Laur Singh anur

M.Sc. Microbiology Syllabus as approved in BOS Meeting held on 04.06.2022

V.B.S. PURVANCHAL UNIVERSITY, JAUNPUR 222003 Syllabus

Master of Science in Microbiology (M. Sc. Microbiology) Designed as per Syllabus Development Guidelines of National Education Policy-2020 (NEP-2020)

Programme Structure:

The M.Sc. Microbiology programme is a two-year course divided into four-semesters. A studentis required to complete hundred credits for the completion of course and the award of degree. A student has to accumulate twenty-eight credits in first semester and twenty for credits in each of the remaining (second, third and fourth) semesters.

Part - I First Year	Semester I	Semester II
Part - II Second Year	Semester III	Semester IV

SEMESTER-WISE DETAILS OF M.Sc. MICROBIOLOGY COURSE

		Semester I		
#	Course	Name of the course	Credits	Teaching Hours
Core	B080701T	Bacteriology	4	60
Paper:	B080702T	Bacteriological Techniques	4	60
Theory	B080703T	Cell Biology and Biochemistry	4	60
	B080704T	Molecular Biology & Microbial Genetics	4	60
Minor Elective: Theory	To be offered by other faculty	Minor Elective (Any one out of all the available Minor Elective papers offered from other Faculties)	4	60
Practical	B080705P	Practical I	4	120
Industrial Training/ Survey/ Research	B080706R	Industrial Training/ Survey/ Research Project I	4	
Project		Total Credits	28	

		Semester II		
	Course	Name of the course	Credits	Teaching Hours
Core	B080801T	Immunology and Immunotechnology	4	60
Paper:	B080802T	rDNA Technology	4	60
Theory	B080803T	Virology	4	60
Major	B080804T	Instrumentation and Analytical Techniques	4	60
Elective: Theory (Any one of the two papers)	B080805T	Extremophiles & their Application		
Practical	B080806P	Practical II	4	120

Industrial Training/ Survey/ Research Project	B080807R	Industrial Training/ Survey/ Research Project II	4	
		Total Credits	24	

		Semester III		
#	Course	Name of the course	Credits	Teaching
Core	B080901T	Industrial Microbiology	4	60
Paper:	B080902T	Microbial Physiology & Metabolism	4	60
Theory	B080903T	Environmental Microbiology	4	60
Major	B080904T	Biostatistics & Bioinformatics	4	60
Elective: Theory (Any one of the two papers)	B080905T	Microbial Biodiversity		
Practical	B080906P	Practical III	4	120
Industrial Training/ Survey/ Research Project	B080907R	Industrial Training/ Survey/ Research Project III	4	
		Total Credits	24	

		Semester IV		
#	Course Code	Course	Credits	Teaching Hours
Major	B081001T	Food Microbiology	4	60
Elective:	B081002T	Agricultural Microbiology	4	60
Theory	B081003T	Clinical Microbiology	4	60
(Any four	B081004T	Entrepreneurship, IPR & Biosafety	4	60
out of eight papers):	B081005T	Microbial Pathogenicity	4	60
papers).	B081006T	Plant Pathogen Interaction	4	60
	B081007T	Mycology & Phycology	4	60
	B081008T	Bioprocess Technology	4	60
Practical	B081009P	Practical IV	4	120
Industrial Training/ Survey/ Research	B081010R	Industrial Training/ Survey/ Research Project IV	4	
Project		Total Credits	24	

Note: 1. Up to first three semesters the marks allocated for continuous internal assessment (25 marks) will be

Industrial Training/ Survey/ Research Project	B080807R	Industrial Training/ Survey/ Research Project II	4	
		Total Credits	24	

		Semester III		
#	Course Code	Name of the course	Credits	Teaching Hours
Core	B080901T	Industrial Microbiology	4	60
Paper:	B080902T	Microbial Physiology & Metabolism	4	60
Theory	B080903T	Environmental Microbiology	4	60
Major	B080904T	Biostatistics & Bioinformatics	4	60
Elective: Theory (Any one of the two papers)	B080905T	Microbial Biodiversity		
Practical	B080906P	Practical III	4	120
Industrial Training/ Survey/ Research Project	B080907R	Industrial Training/ Survey/ Research Project III	4	
		Total Credits	24	

		Semester IV		
#	Course Code	Course	Credits	Teaching Hours
Major	B081001T	Food Microbiology	4	60
Elective:	B081002T	Agricultural Microbiology	4	60
Theory	B081003T	Clinical Microbiology	4	60
(Any four out of eight	B081004T	Entrepreneurship, IPR & Biosafety	4	60
papers):	B081005T	Microbial Pathogenicity	4	60
pupersy.	B081006T	Plant Pathogen Interaction	4	60
	B081007T	Mycology & Phycology	4	60
	B081008T	Bioprocess Technology	4	60
Practical	B081009P	Practical IV	4	120
Industrial Training/ Survey/ Research Project	B081010R	Industrial Training/ Survey/ Research Project IV	4	
110,000		Total Credits	24	

Note:

1. Up to first three semesters the marks allocated for continuous internal assessment (25 marks) will be

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evaluated on the basis of class attendance and a seminar. The seminar will be an integral part of the sessional and will be evaluated by all the faculty members of the Department.

2. The detailed syllabus is given in the following pages. The numbers given in front of each topic/group of topics represent the number of periods (60 minutes each) allocated for teaching that topic(s).

M. Sc. Microbiology Programme Objectives (POs)

At the time of completion of the programme the student will have developed extensive knowledgein various areas of Microbiology. Through the stimulus of scholarly progression and intellectual development the programme aims to equip students with excellence in education and skills, thus enabling the student to pursue a career of his/her choice. By cultivating talents and promoting all round personality development through multi-dimensional education a spirit of self-confidence and self-reliance will be infused in the student. The student will be instilled with values of professional ethics and be made ready to contribute to society as responsible individuals.

M. Sc. Microbiology Programme Specific Outcomes (PSOs)

After completing the two years degree course in M. Sc. Microbiology, the students will be:

PSO1: Able to understand and explain the technical aspects associated with existing microbiological challenges.

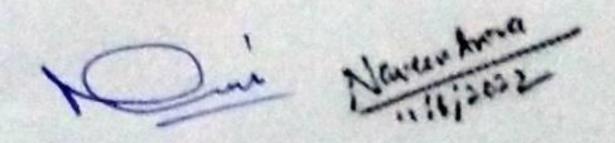
PSO2: Able to explain about various applications of Microbiology such as Environmental Microbiology, Industrial Microbiology, Food Microbiology, and Clinical Microbiology.

PSO3: Able to design and execute experiments related to Basic Microbiology, Immunology, Molecular Biology, Recombinant DNA Technology, and Microbial Genetics.

PSO4: Able to execute a short research project incorporating techniques of Basic and Advanced Microbiology

PSO5: Equipped to take up a suitable position in academia or industry.

		Programme/Class: Bachelor's Degree with Research / M. Sc. Microbiology (I)		
Subject: Mic	robiology	Year: First (1)	Semester: 1	First (I)
The state of the s	er (Compulsory)	Course Code: B080701T	BACTERIO	NAME AND ADDRESS OF THE OWNER, WHEN PERSON ASSESSED.
AND DESCRIPTION OF THE PARTY OF	arks: 100	75 (UE) + 25 (CIE)	Credits	THE RESERVE OF THE PARTY OF THE
		ials-Practical (in hours per week L-		. 04
Course		tive of the course is to build a strong for		a of bacterial
Objectives	cell structure, divi	sion, survival and propagation	oundation in the are	a of bacteria
Course	the state of the s	ompletion of the course, the student:		
Learning				
Outcomes				
COI		scribe the morphological features, cel cterial cell in detail; will be able to di		
		n-negative bacteria.	ricicinate between	Grani-
CO2		knowledge about cell wall structure as	nd extracellular apr	pendages in
		and is acquainted with current method		
		oplasts, sphaeroplasts and L-forms.	and distantante i	
CO3		d detailed information regarding bacte	erial cell division a	nd
		on. Can enlist the salient features of t		
CO4				a, and will
		atures of some model archaeal organis		,
CO5		e basic concept of bacterial systematic		species.
		그는 그들은 그는 그는 그는 그는 그는 그는 그는 그는 그들은 그들은 그들은 그는 그는 그를 보고 있다.		
		standing of phenetic and phylogenetic		
	Develop an under	standing of phenetic and phylogenetic		
	Develop an under	standing of phenetic and phylogenetic		polyphasic
Contents	Develop an unders approach of taxon	standing of phenetic and phylogenetic	classification with	polyphasic Duration:
Contents	Develop an unders approach of taxon	standing of phenetic and phylogenetic omy	bacterial cell	Duration: 60 hours
Contents	Bacterial cell structure organization: nucle	standing of phenetic and phylogenetic omy	bacterial cell	Duration: 60 hours
Contents	Bacterial cell structure organization: nucle cell inclusions. De	eture and appendages: Overview of euceoid, ribosomes, intracytoplasmic me	bacterial cell mbranes and tion of various	Duration: 60 hours
Contents	Bacterial cell structure appears appears and appears and appears appears appears and appears a	ture and appendages: Overview of euceoid, ribosomes, intracytoplasmic metailed account of biogenesis and func	bacterial cell mbranes and tion of various and mechanism	Duration: 60 hours
Contents	Bacterial cell structure appears of movement; pili	standing of phenetic and phylogenetic omy ture and appendages: Overview of eu- eoid, ribosomes, intracytoplasmic me- tailed account of biogenesis and func- ndages: flagella- structure, assembly a	bacterial cell mbranes and tion of various and mechanism ir role. External	Duration: 60 hours
Ontents JNIT I	Bacterial cell structure appears of movement; pilicell surface structure	ture and appendages: Overview of euceoid, ribosomes, intracytoplasmic metailed account of biogenesis and functudages: flagella- structure, assembly and fimbriae- types, structure and the	bacterial cell mbranes and tion of various and mechanism ir role. External and S-layer.	Duration: 60 hours
Ontents JNIT I	Bacterial cell structure appears of movement; pilicell surface structure Bacterial cell wall gram positive bact	eture and appendages: Overview of euceoid, ribosomes, intracytoplasmic metailed account of biogenesis and functural findages: flagella- structure, assembly and fimbriae- types, structure and the ares: capsule, glycocalyx, slime layer and cell membrane: Overview of granterial cell wall, outer membrane lipoper	bacterial cell mbranes and tion of various and mechanism ir role. External and S-layer. m negative and olysaccharide	Duration: 60 hours 12 Hours
Contents JNIT I	Bacterial cell structure appears of movement; pilicell surface structure Bacterial cell wall gram positive bacte (LPS). Detailed ac	cture and appendages: Overview of euceoid, ribosomes, intracytoplasmic mentailed account of biogenesis and functured findages: flagella-structure, assembly and fimbriae-types, structure and the ares: capsule, glycocalyx, slime layer and cell membrane: Overview of granterial cell wall, outer membrane lipoper count of cell wall synthesis and its	bacterial cell mbranes and tion of various and mechanism ir role. External and S-layer. m negative and olysaccharide	Duration: 60 hours 12 Hours
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Ontents UNIT I	Bacterial cell structure appears of movement; pilicell surface structure Bacterial cell wall gram positive bacterial cell divisions. Detailed actincluding different Bacterial cell divisions.	eture and appendages: Overview of euceoid, ribosomes, intracytoplasmic metailed account of biogenesis and functural findages: flagella- structure, assembly and fimbriae- types, structure and the cres: capsule, glycocalyx, slime layer and cell membrane: Overview of graderial cell wall, outer membrane lipope count of cell wall synthesis and its antibiotics.	bacterial cell mbranes and tion of various and mechanism ir role. External and S-layer. m negative and olysaccharide inhibitors	Duration: 60 hours 12 Hours
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Ontents UNIT I	Bacterial cell structure apper of movement; pilicell surface structure Bacterial cell wall gram positive bacterial cell divise Bacte	eture and appendages: Overview of euceoid, ribosomes, intracytoplasmic metailed account of biogenesis and functuring flagella- structure, assembly and fimbriae- types, structure and the tres: capsule, glycocalyx, slime layer and cell membrane: Overview of graderial cell wall, outer membrane lipope count of cell wall synthesis and its antibiotics. ion and reproduction: Genome organother forms of reproduction in bacter	bacterial cell mbranes and tion of various and mechanism ir role. External and S-layer. m negative and olysaccharide inhibitors nization of E.coli, eria, bacterial cell ring, endospore	Duration: 60 hours 12 Hours
Ontents UNIT I	Bacterial cell structure apper of movement; pilicell surface structure Bacterial cell wall gram positive bacterial cell divise Bacte	eture and appendages: Overview of euceoid, ribosomes, intracytoplasmic mentailed account of biogenesis and functured and fimbriae-types, structure and the ares: capsule, glycocalyx, slime layer and cell membrane: Overview of grant erial cell wall, outer membrane lipope count of cell wall—synthesis and its antibiotics. ion and reproduction: Genome organ other forms of reproduction in bacter maintenance and disassembly of Z	bacterial cell mbranes and tion of various and mechanism ir role. External and S-layer. m negative and olysaccharide inhibitors nization of E.coli, eria, bacterial cell ring, endospore	Duration: 60 hours 12 Hours
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Ontents UNIT II NIT III	Bacterial cell structure apper of movement; pilicell surface structure Bacterial cell wall gram positive bacterial cell divise Binary fission and cycle, assembly, structure and stag subtilis. Archaebacteria and	eture and appendages: Overview of euceoid, ribosomes, intracytoplasmic mentailed account of biogenesis and functure and fimbriae-types, structure and the ares: capsule, glycocalyx, slime layer and cell membrane: Overview of grant erial cell wall, outer membrane lipoper count of cell wall synthesis and its antibiotics. ion and reproduction: Genome organ other forms of reproduction in bacter maintenance and disassembly of Z es involved in endospore developed.	bacterial cell mbranes and tion of various and mechanism ir role. External and S-layer. m negative and olysaccharide inhibitors nization of E.coli, eria, bacterial cell ring, endospore ment in Bacillus tremophiles like-	Duration: 60 hours 12 Hours
Ontents UNIT II NIT III	Bacterial cell structure apper of movement; pilicell surface structure Bacterial cell wall gram positive bacterial cell divise Binary fission and cycle, assembly, structure and stag subtilis. Archaebacteria and hyperthermophiles.	eture and appendages: Overview of euroid, ribosomes, intracytoplasmic mentailed account of biogenesis and functure and fimbriae-types, structure, assembly and fimbriae-types, structure and the ares: capsule, glycocalyx, slime layer and cell membrane: Overview of granerial cell wall, outer membrane lipoper count of cell wall synthesis and its antibiotics. ion and reproduction: Genome organ other forms of reproduction in bacternaintenance and disassembly of Z es involved in endospore developed. Extremophiles: Introduction to expsychrophiles, halophiles, acidophiles, acid	bacterial cell mbranes and tion of various and mechanism ir role. External and S-layer. m negative and olysaccharide inhibitors nization of E.coli, eria, bacterial cell ring, endospore ment in Bacillus tremophiles like- les, methnogenic	Duration: 60 hours 12 Hours
Ontents UNIT II NIT III	Bacterial cell structure apper of movement; pilicell surface structure Bacterial cell wall gram positive bacterial cell divise Binary fission and cycle, assembly, structure and stag subtilis. Archaebacteria and hyperthermophiles extremophiles etc.	eture and appendages: Overview of euceoid, ribosomes, intracytoplasmic mentailed account of biogenesis and functure and fimbriae- types, structure and the ares: capsule, glycocalyx, slime layer and cell membrane: Overview of graderial cell wall, outer membrane lipoper count of cell wall—synthesis and its antibiotics. ion and reproduction: Genome organ other forms of reproduction in bacternaintenance and disassembly of Z es involved in endospore development of the count of cell wall. Introduction to expect the count of cell wall and disassembly of Z es involved in endospore development. Introduction to expect the count of cell wall and disassembly of Z es involved in endospore development. Adaptation mechanisms of extremop	bacterial cell mbranes and tion of various and mechanism ir role. External and S-layer. m negative and olysaccharide inhibitors nization of E.coli, eria, bacterial cell ring, endospore ment in Bacillus tremophiles like- les, methnogenic hiles, Importance	Duration: 60 hours 12 Hours
Ontents UNIT II NIT III	Bacterial cell structure apper of movement; pilicell surface structure Bacterial cell wall gram positive bacterial cell divise Binary fission and cycle, assembly, structure and stage subtilis. Archaebacteria and hyperthermophiles, extremophiles etc. of extremophiles etc.	eture and appendages: Overview of euroid, ribosomes, intracytoplasmic metailed account of biogenesis and functionages: flagella- structure, assembly and fimbriae- types, structure and the res: capsule, glycocalyx, slime layer and cell membrane: Overview of graderial cell wall, outer membrane lipope count of cell wall—synthesis and its antibiotics. ion and reproduction: Genome organ other forms of reproduction in bacter maintenance and disassembly of Z es involved in endospore development of the count of cell wall. Introduction to expression of the count of cell wall in endospore development of the count of cell wall in endospore development of the count of cell wall in endospore development of the count of cell wall in endospore development of the count of cell wall in endospore development of the count of cell wall in endospore development of the count of cell wall in endospore development of the count of cell wall in endospore development of the count of cell wall in endospore development of the count of cell wall in endospore development of the count	bacterial cell mbranes and tion of various and mechanism ir role. External and S-layer. m negative and olysaccharide inhibitors nization of E.coli, eria, bacterial cell ring, endospore ment in Bacillus tremophiles like- les, methnogenic hiles, Importance harmaceuticals &	Duration: 60 hours 12 Hours
Ontents JNIT I	Bacterial cell structure apper of movement; pilicell surface structure Bacterial cell wall gram positive bacterial cell divise Binary fission and cycle, assembly, structure and stage subtilis. Archaebacteria and hyperthermophiles etc. of extremophiles etc. of extremophiles etc.	eture and appendages: Overview of euroid, ribosomes, intracytoplasmic metailed account of biogenesis and functure and fimbriae- types, structure and the pressure capsule, glycocalyx, slime layer and cell membrane: Overview of grant erial cell wall, outer membrane lipoper count of cell wall synthesis and its antibiotics. ion and reproduction: Genome organism other forms of reproduction in bacter and intenance and disassembly of Z es involved in endospore development of the count of expectation and the count of cell wall synthesis and its antibiotics. In an antipication of the count	bacterial cell mbranes and tion of various and mechanism ir role. External and S-layer. m negative and olysaccharide inhibitors nization of E.coli, eria, bacterial cell ring, endospore ment in Bacillus tremophiles like- les, methnogenic hiles, Importance harmaceuticals &	Duration: 60 hours 12 Hours
Ontents UNIT II NIT III	Bacterial cell structure apper of movement; pilicell surface structure Bacterial cell wall gram positive bacte (LPS). Detailed actincluding different Bacterial cell divise Binary fission and cycle, assembly, structure and stage subtilis. Archaebacteria and hyperthermophiles extremophiles etc. of extremophiles etc. of extremophiles etc.	eture and appendages: Overview of euroid, ribosomes, intracytoplasmic metailed account of biogenesis and functionages: flagella- structure, assembly and fimbriae- types, structure and the res: capsule, glycocalyx, slime layer and cell membrane: Overview of graderial cell wall, outer membrane lipope count of cell wall—synthesis and its antibiotics. ion and reproduction: Genome organ other forms of reproduction in bacter maintenance and disassembly of Z es involved in endospore development of the count of cell wall. Introduction to expression of the count of cell wall in endospore development of the count of cell wall in endospore development of the count of cell wall in endospore development of the count of cell wall in endospore development of the count of cell wall in endospore development of the count of cell wall in endospore development of the count of cell wall in endospore development of the count of cell wall in endospore development of the count of cell wall in endospore development of the count of cell wall in endospore development of the count	bacterial cell mbranes and tion of various and mechanism ir role. External and S-layer. m negative and olysaccharide inhibitors nization of E.coli, eria, bacterial cell ring, endospore ment in Bacillus tremophiles like- les, methnogenic hiles, Importance harmaceuticals & of archaeal cell	Duration: 60 hours 12 Hours



