"Dissemination of Education for Knowledge, Science and Culture"
-Shikshanmaharshi Dr. Bapuji Salunkhe.

Shri Swami Vivekanand Shikshan Sanstha's

Vivekanand College, Kolhapur (An Empowered Autonomous Institute)



DEPARTMENT OF MICROBIOLOGY

B. Sc. Part-III
Semester - V & VI
NEP Phase I

SYLLABUS

B.Sc.-III (Sem -V and VI) Microbiology

To be implemented from Academic Year 2025-2026

Programme Specific Outcome

Upon completion of B.Sc. Microbiology programme, student will be able to -

PSO1: Perform the basic techniques related to screening, isolation and cultivation of microorganism from various sources

PSO2: Understand microorganisms and their relationship with the environment

PSO3: Conduct the basic research with this microorganism and perform the diagnostic procedures required in food, milk and pharmaceutical industries.

PSO4: Follow the aseptic techniques and conduct the process of sterilization as well as perform the techniques to control the microorganism

PSO5: Produce and analyze the microbial product at laboratory level.

STRUCTURE OF COURSE UG degree in Major and Minor (B.Sc.- III Semester- V & VI)

(B.Sc III Semester- V & VI) Sr. Teaching F										
No.	Course Abbr.				Scheme		Examination Scheme and Marks			
		Course code	Course Course Name Hour	Hours	/week				Credits	
				TH	PR	ESE	CIE	PR	Marks	
Sem	ester-V									
1	DSC-IX	DSC03MIC51	Immunology	2	-	40	10	-	50	2
2	DSC-X	DSC03 MIC 52	Virology	2	-	40	10	-	50	2
3	DSC-XI	DSC03 MIC 53	Agricultural Microbiology	2	-	40	10	-	50	2
4	DSE – I	DSE03 MIC 51	Food and Industrial Microbiology	2	-	40	10	-	50	2
		DSE03 MIC 52	Fermentation Technology-I							
5	VSC-PR-IV	VSC03MIC59	Soil Microbiology		4	-	-	25	25	2
6	FP	FPR03MIC51	Field Project	2		-	-	50	50	2
7	DSC-PR-V	DSC03MIC59	DSC Microbiology Lab-5	-	12	-	-	75	75	6
8	MIN- IX	MIN03 MIC 51	Agricultural Microbiology	2		40	10	-	50	2
9	MIN-PR-V	MIN03 MIC 59	MIN-Microbiology Lab-5	ı	4	-	-	25	25	2
Sen	nester -V Tot	al		12	20	200	50	175	425	22
Sem	ester-VI									
1	DSC-XII	DSC03 MIC 61	Microbial Genetics - II	2	-	40	10	-	50	2
2	DSC-XIII	DSC03 MIC 62	Microbial Biochemistry	2	-	40	10	-	50	2
3	DSC-XIV	DSC03 MIC 63	Medical Microbiology - II	2	-	40	10	-	50	2
4	DCE II	DSE03 MIC 61	Environmental Microbiology	2	-	40	10	-	50	2
	DSE-II	DSE03 MIC 62	Fermentation Technology-II							
5	VSC-PR-V	VSC03MIC69	Waste Water Management		4	-	-	25	25	2
6	OJT	OJT03 MIC 61	On Job Training	2	-	-	-	50	50	2
7	DSC-PR-VI	DSC03 MIC 69	DSC Microbiology Lab-6		12			75	75	6
8	MIN- X	MIN03 MIC 61	Environmental &Industrial Microbiology	2	-	40	10		50	2
9	MIN- PR-VI	MIN03 MIC 69	MIN-Microbiology Lab-6	-	4	-	-	25	25	2
Sen	nester -VI To	tal		12	20	200	50	175	425	22

Abbreviations: TH-Theory, PR-Practical, PRO- Project, ESE- End Semester Examination, CIE-Continuous Internal Examination

Note: Minimum passing for 10 marks Internal evaluation = 04 marks

Minimum passing for 40 marks Theory paper = 16 marks
Minimum passing for 25 marks Practical = 10 marks
Minimum passing for 50 marks Practical /FP/OJT = 20 marks
Minimum passing for 100 marks Practical/FP = 40 marks

Passing percentage for Democracy, Election and Good Governance (DEGG) and

Environmental Studies papers should be 40%

Separate passing for each Head - ESE, CIE and Practicals

SEMESTER -V

DSC-IX	IMMUNOLOGY	No. of
DSC03MIC51	Theory: 30 Hours (Credits -2)	Hours
		per unit
T		•

Expected course outcome -

Upon successful completion of course, students are expected to be able to -

CO1: Understand the overall organization of the Immune system.

CO2: Explain the salient features of antigen antibody reaction and its use in diagnostics .

CO3: Compare and contrast humoral versus adaptive immune system.

CO4: Provide an overview of the interaction between the immune system and pathogen.

UNIT-I	1.Cells of Immune system –	8
	a. Hematapoiesis- characteristics & types of stem cells.	O
	b. Classification of cells of immune system – lymphoid &	
	myeloid cells.	
	c. Structure & function of lymphoid cells – T cell & T cell	
	subsets, NK cells, B cells & dendritic cells.	
	d. Structure & function of myeloid cells- Granulocytes,	
	monocytes & macrophages.	
	2. Membrane receptors for antigen and their role in	
	antigenrecognition	
	a.Bcellsurfacereceptorforantigen(BCR)	
	b.Tcellsurfacereceptorforantigen(TCR)	
	c. NKreceptors	
	3. Molecular mechanism of antibodyproduction.	
	a. Processing and presentation of antigen by Antigen Presenting Cell	
	b.Interaction of APC with T _H cell	
	c.Interaction of B cell and T _H Cell	
	d.Clonalproliferation and differentiation of activated B cell.	
	e.Role of follicular dendritic cells in selection of high affinity B cell.	
	f.Role of cytokines in proliferation and differentiation.	
UNIT-II	1. Cytokines -	7
	a. Properties, types and function of cytokines	

	produced by TH cell and Macrophages	
	2. Interferon -	
	a. Natureandtypesof Interferons	
	b. Induction of Interferon	
	c.Mechanismofaction.	
	3. Immunologicaltolerance:	
	a.	
	Toleranceinductioninadultsandneonatesbydrugan	
	dmonoclonalantibody	
	b. Cellularmechanismofimmunologicaltolerance.	
	c. Terminationoftolerance.	
UNIT-III	1. Complement –	8
	a.Natureand Properties of complement	
	b.Complementactivation by classical and alternate	
	pathway.	
	C.	
	Biologicalconsequencesofcomplementactivatio	
	n.	
	2.Monoclonal antibodies-	
	a. Basicconcepts-	
	Mouse, Humanand Humanized antibodies.	
	b.Productionofmonoclonalantibodiesbyhybridomatechno	
	logy.	
	c. Production of Humanized Monoclonal	
	antibodies by recombinantDNA technology.	
	d.	
	Applicationsofmonoclonalantibodiesindiagnosis, tre	
	atmentandresearch.	
UNIT-IV	1. New diagnostic techniques:	7
	a. RIA	
	b. Dot BlotTechnique	
	2. Hypersensitivity-	
	aBasicconcept,GellandCoombsclassification	
	b. TypeI-Anaphylaxis	
	c. TypeII-Bloodtransfusionreactions	

d. TypeIII-Serumsickness	
e .Type IV- Delayed type hypersensitivity -	
Allograftrejection.	
3. Autoimmunedisease:	
aTypes of autoimmune diseases.	
	1

b Treatmentofautoimmunediseases.

