

CVM UNIVERSITY

MASTER OF SCIENCE

(MICROBIOLOGY)

PROGRAMME

Under Choice Based Credit Scheme

Structure with Effect From: 2020-21



M.Sc. Microbiology Programme Details

Programme Objectives (POs):

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

Programme Specific Outcomes (PSOs):

At the end of the two year programme the student will understand and be able to explain different branches of Microbiology. The student will be able to explain about various applications of Microbiology such as Environmental Microbiology, Industrial Microbiology, Food Microbiology, and Medical Microbiology. He/she will be able to design and execute experiments related to Basic Microbiology, Immunology, Molecular Biology, Recombinant DNA Technology, Microbial Genetics, and Bioinformatics tools will be able to execute a short research project incorporating techniques of Basic and Advanced Microbiology under supervision. The student will be equipped to take up a suitable position in academia or industry, and to pursue a career in research if so desired.

Programme Structure:

The M.Sc. Microbiology programme is a two-year course divided into four-semester. A student is required to complete hundred credits for the completion of course and the award of degree. A student has to accumulate twenty-five credits in each of the four semesters.

PART ONE	FIRST YEAR	SEMESTER I	SEMESTER II
PART TWO	SECOND YEAR	SEMESTER III	SEMESTER IV

Course Credit Scheme

Semester I

Course Type	Course Code	Name of Course	T/ P	Credits	Exam Duration (Hrs)	Components of Marks		
						Internal	External	Total
						Total/ Passing	Total/ Passing	Total/ Passing
Core Course	PG01CMIC01	Molecular Biology	T	4	3	30/10	70/28	100/40
	PG01CMIC02	Bioanalytical Techniques and Instrumentation	T	4	3	30/10	70/28	100/40
	PG01CMIC03	Cell Biology	T	4	3	30/10	70/28	100/40
	PG01CMIC04	Practicals based on PG01CMIC01 and PG01CMIC02	P	4	3	30/10	70/28	100/40
	PG01CMIC05	Practicals based on PG01CMIC03 and PG01EMIC0X	P	4	3	30/10	70/28	100/40
	PG01CMIC06	Comprehensive Viva	P	1			50/20	50/20
Elective Course	PG01EMIC01	Fundamentals of Biochemistry and Bioenergetics	T	4	3	30/10	70/28	100/40
	PG01EMIC02	Food Microbiology	T	4	3	30/10	70/28	100/40
	PG01EMIC03	Microbial Physiology	T	4	3	30/10	70/28	100/40
	PG01EMIC04	Virology	T	4	3	30/10	70/28	100/40
Total Credits				25				650

SEMESTER- II

Course Type	Course Code	Name of Course	T / P	Credits	Exam Duration (Hrs)	Components of Marks		
						Internal	External	Total
						Total/Passing	Total/Passing	Total/Passing
Core Course	PG02CMIC01	Fermentation Technology	T	4	3	30/10	70/28	100/40
	PG02CMIC02	Basics of Microbial Genetics	T	4	3	30/10	70/28	100/40
	PG02CMIC03	Immunology	T	4	3	30/10	70/28	100/40
	PG02CMIC04	Practicals based on PG02CMIC01 and PG02CMIC02	P	4	3	30/10	70/28	100/40
	PG02CMIC05	Practicals based on PG02CMIC03 and PG02EMIC0X	P	4	3	30/10	70/28	100/40
	PG02CMIC06	Comprehensive Viva	P	1			50/20	50/20
Elective Course	PG02EMIC01	Biostatistics	T	4	3	30/10	70/28	100/40
	PG02EMIC02	Microtechniques	T	4	3	30/10	70/28	100/40
	PG02EMIC03	Omics and Computational Biology	T	4	3	30/10	70/28	100/40
	PG02EMIC04	Medical Microbiology	T	4	3	30/10	70/28	100/40
Total Credits				25				650

Course Wise Content Details for M.Sc. (Microbiology) Programme

**CHARUTAR VIDYAMANDAL UNIVERSITY
VALLABH VIDHANAGAR
SEMESTER I
M.Sc MICROBIOLOGY
SYLLABUS EFFECTIVE FROM: JUNE-2020-21**

PG01CMIC01: Molecular Biology

Course Objectives:

The objectives of this course are to make students understand how molecular machines are constructed and regulated so that they can accurately copy, repair, and interpret genomic information in prokaryotes and eukaryotic cells. Further, to appreciate the subject of molecular biology as a dynamic and ever-changing experimental science.