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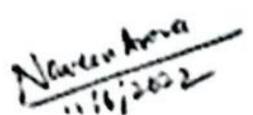
	Analysis, Polyphasic Taxonomy, FAME Analysis, Prokaryotic Species Concept, Phylogenetic trees. General features of Archaea, Actinomycetes, Cyanobacteria, Mollicutes, Rickettsia and Chalamydia. Actinomycetes, Cyanobacteria, Mollicutes, Rickettsia and Chalamydia.
Suggested Readings	 Actinomycetes, Cyanobacteria, Mollicutes, Rickellist and Color and Color

		Programme/Class: Bachelor's Degree with Research M. Sc. Microbiology (I)		
Subject	t: Microbiology	Year: First (1)	Semester:	Contract of the Contract of th
Core Pap	er (Compulsory)	Course Code: B080702T	BACTERIO TECHN	And a second lands of the land
N	larks:100	75 (UE) + 25 (CIE)	Credit	s: 04
Total Numl	ber of Lectures-Tutor	ials-Practical (in hours per week l	L-T-P: 4-0-0	
Course	The primary obj	ective of the course is to build a	i basic foundation	in the area
Objectives	bacteriological te	chniques used for isolation and culti	vation of bacteria.	1000
Course	Upon successful	completion of the course, the student	t;	
Learning				
Outcomes			biological tacksia	ues and their
CO1		the basic concepts of various micro	biological techniq	ues and men
	applications.	1 1 the best mainsiples of star	dization It will als	o develop an
CO2	Will be able to ur	derstand the basic principles of steriout selection of suitable method for s	terilization and dis	infection
CO2	Will loom the foot	tures of bacterial growth and phases	of bacterial growth	with various
CO3	factors affecting g		out	
	lactors affecting g	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,		
CO4	Will understand	learn various conventional and no	on-conventional te	chniques of
	isolation and cul-	tivation of bacteria. He/she will a	lso be able to un	derstand the
		lation of unculturables.		
CO5	Will learn various	techniques for short term and long te	erm storage of micr	oorganisms.
	Contents			Durations
Contents				Duration: 60 hours
	110	C i i constian a	- conification	12 Hours
NIT I	Microscopy: Basic	s of microscopy: image formation, r	nagnification,	12 Hours
	leinds of microscor	cal applications and instrumentation by: Optical Microscopy, Fluorescence	e Confocal	
	and Electron Micro	oscopy. Stains, dyes and staining tec	hniques	
NIT II		ection and Sterilization: Chemical		12 Hours
NII II		aldehyde Phenolic Compounds		12 110015
		ounds, Chlorine, Iodophors and		
	The state of the s	loist Heat, Dry Heat, Mathematica		
	THE RESERVE AND ADDRESS OF THE PARTY OF THE	sses, Arrhenius equation, Del fac		
		edia quality and yield coefficier		
		zation, Sterilization Gases (Eth		
	Formaldehyde, Hyd	drogen Peroxide, Chlorine Dioxide)	and Filtration.	The same of
	filter and steam ster	ilization at industrial scale		
III TIV	Bacterial Growth:	Definition of growth, mathematical	expression of	12 Hours
	0	rve, measurement of growth and		
	THE RESERVE OF THE PARTY OF THE	re, Introduction of continuous cu	ulture; Factors	
	affecting growth.			
IT IV	A PART OF THE PART	ation and Cultivation: Techniques 1	A CONTRACTOR OF THE PROPERTY O	12 Hours
		obic Bacteria; Micromanipulation t		
		ulation systems (Optical tweeze		1 2 2
		iltivation of bacteria and fungi, App		
		turables, Types of media. Technique	es for isolation	
	and cultivation of V	iruses and Fungi		The state of the s

UNITV	Preservation and Maintenance of Microorganisms: Short-Term Preservation Methods- Subculturing, Immersing in Oil, Ordinary Freezing, Deep Freezing, Drying: Long-Term Preservation Methods- Freeze-Drying (Lyophilization and Ultrafreezing: Preservation of Representative Genera And Specific Groups- Anaerobes, Cyanobacteria, Methanogens, Plasmid-Containing Bacteria and Spore formers. Culture Collections and their Functions.
Suggested Readings	 Prescott's Microbiology by J. Willey, L. Sherwood, C. J. Woolverton. 10th edition.McGraw Hill Education. 2017. Brock Biology of Microorganisms by M. Madigan, K. Bender, D. Buckley, W. Sattley, D.Stahl. 15th Edition. Pearson Education. 2018. Alcamo's Fundamentals of Microbiology by J. C. Pommerville. 10th Edition. Jones andBartlett Learning. 2013. Archaea Molecular and Cellular Biology by Ricardo Cavicchioli. American Society ofMicrobiology. 2007. The Physiology and Biochemistry of Prokaryotes by D. White, J. Drummond, C. Fuqua. 4th Edition. Oxford University Press. 2011.

V., b.)		Programme/Class: Bachelor's Degree with Researc M. Sc. Microbiology (I)	h /		
Subject: 6	licrobiology	Year: First (1)	Semester:	Fines (II)	
Core Paper	(Compulsory)	Course Code: B080703T	Semester: I BIOCHEMISTI	FIRST (I)	
Mar	ks:100		BIOLO		
		75 (UE) + 25 (CIE)	The second secon		
Course	The primer	rials-Practical (in hours per week	1 T D. 100		
Objectives	biomolecules an structure compo	d Cell Biology. The course has been sition and functional aspects of the	pasic understanding all developed to understa	oout various and the basic	
Learning Outcomes	Opon successiu	completion of the course, the stude	ent:		
COL	Will learn abou	at structure and functions of protei	ns and linide Davale		
CO2		and seducine ino			
CO2	Will learn abou	t structure, functions and classificat	ion of carbohydrates	and Nuclaia	
CO3	The state of the s				
	madiata antal	individual proteins bind to specific is.	substrates and other n	nolecules to	
CO4		A 700.7			
CO5	Will understan	the basic structure composition and	I functional aspects of	the cell.	
	direct start	d die basie concents of cell to	call communicati	cell cycle,	
Contents	Tragamina ce	ell death and mechanisms of develop	ment of cancer.		
				Duration: 60	
UNIT I	protein sequen	Proteins- Primary; secondary (Randaternary structure; Protein folding cing. Lipids: Classification, structure;	and methods of	hours 12 Hours	
UNIT II	runction of fatt	yacids			
	sugars and glyc Nucleic Acids; structure of tR	Structure and function of nucleoting NA, DNA topology; A, B and Z DN.	des, RNA and 3D	12 Hours	
UNIT III	cofactors and action- Compe inhibition, Allo	structure and function of Enzy prosthetics groups. Enzyme kineti etitive, Uncompetitive, Non-compe- osteric and Regulatory enzymes.	mes, coenzymes, cs: Mechanism of etitive and Mixed	12 Hours	
UNIT IV	interaction: Ce	of Eukaryotic Cell: Structure and fu chloroplast, mechanism of Protein II-cell adhesion, cytoskeleton.	segregation, Cell	12 Hours	
UNIT V	Carcinogenesis	and cell differentiation, Cell cycharacteristics of cancer cells, s, Agents promoting carcinogenesis.	Mechanism of	12 Hours	
Suggested Readings:	1. Princip Macmi 2. Harper P.A. M 3. Bioche	les of Biochemistry (5th Edition) -	Lehninger, Nelson and ition) – R.K. Murray, I Graw Hill International b: Wm. C. Brown Pub	D.K. Garner, I Edition.	





