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University of Mysore
(Estd.1916)

M.Sc. MICROBIOLOGY

**Credit Based Choice
Based Continuous
Evaluation Pattern
System**



UNIVERSITY OF MYSORE

Department of Studies in Microbiology

Manasagangotri, Mysuru-570006

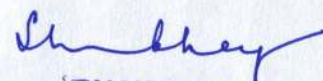
Regulations and Syllabus

Master of Science in Microbiology (M.Sc.)

(Two-year semester scheme)

Under

**Credit Based Choice Based Continuous Evaluation Pattern
System**



**CHAIRMAN
BOARD OF STUDIES
MICROBIOLOGY (PG)
UNIVERSITY OF MYSORE**

UNIVERSITY OF MYSORE
GUIDELINES AND REGULATIONS LEADING TO
MASTER OF SCIENCE IN MICROBIOLOGY
(TWO-YEAR SEMESTER SCHEME) UNDER
CREDIT BASED CHOICE BASED CONTINUOUS EVALUATION PATTERN SYSTEM

Programme Details

Name of the Department	: Department of Studies in Microbiology
Subject	: Microbiology
Faculty	: Science and Technology
Name of the Programme	: Master of Science in Microbiology (M.Sc.)
Duration of the Programme	: 2 years divided into 4 semesters

Programme Outcomes

On completion of M.Sc. (Microbiology), students are able to:

1. Instill the intellectual skills to analyze the molecules using advance biophysical techniques such as HPLC, GC, AAS, PCR etc.
2. Perform the quantitative/ qualitative analysis of biomolecules and understand various biochemical pathways
3. Acquire knowledge and understanding the concepts of microbial genetics, molecular biology, Immunology, Enzymology.
4. Explore the scientific literature effectively and use computational tools such as bio-statistics and bioinformatics
5. Implement the knowledge in industry with regard to scale up, production, down stream processing and quality control of the various microbial products
6. Conduct the basic research related to industry-environmental issues and use of agricultural for sustainable products



Programme Specific Outcomes

1. Understand and be able to explain different branches of microbiology such as bacteriology and virology.
2. To develop laboratory skills applicable to Microbiological research in the field of environmental microbiology, industrial microbiology, food microbiology and microbial pathogenesis.
3. Design and execute experiments related to basic microbiology, immunology, molecular biology, recombinant DNA technology and microbial genetics, which will enable to execute a short research project under the supervision of teachers.
4. Equip to take up a suitable position in academia or industry, and to pursue a career in research if so desired.

University of Mysore
Department of Studies in Microbiology
Credit Based Choice Based Continuous Evaluation Pattern System

SCHEME OF THE STUDY

For B.Sc. (Honors) in Microbiology

Credits to be earned	: 40 credits
Hard core papers	: 16 credits
Soft core paper	: 16 credits
Open elective paper	: 04 credits
Project work / term work	: 04 credits

For M. Sc. in Microbiology

Credits to be earned	: 76 credits
Cumulative total of credits to be completed	: 40 credits (Honors) + 36 credits
Hard core papers	: 36 credits
Soft core papers	: 32 credits
Project work / term work	: 08 credits

M.Sc. in Microbiology (76 credits)
Credit Based Choice Based Continuous Evaluation Pattern System
Proposed Semester-wise distribution of the course structure for the year 2018-2019

First Semester: 20 Credits

No	Paper Code	Title Of The Course Paper	HC/S C/ OE	L:T:P	Credits
1	18251	Virology	HC	3:1:0	4
2	18252	Bacteriology	HC	3:1:0	4
Select 4 out of 5 papers					
3	18253	Mycology	SC	3:1:0	4
4	18254	Microbial Genetics	SC	3:1:0	4
5	18255	Microbial Ecology & Diversity	SC	3:1:0	4
6		Practical I (Virology & Bacteriology)	SC	0:0:2	2
7		Practical II (Mycology & Microbial Genetics)	SC	0:0:2	2

HC=02; SC=04; OE=0

Second Semester: 20 Credit

No	Paper Code	Title Of The Course Paper	HC/SC /OE	L:T:P	Credits
1	18261	Microbial Physiology	HC	3:1:0	4
2	18262	Immunology	HC	3:1:0	4
Select 3 out of 4 papers					
3	18263	Food Microbiology	SC	3:1:0	
4	18264	Soil Microbiology	SC	3:1:0	4
5		Practical III (Microbial Physiology & Immunology)	SC	0:0:2	2

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6		Practical IV (Food Microbiology)	SC	0:0:2	2
7	18265	Microbial Diversity	OE	2:2:0	4

HC=02; SC=03; OE=1

Third semester: 20 credit

NO	Paper Code	Title Of The Course Paper	HC/SC /OE	L:T:P	Credits
1	18271	Molecular Biology	HC	3:1:0	4
2	18272	Genetic Engineering	HC	3:1:0	4
3	18273	Industrial Microbiology	HC	3:1:0	4
Select 3 out of 4 papers					
4	18274	Medical Microbiology	SC	3:1:0	4
5	18275	Clinical & Diagnostic	SC	3:1:0	4
6		Practical V (Molecular Biology & Genetic Engineering)	SC	0:0:2	2
7		Practical VI (Industrial Microbiology & Medical Microbiology)	SC	0:0:2	2
8	18276	Techniques in Microbiology	OE	1:1:0	2

HC=02; SC=03; OE=01

Fourth Semester: 18 credit

NO	Paper Code	Title Of The Course Paper	HC/SC /OE	L:T:P	Credits
1	18281	Agricultural Microbiology	HC	3:1:0	4
Select 2 out of 3 papers					
2	18282	Environmental Microbiology	SC	3:1:0	4

3	18283	Genomics & Proteomics	SC	3:1:0	4
4		Practical VII (Agricultural Microbiology & Environmental Microbiology)	SC	0:0:2	2
5		Project Work	HC	0:2:6	8

HC=01; SC=02; PW=01

Grand Total Credits: 76

FIRST SEMESTER

HARDCORE

COURSE-I: VIROLOGY

48 Hr

Course Outcome

Upon successful completion of the course, the student would have gained knowledge regarding the following aspects;

- classification of viruses
- Tools and methods employed for studying viruses mode of transmission, patterns of infection, virus virulence, and host defense against virus infection
- Knowledge about existing viral vaccines and anti-viral drugs, emerging viruses.

Pedagogy

Virology is a sub-discipline of Microbiology which focuses on the basic knowledge of viruses, their reproduction and pathogenesis within a host cell. The contents of the course are divided into four units. This course will emphasize basic concepts of viruses, their diversity (structural, host range and genetic), various methods employed in virus studies, replication strategies, host virus interactions, emerging viral diseases, antiviral strategies in prevention and control of viral diseases and their applications, evolution and future prospective. These concepts are taught across thirty two hours, which include lectures in which students are provided information and illustrations of various virus aspects, and tutorials in which students present seminars of the topics assigned. The lecture will impart students with knowledge and make them understand how viruses are built, replicate and evolve, cause disease, prevention of infection and their applications. The knowledge gained about viruses helps the students to develop interest in this field and helps the students to work in laboratories that are interested in isolation of viruses from various ecological niche, molecular pathogenesis, vaccines, antiviral drugs, applications such as virus vectors development and in health and diagnostic laboratories.

Course Content:

48 Hr

Signature

UNIT-I

12 Hr

The science of virology: Concept and scope of virology. Definitive properties of viruses: Morphology, Ultra structure, Chemical composition - proteins, nucleic acids, and other contents. Classification and nomenclature of viruses. Evolutionary importance of viruses.

Working with viruses: Visualization and enumeration of virus particles, Biological activity of viruses, Physical and chemical manipulation of the structural components of viruses, Characterization of viral product expressed in the infected cells. Isolation and purification of viruses, Detection of viruses: physical, biological, immunological and molecular methods.

UNIT-II

12 Hr

Virus replication Strategies: Principal events involved in replication: Adsorption, penetration, uncoating nucleic acid and protein synthesis, intracellular trafficking, assembly, maturation and release, viral-host interaction, Host response to viral infection.

Replication patterns of specific viruses: Identification of virus prototypes associated with different virus replication schemes; Details on important viruses namely Herpes virus, Poliovirus, Influenza virus, SV40 and Adeno Virus, Poxviruses, Hepatitis Viruses, Retroviruses.

UNIT-III

12 Hr

Propagation, purification, characterization and identification and genomics of plant viruses: General methods of propagation of plant viruses; purification of plant viruses using centrifugation, chromatography and electrophoresis techniques, methods employed in identification of plant viruses.

Sub-viral particles: Discovery, Structure, Classification, replication and diseases caused by Satellite, Satellites virus, Virusoids, Viroids and Prions.

Anti-viral strategies-prevention and control of viral diseases: Host specific and Non specific defense mechanisms involved in resistance to and recovery from virus infections. Role of interferon in viral infections. Viral Chemotherapy: Nucleoside analogs, reverse transcriptase inhibitors, protease inhibitors, History of vaccines especially smallpox and polio. New methods: subunit vaccines, antiidiotype and DNA vaccines.

UNIT-IV

12 Hr

Microbial viruses: Diversity, classification, characteristics and applications of bacteriophages, and general account on algal, fungal and protozoan viruses.

Viruses and the future: Promises and problems. Emerging diseases, sources and causes of emergent virus diseases.

References:

1. Marc H.V. van Regenmortel , Brian W.J. Mahy (2009) Desk Encyclopedia of General Virology , 1 edition, AcademicPress.
2. Alan J. Cann (2011) Principles of Molecular Virology, 5th edition ,Elsevier
3. Clokie, Martha R. J., Kropinski, Andrew (2009) Bacteriophages, Methods and Protocols, Volume 1: Isolation, Characterization, and Interactions, HumanaPress

4. Edward K. Wagner, Martinez J. Hewlett , David C. Bloom , David Camerini (2007), Basic Virology, 3rd Edition, John Wiley & Sons.
5. Hunter-Fujita, Frances R., Philip F. Entwistle, Hugh F. Evans, and Norman E. Crook. Insect viruses and pest management. John Wiley & Sons Ltd, 1998.
6. Jane S. Flint, Lynn W Enquist, Anna Marie Shalka (2004) Principles of Virology: Molecular Biology, Pathogenesis, and Control of Animal Viruses, American Society for Microbiology
7. John Carter, Venetia A. Saunders, (2007), Virology: Principles and Applications. John Wiley & Sons, west Sussex, England.
8. Lobočka, Malgorzata, and Waclaw T. Szybalski, eds. (2012) Bacteriophages. Part 2 , Academic Press
9. Matthews, Richard Ellis Ford, and Roger Hull. (2002) Matthews' plant virology. 4th edition, Gulf Professional Publishing.
10. Nigel Dimmock, Andrew Easton, Keith Leppard, 2009, Introduction to Modern Virology, 6th Edition, Wiley-Blackwell.

COURSE-II: BACTERIOLOGY

48 Hr

Course Outcome:

Upon successful completion of the course, the student:

- Are able to describe the morphological features, cell arrangement and structural components of bacterial cell in detail and differentiate between Gram-positive and Gram-negative bacteria.
- Will have gained knowledge about cell wall structure and extracellular appendages in different bacteria and will be acquainted with current methodologies available for production of protoplasts and L-forms
- Know key features of some model archaeal organisms.
- Understand Quorum sensing in bacteria and its significance in competence, sporulation and antibiotic resistance.

Pedagogy:

Bacteriology is the sub-disciplinary course of Microbiology. Bacteriology is the study of bacteria and their importance in medicinal and other areas such as agriculture, industries. Bacteria are single celled microorganisms which can live as independent organisms or dependently as parasites. Superficially bacteria appear to be relatively simple forms of life; in fact they are sophisticated and highly adaptable.

The discipline of bacteriology evolved from the need to test and apply the germ theory of disease and from economic concern relating to the spoilage of food and wine. The contents of this course are divided into various units and each unit focuses on various aspects of bacteriology such as serial dilution technique, microscopy, staining technique, classification and taxonomy and economic importance of bacteria.

The knowledge gained under this subject helps the student to work in laboratories, pharmacological industries, and in any industries where microorganisms are used as chief source.

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Course Content

UNIT-I

12 Hr

Introduction: Important events in development of bacteriology, Scope and relevance of bacteriology. Economic importance of bacteria.

Cell Structure: An overview of bacterial size, shape and arrangement, structure, chemical composition of cell wall of Archaeobacteria, gram-negative bacteria, gram-positive bacteria and acid fast bacteria, cell wall deficient organisms including L-form structure, composition and function of cell membrane, capsule, flagella, pili, Inclusion bodies, ribosomes, mesosomes, reserve food materials, magnetosomes and phycobilisomes, endospores, bacterial nucleic acids – chromosome, plasmid, transposons, integrons and antibiotic resistance cassettes.

Microscopy: Working Principles of bright field microscope, fluorescent microscope, dark field microscope, phase contrast microscope, stereo microscope, confocal microscopy and electron microscope. Preparation of sample for electron microscopic studies. Application and importance of above microscopes. Measurement of microscopic objects.

UNIT-II

12 Hr

Bacterial classification and taxonomy: Criteria for the classification of bacteria. Phenetic, Phylogenetic, Genotypic, Numerical taxonomy. Techniques for determining microbial taxonomy and Phylogeny. ICNB rules. Classification systems of major categories and groups of bacteria according to Bergey are manual of Systematic Bacteriology and Determinative Bacteriology. Non-culturable methods for the identification of pathogenic microorganisms.

UNIT-III

12 Hr

Growth, Cultivation and control of Bacteria: Nutrient requirements, nutritional types of bacteria, culture media, classification of media. Growth: Nutritional uptake, Growth kinetics, generation time, growth curve, factors affecting growth. Methods for measurement of microbial growth – direct microscopy, viable count estimates, turbidometry, and biomass. Aerobic, anaerobic, batch, continuous and synchronous cultures. Methods of pure culture isolation, Enrichment culturing techniques, single cell isolation, and pure culture development. Preservation and Maintenance of Microbial cultures: Repeated sub culturing, preservation at low temperature, sterile soil preservation, mineral oil preservation, deep freezing and liquid nitrogen preservation, lyophilization. Advantages and disadvantages of each method. Control of

microorganisms: Antimicrobial agents, physical and chemical methods. Principles, functioning and types of Biosafety cabinets.

UNIT-IV

12 Hr

Characteristics and Salient features of major groups of Bacteria: Archaeobacteria: general characteristics and classification; extremophiles, halophiles, thermophiles and barophiles; General characteristics, classification, diversity and distribution, economic importance of **Actinomycetes, Cyanobacteria. Bioluminescent bacteria;** characteristics and examples, mechanism of bioluminescence. General characteristics, life cycle, growth, multiplication and significance of **Mycoplasma, Rickettsiae and Chlamydia**

References:

1. Sherwood, and Woolverton Willey (2007), Prescott, Harley, and Klein's Microbiology (7th International Edition), McGraw-Hill
2. Mara, Duncan, and Nigel J. Horan, (2003). Handbook of water and wastewater Microbiology, Academic Press.
3. Michael T. Madigan, David P. Clark, David Stahl, John M. Martinko, 2012, Brock Biology of Microorganisms 13th Edition, Benjamin Cummings
4. Jacquelyn G. Black (2012) Microbiology: Principles and Explorations ,8th edition, Wiley
5. Michael J. Leboffe, Burton E. Pierce , David Ferguson (2012) Microbiology Laboratory Theory & Application, Brief, 2nd Edition, Morton Publishing Company
6. Jeffrey C. Pommerville (2010) Alcamo's Fundamentals of Microbiology, 9th Revised edition, Jones and Bartlett Publishers, Inc
7. Jeffrey C. Pommerville (2010) Alcamo's Laboratory Fundamentals of Microbiology, Jones and Bartlett Publishers, Inc
8. Alfred Brown (2011) Benson's Microbiological Applications Short Version (Brown, Microbiological Applications), 12th edition, McGraw-Hill Science/Engineering/Math.
9. Jerome J. Perry , James Staley , Stephen Lory (2002), Microbial Life, Sinauer Associates.
10. Stuart Hogg (2013) Essential Microbiology, 2nd Edition, Wiley-Blackwell

COURSE- III: MYCOLOGY

48 Hr

Course outcome:

- To understand the world of fungi and lichens,
- Appreciate the adaptive strategies of the fungi and lichens,
- understand the economic and pathological importance of fungi, and identify common plant diseases and devise control measures
- In tutorial classes they do group discussions on use full and pathogenesis of different fungi involved in daily life.

- In practical classes they mount the fungi, learn microscopic views and the key characteristics to identify different species of fungi.

Pedagogy:

Mycology is the sub-branch of Microbiology, which is concerned with the study of fungi. It includes the study of taxonomic classification, fungal genetics, and biochemical properties. Fungi are fundamental for life as symbionts, also takes part in biodegradation process. They are socially and economically important as they are capable of causing diseases in plants, animals and human beings.

Study of fungi is highly important as it plays major role in production of food supplements like SCP, fermentation industries, vitamins, enzymes, organic acids. Another notable element is production of secondary metabolites like antibiotics which acts against other microbes. In agriculture, knowledge pertaining to fungi should be maximum as it causes plant diseases leading to economic loss. Fungal infections have more devastating effects on human health and hence clinical significance of fungi has gained more attention, due to its wide applications and effects, the study of fungi is highly recommended.

Course Content

UNIT-I **12 Hr**

Introduction: History and Development of Mycology, scope of mycology. Recent developments in Mycology.

Fungal taxonomy: Taxonomic problems associated with variation in fungi, Classification of fungi (Alexopoulos and Mims).

UNIT-II **12 Hr**

General characteristics of fungi and reproduction: Morphology and somatic structures: The thallus, organization, fungal cell, nuclear components, specialized somatic structures; Aggregation of hyphae, tissues, mycangia, General aspects of fungal nutrition and reproduction (Asexual, Sexual reproduction, Heterothalism and Parasexuality)

UNIT-III **12 Hr**

Salient features of fungal major groups: Chytridiomycota, Zygomycota, Basidiomycota, Ascomycota, Deuteromycota, Oomycota, Hypochytrio mycota, Labyrinthulomycota, Plasmodiophoromycota and Myxomycota. Symbiotic fungi-Lichens.

UNIT-IV **12 Hr**

Economic importance of fungi: Fungi as biocontrol agent, Economic importance of Fungi in Agriculture, Industry and medicine. Fungi as SCP, Fungi as parasites of human and plants. Role of fungi in bio deterioration of wood and paper. Mycorrhiza – ectomycorrhiza, endomycorrhiza, vesicular arbuscularmycorrhiza. Fungi as insect symbiont.

Reference:

1. Alexopoulos C J and Mims C W, 1979 Introductory Mycology 3rd edn, Wiley Eastern.,NewDelhi.
2. Deacon, J W, 1997- Modern Mycology 3rd Edition, Blackwell Science publishers,London.
3. Mehrotra, RS &Aneja, K R, 1998. An Introduction to Mycology. New Age International Pvt. Ltd. NewDelhi.
4. Odum, E.P. 1971. Fundamentals of Ecology; Third Edition. Toppan Co. Ltd. Tokyo, Japan.
5. Mercedes S. Foster& Gerald F. Bills (2011) Biodiversity of Fungi: Inventory and Monitoring Methods. AcademicPress
6. Michael John Carlile, Sarah C. Watkinson, G. W. Gooday (2007) The fungi. Academic Press. London, U. K
7. Kevin Kavanagh (2011) Fungi: Biology and Applications. John Wiley & Sons, Sussex,U.K.
8. David Moore, Geoffrey D. Robson, Anthony P. J. Trinci (2011) 21st Century Guidebook to Fungi. Cambridge UniversityPress.

COURSE-IV: MICROBIAL GENETICS**48 Hr****Course outcome:**

- Can discuss the importance of mutation analysis, can analyze mutations by complementation and recombination tests, and can design a strategy to create gene replacement in bacteria
- Is able to explain how plasmid copy number is regulated, can differentiate between Hfr strains and strains carrying F plasmid, and can construct a genetic map of bacterial genome using conjugation-based method
- Is able to compare and contrast generalized versus specialized transduction, knows how to construct genetic linkage maps using two-factor and three factor cross, is able to discuss the basis of natural competence in bacteria.
- Is able to list the events in the lytic and lysogenic phases of lambda phage life cycle and the regulatory factors and events involved.
- Can list the outcomes of transposition events, can design strategies to mutagenize bacteria using transposons, can explain the construction of conditional knockouts
- Can differentiate between positive and negative regulation of gene expression, inducible and repressible systems. Can describe the regulation of the lac, trp, gal,ara and tol operons.
- Will have learnt about the model organisms used in biological studies.

Pedagogy:

Microbial genetics deals with the transmission of hereditary characters in microorganisms like bacteria, viruses and algae which play a unique role in developing field of molecular and cell biology and plays wide role in applications in the field of medicine, agriculture, food and pharmaceutical industry. The benefits of microbial

genetics in the field of agriculture are increased in crop yields which reduce the cost for food or drug production, reduce need for pesticide and medical benefits to the worlds growing population by recombinant DNA technology and as vectors.

The importance of genetics study involves; to understand the gene function of microorganisms. Microbes provide relatively simple system for studying genetic phenomenon and thus useful to other higher organisms.

Microbes provide many value added products like antibiotics, growth hormones etc. Microbial genetics will be helpful to increase these products productivity by microbial technology. Understanding the genetics of disease-causing microorganisms especially virus, bacteria will help to develop strategies to control disease. Microbial genetics will be useful to study the gene transfer from one organism to another and its contribution for genetic variations such as virulence, antibiotic resistance etc.

Course Content

UNIT-I

12 Hr

Concepts in Microbial Genetics: History and developments of Microbial genetics. Essentials of microbial genetics: Microbes as Genetic Tools for Basic and Applied Genetic studies. Advantages and disadvantages of Microbes, Generalized reproductive cycles of microbes- *Neurospora*, *Saccharomyces*, *Chlamydomonas* and *Acetabularia*.

UNIT-II

12 Hr

Viral Genetics: Lytic and Lysogenic cycles, Phage Phenotypes, Phenotypic Mixing, Recombination in viruses: Mutations, Recombination and Mapping.

Bacterial Genetics: Bacterial Transformation: Types of transformation mechanisms found in prokaryotes, Bacterial Conjugation: properties of the F plasmid, $F^+ \times F^-$ mating, $F' \times F^-$ conjugation, Hfr conjugation. Transduction: Generalized and specialized transduction, Transposable elements.

UNIT-III

12 Hr

Fungal Genetics: *Neurospora*- Tetrad analysis and linkage detection - 2 point and 3 point crosses, chromatid and chiasma interference, Mitotic recombination in *Neurospora* and *Aspergillus*.

Algal Genetics: *Chlamydomonas*- unordered tetrad analysis - Recombination and Mapping, Nucleocytoplasmic interactions and gene expression in *Acetabularia*. Extra nuclear (Cytoplasmic) inheritance.

UNIT-IV

12 Hr

Mutation and mutagenesis: Nature, type and effects of mutations. Mutagenesis – physical and chemical mutagens, base and nucleoside analog, alkylating agents, interrelating agents, ionizing radiation. Induction and detection of mutation in microorganisms. Site directed mutagenesis and its applications.