Books Recommended:

A. For Immunology

- 1) Kubay, Kindt, Goldsby & Osborne . Immunology-6thedition-
- $2) \ Essential Immunology-11 {}^{th}edition-Delves, Martin, Burton and Roitt.$
- 3) Immunology-AnIntroduction,4thedition-Tizzard.
- ${\it 4)} \ \ Basicand Clinical Immunology 5 {\it the} dition-Stites, Stobo, H.H. Fudenberg.$
- 5) EssentialsofImmunology-S.K.Gupta
- 6) Immunology-M.P.Arora

DSC-X	VIDOLOCY	No. of
DSC03MI	VIROLOGY	Hours
C52	Theory: 30 Hours (Credits -2)	per unit
Expected co	ourse outcome -	
Upon succe	essful completion of course, students will be able to -	
CO1: Descr	ibe various stages involved in multiplication cycle of viruses	
CO2: Unde	rstand methodological approaches in isolation, cultivation & purificati	on of
viru	ses.	
CO3: Distin	nguish characteristics of normal cell and cancerous cell.	
CO4: Expla	in various methods for enumeration of viruses.	
UNIT - I	1. The Structural properties of viruses:	7
	a.Capsids, Nucleic acids andenvelope.	
	b. StructureofT4bacteriophage,TMVandHIV,Viroids&prions.	
TINITE II	c. Onestepgrowthexperiment	
UNIT – II	1. Isolation, cultivation and Purification of viruses	8
	a. Isolation and cultivation of viruses-	
	i. Animalvirus-Tissueculture, chickembryoandliveanimals.	
	ii. Plantvirus-Protoplasts culture technique,Insecttissueculture	
	iii. Bacteriophages - Plaque method.	
	b. Purification of viruses using physico-chemical properties	
	i.Density gradientcentrifugation	
	ii. Precipitation	
	2. Methods of Enumeration of viruses	
	a. Latexdropletmethod(Directmicroscopiccount)	
	b. Plaqueandpockmethod	
UNIT-III	1. Lysogeny-	7
	a.Definitionoflysogenyandtemperatephage,	
	b.types of lysogenic phage.	
	c. lysogenybylambda phage - adsorption &penetration	
	genetic map for lysogenic interaction, expression of λ genes,	
	establishment of repression, maintenance of repression,	
	integration of λ genome in hostchromosome.	
	2. Reproductionofanimalviruses-Adenovirus.	
	3. Reproductionofplantviruses-TMV	
	4. ReproductionofT4phage.	
UNIT-IV	1. Oncogenesis:	8

- a. Definition of oncogenesis
- b. Types of cancer
- c. Characteristicsofcancercells.
- d. Tumor suppressor genes and protooncogenes
- e. Hypothesis about cancer.
- i. Somatic mutation hypothesis
- ii. Viral genehypothesis
- RoleofDNAviruseswithspecialemphasisonPapov aviruses.
- Role of RNA tumor viruses

Provirus theory, Protovirus theory, Oncogene theory.

iii. Defective immunity hypothesis.

BooksRecommended:

- 1. General Microbiology Stanier
- 2. Microbiology Prescott, Klein
- 3. Microbiology -Davis
- 4.General Virology -Luria
- 5. Genetics of Bacteria and their Viruses-William Hayes.
- 6.GeneralMicrobiologyVol.II-PowarandDaginawala
- 7. Virology Biswas and Biswas

DSC-XI		No. of
DSC03MIC53	AGRICULTURALMICROBIOLOGY	Hours
	Theory: 30 Hours (Credits -2)	per
		unit
Expected cours	se outcome -	1
Upon successf	ul completion of course, students will be able to -	
CO1: Understa	and various plant microbe interactions especially rhizosphere and thei	r
applicat	ions especially the biofertilizers and their production techniques	
CO2: Understa	nd various biogeochemical cycles - C, N, P cycle and microbes involve	ed
CO3: Perform	isolation of agriculturally important microorganisms and formulate	
biofertili	zers.	
CO4: Explain r	ole of microorganisms and common symptoms of plant diseases.	
UNIT-I	1. SoilMicrobiology.	8
	a. Physicalcharacters.	
	b. Chemicalcharacters.	
	c. Typesofmicroorganismsinsoilandtheirroleinsoilfertility.	
	d. Microbiologicalinteractions-	
	Symbiosis, Commensalism, Amensalism, Parasitism, Predation.	
	2. Role of microorganisms inelemental cycle	
	a.Carboncycle.	
	b.Nitrogencycle	
	c.Phosphorouscycle	
	d.Sulfur cycle	
UNIT-II	1. Manure and Compost	7
	a. Methods of Production of-	
	i. Greenmanureandfarmyardmanure	
	ii. City compost-Windrow and pitmethod.	
	iii Vermicompost	
	b.Optimalconditionsforcompostingwithreferenceto-	
	Compositionoforganic waste, availability of microorganisms,	
	aeration, C: N:P ratio, moisture content, temperature, pH,Time	
UNIT-III	1. Types, production, methods of application and uses of -	8
	a. Biofertilizers	
	i. Nitrogen fixing - Azotobacter, Rhizobium, Azospirillum.	
	ii. Phosphate Solubilizing Microorganisms.	

	b. Biopesticides	
	i.Bacillusthuringiensis	
	ii. Tricodermaspp.	
	2. Biodegradation of -	
	a. Cellulose	
	b. Pesticides	
T 18 17 7 7 7 7		7
UNIT-IV	1. PlantPathology	7
	a. Commonsymptomsproducedbyplantpathogens	
	b. Modesoftransmissionofplantdiseases.	
	c. Plant diseases-	
	i. CitrusCanker	
	ii. Tikka disease ofgroundnut	
	iii.BacterialBlightofPomegranate.	

BooksRecommended

- 1. Soil Microbiology An exploratory approach MarkCoyne.
- 2. Agricultural Microbiology N. Mukherjee and J.Ghosh.
- 3. IntroductiontoSoilMicrobiology-MartinAlexanderIIndEdition.
- 4. AgriculturalMicrobiology-RangaswamyandBhagyarajIIndEdition
- 5. Plantdiseases-R.S.Singh.
- 6. Diseases of cropplants in India-G. Rangaswamy.
- $7.\,Soils and Soils Fertility-6 {}^{th}edition-$

FrederickR.Troeh(BlackwellpublishingCo.)

- $8.\,Soil Microbiology-Singh, Purohit, Parihar. (Agrobios India, 2010)\\$
- 9. Soil Microbiology and Biochemistry Ghulam Hassan Dar (New India Publishing Agency, 2010)

DSE-I DSE03MIC51	FOOD AND INDUSTRIAL MICROBIOLOGY Theory: 30 Hours (Credits -2)	No. of Hours per unit
Expected cours	se outcome -	-
Upon successf	ul completion of course, students will be able to –	
CO1: Know me	ethods used for industrial production of various products using microorga	anisms.
CO2: Explain v	various techniques for product recovery after fermentation.	
CO3: Understa	and the cause of spoilage of food and methods for preservation of food.	
CO4: Design tl	he methods for preservationofindustriallyimportantmicroorganisms.	
UNIT-I	1. Food Microbiology	7
	a. Foodasasubstrateformicroorganisms.	
	b. Foodborne diseases-	
	i. Roleofmicroorganismsinfoodborne diseases	
	ii. Food poisoning - Staphylococcal	
	Fungal (aflatoxin)	
	iii. Food infections -Salmonellosis.	
	c. Food spoilage and its causes.	
	d. General principles of food preservation.	
UNIT-II	1. Industrial Microbiology	8
	a. Strain Improvement	
	b. Scale up of fermentations	
	c. Microbiologicalassays	
	2. Preservationofindustriallyimportantmicroorganisms-	
	a.Methods of preservation.	
	b.Culturecollectioncenters	
UNIT-III	1. Industrial production of-	7
	a. Amylase-	
	Organismsused, i noculum preparation, fermen	
	tationmedia, fermentation conditions,	
	extraction andrecovery.	
	b. Grape wine - Definition, types, production of table wine (Red and	
	White), microbial defects of wine	
	c. Penicillin-	
	Organismsused, i noculum preparation, fermenta	
	tionmedia, fermentationconditions,	

	extractionand	
	recovery.Conceptofsemisyntheticpenicillin	
	d. Citric acid -	
	u. Chile aciu -	
	Organismsused, i noculum preparation, fermen	
	tationmedia, fermentation conditions,	
	extraction and recovery.	
	e. SCP by using yeast.	
	2. Microbial Production of –	
	a. Vitamins - Vit. B ₁₂	
	b. Amino acids – Lysine	
	3. Probiotics - Concept, Production by using <i>Lactobacillus</i> & applications	
UNIT-IV	1. Downstream processing &product recovery-	8
	a. Centrifugation	
	b. Flocculation	
	c. Filtration	
	d. Solvent extraction	
	e. Distillation	
	f. Precipitation	
	g. Crystallization	
	h. Chromatography.	
	2. Testing of sterility, pyrogen, carcinogenicity, toxicity and allergens	