- 5. Biochemistry (2nd edition) D. Voet and J.G. VoetPub: John Willy and Son
- Molecular biology of the cell, (4th Edition) Bruce Albert, Alexander Johnson, Julian Lewis, Martin Raff, Keith Roberts and Peter Walter, Pub: G.S. Garland science Taylor and Francis Group New York – NY 10001-
- 7. Molecular Cell Biology, (5th Edition) H. Lodish, A. Berk P. Matsudaira Chris A.Kaiser, M.Krieger. M. P. Scott, L. Zipursky, J. Darnell. Pub: W.H. Freeman and Com., NY.
- 8. Cell and Molecular Biology: Concepts and Experiments: Gerald Karp, VIthEds

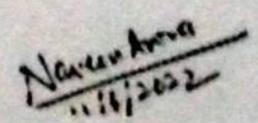
		Programme/Class: Bachelor's Degree with Researc M. Se. Mierobiology (I)	h /	
Subject: N	Aicrobiology	M. Sc. Microbiology (I) Year: First (1)	Comestan	Eine (I)
AND PROPERTY OF THE PERSON NAMED IN COLUMN TWO IS NOT THE OWNER, THE PERSON NAMED IN COLUMN TWO IS NOT THE OWNER.	(Compulsory)	Course Code: B080704T	Semester: MOLECULAR	
			MICROBIAL	
and the second second second second	ks:100	75 (UE) + 25 (CIE)	Credits	
Fotal Number	of Lectures-Tuto	rials-Practical (in hours per week	L-T-P: 4-0-0	
Course Objectives	The primary objection its multiplication systems	ective of this course is to develop a , expression and regulation in prok	n understanding of st aryotic and eukaryoti	ructure of gene ic microbial
Course Learning Outcomes	Upon successful	completion of the course, the stude	ent:	
COL	Will learn about	mechanism of Conjugation and bac	terial gene mapping	
CO2		mechanism of transformation and t		
CO3 ·	Will learn how t	the DNA replicates in Prokaryotic a NA repair. Will also learn about co	ind eukaryotic cell al	
CO4	mechanisms of	the basic concepts of transformation ost transcriptional RNA processing	,	277.52
CO5	genetic elements	the mechanism of gene regulations	on and learn about	
Contents:				Duration: 60 hours
UNIT I		Conjugation, discovery, formation of combination, concept of transferoson		12 Hours
UNIT II	positive and	discovery, mechanism of transformation description of transformation of transformati	ction- discovery,	12 Hours
UNIT III	repair and record	genetic material in prokaryotes and nbination, DNA mismatch repair, Doination as a molecular biology ting, regulating and targeting genom	ouble Strand Break ool, CRISPR-Cas	12 Hours
UNIT IV	transcription fa and Eukaryotes and rRNA).	f transcription in prokaryotes ctors. RNA polymerases., Translation. Post transcriptional RNA procession	on: In Prokaryotes ng (mRNA, tRNA	12 Hours
UNIT V	of gene express	gene expression, Positive and negational and trp-operon, attenuation. An oversion in prokaryotes and eukaryotes, a lecules. Transposable genetic elements	rview of regulation cis acting sites and nts.	12 Hours
Suggested Readings	2. Principle 3. Modern Griffith 4. Gene by 5. Molecu 6. Genetic	s: Analysis and Principles by Robert es of Genetics by Eldon J. Gardner, Genetic Analysis: Integrating Genes s 7th y Benjamin Lewin, IXthEds,Oxford lar Biology of gene by Watson, 12th es Strickberger 13thEds cular Biology (8th Edition) – DeRo	12thEds es and Genomes by A Univ. Press, U.K Eds	Anthony J.F.

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Publishers Pvt Ltd. N. Delhi

	Programme/Class: Bachelor's Degree with Research M. Sc. Microbiology (I)	1
Subject: Microbiology	Year: First (1)	Semester: First (I)
Minor Elective (Optional)	Course Code: to be provided by other faculty	MINOR (OTHER FACULTY)
Marks: 100	75 (UE) + 25 (CIE)	Credits: 04
Total Number of Lectures-Tuto	rials-Practical (in hours per week	L-T-P: 4-0-0
		Duration: 60 hours





		Programme/Class: Bachelor's Degree with Research / M. Sc. Microbiology (I)	
Core Pape M 75 (UI	er (Compulsory) arks:100 E) + 25 (CIE)	Year: First (1) Course Code: B080705P Credits: 04	Semester: First (I) Practical I Duration: 120 hours
Course Objectives	and biochemical identification methods with sterilization to	als-Practical (in hours per week L-7) ve of the course is to impart hands-o techniques. Students will be trained nods, as well as working in biosafety ca echniques when handling bacterial ce that to present the results both, qualitation	n training in basic microbiological d in basic bacterial culturing and abinet. Student will become familiar
Course Learning Outcomes	Upon successful c	the present the results both, qualitation of the course, the student:	vely and quantitatively.
CO1: CO2: CO3:	Will be able to use	e different sterilization procedures and ork in Biosafety Cabinet.	learn handling of micropipette.
CO3: CO4: CO5:	variety of cultural	th identification and classification of gi and biochemical tests.	ven bacterial isolate by performing

Contents:

CO5:

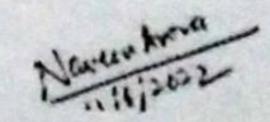
1. To train students in handling, upkeep and calibration of micropipette for measuring small volumes

Can determine concentration of sugar and protein in a given sample after drawing a standard

- To give hands-on training in sterilization techniques and their application in microbiology lab
- To train student in working with a biosafety cabinet in a BSL2 lab
- To purify and identify the given bacterial sample by determining their:- Colony morphology, staining characteristics and biochemical characteristics
- To analyze the given 16srRNA sequences by using BLAST and construct a phylogenetic tree based on the comparison results.
- To draw the titration curve of amino acid and determine its pl.
- To prepare standard curve of BSA and determine the concentration of unknown protein sample using Bradford / Lowry method using regression equation.
- Quantitative estimation of carbohydrate (anthrone/phenol-H2SO4/Dinitrosalisylic acid method).
- Quantitative estimation of proteins by biuret.
- 10. Saponification and acid value of fats
- 11. Estimation of DNA by diphenylamine method
- 12. Estimation of RNA by orcinol method
- 13. To prepare standard curve of ammonia and determine its uptake by bacterial cells with respect to time and temperature
- 14. To determine the specific growth rate of E. coli in different media.
- 15. Staining techniques for bacterial cells: simple, differential, negative, specialized
- 16. Measurement of growth and preparation of growth curve
- 17. To study glucose uptake by E. coli.
- 18. Effect of temperature, pH, salt concentration, antibiotics on growth.
- 19. Calculation of generation time and specific growth rate.
- 20. Microscopic measurements (micrometry)

Suggested Readings:	 Microbiology: A laboratory manual by JG Cappucino, C.T. Welsh. 11th edition. Pearson. 2017.
	2. Biochemistry Lab Manual by D.A. Thompson. 3rd edition. Create Space
The state of the s	10

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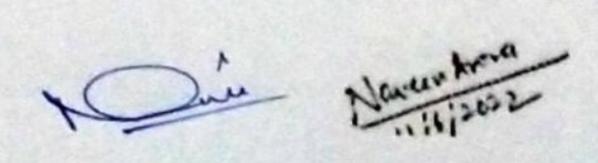


Independent Publishing Platform. 2013.

- 3. Biochemical calculations: how to solve mathematical problems in general biochemistry by Irwin H. Segel, Wiley, 2nd Edition 2004
- 4. Practical Biochemistry (3rd Edition) David Plummer, Pub: Tata McGraw Hill
- 5. Practical Biochemistry (5th Edition) K. Wilson and J. Walker. Pub: Cambridge Univ. Press, (U.K.)

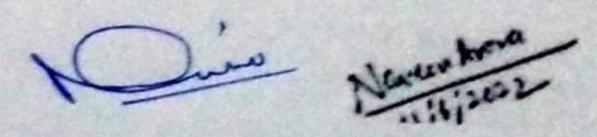
		Programme/Class: Bachelor's Degree with Research / M. Sc. Microbiology (I)		
	Aicrobiology	Year: First (1)	Semester: First (I)	
Core Paper (Compulsory)		Course Code: B080706R	Industrial Training	
	rks:100	Cred	/Surveys/Research Project I ts: 04	
Course Details This research proje in the form of indus		t can be interdisciplinary / multi-disciplinary / m	linary. This research project can also be	
	carried out in both	omit the final report (project report) the semesters at the end of the year, external examiner nominated by the	dissertation) of the research project	

Subject: N	licrobiology	Programme/Class: Bachelor's Degree with Research M. Sc. Microbiology (I)	ch /	
Core Paper	(Compulsory)	Year: First (1)	Com	
Viar	20.100	Course Code: B080801T	Semester: S	econd (II)
Total Number	of Leature To	75 (UE) + 25 (CIE) rials-Practical (in hours per weel I facilitate in understanding of	VIROL	OGY
Course	The control	rials-Practical (in hours per week	Credit	s: 04
Objectives:	viral reproduction a host, and are r	ses and principles in viruses to illustrate. on. The course will teach the strate.	trate viral complexity	, to understand
Course Learning Outcomes:		d addresses the interplay between valued completion of the course, the stud		oiology of viral organisms
COI	Is able to descri	ho al. 10		
CO2	Is able to descri	be classification of viruses		
	entry, virus asse	be tools for studying virus structure embly and release	e, process of virus att	a a bus a state of
CO3	Is able to door	and release	process of virus att	achment and
	DNA viruses	be steps in replication of genome of	f RNA viruses rotro	
CO4	Is able to deser		ruses, retrov	iruses, and
CO5	Is able to descri	ibe steps in virus infection, transmi lost defense against virus infection be methods of making virus vaccine and emerging viruses		
Contents:		and emerging viruses		
LIMITEL				Duration:
UNITI	Introduction to	Virology: The big picture of all	1 vinues	60 hours
LINUTE	virus infection. double strande stranded (ssDN stranded RNA RNA with DN ambisense RNA	Koch's Postulates for viruses, virused DNA (dsDNA), gapped DNA (A) genomes, double stranded RNA (ssRNA), (+) strand RNA, single stranded intermediate, single stranded genomes.	us cycle, studying us genome types, genomes, single- (dsRNA), single tranded (+) sense RNA (-) sense,	12 Hours
UNIT II	Triangulation into the nucleoncentrating concentrating concentrations and concentrations are concentrations.	and Assembly: Metastability, the ogy. Helical symmetry, Icosah number, Quasi-equivalence. Virus of infection, Cellular receptor for leus, virus disassembly, metast omponents for assembly, getting the iruses make sub-assemblies, sequent taging signals, packaging of segregations.	attachment and viruses. Getting table structures, nings to the right	12 Hours
UNIT III	acquisition of a	n envelope, budding strategies. NA synthesis, Reverse Transcription		



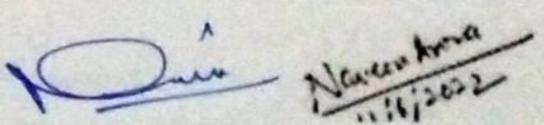
	synthesis in retroviruses. Regulation of translation in virus infected cells. Basic rules of genome replication in DNA viruses, viral origins of DNA replication. Generic steps in transcription, host polymerases, initiation, splicing, alternate splicing, promoter structure, steps in regulation of transcription, enhancers, virus coded transcriptional regulators, transcriptional cascade, export.	
UNIT IV	Virus Infections basics, interaction with host, acute and persistent infections: Fundamental questions of viral pathogenesis. Virion defenses to hostile environment, viral spread, viremia, determinants of tissue tropism. Virus shedding, transmission of infection, host defense, innate immune response, virus virulence, identifying virulence genes. Toxic viral proteins, cellular virulence genes, immunopathology, systemic inflammatory response syndrome. Immune complexes, virus induced auto-immunity, general pattern of infection. Inapparent acute infections, defense against the acute infection. Influenza, Polio, Measles, Rotavirus, persistent infections, chronic and latent Infections.	12 Hours
UNIT V	Anti-Viral drugs, virus evolution and emerging viruses: Anti-viral drugs, search for anti-viral drugs, the quasi-species concept, error threshold, genetic bottlenecks, Muller ratchet, genetic shift and drift. Theories on origin of virus, evolution of new viruses, emerging viruses.	12 Hours
Suggested Readings	 Principles of Virology: Molecular Biology, Pathogenesis and Control Animal Virusesby S.J. Flint, L.W. Enquist, V.R. Racaniello, A.M. 4th edition. ASM Press. 2015. Introduction to Modern Virology by N. Dimmock, A. Easton Leppard. 7th edition. Blackwell Publishing. 2016. Basic Virology by Edward K. Wanger, M. Hewiett, D. Bloom, Camerini. 3th edition. Blackwell Publishing. 2007. Principles of Molecular Virology by A.J. Cann. 6th edition. Elsevier Acad 2015. e protein-only hypothesis. 	I. Skalka. , K. D.

		Programme/Class: Bachelor's Degree with Research M. Sc. Microbiology (I)	h /	
Subject: Mic	crobiology	Year: First (1)	Comment	
Core Paper (C		Course Code: B080802T	Semester: Se IMMUNOL	OGY &
Marks		75 (UE) + 25 (CIE)	IMMUNOTECI	
otal Number	of Lectures-Tuto	rials-Practical (in hours per weel	Credits:	TOTAL SALES OF SALES
Course Objectives	immune system system of the	of this course is to understand the their structure and organization, and body. It would also make the stu- which underlie the host defen-	dents understand the	s the defense
Course Learning Outcomes		ul completion of the course, the stud	lent:	
COI	Will be able to response.	o understand the fundamental base:	of immune system a	and immune
CO2	Will be able to	o gather information about the stru f the immune system and Immunolog	cture and organization	n of various
CO3	Will be able	to understand the genetic organ mmune cell receptors and the bases	ization of the genes	meant for
CO4	Will be able immune respo	to understand the operation and the	mechanisms which	underlie the
CO5	Will be able to	o apply the knowledge gained to un sensitivity (allergy), organ transplan	derstand the phenome tation and certain im	ena like host munological
Contents				Duration: 60 hours
UNITI	(b) concept of structure of ly specificity, di T lymphocyte Basophils, Ne alternative pa		on, B lymphocytes, cells, Eosinophils, stem: classical and	12 Hours
UNITII	Superantigen, and variable antibody into antibody, pri acidity, equili complement immunofluor	rescence, biotin-avidin assay.	body – (i) constant idiotype. Antigen n of antigen and ibody affinity and utination reactions, immunoblotting,	12 Hours
UNIT III	Generation of concept of art antigen: speciarrangement	of diversity in immune response: clone ntigen specific receptor, BCR, TCR, ecific receptors on T and B lyne nts, class switch, comparison of rec	the genes encoding aphocytes, genetic ceptor on B and T	12 Hours
	immunologic	, mechanism of immune response cal diversity of MHC genes and products in immu		



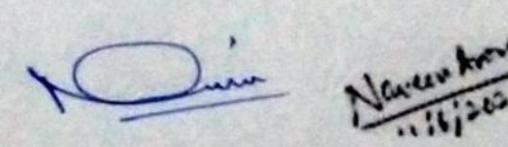
the property of the property of the property of the property of		
UNIT V	recognition of antigen and MHC products, structure of MHC gene complex, polymorphism of MHC genes and products. Graft rejection and GVHD; HLA-matching; Use of CRISPR-Cas for generating transgenic animals for xenotransplantation, Activation of T and B cells by antigen; Antigen processing, antigen presentation on T cells, products and factors released by T cell activation; interleukins, interferons. Cell mediated cytotoxicity, mechanism of T cell and NK cell mediated lysis, ADCC, macrophage cytotoxicity. Monoclonal antibody: production, application. Immunodeficiency: T cell, B cell, combined B and T cell deficiencies, defect in phagocytes and complement components, secondary immunodeficiency, AIDS, Autoimmunity. Immunization: active and passive, Vaccines- types and importance, Tumor antigens, immune response to tumors and immunotherapy of tumors	12 Hours
Suggested Readings	 Kuby Immunology by J.A. Owen, J. Punt, S.A. Stranford. 7th editi Freeman.2013. Cellular and Molecular Immunology by A.K. Abbas, A.H. Lich Pillai. 9th edition.Saunders Elsevier. 2018. Janeway's Immunobiology by K. Murphy, W. Casey. 9th edition. Science Publishing.2017. Review of Medical Microbiology and Immunology by W.Levin 15thedition.LangePublication. 2018. Fundamental Immunology by W.E. Paul. 7th edition. Lippincott Wilkins. 2013. Roitt's Essential Immunology by P.J. Delves, S.J. Martin, D.R. B Roitt. 13thedition. Blackwell Publishing. 2017. 	tman, S. Garland son. illiams and