References:

1. Larry Snyder, Joseph E. Peters, Tina M. Henkin, Wendy Champness (2013) Molecular Genetics of Bacteria, 4th Edition; ASM Press
2. D. Peter Snustad, Michael J. Simmons (2011) Principles of Genetics, 6th Edition; Wiley
3. Stanley R. Maloy, Jhon E. Cronan, Jr. David Freifelder (1994) Microbial Genetics (Jones and Bartlett Series in Biology), 2nd edition; Jones and Bartlett Publishers
4. Uldis N. Streips, Ronald E. Yasbin (2002) Modern Microbial Genetics, 2nd edition; Wiley-Liss
5. Nancy Jo Trun, J. E. Trempy (2003) Fundamental Bacterial Genetics; Wiley-Blackwell
6. John R. S. Fincham (1996) Microbial and Molecular Genetics; Hodder Arnold
7. Venetia A. Saunders (1987) Microbial genetics applied to biotechnology : principles and techniques of gene transfer and manipulation; Springer
8. Sriram Sridhar (2005) Genetics and Microbial Biotechnology; Dominant Publishers & Distributors
9. Dr. Evelyn J. Biluk (2012) Microbiology Study Guide: Microbial Genetics, Controlling Microbial Growth, and Antimicrobial Agents; CreateSpace Independent Publishing Platform
10. Royston C. Clowes, William Hayes (1968) Experiments in Microbial Genetics; Blackwell Science Ltd
11. Jocelyn E. Krebs, Elliott S. Goldstein, Stephen T. Kilpatrick (2012) Lewin's GENES XI, 11 edition; Jones & Bartlett Learning
12. James D. Watson, Tania A. Baker, Stephen P. Bell, Alexander Gann, Michael Levine, Richard Losick (2013) Molecular Biology of the Gene, 7 edition; Benjamin Cummings

SOFTCORE

COURSE-V: MICROBIAL ECOLOGY AND DIVERSITY

48 Hr

Course Outcome

- To understand the ubiquitous nature of microbes.
- To provide knowledge on characteristics of Microbes Outcome
- Students able to differentiate various groups of Microbes
- Get knowledge on adaptability of extremophiles
- Knowledge about microbial taxonomy.
- To create awareness on evolutionary relationship of ecosystem
- To learn about individual ecosystem and its interactions.
- To understand the concepts of community ecology Outcome
- Able to understand the role of microbes in ecology

Pedagogy

Microbial ecology and diversity is a sub discipline of microbiology (environmental Microbiology) which focuses on the huge diversity of microbes, its interaction among themselves and the ecosystem. Microbes in their natural environments, microbes interact with each other, with plants and animals. Such interactions are essential for ecosystem function and may relate to plant and animal health, biogeochemical cycles and numerous other processes.

Overall this course enables students to learn how the microbial world rules over the entire ecosystem focusing on their interactions which form the basis of survival. The study helps us improve our lives via the use of microbes in environmental restoration, food production, bio-engineering of useful products such as antibiotics, food supplements and chemicals. This course is for all biology, allied health, environmentalists and microbiology students.

The knowledge gained under this subject helps the students to work in laboratories like pharmacological industries, clinical health and diagnostic laboratories, environmental research fields, microbial research and any industry where microorganisms are involved. The need of the hour is to focus on the importance of conservation of microbial diversity mainly the role of culture centers in conservation.

Course Content

UNIT-I

12 Hr

Introduction to microbial ecology: Structure of microbial communities. Interaction among microbial populations. Interaction between microorganisms and plants. Biotransformation, biodegradation, bioremediation and phytoremediation. Ecological and Evolutionary diversity (Genetic diversity) of microbial world
Development of Microbial communities: Dynamics of community, ecological succession, structure, dispersion, microbial communities in nature and ecosystem models

UNIT-II

12 Hr

Physiological Ecology of microorganisms: Adaptation to environmental conditions - abiotic limitations to microbial growth.
Viral Diversity: Group I – T2 Bacteriophage, Group II – Banana bunchy top virus, Group III – Reovirus, Group IV- TMV, Group V – Rhabdovirus, Group VI – HIV, Group VII – Hepatitis virus.
Sub-viral particles: Discovery, Structure, Classification, replication and diseases caused by Satellite, Satellites virus, Virusoids, Viroids and Prions.

UNIT-III

12 Hr

Bacterial Diversity: Archaeobacteria, Photosynthetic Eubacteria, Chemoautotrophic and Methophilic Eubacteria, Gliding Eubacteria, Spirochetes, Rickettsiae and Chlamydiae, Actinomycetes, Mollicutes, Protists
Fungal Diversity: salient features of the following group: Zygomycota (*Rhizopus*), Ascomycota (*Neurospora*), Basidiomycota (*Agaricus*), Deuteromycota (*Penicillium*), Chytridiomycota (*Allomyces*) Myxomycota and Yeast.

UNIT-IV

12 Hr

Importance and Conservation of Microbial Diversity: Importance of microbial diversity in environment, pharmaceuticals & human health. Metagenomics. Importance of conservation. *In situ* conservation and *Ex situ* conservation. Role of culture collection centers in conservation.

References

1. OladeleOgunseitan (2008) Microbial Diversity: Form and Function in Prokaryotes; Wiley-Blackwell
2. Ronald M. Atlas, Richard Bartha (1997) Microbial Ecology: Fundamentals and Applications (4th Edition); Benjamin Cummings
3. David L. Kirchman (2012) Processes in Microbial Ecology; Oxford University Press
4. David L. Kirchman (2008) Microbial Ecology of the Oceans; Wiley-Liss
5. McArthur, J. Vaun (2006) Microbial Ecology An Evolutionary Approach; Academic Press
6. Atlas, Ronald M., Bartha, Richard (1997) Microbial Ecology Fundamentals and Applications; Addison-Wesley
7. Nelson, Karen E. (1997) Advances in Microbial Ecology; Springer
8. Pierre Davet (2004) Microbial Ecology of the Soil and Plant Growth; Science Publ Inc
9. Osborn, A. M., Smith, Cindy (2005) Molecular Microbial Ecology; Taylor & Francis Group
10. OladeleOgunseitan (2004) Microbial Diversity: Form and Function in Prokaryotes; Wiley-Blackwell
11. Satyanarayana, T., Johri, B. N. (2005) Microbial Diversity: Current Perspectives and Potential Applications; I.K. International Publishing House Pvt., Limited
12. James W. Brown (2014) Principles of Microbial Diversity; ASM Press
13. Colwell, R. R., Simidu, Usio, Ohwada, Kouicki (1996) Microbial Diversity in Time and Space; Springer

Practical's I (Virology and Bacteriology)

1. Laboratory safety rules
2. Microscopic measurement of microorganisms by micrometry
3. Culturing and maintenance of bacterial cultures
4. Isolation and enumeration of bacteria from soil
5. Isolation and enumeration of bacteria from water
6. Cultural characteristics of bacteria
7. Staining techniques – simple (positive and negative), differential (Grams and acid fast), structural (endospore and capsule)
8. Motility test (hanging drop method and soft agar method)
9. Biochemical tests for the identification of bacteria – catalase, oxidase, IMViC, Urease, TSIA, Nitrate reduction, gelatin, starch, casein, chitin and esculin hydrolysis.
10. Determination of growth curve in E.coli.
11. Diauxic growth curve in E.coli
12. Isolation of coliphages from sewage
13. Study of morphological changes due to viral infection in plants

Practicals II (Mycology and Microbial Genetics)

24

1. Isolation of slime molds.
2. Isolation of aquatic fungi.
3. Isolation of soil fungi.
4. Isolation of fungi from air.
5. Isolation of fungi from cereals and cereal based products.
6. Study of the following representative genera: *Aspergillus*, *Penicillium*, *Fusarium*, *Neurospora*, *Saccharomyces*, *Erysiphae*, *Polyporus*, *Agaricus*, *Puccinia*, *Ustilago*, *Alternaria*, *Drechslera*, *Saprolegnia*, *Rhizopus*, *Trichoderma* and symbiotic fungi-Lichens.
7. Measurement of concentration of fungal conidia by Haemocytometer.
8. Measurement of fungal cells by Micrometer.
9. Replica plating technique for transfer of bacterial colonies.
10. Ultra-violet killing curve and determination of mutant types in *Saccharomyces cerevisiae*.
11. Induction of mutation
12. Isolation of streptomycin resistant strain of *E. coli* by gradient plate method.
13. Ames test
14. Isolation of genomic DNA from bacteria by heat lysis method.
15. Isolation of genomic DNA from yeast by DNA spooning method.

SECOND SEMESTER

HARDCORE

COURSE-I: MICROBIAL PHYSIOLOGY

48 Hr

Course Outcome:

- Learn the methods of measuring microbial growth, calculating growth kinetic parameters with understanding of steady state and continuous growth.
- Knowledge of primary, secondary and group translocation transport systems existing in bacteria, simultaneously learning membrane transport proteins and kinetics of solute transport.
- Understand the central metabolic pathways for carbon metabolism in bacteria and their regulation in diverse physiological conditions. This allows students to apply the acquired knowledge in engineering metabolic pathways for developing industrially useful strains as well as lipid and nucleotide metabolism in bacteria.
- Learn and understand the basic concepts of enzyme biochemistry, kinetics and regulation.
- Knowledge about stress response in bacteria

Pedagogy:

Microbial physiology is defined as the study of microbial cell functions which includes the study of microbial growth, microbial metabolism and microbial cell structures. The contents of the course are divided into four main chapters or units those are: A) Microbial physiology, B) Carbohydrate metabolism, C) lipid metabolism, D) Microbial photosynthesis and each unit focuses on various aspects of microbial physiology.

Microbial physiology is an important research field, not only in fundamental research on microbial species but also in all applied aspects of microbiology including physiology of the aerobic microorganisms and anaerobic microbial communities that play an important role in environmental biotechnological processes, such as waste water treatment, soil remediation, production of chemicals, biofuels and recovery of metals.

Course Content

UNIT-I

12 Hr

Microbial Physiology: Microbial Energetics, The role of ATP in metabolism. Microbial enzymes: Structure and Classification, Mechanism of Enzyme actions: Lock and Key model, induced fit Theory, Factors affecting rates of enzyme mediated reactions (pH, temperature and substrate and enzyme concentration), Enzyme Inhibition and Enzyme regulation.

UNIT-II

12 Hr

Metabolism of Carbohydrate: Glycolysis, Citric acid Cycle and different types of Phosphorylation, Fates of pyruvate, Fermentation. Utilization of sugars other than glucose: Lactose, Galactose, Maltose, Mannitol. Degradation of cellulose, Starch and Glycogen.

UNIT-III

12 Hr

Metabolism of other Substrates: Lipid metabolism: β -oxidation, Biosynthesis of fatty acids, degradation of fatty acids. **Nitrogen metabolism:** Nitrogen metabolism, Biological nitrogen fixation process, symbiotic and non-symbiotic nitrogen fixation. Urea cycle, degradation and biosynthesis of essential and non-essential amino acids. **Nucleic acid metabolism:** Biosynthesis and degradation of purines and pyrimidines.

UNIT-IV

12 Hr

Microbial Photosynthesis: Photosynthetic Pigments and apparatus in bacteria. Oxygenic and Anoxygenic. Photosynthesis. Autotrophic CO₂ fixation and mechanism of Photosynthesis. Utilization of light energy by Halobacteria.

Autotrophic Mechanisms in bacteria: Hydrogen bacteria, Nitrifying bacteria, Purple sulphur bacteria, Non-sulfur bacteria, Green sulfur bacteria, Iron bacteria, Methylophs. **Microbial Stress Responses:** Oxidative stress, Thermal stress, Starvation stress, Aerobic to anaerobic transitions. Biofilm and quorum sensing

References:

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8. Robert K. Poole (2014) Advances in Microbial Systems Biology, Volume 64 (Advances in Microbial Physiology); Academic Press
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10. Rose, Anthony H. (1976) Chemical Microbiology An Introduction to Microbial Physiology; Basic Books

Course-II: Immunology

48 Hr

Course Outcome:

- Will be able to understand the fundamental bases of immune system and immune response
- Will be able to gather information about the structure and organization of various components of the immune system
- Will be able to understand the genetic organization of the genes meant for expression of immune cell receptors and the bases of the generation of their diversity
- Will be able to understand the operation and the mechanisms which underlie the immune response
- Will be able to apply the knowledge gained to understand the phenomena like host defense, hypersensitivity (allergy), organ transplantation and certain immunological diseases

Pedagogy:

Immunology is the branch of biology which deals with various aspects that forms an integrated network of cells, molecules, and organs within the immune system. This course helps students to learn and understand basic concepts as well as its application in various fields of biology.

The content of the course consist of four units where each unit focuses on basic aspects of immunology and its application. The course begins with the brief introduction regarding overview of immune system followed by the mechanism of immunological reactions, immunotechniques, immunodiagnosis and its application in the field of medicine. At the end of each unit a student is able to understand the development of immune system immune response against diseases, and autoimmunity.

Studying this subject will equip students with basic practical skills to work in vast fields like pathology, pharmacy industries, diagnostics and hospitals.

Course Content

UNIT-I

12 Hr

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Introduction to Immunology: An overview of immune system, Phagocytes, Natural killer cells, mast cells, basophils, Dendritic cells and other cells of the innate immune system. Immunity: Types- Innate immunity: (nonspecific) physical, biochemical and genetic factors involved in governing innate immunity, molecules of innate immunity – complement, acute phase proteins and interferons; Chemokines and Cytokines . Acquired immunity: (specific) natural, artificial, passive immunity, humoral or antibody mediated immunity, cell mediatedimmunity.

Antigens and Antibodies: Antigen processing and presentation, properties of antigen, Super antigen, Hapten; Haptens and the study of antigenicity Microbes as antigen Antigen recognition and MHC molecules. Antibodies (Immunooglobulins) – structure and function, clonal selection, monoclonal antibodies and its clinical applications, Antibody engineering (Construction of monoclonal antibodies Lymphoma and other diseases by genetically engineeredantibodies).

UNIT-II

12 Hr

Hypersensitivity: Hypersensitivity reactions, Types and their roles in Immunopathologicalprocesses.

Autoimmune processes: Immunologic tolerance, genetic predisposition to the development of autoimmune processes. Autoimmune disorders- Immunopathogenesis of celiac disease, myasthenia gravis, sclerosis multiplex, psoriasis vulgaris, Rheumatoid arthritis) Immunodeficiency diseases, Hormones and environmental factors in induction of autoimmune processes.

UNIT-III

12 Hr

Transplantation of tissues and organs: Nomenclature of transplantations. Recognition of self and non- self-Transplantation reactions HvG and GvH. Exception from rejections.Kidney and bone marrow transplantations.

Tumours and immune system: Etiology of malignant transformations of cells (physical, chemical and biological factors involved in). Immunological surveillance. Escape mechanisms of tumor cells from immunological surveillance. Metastatic processes. Immunodiagnosis and Immunotherapy.

UNIT-IV

12 Hr

Vaccines and Vaccination: Vaccines – definition, types, Antigens used as Vaccines, effectiveness of vaccines, Vaccine safety, current vaccines, adjuvants, active immunization and passive immunization .**Manipulation of immune mechanisms:** Immunoprevention, Immunoprophylaxis, Immunostimulatory and Immunosuppressivedrugs.

Immunotechniques and Immunodiagnosis: Antigens and Antibody reactions *in vitro*; Agglutination, complement fixation, ELISA, Immunodiffusion,

Immuno-electrophoresis, Immunofluorescence, Immunoprecipitation,
Radioimmunoassay and serotyping.

References:

1. Abul K. Abbas (2014) Cellular and Molecular Immunology, ;Saunders
2. Judy Owen , Jenni Punt, Sharon Stranford (2013) Kuby Immunology; W. H.Freeman
3. Abul K. Abbas , Andrew H. H. Lichtman , Shiv Pillai (2012) Basic Immunology: Functions and Disorders of the Immune System, ;Saunders
4. Peter Parham (2009) The Immune System, 3rd Edition; Garland Science
5. Abul K. Abbas , Andrew H. H. Lichtman , Shiv Pillai (2011) Cellular and Molecular Immunology; Saunders
6. George Pinchuk (2001) Schaum's Outline of Immunology; McGraw-Hill
7. William E. Paul (2012) Fundamental Immunology; LWW
8. Helen Chapel , Mansel Haeney, Siraj Misbah, Neil Snowden (2014) Essentials of Clinical Immunology; Wiley-Blackwell
9. Louise Hawley, Benjamin Clarke, Richard J. Ziegler (2013) Microbiology and Immunology; LWW
10. Delves, Peter J., Martin, Seamus J., Burton, Dennis R. (2011) Roitt's Essential Immunology; Wiley & Sons, Incorporated, John.

SOFTCORE

COURSE –III: FOOD MICROBIOLOGY

48Hr

Course Outcome:

- Learn about production and evaluation of the quality of starter cultures and fermented milk products and understands the use and production of probiotics, prebiotics and nutraceuticals.
- Knowledge about fermentation protocols for production of microbial biomass such as edible yeasts, mushrooms, single cell proteins and single cell oils. The student also learns about production of microbial carotenoid pigments such as lycopene and β -carotene.
- To know microbes causing food intoxications and food-borne infections.
- Know traditional food preservation techniques, chemical preservation and irradiation.
- Is able to utilize modern techniques viz. high-pressure processing (HPP), bacteriocins, manosonication (MS) and pulsed electric field (PEF) for effective food preservation. The student can also calculate kinetics of inactivation, process and product parameters.
- Gain knowledge about conventional methods for food quality analysis and is able to use the most recent and non-invasive techniques of quantification and detection of food borne microbes and pathogens such as ESS and various new imaging techniques.
- Understands the relevance of microbial standards for food safety, quality assurance programs that revolutionize food safety.

Pedagogy:

Food microbiology is a sub-discipline of Microbiology which focuses on the study of the microorganisms that ferment, inhibit or contaminate food. It also includes the study of microorganisms that cause food spoilage and those with other useful roles.

The course emphasizes basic concepts of food microbiology, contamination and food spoilage, dairy microbiology, food poisoning and intoxication, food produced by microbes, detection of food borne microorganisms and microbial indicators of food safety quality control, food law and legislation.

The study encompasses food borne diseases of microbial origin, microbial food spoilage, beneficial uses of microbes in food, control of microbial growth in foods, destruction of microbes in foods, microbial food fermentation, pro-biotic bacteria, regulatory aspects to ensure consumers related to microbial hazards in food.

The lecture will impart students with knowledge, how microorganisms are useful to produce food, how they contaminate, spoil and cause diseases and how to detect their presence in the food. The knowledge gained about food microbiology helps the students to develop interest in this field and helps the students to work in the food industries that are interested in isolation, detection of food borne pathogens and production of food products from microorganisms.

Course Content

UNIT-I

12 Hr

Introduction to food microbiology: Definition, concepts and scope. Food as substrate for microbes. Factors influencing microbial growth in food-Extrinsic and intrinsic factors. Principles of food preservation- Chemical preservatives and Food additives Asepsis-Removal of microorganisms, (anaerobic conditions, high temperatures, low temperatures, drying).Canning, processing for Heat treatment.

UNIT-II

Contamination and food spoilage: Cereals, sugar products, vegetables, fruits, meat and meat products, Fish and sea foods- poultry- spoilage of cannedfoods.

Dairy Microbiology: Microbiology of raw milk, Milk as a vehicle of pathogens, Prevention of contamination of raw milk, Microbiology of processed milk, Spoilage and defects fermented milk and milk products, Microbiological standards for milk and milk products. Cream and butterbacteriology.

UNIT-III

12 Hr

Food poisoning and intoxication: Significance of food borne diseases, Staphylo Food poisoning and intoxication: Significance of food borne diseases, Staphylococcal, Gastroenteritis and enterotoxins: Types and incidence, Prevention of Staphylococcal and other food poisoning syndromes, *Clostridium perfringens* food poisoning and Botulism, *Bacillus cereus* food poisoning, Food borne Listeriosis by *Listeria monocytogenes*, Food borne Gastroenteritis by *Salmonella* and *Shigella*, *Vibrio*, *Campylobacter* and *Yersinia*, fungal spoilage and Mycotoxins.

Food produced by Microbes: Microbial cells as food (single cell proteins) – mushroom cultivation. Bioconversions- production of alcohol-fermented beverages- beer and wine. Genetically modified foods.

UNIT-IV

12 Hr

Detection of food-borne microorganisms: Culture, Microscopic and Sampling methods.. Chemical: Thermostable nuclease *Limulus* Lysate for Endotoxins, Nucleic Acid (DNA) probes, DNA Amplification (PCR), Adenosine- Triphosphate Measurement, Radiometry, Fluoro-and Chromogenic substrates. Immunologic Methods: Fluorescent Antibody, Enrichment Serology, Salmonella 1-2. Test, Radioimmunoassay, ELISA.

Microbial indicators of food safety and quality control: Principles of quality control and microbiological criteria, Indicators of product quality and microbiological safety of foods, Hazard analysis, critical control points (HACCP), Good manufacturing process (GMP) Microbiological standards Codex Alimentarius and Foodlegislation.

References:

1. Adams M. R. and Moss M. O. 2007. Food Microbiology 3rd Edition. Royal Society of Chemistry.UK.
2. Ahmed E.Y. and Carlstrom C. 2003 Food Microbiology: A Laboratory Manual, John Wiley and Sons, Inc. NewJersey.
3. Bibek Ray, ArunBhunia. 2013. Fundamental Food Microbiology, Fifth Edition. CRCPress
4. C Blackburn.2006. Food Spoilage Microorganisms. WoodheadPublishing.
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7. Frazier W.C. and Westhoff C.D. 2008 Food Microbiology. Tata McGraw Hill Publishing Company Limited, New Delhi. IndianEdition.
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10. Pina M. Fratamico, Arun K. Bhunia, and James L. Smith. 2008. Foodborne Pathogens: Microbiology and Molecular Biology. Caister AcademicPress.
11. Pitt, John I., Hocking, Ailsa D. 2009. Fungi and Food Spoilage 3rd Edition.Springer.
12. Sperber, William H., Doyle, Michael P. (Eds.). 2010. Compendium of the Microbiological Spoilage of Foods and Beverages.Springer.
13. Stephen J. Forsythe. 2010. The Microbiology of Safe Food, 2nd Edition.Wiley-Blackwell.

Course-IV: SOIL MICROBIOLOGY

48 Hr

Course Outcome:

- Students will learn that the soil is an excellent habitat for multitude of microorganisms balancing the soil ecosystem.

- The knowledge acquired in Soil Microbiology will enhance the students' competency in the performance of their duties as future employees in the field of Agronomy/Soil Science.
- Attainment of course objectives will mean realization of the various beneficial effects of soil microorganisms on soil health, which is instrumental in the production of food and fiber. Conversely, students learned that some soil microbes are deleterious to agronomic crops
- Students will learn that some soil animals and what they eat are of ecological importance; thus, plant-eating insects and mollusks may add organic matter to the soil; insects, arachnids, and worms that consume dung and plant litter mix it with soil and speed up its decay; and, plant parasitic nematodes reduce soil's productivity.

Pedagogy:

Soil microbiology is the study of all microorganisms that exist in the soil, specifically the ways they function and affect soil properties. Our soils are pulsating with life, serving as excellent hosts for the growth and development of various organisms. In fact, there are more microbes in one teaspoon of soil than there are people on the planet. This collection of organisms consists of bacteria, fungi, and algae that serve many vital roles in the overall nourishment of soils.

Bacteria are largely responsible for the process of nitrogen fixation; converting atmospheric nitrogen into compounds that can be used by plants. Although not as commonly abundant as bacteria, fungi also assist with extremely significant functions of soil health. While one of their main activities is decomposition of organic matter, fungi also perform necessary services related to water and nutrient cycling. Fungi are responsible for binding soil particles together, assembling a system to increase water filtration and water holding capacities. In a similar manner as fungi, earthworms also break down organic matter, such as dead leaves, and produce natural fertilizers. They too support soil fertility with the transportation of water throughout the soil, as well as air.