Course Learning Outcomes:

Unit 1: Students should be able to acquire basic knowledge on DNA structure, different conformations of DNA, supercoiling and DNA-protein interactions.

Unit 2: Students should be clear about organization of prokaryotic and eukaryotic genomes and should learn various molecular events that lead to duplication of DNA.

Unit 3: Students should have understood the process of transcription in prokaryotic and eukaryotic cells. They should have clear understanding of pre and post transcriptional modifications happening in the cells.

Unit 4: Student should have learnt protein synthesis in prokaryotic and eukaryotic cell along with processing of proteome in cell.

Contents:

UNIT -1

DNA structure

Chemistry of DNA, DNA structure, Different conformations of DNA (B, A and Z), Denaturation and Renaturation (Cot curves) of DNA. DNA topology: Supercoiling, Biology of Supercoiled DNA, DNA topoisomerases and their mechanism of action. DNA- protein interactions: General features, Sequence specific DNA binding protein motifs, ssDNA binding proteins.

UNIT –II

Organization of genome and its replication

Packaging of DNA and organization of chromosome in bacterial cells; Packaging of DNA in eukaryotic nucleosome and chromatin condensation, assembly of nucleosomes upon replication, chromatin modification.

Mechanism of DNA polymerase catalyzed synthesis of DNA, Types of DNA polymerases in bacteria, Initiation of DNA replication and its regulation in prokaryotes, assembly of replisome and progress of replication fork, termination of replication. DNA replication in eukaryotes and archaea. Inhibitors of DNA replication.

UNIT -III

Transcription

RNA polymerases, features of prokaryotic and eukaryotic promoters, assembly of transcription initiation complex in prokaryotes and eukaryotes, and its regulation; synthesis and processing of prokaryotic and eukaryotic transcripts.

UNIT-IV

Translation & Processing of proteome

Structure and role of t-RNA in protein synthesis, ribosome structure, basic features of genetic code and its deciphering, translation (initiation, elongation and termination in detail in prokaryotes as well as eukaryotes).

Post-translational processing of proteins (protein folding, processing by proteolytic cleavage, processing by chemical modification, Inteins), Protein degradation.

References:

1. Lewin's Genes X: Jocelyn E. Krebs
2. Molecular Biology of the Gene 6th Edition: Watson et al
3. Molecular Genetic of Bacteria 3rd Edition: Snyder and Champness
4. Molecular Biology: Genes to Proteins, 4th Edition: Burton E Tropp
5. Principles of Genetics 6th Edition: Snustad and Simmons
6. Genomes, 3rd Edition: T.A. Brown

CHARUTAR VIDYAMANDAL UNIVERSITY
VALLABH VIDHANAGAR
SEMESTER I
M.Sc MICROBIOLOGY
SYLLABUS EFFECTIVE FROM: JUNE-2020-21

PG01CMIC02: Bioanalytical Techniques and Instrumentation

Course Objectives:

The course will enable the students to understand the principle and working of visualization techniques, separation techniques, spectroscopic techniques for analysis of the samples and principles and applications of tracer techniques in biology. Principles and applications of different types of microscopy, principle & application of cytophotometry and flow cytometry, centrifugation, electrophoresis chromatography, spectroscopy, radioactivity, radiation counters, x-ray diffraction will be known to the students.

Course Learning Outcomes:

Unit 1: Deals with the knowledge of different types of microscopes such as Light microscope, Compound microscope, Dark field, Bright field, Stereo microscope, Confocal, Phase contrast microscope, Fluorescent microscope, Transmission Electron Microscopy (TEM) and Scanning Electron Microscopy (SEM). It also deals with the principle and application of cytophotometry and flow cytometry.