		Programme/Class: Bachelor's Degree with Research M. Sc. Microbiology (I)	h /	
Subject: M	icrobiology	Year: First (1)	Semester: Sec	ond (II)
	Compulsory)	Course Code: B080803T	RECOMBINA	
Core raper (Compaisory		TECHNOL	Control of the Contro
Mark	s:100	75 (UE) + 25 (CIE)	Credits:	04
Total Number	of Lectures-Tuto	rials-Practical (in hours per week	L-T-P: 4-0-0	.1 1
Course Objectives	familiar with the various applicate about the meth analyses of transwith how recommends	f this course is to make the student anipulate/ analyze DNA, RNA and protein emethods and protein entire tions of the polymerase chain reached to currently used to carry out generated and protein expression. In a binant DNA technology has been employed to carry of the polymerase chain reached to carry out generated and protein expression. In a binant DNA technology has been employed to carry out generated and protein expression.	ake and screen librar ake and screen librar ation. The student with mome-wide analyses the student will be mapped to the student will be mapped to the student will be mapped to the study of the student of the stud	ies, and the libe taught and global ade familiar
Course Learning Outcomes	Upon successfu	il completion of the course, the stud	ent:	
CO1		ar with the use of various cloning v		
CO2	genomic and cl	describe artificial transformations a DNA libraries.		
CO3	Will be able to	understand the Screening and char-	acterization of cloned	DNA.
CO4	Will have learn	nt about various types of PCR and the	neir applications.	
CO5	Will be aware proteomics, ge	of DNA sequencing, RNA Interference nomics, transcriptomics and metabo	nce with a brief accou lomics.	
Contents				Duration: 60 hours
UNITI	bacterial artifications shuttle vectors for recombinations	ystems, cloning vectors (plasmids, cial chromosomes and yeast artifics, expression vectors, screening and ints. HACS. Enzymes used for nucleases, methylases, polymeras	selection methods nanipulating DNA	12 Hours
UNIT II	Preparation of DNA (plasmic	competent cells and their transform d, cosmid, phage and genomic DN nd eukaryotes. Construction of ge	A) and RNA from	12 Hours
UNIT III	Restriction m Northern Hy homologous E	apping and RFLP analysis. South bridization probe preparation, expression of cloned genes in cultudes probes. <i>In situ</i> hybridization. Anti-	heterologous and red cells, synthetic	12 Hours
UNITIV	PCR and its interaction: g protein interaction gene targeting	application. Site directed mutageneral mobility shift assay, DNA footion. Principles and method of general Real time PCR.	etic engineering and	12 Hours
UNIT V	DNA seque	ncing: Sanger's Method, Autor of recombinant DNA technology in	nated sequencing. agriculture, health	12 Hours



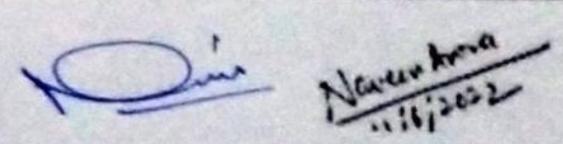
	and industry. RNA Interference. Brief account of proteomics, genomics, transcriptomics and metabolomics.
Suggested Readings	 Molecular Biology by D.P. Clarke, N. Pazdernik. 2nd edition. Academic Press. 2012. Molecular Cloning: A laboratory manual by J. Sambrook, D. Russell. 4th edition. ColdSpring Harbor laboratory Press. 2012. DNA Technology: The Awesome Skill by I. Edward Alcamo. Harcourt Academic Press. 2001. Molecular Biology of the Gene by J. Watson, T. Baker, S. Bell, A. Gann, M. Levine, R. Losick. 7th edition. Pearson. 2014. Gene Cloning and DNA Analysis: An Introduction by T.A. Brown. 7th edition. Wiley-Blackwell Publishers. 2016.

Subject: Microbiology Major Elective (Optional) Course Code: B080804T Marks: 100 Total Number of Lectures-Tutorials-Practical (in hours per week L-T-P: 4-0-0 Course Objectives To introduce the student to the variety of biophysical and biochem currently available to probe the structure and function of macromolecules, make them aware of the physical principles behind and the instrumentation involved, make them familiar with vario analyzing the output data, and to build a strong foundation in the area structure, division, survival and propagation. Course Learning Outcomes COI Be able to carry out the analysis of the data from CD and Fluorescence to monitor the stability of the protein under different environmental of the monitor than the databases. CO3 Be able to evaluate the quality and highlights of the structure reported journals/structural databases. CO4 Be able to design a multi-step purification protocol for a target protein PAGE under native and SDS conditions Will be aware of the use of tracer techniques and safety precautions Contents: UNIT I Spectroscopy: Biological application and interpretations of Nuclear Magnetic Resonance (NMR) & Electron Spin Resonance (ESR)., Absorption spectroscopy, Infrared and Raman spectroscopy, Optical Rotatory Dispersion (ORD), Circular Dichroism (CD)., Basics of X-ray Crystallography. UNIT II Basics principles and applications of various chromatography methods: Partition and Absorption chromatography, gel filtration, ion-exchange and affinity chromatography. Biological applications of HPLC. UNIT III Basics of centrifugation based methods: viscosity, diffusion,		1/	Programme/Class: Bachelor's Degree with Researc M. Sc. Microbiology (I)		
Marks: 100 75 (UE) + 25 (CIE) Credi Total Number of Lectures-Tutorials-Practical (in hours per week L-T-P: 4-0-0 Course Objectives To introduce the student to the variety of biophysical and biochem currently available to probe the structure and function of macromolecules, make them aware of the physical principles behind and the instrumentation involved, make them familiar with vario analyzing the output data, and to build a strong foundation in the area structure, division, survival and propagation. Course Learning Outcomes COI Be able to carry out the analysis of the data from CD and Fluorescence to monitor the stability of the protein under different environmental or to monitor the stability of the protein under different environmental or be able to evaluate the quality and highlights of the structure reported journals/structural databases. CO3 Be able to design a multi-step purification protocol for a target protein PAGE under native and SDS conditions CO5 Will be aware of the use of tracer techniques and safety precautions Co65 Will be aware of the use of tracer techniques and safety precautions Co7 Spectroscopy: Biological application and interpretations of Nuclear Magnetic Resonance (NMR) & Electron Spin Resonance (ESR), Absorption spectroscopy, Infrared and Raman spectroscopy, Optical Rotatory Dispersion (ORD), Circular Dichroism (CD)., Basics of X-ray Crystallography. UNIT II Basics principles and applications of various chromatography methods: Partition and Absorption chromatography, Biological applications of HPLC. UNIT III Basics of centrifugation based methods: viscosity, diffusion, of HPLC.	econd (II)	Semester: See		icrobiology	Subject: Mi
Total Number of Lectures-Tutorials-Practical (in hours per week L-T-P; 4-0-0 Course Objectives To introduce the student to the variety of biophysical and biochem currently available to probe the structure and function of macromolecules, make them aware of the physical principles behind and the instrumentation involved, make them familiar with vario analyzing the output data, and to build a strong foundation in the area structure, division, survival and propagation. Course Learning Outcomes COI Be able to carry out the analysis of the data from CD and Fluorescence to monitor the stability of the protein under different environmental or monitor the stability of the protein under different environmental or journals/structural databases. CO3 Be able to design a multi-step purification protocol for a target protein PAGE under native and SDS conditions CO4 Be able to understand and correctly interpret the migration of protein PAGE under native and SDS conditions CO5 Will be aware of the use of tracer techniques and safety precautions CO6 Contents: UNIT I Spectroscopy: Biological application and interpretations of Nuclear Magnetic Resonance (NMR) & Electron Spin Resonance (ESR)., Absorption spectroscopy, Infrared and Raman spectroscopy, Optical Rotatory Dispersion (ORD), Circular Dichroism (CD)., Basics of X-ray Crystallography. UNIT II Basics principles and applications of various chromatography methods: Partition and Absorption chromatography, gel filtration, ion- exchange and affinity chromatography. Biological applications of HPLC. UNIT III Basics of centrifugation based methods: viscosity, diffusion,	TICAL QUES	701067			
To introduce the student to the variety of biophysical and biochem currently available to probe the structure and function of macromolecules, make them aware of the physical principles behind and the instrumentation involved, make them familiar with vario analyzing the output data, and to build a strong foundation in the area structure, division, survival and propagation. Course Learning Outcomes COI Be able to carry out the analysis of the data from CD and Fluorescence to monitor the stability of the protein under different environmental of the monitor the stability of the protein under different environmental of journals/structural databases. CO3 Be able to design a multi-step purification protocol for a target protein PAGE under native and SDS conditions Will be aware of the use of tracer techniques and safety precautions Cotontents: UNIT I Spectroscopy: Biological application and interpretations of Nuclear Magnetic Resonance (NMR) & Electron Spin Resonance (ESR)., Absorption spectroscopy, Infrared and Raman spectroscopy, Optical Rotatory Dispersion (ORD), Circular Dichroism (CD)., Basics of X-ray Crystallography. UNIT II Basics principles and applications of various chromatography methods: Partition and Absorption chromatography, gel filtration, ion- exchange and affinity chromatography. Biological applications of HPLC. UNIT III Basics of centrifugation based methods: viscosity, diffusion,	: 04	Credits:			
To introduce the student to the variety of biophysical and biochem currently available to probe the structure and function of macromolecules, make them aware of the physical principles behind and the instrumentation involved, make them familiar with vario analyzing the output data, and to build a strong foundation in the area structure, division, survival and propagation. Course Learning Outcomes COI Be able to carry out the analysis of the data from CD and Fluorescence to monitor the stability of the protein under different environmental of the monitor the stability of the protein under different environmental of pournals/structural databases. CO3 Be able to design a multi-step purification protocol for a target protein PAGE under native and SDS conditions Will be aware of the use of tracer techniques and safety precautions CO5 Will be aware of the use of tracer techniques and safety precautions Contents: UNIT I Spectroscopy: Biological application and interpretations of Nuclear Magnetic Resonance (NMR) & Electron Spin Resonance (ESR)., Absorption spectroscopy, Infrared and Raman spectroscopy, Optical Rotatory Dispersion (ORD), Circular Dichroism (CD)., Basics of X-ray Crystallography. UNIT II Basics principles and applications of various chromatography methods: Partition and Absorption chromatography, gel filtration, ion- exchange and affinity chromatography. Biological applications of HPLC. UNIT III Basics of centrifugation based methods: viscosity, diffusion,		L-T-P: 4-0-0	ials-Practical (in hours per weel	of Lectures-Tutor	Total Number
Course Learning Outcomes COI Be able to carry out the analysis of the data from CD and Fluorescence to monitor the stability of the protein under different environmental or monitor the stability of the protein under different environmental or monitor the stability of the protein under different environmental or monitor the stability of the protein under different environmental or Be able to evaluate the quality and highlights of the structure reported journals/structural databases. CO3 Be able to design a multi-step purification protocol for a target protein PAGE under native and SDS conditions CO5 Will be aware of the use of tracer techniques and safety precautions Contents: UNIT I Spectroscopy: Biological application and interpretations of Nuclear Magnetic Resonance (NMR) & Electron Spin Resonance (ESR)., Absorption spectroscopy, Infrared and Raman spectroscopy, Optical Rotatory Dispersion (ORD), Circular Dichroism (CD)., Basics of X-ray Crystallography. UNIT II Basics principles and applications of various chromatography methods: Partition and Absorption chromatography, gel filtration, ion-exchange and affinity chromatography. Biological applications of HPLC. UNIT III Basics of centrifugation based methods: viscosity, diffusion,	ne biological ach technique s methods of	ical and biochemica d function of the principles behind eac miliar with various	the student to the variety of biophy able to probe the structure a make them aware of the physical mentation involved, make them fat the total total and to build a strong four	To introduce the currently availar macromolecules and the instrumanalyzing the out	Course
to monitor the stability of the protein under different environmental colors. CO2 Be able to evaluate the quality and highlights of the structure reported journals/structural databases. CO3 Be able to design a multi-step purification protocol for a target protein PAGE under native and SDS conditions CO5 Will be aware of the use of tracer techniques and safety precautions Contents: UNIT I Spectroscopy: Biological application and interpretations of Nuclear Magnetic Resonance (NMR) & Electron Spin Resonance (ESR)., Absorption spectroscopy, Infrared and Raman spectroscopy, Optical Rotatory Dispersion (ORD), Circular Dichroism (CD)., Basics of X-ray Crystallography. UNIT II Basics principles and applications of various chromatography methods: Partition and Absorption chromatography, gel filtration, ion- exchange and affinity chromatography. Biological applications of HPLC. UNIT III Basics of centrifugation based methods: viscosity, diffusion,		ent will:			Learning
Be able to evaluate the quality and highlights of the structure reported journals/structural databases. Be able to design a multi-step purification protocol for a target protein Be able to understand and correctly interpret the migration of protein PAGE under native and SDS conditions Will be aware of the use of tracer techniques and safety precautions Cost Will be aware of the use of tracer techniques and safety precautions Contents: UNIT I Spectroscopy: Biological application and interpretations of Nuclear Magnetic Resonance (NMR) & Electron Spin Resonance (ESR)., Absorption spectroscopy, Infrared and Raman spectroscopy, Optical Rotatory Dispersion (ORD), Circular Dichroism (CD)., Basics of X-ray Crystallography. UNIT II Basics principles and applications of various chromatography methods: Partition and Absorption chromatography, gel filtration, ion- exchange and affinity chromatography. Biological applications of HPLC. UNIT III Basics of centrifugation based methods: viscosity, diffusion,	experiments iditions	D and Fluorescence e t environmental cond	out the analysis of the data from C ability of the protein under differen	Be able to carry to monitor the st	CO1
Be able to understand and correctly interpret the migration of protein PAGE under native and SDS conditions Will be aware of the use of tracer techniques and safety precautions Contents: UNIT I Spectroscopy: Biological application and interpretations of Nuclear Magnetic Resonance (NMR) & Electron Spin Resonance (ESR)., Absorption spectroscopy, Infrared and Raman spectroscopy, Optical Rotatory Dispersion (ORD), Circular Dichroism (CD)., Basics of X-ray Crystallography. UNIT II Basics principles and applications of various chromatography methods: Partition and Absorption chromatography, gel filtration, ion- exchange and affinity chromatography. Biological applications of HPLC. UNIT III Basics of centrifugation based methods: viscosity, diffusion,	Be able to evaluate the quality and highlights of the structure reported/deposited in		Be able to evalu	CO2	
PAGE under native and SDS conditions Will be aware of the use of tracer techniques and safety precautions Contents: UNIT I Spectroscopy: Biological application and interpretations of Nuclear Magnetic Resonance (NMR) & Electron Spin Resonance (ESR)., Absorption spectroscopy, Infrared and Raman spectroscopy, Optical Rotatory Dispersion (ORD), Circular Dichroism (CD)., Basics of X-ray Crystallography. UNIT II Basics principles and applications of various chromatography methods: Partition and Absorption chromatography, gel filtration, ion- exchange and affinity chromatography. Biological applications of HPLC. UNIT III Basics of centrifugation based methods: viscosity, diffusion,	Be able to design a multi-step purification protocol for a target protein		Be able to desig	CO3	
UNIT I Spectroscopy: Biological application and interpretations of Nuclear Magnetic Resonance (NMR) & Electron Spin Resonance (ESR)., Absorption spectroscopy, Infrared and Raman spectroscopy, Optical Rotatory Dispersion (ORD), Circular Dichroism (CD)., Basics of X-ray Crystallography. UNIT II Basics principles and applications of various chromatography methods: Partition and Absorption chromatography, gel filtration, ion- exchange and affinity chromatography. Biological applications of HPLC. UNIT III Basics of centrifugation based methods: viscosity, diffusion,	Be able to understand and correctly interpret the migration of protein molecule on PAGE under native and SDS conditions			CO4	
UNIT I Spectroscopy: Biological application and interpretations of Nuclear Magnetic Resonance (NMR) & Electron Spin Resonance (ESR)., Absorption spectroscopy, Infrared and Raman spectroscopy, Optical Rotatory Dispersion (ORD), Circular Dichroism (CD)., Basics of X-ray Crystallography. UNIT II Basics principles and applications of various chromatography methods: Partition and Absorption chromatography, gel filtration, ion- exchange and affinity chromatography. Biological applications of HPLC. UNIT III Basics of centrifugation based methods: viscosity, diffusion,		fety precautions	f the use of tracer techniques and sa	Will be aware or	CO5
Magnetic Resonance (NMR) & Electron Spin Resonance (ESR)., Absorption spectroscopy, Infrared and Raman spectroscopy, Optical Rotatory Dispersion (ORD), Circular Dichroism (CD)., Basics of X- ray Crystallography. UNIT II Basics principles and applications of various chromatography methods: Partition and Absorption chromatography, gel filtration, ion- exchange and affinity chromatography. Biological applications of HPLC. UNIT III Basics of centrifugation based methods: viscosity, diffusion,	Duration: 60 hours				Contents:
methods: Partition and Absorption chromatography, gel filtration, ion- exchange and affinity chromatography. Biological applications of HPLC. UNIT III Basics of centrifugation based methods: viscosity, diffusion,	12 Hours	esonance (ESR)., troscopy, Optical	nance (NMR) & Electron Spin R troscopy, Infrared and Raman spe- sion (ORD), Circular Dichroism (C	Absorption special Rotatory Disper	UNIT I
	12 Hours	hy, gel filtration,	on and Absorption chromatograp	methods: Partiti	UNIT II
sedimentation equilibrium, dialysis, solvent fractionation, centrifugation, Biological applications and interpretations of Density Gradient methods, Ultracentrifugation methods	12 Hours	t fractionation,	equilibrium, dialysis, solver Biological applications and interpre	Basics of cen sedimentation centrifugation, E	UNIT III
UNIT IV Basics of electrophoresis: electrophoretic mobility and affecting factors, Biological applications and interpretation of different types of electrophoresis: PAGE, gradient gel, Agarose Gel Electrophoresis, 2D Electrophoresis, iso-electric focusing	12 Hours	different types of ectrophoresis, 2D	al applications and interpretation of PAGE, gradient gel, Agarose Gel E iso-electric focusing	factors, Biologic electrophoresis: Electrophoresis,	UNIT IV
UNIT V Radioactive methods: Basics of radioactive isotopes and radioactive decay, sample preparation, counting, Safety precautions during handling, biological applications.	12 Hours	Radioactive methods: Basics of radioactive isotopes and radioactive lecay, sample preparation, counting, Safety precautions during		decay, sample	UNIT V