Books Recommended:

A. For Food microbiology and industrial microbiology

- 1. Principles of fermentation technology- Peter F. Stanbury & Allan Whitaker (PergamonPress).
- $2.\ Principles of Microbial technology-Peppler, Vol. I\&II.$
- 3. Industrial Microbiology Casida
- 4. IndustrialMicrobiology-A.H.Patel
- 5. Industrial Microbiology-Prescott & Dunn
- 6. Industrial Microbiology -Miller
- $7.\ Pharmaceutical Microbiology-Huggo \& Russel$

DSE-I	FERMENTATION TECHNOLOGY-I	No.of
DSE03MIC52	Theory:30Hours(Credits-2)	Hours
	, , , ,	perunit/
		credit
	nes-Oncompletionofcourse,studentswillbeableto- thodsusedforindustrialproductionofvariousproductsusing micr	oorganisms.
CO2: Explain	various techniques for product recovery after fermentation.	
CO3: Apply n	ethods used for recovery of fermentation products.	
CO4:Identify	he industrially important microorganisms using screening technic	que.
	1. Basic concepts of fermentation.	8
UNIT I	a. Definition, concept of primary and secondary metabolites	
	b. Types of fermentations – Batch, continuous, dual and multiple.	
	c. Typical Fermenter design - Parts and their functions	
	d. Factors affecting fermentation process.	
	2. Fermentation Media.	
	Fermentation media –	
	 i) Water, carbon source, nitrogen source, precursors, growth factors, antifoam agents, chelating agents. 	
	ii) Use of wastes as Fermentation media – Molasses, sulphite waste liquor & corn steep liquor.	
	1. Screening:	
UNIT-II	Primary and secondary screening.	7
	2. Production strains	
	i) Concept	
	ii) Preparation of inoculum	
	iii) Strain improvement	

UNIT-III	 Scale upoffermentations Microbiologicalassays Testingofsterility,pyrogen,carcinogenicity,toxicityandallerg ens. 	7
UNIT-IV	1. Downstreamprocessing&productrecovery-	8
	1. Centrifugation	
	2. Flocculation	
	3. Filtration	
	4. Solventextraction	
	5. Distillation	
	6. Precipitation	
	7. Crystallization	
	8. Chromatography.	

Books Recommended:

- 1. Moo-Young M. ed. (1985) Comprehensive Biotechnology vol: I & II, Pergamon Press N.Y.
- 2. Ratledge C and Kristiansen B. eds. (2001) Basic Biotechnology 2nd ed. Cambridge Univ
- 3. Old R.W and Primose S.D (1995) Principles of Gene Manipulation 5th ed. Blackwell Scientific Pub. Oxford.
- 4. Bailey J.E and Ollis D.F. (1986) Biochemical Engineering Fundamentals 2nd ed. McGraw Hill Book Company, N. Delhi.
- 5. Aiba S, Humphrey A. E. and N. F. Millis (1973) Biochemical Engineering, 2nd Edition University of Tokyo Press, Tokyo, Japan.
- 6. Stanbury P.F., Whitaker A, and Hall S.J. (1997) Principles of Fermentation Technology
- 7. Mukhopadhaya S.N. (2001) Process Biotechnology Fundamentals. Viva Books Pvt. Ltd. N.Delhi.
- 8. Rehm H.J and Reed G. (1985) Biotechnology vol. I & II. VCH, Basel.
- 9. Stainer R. Y. Ingrahm J. L., Wheelis M. L. and Painter P. R. (1987) General Microbiology 5th Edition, Macmillan Press Ltd. London.

MIN-I		No. of
MIN03MIC11	AGRICULTURALMICROBIOLOGY	Hours
	Theory: 30 Hours (Credits -2)	per
		unit
Expected cours	e outcome -	
Upon successfu	al completion of course, students will be able to -	
CO1: Understa	nd various plant microbe interactions especially rhizosphere and the	ir
applicatio	ons especially the biofertilizers and their production techniques.	
CO2: Understa	nd various biogeochemical cycles - C, N, P cycle and microbes involv	red
CO3: Perform i	solation of agriculturally important microorganisms and formulate	
biofertili	zers.	
CO4: Explain r	ole of microorganisms and common symptoms of plant diseases.	
UNIT-I	1. SoilMicrobiology.	8
	a.Physicalcharacters.	
	b.Chemicalcharacters.	
	c. Types of microorganisms in soil and their role in soil fertility.	
	d.Microbiologicalinteractions-	
	i.Symbiosis,	
	ii.Commensalism,	
	iiiAmensalism,	
	iv.Parasitism,	
	v.Predation.	
	2. Role of microorganisms inelemental cycle	
	a.Carboncycle.	
	b.Nitrogencycle	
	c.Phosphorouscycle	
	d.Sulfur cycle	
UNIT-II	1. Manure and Compost	7
	a. Methods of Production-	
	i. Greenmanureandfarmyardmanure	
	ii. City compost-Windrow and pitmethod.	
	iii Vermicompost	
	b.Optimalconditionsforcompostingwithreferenceto-	
	Compositionoforganic waste, Availability of microorganisms,	
	aeration, C: N:P ratio, Moisture content, Temperature, pH,Time.	
	aeranon, C. IV.1 Tano, Moisture Content, Temperature, pri, Time.	

UNIT-III	1. Types, production, methods of application and uses of -	8
	a. Biofertilizers	
	i. Nitrogen fixing - Azotobacter, Rhizobium, Azospirillum.	
	ii. Phosphate Solubilizing Microorganisms.	
	b. Biopesticides	
	i.Bacillusthuringiensis	
	ii. Tricodermaspp.	
	2. Biodegradation by bacteria & fungi-	
	a. Cellulose	
	b. Pesticides	
UNIT-IV	1. PlantPathology	7
	a. Commonsymptomsproducedbyplantpathogens	
	b. Modesoftransmissionofplantdiseases.	
	c. Plant diseases-	
	i. CitrusCanker	
	ii. Tikka disease ofgroundnut	
	iii.BacterialBlightofPomegranate.	
	iv. Control of plant disease caused by bacteria.	

BooksRecommended

- 6. Soil Microbiology An exploratory approach MarkCoyne.
- 7. Agricultural Microbiology N. Mukherjee and J.Ghosh.
- 8. IntroductiontoSoilMicrobiology-MartinAlexanderIIndEdition.
- $9. \ Agricultural Microbiology-Rangas wamyand Bhagyaraj II^{nd} Edition$
- 10.Plantdiseases-R.S.Singh.
- 6. Diseases of cropplants in India-G. Rangaswamy.
- 7. SoilsandSoilsFertility-6thedition-

FrederickR.Troeh(BlackwellpublishingCo.)

- 8. SoilMicrobiology-Singh, Purohit, Parihar. (Agrobios India, 2010)
- 9. Soil Microbiology and Biochemistry Ghulam Hassan Dar (New India Publishing Agency, 2010)

PRACTICAL

SEMESTER V

DSC-PR-V	DSC Microbiology Lab-V		
DSC03MIC59	(Credits -4)		
	PRACTICALS BASED ON IMMUNOLOGY		
	 Major: Determination of Antibacterial activity of the serum. Enzyme Linked Immunosorbent Assay (ELISA)- DOT Determination of C- Reactive Protein (CRP) in Blood. Detection of Rheumatoid factor in blood .(Slide agglutination test) Minor: Sample handling Determination of Total Blood cell count Determination of Differential blood cell count. Separation of serum and plasma from blood. EstimationofhaemoglobinbySahli'smethod. DeterminationofESRofthebloodsample(Westergrenmethod) 		
	PRACTICALS BASED ON VIROLOGY		
	 Isolation of coliphages from sewage. Isolation of high titre of bacteriophage. Enumeration of bacteriophage in a sample by plaque forming unit (PFU) method. Determination of one step growth curve of bacteriophage. Determination of cross infectivity of <i>E.coli</i> phage. Minor: Demonstration of viruses inoculation by chick embryo technique Demonstration of PCR 		