Course content

UNIT-I

Soil Microbiology: Historical accounts and the "Golden Age" of soil microbiology and significant contributions of pioneer soil microbiologists. **Soil Microbial diversity:** Diversity and abundance of dominant soil microorganisms, Methods of isolation of soil microflora, soil organic matter decomposition.

Unit-II

12 Hr

Biogeochemical cycles: carbon, sulphur and iron cycles in soil.

Soil microbe interaction - Antagonism, commensalism, mutualism, symbiosis, predators and parasite relationship and competition. Interaction of soil microflora with vascular plants - Rhizosphere, rhizoplane microorganisms, *Rhizobium*, *Azotobacter*, *Azospirillum*, *Cyanobacteria* and *Azolla*.

Unit-III

12 Hr

Applied soil microbiology: soil microbial inoculants, Manipulations of soil microorganisms for agriculture, Soil environmental contaminants and Bioremediation, Microbial products- Plant growth promoting Hormones, Antibiotics, Toxins and Enzymes

Unit-IV

12 Hr

Soil-Borne Diseases and Human Health: *Clostridium tetani*(tetanus), Toxoplasmosis, Aspergillosis, Actinomyces.

Soil microorganisms in agro ecosystems: Types of microbial communities; soil microbial

diversity: significance and conservation; effect of agricultural practices on soil organisms. Biological nitrogen-fixation: The range of nitrogen fixing organisms; mechanism of nitrogen fixation (biochemistry of nitrogenase); genetics of nitrogen-fixation; *Rhizobium*-Legume Association; Symplasmids, N₂ fixation by non-leguminous plants.

References:

1. Agrios, G. N. 2000. Plant pathology. Harcourt Asia Pvt.Ltd.
2. Bergersen, F.J. and Postgate, J.R. 1987. A Century of Nitrogen Fixation Research Present Status and Future Prospects. The Royal Soc.,London.
3. Buchanan, B.B., Gruissem, W. and Jones, R.L. 2000. Biochemistry and Molecular Biology of Plants. I.K. International Pvt. Ltd.
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5. Dixon, R.O.D. and Wheeler, C.T. 1986. Nitrogen Fixation in plants. Blackie USA, Chapman and Hall, NewYork.
6. Kannaiyan, S. 1999. Bioresources Technology for sustainable agriculture. Assoc. Pub. Co. New Delhi.
7. Mehrotra, R.S. 2000. Plant pathology. Tata McGraw-Hill Publishing Company Limited.
8. Metcalf, R.L. and Luckmann, W.H. 1994. Introduction to insect pest management 3ed edn. John Willey and Sons,Inc.
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10. Somasegaran, PandH.J.Hoben, 1994. Handbook for Rhizobia; methods in legume eRhizobium Technology. Springer-Verlan, New York.

PRACTICAL-III (Microbial Physiology and Immunology)

1. Population growth of yeast – *S.cerevisiae*.
2. Population growth of bacteria – *Ecoli*.
3. Sugar fermentation tests.
4. Catalase activity.
5. Hydrolytic rancidity.
6. Casein hydrolysis.
6. Carbohydrate catabolism by microbes
7. Study of acid and pH stress tolerance by microbes.

8. Effect of molecular oxygen on microbial growth.
9. Effect of osmotic pressure on microbial growth.
10. Effect of relative humidity on microbial growth.
11. Effect of different wavelengths of light on microbial growth.
12. Immunological Methods used for organism detection – production of antibodies for use in laboratory testing.
13. Serological Diagnosis of Infectious diseases – Serologic test Methods.
14. Precipitin test, ELISA, Ouchterlony Immunodiffusion test, Immunoelectrophoresis, Complement fixation test.
15. Isolation of Antigens and raising antibodies from animals (from different Models),
16. Development of polyclonal antibodies, purification of antibodies.
17. WIDAL Test.
18. VDRL Test (RPR).
19. HBs Ag Test.
20. HCG test (Agglutination inhibition test).
21. Detection of RA factor.
22. CRP test.
23. ASO Test (Anti streptolysin 'O' Test).

PRACTICAL-IV (FOOD AND DAIRY MICROBIOLOGY)

1. Bacterial examination of drinking water by membrane filter technique.
2. Study of important microbes in the degradation of wastes.
3. Determination of TDT.
4. Determination of TDP.
5. Detection and quantification of Aflatoxin B1.
6. Detection of food-borne bacteria by immunoassays.
7. Detection and enumeration of Microorganisms present in Utensils.
8. Isolation and identification of pathogenic microorganisms from canned food.
9. Enumeration of bacteria in raw and pasteurized milk by SPC method.
10. Determination of quality of a milk sample by MBRT.
11. Detection of number of bacteria in milk by breed-count method
12. Litmus milk test.
13. Microbial quality of milk products.
14. Microbiological examination of Ice-cream and Dairy products
15. Soil microbes interaction *In vitro* by dual culture method
16. Isolation, identification and enumeration of Rhizosphere and Rhizoplane microorganism
17. Isolation of *Rhizobium* from roots of leguminous plant.

OPEN ELECTIVE

COURSE –VII: MICROBIAL DIVERSITY

48 Hr

Course Outcome:

- To gain insights into common groups of bacteria and archaea in different ecosystems, and their role in biogeochemical key processes in these environments.

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- To understand cultivation-independent methods for studies of the composition of microbial communities and for the function and occurrence of individual groups.
- Genomic and meta genomic based methods to study microbial diversity in nature
- To understand interactions within microbial communities and between microorganisms and plants and animals.
- Use bioinformatic tools and databases that are used to study microbial diversity.
- Acquire a fairly good understanding of the Diversity of the microbes and activities/importance of microbes.; practical skills of handling microorganisms in the laboratory for study

Pedagogy:

Microbial diversity is a sub discipline of microbiology focuses on the huge diversity of microbes, its interaction with the ecosystem. Such interactions are essential for ecosystem function and may relate to plant and animal health, biogeochemical cycles and numerous other processes.

Overall this course enables students to learn how the microbial world rules over the entire ecosystem focusing on their interactions which form the basis of survival. The study helps us improve our lives via the use of microbes in environmental restoration, food production, bio-engineering of useful products such as antibiotics, food supplements and chemicals.

The knowledge gained under this subject helps the students to work in laboratories like pharmacological industries, clinical health and diagnostic laboratories, environmental research fields, microbial research and any industry where microorganisms and involved. The need of the hour is to focus on the importance of conservation of microbial diversity mainly the role of culture centers in conservation.

COURSE CONTENT

UNIT-I

12 Hr

Viral Diversity: Morphology, ultra structure, chemical composition of virus, classification of viruses, Group I – T2 Bacteriophage, Group II – Banana bunchy top virus, Group III – Reovirus, Group IV- TMV, Group V – Rhabdovirus, Group VI – HIV, Group VII – Hepatitis virus.

Sub-viral particles: Discovery, Structure, Classification, replication and diseases caused by Satellite, Satellites virus, Virusoids, Viroids and Prions.

UNIT-II

12 Hr

Bacterial Diversity: Archaeobacteria, Photosynthetic Eubacteria, Chemoautotrophic and Methophilic Eubacteria, Gliding Eubacteria, Spirochetes, Rickettsiae and Chlamydiae, Actinomycetes, Mollicutes, Protists. Classification based on Bergey's manual (Determinative & Systematic).

UNIT-III

12 Hr

Fungal Diversity: Classification, Distribution, Importance, Structure, reproduction and general characteristics of the fungal divisions: Zygomycota (*Rhizopus*), Ascomycota

(*Neurospora*), Basidiomycota (*Agaricus*), Deuteromycota (*Penicillium*), Chytridiomycota (*Allomyces*), Myxomycota and Yeast.

UNIT-IV

12 Hr

Importance and Conservation of Microbial Diversity: Importance of microbial diversity in agriculture, forestry, environment, industrial & food biotechnology, animal & human health. Metagenomics. Importance of conservation. *In situ* conservation and *Ex situ* conservation. Role of culture collection centers in conservation.

References

1. Alexopoulos, C. J. and Mims, C. W. 1979. Introductory Mycology. III edition, Wiley Eastern, New Delhi.
2. Dimmock, N. J., Easton, A. J. and Leppard, K. N. 2001. Introduction to Modern Virology. 5th edn. Blackwell publishing, USA.
3. Ghosh, A. 2003. Natural Resource Conservation and Environment Management. Aph Publishing Corp. Calcutta.
4. Landecker, E. M. 1972. Fundamentals of Fungi. Prentice-Hall, Angelwood Cliff, New Jersey.
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6. Pelczar, (Jr.) M. J., Chan, E. C. S. and Kreig, N. R. 1993. Microbiology. McGraw Hill, New York
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10. Stainer, R. Y., Ingraha, J. L., Wheelis, M. L. and Painter, P. K. 1986. General Microbiology. McMillan Edun. Ltd. London.
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12. Wagner, E.K. and Hewlett, M.J. 1999. Basic Virology. Blackwell Science. Inc.

THIRD SEMESTER

HARDCORE

COURSE-1: MOLECULAR BIOLOGY

48Hr

Course Outcome:

- Understand the structure of DNA and RNA, organization of eukaryotic/prokaryotic genome, replication, , transcription, Mutations and DNA repair
- Insights into gene regulation in both prokaryotic and eukaryotic organisms
- Is able to describe post-transcriptional processes, RNA editing, RNAi and miRNA
- Is able to describe translation mechanism in prokaryotes and eukaryotes, regulation of translation, and post-translational processing
- Is able to describe post-translational processes.

Pedagogy:

Molecular biology is the root branch of biology which deals with biomolecules, its modifications and other molecular level mechanisms occurring in the body of living organisms. This field is developed out of related fields like genetics, biochemistry, biophysics and microbiology. Molecular biology gives a wide information on basic concepts of DNA structure and replication, DNA damage and recombination, synthesis of proteins by transcription, translation and regulation of gene expression in bacteria, bacteriophage, eukaryotes. Each unit is well presented with basic descriptions of cellular mechanisms of both prokaryotes and eukaryotes. In this discipline the major interest is drawn towards the differences in the molecular mechanisms in prokaryotes and eukaryotes.

The students view is synchronized into the world of biomolecules for the better understanding of molecular mechanism, cell to cell interaction, cell replication, mutations. This discipline allows the students to understand the molecular mechanisms so that they can study the cause of evolutionary existence of life and also the various diseases that result due to the changes in the biomolecules.

COURSE CONTENT**UNIT-I****12Hr**

Concepts in Molecular Biology: Microbes in molecular biology.

Organization of Genomes: Prokaryotic genome- Genetic and Physical organization of bacterial genome, Eukaryotic genome – Genetic and Physical organization of nucleargenome **DNA structure and Replication:** DNA as Genetic material, Chemistry of DNA, Modes of DNA Replication, Enzymes of DNA replication, Molecular mechanism of DNA replication, Differences in prokaryotic and eukaryotic DNA replication.

UNIT-II**12Hr**

DNA damage and recombination: Types of DNA damage - deamination, oxidative damage, alkylation and pyrimidine dimers; DNA repair – mismatch, short patch repair, nucleotide/base, excision repair, recombination repair and SOS repair. Molecular basis of mutation, Recombination; Site specific recombination, Homologous recombination, transposition

UNIT-III**12Hr**

Gene Expression: Structure of RNA- Classes of RNA, Chemistry of RNA.

Transcription: Transcription in prokaryotes and eukaryotes, Eukaryotic transcription factors. RNA processing, Ribozymes, Antisense RNA, Inhibitors of transcription and their mechanism of action.

Translation: Role of ribosome and different types on RNA in protein synthesis, basic feature of genetic code, mechanism of initiation, elongation and termination, Translational control and posttranslational events.

UNIT-IV**12Hr**

Regulation of Gene expression: Regulation of gene expression in prokaryotes and Eukaryotes. Regulation of gene expression in bacteriophages, gene silencing – gene regulation after transcription.

References:

1. Benjamin, L. 1990. Gene 4th edn. Oxford Univ. Press, Oxford.
2. Brown, T. A. 1991. Essential Molecular Biology. A Practical Approach Vol-I & Vol.-II, Oxford Univ. Press, Oxford.
3. Flint, S.J., Enquist, L.W., Drug, R.M., Racaniello, V.R. and Skalka, A.M. 2000. Principles of Virology- Molecular Biology, Pathogenesis and Control. ASM Press, Washington, D.C.
4. Garrett and Grisham. 1999. Biochemistry. 2nd edn. Saunders college pub. USA.
5. Hartl, D.L. 1994. Genetics. Jones and Bartler Publishers, London.
6. Lewin, B. 2000. Genes VII. Oxford Univ. Press.
7. Lodish, H., Berk, A., Zipursky, S. A., Matsudaira, P., Baltimore, D. and Darnell, J. 1999.
8. Molecular Cell Biology, W.H. Freeman and Company, New York.

COURSE-II: GENETIC ENGINEERING

48Hr

Course Outcome:

- Students will become familiar with the tools and techniques of genetic engineering DNA manipulation enzymes, genome and transcriptome analysis and manipulation tools, gene expression regulation, production and characterization of recombinant proteins.
- This course exposes students to the applications of genetic engineering in biological research.
- Students will be able to perform basic genetic engineering experiments at the end of course.
- Students will acquire knowledge of advances in biotechnology- healthcare, agriculture and environment cleanup via recombinant DNA technology.
- Given the impact of genetic engineering in modern society, the students should be endowed with strong theoretical knowledge of this technology. In conjunction with the practicals in molecular biology & genetic engineering, the students should be able to take up biological research as well as placement in the relevant biotech industry

Pedagogy:

Genetic Engineering is an inter-disciplinary subject of biology which focuses on gene manipulation techniques using living systems and the applications of manipulated genes. This course helps students learning the components, techniques of gene manipulation in organisms and use of these techniques to create novel products (vaccines, enzymes, GMOs).

The contents of the course are divided into four units. Each unit focuses on tools, techniques used in gene manipulation, applications of recombinant DNA, ethics concerned with gene manipulation and bioinformatics. Overall, this course teaches students the importance and scope of genetic engineering in the current world.

The lecture will impart knowledge of using these techniques in various fields such as agriculture to create transgenic plants, in therapeutics or medicine to create vaccines, to cure genetic diseases. In industries to increase efficiency of production of various microbial products, in forensic science to identify suspects, paternity issues etc., this subject has wide scope and great significance in the world.

COURSE CONTENT

UNIT-I

12Hr

Introduction to Genetic Engineering: Historical perspectives and milestones in Recombinant DNA Technology. Importance of gene cloning and future perspectives.

Tools in Genetic Engineering: Enzymes in genetic engineering. Cloning vectors: Ti Plasmid, pBR322, pUC –series. Phage vectors-M13 phage vectors, Cosmids-Types, Phasmids or Phagemids, Shuttle vectors. YAC and BAC vectors, Adenovirus vector, Synthetic construction of vectors, Ti cloningvector

UNIT-II

12Hr

rDNA Technology: The basic principles of gene cloning strategies: Preparation, Manipulation and Insertion of desired DNA into vector. Introduction of DNA into host cells – Transformation, Transduction, Transfection, Microinjection, Biolistics, Electroporation, Liposome fusion. Shotgun cloning. Genomic and c-DNA Libraries. Cloning and expression in bacteria, yeasts, Identification and Selection of recombinants.

UNIT-III

12Hr

Analysis of gene and gene products: Isolation and purification of nucleic acids, staining, Molecular markers in genome analysis: RFLP, RAPD, AFLP and ISSR analysis, DNA sequencing. Blotting techniques- Southern, Northern and Western blotting techniques. PCR –principles, types, and applications.

Introduction to Bioinformatics and Molecular Databases, Primary Databanks – NCBI, EMBL, DDBJ; Secondary Databases – UNIPROT; Structural Database –PDB; Database similarity search (FASTA, BLAST); Alignment: Pairwise and Multiple sequence alignment; Genome Annotation and Gene Prediction; Primer Designing; Phylogenetics analysis and Tree construction; Protein Sequence Analysis; DNA microarrays. DNA sequencing methodology – Sangers dideoxymethod.

UNIT-IV

12Hr

Applications of gene cloning and Ethics in Genetic Engineering: Applications of gene cloning in Biotechnology, Medicine, Agriculture, Forensic Science, Antisense technology. RNAi and Gene silencing, Gene therapy.

Safety of recombinant DNA technology: Restriction and regulation for the release of GMOs into Environment. Ethical, Legal, Social and Environmental Issues related to rDNA technology.

References:

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4. Harvey Lodish, Arnold Berk, Chris A. Kaiser, Monty Krieger (2007) Molecular Cell Biology 6th Ed. W.H. Freeman and Company, New York.
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10. P. K. Gupta. (2008) Molecular Biology and Genetic Engineering. Deep and Deep Publications. India.
11. VK Gupta, MSchmoll, M Maki, MTuohy, MAMazutti. (2013) Applications of Microbial Engineering. CRC Press.

COURSE-III: INDUSTRIAL MICROBIOLOGY

48Hr

Course Outcome:

- Get equipped with a theoretical and practical understanding of industrial microbiology
 - Appreciate how microbiology is applied in manufacture of industrial products
 - Know about design of bioreactors, factors affecting growth and production, heat transfer, oxygen transfer
 - Understand the rationale in medium formulation & design for microbial fermentation, sterilization of medium and air
 - Appreciate the different types of fermentation processes
 - Understand the biochemistry of various fermentations
 - Identify techniques applicable for improvement of microorganisms based on known biochemical pathways and regulatory mechanisms
- Comprehend the techniques and the underlying principles in downstream processing.

Pedagogy:

Industrial microbiology is a branch of applied microbiology. Which deals with the microorganisms and fermentation technology used for production of high value added products such as therapeutic agents, fuels, food items, chemicals, sweeteners, detergents, beverages, enzymes, vitamins, and proteins. The course imparts detailed fundamental principles and of industrial microbial processes.

The course contained four units and focuses on basic industrial equipment's, isolation and screening of microorganisms, media formulation, production and key factors for optimum maintenance, recovery process and production economics, commercial value

and their applications. The course provides the basic knowledge of the industrial processes and biosynthesis of potent microbial agents.

This course makes the students as entrepreneurs and gives so many jobs for the people, and helps the students to work in the pharmaceutical, chemical, food ,beverages and dairy industries and biotechnological sectors includes biomedical, bio prospecting and biomass industries.

Course Content

UNIT-I

12Hr

Introduction: Fermenter design and types of fermenters, achievement and maintenance of aseptic conditions, Types of fermentation processes (Surface, submerged, Batch, Continuous, solid-substrate, Dual, Fed batch fermentation and its applications),

Industrial Microorganisms: Screening, Isolation. Identification and characterization of industrially important microbes. Strain improvement- mutation, recombination- gene regulation and genetic manipulation. Preservation of industrially important microbes. Culture collection centers.

UNIT-II

12Hr

Media for Industrial Fermentations: Media formulation, growth factors, carbon, nitrogen, Energy and Mineral sources, buffers, inhibitors, precursors, inducers, Oxygen requirements Antifoam agents and others, Sterilization: Sterilization of bioreactor, media, air and exhaust air and filter sterilization

Downstream processing and fermentation economics: Steps in recovery and purification Methods of cell separation – filtration and centrifugation, cell disruption, liquid liquid extraction, chromatography, membrane processes. Fermentation economics- expenses for industrial organisms, strain improvement, media sterilization, heating, cooling, aeration and agitation. Cost of Plant and equipments, batch process cycle time, continuous culture, recovery and effluent treatment, cast recovery due to waste usages and recycling.

UNIT-III

12Hr

Industrial production of energy fuels: Industrial alcohol production: Biosynthesis, methods of production, recovery and applications of ethanol, acetone – butanol and glycerol through microbial process.

Industrial production of Organic acids and Enzymes: biosynthesis, media, production process, product recovery and application of citric acid and lactic acid, Enzymes: Fungal and Bacterial Amylase; Bacterial proteases.

UNIT-IV

12Hr

Industrial production of food additives: amino acid production, methods of production, product recovery of L-Glutamic acid and L-lysine. Commercial uses of Amino acids Vitamins: Commercial production of Vitamin B₁₂, and Riboflavin. Alcoholic beverages (Beer, Wine,)

Industrial production of health care product: Industrial production of β -lactum antibiotic

(Penicillin): Biosynthesis, production and recovery. Streptomycin. Biosynthesis, production and recovery. Antitumor and anticholesterol agents, SCP and SCO, I P R: Patent Laws: Patent regulations of processes, products and microorganisms.

References:

1. Barsanti, L and Gualtieri, P. 2005. Algae: Anatomy, Biochemistry, and Biotechnology. Taylor and Francis New York.
2. Casida, L.E. 1997. Industrial Microbiology. New Age International Publishers.
3. Crueger, W. and Crueger, A. 2003. Biotechnology- A text book of Industrial Microbiology. Panima Publishing corporation.
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18. Peppler, H.J. and Perlman, D. 2005. Microbial Technology: Fermentation Technology Second Edition Volume 2. Elsevier India Private Limited.
19. Puri, R.S. and Viswanathan, A. 2009. Practical Approach to Intellectual Property Rights. I.K. International Publishing House. New Delhi.
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Course-IV: MEDICAL MICROBIOLOGY

48Hr

Course Outcome:

- This course provides knowledge regarding basic principles of medical microbiology and infectious disease.
- It covers mechanisms of infectious disease transmission, principles of aseptic practice, and the role of the human body's normal microflora
- It also provides opportunities to develop informatics and diagnostic skills, including the use and interpretation of laboratory tests in the diagnosis of infectious diseases.
- To understand the importance of pathogenic bacteria in human disease with respect to infections of the respiratory tract, gastrointestinal tract, urinary tract, skin and soft tissue and the methods to control diseases, e.g. chemotherapy & vaccines.

Pedagogy

Medical microbiology, the large subset of microbiology that is applied to medicine, is a branch of medical science concerned with the prevention, diagnosis and treatment of infectious diseases. In addition, this field of science studies various clinical applications of microbes for the improvement of health. It is the study of microbes, such as bacteria, viruses, fungi and parasites, which cause human illness and their role in the disease.

Clinical microbiology laboratories perform aerobic and anaerobic bacteriology, parasitology, mycobacteriology, mycology, and virology. Clinical microbiology is also a rather complex discipline because it utilizes many different types of methodologies and constantly undergoes changes in testing methods. Methods used to diagnose microbial diseases, and the microbiology laboratory may comprise several disciplines (e.g., classical culture methods, antigen detection methods, molecular methods, and serological methods are often performed under the purview of microbiology). The wide variety of pathogens and testing methods that are available makes microbiological testing challenging. In the clinical microbiology laboratory, as in every other discipline, the frequency of analytical errors has been reduced considerably with the implementation of quality control and quality assurance programs.

COURSE CONTENT

UNIT-I

12Hr

Introduction to Medical Microbiology: History, Development and scope of Medical Microbiology. Concept of Disease, disorder, syndrome, Communicable diseases- Microbial infections and diseases. Factors responsible for microbial pathogenicity.

Microbial infections: Types of infections, modes of transmission, portal of entry: Urinary tract infection, sexually transmissible infection, Infection of the central nervous system, Infections of circulatory system, Oral cavity and respiratory infection, gastrointestinal infection.