Readings:	McGrawHill.1994. 2. Principles of Fluorescence Spectroscopy by J. Lakowicz, R. Joseph. 2 nd edition.Springer.1999.
	3. Molecular Fluorescence: principles and Applications by B. Valeur. 2 nd edition. Wiley.2013.
	4. NMR - Conformation of Biological Molecules by G. Govil, R.V.
	Hosur. 1st edition.Springer- Verlag, 2011. 5. Biomolecular crystallography: Principles, practice and application to
	structural biology by B. Rupp. 1 st edition. Garland Science. 2009. 6. Optical methods in Biology by E.M. Slayter. 1 st edition. John Wiley. 1970.
	7. NMR of proteins and nucleic Acids by K. Wuthrich. Ist edition. Wiley IntersciencePublications, 1988.
	Biophysical chemistry, Part 2: Techniques by C. R. Cantor, P. R. Schimmel. 1st edition, W.H Freeman and Co. 2008.

		Programme/Class: Bachelor's Degree with Research M. Sc. Microbiology (I)		
Subject: 1	Microbiology	Year: First (1)	Semester: Se	cond (II)
Major Elective (Optional)		Course Code: B080805T	EXTREMOPH THEIR APPLI	
Ma	rks:100	75 (UE) + 25 (CIE)	Credits	The state of the s
		ials-Practical (in hours per week I	-T-P: 4-0-0	
Course Objectives	which microorg extremophiles in about the applica era in the bio understanding for	anisms adopt to extreme environthe evolution related to the origin of ation of extremophiles in the industrial technology. The study of extremor astrobiology that will help to uny bodies in our own solar system and	nderstanding about no nments and the cr of life. The students ial processes that has ne environment will inderstand what form	itical role of will also learn opened a new I develop an
Course Learning Outcomes		completion of the course, the studen		£
COI	extremophiles in	ed to extremophiles and will underst the evolution related to the origin of	f life.	
CO2		Will develop an understanding about mechanisms by which thermophiles and psychrophiles adopt to extreme environments.		and
CO3	Will develop an understanding about mechanisms by which Halophiles Acidophiles and Alkaliphiles: adopt to extreme environments.		cidophiles	
CO4		the application of extremophiles in t	the industrial process	es.
CO5		erstand what form life takes on anoth		
Contents				Duration: 60 hours
UNIT I	and general hyperthermophi	extremophiles and Origin of Life; Iso properties of extremophiles I les, psychrophiles, halophiles, acide philes, Natural habitats of extremoph	like thermophiles, ophiles, alksliphiles	12 Hours
UNIT II	High Temperate The Physiologic Solutes from (H	Microbial Life at high temperature vival: Membrane Adaptations of (Hypures, Temperature-Dependent Mole al Role, Biosynthesis, and Mode of Ayper); Psychrophiles Mechanism of ure, Membrane Adaptations, Colder, Perception and Transduction of L	per)Thermophiles to ecular Adaptations, action of Compatible bacterial adaptation Adapted, The Cold-	12 Hours
UNIT III	a Haloarchaeal Regulatory M Physiology and	diversity in Highly Saline, Response Genome: a Role for General Stress lechanisms; Acidophiles Acidur Ecology of Acidophilic Microorgan laptations that Support Alkaliphily.	Proteins and Global ric Proteobacteria,	12 Hours
UNIT IV	Radiation-resist	ant extremophiles and their potential Exobiology: Astrobiology and the S	in biotechnology	12 Hours
UNIT V	Extremophiles a	as a source of novel enzymes for in	dustrial application,	12 Hours



	Versatile applications of natural compounds from extremophiles, Polysaccharides from extremophilic microorganisms. Importance of extremophilic microbial diversity in environment, pharmaceuticals & human health,
Suggested Readings	 Extremophiles: From Biology to Biotechnology, Edited by- Ravi Durvasula and D. V. Subba Rao, CRC Press, Taylor & Francis Group, ISBN 9781498774925 Physiology and biochemistry of extremophiles / Edited by C. Gerday and N. Glansdorff, ASM Press, American Society for Microbiology, ISBN-10: 1-55581-422-0

Core Pape	Microbiology	M. Sc. Microbiology (I) Year: First (1)	Company Control (197)
Core Pape	Company of the Compan	I car. First (1)	Semester: Second (II)
CONTRACTOR AND ADMINISTRATION OF THE PARTY O	r (Compulsory)	Course Code: B080806P	Practical II
MI	rks:100	Credits: 04	Duration: 120 hours
75 (UE) + 25 (CIE)			
Annual Control of the	A STATE OF THE PARTY OF THE PAR	s-Practical (in hours per week L-T	-P: 0-0-8
Course Objectives	The course will ena	ble students to learn basic techniques tudents will also explore the immunol	used in separation and analysis of
Course Learning Outcomes	Upon successful co	mpletion of the course, the student:	
COI:	Will be able to use	chromatographic and centrifugation p	rocedures.
CO2:		electrophoretic techniques.	
CO3:	Will be aware of va	rious immunological techniques	
CO4:	Will be able to make and transform desired plasmid DNA into bacterial cells along with to other techniques used in cloning and rDNA technology.		
CO5:	Will be able to isolate Bacteriophages		
2. Separation 3. Column 4. Agurose 5. SDS-PA 6. Double in 7. Determin 8. Isolation 9. Determin 10. Immunol 11. Raising of 12. ELISA 13. Isolation 14. Compete 15. Transfer	on of amino acids by Thi chromatography gel electrophoresis for s GE for separation of Pro mmone diffusion nation of Blood group of Macrophages sation of hypersensitivity bloting of Ab in mice/rabbit.	lasmid DNA, genomic DNA and RNA E. coli transformant of E. coli cells using anti enzyme activity	

4º edition. Cold Spring Harbor laboratory Press. 2012.

2. Molecular Cloning: A laboratory manual by Joseph Sambrook, David Russell,

Pearson. 2017.

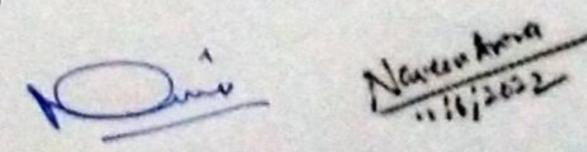
Readings:

		Programme/Class: Bachelor's Degree with Research M. Sc. Microbiology (I)	
Subject: 1	Microbiology	Year: First (1)	Semester: Second (I)
Subject: Microbiology Core Paper (Compulsory)		Course Code: B080807R	Industrial Training /Surveys/Research Project II
Marks:100			lits: 04
Course Details	this response project can a		etc.
	* Students will sul	bmit the final report (project report) the semesters at the end of the year,	dissertation) of the research project which will be assessed jointly by the university at the end of the year out

		Programme/Class: M. Sc. Microbiology (II)			
	Microbiology	Year: Second (2)	Semester: Tl	nird (III)	
Core Paper (Compulsory)		Course Code: B080901T	MICROI PHYSIOLOG METABO	GY AND	
Ma	rks:100	75 (UE) + 25 (CIE)	Credits:	04	
Total Numb		rials-Practical (in hours per week l		4	
Course Objectives	relation to its sur courses taught late	ye of this paper is to develop clear ungy along with diverse metabolic provided and propagation, and to enabor such as Microbial Pathogenicity and	athways existing in le students to better	bacteria in understand	
Course Learning Outcomes		ompletion of the course, the student:			
CO1		d with methods of measuring micro with understanding of steady state and			
CO2		n in-depth knowledge of phototrophic			
CO3	differences with conditions. This a	entral metabolic pathways for carbon eukaryotic systems and their regul llows students to apply the acqui is fordeveloping industrially useful str	lation in diverse p red knowledge in	hysiological	
CO4	Will have gathered	l understanding of inorganic and organows role of glutathione in cellular re	nic nitrogen assimila	ation and its iochemistry	
CO5	Will understand de along with biocher	tails of lipid and nucleotide metabolis	sm in E. coli and its	regulation	
Contents				Duration: 60 hours	
UNIT I	Membrane transpo channels, ABC Catabolite repressi	ntroduction, primary and secondary to ort proteins: porins and aquaporins, a transporter, group translocation P on, inducer exclusion and expulsion	mechanosensitive EP-PTS system.	12 Hours	
UNIT II	properties of pig photosystems, p cyanobacterial pl hydrogen-, iron- ar	green and purple bacteria, structura ment, oxygenic and anoxygenic hotodynamic death and photo notosynthesis, photorespiration C d sulfur, bacteria, methanogens and r	photosynthesis, ophosphorylation. themolithotrophy, nethylotrophs.	12 Hours	
UNIT III	Central Metabolic I Gluconeogenesis, Pathway, Citric Ad regulation. Examp pathways to develo	Pathways and Regulation: Glycolysis a Pentose-Phosphate Pathway, I id Cycle, alternate TCA, Glyoxylate les of pathway engineering of compine pindustrial useful strains: Co-metabonic and citric acid production.	and its regulation, Entner-Doudoroff Pathway and its earbon metabolic	12 Hours	
UNIT IV	Biochemistry of ni gene and genetic r regulation of nitrog	trogenase complex, nitrogenase types egulation of nitrogenase, symbiotic re enase by oxygen and combined N-so ainst oxygen, nitrate reduction (a	nitrogen fixation, ources, protection	12 Hours	

	dissimilatory) and sulfate reduction, methanogenesis and acetogenesis. Hydrocarbon transformation	
UNITV	Metabolism of lipids and nucleotides: Biosynthesis and degradation of lipids and its regulation in <i>E. coli</i> , lipid accumulation in yeast. Purine and pyrimidine biosynthesis, deoxyribonucleotide synthesis, regulation of purine and pyrimidine biosynthesis, inhibitors of pucleotide biosynthesis.	12 Hours
Suggested Readings	 Biochemistry by Geoffrey L. Zubay. 4th Edition. Brown Co, USA. 1th Lord Distriction Physiology by A.G. Moat, J. W. Foster, M. P. Spector. 3th John Wiley& Sons. 2002 Lehninger Principles of Biochemistry by D. L. Nelson, M. M. Cox. of Edition. W. H.Freeman. 2012 The Physiology and Biochemistry of Prokaryotes by D. White, J. Dru C. Fuqua. 4th Edition. Oxford University Press. 2011. Microbial Biochemistry by G. N. Cohen. 2nd Edition. Springer. 2016. Lippincott's Illustrated Reviews: Biochemistry edited by D. R. Fer Edition. Lippincott Williams & Wilkins. 2013 Biochemical Calculations: by Irwin H. Segel. 2nd Edition. Wiley. 2008. Understanding Enzymes by T. Palmer, E. Horwood. 3th Edition. Wiley. 	Edition. 6th ummond, 4. rier. 6th