PRACTICALS BASED ON AGRICULTURAL MICROBIOLOGY
Major:
1. Isolation of Azotobacter fromsoil.
2. IsolationofXanthomonasfrominfectedcitrusfruit.
3. IsolationofRhizobiumfromrootnodules.
4. Isolationofphosphatesolubilizingbacteriafromsoil.
5. Isolation of Trichoderma from soil .
Minor:
1. Determination of texture of soil.
2. Determination of color of soil.
3. Determination of pH of soil.
4. EstimationofCalciumfromsoil(EDTAmethod)
5. EstimationofMagnesiumfromsoil(EDTAmethod)
6. Determinationoforganiccarboncontentofsoil(WalkleyandBlackmethod)

DSE-PR-V	DSC Microbiology Lab-V.
DSE03MIC59	PRACTICALS BASED ON FOOD AND
	INDUSTRIALMICROBIOLOGY
	(Credits -4)
	Major:
	1. Production of wine.
	2. Examination of wine for pH, color and alcohol content.
	3. Determination of microflora of vegetables and fruits.
	4. Isolation and detection of aflatoxins from given food sample.
	5. Detection for the presence of yeast and mold from given sample.
	6. Sterility testing of pharmaceutical product.
	7. Rapid detection of food pathogens (E.coli&Staphylococcus) from given
	food sample
	Minor:
	1. Citric acid fermentationand recovery.
	2. Estimation of citric acid by titration.
	3. Amylase production by using <i>Bacillus</i> species.
	4. Isolation of lactic acid bacteria from fermented food.

5. Sauerkraut production.
6. Examination of milk by Directmicroscopic count (DMC)

VSC-PR-IV VSC03MIC 59	PRACTICALS BASED ON SOIL MICROBIOLOGY (Credits -2)	No. of Hours per unit
Expected cour	rse outcome -	'
On completio	on of course, student will be able to –	
CO1: Explain	physical and chemical characteristics of soil.	
CO2: Describ	e the role of microorganisms in soil fertility.	
CO3: Describ	be the role of microorganisms in various elemental cycles.	
CO4: Define	soil quality and its relation to Soil Microbiology	
	1) Determination of color & pH of soil.	
	2) Determination of temperature & humidity of soil.	
	3) Determination of water content of soil.	
	4) Determination of total bacterial count of soil.	
	5) Determination of texture of soil.	
	6) Estimation of total nitrogen content of soil.	
	7) Estimation of total phosphorous content of soil.	
	8) Estimation of potassium content of soil.	

MIN-PR-I	PRACTICALS BASED ON AGRICULTURAL MICROBIOLOGY
MIN03MIC19	(Credits -4)
	Major:
	1. Isolation of Azotobacter fromsoil.
	2. Isolation of Xanthomonas from infected citrus fruit.
	3.IsolationofRhizobiumfromrootnodules.
	4. Isolation of phosphatesolubilizing bacteria from soil.
	5. Isolation of Trichoderma from soil .
	Minor:
	1.Determination of texture of soil.
	2.Determination of color of soil.
	3.Determination of pH of soil.
	4.EstimationofCalciumfromsoil(EDTAmethod)
	5.EstimationofMagnesiumfromsoil(EDTAmethod)
	6.Determination of organic carbon content of soil (Walkley and Blackmethod)

SEMESTER VI

DSC XII	MICROBIAL GENETICS	No. of
DSC03MIC61	Theory: 30 Hours (Credits -2)	Hours per unit
Expected cours	se outcome -	F
Upon successf	ul completion of course, students will be able to –	
CO1: Understa	nd molecular mechanism involved in gene regulation	
CO2: Understa	nd the basic concept of operon and mutation.	
CO3: Discuss t	he principle, working and applications of molecular biology te	chniques
including	g PCR and DNA sequencing.	
	echniques used to manipulate genes & formation of clones	
UNIT-I	1. One cistron - one polypeptidehypothesis.	
	2. Molecular mechanism of geneexpression	7
	a. Concept ofoperon	
	b. Pribnowbox	
	c. Genetic regulation in tryptophanoperon	
UNIT-II	1. Mutations	
	a. Expression of mutations-	8
	i.Time course of phenotypicexpression.	
	ii.Conditional expression ofmutation.	
	b. Suppressor mutations (with examples) -	
	Genetic andnon-genetic.	
	2. Methods of isolation and detection of mutants based on-	
	a. Relativesurvival	
	b. Relativegrowth	
	c. Visualdetection	
UNIT - III	1.Genetic complementation - Cis-transtest	
	2.Extrachromosomal inheritance -	7
	a. Kappaparticles.	
	b. Transposable elements –	
	general properties andtypes.	
	3.Techniques in Molecular Biology-	
	a. DNA sequencing (Sanger's method)	
	b. DNA Fingerprinting	
	c. PCR	
	d. Blotting techniques- Southern, Western, Northern	

UNIT-IV	1.Geneticengineering	8
	a. Introduction	
	b. Tools of genetic engineering	
	i. Enzymes	
	ii. Vectors-phage, plasmid and cosmid	
	iii. DNAprobe - methods of preparation and detection.	
	iv. Linkers andadaptors	
	v. Cloning organisms - (Bacteria and Yeasts)	
	vi. Genomic library and cDNAlibrary	
	c. Techniques-	
	i. Isolation of desired DNA segment- Shotgun Method,	
	cDNA synthesis,	
	Chemical synthesis	
	ii. Construction of r-DNA using appropriate vector- Use of	
	restrictionenzymes,Linkers, Adaptors	
	Homopolymer tails	
	iii. Transfer to cloning organisms (Bacteria and Yeasts)	
	iv. Selection of recombinant microorganism Blue and white	
	screening,	
	Colony hybridizationtechnique.	
	d. Application of genetic engineering in-	
	i. Medicine-	
	ii. Agriculture	
	iii. Industry	
	iv. Environment	
	v. Understanding biology	

BooksRecommended:

- 1.Genetics -Stickberger.
- 2. Genes Benjamin Lewin IXed.
- $3. Principles \ of \ gene \ manipulation \ \ Primrose \ and Old$
- 4.Genetic Engineering Second Ed. Desmond S. T.Nicholl
- 5.Recombinant DNA J. D.Watson
- 6.Biochemistry -Lehninger
- 7. Molecular Biology of Gene J. D. Watson

DSC XIII	MICROBIAL BIOCHEMISTRY	No. of
DSC03MIC62		Hours
	Theory: 30 Hours (Credits -2)	per unit
Expected course	outcome -	
Upon successfu	l completion of course, students are expected to be able to -	
CO1: Explain M	letabolic pathways and Bioenergetics	
CO2: Understan	nd Various downstream processing	
CO3: Understan	nd Basic concept related to enzyme	
CO4: Determin	e enzyme production and its activity	
	1. Enzymes-	7
UNIT I	a. Definition, properties, structure, specificity, classification	
	and mechanism of action (Lock &Key, Induced	
	fithypothesis)	
	b. Allostericenzymes-	
	Definition, properties, models explainin	
	gmechanismofaction.	
	c. Ribozymes -concept, significance.	
	dIsozymes- definition, properties, example.	
	e. Factorsaffectingcatalyticefficiencyofenzymes	
	i. Proximity andorientation	
	ii. Strain and distortion.	
	iii. Acid basecatalysis	
	iv. Covalentcatalysis	
	f. Enzymekinetics-DerivationofMichaelis-	
	Mentenequation,	
	LineweaverBurkPlot,SignificanceofKm& Vmax.	
	g. Regulationofenzymesynthesis.	
	i. Positive control -Ara operon	
	ii. Negative control -Lac operon	
	iii. Cataboliterepression	
UNIT II	1 Extraction &purification ofenzymes.	8
	a. Methodsofextractionofintracellularandextracellularenzymes.	
	i. Choiceofsourceandbiomassdevelopment	
	ii. Methodsofhomogenization-celldisruptionmethods	

	iii. I diffeddololei Zyffesofid ebasisof	
	Molecularsize	
	 Solubilitydifferences 	
	Electricalcharge	
	Adsorption characteristic differences	
	2. Assayofenzymes-Basedonsubstrateandproductestimation.	
	3. Immobilizationofenzymes-Methods&applications	
	4. Confirmation of purified enzymes	
		7
UNIT III	1. Basic concepts of-	
	a.Glyoxylatebypass	
	b. Phosphoketolase pathway	
	c.Bioluminescence-Occurrence,mechanism&applications.	
	2. Assimilation of-	
	a.Carbon	
	b.NitrogenwithrespecttoN2andNH3(GOGAT)	
	c.Sulphur	
UNIT IV	1. Prokaryotic Biosynthesis of-	8
	a.RNA	
	b.DNA	
	c.Proteins	
	d. Peptidoglycan	

iii. Purification of enzymes on the basis of-

Books Recommended:

B. For Microbial Biochemistry

- 1. Enzymology-Prise&Stevens
- $2. \ Enzymes-Biochemistry, Biotechnology, clinical chemistry-Trevor Palmer.\\$
- 3. Enzymes -Dixon and Webb
- $4. \ \ Lehnigers Principles of Biochemistry by David Nelson \& Michale Cox, Fifthe dition.$
- 5. General Microbiology -Stanier
- $6. \ Principles \& techniques of Biochemistry-Wilson \& Walker, 6 the dition.$

DSC XIV DSC0 MEDICAL MICROBIOLOGY Theory: 30 Hours (Credits -2) Expected course outcome - Upon successful completion of course, students are expected to be able to - CO1: Correlate disease symptoms with causative agent, isolate and identify pathogens. CO2:Understand mechanism of action of antimicrobial drugs and their uses as prophylactic agents. CO3:Explain pathogenicity of organisms associated with human infections. CO4:Explain different antimicrobial agents with respect to their mode of action uses. I. Morphology, culturalandbiochemicalcharacteristics, antigenic structure, modesoftransmissionand pathogenesis, symptoms, laboratory diagnosis, preventionand controlof diseases caused by a. Mycobacteriumlepme b. Clostridiumperfringens, c. Trepomenapallidum UNIT II Morphology, cultural and biochemical characteristics, antigenic structure, modes oftransmission and pathogenesis, symptoms, laboratory diagnosis, preventionand controlof diseases caused by a. Pseudomonasacruginesa b. Virio delen c. Supukcous mutans d. Hillachatory ploi UNIT II Morphology, cultural and biochemical characteristics, antigenic structure, modes of transmission and pathogenesis, symptoms, laboratory diagnosis, preventionand controlof diseases caused by a. Protozoa: Plasmodium falciparum (malaria) b. Virusesi) Hepatitis A & B virus ii) Rabiesvirus			No. of
Theory: 30 Hours (Credits -2) Theory: 30 Hours (Credits -2) Expected course outcome - Upon successful completion of course, students are expected to be able to - COI: Correlate disease symptoms with causative agent, isolate and identify pathogens. CO2:Understand mechanism of action of antimicrobial drugs and their uses as prophylactic agents. CO3:Explain pathogenicity of organisms associated with human infections. CO4:Explain different antimicrobial agents with respect to their mode of action uses. 1. Morphology.culturalandbiochemicalcharacteristics, antigenicstructure, modesoftransmissionandpathogenesis, symptoms, laboratory diagnosis, preventionand control of diseases caused by - a. Mycobacteriumlepme b. Clostridiumperfringens, c. Treponemapallidum UNIT II Morphology, cultural and biochemical characteristics, antigenic structure, modes of transmission and pathogenesis, symptoms, laboratory diagnosis, preventionand control of diseases caused by - a. Pseudomonasaeruginusa b. Vibrod solem c. Stepharame mutars d. Hibiochacter pylori 1. UNIT III Morphology, culturaland biochemical characteristics, antigenic structure, modes of transmission and pathogenesis, symptoms, laboratory diagnosis, preventionand control of diseases caused by of transmission and pathogenesis, symptoms, laboratory diagnosis, prevention and control of diseases caused by a. Protozoa: Plasmodium falciparum (malaria) b. Virusesi) Hepatitis A & B virus	DSC XIV		Hours
Expected course outcome - Upon successful completion of course, students are expected to be able to - CO1: Correlate disease symptoms with causative agent, isolate and identify pathogens. CO2:Understand mechanism of action of antimicrobial drugs and their uses as prophylactic agents. CO3:Explain pathogenicity of organisms associated with human infections. CO4:Explain different antimicrobial agents with respect to their mode of action uses. 1. Morphology,culturalandbiochemicalcharacteristics,antigenicstructure,m odesoftransmissionandpathogenesis, symptoms,laboratorydiagnosis,preventionandcontrolofdiseasescausedb y- a. Mycobacteriumlepme b.Clostridiumperfringens, c. Treponemapallidum UNIT II Morphology, cultural and biochemical characteristics, antigenic structure, modes oftransmission and pathogenesis, symptoms,laboratorydiagnosis,preventionandcontrolofdiseasescausedb y- a. Pseudomonasaeruginosa b. Vibrio dukm c. Steptucaus mutans d.Hilichacteryplori 1. Morphology, culturalandbiochemicalcharacteristics, antigenicstructure, modes oftransmissionandpathogenesis, symptoms,laboratorydiagnosis,preventionandcontrolofdiseasescausedby- a. Protozoa: Plasmodium fidciparum(malaria) b. Virusesii) Hepatitis A & B virus	DSC0	MEDICAL MICROBIOLOGY	per
Expected course outcome - Upon successful completion of course, students are expected to be able to - CO1: Correlate disease symptoms with causative agent, isolate and identify pathogens. CO2:Understand mechanism of action of antimicrobial drugs and their uses as prophylactic agents. CO3: Explain pathogenicity of organisms associated with human infections. CO4: Explain different antimicrobial agents with respect to their mode of action uses. I. Morphology.culturalandbiochemicalcharacteristics, antigenicstructure, modesoftransmissionandpathogenesis, symptoms, laboratorydiagnosis, preventionand controlof diseases caused by - a. Mycobacteriumlepnae b. Clostridiumperfringens, c. Treponemapallidum UNIT II Morphology, cultural and biochemical characteristics, antigenic structure, modes oftransmission and pathogenesis, symptoms, laboratorydiagnosis, preventionand controlof diseases caused by - a. Pseudomonasaeruginosa b. Vibrio dyslen c. Sinykaracs mutars d. Hekiobacterysleri 1. UNIT III Morphology, culturaland biochemical characteristics, antigenic structure, modes oftransmissionand pathogenesis, symptoms, laboratory diagnosis, preventionand controlof diseases caused by - a. Protozoa: Plasmodium falciparum (malaria) b. Virusesti) Hepatitis A & B virus	3MIC	Theory: 30 Hours (Credits -2)	unit
Upon successful completion of course, students are expected to be able to – CO1: Correlate disease symptoms with causative agent, isolate and identify pathogens. CO2:Understand mechanism of action of antimicrobial drugs and their uses as prophylactic agents. CO3:Explain pathogenicity of organisms associated with human infections. CO4:Explain different antimicrobial agents with respect to their mode of action uses. I. Morphology,culturalandbiochemicalcharacteristics,antigenicstructure,m odesoftransmissionandpathogenesis, symptoms,laboratorydiagnosis,preventionandcontrolofdiseasescausedb y- a. Mycobacteriumlepme b. Clostridiumperfringens, c. Treponemapallidum UNIT II Morphology, cultural and biochemical characteristics, antigenic structure, modes oftransmission and pathogenesis, symptoms,laboratorydiagnosis,preventionandcontrolofdiseasescausedb y- a. Pseudomonasaeruginosa b. Vinio drulem c. Sinphoxoas multurs d. Hidiatracter pykni 1. UNIT III Morphology, culturalandbiochemicalcharacteristics, antigenicstructure, modes oftransmissionandpathogenesis, symptoms,laboratorydiagnosis, preventionandcontrolofdiseasescausedby- a. Protozoa: Plasmodium falciparum(malaria) b. Virusesi) Hepatitis A & B virus	63		
