>	
	1.2	Catabolism of Lipids	05
		<ul> <li>a) Oxidation of saturated fatty acid- β oxidation pathway, Energetics of β oxidation of Palmitic acid</li> <li>b) Oxidation of propionic acid.</li> <li>c) Degradation of poly beta hydroxy butyrate</li> </ul>	
	1.3	Anabolism of Lipids	05
		<ul> <li>a) Biosynthesis of straight chain even carbon saturated fatty acid (palmitic acid)</li> <li>b) Biosynthesis of phosphoglycerides in bacteria</li> <li>c) Biosynthesis of PHB</li> </ul>	
	1.4	Catabolism of aliphatic hydrocarbons	03
	20	<ul> <li>a) Oxidation of saturated aliphatic hydrocarbon (nalkane)</li> <li>b) Omega oxidation pathway-</li> <li>c) Pathway in Corynebacterium and yeast</li> <li>d) Pathway in Pseudomonas</li> </ul>	
11		Metabolism Of Proteins And Nucleic Acids	15
	2.1	Protein catabolism	05
Sall		<ul> <li>a) Enzymatic degradation of proteins</li> <li>b) Metabolic fate of amino acids (schematic only</li> <li>c) Metabolism of single amino acids –  i. Deamination reactions</li> <li>ii. Decarboxylation</li> <li>iii. Transamination</li> <li>e) Fermentation of single amino acid -Glutamic acid by Clostridium</li> <li>f) Fermentation of pair of amino acids -Stickland reaction</li> </ul>	



	2.2	Amino acid synthesis	04
		a) Schematic representation of amino acid families	
		b) Synthesis of amino acids of Aspartate family	
	2.3	. Nucleic acid Catabolism	03
		a) Degradation of purine nucleotides up to uric acid	
		formation	
		b) Recycling of purine and pyrimidine nucleotides by	
		salvage pathway	
	2.4	Anabolism of Nucleic Acids	03
	2.4	a) Metabolic origin of atoms in purine and pyrimidine	03
		ring	
		b) Biosynthesis of pyrimidine nucleotides.	
		c) Biosynthesis of purine nucleotides.	
		d) Formation of deoxyribonucleotides.	
		e) Synthesis of nucleotide diphosphates and	
		triphosphates.	
		f) Role of nucleotides (high energy triphosphates)	
			4 =
III	0.4	Metabolic Regulation	15
	3.1	Overview and major modes of regulation	01
		Examples of cellular control mechanism acting at various	
		levels of metabolism (tabulation only)	
	3.2	Allosteric proteins	03
	J.2	a) Definition	- 00
		b) Allosteric enzymes - Role of allosteric enzymes	
		using ATCase as example (no kinetic study)	
		c) Regulatory allosteric proteins	
		<ol> <li>Interaction of proteins with DNA</li> </ol>	
		ii. Structure of DNA Binding proteins	
		iii. Examples - Lac repressor, Trp repressor,	
	7	CAP protein	
		iv. Definition and examples of alarmones	
	2.2	Description of more expression (Transcription)	00
	3.3	Regulation of gene expression (Transcription)	06
	2,	<ul><li>a) Introduction to operon model</li><li>b) Common patterns of regulation of transcription –</li></ul>	
		General concept of positive and negative	
		regulation of operons	
07		i. Lac operon - Mechanism of regulation - Induction	
		- Catabolite repression	
		ii Trp operon - End Product Repression	
		- Attenuation	
		c) Regulation of gene expression	
		i. Multiple Sigma Factors	
		ii. Riboswitches	