		Programme/Class: M. Sc. Microbiology (II)			
Subject: N	Aicrobiology	Year: Second (2)	Semester: Thi	rd (III)	
A SHARL SHAR	Core Paper (Compulsory) Course Code: B080902T INDU		INDUSTR MICROBIO	USTRIAL	
Marks:100		75 (UE) + 25 (CIE)	Credits:	04	
		rials-Practical (in hours per week)	L-T-P: 4-0-0		
Course	The course will e	nable students to apply the learning	of microbiology conc	epts toward	
Objectives	for development product recovery	of microbial population for industrial a of microbial strains, process optimize will be covered for industrially a ins. Acquires knowledge about the	ation, large scale pro- elevant microbial pr	duction and oducts and	
Course Learning Outcomes	Upon successful	completion of the course, the student			
COI		the biochemical and industrial con of fermentation systems used in the f			
CO2		Will attain knowledge about designing of industrial strains and various media optimizationstrategies. Develop an understanding about design and use of various types			
CO3	Will acquire kno microrganisms	wledge about various food products	by the application	of	
CO4	The state of the s	wledge about various pharmaceutica	l products by the ap	plication	
CO5	Will understand	the production of commercial product	s by recombinant mic	roorganism	
Contents				Duration: 60 hours	
UNIT I	Ideal bioreactor,	ction to the fermentation; Introduct Reactor with non-ideal mixing. Multi cell reactor technology.		12 Hours	
UNIT II	metabolites. St recombination an	ning for new metabolites - primar rain development through selected d other genetic and biochemical methods pes and availability.	ection, mutation,	12 Hours	
UNIT III	Production of ale	immobilization technology for en cohol (ethanol), Organic acid (citric e, glutamic acid), nucleotides and rel	acid. lactic acid),	12 Hours	
UNIT IV	Production of e microbial food, antibiotics (strep) vitamins, steroids	enzymes (protease, amylase, lipase single cell protein and mushroom tomycin, tetracycline, penicillin, amp s and alkaloids.	e), Production of m. Production of icillin), hormones,	12 Hours	
UNIT V	hormones, interfe	mmercial products by recombinant onucleases, biopolymers, human eron and vaccines. Microorganisms in disteroids; Microorganisms as biosens	insulin, growth biotransformation fors & biochips	12 Hours	
Suggested Readings	2. Biopro edition	les of Fermentation Technology by Independent des dition. Butterworth-Heinemann. 2006 cess Engineering: Basic Concepts by PearsonEducation India. 2015. Industrial Microbiology & Biotech	016. M. L. Shuler, F. Ka	argi, 2 nd	



edition. CRC Press, USA. 2007.

4. Fermentation Microbiology and Biotechnology edited by E.M.T. El-Mansi, C.F. Bryce,
A.L. Demain, A.R. Allman. 3rd edition. CRC Press. 2012.

Microbial Biotechnology: Fundamentals of Applied Microbiology by A.N. Glazer, H.Nikaido. 2nd edition. Cambridge University Press. 2007.

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		Programme/Class:			
Bublanti N	Alarabialagy	M. No. Microbiology (11) Year: Second (2)	Semester: Thir	d (III)	
Core Paper (Compulsory)		Course Code: B080903T	ENVIRONMENTA MICROBIOLOG		
Marks; 100		75 (UE) + 25 (CIE)	Credits; 0	Charles and the same of the same	
Fotal Numbe	r of Lectures-Tuto	rials-Practical (in hours per week l	T-P: 4-0-0	135965	
Course Objectives	The major object andfunctioning o population in agr	ive of this paper is to impart knowle f microbial communities of diverse en loulture, mineral recovery, managem processes of various types of wastes in processes of various types of wastes in the contract of the contra	dge about structure, convironment. The use of ent of various types of	f microbia pollutant:	
Course Learning Outcomes	The state of the s	completion of the course, the student:			
coi	Will have an ov microbiology wi pollution.	erview of the till date development th special emphasis on the role of mic	s in the field of envi crobes in mitigating en	ronmental vironment	
CO2	The state of the s	me acquainted with various cultur in understanding microbial diversity.	al, biochemical and	molecular	
CO3		Will be able to describe the role of soil microbes in nutrient transformation, plant- microbe interactions and biotechnology. Also knows about potability of water and its			
CO4	Is able to describ	e the role of microbes in solid and lic			
CO3	Understands the petroleum hydro	role of microbes in bioremediation ocarbons, pesticides, plastic and elected in mineral and oil recovery.	of environmental pollu	itants like	
Contents				Duration 60 hours	
UNITI	microbial ecolog	field of environmental microbiology and emergence of field of environmental microbes in solving environmental microbiology and environmental microbiology a	mental microbiology,	12 Hours	
UNITII	the environment Analysis by FAN analysis, slot-blo	nt and culture-independent approach ity in the environment: Understanding by culture-dependent and culture-index ME, measuring metabolic capabilities of hybridization of community DNA, a intact cells, metagenomic analysis	microbial diversity in ependent approaches, using BIOLOG, G+C and fluorescent in situ	12 Hours	
UNIT III	transformation pro	microbiology: Importance of soil mi ocesses, plant-microbe symbiosis, microbi nological applications, drinking water mi	al antagonism, biofilms	12 Hours	
UNITIV	Liquid and solid secondary and (distillery, textil (metals, sedime composting, land	d waste management: Treatment of tertiary treatments), treatment of interest and paper), methodsto detect ats, toxin and organic matters). So ifill development, incineration method riculture, biogas production, p	dustrial effluents various pollutants olid waste types,	12 Hours	

	microorganisms as a tool for bioremediation, challenges in waste management.
UNITV	Lignocellulolytic microorganisms, enzymes and their biotechnological applications in: biopulping, biobleaching, textiles biofuels, animal feed production. Bioremediation of environmental pollutants: Petroleum hydrocarbons and pesticides, use of biosensors for their detection. Microbial enhanced oil recovery, bioleaching of copper, goldand uranium, electronic waste management.
Suggested Readings	 Microbial Ecology by R.M. Atlas, R. Bartha. 3rd edition. Benjamin Cummings Publishing Co, USA. 1993. Environmental Microbiology by A.H. Varnam, M.G. Evans. Manson Publishing Ltd.2000. Manual of Environmental Microbiology edited by C.J. Hurst, R.L. Crawford J.L. Garland, D.A. Lipson, A. L. Mills, L.D. Stetzenbach. 3rd edition. Blackwell Publishing. 2007. Environmental Microbiology edited by R. Mitchell, J-D Gu. 2nd edition. Wiley-Blackwell.2009. Environmental Microbiology by R. Maier, I. Pepper, C. Gerba. 2nd edition. AcademicPress. 2009. Environmental Microbiology: Principles and Applications by P.K. Jjemba, SciencePublishing Inc. 2004. Lignocellulose Biotechnology: Future Prospects by R.C. Kuhad, A. Singh. I.K.International. 2007. Environmental Microbiology of Aquatic & Waste systems by N. Okafor. Istedition, Springer, New York. 2011.

		Programme/Class: M. Sc. Microbiology (II)			
Subject: Microbiology				hird (III)	
Major Elective (Optional) Marks:100		Course Code: B080904T	BIOSTATIST	emester: Third (III) IOSTATISTICS AND BIOINFORMATICS	
				edits: 04	
Total Numb	er of Lectures-Tuto	rials-Practical (in hours per week L			
Course Objectives	The course will introduce the student to the variety of computational methods currently available for predicting functional behavior of biological systems. The algorithms behind each method and the shortcomings in present methods will be discussed. Students should				
	be able to analyze	the output data to predict a biologicall	v relevant function	idents should	
		Section A: Biostatistics	y relevant function.		
Course Learning Outcomes	Upon successful c	ompletion of the course, the student;			
CO1	Will understand va	arious methods of collection and repres	sentation of biologic	cal data	
CO2	Will be able to understand the concepts of statistical population and samples and will become aware of Measures of Central tendencies and Dispersion.				
CO3	Will learn about sample size calculation and distribution. He/ She will also learn about principles of probability.				
CO4	Will be able to understand the concepts of correlation and regression.				
CO5	Will learn about ba	asic idea of significance,			
Contents:				Duration 30 hours	
UNIT I	Scope of biostatis tabulation and diag	stics, variables in biology. Collection grammatic presentation of statistical da	n, classification,	06 hours	
UNIT II	Concepts of statis	stical population and samples. Meas	sures of Central	06 hours	
UNIT III	Sample size calculation. Simple measure of Skewness and Kurtosis Probability: definition, simple theorems of probability and simple application of probability. Binomial and Poisson distributions.				
UNIT IV	Correlation, corre	relation, correlation coefficient, standard error of estimate and of hour ression. Linear regressions, least square method of fitting.			
UNIT V	Testing level of sig e.g. Chi square test	nificance, random variations. Statistica, students 't' test.	al analysis test	06 hours	
		Section B: Bioinformatics			
Course Learning Outcomes	Upon successful co	empletion of the course, the student:			
CO1	Will be able to gair	elementary knowledge of computers			
002	Will be able to gain elementary knowledge of computers Will be able to access and derive information from various primary and secondary databases				
203	Will be able to create and usefully interpret the results of a multiple sequence alignment.				
04	Can create and correctly interpret phylogenetic trees to gain insight into evolutionary pathof the target molecule				
05		various protein databases and will lear	n about primer des	ignine	
Contents			The des	Duration 30 hours	
JNITI		of applications of common spress, DOS and Windows based software		06 hours	

	office.			
UNIT II	Biological Databases: Introduction. Types of databases in terms of biological information content. Protein and gene information resources. Different formats of molecular biology data.			
UNIT III	Molecular Phylogenetics: Sequence Alignment: Methods and algorithms of pairwise and multiple sequence alignment. Global and local alignment. Alignment scoring matrices. Database similarity searching.			
UNIT IV	Methods and tools for phylogenetic analysis. Creation evaluation and interpretation of evolutionary trees. Advantages and disadvantages of phenetic and cladistic approaches.			
UNIT V	Protein database: Retrival of protein sequence from PDB, Primer designing.	06 hours		
Readings	 Haubold, Wiele. 1stedition. Springer International. 2006. Introduction to Bioinformatics by A. Lesk. 3rd edition. OUP India. 2009 Statistical methods in Bioinformatics: An introduction by W. Ewens, C. Grant. 2nd Edition. Springer-Verlag. 2006. Bioinformatics: Sequence and genome analysis by D. Mount. 2nd edition Spring HarborLab Press. 2004. Bioinformatics: A practical guide to the analysis of genes & proteins. by Baxevanis, Outlette. 2nd edition. John Wiley and Sons. 2001. An Introduction to Protein Informatics by K-H Zimmermann. 1st edition SpringerInternational. 2007. Fundamental Concepts of Bioinformatics by Krane. 1st edition. Pearson Education. 2003. Discovering Genomics, Proteomics and Bioinformatics by Campbell. edition. CampbellPearson Education. 2007. Structural bioinformatics: an algorithmic approach by F. J. Burkowski. 1 Chapman &Hall/CRC. 2009. Structural Bioinformatics edited by J. Gu, P.E. Bourne. 2nd Edition. Wile Blackwell. 2009. 	G.R. on. Cold Edited n, 2 nd st edition,		

		Programme/Class: M. Sc. Microbiology (II)			
Subject: Microbiology		Year: Second (2)	Semester: Thi	Semester: Third (III)	
Major Elective (Optional) Marks:100		Course Code: B080905T	MICROBIAL DIV	Control of the Contro	
		75 (UE) + 25 (CIE)		Credits: 04	
Total Numb	er of Lectures-Tutor	rials-Practical (in hours per we			
Course Objectives	The objective of the diversity in the mice	is course is to introduce the stude crobial world. The course will dev robial world and the power of the	ents with enormous range velop an understanding ab		
Course Learning Outcome s	Upon successful co	ompletion of the course, the stude	nt:		
COI	Will establish a po	int of view to examine microbial	diversity		
CO2		Vill learn how to construct and interpret evolutionary trees from DNA sequences.			
CO3	Will develop a ba	sic understanding about the maj	or branches of the "tree		
CO4	Will learn how new organisms are identified (usually without being cultivated) and will progress in steps to broad surveys of entire microbial communities.			d) and will	
CO5	Will understand ho	w specific kinds of organisms cor	ntribute to the ecosystem.		
Contents				Duration: 60 hours	
UNITI	Introduction to Microbial Diversity: Facets of microbial diversity, the fundamental similarity of all living things, Taxonomy and phylogeny, Phylogenetic Information, Obtaining the required sequence data, assembling sequences in a multiple-sequence alignment.				
UNIT II	Constructing a P Alternatives to Sn	Constructing a Phylogenetic Tree. Tree Construction Complexities, alternatives to Small-Subunit rRNA Analysis, SSU rRNA cannot be used to distinguish closely related organisms.			
UNIT III	General properti Phototrophic Bact Gram-Positive Bac Actinobacteria (h Bacteroids: Deine	les of Primitive Thermoph ceria: Proteobacteria (purple b cteria, Firmicutes (low G+C gr igh G+C gram-positive bacte ococci, Chlamydiae and Plan karyotic Microorganisms	acteria and relatives), ram-positive bacteria), ria): Spirochetes and	12 Hours	
UNIT IV	Microbial Populations: Identification of Uncultivated Organisms, Sequence-Based Microbial Surveys, Fluorescent In Situ Hybridization Surveys, Molecular Fingerprinting of Microbial Populations, Linking Phenotype and Phylotype.			12 Hours	
UNIT V	The Phylogenetic Perspective, Genomics, Comparative Genomics, and Metagenomics, Origins and Early Evolution, the timescale, Ancient microbial fossils, The last common ancestor, The RNA world hypothesis, The emergence of life.				
Suggested Readings	Principles of microbial diversity / James W. Brown, Department of Biological Sciences, North Carolina State University, Raleigh, North Carolina. ISBN 978-1-55581-442-7 Microbial Diversity: Form and Function in Prokaryotes by Oladele Ogunseitan; 2005; Blackwell Science Ltd.; ISBN 0-632-04708-9				

		Bachelor's Degree with Research / M. Sc. Microbiology (I)			
Subject	: Microbiology	Year: Second (2)	Semester: Third (III)		
Core Pap	er (Compulsory)	Course Code: B080906P	Practical III		
Marks:100 75 (UE) + 25 (CIE)		04 credits	Duration: 120 hours		
Total Numb	er of Lectures-Tutori	als-Practical (in hours per week L-	Г-Р: 0-0-8		
Course Objectives	The course will enable students to apply the learning of microbiology concepts toward exploitation of microbial population for industrial and human benefits.				
Course Learning Outcomes	Upon successful completion of the course, the student:				
COL	Will be able to ana	lyze the water quality and potability by	v using various techniques		
CO2	Will be able to use	various bioinformatics tools.	y asing rainous teeminques.		
CO3	Will be aware of various biochemical tests used in bacterial identification.				
CO4	Will be able to use special staining procedures.				
CO5		se microorganisms for production of	of various useful and industrially		

Drougamma/Class

Contents

- 1. Analysis of water quality: DO, BOD, alkalinity, free CO2, free chloride, TS, TSS, TDS, nitrate, phosphate
- 2. Determination of most probable number (MPN) for coliform bacteria
- 3. Isolation of bacterial strains from different soil samples
- 4. Analysis of sequence data and searching of research papers from various national and international journals
- Retrieval of gene and protein sequences from data bank
- 6. Sequence comparisons and alignment (8P)
- 7. Visualisation and other utilities (PDB viewer)

important products.