UNIT-II

12Hr

Nosocomial infection: Incidence of nosocomial infections, types of nosocomial infections, emergence of antibiotic resistant microorganisms, hospital infection control programmes, preventing nosocomial infections and surveillance, General concepts for specimen collection and handling of specimen, specimen processing and biosafety.

Chemotherapeutic agents: antibiotics (Classification based on chemical structure, mode of action and range of effectiveness). Recent trends-Drug resistance and its consequences, antibiotic policy, NCCLS (CLSI) guidelines and standards, WHO guidelines.

UNIT-III

12Hr

Epidemiology, Pathogenesis, Spectrum of disease, Laboratory diagnosis and Prevention: Diseases caused by Viruses: Chicken pox, Rabies virus, hepatitis, encephalitis, AIDS, Herpes simplex infections, Influenza, Dengue

Diseases caused by Bacteria: Tuberculosis, Leprosy, Cholera, Typhoid, Botulism, Shigellosis, Helicobacter pylori infection, Salmonellosis, Tetanus. Diseases caused by Fungi: Candidiasis, Histoplasmosis, Blastomycosis, Coccidiomycosis, Dermatophytosis, Aspergillosis and Cryptococcosis, Anthrax

UNIT-IV

12Hr

Diseases caused by Mycoplasma: *Mycoplasma pneumoniae*, *M. urealyticum*, *M. hominis*.

Diseases caused by Protozoa: Giardiasis, Trichomoniasis, Cerebral Malaria, Toxoplasmosis, Cryptosporidium.

Disease caused by Chlamydiae: Psittacosis, Lymphogranuloma Venereum, Trachoma and Inclusion conjunctivitis.

Emergent Diseases: Hemorrhagic fever, Swine flu, SARS, Chikungunya, Ebola, Hanta, Leptospirosis, Marburg

References:

1. Robert W. Bauman Ph.D. (2011) Microbiology with Diseases by Body System (3rd Edition); Benjamin Cummings
2. Patrick R. Murray PhD, Ken S. Rosenthal PhD, Michael A. Pfaller MD (2012) Medical Microbiology; Saunders
3. Brooks, Geo F., Carroll, Karen C., Butel, Janet S. (2012) Jawetz Melnick & Adelbergs Medical Microbiology ; McGraw-Hill Medical Publishing Division
4. Kenneth Ryan, C. George Ray , Nafees Ahmad , W. Lawrence Drew, Michael Lagunoff , Paul Pottinger, L. Barth Reller, Charles R. Sterling (2014) Sherris Medical Microbiology, Sixth Edition; McGraw-Hill Medical
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8. Connie R. Mahon , Donald C. Lehman , George Manuselis Jr. (2010) Textbook of Diagnostic Microbiology ; Saunders
9. Ananthanarayan , Paniker (2009) Textbook of Microbiology , 8th Edition; University Press
10. Jawetz (2010) Medical Microbiology , 25th Edition; Tata McGraw - Hill Education

COURSE-V: CLINICAL & DIAGNOSTIC MICROBIOLOGY

48Hr

Course Outcome:

- Various bacterial, viral, fungal and protozoal disease their causative agent, mode of infection, epidemiology, treatment, lab diagnosis, prophylaxis.
- students will develop skill regarding Isolate and identify microorganism from laboratory sample,
- Antibiotics sensitivity and resistance test
- Detection of parasite
- Handling of blood and body fluids

Pedagogy:

Clinical and Diagnostic Microbiology is a specialty within the sciences which focuses on applying microbiology to medical application. Similarly to being concerned with the identification of a disorder-inflicting organism, diagnostic microbiology can also be a part of modifying a treatment plan. Microbes including bacteria, protozoans, and fungi play a vital factor in many disease processes. The various laboratory techniques like microscopy, immunological assessments, radiology, biomarker tests, ELISA, serology checks, vaccines vectors are the primary diagnostic tests which are currently in use. Many microbes have developed resistance to medications. Hence, it's far essential for the scientists to give smarter methods of diagnosing those microbes and their pathogenic mechanisms.

Course Content

UNIT-I

12Hr

Introduction to clinical Microbiology: Role of Microbiologist in Diagnostic laboratory, General concepts for specimen collection, handling, transportation, processing, specimen workup, Laboratory safety and infection control.

Scientific and Laboratory basis for Clinical/Diagnostic Microbiology: Microscopic examination of infectious diseases, Growth and biochemical characteristics, Rapid methods of identification.

UNIT-II

12Hr

Immunotechniques and Immunodiagnosis: Antigens and Antibody reactions *in vitro*; Agglutination, complement fixation, ELISA, Western Blotting Immunodiffusion, Immunoelectrophoresis, Immunofluorescence, Immunoprecipitation, Radioimmunoassay and serotyping.

Vaccines and Vaccination: Vaccines – definition, types, Antigens used as Vaccines, effectiveness of vaccines, Vaccine safety, current vaccines, adjuvants, active immunization and passive immunization.

UNIT-III

12Hr

Recent Diagnostic tools and techniques: Principle, working and application of a) Autoanalyser b) Biosensor glucometer c) Diagnostic kits- ELISA, Western Blot
Enzymes in Disease diagnosis and therapy: Lactate dehydrogenase, Aspartate aminotransferase, Alkaline phosphatase, Creatine kinase, Acid phosphatase, Cholinesterase.

UNIT-IV

12Hr

Antimicrobial Chemotherapy: Development of chemotherapy; General characteristics of drugs and their testing; Mechanism of action. Antibacterial drugs; antifungal drugs, antiviral and antiprotozoan drugs; antibiotic sensitivity testing, MIC, Drug resistance; mechanism of drug resistance; multi drug resistance.

Reference

1. GouraKudesia (2009) Clinical and Diagnostic Virology. Cambridge University Press.UK.
2. J. Andre Knottnerus and Frank Buntinx (2008) The Evidence Base of Clinical Diagnosis: Theory and Methods of Diagnostic Research, 2nd Edition. WileyPublication.
3. Huggett and Justin O'Grady *LGC (2014)* Molecular Diagnostics: Current Research and Applications. Caister AcademicPress.
4. Vinay Kumar et al., (2010) Robbins and Cotran pathologic basis of disease. Philadelphia, PA: Saunders/Elsevier.
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9. Stephen B. Hulley; Steven R. Cummings; Warren S. Browner; Deborah G. Grady; Thomas B. Newman (2007) Designing clinical research (3rd edition). Philadelphia, PA: Lippincott Williams & Wilkins.
10. HuwLlewelyn , Hock AunAng, Keir E Lewis and Anees Al-Abdullah (2009). Oxford Handbook of Clinical Diagnosis. Oxfordpublications.
11. Peter Hu MadhuriHegde and Patrick Alan Lennon (2012). Modern Clinical Molecular Techniques. Springerpublications.
12. HenrikWinther and Jan T. Jorgensen (2010). Molecular Diagnostics. Springerpublications.
13. Prakash S. Bisen, MousumiDebnath and GBKS Prasad (2010) Molecular Diagnostics: Promises and Possibilities. Springerpublications

PRACTICAL V (Molecular Biology and Genetic engineering)

1. Isolation of Genomic DNA from *E. coli*.
2. Determination of purity and concentration of isolated DNA using spectrophotometer
3. Separation of proteins by SDS PAGE.
4. Salt fractionation of Yeast protein and quantification.
5. Isolation of plasmids from bacteria by agarose gelelectrophoresis.
6. Estimation of DNA
7. Estimation of RNA
8. Estimation of protein by Lowry's method

9. Digestion of the gene of interest with suitable restriction enzymes.
10. Ligation of the digested gene in a vector.
11. Preparation of competent *E. coli* cells for Bacterial transformation.
12. Transformation of the vector into the host cell and selection of the desired clones.
13. Induction of gene expression and purification of the induced protein from the host.
14. Amplification, Purification and separation of PCR product.
15. Determination of DNase activity on isolated DNA.
16. Determination of RNase activity on isolated RNA.
17. Determination of Proteinase activity on proteins.

PRACTICAL VI (Industrial and Medical Microbiology)

1. Study design of Fermentor and Parameters
2. Isolation of antibiotic/ amino acid/organic acid producing microbes and their preservation.
3. Batch fermentation of Citric acid production, recovery and estimation of citric acid.
4. Production of any vitamin and its quantification by bioassay.
5. Antibiotic fermentation and estimation of penicillin.
6. Preparation of wine and estimation of alcohol by specific gravity method.
7. Alcoholic fermentation and determination of total acidity and non-reducing sugars
8. Preparation of banana juice using Pectinase
9. Pathogenic fungi of the skin (Dermatophytes).
10. Microbial flora of mouth – teeth crevices.
11. Microbial flora of saliva.
12. Microorganisms of respiratory tract – examination of sputum/ AFB acid – fast bacteria.
13. Estimation of bacteria in urine by calibrated loop direct streak method.
14. Antimicrobial assay – sensitivity test (MIC) for pathogenic bacteria.
15. Laboratory diagnosis of important human diseases: Diphtheria, Tuberculosis, Typhoid, Wound infections, Malaria, Leprosy, AIDS and Hepatitis.

OPEN ELECTIVE

Course-VIII : MICROBIAL TECHNOLOGY

48Hr

Course Outcome:

- To acquire knowledge on food product analysis
- To enable them to know about preservation of pharmaceutical products
- Learn to assess the microbial quality of marine foods Outcome
- Acquire Knowledge on food product analysis
- Impart knowledge of preservation technology.
- Knowledge on quality analysis of marine food products

Pedagogy:

It is a sub-discipline of Microbiology which focuses on microbiological techniques or methods used for the study of microbes, including bacteria, fungi and protists. This course helps students learning fundamental procedures and safety guidelines followed in the microbiology laboratory.

This course teaches students the basic skills necessary to be successful in the laboratory as well as provides easy to follow, step-by-step, directions on how to perform each technique based in microbiology. The lecture will impart the students with knowledge and skills about how to culture, stain, identify, preserve and control of microorganisms. The skills and knowledge gained about techniques in microbiology helps the students to work in the laboratories like food and dairy industries, pharmacological industries, clinical, health and diagnostics laboratories, and any industries where microorganisms are used.

COURSE CONTENT

UNIT-I

12Hr

Microscopy: Light microscopy- Simple microscopy (dissection microscope), Compound microscopy (Bright field, Dark field, phase contrast, and Fluorescence microscopy) and stereomicroscopy. Electron microscopy: Principles, construction and mode of operation of scanning and Transmission electron microscopy, limitations. Preparation of specimens for electron microscopic studies (Ultra-thin sectioning, negative staining, shadow casting and freeze etching).

UNIT-II

12Hr

Microbiological stains and staining techniques: Types of stains and principles of staining. Stains for bacteria, fungi, algae and protozoa, spirochetes, stains for azotobacter cysts, stains for mycoplasma. Preparation of bacterial smears for light microscopy: Fixation, simple staining, Differential staining, Structural staining (Capsule, Flagella, Cell wall and Endospore of bacteria), and nuclear staining.

UNIT-III

12Hr

Culture media for Microbes Types of media- general purpose media, special purpose mediaselective, elective, diagnostic, resuscitation media, Media for fungi, algae, bacteria, mycoplasma andviruses.

Sterilization techniques: Principles, types of Sterilization, and their mode of action. Physical methods: Heat-dry heat (Hot-Air oven), Incineration, Moist heat (Autoclave and Pressure cooker), Tyndalization (Fractional Sterilization), Filtration-Types of filters, Laminar airflow. Radiation methods (UV radiation, x- rays and cathoderays).

UNIT-IV

12Hr

Control of Microorganisms: Chemical methods: Definition of terms- Disinfectants, Antiseptics, Sanitizers, Microbicides (bactericide, fungicide and Sporicide),

Microbistatic (bacteristatic and fungi static agents). Use and mode of action of Alcohols, Aldehydes, Halogens, Phenols, Heavy metals, and Detergents.

Pure culture techniques: Different types of inoculation techniques - Spread plate, Pour plate and Streak plate methods

References:

1. Alcomo, I.E. 2001. Fundamentals of Microbiology. VI Edition, Jones and Bartlett Publishers, Sudbury, Massachusetts.
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**FOURTH SEMESTER
HARDCORE**

Course-I: AGRICULTURAL MICROBIOLOGY

48Hr

Course Outcome:

- Approaches used in agriculture to control disease in plant
- Microbial ecology and microbial interaction
- Pathogenic interactions with plant

- Microbial biocontrol agents

Pedagogy:

Agricultural microbiology is a branch of microbiology dealing with plant associated microbes. It also deals with microbiology of soil fertility, such as microbial degradation of organic matter and soil nutrient transformations. It aims to address problem in agricultural practices usually caused by lack of biodiversity in microbial communities.

An understanding of microbial strains relevant to agricultural applications is useful in the enhancement of factors such as soil nutrient, plant pathogen resistance, crops robustness fertilization uptake effectively. The many symbiotic relationship between plant and microbes can ultimately be exploited for greater food production necessary to feed expanding new population safer to minimize the ecological disruption. The microbes are also used as bio fertilizers, bio pesticides, and fungicides. Agricultural microbiology also explains about the plant pathogen and the control measures. The use of techniques for the proper harvest and storage of the crops and its prevention from the contamination by microorganisms.

The syllabus includes four disciplines which deal with the introduction to agricultural microbiology, the plant pathology, parasitism and disease development, the defense mechanism of plant, plant disease and their management, the microbes and plant interaction and the bio pesticides.

Course Content

UNIT-I

12Hr

Introduction to Agricultural Microbiology:, Introduction to agricultural microbiology, concepts and scope of agricultural microbiology, Agronomy and production of important crop plants, Green revolution. **Plant Pathology:** Concept of disease, History of Plant Pathology, Significance of plant diseases, Symptoms and types of plant diseases.

Plant Pathology in Practice: Plant Clinic and Plant Doctor Concept. Diagnosis of Plant Diseases – Infectious diseases, Non-infectious diseases, Kochs' rules;

UNIT-II

12Hr

Parasitism and Disease Development Parasitism and pathogenicity, Host range of pathogens, Disease triangle, Diseases cycle / Infection cycle, Relationship between disease cycles and epidemics; Pathogens Attack Plants – Mechanical forces, Microbial enzymes and toxins, Growth regulators. Effect on physiology of Host – Photosynthesis, Translocation and transpiration, Respiration, Permeability, Transcription and translation. Environment and Plant Disease– Effect of Temperature, Moisture, Wind, Light, Soil, pH and structure, Nutrition and Herbicides.

Defense Mechanisms of Plant: Disease Pre-existing structural and chemical defenses, Induced structural and biochemical defenses. Microbe mediated strategies for abiotic stress management.

UNIT-III

12Hr

Plant Disease & their management: Tobacco Mosaic Disease, Sandal Spike Disease, Bacterial blight of Paddy, Citrus canker, Angular leaf spot of cotton, Late Blight of

Potato, Downy Mildew of Bajra, Blast of paddy, Tikka disease of ground nut, Rust of coffee, Grain and Head smut of Sorghum. Powdery mildew of Cucurbits, Wilt of Tomato, and Root Knot of Mulberry. Bunchy top of Banana.

UNIT-IV

12Hr

Microbes and Plant interaction-Mycorrhizae-Biology and their applications, Biofertilizers - microbial inoculants. Production and application of *Rhizobium*, *Azospirillum*, *Azotobacter*, phosphor bacteria and Cyanobacteria. PGPR's plant growth promoting *Rhizobacteria* and their uses.

Biopesticides: Definition, types-bacterial, viral, fungal and protozoan, mode of action, target pests, use of transgenic plants. mode of action, Bacteria-endo and ecto-toxins production by *Bacillus thuringiensis*, and *Pseudomonas*. Fungi- *Beauveria*, *Cephalosporium*, and *Trichoderma*.

References:

1. George. N. Agrios (2005), Plant pathology, Elsevier academic press, 5th edition, U.K.
2. Mehrotra. R.S. and Ashok Aggarwal (2002), Plant pathology, Tata MC Graw-Hill publishers, 2nd edition, Delhi.
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4. Bagyaraj D.G. and Rangaswami. G. (2005). Agricultural Microbiology, Prentice- Hall of India, 2nd edition, New Delhi.
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8. Roger Hull (2013). Plant virology, Elsevier academic press, 1th edition, U.K.
9. Hermann H. Prell, Peter R. Day. (2001). Plant-Fungal Pathogen Interaction: A Classical and Molecular View, 1st edition, Springer-Verlag Berlin Heidelberg, Germany.
10. Geoffrey Clough Ainsworth (1981). Introduction to the History of Plant Pathology 1st edition, Cambridge university press, U.K.
11. Vidhyasekaran, P. (2007). Fungal Pathogenesis in Plants and Crops: Molecular Biology and Host Defense Mechanisms, 2nd edition, APS press, U.S.A

SOFTCORE

COURSE -II: ENVIRONMENTAL MICROBIOLOGY

48Hr

Course Outcome:

- Will have an overview of the till date developments in the field of environmental microbiology with special emphasis on the role of microbes in mitigating environment pollution.
- Will have become acquainted with various cultural, biochemical and molecular techniques used in understanding microbial diversity.
- Will be knowledgeable about the diversity, adaptations and biotechnological applications of microbes of extreme environment.
- Is able to describe the role of microbes in solid and liquid waste management, gaining knowledge of various methods employed in sewage treatment and solid waste treatment.
- Understands the role of microbes in bioremediation of environmental pollutants like petroleum hydrocarbons, pesticides, plastic and electronic waste; also understands utility of microbes in mineral and oil recovery.

Pedagogy:

Microbial communities control nutrient cycles and biogeochemical transformations in natural, managed and engineered ecosystems. Microorganisms recycle organic matter, transform contaminants, and maintain ecosystem health. Understanding the ecology of natural microbial communities will deepen our understanding of how ecosystems function. Since microbial communities are critical for ecosystem function, microbial ecology can also assist the development of models to predict how ecosystems will respond to future environmental conditions.

Environmental Microbiology introduces students to the diversity of microbial populations and their important roles in environmental processes in air, water, soils, and sediments. Microbial community ecology and interactions with plants and animals will also be discussed. Students will learn how microbial activities sustain natural ecosystems. Techniques for characterizing microorganisms and investigating microbial processes will also be discussed. The lectures impart knowledge to utilize bio wastes from industrial and agricultural fields. This recycling of bio wastes leads to costless and environment clean .

COURSE CONTENT

UNIT-I

12Hr

Air Microbiology: Airspora of indoor and outdoor environment, factors affecting airspora, Techniques of trapping air borne microorganisms.

Aquatic Microbiology: Distribution of microorganisms in the aquatic environment, Water pollution sources, Biological indicators of water pollution, Determination of sanitary quality of water, Waste water microbiology-Primary, secondary, tertiary treatment and reclamation of waste water

UNIT-II

12Hr

Soil Microbiology: Characteristics and classification of soil. Interactions between microorganisms: Mutualism, commensalism, ammensalism synergism, parasitism, predation, competition. Rhizosphere, rhizosphere, microflora and its beneficial activity. Role of microorganism in nitrogen, phosphorous and sulphur cycle. Detrimental effects of diverted biogeochemical cycles. Biological nitrogen fixation in detail:

Symbiotic, asymbiotic and associated nitrogen fixation. Structure, function and genetic regulation of nitrogenases. Viable but nonculturable bacteria.

UNIT-III

12Hr

Microbes in extreme environment: Microbes of extreme environments, Thermophiles, acidophiles, alkaliphiles, halophiles, barophiles and their survival mechanisms.

Space microbiology: Historical development of space microbiology, Life detection methods

a) Evidence of metabolism (Gulliver) b) Evidence of photosynthesis (autotrophic and heterotrophic).

UNIT-IV

12Hr

Microbes in the degradation of wastes: Treatment of solid and liquid industrial wastes, Microbial degradation of pesticides, Xenobiotics, degradation of lignin, cellulose and pectin. Bioremediation. Geomicrobiology: Microbes in metal extraction, mineral leaching and mining, copper extraction by leaching and microbes in petroleum product formation. Global Environmental Problems: Global Warming, Acid rain, Ozone depletion. Bio deterioration of wood and metals.

COURSE-III: GENOMICS AND PROTEOMICS

48Hr

Course Outcome:

- The aim of this course is to teach genomics, transcriptomics, proteomics, metabolomics and phenomics using model organisms representing plants and animals.
- The course will cover recent developments in genomics, gene expression and small RNAs, synthetic biology, epigenetics, proteomics, fast-forward genetics and next-generation mapping.
- An objective of the course is to develop skills in experimental design within the context of learning about biology including: regulation of transcription and translation, stress response, signal transduction and the engineering and regulation of metabolic pathways.

Pedagogy:

Genomics is the study of entire genomes, including the complete set of genes, their nucleotide sequence and organization, and their interactions within a species and with other species. The advances in genomics have been made possible by DNA sequencing technology. Genomic information is used to create similar maps of the DNA of different organisms. *Proteomics* generally refers to the large-scale experimental analysis of proteins and proteomes

Bioinformatics helps the students to understand Genomics and proteomics which uses the computational knowledge to extract the knowledge from biological database. This helps in data analysis, visualization, prediction, primer designing, data storage etc., through web based tools like NCBI. Students are able to understand and use the knowledge of bioinformatics and do Insilco analysis to verify and test their hypothesis

before they start their wet lab experiments. Bioinformatics helps in drug discovery and students will be placed in pharmaceutical and drug companies.

COURSE CONTENT

UNIT-I

12Hr

Genome - Overview Of Genome; Sequence Of Genome Acquisition And Analysis - Homologies - Snps - Genetic Analysis, Linkage Mapping, High **Resolution Chromosome Mapping And Analysis** - Physical Mapping, Yac, Hybrid Mapping, Strategies, Sequence Specific Tags (Sst), Sequence Tagged Sites(Sts), Ish, Fish, Rflp,Rapd.

UNIT-II

12Hr

DNA Sequencing - Methods, MaxamAnd Gilbert Method, Ladder, Fluorescent, Shot Gun, Mass Spectrometry, Automation Sequencing – Find Gene Mutations, Implications of DNA – Sequencing And Sequencing Genomes.

Genome Data Bank, Metabolic Pathway Data - Construction And Screening Of cDNA, Libraries And Microarrays - Application Of DNA Arrays - PCR - Variations In PCR - Gene Disruptions – Sage And Sade, Pharmacogenomics.

UNIT-III

12Hr

Protein Sequence Analysis - Introduction - Sequence Data Banks - Wbrf – Pir - Swissport - Databases, Data Mining - Algorithms Of Proteomics And Its Applications - Protein Expression

Profiling - Protein - Protein Interaction - Protein Modifications. Automation - Nucleic Acid Data Bank – EMBL Nucleotide Sequence Data Bank - Aids Virus Sequence Data Bank - RNA DataBank.

UNIT-IV

12Hr

Tools For Data Bank - Pairwise Alignment - Needleman And WunschAlgorithm – Smith Waterman - Multiple Alignment - Clustral - Pras - Blast - Fast, Algorithms To Analyse Sequence Data - Pdb, Cambridge Structure Data Base (Lsd), 2d Electrophoresis, Ief, Hplc, Protein Digestion Technique, Mass Spectrometry, Maldi, Tof, Peptides, Mass Finger, Printing,Protein.