	3.4	Regulation of enzyme activity (Post translational regulation)	04
		<ul> <li>a) End-Product Inhibition and Mechanism of End Product Inhibition in branched pathways with examples <ol> <li>Isofunctional enzymes</li> <li>Concerted feedback inhibition</li> <li>Sequential feedback inhibition</li> <li>Cumulative Feedback inhibition</li> <li>Combined activation and inhibition</li> </ol> </li> <li>b) Covalent modifications of enzymes <ol> <li>General examples without structure</li> <li>Monocyclic cascade &amp;inter-convertible enzyme definition</li> <li>Glutamine synthetase system of E.coli</li> <li>Regulation by proteolytic cleavage</li> </ol> </li> </ul>	
	3.5	Population of EMP and TCA	01
	3.3	Regulation of EMP and TCA Schematic and Role of Pyruvate dehydrogenase Complex	UI
		The same and the s	
IV		Prokaryotic Photosynthesis & Inorganic Metabolism	15
	4.1	Prokaryotic photosynthesis	09
	RR	<ul> <li>a) Early studies on photosynthesis</li> <li>i. Light and dark reactions</li> <li>ii. Bacterial photosynthesis</li> <li>iii. Hill reaction</li> <li>b) Phototrophic prokaryotes -Oxygenic, Anox phototrophs examples only</li> <li>c) Photosynthetic pigments</li> <li>d) Location of photochemical apparatus</li> <li>e) Photophosphorylation</li> <li>f) Light reactions in <ul> <li>i. Purple photosynthetic bacteria</li> <li>ii. Green sulphur bacteria</li> <li>ii. Cyanobacteria (with details)</li> </ul> </li> <li>g) Dark reaction <ul> <li>i. Calvin Benson cycle</li> <li>ii. Reductive TCA</li> </ul> </li> </ul>	
	4.2	Inorganic Metabolism	06
<i>S</i> -2		a) Assimilatory pathways- i. Assimilation of nitrate, ii. Ammonia fixation – Glutamate dehydrogenase, iii. Glutamine synthetase, GS-GOGAT, Carbamoyl phosphate synthetase iv. Biological nitrogen fixation (Mechanism for N2 fixation and protection of nitrogenase) v. Assimilation of sulphate	03



b) Dissimilatory path	nways- 2
, ·	ectron acceptor  Paracoccus denitrificans)  electron acceptor
c) Lithotrophy– Enli	st organisms and products formed 1 ogen, carbon monoxide,

- a) Stanier, R. Y., M. Doudoroff and E. A. Adelberg. General Microbiology, 5th edition, The Macmillan press Ltd
- b) Conn, E.E., P. K. Stumpf, G. Bruening and R. Y. Doi. 1987. Outlines of Biochemistry, 5th edition, 1987. John Wiley & Sons. New York.
- c) Gottschalk, G., (1985), Bacterial Metabolism, 2nd edition, Springer Verlag
- d) White, D., (1995), The Physiology and Biochemistry of Prokaryotes, 3rd edition, Oxford University Press
- e) Nelson, D. L. and M.M. Cox (2005), Lehninger, Principles of biochemistry. 4th edition, W. H. Freeman and Company.
- f) Salle, A.J. Fundamental Principles of Bacteriology, 7thedn McGraw Hill Book Co.
- g) Cohen, G.N. (2011). Microbial Biochemistry. 2ndedn, Springer
- h) Madigan, M.T. and J.M. Martinko 2006. Brock Biology of Microorganisms. Pearson Prentice Hall;
- i) Biochemistry 3rd edition, Mathew, Van Holde and Ahern, Pearson Education
- j) Zubay, G. L (1996), Biochemistry, 4th edition, Wm. C. Brown publishers
- k) Principles of Biochemistry, Lehninger, 5thednW. H. Freeman and Company



**Course Title: Industrial Microbiology** 

Academic year: 2020-21

COURSE	DESCRIPTION
OUTCOME	Describing the state of the sta
CO 1	Understand and outline the processes of fermentation for the bulk
	production of primary and secondary metabolites and summarize the
	significance of each step
CO 2	Outline the production of commercially important fermentation
	products like alcoholic beverages, SCP, probiotics etc
CO 3	Extrapolate the examples studied to design and execute conventional
	fermentation processes and be able to collaborate to set up an
	enterprise
CO 4	Explain the principles underlying Bioassays and differentiate and
	compare the methods of Biological assays
CO 5	Test and evaluate activity of fermentation products using
	microbiological assays
CO 6	Summarize factors responsible for contamination during production of
	sterile products, execute preventive measures against contamination
CO 7	Evaluate effectiveness of sterilization procedures and assess the
	Microbiological Quality of pharmaceutical products
CO 8	Outline the salient features of quality management and Good
(-)	Manufacturing Practices