- 8. Biochemical tests for characterization of microbes (based on metabolic properties:
 - a. Carbohydrate fermentation
 - b. H2S production
 - c. Nitrate reduction
 - d. Urease activity
 - e. IMViC test
 - f. Gelatine liquefaction
 - g. Starch hydrolysis
 - h. Glycine decarboxylation
 - i. Catalase oxidase peroxidase test
- 9. Staining of polyphosphate bodies, polyhydroxybutyrate and endospore
- 10. Isolation of protease, amylase and lipase producing bacterial strains and estimation of enzyme activity
- 11. Mushroom production
- 12. Cell and enzyme immobilization.
- 13. Production of alcohol from molasses/cane sugar.
- 14. Production of vinegar.
- 15. Production of citric acid.
- 16. Isolation of cellulose producing strain.

Suggested Readings:	 Microbiology: A laboratory manual by JG Cappucino, N Sherman. 10th edition. Pearson. 2014.
readings.	2. Environmental Microbiology: A lab manual by I. Pepper, C. Gerba, I
	Brendecke. 46 th edition. Academic Press. 2011.
	 Sequence - Evolution - Function: Computational Approaches in Comparative Genomics by E.V. Koonin, M.Y. Galperin. Kluwer Academic, USA. 2003.
	4. Bioinformatics: A Practical Guide to the Analysis of Genes and Proteins edited



by A. D. Baxevanis, B.F. Francis Ouellette . 3rd edition. Wiley and Sons. 2004

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Naue kon

		Programme/Class: Bachelor's Degree with Research M. Sc. Microbiology (I)	
Subject:	Microbiology	Year: Second (2)	Semester: Third (III)
Core Paper (Compulsory)		Course Code: B080907R	Industrial Training /Surveys/Research Project III
Marks:100		Credits: 04	
Course Details		t can be interdisciplinary / multi-discip rial training / internship / survey work	linary. This research project can also be etc.
	carried out in both	the semesters at the end of the year,	dissertation) of the research project which will be assessed jointly by the university at the end of the year out

		Programme/Class: M. Sc. Microbiology (II)			
Subject:	Microbiology	Year: Second (2)	Semester: Fourth	(IV)	
AND RESIDENCE OF THE PROPERTY	tive (Optional)	Course Code: B081001T	FOOD MICROBIO	Address Control of the Control of th	
A CONTRACTOR OF THE PARTY OF TH	rks:100	75 (UE) + 25 (CIE)	Credits: 04	LOGI	
The second second second second second		ials-Practical (in hours per week			
Course Objectives	The course will phenotypic andb fungi and bacteria fermented milk p compounds and	l enable students to understant iochemical identification of food. The course will teach the strategoroducts, plant-based products, fish malt beverages, wines, distilled spoilage, preservation and various f	d the taxonomical class associated molds, yeasts, gies to develop fermented a products, meat products liquors and vinegar. The	yeast-like and non- bioactive e role of	
Course Learning Outcomes		completion of the course, the studen		nscussed.	
CO1	Will know about	microbial spoilage of various kinds	of food.		
CO2	Will be aware of	general principles of food preservat	ion.		
CO3		on regarding fermented food produ		1	
CO4	Knows about indi-	cator Microorganisms and microbia ns that revolutionize food safety.		quality	
CO5		about food borne microorganisms	and food poisoning.		
Contents:				Duration 60 hours	
UNITI	Food as a substrat and Seafood; Milk Cereals	te for microorganisms, Microbial s and Dairy Products; Fruits and Veg	poilage of Meat, Poultry, getables; Nuts, Seeds, and	12 Hours	
UNIT II	preservation. New d	Various classical, physical, chemical, a levelopments in food preservation tech ation of such techniques.	and biological methods of iniques. Analysis of	12 Hours	
UNIT III	Fermented Dairy adaptations and ch and non-fermented foods, Fermented	Products, Microbial habitat of nanges in microbiome of vegetable milk products, fresh meats, poultry Vegetables, Fermented Meat, Pou Beer, Wine, Vinegar, Probiotics a	s, fruits, milk, fermented and non-dairy fermented ultry, and Fish Products,	12 Hours	
JNIT IV	(DART) and prev	ganisms as an indicator of good quailing food standards in India (find Critical Control Point System	uality, Food adulteration	12 Hours	
JNIT V	Food borne infecti	ons including bacterial, viral and food borne parasites. In depth stu	ungalinfections. Study of idy of various types and	12 Hours	
uggested leadings	1. Food M edition. I	icrobiology by W.C. Frazier, D.C. McGrawHill Education. 2013.			
	edition.S 3. Fundame press. 20 4. Food Mic Royal So	crobiology by M. R. Adams, M. C cietyof Chemistry. 2015.	and A. Bhunia. 5 th edition D. Moss, P. McClure. 4 th e	n. CRC	
	5. Food Mi	crobiology: Fundamentals and F	rontiers by M. P. Doyle,	L. R.	

Beuchat. 3rdedition. ASM press. 2007.

6. Food Microbiology: An Introduction by T. Montville, K. Matthews, K.Kniel. 4th edition. ASM press. 2017.

		Programme/Class:		
Subjects N	dicrobiology	M. Sc. Microbiology (II)		
and the state of t	tive (Optional)	Year: Second (2) Course Code: B081002T	Semester: Fo	The second secon
			AGRICULT MICROBIO	
Mark to the Indian State	ks:100	75 (UE) + 25 (CIE)	Credits	THE RESIDENCE OF CHILDREN STREET, SANSAGE STRE
Total Number	er of Lectures-Tute	rials-Practical (in hours per week	L-T-P: 4-0-0	
Course Objectives	causing change translocation. Th	facilitate in understanding of major athogens that interact with various p in physiology, photosynthesis, he course will cover the application d microbial biopesticides along with	respiration, transpons of Plant Growth	the plant by piration and Promoting
Course Learning Outcomes	Upon successful	completion of the course, the studen	ıt:	
COI	Will have acquire processes.	red knowledge about the role of so	il microorganism in	various soil
CO2	Will learn about action. Will also	plant growth promotion attributes of learn about biostimulants.	f PGPR and their me	echanism of
CO3	Understands abo	ut various types of plant microbe into	eractions.	
CO4	The state of the s	ed to mechanism of action of variou		ides.
CO5		ed to mechanism of action of variou		
Contents:				Duration: 60 hours
UNITI	soil health, root rhizospheric mi microbial transfe	isms, major groups, decomposition of exudates and rhizospheric effect, croflora in plant productivity, mi ermation of phosphorus and sulphur izers in agriculture and forestry, b	manipulation of crobial biomass, minor nutrients,	12 Hours
UNIT II	The state of the s	romoting Rhizobacteria and their omposition of soil organic matter-		12 Hours
UNIT III	Mycorrhizal asso	relationships: Association and ciation: Their types and role in plant	nutrition.	12 Hours
UNIT IV	Mechanism of a		sticides (Bacillus	12 Hours
UNIT V	virus) and funga	es (nuclear polyhedrosis virus, cytopl l biopesticides (Metarrhinium aniso llum lecani, Hirsutella thomsonii)		12 Hours
Suggested Readings	2. Principles of Pvt. Ltd. 3. Plant Diseased 4. Agriculture Prentice Harman Pr	ology by Agrios GN. Fifth edition, E of plant pathology by R.S. Singh, Ox ases by R.S. Singh, CBS Publisher. Microbiology by Rangaswami, G, all of India Pvt. Ltd., New Delhi. In Agriculture Microbiology by Subb at pathology by M. Dickinson, Bios	and Bagyaraj, DJ, e	hing Comparedition 2nd,

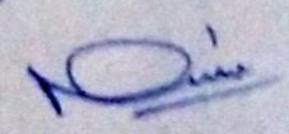
		Programme/Classi		
Subject 1	Alamakia k	M. Sc. Microbiology (II)		
Subject: Microbiology Major Elective (Optional)		Year: Second (2) Course Code: B081003T	MICROBIOLOGY	
	ks:100	75 (UE) + 25 (CH)	Chaditas	SECUL AND SECULATION OF THE PARTY OF THE PAR
Total Numbe	r of Lectures-Tuto	rials-Practical (in hours per week	L-T-D: 4-0-0	V1
Course Objectives	with their mecha mechanism of ac microbes to coun	acilitate in understanding of major granism of action. The course will detion of various antibiotics along witteract the action of various antimicrous with current scenario by an introducts with current scenario	oups of Human path evelop an understar h the mechanisms d bial agents. The cou	iding about eveloped in rse will also
Course Learning Outcomes	Upon successful	completion of the course, the student		
COI	the methods for c	ed knowledge about the normal micro collection and transportation of pathol	flora of human body logical specimens.	along with
CO2	Will have learnt	about principles of pathogenicity.		
CO3		ut various bacterial and fungal disea	ses along with their	symptoms
CO4	Will have learn al of action.	bout various viral diseases along with	their symptoms and i	nechanism
CO5		ntroduced to mechanism of action of of development of antimicrobial resis		alond with
Contents				Duration: 60 hours
UNIT I	Human body. Co	cally important microorganisms; nor ollection, transportation and examina- ion and identification of pathogenic of	ation of pathologic	12 Hours
UNIT II	pathogens, bacter	irulence factors, spreading and ial toxins-their types, mycotoxins, in Epidemiology of infection diseases.		12 Hours
UNIT III	Brief account of bar pertusis), food and and gas gangrene) Bacterial zoonoses(diseases (malaria, symptomology, pat Candidiasis, histopl	cterial diseases spread through air (dipthe water (typhoid, cholera and dysentery) s and contact (leprosy, conjunctivitis and brucellosis, bubonic plague and salmone filarial and kalazar). Etiology, epidemi hology, disease diagnosis and treatmen asmosis, aspergillosis, cryptooccosis and	d venereal diseases). closis) and protozoal ology, pathogenesis, t of fungal diseases: dermatomycosis.	
JNIT IV	(pneumotropic): (dermotropic); der poliomyelitis and s fever viral zoonose	And Andrews the Company of the Compa	sles and rubella cerotropic): rabies, phalitis and yellow	12 Hours
INIT V	of their action with resistance in bacter infections and eme in biological science	otherapy, role of antimicrobial agent special reference to antibiotics. Mole ria, and drug sensitivity test. Introduct erging microbial infection diseases. I ces and disposal of biomedical waste,	cular basis of drug tion to Nosocomial Biosafety practices bio-terrorism.	12 Hours
uggested eadings	Medical N Kobayashi,	Microbiology by Murray, PR, R GS & Pfaller, MA (ed III) Mosby In	c. KS,	5.

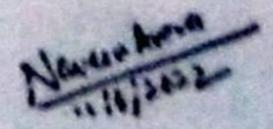
- Essentials of Medical Microbiology By Volk WA, Gebhardt, BM, Hammarskjold, ML & Kadner RJ (Ed V) Lipincott-Raven Publisher, Philadelphia
- Jawetz, Melnick & Adelberg's medical microbiology by Brooks, GF, Carroll, KC, Butel, JS, Morse, SA, Edition 24th, McGraw-Hill Medical,
- Medical Microbiology by Cruikshank, Edition 12th, Churchill Livingstone Pub.

Naukhan Naukhan

		Programme/Class: M. Sc. Microbiology (II)		
Subject: N	Microbiology	Year: Second (2)	Semester: Four	th (IV)
Major Elective (Optional) Marks:100		Course Code: B081004T	IPR AND BIOS	THE THE PARTY OF T
		75 (UE) + 25 (CIE)	Credits: 0	ASSESSMENT OF THE PARTY OF THE
Total Number	er of Lectures-Tutor	ials-Practical (in hours per week	The same of the sa	
Course Objectives	The course will International fram	develop an understanding about I nework for the protection of IP. To esafety when dealing with different	ntellectual Property The course will also	aware the
Course Learning Outcomes		ompletion of the course, the student		
COI	Will have acquire Property.	ed basic knowledge about IPR and	different types of I	ntellectual
CO2:		bout international framework for the	protection of IP	
CO3	Understands the b	asics of patents.		
CO4		to various levels of Biosafety.		
CO5		ced to GRAS organisms and	biosafety levels of	f specific
Contents:				Duration: 60 hours
UNIT I	copyright & rela	ntellectual property; types of IP: pated rights, industrial design, tractations, protection of new GMOs	patents, trademarks, ditional knowledge,	12 Hours
UNIT II	International fram	ework for the protection of IP; IP as icrobiology and few case studies;	a factor in R&D IPs	12 Hours
UNIT III		istory of GATT, WTO, WIPO and	d TRIPS. Basics of	12 Hours
UNIT IV		osafety; historical background; introdimary containment for biohazards; b		12 Hours
UNIT V	GRAS organism	s, Biosafety levels of specific safety levels for infectious agents an	e microorganisms;	12 Hours
Suggested Readings:	Fundamentals of I	Intellectual Property Rights (Third ty Rights by Neeraj Pandey and Khuntellectual Property Rights: For Stukrishna B & Anil Kumar H.S.	ushdeep Dharni	