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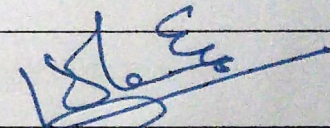
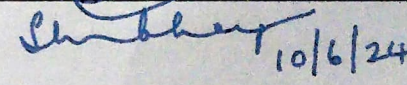
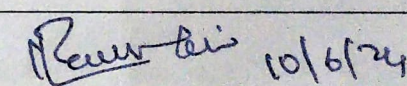
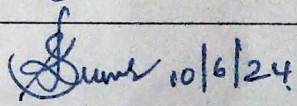
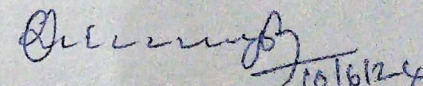
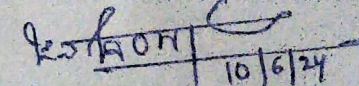
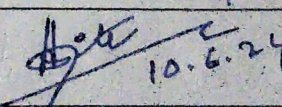
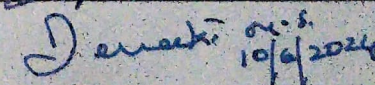
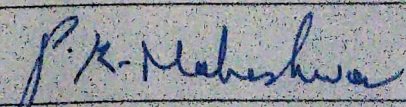
COURSE-IV: PRACTICAL VII (Agricultural Microbiology & Environmental Microbiology)

1. Isolation, culturing and seed inoculation of *Rhizobium* and testing of nodulation ability and beneficial effects.
2. Isolation and testing the efficiency of various biofertilizers like *Rhizobium*, *Azotobacter*, *Azospirillum*.
3. Mass multiplication techniques of *Azolla*.
4. Estimation of total phenols in diseased and healthy plant tissues.
5. Seed health testing by SBM.
6. Collection and Identification of following disease: Tobacco mosaic disease, Bunchy top of Banana, Bean Mosaic, Sandal spike, Bacterial blight of paddy. Citrus canker, Downy mildew of Bajra, Powdery mildew of mulberry, Head smut of sorghum, Leaf rust of coffee, Blast disease of paddy, Tikka disease of groundnut, Leaf spot of paddy and Grassy shoot of sugarcane.
7. Isolation and identification of micro flora of soil, sewage and air
8. Microbes as indicators of water pollution – Determination of indices of water quality.
9. Determination of BOD of pollution water.
10. Determination of COD of polluted water.
11. Degradation of cellulose by *Chaetomium globosum*.
12. Bacterial examination of drinking water by membrane filter technique.
13. Study of associated soil microorganisms with plants, Actinorhiza, Mycorrhiza.
14. Study of important microbes in the degradation of wastes.
15. Isolation of cellulose degraders, chitinase and pesticide degraders
16. Determination of TS and MLSS


UNIVERSITY OF MYSORE
(Re-accredited by NAAC with 'A' Grade)
(NIRF-2023: Ranked 44 in University Category & 71 in Overall Category)
DEPARTMENT OF STUDIES IN MICROBIOLOGY

Proceedings of the BOS Meeting of Microbiology (CB)

As per the direction of the Registrar, University of Mysore, Mysuru, Board of Studies Meeting of Microbiology (CB) of University of Mysore was conducted on 10/06/2024 at Department of studies in Microbiology, Manasagangothri, University of Mysore, Mysuru. The information about the discussion carried out in the meeting is enclosed with this letter. The details of the BoS members who participated.

Sl.No	Name of the Bos Member	Designation	Signature
1.	Prof. M.Y. Sreenivasa	Chairman	
2.	Prof. Shubha Gopal	Member	 10/6/24
3.	Prof. K. Ramachandrakini	Member	 10/6/24
4.	Prof. V. Shyam Kumar	Member	 10/6/24
5.	Prof. Virupakshaiah, DBM	Member	 10/6/24
6.	Prof. Srinivas C	Member	Attended online
7.	Dr G. S Siddegowda	Member	 10/6/24
8.	Dr. H S. Jayanth	Member	 10.6.24
9.	Dr. N.S. Devaki	Member	 10/6/2024
10.	Dr. N.Anuroopa	Member	Absent
11.	Dr. P.K. Maheshwar	Member Special Invitee	

Details of the Discussion & Decisions Taken at BoS Meeting of Microbiology (CB)

1. The Chairman BoS in Microbiology welcomed the BoS members and placed the BoS Meeting Agenda before the members.
2. Discussed and approved the changes in the course of B.Sc., Microbiology syllabi and Scheme of examination 2024-2025
3. Discussed and approved the course of M.Sc., Microbiology syllabi and Scheme of examination 2024-2025
4. Prepared and approved of Question paper pattern & scheme of Practical Examination 2024-2025
5. Prepared and approved the Panel of examiners for UG & PG examinations 2024-2025
6. Any other Matter:
 - Opinion regarding: Guide Recognition to Dr. Ragavendra M P, Professor, Maharani, Science College, Mysuru (Ref.No.Dor/9.5/Ph.D/ROG-Mic/18/14/2024-25 Dtd:10-6-2024)
 - Decision: The members approved the Guide Recognition to Dr. Ragavendra M P.
7. The BoS meeting is concluded with the vote of thanks by the Chairman.

[Signature]
10/6/24

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P. Maheshwari

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10/6/24



UNIVERSITY OF MYSORE, MYSURU

CHOICE BASED CREDIT SYSTEM (CBCS)

&

CONTINUOUS ASSESSMENT AND GRADING PATTERN (CGPA)

Based

M. Sc. MICROBIOLOGY SYLLABUS

For

2024-25 onwards

**Board of Studies in Microbiology
Department of Studies in Microbiology
University of Mysore
Manasagangotri, Mysuru 570 006
Karnataka, India**

Introduction

Microbiology is an important and wide-ranging discipline within the life sciences, covering a range of subjects relevant to human health, diseases, environmental studies and industrial and biotechnological application. Microbiology has vast scope in understanding the life through intervention of microorganism. There is an increase in demand for microbiologist globally. A microbiologist can innovate new diagnostic kits, teach, research, discover new drugs etc., it encompasses many disciplines of science like medicine, dairy agriculture, pharmacy, nanotechnology etc., Knowledge and skills in Microbiology that will empower the students, through awareness of the significance of microorganisms in plant, animal and human health, environment, industry and general human welfare by a problem based and skill-oriented curriculum. The syllabus is highly oriented towards the complete knowledge of the subject, which includes the basic as well as contemporary applied aspects of Microbiology including molecular biology and genetic engineering.

Program Pedagogy:

The seminar presentation will improve the oration skills of students and group discussion will kindle their logical ability to analyze the problems. Assignments improve students in gathering the information and enhancing their writing ability. In practical laboratory they will be enhancing their skills towards various techniques used in the laboratory. As a part of curriculum, students work on project, which will give a hands-on experience on different techniques and will be a platform for the students to work and interact with different scientists and research institutions. This will pave the way for the students to know about recent research work going on in the field and help the student in working in different amenities.

Program outcome:

- The students get to know about different beneficial and harmful microorganisms, which might be useful / pathogenic to humans, animals and plants.
- Microbiology is concerned with diversified forms of microorganism, classification, structure, reproduction, physiology, metabolism and most importantly their economic importance.
- Industrial productions of organic acids, enzymes and pest control using microbes and improving soil quality and agricultural output and cleaning the environment through sustainable microbiological applications.
- To enable them to employ the acquired theoretical knowledge in the sector of Disease diagnosis, treatment and prevention.
- To enrich the post graduate students with fundamentals of microbiology and advanced technologies, which enables them use this knowledge in industry, hospitals, community and institutes or any other profession they would like to pursue.

Program specific outcome:

Understand the basic knowledge and concepts of microbiology and other related areas. Hands on skills in Industry and/or Institutes, for better placement in drug manufacturing companies, public health entities, blood service, industrial laboratories, cancer research institutes, R&D, educational institutes, environmental pollution control, agriculture and fisheries, food and dairy industry, forensic science, hospitals, public health laboratories, etc. There is requirement for microbiologist in quality control and safety sections of food, pharmaceuticals, health and beauty care, etc.

SCHEME OF THE STUDY

For M.Sc. in Microbiology

Credits to be earned	40 credits
Core papers	28 credits
Open elective paper	04 credits
Total credits	72 credits

Honors in Microbiology

Credit Based Choice Based Continuous Evaluation Pattern System Proposed Semester-wise distribution of the course structure for the year 2024-2025

Semester-I Credits: 18

No	Paper Code	Title of The Course Paper	Credit Pattern in L:T:P	Credits
1	MB 1.1 Hard core	Virology	3:1:0	4
2	MB 1.2 Hard core	Bacteriology	3:1:0	4
3	MB 1.3 Hard core	Mycology	3:1:0	4
		Select 3 among 4 papers		
4	MB 1.4 Soft core	Microbial Genetics	3:0:0	2
5	MB 1.5 Soft core	Microbial Ecology & Diversity	3:0:0	2
6	MB 1.6 Soft core	Practical I (Virology & Bacteriology)	0:0:2	2
7	MB 1.7 Soft core	Practical II (Mycology & Microbial Genetics)	0:0:2	2
		Total credits to be earned		18

HC=03; SC=03; O.E=0.

Semester-II Credits: 18

No	Paper Code	Title of The Course Paper	Credit Pattern in L:T:P	Credits
1	MB 2.1 Hard core	Microbial Physiology	3:1:0	4
2	MB 2.2 Hard core	Immunology	3:1:0	4
		Select 3 among 4 papers		
3	MB 2.3 Soft core	Food Microbiology	3:0:0	2
4	MB 2.4 Soft core	Soil Microbiology	3:0:0	2
5	MB 2.5 Soft core	Practical III (Microbial Physiology & Immunology)	0:0:2	2
6	MB 2.6 Soft core	Practical IV (Food Microbiology)	0:0:2	2
7	MB 2.7 OE	Techniques in Microbiology	4:0:0	4
		Total credits to be earned		18

HC=02; SC=03; O.E=1.

M.Sc. Microbiology

Credit Based Choice Based Continuous Evaluation Pattern System Proposed Semester wise distribution of the course structure

Semester-III Credits: 18

No	Paper Code	Title of The Course Paper	Credit Pattern in L:T:P	Credits
1	MB 3.1 Hard core	Molecular Biology	3:1:0	4
2	MB 3.2 Hard core	Genetic Engineering	3:1:0	4
3	MB 3.3 Hard core	Industrial Microbiology	3:1:0	4
		Select 3 among 6 papers		
4	MB 3.4 Soft core	Medical Microbiology	3:0:0	2
5	MB 3.5 Soft core	Clinical & Diagnostic	3:0:0	2
6	MB 3.6 Soft core	Practical V (Molecular Biology & Genetic Engineering)	0:0:2	2
7	MB 3.7 Soft core	Practical VI (Industrial Microbiology & Medical Microbiology)	0:0:2	2
8	MB 3.8 Soft core	MOOCS/ SWAYAM	2:0:0	2
9	MB 3.9 OE	Microbial Diversity	3:0:0	2
		Total credits to be earned		18

HC=03; SC=03; O.E=01.

Semester-IV Credits: 18

No	Paper Code	Title of The Course Paper	Credit Pattern in L:T:P	Credits
1	MB 4.1 Hard core	Agricultural Microbiology	3:1:0	4
2	MB 4.2 Hard core	Environmental Microbiology	3:1:0	4
		Select 3 among 7 papers		
3	MB 4.3 Soft core	Microbial Nanotechnology	4:0:0	4
4	MB 4.4 Soft core	Genomics & Proteomics	4:0:0	4
5	MB 4.5 Soft core	Practical VII (Agricultural Microbiology)	0:0:2	2
6	MB 4.6 Soft core	Practical VIII (Environmental Microbiology)	0:0:2	2
7	MB 4.7 Soft core	Practical IX (Microbial Nanotechnology)	0:0:2	2
8	MB 4.8 Soft core	Project Work	0:0:6	6
9	MB 4.9 Soft core	MOOCS/ SWAYAM	2:0:0	2
		Total credits to be earned		18

HC= 02; SC=03

Grand Total Credits: 72

SEMESTER I
MB1.1 Hardcore: VIROLOGY

Course Pedagogy:

- Knowledge on history, general characters of viruses and viral classification
- Understanding the replication strategies of viruses; Cultivation and detection of viruses.
- Comprehend evolutionary importance of viruses.
- Knowledge on some common plant and animal diseases caused by different viruses, viral transmission and control.

Course Outcome:

After the completion of the course students would be able

- To study the nature of viruses.
- Techniques employed for culturing and detection of plant and animal viruses
- To gain knowledge about newer emerging viral
- To unravel the mechanisms by which viruses infect cells and cause disease.
- Viruses used as cloning vectors for gene transfer, therapeutic agents.

THEORY

48 hours

UNIT I

12

hours

Viral Diversity: Classification – LHT, Baltimore & ICTV; and nomenclature of viruses. Replication patterns of the following groups; Group I – T2 Bacteriophage, Group II – Banana bunchy top virus, Group III – Reovirus, Group IV-TMV, Group V – Rhabdovirus, Group VI – HIV and Group VII – Hepatitis B virus. Microbial viruses: General account on algal, fungal, protozoan viruses, Giant viruses and Bacteriophages.

UNIT II

12

hours

Plant viruses: Propagation, Cultivation, Isolation and purification using centrifugation, chromatography and electrophoresis techniques. Detection and diagnosis of Plant Viruses
Cultivation and detection of animal viruses: Animal Inoculation, Inoculation into embryonated egg and Cell Culture. Direct methods of detection- light microscopy (inclusion bodies), electron microscopy (SEM, TEM, AFM and Cryo EM) and fluorescence microscopy. Immunodiagnosis: hemagglutination and hemagglutination inhibition test, complement fixation, neutralization, western blot, flow cytometry. Nucleic acid based diagnosis: nucleic acid hybridization, PCR, qRT, Microarray and nucleotide sequencing.
Infectivity assay for animal and bacterial viruses: Plaque assay, Transformation assay, Fluorescent focus assay, Infectious centre assay, end point dilution methods, LD50, ID50, EID50, TCID50.

UNIT III

12

hours

Sub-viral particles: Discovery, Structure, Classification, replication and diseases caused by Satellite virus, Virusoids, Viroids and Prions. Anti-viral strategies-prevention and control of viral diseases: Host specific and nonspecific defense mechanisms. Role of interferon in viral infections.
Viral Chemotherapy: Nucleoside analogs, reverse transcriptase inhibitors, protease inhibitors.
Conventional viral vaccines: killed and attenuated vaccines, Modern vaccines: peptide vaccines, edible vaccines, immune-modulators (cytokines), anti-idiotype, DNA and m-RNA vaccines.

UNIT IV

12

hours

Viral transformation and oncogenesis: Oncogenic viruses and viral transformation mechanism by EBV, HPV, and HTLV-1. **Viruses and the future:** Promises and problems. Evolutionary importance of viruses: Antigenic shift, antigenic drift. Emerging and life threatening diseases – COVID-19 and variants, KFD virus, and ZIKA. Sources and causes of emerging viral diseases. Viruses as threat of bioterrorism, as therapeutic agents, as gene delivery system, viruses to destroy other viruses. **Virus and nano-technology.**

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10. Moulay Mustapha Ennaji (2020), Emerging and Reemerging Viral Pathogens: Volume 1: Fundamental and Basic Virology Aspects of Human, Animal and Plant Pathogens 1st Edition. Academic Press.
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MB1.2Hardcore: BACTERIOLOGY

Course Pedagogy:

- To study the scope, history, economic importance, cell structure, growth, cultivation and control of bacteria.
- Working principles of microscopy and staining.

Course Outcome:

After the completion of the course students would be able:

- To know bacterial classification, nutrition, cultivation, preservation of microbial culture.
- To describe the morphological features, cell arrangement and structural components of bacterial cell.
- To enlist the characteristics of Archaea.
- To use different microscopes for studying bacterial morphology.
- To work in medical laboratories, pharmacological, food and fermentation industries.

THEORY 48 hours

UNIT I

12 hours

Introduction: Important events in development of bacteriology, Scope and relevance of bacteriology. Economic importance of bacteria.

Cell Structure: An overview of bacterial size, shape and arrangement, structure, chemical composition of cell wall of Archaeobacteria, gram-negative bacteria, gram-positive bacteria and acid fast bacteria, cell wall deficient organisms including L-form structure, composition and function of cell membrane, capsule, flagella, pili, Inclusion bodies, ribosomes, mesosomes, reserve food materials, magnetosomes and phycobilisomes, endospores, bacterial nucleic acids – chromosome, plasmid, transposons, integrons and antibiotic resistance cassettes.

Microscopy: Working Principles of brightfield microscope, fluorescent microscope, darkfield microscope, phase contrast microscope, stereomicroscope, confocal microscopy and electron microscope. Preparation of sample for electron microscopic studies. Application and importance of above microscopes. Measurement of microscopic objects.

UNIT II

12 hours

Bacterial classification and taxonomy: Criteria for the classification of bacteria. Phenetic, Phylogenetic, Genotypic, Numerical taxonomy. Techniques for determining microbial taxonomy and Phylogeny. ICNB rules. Classification systems of major categories and groups of bacteria according to Bergey are manual of Systematic Bacteriology and Determinative Bacteriology. Non-culturable methods for the identification of pathogenic microorganisms.

UNIT III

12 hours

Growth, Cultivation and control of Bacteria: Nutrient requirements, nutritional types of bacteria, culture media, classification of media. Growth: Nutritional uptake, Growth kinetics, generation time, growth curve, factors affecting growth. Methods for measurement of microbial growth – direct microscopy, viable count estimates, turbidometry, and biomass. Aerobic, anaerobic, batch, continuous and synchronous cultures. Methods of pure culture isolation, Enrichment culturing techniques, single cell isolation, and pure culture development. Preservation and Maintenance of Microbial cultures: Repeated sub culturing, preservation at low temperature, sterile soil preservation, mineral oil preservation, deep freezing and liquid nitrogen preservation, lyophilization. IUBS – International Union of Biological Sciences. World federation for culture collections – guidelines, statutes and bylaws.

Control of microorganisms: Antimicrobial agents, physical and chemical methods. Principles, functioning and types of Biosafety cabinets.

UNIT IV

12 hours

Characteristics and Salient features of major groups of Bacteria: Archaeobacteria: general characteristics and classification; extremophiles, halophiles, thermophiles and barophiles; General characteristics, classification, diversity and distribution, economic importance of **Actinomycetes, Cyanobacteria. Bioluminescent bacteria;** characteristics and examples, mechanism of bioluminescence. General characteristics, life cycle, growth, multiplication and significance of **Mycoplasma, Rickettsiae and Chlamydia.**

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10. Stuart Hogg (2013) *Essential Microbiology*, 2nd Edition, Wiley-Blackwell

MB1.3Hardcore: MYCOLOGY

Course Pedagogy:

- It includes the study of taxonomic classification, fungi as symbionts.
- Fungi in production of food supplements like SCP, vitamins, enzymes, organic acids and production of secondary metabolites like antibiotics.
- In practical classes they mount the fungi, learn microscopic views and the key characteristics to identify different species of fungi.

Course outcome:

After the completion of the course students would be able to

- To understand the general characteristics and reproduction in fungi and lichens.
- To understand the economic and pathological importance of fungi.
- To identify common fungal plant diseases and devise control measures and work as a plant doctor.

THEORY

48 hours

UNIT I

12 hours

Introduction: History and Development of Mycology, scope of mycology. Recent developments in Mycology.

Fungal taxonomy: Taxonomic problems associated with variation in fungi, Classification of fungi (Alexopoulos and Mims).

UNIT II

12 hours

General characteristics of fungi and reproduction: Morphology and somatic structures: The thallus, organization, fungal cell, nuclear components, specialized somatic structures; Aggregation of hyphae, tissues, mycelia, General aspects of fungal nutrition and reproduction (Asexual, Sexual reproduction, Heterothallic and Parasexuality)

UNIT III

12 hours

Salient features of fungal major groups: Chytridiomycota, Zygomycota, Basidiomycota, Ascomycota, Deuteromycota, Oomycota, Hypochytridiomycota, Labyrinthulomycota, Plasmodiophoromycota and Myxomycota. Symbiotic fungi - Lichens.

Opportunistic fungal infections: *Candida albicans*, *Aspergillus fumigatus* and Mucormycosis.

UNIT IV

12 hours

Economic importance of fungi: Fungi as biocontrol agent, Economic importance of Fungi in Agriculture, Industry and medicine. Fungi as SCP, Fungi as parasites of human and plants. Role of fungi in bio-deterioration of wood and paper. Mycorrhiza - ectomycorrhiza, endomycorrhiza, vesicular-arbuscular mycorrhiza. Fungi as insect symbionts.

Important metabolites of Fungi - aflatoxin, Ochratoxin, Ergot alkaloids, T-2 toxin, DON, Fumonisin. Impact of mycotoxins on human health. Importance of secondary metabolites of fungi as nephrotoxins, neurotoxins, hepatotoxins, mutagens/carcinogens.

Reference:

1. Alexopoulos CJ and Mims C W, 1979 Introductory Mycology 3rd edn, Wiley Eastern., New Delhi.
2. David Moore, Geoffrey D. Robson, Anthony P. J. Trinci (2011) 21st Century Guidebook to Fungi. Cambridge University Press.
3. Deacon, JW, 1997-Modern Mycology 3rd Edition, Blackwell Science publishers, London.
4. Kevin Kavanagh (2011) Fungi: Biology and Applications. John Wiley & Sons, Sussex, U.K.
5. Mehrotra, RS & Aneja, KR, 1998. An Introduction to Mycology. New Age International Pvt. Ltd. New Delhi.
6. Mercedes S. Foster & Gerald F. Bills (2011) Biodiversity of Fungi: Inventory and Monitoring Methods. Academic Press
7. Michael John Carlile, Sarah C. Watkinson, G.W. Gooday (2007) The fungi. Academic Press. London, U.K
8. Odum, E.P. 1971. Fundamentals of Ecology; Third Edition. Toppan Co. Ltd. Tokyo, Japan.

MB1.4Soft-core:MICROBIALGENETICS

Course Pedagogy:

- Describe the fundamental molecular principle of genetics.
- Understand the relationship between phenotype and genotype.
- Describe the basics of genetic mapping.
- Understand how gene expression is regulated

Course Outcome:

After the completion of the course students would be able

- To understand the genetic constituents of bacteria with special emphasis on inheritance.
- To extend the knowledge on molecular basis of mutation at microbial level.
- To focus on gene regulation and expression mechanisms.
- To understand the principle role of plasmids and gene transfer methods and mapping.

THEORY

48 hours

UNIT I

12 hours

Concepts in Microbial Genetics: History and developments of Microbial genetics. Essentials of microbial genetics: Microbes as Genetic Tools for Basic and Applied Genetic studies. Advantages and disadvantages of Microbes, Generalized reproductive cycles of microbes- *Neurospora*, *Saccharomyces*, *Chlamydomonas* and *Acetabularia*.

UNIT II

12 hours

Viral Genetics: Lytic and Lysogenic cycles, Phage Phenotypes, Phenotypic Mixing, Recombination in viruses: Mutations, Recombination and Mapping (rIII loci)

Bacterial Genetics: Bacterial Transformation: Types of transformation mechanisms found in prokaryotes, Bacterial Conjugation: properties of the F plasmid, F⁺ x F⁻ mating, F' x F⁻ conjugation, Hfr conjugation, gene mapping in bacteria. Transduction: Generalized and specialized transduction, Transposable elements. Regulation of competence in *Bacillus*.

UNIT III

12 hours

Fungal Genetics: *Neurospora*- Tetrad analysis and linkage detection - 2 point and 3 point crosses, chromatid and chiasma interference, Mitotic recombination in *Neurospora* and *Aspergillus*.

Algal Genetics: *Chlamydomonas* - unordered tetrad analysis - Recombination and Mapping, Nucleocytoplasmic interactions and gene expression in *Acetabularia*. Extranuclear (Cytoplasmic) inheritance.