Course Code	Unit	Course/ Unit Title	Credits/ Lectures					
RUSMIC 604		INDUSTRIAL MICROBIOLOGY						
I		Industrial Fermentations: I	15					
		a) Types of alcoholic beverage.	1					
		b) Beer –Ale and Lager	3					
		c) Wine –Red and white & Champagne	4					
		d) Vinegar (acetator& Generator)	2 3					
		e) Bioethanol production-	3					
		-From feedstock to fermentable sugars						
		- Zymomonas mobilis as an alternate ethanol						
		producer	0					
		f) Acetone Butanol Fermentation	2					
II		Industrial Fermentations: II	15					
	2.1	Production of secondary metabolites-	04					
		Antibiotics- Penicillin& Semisynthetic Penicillins						
	2.2	Production of primary metabolites-						
		a) Vitamin B <sub>12</sub> from <i>Propionibacterium</i> & <i>Pseudomonas</i>	03					
		<ul> <li>b) Amino acids- Methods for manufacture, Glutamic Acid (direct)</li> </ul>	01					
		c) Organic acids- Citric acid	02					
		d) Enzymes- Uses of enzymes in industry, Production	04					
		of Fungal amylase by solid substrate fermentation,						
		Stabilization of enzymes- Immobilization techniques	•					
		e) Biotransformation of steroids	01					
III		Industrial Fermentations: III	15					
111	3.1		03					
	3.1)	<ul><li>a) Mushroom cultivation</li><li>b) SCP- Substrates used, Organisms and safety</li></ul>	03 03					
<u> </u>		c) Fermented foods- Bread, fermented cassava, tea	03					
	71	and coffee	03					
		d) Mold modified foods- Types (list only), Production of						
" LA.		Soya sauce	02					
07		e) Lactic acid starter cultures, prebiotics and probiotics	04					
IV		Bioassays & Quality Assurance	15					
	4.1	Bioassays	05					
		a) Comparison of Chemical and Biological assays						
		b) Microbiological assays- Test organisms, types of						
		assay methods and factors affecting.						
		c) Modern methods for assay of fermentation products						



4.3	2 QA, Q	C, GMP	07
		Definitions- Manufacture, Quality, Quality Control, In- Process Control, Quality Assurance, Good Manufacturing Practices. Chemicals & Pharmaceutical production: The five variables, Raw materials, in process Items, Finished Products, Labels and Labelling, Packaging materials, Documentation, Regulations. Control of Microbial contamination during manufacture: Premises and contamination control Manufacture of sterile products, Clean and Aseptic Area, Important publications related to QA	K.GK.
4.5	3 Sterili	zation Control and Sterility Assurance	03
	a)	Bio-burden determinations	
	p)	Environmental monitoring	
	c)	Sterilization Monitors – Physical, Chemical and	
	٠,١	Biological indicators	
	d)	Sterility Testing	

- a) Crueger W. and Crueger A. (2000) "Biotechnology -"A Textbook of Industrial Microbiology",
   2nd Edition, Panima Publishing Corporation, New Delhi.
- b) Casida L. E., "Industrial Microbiology 2009 Reprint, New Age International (P) Ltd, Publishers, New Delhi
- c) H. A. Modi, 2009. 'Fermentation Technology "Vol: 1 & 2, Pointer Publications, India
- d) Prescott and Dunn's 'Industrial Microbiology' (1982) 4th Edition, McMillan Publishers
- e) Hugo & Russell's, Pharmaceutical Microbiology Blackwell Science, Seventh Edition
- f) Peppler, H. J. and Perlman, D. (1979), "Microbial Technology". Vol 1 & 2, Academic Press.
- g) Michael J. Waites, 2001 —Industrial Microbiology: An Introduction, Blackwell Science Publications
- h) Naduka Okafor, -Modern Industrial Microbiology, Science Publications, 2007
- i) https://www.dairyscience.info/index.php/science-and-technology-of-wine/124-thescience-and-technology-of-wine-making.html
- j) Andrew G. Reynolds, "Managing Wine Quality, Vol 1 and 2
- k) Sindhu Raveendran et.al. "Applications of Microbial Enzymes in Food Industry" Food Technology and Biotechnology



- I) O. P. Ahlawat, R. P. Tewari "Cultivation Technology of Paddy-straw Mushroom" (2007), ICAR-National Research Centre for Mushroom
- m) Anupam Mishra, at. al "Training manual on cultivation of tropical mushroom and its value addition", ICAR- Agricultural Technology Application Research Institute
- n) Barbara Speranza, Antonio Bevilacqua, Maria Rosaria Corbo, Milena Sinigaglia "Starter Cultures in Food Production"
- o) R. W. Hutkins, "Microbiology and Technology of Fermented Foods (2006) Blackwell Publications p067-105
- p) https://www.dairyscience.info/index.php/cheese-starters/49-cheese-starters.html
- q) Marth and Steele, "Applied Dairy Microbiology", Lactic acid starter cultures
- r) Probiotics and Prebiotics
- ANN ARAM RUHA AN s) https://www.spg.pt/wp-content/uploads/2015/11/2011-Probiotics FINAL 20110116.pdf