Unit 2: Enrich the concept and application for separation of molecules by different types of centrifugation techniques. Knowledge of separation by horizontal and vertical gel electrophoresis is also anticipated. The separation of molecules by different types of chromatographic techniques will be learnt.

Unit 3: Explore the consideration of principle and analysis of samples by different spectroscopic techniques such as UV, Visible, IR (including FTIR and ATR), AAS, NMR, Mass, MALDI-TOF, fluorescence, CD spectroscopy etc. will be learnt.

Unit 4: Gather the concept of radioactivity autoradiography, different types of counters used to trace the radiation will be studied. The principle and application of x-ray diffraction methods to study the structure of biopolymer will be known.

Contents:

Unit I

Visualization techniques:

Principle of working and applications of bright field & dark field microscopy, phase contrast microscopy, fluorescence microscopy, confocal microscopy, scanning and transmission electron microscopy, scanning tunneling microscopy, atomic force microscopy. Principle and applications of cytophotometry and flow cytometry.

Unit II

Separation techniques:

Basic principle and application of analytical and preparative centrifugation, settling time & velocity, types of rotor, sedimentation coefficient, relative centrifugal force (RCF) differential, density and ultracentrifugation.

Principle and applications agarose and 2D gel electrophoresis. Capillary electrophoresis and its applications. Native-PAGE, SDS-PAGE

Principle, methodology and applications of gel-filtration, ion-exchange and affinity chromatography; Thin layer and High-Performance Thin Layer Chromatography. Gas chromatography, High performance liquid chromatography and FPLC.

Unit III

Spectroscopy

Basic principle of electromagnetic radiation, instrumentation and applications of UV, Visible, IR (including FTIR and ATR), AAS, NMR, Mass, MALDI-TOF, fluorescence and CD spectroscopy.

Unit IV

Principle and applications of tracer technique in biology:

Concept of radioactivity, rate of radioactive decay; units of radioactivity- uses of radioisotopes in life sciences and biotechnology; autoradiography; cerenkov radiation; radiation dosimetry; ionization and scintillation-based detection of radioactivity.

Principle of biophysical methods used for analysis of biopolymer structure: X-ray diffraction.

References:

1. Instrumental method of chemical analysis: Sharma B K
2. Instrumental methods of analysis: D A Skoog
3. An introduction to practical Biochemistry: Plummer
4. Instrumentation: Chatwal and Anand
5. Modern experimental Biology: Boyer
6. Freifelder D. M. Physical Biochemistry- Application to Biochemistry and Molecular Biology, 2nd ed., W.H. Freeman, 1982.
7. Wilson & Walker. Principles and Techniques in Practical Biochemistry. 5th ed. Cambridge Univ. Press, 2000.
8. West & Todd. Biochemistry. 4th ed. Oxford and IBH.

9. Horst Friebolin. Basic One and Two-dimensional spectroscopy. VCH Publ, 1991.
10. Murphy D. B. Fundamental of Light Microscopy & Electron Imaging. 1st ed. Wiley-Liss, 2001.
11. R. Marimuthu – Microscopy and Microtechnique, MJP Publishers, 2015.

CHARUTAR VIDYAMANDAL UNIVERSITY
VALLABH VIDHANAGAR
SEMESTER I
M.Sc MICROBIOLOGY
SYLLABUS EFFECTIVE FROM: JUNE-2020-21

PG01CMIC03: Cell Biology

Course Objectives:

The major objective of this paper is to develop clear understanding of various aspects of cell biology along with diverse metabolic pathways existing at cellular level in relation to survival and propagation. This course enables the students to understand the structure and function of cell organelles, protein transport mechanism, intracellular signalling mechanism and acquainted with cell cycle, its regulation and apoptosis.

Course Learning Outcomes:

Unit 1: The students will understand the evolution of the cell, Cell as a unit of living organisms. They will learn structural details of prokaryotic and eukaryotic cells, their cell wall, cell membrane and other outer appendages.