UNIT IV

12 hours

Mutation and mutagenesis: Nature, type and effects of mutations. Concept of gene: muton, recon and cistron. Mutagenesis - physical and chemical mutagens, base and nucleoside analog, alkylating agents, interrelating agents, ionizing radiation. Induction and detection of mutation in microorganisms. Site directed mutagenesis and its applications.

References:

1. D.PeterSnustad,MichaelJ. Simmons(2011)PrinciplesofGenetics, 6thEdition;Wiley
2. Dr.EvelynJ.Biluk(2012)MicrobiologyStudyGuide:MicrobialGenetics, ControllingMicrobialGrowth,and AntimicrobialAgents; CreateSpaceIndependentPublishingPlatform
3. James D. Watson, Tania A. Baker, Stephen P. Bell, Alexander Gann, Michael Levine, Richard Losick(2013)Molecular Biology ofthe Gene,7thedition; BenjaminCummings
4. JocelynE.Krebs,ElliottS.Goldstein, StephenT.Kilpatrick(2012)Lewin'sGENES XI,11thedition;Jones& BartlettLearning
5. JohnR.S. Fincham(1996)Microbial andMolecularGenetics; HodderArnold
6. LarrySnyder,JosephE.Peters,TinaM.Henkin, WendyChampness(2013)MolecularGeneticsofBacteria,4th Edition; ASMPress
7. NancyJoTrun,J.E.Trempey(2003)Fundamental BacterialGenetics;Wiley-Blackwell
8. RoystonC.Clowes,WilliamHayes(1968)ExperimentsinMicrobialGenetics;BlackwellScienceLtd
9. SriramSridhar(2005)GeneticsandMicrobialBiotechnology;DominantPublishers&Distributors
10. Stanley R. Maloy, Jhon E. Cronan, Jr. David Freifelder (1994) Microbial Genetics (Jones and BartlettSeriesin Biology),2nd edition; Jones and BartlettPublishers
11. UldisN.Streips,Ronald E.Yasbin(2002)ModernMicrobialGenetics,2nd edition;Wiley-Liss
12. VenetiaA.Saunders(1987)Microbialgenetics appliedtobiotechnology:principlesandtechniquesofgenetransferand manipulation;Springer

MB1.5 Soft-core: MICROBIAL ECOLOGY AND DIVERSITY

Course Pedagogy:

- To understand the ubiquitous nature of microbes.
- To give basic knowledge on extremophiles.
- To provide knowledge on characteristics of Microbes.

Course Outcome:

After the completion of the course students would be able

- Students able to differentiate various groups of Microbes.
- Get knowledge on adaptability of extremophiles.
- Knowledge about microbial taxonomy.

THEORY

48 hours

UNIT I

12 hours

Introduction to microbial ecology: Structure of microbial communities. Interaction among microbial populations. Interaction between microorganisms and plants. Biotransformation, biodegradation, bioremediation and phytoremediation. Ecological and Evolutionary diversity (Genetic diversity) of microbial world

Development of Microbial communities: Dynamics of community, ecological succession, structure, dispersion, microbial communities in nature and ecosystem models

UNIT II

12 hours

Physiological Ecology of microorganisms: Adaptation to environmental conditions - abiotic limitation to microbial growth.

Viral Diversity: Group I – T2 Bacteriophage, Group II – Banana bunchy top virus, Group III – Reovirus, Group IV – TMV, Group V – Rhabdovirus, Group VI – HIV, Group VII – Hepatitis virus.

Sub-

viral particles: Discovery, Structure, Classification, replication and diseases caused by Satellite, Satellite virus, Virusoids, Viroids and Prions.

UNIT III

12 hours

Bacterial Diversity: Archaeobacteria, Photosynthetic Eubacteria, Chemotrophic and Methophilic Eubacteria, Gliding Eubacteria, Spirochetes, Rickettsiae and Chlamydiae, Actinomycetes, Mollicutes, Protists

Fungal Diversity: salient features of the following group: Zygomycota (*Rhizopus*), Ascomycota (*Neurospora*), Basidiomycota (*Agaricus*), Deuteromycota (*Penicillium*), Chytridiomycota (*Allomyces*) Myxomycota and Yeast.

UNIT IV

12 hours

Importance and Conservation of Microbial Diversity: Importance of microbial diversity in environment, pharmaceuticals & human health. Metagenomics. Importance of conservation. *In situ* conservation and *Ex situ* conservation. Role of culture collection centers in conservation.

References

1. Atlas, Ronald M., Bartha, Richard (1997) Microbial Ecology Fundamentals and Applications; Addison-Wesley
2. Colwell, R.R., Simidu, Usio, Ohwada, Kouicki (1996) Microbial Diversity in Time and Space; Springer
3. David L. Kirchman (2008) Microbial Ecology of the Oceans; Wiley-Liss
4. David L. Kirchman (2012) Processes in Microbial Ecology; Oxford University Press
5. James W. Brown (2014) Principles of Microbial Diversity; ASM Press
6. McArthur, J. Vaun (2006) Microbial Ecology An Evolutionary Approach; Academic Press
7. Nelson, Karen E. (1997) Advances in Microbial Ecology; Springer
8. Oladele Ogunseitan (2004) Microbial Diversity: Form and Function in Prokaryotes; Wiley-Blackwell
9. Oladele Ogunseitan (2008) Microbial Diversity: Form and Function in Prokaryotes; Wiley-Blackwell
10. Osborn, A.M., Smith, Cindy (2005) Molecular Microbial Ecology; Taylor & Francis Group
11. Pierre Davet (2004) Microbial Ecology of the Soil and Plant Growth; Science Pub Inc
12. Ronald M. Atlas, Richard Bartha (1997) Microbial Ecology: Fundamentals and Applications (4th Edition); Benjamin Cummings
13. Satyanarayana, T., Johri, B.N. (2005) Microbial Diversity: Current Perspectives and Potential Applications; I.K. International Publishing House Pvt., Limited

MB1.6 Softcore: Practical II (Virology and Bacteriology)

1. Laboratory safety rules
2. Microscopic measurement of microorganisms by micrometry
3. Culturing and maintenance of bacterial cultures
4. Isolation and enumeration of bacteria from soil
5. Isolation and enumeration of bacteria from water
6. Cultural characteristics of bacteria
7. Staining techniques—simple (positive and negative), differential (Grams and acid fast), structural (endospore and capsule)
8. Motility test (hanging drop method and soft agar method)
9. Biochemical tests for the identification of bacteria—catalase, oxidase, IMViC, Urease, TSIA, Nitrate reduction, gelatin, starch, casein, esculin hydrolysis and Litmus Milk test.
10. Determination of growth curve in *E. coli*.
11. Diauxic growth curve in *E. coli*
12. Isolation of coliphages from sewage
13. Study of morphological changes due to viral infection in plants

MB1.7 Softcore: Practical III (Mycology and Microbial Genetics)

1. Isolation of slimemolds.
2. Isolation of aquatic fungi.
3. Isolation of soil fungi.
4. Isolation of fungi from air.
5. Isolation of fungi from cereals and cereal based products.
6. Study of the following representative genera: *Aspergillus*, *Penicillium*, *Fusarium*, *Neurospora*, *Saccharomyces*, *Erysiphae*, *Polyporus*, *Agaricus*, *Puccinia*, *Ustilago*, *Alternaria*, *Drechslera*, *Saprolegnia*, *Rhizopus*, *Trichoderma* and symbiotic fungi-Lichens.
7. Measurement of concentration of fungal conidia by Haemocytometer.
8. Measurement of fungal cells by Micrometer.
9. Replica plating technique for transfer of bacterial colonies.
10. Ultra-violet killing curve and determination of mutant types in *Saccharomyces cerevisiae*.
11. Induction of mutation
12. Isolation of streptomycin resistant strain of *E. coli* by gradient plate method.
13. Study of Conjugation in *E. coli*
14. Isolation of genomic DNA from bacteria by heat lysis method.
15. Isolation of genomic DNA from yeast by DNA spooning method.
16. Extraction of mycotoxins and detection by TLC.

SEMESTER II
MB2.1 Hardcore: MICROBIAL PHYSIOLOGY

Course Pedagogy:

- To develop understanding about microbial metabolism, growth and energy generation.
- Gain knowledge of various fermentation pathways, microbial communication and energetics.
- To acquire knowledge on microbial stress response.

Course Outcomes:

After the completion of the course students would be able.

- To acquaint with basics of metabolism and growth under normal and stressed conditions.
- To understand major fermentation, aerobic and anaerobic pathways for energy generation in microbial cells.
- To know the concepts of microbial cross-talk.

THEORY

48 hours

UNIT I

12 hours

Microbial bioenergetics: The role of ATP in metabolism. Microbial enzymes and mechanism of Enzyme action and kinetics. Inhibition and regulation – allosteric, feedback, competitive, non-competitive.

Metabolism of Carbohydrate: Glycolysis, Citric acid Cycle and different types of Phosphorylation, Fates of pyruvate, Fermentation. Utilization of sugars other than glucose: Lactose, Galactose, Maltose, Mannitol. Degradation of cellulose, Starch and Glycogen.

UNIT II

12 hours

Lipid metabolism: β -oxidation, Biosynthesis of fatty acids, degradation of fatty acids.

Nitrogen metabolism: Nitrogen metabolism, Biological nitrogen fixation process, symbiotic and non-symbiotic nitrogen fixation. Degradation and biosynthesis of essential and non-essential amino acids. **Nucleic acid metabolism:** Biosynthesis and degradation of purines and pyrimidines.

UNIT III

12 hours

Microbial Photosynthesis: Photosynthetic Pigments and apparatus in bacteria. Oxygenic and Anoxygenic. Photosynthesis. Autotrophic CO₂ fixation and mechanism of Photosynthesis. Utilization of light energy by Halobacteria.

Autotrophic Mechanisms in bacteria: Hydrogen bacteria, Nitrifying bacteria, Purple sulphur bacteria, Non-sulfur bacteria, Green sulfur bacteria, Iron bacteria, Methylo-trophs.

UNIT IV

12 hours

Microbial Signaling and Stress response: Two Component signal transduction in prokaryotes: Chemotaxis, Quorum sensing, biofilms, response to anti-microbials, sporulation inducing signals and events in sporulation; Dormancy, osmolarity porin regulation in *E. coli* (Omp system), phosphate assimilation in *E. coli* (Pho system), Nitrogen fixation in *Klebsiella* and *Rhizobium* (Ntr system). Oxidative stress, Thermal stress, Starvation stress, Aerobic to anaerobic transitions.

References:

1. Albert G. Moat, Michael P. Spector, John W. Foster (2009) *Microbial Physiology*; BWSM
2. Byung Hong Kim, Geoffrey Michael Gadd (2008) *Bacterial Physiology and Metabolism*; Cambridge University Press
3. Daniel R. Caldwell (1999) *Microbial Physiology and Metabolism*; Star Pub Co
4. David White, James Drummond, Clay Fuqua (2011) *The Physiology and Biochemistry of Prokaryotes*, Oxford University Press
5. Frederick C. Neidhardt, John L. Ingraham, Moselio Schaechter (1990) *Physiology of the Bacterial Cell: A Molecular Approach*; Sinauer Associates Inc
6. Robert K. Poole (2014) *Advances in Microbial Systems Biology, Volume 64 (Advances in Microbial Physiology)*; Academic Press
7. Rose, Anthony H. (.) *Advances in Microbial Physiology, Vol. 9*; Elsevier Science & Technology Book
8. Rose, Anthony H. (1976) *Chemical Microbiology: An Introduction to Microbial Physiology*; Basic Books

MB 2.2 HARDCORE: IMMUNOLOGY

Course Pedagogy:

- To provide overview of immune system, antigen, antibody structure and interactions.
- Understanding of innate and adaptive immunity along with major cells and molecules involved.
- To integrate immunology with health and enrich the knowledge for autoimmune disorders, hypersensitivity reaction.

Course Outcomes:

After the completion of the course students would be able

- To gain knowledge of immune system, cells involved along with complement system and autoimmunity.
- To evaluate the usefulness of immunology in different pharmaceutical companies
- To understand immune system, antigen antibody interactions.
- To gain theoretical knowledge of various diseased conditions generated due to interplay of immune system components

THEORY 48 hours

UNIT I

12 hours

Introduction to Immunology: An overview of immune system, Phagocytes, Natural killer cells, mast cells, basophils, Dendritic cells and other cells of the innate immune system. Immunity: Types- Innate immunity: (nonspecific) physical, biochemical and genetic factors involved in governing innate immunity, molecules of innate immunity – complement, acute phase proteins and interferons; Chemokines and Cytokines. Acquired immunity: (specific) natural, artificial, passive immunity, humoral or antibody mediated immunity, cell mediated immunity.

UNIT II

12 hours

Antigens and Antibodies:

Antigens: Properties of antigen, Super antigen, Hapten. **Major Histo-compatibility Complex (MHC) and Antigen presentation:** Types, Structure and functions of MHC molecules, Presentation of Bacterial and Viral Antigens: Phagocytosis, Processing and presentation of antigens by Class I and class II MHC molecules.

Antibodies (Immunoglobulins) – Structure and function. Ig gene organization and generation of Ab diversity. Monoclonal antibodies production and its clinical applications; Antibody engineering.

Hypersensitivity: Hypersensitivity reactions, Types and their roles in Immunopathological processes. Autoimmune processes: Immunologic tolerance, genetic predisposition to the development of autoimmune processes. Autoimmune disorders- Immunopathogenesis of sclerosis multiplex, psoriasis vulgaris, Rheumatoid arthritis), Immunodeficiency diseases

UNIT III

12 hours

Transplantation of tissues and organs: Nomenclature of transplantations. Recognition of self and non-self- Transplantation reactions, HvG and GvH. Exception from rejections. HLA Typing: Antibody dependent cell mediated cytotoxicity, mixed lymphocyte reactions. Kidney and bone marrow transplantations.

Immuno stimulatory and immune suppressive drugs and their mechanism.

Antigens and Antibody reactions: Agglutination, complement fixation test, ELISA, immunodiffusion, immunoelectrophoresis, immunofluorescence, immunoprecipitation, radioimmunoassay, Western blotting, flow-cytometry and immunohistochemistry.

UNIT IV

12 hours

Immune response to infectious diseases and Vaccines:

Viral Diseases: Neutralization of Viruses, Cell mediated immunity to control viral pathogens, Viruses can evade defense mechanisms. **Bacterial Diseases:** Immune response to extracellular and intracellular bacteria, bacteria can evade defense mechanisms, Immune response to Bacterial pathogenesis. **Parasitic Diseases:** Immune response to Malaria, Trypanosoma, Leishmaniasis. **Fungal Diseases:** Innate and Acquired Immunity to control fungal infections.

Vaccines – Definition, active and passive immunization, designing vaccines for active immunization. Live attenuated vaccines, Inactivated or killed vaccines, Subunit vaccines (Toxoids, Bacterial polysaccharide capsules, viral glycoproteins, Recombinant vaccines, multivalent subunit vaccines), DNA vaccines. Effectiveness of vaccines, Vaccine safety, current vaccines and National vaccination schedule.

References:

1. AbulK. Abbas (2014) Cellular and Molecular Immunology ; Saunders
2. AbulK. Abbas, Andrew H.H. Lichtman, Shiv Pillai (2011) Cellular and Molecular Immunology; Saunders
3. AbulK. Abbas, Andrew H.H. Lichtman, Shiv Pillai (2012) Basic Immunology: Functions and Disorders of the Immune System, ; Saunders
4. Delves, Peter J., Martin, Seamus J., Burton, Dennis R. (2011) Roitt's Essential Immunology; Wiley & Sons, Incorporated, John.
5. George Pinchuk (2001) Schaum's Outline of Immunology; McGraw-Hill
6. Helen Chapel, Mansel Haeney, Siraj Misbah, Neil Snowden (2014) Essential of Clinical Immunology; Wiley-Blackwell
7. Judy Owen, Jenni Punt, Sharon Stranford (2013) Kuby Immunology; W.H. Freeman
8. Louise Hawley, Benjamin Clarke, Richard J. Ziegler (2013) Microbiology and Immunology; LWW
9. Peter Parham (2009) The Immune System, 3rd Edition; Garland Science
10. William E. Paul (2012) Fundamental Immunology; LWW

MB2.3:Soft-core:FOODMICROBIOLOGY

Course Pedagogy:

- The course aims to provide instruction in the general principles of food microbiology.
- The course covers the biology and epidemiology of foodborne microorganisms of public health significance, including bacteria, yeasts, fungi, protozoa and viruses.
- Understand food spoilage microorganisms; the microbiology of food preservation and food commodities; fermented and microbial foods; principles and methods for the microbiological examination of foods; microbiological quality control, and quality schemes.
- To supplement the academic input of students by way of seminars, conferences, guest lectures and industry oriented projects/visits.

Course Outcome:

After the completion of the course students would be able

- To understand the principles of microorganisms during various food-processing and preservation steps.
- To comprehend the interactions between microorganisms and the food environment, and factors influencing their growth and survival.
- To understand the significance and activities of microorganisms in food.
- To recognize the characteristics of food-borne and spoilage microorganisms, and methods for their isolation, detection and identification.
- To analyze the importance of microbiological quality control programme's in food production.
- To describe the rationale for the use of standard methods and procedures for the microbiological analysis of food.

THEORY

48 hours

UNIT I

12 hours

Introduction to food microbiology: Definition, concepts and scope. Food as substrate for microbes. Factors influencing microbial growth in food – Extrinsic and intrinsic factors. Principles of food preservation – Chemical preservatives and Food additives. Asepsis – Removal of microorganisms, (anaerobic conditions, high temperatures, low temperatures, drying). Canning, processing for Heat treatment.

UNIT II

12 hours

Contamination and food spoilage: Cereals, sugar products, vegetables, fruits, meat and meat products, Fish and seafoods – poultry – spoilage of canned foods.

Dairy Microbiology: Microbiology of raw milk, Milk as a vehicle of pathogens, Prevention of contamination of raw milk, Microbiology of processed milk, Spoilage and defects fermented milk and milk products, Microbiological standards for milk and milk products. Cream and butter bacteriology. Prebiotics and Probiotics.

UNIT III

12 hours

Food poisoning and intoxication: Significance of food borne diseases, Food poisoning and intoxication: Botulism, Listeriosis, *Bacillus cereus* food poisoning, Foodborne Gastroenteritis by *Salmonella*, *Shigella*, *Vibrio*, *Campylobacter* and *Yersinia*, Staphylococcus and Staphylococcal enterotoxins, fungal spoilage and Mycotoxins. **Introduction to bio warfare:** Food and water as media to transmit food borne threats to health; policies and practices.

Microbes as alternate food – single cell proteins, sea weed (algae), mushroom cultivation. Bioconversions – production of alcohol-fermented beverages – beer and wine. Genetically modified foods.

UNITIV

12 hours

Detection of food-borne microorganisms: Culture, Microscopic and Sampling methods. Chemical:Thermostable nuclease *Limulus* Lysate for Endotoxins, Nucleic Acid (DNA) probes, DNA Amplification(PCR),Adenosine-TriphosphateMeasurement,Radiometry,Fluoro- andChromogenicsubstrates.ImmunologicMethods:FluorescentAntibody,EnrichmentSerology,Salmonella1-2.Test,Radioimmunoassay,ELISA.

Microbialindicatorsoffoodsafetyandqualitycontrol:Principlesofqualitycontrolandmicrobiologicalcriteria,Indicatorsofproductqualityandmicrobiologicalsafetyoffoods,Hazardanalysis,criticalcontrolpoints(HACCP),Goodmanufacturingprocess(GMP)MicrobiologicalstandardsCodexAlimentarius andFood legislationwithrespect toFSSAI,NABLandISO

References:

1. AdamsM.R.andMoss M.O.2007.Food Microbiology3rdEdition.Royal SocietyofChemistry.UK.
2. AhmedE. Y.andCarlstromC.2003FoodMicrobiology: ALaboratoryManual,JohnWileyandSons,Inc.New Jersey.
3. BibekRay,ArunBhunia.2013.FundamentalFoodMicrobiology,FifthEdition.CRCPress
4. CBlackburn.2006.FoodSpoilageMicroorganisms.WoodheadPublishing.
5. DongyouLiu.2009.MolecularDetection of FoodbornePathogens.CRC Press.
6. ElmerH.Marsh,James Steele.2001.Applied DairyMicrobiology,SecondEdition. CRCPress.
7. Frazier W.C. and Westhoff C.D. 2008 Food Microbiology. Tata McGraw Hill Publishing CompanyLimited,New Delhi.IndianEdition.
8. Jay,JamesM., Loessner, MartinJ., Golden,David A.2004.ModernFoodMicrobiology.7thed. Springer
9. Marshall,RichardJ.(Ed.).2007.FoodSafety. Springer.
10. Pina M. Fratafico, Arun K. Bhunia, and James L. Smith. 2008. Foodborne Pathogens: Microbiology andMolecularBiology. CaisterAcademicPress.
11. Pitt,JohnI.,Hocking,Ailsa D.2009.Fungiand Food Spoilage3rdEdition.Springer.
12. Sperber, William H., Doyle, Michael P. (Eds.). 2010. Compendium of the Microbiological Spoilage ofFoodsand Beverages. Springer.
13. StephenJ.Forsythe.2010.TheMicrobiologyofSafeFood,2ndEdition.Wiley-Blackwell.

MB2.4:Softcore:SOILMICROBIOLOGY

Course Pedagogy:

- Lectures are held with the help of slides, the laboratory lessons will be performed in a laboratory designed and equipped for microbiological practices.
- The laboratory practices will be performed in groups of students. The e-learning site will be used to provide teaching material and to communicate with the students.
- The interaction between teacher and student takes place through tutorials, seminars and Intermediate written tests.

Course Outcome:

After the completion of the course students would be able

- To have knowledge about soil as an excellent habitat for multitude of microorganisms balancing the soil ecosystem.
- To be employable in the field of Agronomy/Soil Science
- To acquire skills and knowledge on the importance of microorganisms in biogeochemical cycles biological fertility of soil.

THEORY

48 hours

UNIT I

12 hours

Soil Microbiology: Historical accounts and the “Golden Age” of soil microbiology and significant contributions of pioneer soil microbiologists.

Soil Microbial diversity: Soil as habitat for microbes; soil pH, temperature and soil atmosphere. Diversity and abundance of dominant soil microorganisms, Methods of isolation of soil microflora, soil organic matter decomposition,

UNIT II

12 hours

Biogeochemical cycles: Organic matter decomposition, humification. Carbon, sulphur, nitrogen and iron cycles in soil.

Soil microbe interaction - Antagonism, commensalism, mutualism, symbiosis, predators and parasite relationship and competition. Interaction of soil microflora with vascular plants - Rhizosphere, rhizoplane microorganisms, *Rhizobium*, *Azotobacter*, *Azospirillum*, *Cyanobacteria* and *Azolla*.

UNIT III

12 hours

Techniques to study soil organisms: Microbial biomass estimation; fumigation-incubation technique, fumigation-extraction method, substrate-induced respiration method and Using ATP or enzyme activity.

Applied soil microbiology: soil microbial inoculants, Manipulations of soil microorganisms for agriculture, Soil environmental contaminants and Bioremediation, Microbial products- Plant growth promoting Hormones, Antibiotics, Toxins and Enzymes

UNIT IV

12 hours

Soil-Borne Diseases and Human Health: *Clostridium tetani* (tetanus), Toxoplasmosis, Aspergillosis, Actinomycosis.