RUSMICP602  Practical Based on 603  1. Qualitative detection of Lipase 2. Estimation of proteins by Lowry's method 3. Qualitative detection of Protease 4. Assay of enzyme Protease 5. Study of breakdown of amino acids — Lysine decarboxylase and Deaminase activity 6. Estimation of uric acid 7. To study catabolite repression 8. Study of Hill reaction 9. Study of photosynthesis in microalgae 10. Study of Lithotrophs — Nitrosification and Nitrification  RUSMICP602  Practical Based on 604  1. Alcohol tolerance for yeast. 2. Sugar tolerance for yeast. 3. Inoculum Development for alcohol fermentation 4. Alcohol fermentation: -Efficiency of fermentation 5. Chemical estimation —Sugar by Cole's Ferricyanide method 6. Chemical estimation —Alcohol Estimation—Dichromate method 7. GC demonstration of ethanol 8. Production of fungal amylase using solid substrate fermentation 9. Immobilization of yeast invertase	COURSE CODE
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or ministration of journal ministration	
10. Mushroom cultivation	
11. Production of Spirulina SCP	
12. Bioassay of an antibiotic Ampicillin	"67"
13. Bioassay of Cyanocobalamin.	
14. Chemical assay of Ampicillin	
15. Sterility testing of water for injection or DPT vaccine.	6-1



## **Modality of Assessment:**

### **Theory Examination Pattern:**

## A. Internal Assessment- 40%- 40 Marks per paper

Sr No	Evaluation type	Marks
1	One Assignment/Case study/Project/ Presentation	15
2	One class Test (multiple choice questions / objective)	20
3	Active participation in routine class instructional deliveries	05
	TOTAL	40

## B. External Examination- 60%- 60 Marks per paper

## **Semester End Theory Examination:**

- 1. Duration These examinations shall be of **two hours** duration.
- 2. Theory question paper pattern:
  - a. There shall be four questions each of 15 marks on each unit.
  - b. All questions shall be compulsory with internal choice within the questions.

### Paper Pattern:

Questions	Options	Marks	Total marks	Questions on
Q.1) A)	Any 2 out of 3	10		11.74
Q.1) B)	Any 1 set out of 2 (i & ii or i & ii)	03 & 02	15	Unit I
Q.2) A)	Any 2 out of 3	10		
Q.2) B)	Any 1 set out of 2 (i & ii or i & ii)	03 & 02	15	Unit II
Q.3) A)	Any 2 out of 3	10		
Q.3) B)	Any 1 set out of 2 (i & ii or i & ii)	03 & 02	15	Unit III
Q.4) A)	Any 2 out of 3	10		
Q.4) B)	Any 1 set out of 2 (i & ii or i & ii)	03 & 02	15	Unit IV



#### **Practical Examination Pattern:**

### A. Internal Examination: 40%-80 Marks

Practical		I	II			
Particulars	Paper I	Paper II	Paper III	Paper IV		
Journal	05	05	05	05		
Experimental tasks	10	10	10	10		
Participation	05	05	05	05		
Total	20	20	20	20		

## B. External Examination: 60%- 120 Marks

### **Semester End Practical Examination:**

Particulars	Practical I	Practical II
Laboratory work	50	50
Spots/Quiz/Viva	10	10
Total	60	60

### PRACTICAL BOOK/JOURNAL

#### Semester VI:

The students are required to present a duly certified journal for appearing at the practical examination, failing which they will not be allowed to appear for the examination.

In case of loss of Journal and/ or Report, a Lost Certificate should be obtained from Head/ Co-ordinator / In charge of the department; failing which the student will not be allowed to appear for the practical examination.

### **Overall Examination and Marks Distribution Pattern**

#### Semester VI

Confector 11													
Course		60′	1		602	2		603	3		60	4	Grand Total
	In	Ex	Total	In	Ex	Total	In	Ex	Total	In	Ex	Total	
Theory	40	60	100	40	60	100	40	60	100	40	60	100	400
Practical	20	30	50	20	30	50	20	30	50	20	30	50	200

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