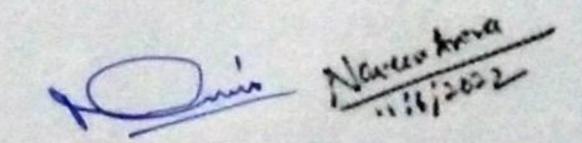
		Programme/Class: M. Sc. Microbiology (II)				
Sebject	Microbiology	Year: Second (2)	Semester: Fo	urth (IV)		
Major Elective (Optional)		Course Code: B081005T	PATHOGE	HAL		
M	arks:100	75 (UE) + 25 (CIE)	Credita	minimal markethine and the		
Indianate Consumption States of States	Color of the Color	ials-Practical (in hours per week I	Principle of Market Advisory Control of the Control			
Course		this course is to make the students				
Objectives	The students would and role of newer enable students to	nicrobes pathogenic or disease-cause levance to India and the various tools id also learn the mechanisms of resist vaccines in controlling infectious disto describe the molecular diagnosmay be used for diagnosis of disease	for their local or glob tance of bacteria to a seases. The course a stic methods and a	nal spread. intibiotics vould also intomated		
Course Learning Outcomes	Upon successful o	ompletion of the course, the student	will be able:			
COI	To understand class	ssical and molecular determinants of	disease causing mis	coher		
CO2		aracteristics of newer disease-causin				
CO3	To study and crit	ique the various molecular tools av	THE RESIDENCE OF REPARTMENT AND ADDRESS OF THE PARTMENT OF THE			
CO4	To study and evalu	To study and evaluate mechanisms underlying resistance of bacteria to antibiotics, sprea of resistance and the use of newer vaccines to control infectious diseases				
COS	To gather informa	ition as to how the infectious disease and what automated equipment are	es may be diagnosed			
Contents:				Duration: 60 hours		
UNITI	virulence; Quantita (MLD), LD50, ID	microbial pathogenicity: Define patative measures of pathogenicity: minimo. TCIDso Virulence determinants and invasiveness. Facultative/ oblig	mal lethal dose	12 Hours		
UNIT II	multiplicity of virulence genes, determinants by two variation; clonal a	rulence determinants, coordinated and environmental regulation to component signal transudation system (TTSS, T3SS), Role of biofilm al pathogenicity	regulation of of virulence tems, antigenic athogens, type	12 Hours		
UNIT III	epidemiology. Bid serotyping, phage to Molecular typing: typing, PFGE, AFL of geographical inf	bial epidemiology: Objectives ochemical and Immunological tool typing, multilocus enzyme electropho RAPD, rep (REP, ERIC, BOX)-I P, MLST, VNTR and whole genome formation system (GIS) for microbial	oresis (MLEE); PCR, IS based e sequence, use epidemiology.	12 Hours		
UNIT IV	pumps, extended	stance (AMR): Recent concepts – m spectrum β-lactamases (ESBL), scillin-resistant S. aureus (MRSA), re	X-MDR M.	12 Hours		
JNITV	Rapid diagnostie pr	inciples: Nucleic acid probes in diag	nostic microbiology.	12 Hours		





	sequencing and mutation detection, automated instruments for detection/diagnosis of infectious agents (BACTAC and Vitek-2, GeneXpert).
Suggested Readings	 Jawetz, Melnick, & Adelberg's Medical Microbiology by Carroll KC, Hobdon JA, Miller S, Morse SA, Mietzner TA. 27th edition. Lange Publication, 2016.
	 Beginner's guide to comparative genome analysis using next generation sequence data by Edward DJ and Holt KE in Microbial Informatics and Experimentation, 3:2, https://doi.org/10.1186/2042-5783 3-2, 2013.
	 Bacterial Pathogenesis: A molecular approach by Wilson BA, Salyers AA, Whitt DD, Winkler ME. 3rd edition. American Society for Microbiology Press, Washington, DC USA, 2011.
	 Bacterial Pathogenesis: Molecular and Cellular Mechanisms by Locht C, Simonet M, Caister Academic Press, 2012.
	 Molecular Microbiology: Diagnostic Principles and Practice by Persing DH, Tenover FC, Hayden R, Leven M, Miller MB, Nolte FS, Tang YW, Belkum AAV. 3rd edition. Washington, American Society for Microbiology Press, 2016
	 Infectious Disease Epidemiology: Theory and Practice by Nelson KE, Williams CM. 4th edition. Jones and Bartlett, 2019.

		Programme/Class: M. Sc. Microbiology (II)		
Subject:	Microbiology	Year: Second (2)	Samastar: For	wth (TV)
Major Elective (Optional) Marks: 100		Course Code: B081006T	Semester: Fourth (IV) PLANT-PATHOGEN INTERACTIONS	
		75 (UE) + 25 (CIE)	Credits:	
Total Numbe	r of Lectures-Tutor	rials-Practical (in hours per week L-	T-P: 4-0-0	
Course Objectives	The course will it plants and effect translocation. The molecular interact transgenic plants.	facilitate in understanding of how pate t plant physiology, photosynthesis, re- involvement of various enzymes and tion will help in designing biocontrol so The course covers the novel molecular g of plant diseases.	hogens interact with respiration, transpiration toxins and understand strategies and development	nding the
Course Learning Outcomes		completion of the course, the student:		
CO1	Will have gained resistance mecha	insight into genetics of host-pathogen	interactions, resistar	nce genes,
CO2	Will have been i methods of disea	ntroduced to plant disease control, phy	sical, chemical and	biological
CO3	Understands abo	out crown gall, symptoms of viral	diseases and thei	r control,
CO4	Will have attained knowledge about designing of molecular diagnosis of plant disease anddevelopment of transgenic plants with applications and constraints.			
CO5	Will be able to d	lescribe various important milestones in Indian farming.	in disease control ar	nd disease
Contents:		ant in morali fariting.		Duration 60 hours
UNIT I	Genetic basis interactions, res	of plant diseases: Genetics of istancegenes, resistance mechanisms in	host-pathogen plants.	12 Hours
UNIT II	Disease control chemical metho concepts and pra	Principles of plant disease control, ds of disease control, biocontrol, biocontrol, biocontrol, ctices, fungal agents, <i>Trichoderma</i> bis – uses and practical constraints.	physical and ontrol agents -	12 Hours
UNIT III	Some important gall, symptoms	plant diseases and their etiological sof viral diseases and their control, diseases, vegetables and crops.	studies: Crown seases of some	12 Hours
UNIT IV	Molecular appr plant protection and constraints.	oach: Molecular diagnosis, transgenie , futuristic vision of molecular diagnos	c approach for sis, applications	12 Hours
UNIT V		ting: History and important milestones ng and its relevance in Indian farming.	The second secon	12 Hours
Suggested Readings:	2. Plant Patho Hill. 2017 3. Bacterial p Cambridge 4. Molecular London. 20 5. The essent	logy by G. N. Agrios. 5th edition. Acade logy by R.S. Mehrotra, and A. Aggarw lant pathology: cell and molecular aspuniversityPress.1993. plant pathology by M. Dickinson. BIOS 1003. ials of Viruses, Vectors and Plant diseaseternLimited.1993.	al, 3 rd edition. Tata leads by D. C. Sign	e. rs,



- Biocontrol of Plant Diseases (Vol. I) by K.G. Mukerji and K.L.Garg. CRC PressInc., USA. 1988.
- 7. Molecular Biology of Filamentous Fungi by U.
 Stahl and P. Tudzyski. VCHVerlagsgesellschaftmbH. 1992.

		Programme/Class:		
Subject: M	icrobiology	M. Sc. Microbiology (II) Year: Second (2)		
Major Elective (Optional) Marks: 100		Course Code: B081007T	Semester: Fourth (IV) MYCOLOGY AND PHYCOLOGY	
		75 (UE) + 25 (CIE)		
otal Number of Lectures-Tutorials-Practical (in hours per week L-T-P: 4-0-0			04	
bjectives	characteristics	vill develop a basic understand of basic groups of algae and fur d economic importance.	ling about classifica	tion and symbiotic
Course Learning Outcomes	Upon successfi	ul completion of the course, the stud	lent:	
COI	Will gain kno characteristics	wledge about occurrence and distr of different classes of fungi.	ibution of fungi along	g with the
CO2	Will have bee	n introduced to different classes of fur	ngi	
CO3	Understands a	bout economic importance of fungi	ngi.	
CO4	Will attain kn	owledge about salient features of alga	P	
CO5	Is able to kno	w about economic importance of algae		
Contents:				Duration: 60 hours
UNITI	heterothallism, fungi and eco relationship wi fungi: asexual,	fungi: Contributions of Mycologists in ace and distribution, somatic structure, he sex hormones in fungi, physiological osystem; saprophytic parasitic, mutu th plants and animals. Classification of sexual and parasexual.	specialization in fungi, alistic and symbiotic fungi. Reproduction in	12 Hours
UNIT II	Study of the division of basidiomycotin Dictyostelium,	ifferent classes of fungi: Salient feature myxomycota, mastigomycota, zygor na and euetromycotina. Structure a Allomyces, Pilobolus, Claviceps and Fu	nycota, ascomycotina, and reproduction of:	
UNIT III	Economic implement and and ect-extile attack of funging environment, in textile. Mycote	ndo VAM, Fungi as insect symbionts, fur on other microorganisms, potential ap- ndustry, food. Role of fungi in bio deter oxins, quorum sensing in fungi.	e of Mycorrhiza: ecto-, ngi as biocontrol agents, plication in Agriculture, rioration of wood, paper,	
UNITIV	morphology a ecology, medi growth, strain cycle pattern i	es of Algae: Contributions of Phycologis nd classification of algae. Isolation from a and methods used for cultivating alga selection and large scale cultivation. On the different classes of algae	om soil and water, algal e. Measurement of algal General features and life	
UNIT V	removal, alga	otechnology: Algae as source of food a el and bioactive compounds Uses of l blooms and toxins.	f algae in heavy metal	
Suggested Readings	2. Charl 3. E.Mo 4. L. Ba 5. Ayha ofBio 6. Linda	poulos, C.J. and C.W. Mims 1979. Introduced in Ltd., NewDel ile M. & Watkinson S.C. The Fungi, Purore –Landeekeer: Fundamentals of the forsanti, Paolo Gualtieri: Algae: anatomy, nDemirbas, M. FatihDemirbas: Algae Endiesel (2010) a E. Graham, James Graham, James M. ett J.H., Publisher: Edward, Arnold Crar	blisher: Academic Press. fungi, Publisher: Prentice biochemistry, and biotec nergy: Algae as a New So Graham: Algae (2009)	Hall. hnology ource

		Programme/Class: M. Sc. Microbiology (II)		
Subject:	Microbiology	Year: Second (2)	Semester: Fourt	h (IV)
The second secon	tive (Optional)	Course Code: B081008T		
Ma	rks:100	75 (UE) + 25 (CIE)	Credits: 04	1
	r of Lectures-Tuto	rials-Practical (in hours per week	L-T-P: 4-0-0	
Course Objectives	The course will develo	evelop a basic understanding about op an idea about microbial growth k ol systems used to control various p	biochemical engineering inetics, transport pheno	menon
Course Learning Outcomes		completion of the course, the studen		
COI	behavior.	edge about upstream and downstream		
CO2	determination.	ntroduced to volumetric mass transfer		
CO3		out rheological properties of fermentation		y balance.
CO4	Will attain know	ledge about various types of bioreacto	ors.	
CO5	Will be able to k	now about various control systems	used to control various	Duration:
Contents:				60 hours
UNIT I	continuous and fe	ne bioprocess technology; Microbial ed batch culture.		
UNIT II	industrial fermer	menon in bioprocess: Introduction, on tation, oxygen supply and oxygen transfer rate, determination of Kla va	transfer rate, factors	
UNIT III	Non-Newtonian and energy balan	fluids, heat transfer and heat transfer are.	r correlation, and mass	
UNIT IV	Introduction to b	oioreactor: Ideal bioreactor, Reactor ctors, Multiphase bioreactors, anima	with non-ideal mixing, and plant cell reactor	12 Hours
UNITV	Instrumentation A. Methods of ma. Temperature b. Flow c. Pressure d. DO and free Company of the	chemical factors ms		12 Hour
Suggested Readings	1. Principle J. Hall; 7506-43	les of Fermentation Technology by F Second Edition; Butterrworth He 301-6. Biotechnology to Work: Bioproc	ess Engineering; Con	ISBN: 0-
	ISBN:	cess Engineering, National Research 0-309-58487-6. cyclopedia of bioprocess technology		

bioseparation by Michael C. Flickinger, Stephen W.Drew; John Wiley & Sons, Inc.; ISBN 0-471-13822-3

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Programme/Class: Bachelor's Degree with Research / M. Sc. Microbiology (I) Subject: Microbiology Year: Second (2) Semester: Fourth (IV) Course Code: B081009P Practical III Core (Compulsory) **Duration: 120 hours** 04 credits

Total Number	er of Lectures-Tutorials-Practical (in hours per week L-T-P: 0-0-8			
Course Objectives	The course will enable students to apply the learning of microbiology concepts toward the exploitation of microbial population for industrial and human benefits.			
Course Learning Outcomes	Upon successful completion of the course, the student:			
COI	Will be able to measure various bioprocess parameters.			
CO2	Will be able to microbiological quality of milk.			
CO3	Will be aware of isolation procedure of various PGPR and their efficiency assessment			
CO4	Will be able to determine Determination of antibiotic sensitivity and MIC by different procedures.			

Contents:

Measurement of Ks value

Marks:100

- Determination of specific growth rate and generation time
- Estimation of KLa value by sufite oxidation method
- Milk quality test- methylene blue reduction test,
- Ames test
- Isolation of PGPR from soil.
 - a. Isolation of Azotobacter
 - b. Isolation of Azospirillum
 - c. Isolation of Pseudomonas
 - d. Isolation of Rhizobium
- 7. Determination of plant growth promotion activity of bacterial isolates
 - a. IAA Production
 - b. Ammonia Production
 - c. Siderophore production
 - d. Phosphate solubilization
 - e. Ammonia production
 - f. HCN Production
- 8. Determination of antibiotic sensitivity by
 - Well diffusion method.
 - b. Disk diffusion method.
 - c. Plug diffusion method..
- 9. Determination of MIC for selected antibiotics

Suggested	1. Microbiology: A laboratory manual by JG Cappucino, C.T. Welsh. 11th edition.
Readings:	Pearson. 2017. 2. Environmental Microbiology: A lab manual by I. Pepper, C. Gerba, J. Brendecke. 46 th edition. Academic Press. 2011.

		Programme/Class; Bachelor's Degree with Research M. Sc. Microbiology (I)	
Subject: Microbiology		Year: Second (2)	Semester: Third (IV)
Core Paper (Compulsory)		Course Code: B081010R	Industrial Training /Surveys/Research Project IV
Marks:100		Credits: 04	
Course Details			
	* Students will sul carried out in both	mit the final report (project report, the semesters at the end of the year,	dissertation) of the research project which will be assessed jointly by the university at the end of the year out