Soil microorganisms in agro ecosystems: Types of microbial communities; soil microbial diversity: significance and conservation; effect of agricultural practices on soil organisms. Biological nitrogen-

fixation: The range of nitrogen fixing organisms; mechanism of nitrogen fixation (biochemistry of nitrogenase);

genetics of nitrogen-fixation; *Rhizobium*-Legume Association; Sym plasmids, N₂ fixation by non-leguminous plants.

References:

1. Agrios, G.N. 2000. Plant pathology. Harcourt Asia Pvt. Ltd.
2. Bergersen, F.J. and Postgate, J.R. 1987. A Century of Nitrogen Fixation Research Present Status and Future Prospects. The Royal Soc., London.
3. Buchanan, B.B., Gruissem, W. and Jones, R.L. 2000. Biochemistry and Molecular Biology of Plants.
4. Burges, H.D. 1981. Microbial control of insect pests, Mites and plant diseases. Academic, London.
5. Dixon, R.O.D. and Wheeler, C.T. 1986. Nitrogen Fixation in plants. Blackie USA, Chapman and Hall, New York.
I.K. International Pvt. Ltd.
6. Kannaiyan, S. 1999. Bioresources Technology for sustainable agriculture. Assoc. Pub. Co. New Delhi.
7. Mehrotra, R.S. 2000. Plant pathology. Tata McGraw – Hill Publishing Company Limited.
9. Metcalf, R.L. and Luckmann, W.H. 1994. Introduction to insect pest management 3
edn. John Wiley and Sons, Inc.
10. Motsara, I.M.R., Bhattacharyya, P. and Srivastava, B. 1995. Biofertilizer Technology, Marketing and usage - A source Book-cum-glossary-FDCO, New Delhi.
11. Somasegaran, P. and Hoben, J.H. 1994. Handbook for Rhizobia; methods in legume *Rhizobium* Technology. Springer-Verlan, New York.

MB2.5 Softcore: PRACTICAL III (Microbial Physiology and Immunology)

1. Population growth of yeast – *S. cerevisiae*.
2. Population growth of bacteria – *E. coli*.
3. Sugar fermentation tests.
4. Catalase activity.
5. Hydrolytic acidity.
6. Casein hydrolysis.
7. Carbohydrate catabolism by microbes
8. Study of acid and pH stress tolerance by microbes.
9. Effect of temperature on microbial growth
10. Effect of molecular oxygen on microbial growth.
11. Effect of osmotic pressure on microbial growth.
12. Effect of relative humidity on microbial growth.
13. Effect of different wavelengths of light on microbial growth.
14. Precipitin test, ELISA, Ouchterlony Immunodiffusion test, and Radial Immunodiffusion
15. Determination of Blood grouping and rh factor
16. WIDAL Test.
17. VDRL Test (RPR).
18. HBsAg Test.
19. HCG test (Agglutination inhibition test).
20. Detection of RA factor.
21. CRP test.
22. ASO Test (Anti-streptolysin 'O' Test).

MB2.6 Softcore: PRACTICAL IV (FOOD MICROBIOLOGY)

1. Detection and enumeration of Microorganisms present in Utensils.
2. Isolation and identification of pathogenic microorganisms from canned food.
3. Enumeration of bacteria in raw and pasteurized milk by SPC method.
4. Determination of quality of a milk sample by Dye reduction test (MBRT, Rezasurin).
5. Detection of number of bacteria in milk by breed-count method
6. Microbial quality of milk and milk products.
7. Microbiological examination of Fermented products
8. Evaluation of antimicrobial activity of food preservatives
9. Isolation and identification of common food borne pathogens (Enterobacteriaceae – *E. coli*, *Enterobacter aerogenes*, *Salmonella*, *Shigella*, *Staphylococcus*, *Listeria*, *Vibrio*, *Aspergillus*, *Penicillium*, and *Fusarium*)
10. Bacterial examination of drinking water by membrane filter technique.
11. Determination of TDT.
12. Determination of TDP.
13. Detection and quantification of Aflatoxin B1.

MB2.7:OPEN ELECTIVE: MICROBIAL TECHNOLOGY

Course Pedagogy:

- The course will impart a comprehensive knowledge and understanding of techniques used in Microbiology, like microscopy, staining technique, culture media, sterilization methods and control of microorganisms.

Course Outcome:

After the completion of the course students would be able

- To acquire knowledge of culturing methods and identification of microorganisms.
- To enable them to isolate pure culture and preserve them and control measures.

THEORY

48 hours

UNIT I

12 hours

Microscopy: Light microscopy- Simple microscopy (dissection microscope), Compound microscopy (Bright field, Dark field, phase contrast, and Fluorescence microscopy) and stereomicroscopy. Electron-microscopy: Principles, construction and mode of operation of scanning and Transmission electron microscopy and limitations. Preparation of specimens for electron microscopic studies (Ultra-thin sectioning, negative staining, shadow casting and freeze etching). Confocal/ Laser scanning, programmable array microscopes

UNIT II

12 hours

Microbiological stains and staining techniques: Types of stains and principles of staining. Stains for bacteria, fungi, algae, protozoa and spirochetes. Stains for Azotobacter cysts, stains for mycoplasma. Preparation of bacterial smears for light microscopy: Fixation, simple staining, Differential staining, Structural staining (Capsule, Flagella, Cellwall and Endospore of bacteria), and nuclear staining.

UNIT III

12 hours

Culture media for Microbes Types of media- general purpose media, special purpose media, selective, elective, diagnostic, resuscitation media, Media for fungi, algae, bacteria, mycoplasma and viruses.

Sterilization techniques: Principles, types of Sterilization, and their mode of action. Physical methods: Heat-dry heat (Hot-Air oven), Incineration, Moist heat (Autoclave and Pressure cooker), Tyndalization (Fractional Sterilization), Filtration-Types of filters, Laminar airflow. Radiation methods (UV radiation, x-rays and cathoderays). Biosafety cabinets – Level I – IV, Containment labs – containment, high containment and maximum containment labs

UNIT IV

12 hours

Control of Microorganisms: Chemical methods: Definition of terms- Disinfectants, Antiseptics, Sanitizers, Microbicides (bactericide, fungicide and Sporicide), Microbistatic (bacteristatic and fungistatic agents). Use and mode of action of Alcohols, Aldehydes, Halogens, Phenols, Heavy metals, and Detergents.

Pure culture techniques: Different types of inoculation techniques - Spread plate, Pour plate and Streak plate method

References:

1. Alcom, I.E. 2001. Fundamentals of Microbiology. VI Edition, Jones and Bartlett Publishers, Sudbury, Massachusetts.
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9. Schaechter, M. Ingraham,J.L. and Neidhardt,F.C.2006. Microbe. ASM Press, Washington. D.C.
10. Stainer, R. Y., Ingraha, J L, Wheelis, M. L. and Painter, P. K. 1986. General Microbiology. McMillanEdun. Ltd. London.
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13. Talaro, K and Talaro, A.1996. Foundations in Microbiology, II edition, WCB publishers.
14. Tortora, G.J., Funke, B.R. and Case, C.L. 2004. Microbiology-An Introduction. Benjamin Cummings. San Francisco.

SEMESTER III
MB 3.1Hard-core: MOLECULAR BIOLOGY

Course Pedagogy:

- To extend the knowledge on structure and functions of genetic material
- To focus on genome organization, transcription and translation process in Prokaryotes.
- To understand the principles of oncogenes

Course Outcome:

After the completion of the course students would be able

- To have elaborate knowledge on nucleic acids
- To have better understanding of gene expressions
- To get thorough knowledge on Tumor viruses and oncogenes

THEORY

48hours

UNIT I

12 hours

Concepts in Molecular Biology: Microbes in molecular-biology.

Organization of Genomes: Prokaryotic genome- Genetic and Physical organization of bacterial genome, Eukaryotic genome–Genetic and Physical organization of nuclear genome

DNA structure and Replication: DNA as Genetic material, Chemistry of DNA, Modes of DNA Replication, Meselson and Stahl's Experiment, θ model, replication fork. Enzymes of DNA replication, preprimosome, primosome and replisome complex. Molecular mechanism of DNA replication, Differences in prokaryotic and eukaryotic DNA replication.

UNIT II

12 hours

DNA damage and recombination: Types of DNA damage - deamination, oxidative damage, alkylation and pyrimidine dimers; DNA repair – mismatch, short patch repair, nucleotide/base, excision repair, recombination repair and SOS repair. Recombination; Site specific recombination, Homologous recombination, transposition.

UNIT III

12 hours

Gene Expression: Structure of RNA- Classes of RNA, Chemistry of RNA.

Transcription: Transcription in prokaryotes and eukaryotes, Eukaryotic transcription factors. RNA processing, Ribozymes, Antisense RNA, mi RNA, Si RNA, RNAi and other small RNAs. Inhibitors of transcription and their mechanism of action.

Translation: Role of ribosome and different types of RNA in protein synthesis, deciphering the genetic code, basic feature of genetic code, mechanism of initiation, elongation and termination, Non ribosomal proteinsynthesis. Translational control and posttranslational events. Protein targeting, protein degradation, protein folding. Small peptides and therapeutic peptides.

UNIT IV

12 hours

Regulation of Gene expression: Regulation of gene expression in prokaryotes. Operon concept: lac, trp and arabinose. Regulation of gene expression in Eukaryotes. 2 component regulatory system (Sensor Kinases, response regulators, enhancers and silencers): constitutive, regulatory genes. Regulation of gene expression in bacteriophage. Gene silencing – gene regulation after transcription.

Recent trends in molecular biology research: Targeted genome editing: ZFNs, TALENs, CRISPRs gene editing, Knock-ins and Knock-outs. **Oncogenes**, proto-oncogenes, activation of proto-oncogenes

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2. Brown, T. A. 1991. Essential Molecular Biology. A Practical Approach Vol-I & Vol.-II, Oxford Univ. Press. Oxford.
3. Flint, S.J., Enquist, L.W., Drug, R.M., Racaniello, V.R. and Skalka, A.M. 2000. Principles of Virology-Molecular Biology, Pathogenesis and Control. ASM Press, Washington, D.C.
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5. Hartl, D.L. 1994. Genetics. Jones and Bartler Publishers, London.
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8. Molecular Cell Biology, W.H. Freeman and Company, New York.

MB 3.2 Hardcore: GENETIC ENGINEERING

Course Pedagogy:

- To learn about genetic engineering, principals involved in manipulating genes and DNA.
- To know about cloning strategies and expression systems.
- To acquire basic understanding of techniques in genetic engineering.
- To provide basic knowledge on intellectual property rights and their implications in biological research and product development

Course Outcome:

After the completion of the course students would be able

- To acquire knowledge on the concepts and terminology in genetic engineering.
- Familiar with various cloning strategies in prokaryotes.
- To have awareness of IPR ,the social and ethical issues concerning cloning by genetic engineering

THEORY

48hours

UNIT I

12 hours

Introduction to Genetic Engineering: Milestones in the development of genetic engineering. Genetic engineering as tool in biotechnology.Importance of gene cloning and future perspectives.

Tools in Genetic Engineering: Enzymes in genetic engineering. Cloning vectors; Plasmids (pUCseries,pBR 322), Phage vectors (M13, λ gt 10 and λ ZAP series), Ti vector. YAC, BAC vectors and specialist –purpose vectors; Expression vectors (pET vectors, pLITMUS). Synthetic construction of vectors.

UNITII

12 hours

rDNA Technology: The basic principles of gene cloning strategies: Preparation, Manipulation and Insertion of desired DNA in to vector. Introduction of DNA in to host cells– Transformation, Transduction, Transfection, Microinjection, Biolistics, Electroporation, Liposome fusion.Preparation andapplications of DNA libraries and cDNA libraries.Identification and Selection of recombinants.Applications of gene cloning in Biotechnology, Medicine, Agriculture, Forensic Science, Genetherapy.

UNIT III

12 hours

Analysis of gene and gene products: Molecular markers. DNA based and PCR - based markers, RFLP,RAPD, AFLP STS, EST, SSCP, VNTR, Microsatellites and mini-satellites. **DNA analysis:** labeling of DNA and RNA probes. Southern and fluorescence in situ hybridization, chromosome walking. PCR –types and applications.

Techniques for gene expression: Northern and Western blotting, Gel retardation technique, DNA foot printing, Primer extension, Reporter assays. DNA sequencing and sequence assembly. Sanger's methods, Next Generation Sequencing, techniques of Site-directed mutagenesis, Shot gun sequencing, chemical synthesis of oligonucleotides. Protein analysis; PAGE, 2D-GEL, **Protein sequencing**-N-terminal sequencing by Edman degradation method, C-terminal amino-acid analysis by carboxy peptidase digestion and Dansyl Chloride method.

UNIT IV

12 hours

Bioinformatics and Molecular Databases: Primary Databanks–NCBI, EMBL, DDBJ, KEGG; Secondary Databases–UNIPROT; Structural Database–PDB; Alignment: Pairwise and Multiple sequence alignment; Genome Annotation and Gene Prediction; Primer designing; Phylogenetic analysis and tree construction.

Safety of recombinant DNA technology: Restriction and regulation for the release of GMOs into Environment. Ethical, Legal, Social and Environmental Issues related to rDNA technology.

Introduction to IPR: Kinds of IPR; patents, copy right, design, trademark, geographical indicators,

Industrial design and trade secrets. India's new National IP Policy.

References:

1. Brown, T.A. (2010) Gene Cloning and DNA Analysis-An Introduction 6thedn. Blackwell Science.
2. Brown, T.A. (2011) Introduction to Genetics: A Molecular Approach 1stEd.
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4. Harvey Lodish, Arnold Berk, Chris A. Kaiser, Monty Krieger (2007) Molecular Cell Biology 6thEd. W.H. Freeman and Company, New York.
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12. J.F. Sambrook and D.W. Russell, ed. (2001), Molecular Cloning; A Laboratory Manual, 3rd ED, Vols 1, 2 & 3, Cold Spring Harbor Laboratory Press.

Course Pedagogy:

- To give knowledge on strain improvement methods
- To learn different fermentation techniques, bioreactor design, inoculum development.
- To understand techniques involved in down-stream fermentation process

Course Outcome:

After the completion of the course students would be able

- To get knowledge on strain improvement
- To understand methods of manipulating the metabolic pathways to get desired yield.
- To understand industrial production and purification of antibiotics, enzymes, amino-acids and steroids.
- To work in fermentation industry
- To understand the application of the bio-molecules in benefit to mankind

THEORY

48hours

UNIT I

12hours

Introduction: Fermenter design and types of fermenters, achievement and maintenance of aseptic conditions, Types of fermentation processes (Surface, submerged, Batch, Continuous, solid-substrate, Dual, Fed batch fermentation and its applications),

Industrial Microorganisms: Screening, Isolation. Identification and characterization of industrially important microbes. Strain improvement- mutation, recombination-gene regulation and genetic manipulation. Preservation of industrially important microbes. Culture collection centers.

UNIT II

12 hours

Media for Industrial Fermentations: Media formulation, growth factors, carbon, nitrogen, Energy and Mineral sources, buffers, inhibitors, precursors, inducers, Oxygen requirements Antifoam agents and others, Sterilization: Sterilization of bioreactor, media, air and exhaust air and filter sterilization. **Downstream processing and fermentation economics:** Steps in recovery and purification Methods of cell separation – filtration and centrifugation, cell disruption, liquid liquid extraction, chromatography, membrane processes. Fermentation economics- expenses for industrial organisms, strain improvement, media sterilization, heating, cooling, aeration and agitation. Cost of Plant and equipment, batch process cycle time, continuous culture, recovery and effluent treatment, cost recovery due to waste usages and recycling.

UNIT III

12 hours

Industrial production of energy fuels: Industrial alcohol production: Biosynthesis, methods of production, recovery and applications of ethanol: crab tree and pasture effect, acetone – butanol and glycerol through microbial process.

Industrial production of Organic acids and Enzymes: biosynthesis, media, production process, product recovery and application of citric acid and lactic acid, Enzymes: Fungal and Bacterial Amylase; Bacterial proteases.

UNIT IV

12 hours

Industrial production of food additives: amino acid production, methods of production, product recovery of L-Glutamic acid and L-lysine. Commercial uses of Amino acids Vitamins: Commercial production of Vitamin B12, and Riboflavin. Alcoholic beverages (Beer, Wine,)

Industrial production of health care product: Industrial production of β -lactum antibiotic (Penicillin): Biosynthesis, production and recovery. Streptomycin: Biosynthesis, production and recovery. Antitumor and anti-cholesterol agents, SCP and SCO, I P R: Patent Laws: Patent regulations of processes, products and microorganisms.

References:

1. Barsanti, LandGualtieri, P.2005. *Algae: Anatomy, Biochemistry, and Biotechnology*. Taylorand Francis NewYork.
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21. Reed, G. 1999. *Prescott and Dunn's Industrial Microbiology*. CBS Publishers and Distributors.

MB 3.4 Soft-core: MEDICAL MICROBIOLOGY

Course Pedagogy:

- To understand the role of normal flora and pathogenic microbes
- To understand the pathogenesis of various diseases
- To understand the various clinical microbiological techniques.

Course Outcome:

After the completion of the course students would be able

- To learn the concept, etiology and epidemiology of infections and mechanisms of infection
- To have knowledge on clinical lab techniques
- To acquire knowledge on control measures of diseases

THEORY

48hours

UNIT I

12 hours

Introduction to Medical Microbiology: History, Development and scope of Medical Microbiology. Concept of Disease, disorder, syndrome, Communicable diseases- Microbial infections and diseases. Factors responsible for microbial pathogenicity.

Microbial infections: Types of infections, modes of transmission, portal of entry: Urinary tract infection, sexually transmissible infection, Infection of the central nervous system, Infections of circulatory system, Oral cavity and respiratory infection, gastrointestinal infection.

UNIT II

12 hours

Nosocomial infection: Incidence of nosocomial infections, types of nosocomial infections, emergence of antibiotic resistant microorganisms, hospital infection control programs, preventing nosocomial infections and surveillance, General concepts for specimen collection and handling of specimen, specimen processing and biosafety.

Chemotherapeutic agents: antibiotics (Classification based on chemical structure, mode of action and range of effectiveness). Recent trends-Drug resistance and its consequences, antibiotic policy, NCCLS (CLSI) guidelines and standards, WHO guidelines. MDR strains.

UNIT III

12 hours

Epidemiology, Pathogenesis, Spectrum of disease, Laboratory diagnosis and Prevention: Diseases caused by **Viruses:** Chicken pox, Rabies virus, hepatitis, encephalitis, AIDS, Herpes simplex infections, Influenza, Dengue

Diseases caused by Bacteria: Tuberculosis, Leprosy, cholera, Typhoid, Botulism, Shigellosis, Helicobacter pylori infection, Salmonellosis, Tetanus. Diseases caused by **Fungi:** Candidiasis, Histoplasmosis, Blastomycosis, Coccidiomycosis, Dermatormycosis, Aspergillosis and Cryptococcosis, Anthrax

UNIT IV

12 hours

Diseases caused by Mycoplasma: *Mycoplasma pneumoniae*, *M. urealyticum*, *M. hominis*.

Diseases caused by Protozoa: Giardiasis, Trichomoniasis, Cerebral Malaria, Toxoplasmosis, Cryptosporidium.

Disease caused by Chlamydiae: Psittacosis, Lymphogranuloma Venereum, Trachoma and Inclusion conjunctivitis.

Emergent Diseases: Hemorrhagic fever, Swine flu, SARS, Chikungunya, Ebola, Hanta, Leptospirosis, Marburg

References:

1. Robert W. Bauman Ph.D. (2011) Microbiology with Diseases by Body System (3rd Edition); Benjamin Cummings
2. Patrick R. Murray PhD, Ken S. Rosenthal PhD, Michael A. Pfaller MD (2012) Medical Microbiology; Saunders
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9. Ananthanarayan, Paniker (2009) Textbook of Microbiology, 8th Edition; University Press
10. Jawetz (2010) Medical Microbiology, 25th Edition; Tata McGraw –Hill Education

MB 3.5 Soft-core: CLINICAL & DIAGNOSTIC MICROBIOLOGY

Course Outcome:

After the completion of the course students would be able

- To develop skill to isolate and identify microorganism from clinical sample.
- To do antibiotics sensitivity and resistance test
- To do detection of parasite/pathogens using diagnostic kits.

Course Pedagogy:

- Knowledge about microbes causing disease.
- Knowledge about various laboratory techniques like microscopy, immunological assessments, radiology, biomarker tests, ELISA, serology checks, vaccines and vaccines schedule.
- Many microbes have developed resistance to medications.

THEORY

48hours

UNIT I

12 hours

Introduction to clinical Microbiology: Role of Microbiologist in Diagnostic laboratory, General concepts for specimen collection, handling, transportation, processing, specimen workup, Laboratory safety and infection control.

Scientific and Laboratory basis for Clinical/Diagnostic Microbiology: Microscopic examination of infectious diseases, Growth and biochemical characteristics, Rapid methods of identification.

UNIT II

12hours

Immunotechniques and Immunodiagnosis: Antigens and Antibody reactions *in vitro*; Agglutination, complement fixation, ELISA, Western Blotting Immuno-diffusion, Immuno-electrophoresis, Immunofluorescence, Immuno-precipitation, Radioimmuno assay and serotyping.

Vaccines and Vaccination: Vaccines – definition, types, Antigens used as Vaccines, effectiveness of vaccines, Vaccine safety, current vaccines, adjuvants, active immunization and passive immunization.

UNIT III

12 hours

Recent Diagnostic tools and techniques: Principle, working and application of a) Autoanalyser b) Biosensor glucometer /labon chip/microfluidics c) Diagnostic kits- ELISA, Western Blot d) Enzymes in Disease diagnosis and therapy: Lactate dehydrogenase, Aspartate aminotransferase, Alkaline phosphatase, Creatine kinase, Acid phosphatase, Cholinesterase.

UNIT IV

12 hours

Antimicrobial Chemotherapy: Development of chemotherapy; General characteristics of drugs and their testing; Mechanism of action. Antibacterial drugs; antifungal drugs, antiviral and antiprotozoan drugs; antibiotic sensitivity testing, MIC, Drug resistance; mechanism of drug resistance; multi drug resistance

Reference

- 1 Alberto M. Marchevsky and Mark Wick. (2011). Evidence Based Pathology and Laboratory Medicine. Springer publication.
- 2 David E. Bruns; Edward R. Ashwood; Carl A. Burtis; Barbara G. Sawyer (2007). Fundamentals of Molecular Diagnostics St. Louis, Mo. : Saunders Elsevier
- 3 Goura Kudesia (2009) Clinical and Diagnostic Virology. Cambridge University Press. UK.
- 4 Henrik Winther and Jan T. Jorgensen (2010). Molecular Diagnostics. Springer publications.
- 5 Huggett and Justin O'Grady *LGC (2014) Molecular Diagnostics: Current Research and Applications*. Caister Academic Press.
- 6 Huw Llewelyn, Hock Aun Ang, Keir E Lewis and Anees Al-Abdullah (2009). Oxford Handbook of Clinical Diagnosis. Oxford publications.
- 7 J. Andre Knottnerus and Frank Buntinx (2008) The Evidence Base of Clinical Diagnosis: Theory and Methods of Diagnostic Research, 2nd Edition. Wiley Publication.
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- 10 Richard A. McPherson and Matthew R. Pincus (2011). Henry's clinical diagnosis and management by laboratory methods. (22nd Ed) Philadelphia, PA : Elsevier/Saunders,
- 11 Stephen B. Hulley; Steven R. Cummings; Warren S. Browner; Deborah G. Grady; Thomas B. Newman (2007) Designing clinical research (3rd edition). Philadelphia, PA: Lippincott Williams & Wilkins.
- 12 Vinay Kumar et al., (2010) Robbins and Cotran pathologic basis of disease. Philadelphia, PA: Saunders/Elsevier.