CO1: Correlate disease symptoms with causative agent, isolate and identify pathogens. CO2:Understand mechanism of action of antimicrobial drugs and their uses as prophylactic agents. CO3:Explain pathogenicity of organisms associated with human infections. CO4:Explain different antimicrobial agents with respect to their mode of action uses. I. Morphology, culturalandbiochemicalcharacteristics, antigenic structure, modes of transmission and pathogenesis, symptoms, laboratory diagnosis, prevention and control of diseases caused by a. Mycobacterium leprate b. Clostridium perfringens, c. Treponemapallidum UNIT II Morphology, cultural and biochemical characteristics, antigenic structure, modes of transmission and pathogenesis, symptoms, laboratory diagnosis, prevention and control of diseases caused by a. Pseudomonasaeruginosa b. Vibrio droken c. Singkucus mulius d. Heliculus terpylori I. Morphology, culturaland biochemical characteristics, antigenic structure, modes of transmission and pathogenesis, symptoms, laboratory diagnosis, prevention and control of diseases caused by a. Protozoa: Plasmodium falciparum (malaria) b. Virusesi) Hepatitis A & B virus	Expected co	urse outcome -	
CO2:Understand mechanism of action of antimicrobial drugs and their uses as prophylactic agents. CO3:Explain pathogenicity of organisms associated with human infections. CO4:Explain different antimicrobial agents with respect to their mode of action uses. 1.	Upon succes	ssful completion of course, students are expected to be able to -	
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CO3:Explain pathogenicity of organisms associated with human infections. CO4:Explain different antimicrobial agents with respect to their mode of action uses. 1. When the different antimicrobial agents with respect to their mode of action uses. 1. When the different antimicrobial agents with respect to their mode of action uses. 1. When the different antimicrobial agents with respect to their mode of action uses. 1. When the different antimicrobial agents with respect to their mode of action uses. 1. When the different antimicrobial agents with respect to their mode of action uses. 8. When the different antimicrobial agents with respect to their mode of action uses. 8. While the different antimicrobial agents with respect to their mode of action uses. 8. While the different antimicrobial agents with respect to their mode of action uses. 8. While the different antimicrobial agents with respect to their mode of action uses. 8. While the different antimicrobial agents with respect to their mode of action uses. 8. While the different antimicrobial agents with respect to their mode of action uses. 8. While the different antimicrobial agents with respect to their mode of action uses. 8. While the different antimicrobial agents with respect to their mode of action uses. 8. While the different antimicrobial agents with respect to their mode of action uses. 8. While the different antimicrobial agents with respect to their mode of action uses. 8. While the different antimicrobial agents with respect to their mode of action uses. 8. While the different antimicrobial agents with respect to their mode of action uses. 8. While the different antimicrobial agents with respect to their mode of action uses. 8. While the different antimicrobial agents with respect to their mode of action uses. 9. While the different antimicrobial agents with respect to their mode of action uses. 9. While the different antimicrobial agents with respect to their mode of action uses. 9. While the differen	CO2:Unders	stand mechanism of action of antimicrobial drugs and their uses as propl	ylactic
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symptoms,laboratorydiagnosis,preventionandcontrolofdiseasescausedb y- a. Mycobacteriumlepme b.Clostridiumperfringens, c. Trepenemapallidum UNIT II Morphology, cultural and biochemical characteristics, antigenic structure, modes oftransmission and pathogenesis, symptoms,laboratorydiagnosis,preventionandcontrolofdiseasescausedb y- a. Pseudomonasaeruginosa b. Vibrio dolem c. Shaptocaus mutans d. Helicobacter pylori 1. UNIT III Morphology, cultural and biochemical characteristics, antigenic structure, modes oftransmissionand pathogenesis, symptoms,laboratorydiagnosis, preventionandcontrolofdiseasescaused by- a. Protozoa: Plasmodium falciparum(malaria) b. Virusesii) Hepatitis A & B virus	UNIT I	Morphology,culturalandbiochemicalcharacteristics,antigenicstructure,m	
a. Mycobacteriumleprae b. Clostridiumperfringens, c. Treponemapallidum UNIT II Morphology, cultural and biochemical characteristics, antigenic structure, modes oftransmission and pathogenesis, symptoms,laboratorydiagnosis,preventionandcontrolofdiseasescausedb y- a. Pseudomonasaeruginosa b. Vibrio dulem c. Sheptoccus multurs d. Hdiaducterpylori 1. UNIT III Morphology, culturalandbiochemicalcharacteristics, antigenic structure, modes oftransmissionandpathogenesis, symptoms,laboratory diagnosis, preventionandcontrolofdiseasescausedby- a. Protozoa: Plasmodium falciparum(malaria) b. Viruses:i) Hepatitis A & B virus		odesoftransmissionandpathogenesis,	
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b.Clostridiumperfringens, c. Treponemapallidum UNIT II Morphology, cultural and biochemical characteristics, antigenic structure, modes oftransmission and pathogenesis, symptoms,laboratorydiagnosis,preventionandcontrolofdiseasescausedb y- a.Pseudomonasaeruginosa b. Vibrio dolena c. Streptococus mutans d.Hdiachacter pylori 1. UNIT III Morphology, cultural and biochemical characteristics, antigenic structure, modes oftransmissionand pathogenesis, symptoms, laboratory diagnosis, preventionand controlof diseases caused by- a. Protozoa: Plasmodium falciparum (malaria) b. Viruses:i) Hepatitis A & B virus		y-	
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a. Pseudomonasaeruginosa b. Vibrio drolem c. Streptococus mutuus d. Helicobacter pylori 1. UNIT III Morphology, culturaland biochemical characteristics, antigenic structure, modes of transmission and pathogenesis, symptoms, laboratory diagnosis, prevention and control of diseases caused by-a. Protozoa: Plasmodium falciparum (malaria) b. Viruses:i) Hepatitis A & B virus		modes of transmission and pathogenesis,	
a. Pseudomonasaeruginosa b. Vibrio drolem c. Streptococcus mutans d. Hdiobacter pylori 1. UNIT III Morphology, culturaland biochemical characteristics, antigenics tructure, modes of transmission and pathogenesis, symptoms, laboratory diagnosis, prevention and control of diseases caused by- a. Protozoa: Plasmodium falciparum (malaria) b. Viruses:i) Hepatitis A & B virus		symptoms,laboratorydiagnosis,preventionandcontrolofdiseasescausedb	
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c. Streptococus mutans d. Helicobacter pylori 1. UNIT III Morphology, cultural and biochemical characteristics, antigenic structure, modes of transmission and pathogenesis, symptoms, laboratory diagnosis, prevention and control of diseases caused by- a. Protozoa: Plasmodium falciparum (malaria) b. Viruses:i) Hepatitis A & B virus		a.Pseudomonasaeruginosa	
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1. UNIT III Morphology,culturalandbiochemicalcharacteristics,antigenicstructure,modes oftransmissionandpathogenesis, symptoms,laboratorydiagnosis,preventionandcontrolofdiseasescausedby- a. Protozoa: Plasmodium falciparum (malaria) b. Viruses:i) Hepatitis A & B virus		cStreptococcus mutans	
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symptoms,laboratorydiagnosis,preventionandcontrolofdiseasescausedby- a. Protozoa: <i>Plasmodium falciparum</i> (malaria) b. Viruses:i) Hepatitis A & B virus	UNIT III	Morphology,culturalandbiochemicalcharacteristics,antigenicstructure,modes	8
a. Protozoa: Plasmodium falciparum (malaria) b. Viruses:i) Hepatitis A & B virus		oftransmissionandpathogenesis,	
b. Viruses:i) Hepatitis A & B virus		symptoms, laboratory diagnosis, prevention and control of diseases caused by-	
		a. Protozoa: <i>Plasmodium falciparum</i> (malaria)	
ii) Rabiesvirus		b. Viruses:i) Hepatitis A & B virus	
l l		ii) Rabiesvirus	