Unit 2: The students can gain knowledge for molecular organization of Mitochondria, Chloroplast. Will know the ultrastructure and functions of Nucleus, Endoplasmic reticulum, Golgi complex, Lysosomes and other microbodies. They will also gain the knowledge of Protein sorting: organelle biogenesis and protein secretion, synthesis and its intracellular traffic, vesicular traffic in the secretory pathways

Unit 3: Will get the information for cytoskeleton topography which include the role of Microtubule and its dynamics, motor proteins, Microfilament and its functions, Intermediate filaments and their functions, Cilia and centrioles

Unit 4: Will be acquainted with overview of the Cell cycle and its control, the molecular mechanisms for regulating mitotic events, checkpoints in cell cycle regulation and signalling pathways which regulate apoptosis process

Contents

Unit I

The origin and Evolution of cells: Evolution of metabolism, Diversity of cell size and shapes, Structure of Prokaryotic and Eukaryotic cells, Single cell to multicellular organism

The Structure of cell membrane: The fluid Mosaic Model, Membrane lipids and Proteins, The Glycocalyx, Transport across plasma membrane.

Endocytosis: Phagocytosis and Receptor mediated endocytosis)

Cell walls and extracellular matrix & Cell Matrix Interactions

Cell-Cell interactions: Adhesion protein, Tight junctions, gap junctions and plasmodesmata.

Unit II

Cell Organelles: Molecular organization of Mitochondria, Chloroplast, Ultrastructure and Functions of Nucleus

Molecular Organization and functions of Endoplasmic reticulum, Golgi complex, Lysosomes (Protein sorting and transport, Types of vesicular transport and their functions), Microbodies: Peroxisomes, Ribosomes.

Unit III

The cytoskeleton: The nature of cytoskeleton, Intermediate filaments, Microtubules: Organization of tubules, assembly and organization within the cells, microtubule motors and movements, cilia and flagella: structure and function.

Cell signalling: Signalling molecules and their receptors, Functions of cell surface receptors, pathways of intracellular signal transduction, signal transduction and cytoskeleton.

Unit IV

Cell growth and division: Overview of the Cell cycle and its control, the molecular mechanisms for regulating mitotic events, Cell cycle control in mammalian cells, Checkpoints in cell cycle regulation, regulators of cell cycle progression-MPF, cyclins and CDKs, Inhibitors of cell cycle progression; M-phase and cytokinesis.

Programmed Cell Death: Difference between necrosis, apoptosis and necroptosis, Caspases, Central regulators of apoptosis (Bcl-2 family), signalling pathways that regulate apoptosis.

References:

- The cell: A molecular approach-Geoffrey M Cooper and Robert E. Hausman
- Cell Biology-Karp
- Molecular Biology of the cell- Alberts
- Molecular Cell Biology-Lodish et al.

CHARUTAR VIDYAMANDAL UNIVERSITY
VALLABH VIDHANAGAR
SEMESTER I
M.Sc MICROBIOLOGY
SYLLABUS EFFECTIVE FROM: JUNE-2020-21

PG01EMIC01 Fundamentals of Biochemistry and Bioenergetics

Course Objectives:

The major objective of this paper is to develop clear understanding of various aspects of biochemistry which includes properties of biomolecules, their metabolism and regulation. This course content enables students to better understand concept of bioenergetics and its importance in cellular metabolism. Moreover, useful to understand key role of water in metabolism which maintain acid base equilibrium at cellular level as well as an importance of physiological buffers.

Course Learning Outcomes:

Unit 1: Will have learnt carbohydrates, their types and properties. Further, will be acquainted with central metabolic pathways for carbon metabolism in bacteria enlisting differences with eukaryotic systems and their regulation in diverse physiological conditions.

Unit 2: Understands types of amino acids and their properties. Moreover, will have gathered understanding of inorganic and organic nitrogen assimilation and its regulation. Also knows role of glutathione in cellular redox regulation and biochemistry of glutamate overproducing strains. Will understand biochemical basis of nucleotides and its metabolism.

Unit 3: Will understand details of lipid, its metabolism and regulation along with biochemical basis of lipid accumulation at cellular level.