MB3.6 Soft-core: PRACTICALV (Molecular Biology and Genetic engineering)

1. Isolation of Genomic DNA from *E.coli*.
2. Isolation of plasmids from bacteria by agarose gel electrophoresis.
3. Determination of purity and concentration of isolated DNA using spectrophotometer
4. Estimation of DNA by DPA method
5. Determination of DNase activity on isolated DNA
6. Amplification, Purification and separation of PCR product.
7. Estimation of RNA by Orcinol Method
8. Determination of RNase activity on isolated RNA.
9. Salt fractionation of Yeast protein and quantification
10. Estimation of protein by Lowry's method
11. Determination of Proteinase activity on proteins
12. Separation of proteins by SDS PAGE
13. Digestion of the gene of interest with suitable restriction enzymes.
14. Ligation of the digested gene in a vector.
15. Preparation of competent *E.coli* cells for Bacterial transformation.
16. Transformation of the vector in to the host cell and selection of the desired clones.
17. Induction of gene expression and purification of the induced protein from the host.

MB 3 .7 Soft-core: PRACTICALVI (Industrial and Medical Microbiology)

1. Study design of Fermentor and Parameters
2. Isolation of antibiotic producing microbes and their preservation.
3. Antibiotic fermentation and estimation of penicillin
4. Batch fermentation of Citric acid production, recovery and estimation of citric acid.
5. Preparation of wine and estimation of alcohol by specific gravity method.
6. Alcoholic fermentation and determination of total acidity and non-reducing sugars
7. Clarification of banana juice using Pectinase
8. Mushroom cultivation
9. Isolation of Pathogenic fungi of the skin (Dermatophytes).
10. Isolation and identification of clinically important microbes from throat swab
11. Isolation and identification of clinically important microbes from nasal swab
12. Isolation and identification of clinically important microbes from wound infections
13. Microbial flora of mouth-teeth crevices and saliva.
14. Estimation of bacteria in urine by calibrated loop direct streak method.
15. Antimicrobial assay-sensitivity test (MIC) for pathogenic bacteria.
16. Demonstration of laboratory diagnosis of important human diseases:-Tuberculosis, Typhoid, Malaria, and Hepatitis.

MB 3.8 Open elective: MICROBIAL DIVERSITY

Course Pedagogy:

- To understand the ubiquitous nature and characteristics of microbes
- To impart knowledge on viral, bacterial, fungal diversity.
- Importance and conservation of microbial diversity.

Course Outcome:

After the completion of the course students would be able

- To differentiate various groups of Microbes.
- To learn about conservation methods.
- To have knowledge about the role of culture collection centers in conservation.

THEORY

48 hours

UNIT I

12 hours

Viral Diversity: Morphology, ultra structure, chemical composition of virus, classification of viruses, Group I – T2 Bacteriophage, Group II – Banana bunchy top virus, Group III – Reovirus, Group IV-TMV, Group V– Rhabdovirus, Group VI– HIV, Group VII–Hepatitis virus.

Sub-viral particles: Discovery, Structure, Classification, replication and diseases caused by Satellite, Satellites virus, Virusoids, Viroids and Prions.

UNIT II

12 hours

Bacterial Diversity: Archaeobacteria, Photosynthetic Eubacteria, Chemoautotrophic and Methophilic Eubacteria, Gliding Eubacteria, Spirochetes, Rickettsiae and Chlamydiae, Actinomycetes, Mollicutes, Protists. Classification based on Bergey's manual (Determinative & Systematic).

UNIT III

12 hours

Fungal Diversity: Classification, Distribution, Importance, Structure, reproduction and general characteristics of the fungal divisions: Zygomycota (*Rhizopus*), Ascomycota (*Neurospora*), Basidiomycota (*Agaricus*), Deuteromycota (*Penicillium*), Chytridiomycota (*Allomyces*), Myxomycota and Yeast.

UNIT IV

12 hours

Importance and Conservation of Microbial Diversity: Importance of microbial diversity in agriculture, forestry, environment, industrial & food biotechnology, animal & human health. Metagenomics. Importance of conservation. *In situ* conservation and *Ex situ* conservation. Role of culture collection centers in conservation.

References

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12. Wagner, E.K. and Hewlett, M.J. 1999. Basic Virology. Blackwell Science. Inc.

SEMESTER IV
MB4.1 Hardcore: AGRICULTURAL MICROBIOLOGY

Course Pedagogy:

- To study the microbes associated with the plant and soil fertility.
- To understand about beneficial microbes and their uses in protecting agriculture, preserving food, enhancing the value of food products and providing general benefits to health and wellbeing.
- To classify various aspects of N₂ fixation, P solubilization, PGPR, are easily grasped by students
- To understand microbe and plant interactions
- Enable them to understand plant disease, plant defense mechanism and disease management.

Course Outcome:

After the completion of the course students would be able

- To develop newer approaches for plant disease management.
- Have better knowledge of pathogen interactions and plant defense mechanisms
- To know the application of microbial bio-control agents and to reduce drug resistance and environmental pollution.

THEORY

48hours

UNIT I

12 hours

Introduction to Agricultural Microbiology: Introduction to agricultural microbiology, concepts and scope of agricultural microbiology, Agronomy and production of important crop plants, Green revolution. Plant Pathology: Concept of disease, History of Plant Pathology, Significance of plant diseases, Symptoms and types of plant diseases.

Plant Pathology in Practice: Plant Clinic and Plant Doctor Concept. Diagnosis of Plant Diseases – Infectious diseases, Non-infectious diseases, Koch's rules;

UNIT II

12 hours

Parasitism and Disease Development Parasitism and pathogenicity, Host range of pathogens, Disease triangle, Diseases cycle / Infection cycle, Relationship between disease cycles and epidemics; Pathogens Attack Plants–Mechanical forces, Microbial enzymes and toxins, Growth regulators. Effect on physiology of Host–Photosynthesis, Translocation and transpiration, Respiration, Permeability, Transcription and translation. Environment and Plant Disease– Effect of Temperature, Moisture, Wind, Light, Soil, pH and structure, Nutrition and Herbicides.

Defense Mechanisms of Plant: Disease Pre-existing structural and chemical defenses, Induced structural and biochemical defenses. Microbe mediated strategies for abiotic stress management.

UNIT III

12 hours

Plant Disease & their management: Tobacco Mosaic Disease, Sandal Spike Disease, Bacterial blight of Paddy, Citrus canker, Angular leaf spot of cotton, Late Blight of Potato, Downy Mildew of Bajra, Blast of paddy, Tikka disease of ground nut, Rust of coffee, Grain and Head smut of Sorghum. Powdery mildew of Cucurbits, Wilt of Tomato, and Root Knot of Mulberry. Bunchy top of Banana.

UNIT IV

12 hours

Microbes and Plant interaction- Mycorrhizae- Biology and their applications, Bio fertilizers – microbial inoculants. Production and application of *Rhizobium*, *Azospirillum*, *Azotobacter*, phosphor bacteria and Cyanobacteria. PGPR's plant growth promoting *Rhizobacteria* and their uses.

Biopesticides: Definition, types- bacterial, viral, fungal and protozoan, mode of action, target pests, use of transgenic plants. Mode of action, Bacteria- endo and ecto- toxins production by *Bacillus thuringiensis*, and *Pseudomonas*. Fungi- *Beauveria*, *Cephalosporium*, and *Trichoderma*.

References:

1. George.N. Agrios(2005), Plant pathology, Elsevier academic press, 5thedition,U.K.
2. Mehrotra. R.S. and Ashok Aggarwal (2002), Plant pathology, Tata MC Graw-Hill publishers, 2ndedition, Delhi.
3. Kannaiyan.S. (2002), Biotechnology of Biofertilizers, Alpha science international,1stedition.
4. Bagyaraj D.G. and Rangaswami. G. (2005). Agricultural Microbiology, Prentice- Hall of India, 2ndedition, New Delhi.
5. NeelimaRajvaidya and Dilip Kumar Markandey. (2006). Agricultural Applications of Microbiology, NangiaS.B. And A.P.H.publishing corporation, New Delhi.
6. Oerke,E.C.Dehne,H.C. Schönbeck, F.Weber,A.(1999).Crop Production and Crop Protection, Elsevier academicpress, 5th edition,U.K.
8. RogerHull (2013). Plantvirology, Elsevier academic press,11th edition, U.K.
9. Hermann H. Prell, Peter R. Day. (2001). Plant-Fungal Pathogen Interaction: A Classical and MolecularView,1st edition, Springer-Verlag Berlin Heidelberg, Germany.
10. Geoffrey Clough Ainsworth (1981). Introduction to the History of Plant Pathology 1st edition, Cambridge university press, U.K.
11. Vidhyasekaran,P.(2007).Fungal Pathogenesis in Plants and Crops: Molecular Biology and Host Defense Mechanisms, 2nd edition, APS press, U.S.A

MB 4.2 Soft-core: ENVIRONMENTAL MICROBIOLOGY

Course Pedagogy:

- To give basic idea on environmental sample analysis; Topics covered in detail include soil microbiology, aquatic microbiology, aero microbiology, bio-fertilizers and pesticides, microbial waste recycling and bioremediation etc.
- To understand the basic principles involved in waste water management
- To get the information on usage of Bioremediation- biotechnology
- To inform students about Bio-oxidation & microbial leaching

Course Outcome:

After the completion of the course students would be able

- To apply advanced knowledge on environmental sample analysis
- To use the knowledge for better waste management
- To formulate technique for bioremediation process
- To apply principles of environmental microbiology to solve the current environmental issues
- To be employable in pollution control boards

THEORY

48 hours

UNIT I

12 hours

Environment and Ecosystem: Physical, chemical and biological aspects of environment, natural habitats of microorganisms, microorganisms in ecosystem as producers and decomposers.

Soil Microbiology: Characteristics and classification of soil. Interactions between microorganisms: Mutualism, commensalism, ammensalism synergism, parasitism, predation, competition. Rhizosphere, rhizosphere, microflora and its beneficial activity. Role of microorganism in nitrogen, phosphorous and sulphur cycle. Detrimental effects of diverted biogeochemical cycles. Biological nitrogen fixation in detail: Symbiotic, asymbiotic and associated nitrogen fixation. Structure, function and genetic regulation of nitrogenases.

UNIT II

12 hours

Air Microbiology: Microorganisms in air, sources of air-borne microorganisms. Airspora of indoor and outdoor environment, factors affecting airspora, Techniques of trapping air borne microorganisms. Brief account of air-borne diseases of humans, plants and their significance.

Aquatic Microbiology: Distribution of microorganisms in the aquatic environment, Water pollution sources, Biological indicators of water pollution. Eutrophication- role of nitrogen and phosphorus in eutrophication, process and control of eutrophication. Determination of sanitary quality of water, Waste water microbiology- Primary, secondary, tertiary treatment and reclamation of wastewater.

UNIT III

12 hours

Culture- dependent and independent approaches for microbial diversity in environment.

Culture- dependent approaches and their limitations, and culture-independent molecular approaches for understanding microbial diversity in the environment. Viable but non-culturable bacteria. Introduction to Metagenomics.

Microbes in extreme environment: acidophiles, alkaliphiles, halophiles, barophiles and their survival mechanisms.

Space microbiology: Historical development of space microbiology, Life detection methods a) Evidence of metabolism (Gulliver) b) Evidence of photosynthesis (autotrophic and heterotrophic).

Microbes in the degradation of wastes: Treatment of solid and liquid industrial wastes, Microbial degradation of pesticides, Xenobiotics, degradation of lignin, cellulose, pectin and plastic. Bio-remediation. Geo microbiology: Microbes in metal extraction, mineral leaching and mining, copper extraction by leaching and microbes in petroleum product formation. Global Environmental Problems: Global Warming, Acid rain, Ozone depletion. Bio- deterioration of wood and metals.

References:

1. Microbial Ecology by R.M. Atlas, R. Bartha. 3rd edition. Benjamin Cummings Publishing Co, USA. 1993.
2. Environmental Microbiology by A.H. Varnam, M.G. Evans. Manson Publishing Ltd. 2000.
3. Manual of Environmental Microbiology edited by C.J. Hurst, R.L. Crawford, J.L. Garland,
4. D.A. Lipson, A. L. Mills, L.D. Stetzenbach. 3rd edition. Blackwell Publishing. 2007.
5. Environmental Microbiology edited by R. Mitchell, J-D Gu. 2nd edition. Wiley-Blackwell. 2009.
6. Environmental Microbiology by R. Maier, I. Pepper, C. Gerba. 2nd edition. Academic Press. 2009.
7. Environmental Microbiology: Principles and Applications by P.K. J jemba, Science Publishing Inc. 2004.
8. Lignocellulose Biotechnology: Future Prospects by R.C. Kuhad, A. Singh. I.K. International. 2007.
9. Environmental Microbiology of Aquatic & Waste systems by N. Okafor. 1st edition, Springer, New York. 2011
10. Bioremediation by Baker K.H. and Herson D.S. 1994.. Mac Graw Hill Inc. N.Y.
11. Waste Water Engineering - Treatment, Disposal and Re-use by Metcalf and Eddy, Inc., Tata MacGraw Hill, New Delhi.
12. Pollution: Ecology and Biotreatment by EcEldowney, S. Hardman D.J. and Waite S. 1993. - Longman Scientific Technical.
13. Environmental Microbiology edited by Ralph Mitchell. A John Wiley and Sons. Inc.
14. Waste Water Microbiology 2nd Edition by Bitton.
15. Chemistry and Ecotoxicology of pollution. Edited by Des. W. Connell, G.J. Miller. Wiley Interscience Publications.
16. Environmental Biotechnology. Edited by C. F. Forster and D.A., John Wase. Ellis Horwood Ltd. Publication.
17. Advances in Waste Water Treatment Technologies. 1998. Volumes II and I by R. K. Trivedy. Global Science Publication.
18. Biocatalysis and Biodegradation: Microbial transformation of organic compounds. 2000 by Lawrence P. Wacekett, C. Douglas Hershberger. ASM Publications.
19. A Manual of Environmental Microbiology. 2nd Edition. 2001 by Christon J. Hurst (Chief Editor), ASM Publications.
20. Biodegradation and Bioremediation, Academic Press, San Diego.
21. Biotechnology in the sustainable environment, Plenum Press, N.Y.
22. Basic Principles of Geomicrobiology by A. D. Agate, Pune.

MB 4.3 Soft-core: GENOMICS AND PROTEOMICS

Course Pedagogy:

- The objectives of this course are to provide introductory knowledge concerning genomics, proteomics and their application
- To have knowledge about bioinformatics using web based tools (NCBI, CLUSTALW, MSA etc.)

Course Outcome:

After the completion of the course students would be able

- To acquire knowledge and understanding of the fundamentals of genomics and proteomics, transcriptomics and their applications in various applied areas of biology.
- Do Insilco analysis using web based tools, will help the students in their research

THEORY

32 hours

UNIT I

8 hours

Genome - Overview of Genome; Sequence of Genome Acquisition and Analysis - Homologies - Snps – Genetic Analysis, Linkage Mapping,

High Resolution Chromosome Mapping And Analysis - Physical Mapping, Yac, Hybrid Mapping, Strategies, Sequence Specific Tags (Sst), Sequence Tagged Sites (Sts), Ish, Fish, Rflp, Rapd.

UNIT II

8 hours

DNA Sequencing- Methods, Maxam and Gilbert Method, Ladder, Fluorescent, Shot Gun, Mass Spectrometry, Automation Sequencing – Find Gene Mutations, Implications of DNA – Sequencing and Sequencing Genomes.

Genome Data Bank, Metabolic Pathway Data - Construction And Screening of cDNA, Libraries And Microarrays - Application Of DNA Arrays - PCR - Variations In PCR - Gene Disruptions – Sage And Sade, Pharmacogenomics.

UNIT III

8 hours

Protein Sequence Analysis - Introduction - Sequence Data Banks - Wbrf – Pir - Swissport - Databases, Data Mining – Algorithms Of Proteomics And Its Applications- Protein Expression **Profiling** - Protein- Protein Interaction - Protein Modifications. Automation-Nucleic Acid Data Bank –EMBL Nucleotide Sequence Data Bank- Aids Virus Sequence Data Bank-RNA Data Bank.

UNIT IV

8 hours

Tools For Data Bank - Pairwise Alignment - Needleman And Wunsch Algorithm – Smith Waterman - Multiple Alignment- Clustral - Pras - Blast - Fast, Algorithms To Analyse Sequence Data- Pdb, Cambridge Structure Data Base (Lsd), 2d Electrophoresis, Ief, Hplc, Protein Digestion Technique, Mass Spectrometry, Maldi-Tof, Peptides, Mass Finger Printing Protein.

Metabolomics: Introduction, importance of metabolomics, designing of metabolomic study. Data base for repository of metabolites, CHEBI, EMBL, EBI, reactome data base.

References

1. DevarajanThangadur and JeybalanSangeetha(2015). Genomics and Proteomics Principles, Technologies, and Applications. Apple Academic Press.
2. FerencDarvas, AndrásGuttman, GyörgyDormán(2013). Chemical Genomics and Proteomics (2nd Ed). CRC Press.
3. Ganapathy Subramanian and Nawin Mishra (2012). Science of Proteomics: Historical Perspectives and Possible Role in Human Healthcare. WileyPublications.UK
4. Lynn Jorde, Peter Little, Mike Dunn and Shankar Subramaniam (2014). Encyclopedia of Genetics, Genomics, Proteomics and Bioinformatics. WileyPublication.UK
5. Malcolm Campbell, Laurie J. Heyer (2003). Discovering genomics, proteomics and bioinformatics. Benjamin Cummings publications.
6. MetinAkay (2007). Genomics and Proteomics Engineering in Medicine and Biology. WileyPublications.UK.
7. NachimuthuSaraswathy and PonnusamyRamalingam (2011). Concepts and Techniques in Genomicsand Proteomics .Wood head Publishing groups.
8. Nawin Mishra (2010).Applications of Proteomics: Proteomics, Human Disease, and Medicine. Wileypublication.UK
9. R. S. Dassanayake, Y. I. N. Silva Gunawardene (2011). Genomic and Proteomic Techniques: In Post Genomics Era. Narosa Book Distributors.
10. Ruchi Singh (2014). BIOINFORMATICS: GENOMICS AND PROTEOMICS. Vikas Publications. New Delhi.
11. Suhai, Sándor (2002). Genomics and Proteomics. Springer publications.

MB 4.4 Soft core: Microbial Nanotechnology

Theory

48 hours

Unit I

12 hours

Introduction to Nanotechnology: Characteristic scale for quantum phenomena, nanoparticles, nano-clusters, nanocomposite, nanotubes, nanowires and Characterization of nanoparticles – UV-VIS IR spectroscopy, TEM, SEM, AFM, EDS, XRD. Emergence of bionanotechnology.

Unit II

12 hours

Microbial nanotechnology – Microbial synthesis of Nanoparticles. Synthesis of nanodrugs – metal nanoparticles and drug delivery vehicles – Nanoshells – Tectodentrimers Nanoparticle drug systems – Diagnostic applications of nanotechnology.

Unit III

12 hours

Preparation of nano biomaterials – Polymeric scaffolds collagen, Elastins: Mucopolysaccharides, proteoglycans, cellulose and derivatives, Dextrans, Alginates, Pectins and Chitin. Nanoparticles – types (Silver, Gold and Titanium). Physical and chemical properties and functions. Drug delivery – protein mediated and nanoparticle mediated. Hybrid conjugates of gold nanoparticles – DNA oligomers in nano mechanics and Computing. Nanoparticles as carrier for genetic material.

Unit IV

12 hours

Applications in biology and medicine: Nanotechnologies for biology and medicine – Micro and nano- fluidics - Scanning probe microscopy in biology and medicine – Self-assembly of biological molecules.

Health and safety implications: Health issues – Environmental issues – regulation guidelines. Societal implications- Possible military applications – Potential benefits and risks to developing countries – Intellectual property issues – Criticism of Nanotechnology.

Reference Books:

1. Parthasarathy, B.K. (2007). Introduction to Nanotechnology, Isha Publication.
2. Elisabeth Papazoglou and Aravind Parthasarathy (2007). Bionanotechnology. Morgan & Claypool Publishers.
3. Bernd Rehm (2006). Microbial Bionanotechnology: Biological Self-assembly Systems and Biopolymer-based Nanostructures. Horizon Scientific Press.
4. David E. Reisner, Joseph D. Bronzino (2008). Bionanotechnology: Glo
5. Chaudhery Hussain (2022) Handbook of Microbial Nanotechnology, 1st Edition.

MB4.6 Soft-core: PRACTICAL VII (Agricultural Microbiology)

1. Isolation of Rhizobium from roots of leguminous plants
2. Assay of bio-fertilizers (*Rhizobium*, *Azotobacter*, *Azospirillum*) (Seed treatment/ seedling inoculation and measurement of root and shoot length)
3. Mass multiplication techniques of *Azolla*.
4. Estimation of total phenols in diseased and healthy plant tissues.
5. Isolation of phosphate solubilizing bacteria and fungi.
6. Isolation of phylloplane microorganisms
7. Soil microbes interaction In vitro by dual culture method
8. Isolation, identification and enumeration of Rhizosphere and Rhizoplane microorganism
9. Seed health testing by SBM
10. Collection and Identification of following disease:
 1. Tobacco mosaic disease
 2. Bunchy top of Banana
 3. Bean Mosaic
 4. Sandal spike
 5. Bacterial blight of paddy
 6. Citrus canker
 7. Downy mildew of Bajra
 8. Powdery mildew of mulberry
 9. Head smut of sorghum
 10. Leaf rust of coffee
 11. Blast disease of paddy
 12. Tikka disease of groundnut
 13. Leaf spot of paddy
 14. Grassy shoot of sugarcane

MB 4.7 Soft-core: PRACTICAL VIII (Environmental Microbiology Practical)

1. Microbes as indicators of water pollution– Determination of indices of water quality.
2. Determination of TS from Sewage Water
3. Determination of BOD of pollution water.
4. Determination of COD of polluted water.
5. Degradation of cellulose by *Chaetomium globosum*.
6. Study of Actinorhiza, Mycorrhiza.
7. Isolation of pesticide degrading microbes
8. Isolation of plastic degrading microbes

MB 4.8 Soft-core: PRACTICAL IX: Microbial Nanotechnology Practical

1. Synthesis of gold Nano particles from bacteria and its confirmation by UV-Spectroscopy
2. Synthesis of gold Nano particles from fungi and its confirmation by UV-Spectroscopy
3. Synthesis of silver Nano particles from bacteria and its confirmation by UV-Spectroscopy
4. Synthesis of silver Nano particles from fungi and its confirmation by UV-Spectroscopy
5. Synthesis of Nano metal particles from microbes and its confirmation by UV-Spectroscopy
6. Evaluation of antimicrobial properties of gold Nano particles produced by microbes
7. Evaluation of antimicrobial properties of silver Nano particles produced by microbes
8. Determination of MIC values of synthesized nanoparticles
9. Evaluation of plant growth promotion of Nano metals/composites.
10. Determination of size of nanoparticles by SEM/TEM
11. Characterization of nanoparticles by using XRD, EDX and DLS