	iii) Dengue virus	
	c Fungi:Candidaalbicans	
UNIT IV	1.Chemotherapy	7
	a.Generalprinciplesofchemotherapy	
	b. Modeof action of Penicillin, Streptomycin, Bacitracin,	
	,sulphonamideandQuinolones onmicroorganisms.	
	c.Antiviraldrug:AZT	
	d. Antifungal drugs:Ketoconazole	
	e. Antiprotozoal drugs: Metronidazole	
	f. Mechanismofdrugresistance	
	g. Chemoprophylaxis	
	2. Genetherapy-Concept, advantages & disadvantages.	
	3. Immunoprophylaxis–VaccinesandImmuneSera	
	a. Vaccines-liveattenuated, heat	
	killed, subunit, conjugate and DNA vaccines	
	b. ImmuneSera-exampleswithapplications	

Books Recommended:

B. For Medical Microbiology

- 1) Microbiology -Davis
- 2) Immunology&serology-AshimChakravarty
- $3) \ Medical Microbiology 16 {}^{th} edition by David Greenwood, Richard CBS lack, John Peutherer \\$
- 4) MedicalBacteriology-Dey&Dey
- 5) MedicalBacteriologyincludingMedicalMycology&AIDS-NCDey&T.K.Dey
- ${\bf 6)}\ Principals and Practice of Clinical Bacteriology-A.M. Emmerson$

		No. of
DSE-II	ENVIRONMENTALMICROBIOLOGY	Hours
DSE03MIC61	Theory: 30 Hours (Credits -2)	per
		unit
Expected cours	se outcome -	
Upon successf	ul completion of course, students will be able to -	
CO1: Understa	and the basic principle of environment microbiology and be ab	le to
apply the	ese principles to understanding and solving environmental pro	blems.
CO1: Know th	e Microorganisms responsible for water pollution and their	
transmis	sion	
CO1: Describe	classification of lakes, sources, consequences and control of	
eutrophi	cation.	
CO1: Explain	various bioburden test and clean room concepts.	
	1.General characteristics of waste-	7
UNIT I	a. Liquid waste - pH, electrical conductivity, COD,	
	BOD, total solids, total dissolved solids, total	
	suspended solids, total volatile solids, chlorides,	
	sulphates, oil &grease.	
	b. Solid waste- pH, electrical conductivity, total volatile	
	solids,ash.	
	c. Standards as perMPCB	
	2.Eutrophication	
	a. Classification oflakes	
	b. Sources c. Consequences	
	d. Control	
UNIT II	1.SewageMicrobiology	8
	a. Physico-chemical and biologicalcharacteristics	
	b. Treatment methods-	
	i. Physical treatment: Screening, Sedimentation	
	ii. Biological treatment: Trickling filter, Activated	
	sludge process, Oxidationponds, Anaerobic	
	digestion (Biomethanation), Septic tank.	

iii. Chemical treatment - Chlorination

2. Characteristics and treatment of waste generated by Hospitals

1. Biological safety inlaboratory	
UNIT III a. Good LaboratoryPractices 7	
b. Bio safety levels(BSL)	
2. Environmental monitoring	
a. Definition and purpose	
b. Cleanroom- Concept, classification, prevention of	
contamination in clean rooms	
c. Routine Environmental monitoring	
programme in pharmaceutical industries-	
Air monitoring, Surface monitoring and	
Personnelmonitoring.	
d. Bioburdentest	
3. Environmental Impact Assessment-	
Concept and Brief introduction	
UNIT IV 1. Bioremediation and Bioleaching 8	
a. Bioremediation	
i. Definition	
ii. Types	
iii. Applications.	
b. Bioleaching	
i. Introduction	
ii. Microorganismsinvolved	
iii. Chemistry of Microbialleaching	
iv. Laboratory scale and pilot scaleleaching	
v. In situ leaching - Slope,heap	
vi. Leaching of Copper and Uranium	

BooksRecommended

- 1. BiochemistryandMicrobiologyofPollution-HigginsandBurns.
- 2. Waste Water Treatment-Datta and Rao (Oxford and IBH)
- 3. EnvironmentChemicalHazards-RamKumar(SwarupandSons,NewDelhi).
- 4. EnvironmentPollution-TimmyKatyal(SatkeAnmolPub.NewDelhi).
- 5. EcologyofPollutedWater-Vol.II-AnandKumar(AphPub.Co.NewDelhi).
- 6. EnvironmentPollutionandManagementofwastewatersbyMi crobialTechniques- PathadeandGoel(ABDPub.Jaipur)

	erunit/ edit
	miomo
-	amisms.
•	
A. Industrialproduction- OrganismsusedInoculumpreparation,Fermentation media,Fermentationconditions, ExtractionandRecovery.	7
 Primary metabolite: i) Vitamin: Vitamin B12, riboflavin, β carotene, ii) Aminoacids: Lysine & Glutamic acid iii) Organic acid: Citricacid & Lactic acid acetic acids, lactic acids, kojic acids, Itaconic acids i) 	
 2. Secondary metabolite i) Antibiotics: a. Penicillin & semi-synthetic penicillin b. Streptomycin ii) Alcoholic Beverages: Wine: a) Red Table Wine b) Sparkling Wine- Champagne 3. Enzyme: ii) Amylase iii) Protease iv) lipase 	8
 A. Production of biofuels Bioethanol- microorganisms used, fermentation condition recovery, purification of Ethanol Biogas- Biomass used, Microbiology & Biochemistry of biogas production, Biodiesel production from algae Brobiotics- Concept, Production by using Lactobacillus and applications Production of SCP by using yeast 	8
	mes- mofcourse, students will be ableto- ethods used for industrial product recovery after fermentation. tand the importance of fermentation economics. computer applications in downstream processing. A. Industrial production- Organisms used Inoculum preparation, Fermentation media, Fermentation conditions, Extraction and Recovery. 1. Primary metabolite:

FERMENTATION TECHNOLOGY II

No.of

Hours

DSE-II

DSE03MIC61

UNIT IV	A. Fermentation economics – discovery and process development,	7
	strain improvement, market potential, plant and equipment,	
	operating cost, contract manufacturing, return on investment –	
	recovery cost. Water usage and recycling and effluent treatment.	
	B. Computer applications in downstream processing - Introduction,	
	History, General specific applications, System configuration. Product	
	formulation, monitoring of downstream processing, process	
	integration.	

		No. of
MIN-II	ENVIRONMENTALMICROBIOLOGY	Hours
MIN03MIC21	Theory: 30 Hours (Credits -2)	per uni
Expected course or	utcome -	
Upon successful c	ompletion of course, students will be able to –	
CO1: Understand	the basic principle of environment microbiology and be able to	apply
these principles to	understanding and solving environmental problems.	
CO1: Know the M	icroorganisms responsible for water pollution and their transmi	ission
CO1: Describe clas	ssification of lakes, sources, consequences and control of eutrop	hication.
CO1: Explain vario	ous bioburden test and clean room concepts.	
	1.General characteristics of waste-	7
UNIT I	a. Liquid waste - pH, electrical conductivity, COD,	
	BOD, total solids, total dissolved solids, total	
	suspended solids, total volatile solids, chlorides,	
	sulphates, oil &grease.	
	b. Solid waste- pH, electrical conductivity, total volatile	
	solids,ash.	
	c. Standards as perMPCB	
	2. Eutrophication	
	a. Classification oflakes	

UNIT II

2.SewageMicrobiology

b. Sources

d. Control

c. Consequences

a. Physico-chemical and biological characteristics

8

- b. Treatment methods-
- i. Physical treatment: Screening, Sedimentation
 - ii. Biological treatment: Trickling filter, Activated sludge process, Oxidationponds, Anaerobic digestion (Biomethanation), Septic tank.
 - iii. Chemical treatment Chlorination

	1. Biological safety inlaboratory	
UNIT III	a. Good LaboratoryPractices	7
	b. Bio safety levels(BSL)	
	2. Environmental monitoring	
	a. Definition and purpose	
	b. Cleanroom- Concept, classification, prevention of	
	contamination in clean rooms	
	c. Routine Environmental monitoring programme	
	in pharmaceutical industries-	
	Air monitoring, Surface monitoring and	
	Personnelmonitoring.	
	d. Bioburdentest	
	3. Environmental Impact Assessment-	
	Concept and Brief introduction	
UNIT IV	1. Characteristics and treatment of waste generatedby	8
	Hospitals	
	2. Bioremediation and Bioleaching	
	a. Bioremediation	
	i. Definition ii. Types	
	iii. Applications.	
	b. Bioleaching	
	i. Introduction	
	ii. Microorganismsinvolved	
	iii. Chemistry of Microbialleaching	
	iv. Laboratory scale and pilot scaleleaching	
	v. In situ leaching - Slope,heap	
	vi. Leaching of Copper and Uranium	

BooksRecommended

- $1. \ Environmental Pollution by Chemicals-Walker, Hulchiason.$
- $2.\ Biochemistry and Microbiology of Pollution-Higgins and Burns.$
- $3.\ Environmental Pollution-Laurent Hodge, Holt.$
- 4. WasteWaterTreatment-DattaandRao(OxfordandIBH)
- 5. Sewageandwastetreatment-Hammer
- 6. EnvironmentChemicalHazards-RamKumar(SwarupandSons,NewDelhi).
- $7.\ Environment Pollution-Timmy Katyal (Satke Anmol Pub. New Delhi).$

DSC-PR-VI	DSC Microbiology Lab-VI
DSC03MIC69	(Credits -4)
	PRACTICALS BASED ON GENETICS
	Major:
	1. Effectof U.V. lighton bacteria and graphical presentation of result.
	2. Isolationofauxotrophicmutantsbyreplicaplatetechnique
	3.Transferofgeneticmaterialbytransformationin <i>E. coli</i>
	4.IsolationofchromosomalDNAfrombacteria(J.Marmursmethod)
	5Isolationofstreptomycin-resistantmutants(gradientplatetechnique)
	Minor:
	1.Electrophoretic separation of DNA.
	2.Spectrophotometric analysis of DNA in extracted solution.
	3. Demonstration of PCR.
	4. Isolation of Lac negative mutants of <i>E. coli</i>
	5. Testing of carcinogenicity of a substance by Ames test.
	PRACTICALS BASED ON BIOCHEMISTRY
	Major:
	AssayofamylasebyDNSAmethod(graphicalestimation)
	2. Immobilization of enzymes by sodium alginate method.
	3. Bio-assay of Vitamin B12
	4. Bio-assay of Penicillin.
	5. Protein purification by using ammonium sulfate precipitation.
	Minor:
	1.Seperation and detection of amino acid by TLC
	2. Effect of activator on enzyme activity
	3. Effect of inhibitor on enzyme activity.
	4. Effect of pH on enzyme activity.
	5. Effect of temperature on enzyme activity
]	PRACTICALS BASED ON MEDICAL MICROBIOLOGY
	Major:
	1.Isolation of <i>Pseudomonasaeruginosa</i> from clinical samples (wherever
	possible) and identification of the same by morphological, cultural and

biochemical characteristics.

- 2. Isolation of *Klebsiella pneumoniae* from clinical samples (wherever possible) and identification of the same by morphological, cultural and biochemical characteristics.
 - 3. Isolation of *Candidaalbicans* from clinical samples (wherever possible) and identification of the same by morphological, cultural and biochemical characteristics.
- 4. Determination of MIC of streptomycin against E.coli by broth method .
 - 5. Determination of sensitivity of common pathogens to antibiotics by paper disc method .

Minor:

- 1. Widaltest Quantitative
- 2. Physicalandchemicalexaminationofurine.
- 3. Detection of presence of sugar in urine.
 - 4. Detection of presence of protein in urine (Aceticacidtest)
 - 5. Detection of presence of ketonebodies(Rothra'stest)
 - 6. Detection of presence of bilesalt.

DSE-PR-II	PRACTICALS BASED ON FERMENTATION TECHNOLOGY-II
	Major
	Isolation and identification of probiotic micro flora from natural sources or any commercial formulation
	2. Production of Biogas from organics waste
	3. Production of alcohol from molasses
	4. Screening of organic acid producers & amine producers
	5. Screening of Amylase, Protease & Lipase producers
	6. Screening of Vitamin producers.
	Minor:
	1.Chemical assay of Vitamin C.
	2.Chemical assay of Penicillin.
	3.Estimation of alcohol
	4.Examination of milk by Directmicroscopic count (DMC) 5.Sauerkrautproduction.

DSE-PR-II	PRACTICALS BASED ON ENVIRONMENTAL MICROBIOLOGY
	Major:
	1. Microbial testing of Water:
	Presumptive,confirmedandcompletedtest.
	2. Determination of Most Probable Number of Water
	3. Determination of COD of sewage.
	4. DeterminationofBODofsewage
	Minor:
	1.Determination of color&pH of water.
	2.Determination of total alkalinity of water
	3.Determination of chloride content of water

MIN-PR-II	ENVIRONMENTAL MICROBIOLOGY (Credits -4)
	Major:
	1. Microbial testing of Water:
	a. Presumptive,confirmedandcompletedtest.
	2. Determination of Most Probable Number of Water
	3. Determination of COD of sewage.
	4. DeterminationofBODofsewage
	Minor:
	5. Determination of color& pH of water.
	6. Determination of total alkalinity of water
	7. Determination of chloride content of water

VSC-PR-IV VSC03MIC69	WASTE WATER MANAGEMENT (Credits -2)	No. of Hours per unit					
Expected course	outcome -						
On completion o	f course, student will be able to –						
i. Explain physical and chemical characteristics of waste water							
ii. Examine various types of solid waste in water and categorize it.							
	1) Determination of color & pH and Temperature of Water						
	2) Determination total alkalinity of water						
	3) Determination of electric conductivity of water						
	4) Determination of Specific gravity of water						
	5) Determination of oil and grease content of water						
	6) Determination of Total solid content of water						

Practical Examination

- A) The practical examination will be conducted on three (3) consecutive days for not less than 6hours on each day of the practical examination.
- B) Each candidate must produce a certificate from the Head of the Department in his/her college stating that he/she has completed in a satisfactory manner the practical course on the guidelines laid down from time to time by Academic Council on the recommendation of Board of studies and has been recorded his/her observations in the laboratory journal and written a report on each exercise performed. Every journal is to be checked and signed periodically by a member teaching staff and certified by the Head of the Department at the end of staff and certified by the Head of the Department at the end of the semester. Candidates are to produce their journal at the time of practical examination.
- C) Nature of question paper and distribution of marks for B.Sc. Part III MicrobiologyPractical Examination

PracticalsPaper DSC I, II, III &DSE I/II

Semester V

Section I/II/III/ and DSE-PRIV

Q.1MajorExperiment 10Marks
Q. 2MinorExperiment 05Marks
Q.3Journal 05Marks
Q.4 Seminar 10Marks

Semester VI

Section I/II/III/ and DSE-PRIV

Q.1MajorExperiment 10Marks
Q. 2MinorExperiment 05Marks
Q.3Journal 05Marks
Q.4Seminar 10Marks

3)Draw neat labeled diagrams wherever necessary. 4) Use of calculator is allowed. Time: 2 hours **Total Marks: 40** PAPER DSC IX/X/XI/DSE I/II Q.1.A Select correct alternative. **(8)** i) b) d) a) c) ii) a) b) c) d) iii) a) b) c) d) iv) d) a) b) c) v) b) c) d) a) vi) a) b) c) d) vii) a) b) c) d) viii) a) b) c) d) Q.2. Attempt any Two (16)i) ii) iii) Q.3. Attempt any Four **(16)** i) ii) iii) iv) v) vi)

Instructions: 1) All the questions are **compulsory**.

2 Figures to the right indicate **full** marks.

Instruction to paper setters: Equal weight age should be given to all units.

*Select any one for B.Sc.III ---- (10 marks) 1) Unit test

- 2) Home assignment
- 3) Project
- 4) Seminar

SCHEME OF MARKING (THEROY)

Sem.	Core	Marks	Evaluation	Paper	Answer	Standard of
	Course				Books	passing
			_			
V	DSCIX	40	Semester wise	Each paper of	As per	40%
				40 marks	Instruction	(16 marks)
V	DSCX	40	Semester wise	Each paper of	As per	40%
				40 marks	Instruction	(16 marks)
V	DSCXI	40	Semester wise	Each paper of	As per	40%
				40 marks	Instruction	(16 marks)
V	DSE-I	40	Semester wise	Each paper of	As per	40%
				40 marks	Instruction	(16 marks)
						, ,

SCHEME OF MARKING (CIE) Continuous Internal Evaluation

Sem.	Core Course	Marks	Evaluation	Paper	Answer Books	Standard of passing
VI	DSC XII	40	Semester wise	Each paper of 40 marks	As per Instruction	40% (16 marks)
VI	DSC XIII	40	Semester wise	Each paper of 40 marks	As per Instruction	40% (16 marks)
VI	DSC XIV	40	Semester wise	Each paper of 40 marks	As per Instruction	40% (16 marks)
VI	DSE-II	40	Semester wise	Each paper of 40 marks	As per Instruction	40% (16 marks)

SCHEME OF MARKING (PRACTICAL)

Sem.	Course	Marks	Evaluation	Paper	Sections	Standard of passing
V AND VI	Practical I,II,III&IV	150	Semester Wise	Four	As per Instruction	45%

*A separate passing is mandatory