Unit 4: Will have learnt basic concepts of bioenergetics and its importance in cellular metabolism. The students will be aware with different electron carriers compounds and their role in ATP generation. Moreover, gain in depth knowledge of Water and Acid-Base Equilibrium.

Contents:

Unit I

Carbohydrates and Glycobiology: Monosaccharide - structure of aldoses and ketoses, ring structure of sugars, conformations of sugars, mutarotation, anomers, epimers and enantiomers, structure of biologically important sugar derivatives, oxidation of sugars. Formation of disaccharides, reducing and nonreducing disaccharides. Polysaccharides – homo- and heteropolysaccharides, structural and storage polysaccharides. Structure and role of proteoglycans, glycoproteins and glycolipids (gangliosides and lipopolysaccharides).

Carbohydrate metabolism: Glycolysis, Gluconeogenesis, PP Pathway, Citric acid cycle- steps involved, amphibolic nature, anaplerotic reactions, Coordinated regulation of glycolysis and gluconeogenesis, Glycogen synthesis

Unit II

Amino acids: Structure of amino acids, physical, chemical and optical properties of amino acids, Classification of amino acids, Peptides and Proteins, Secondary, tertiary and Quaternary structure of proteins

Protein metabolism: Nitrogen metabolism, Biosynthesis of amino acids, molecules derived from the amino acids, amino acid oxidation and production of urea

Nucleotides and Nucleic acids: Structure of major species of RNA - mRNA, tRNA and rRNA. Nucleic acid chemistry – UV absorption, effect of acid and alkali on DNA.

Nucleotides metabolism: Biosynthesis and Degradation of Nucleotides

Unit III

Lipids - fatty acids, glycerol, ceramide. Storage lipids - triacyl glycerol and waxes, Structural lipids in membranes – glycerophospholipids, galactolipids and sulpholipids, sphingolipids and sterols, structure, distribution and role of membrane lipids, Lipids as signals, cofactors and pigments

Lipid Metabolism: Biosynthesis of fatty acids, Triacylglycerol, membrane lipids and cholesterol, Fatty acid catabolism

Unit IV

Bioenergetics: The laws of thermodynamics, concept of entropy and free energy; ATP synthesis and hydrolysis, Biological oxidation: oxygenases, hydrolases, dehydrogenases, free energy changes and redox potentials, Gibbs energy

The mitochondrial respiratory chain, order and organization of carriers, proton gradient, iron sulphur proteins, cytochromes and their characterization, ATP- synthetase complex, Chemiosmotic theory of Energy Coupling, Inhibitors of ETC

Water and Acid-Base Equilibrium: Ionization of Water, Weak Acids, and Weak Bases, buffering against pH Changes in Biological Systems: Henderson and Hassebach equation, Buffers and their importance, pKa of amino acid and their relevance, Importance of discontinuous buffer system used in SDS PAGE, Water as a Reactant

References:

- Lehninger's Principles of Biochemistry: D. L. Nelson and M. M. Cox, Macmillan, Worth Pub. Inc., NY.
- Chemistry of Biomolecules by S. P. Bhutani, Ane Books Pvt. Ltd. CRC Press
- Biochemistry: Lubert Stryer WH Freeman & Co., NY.
- Harper's Biochemistry: R. K. Murray and others. Appleton and Lange, Stanford.
- Text book of Biochemistry with clinical correlations by Delvin.

CHARUTAR VIDYAMANDAL UNIVERSITY
VALLABH VIDHANAGAR
SEMESTER I
M.Sc MICROBIOLOGY
SYLLABUS EFFECTIVE FROM: JUNE-2020-21

PG01EMIC02: Food Microbiology

Course Objectives:

The course will enable students to understand the taxonomical classification, phenotypic and biochemical identification of food associated molds, yeasts, yeast-like fungi and bacteria. The course will teach the strategies to develop fermented and non-fermented milk products, fermented plant-based products, malt beverages, distilled liquors, etc. The role of microbes in food spoilage, preservation and various food borne diseases can be discussed.

Course Learning Outcomes:

Unit 1: Will know about production and evaluation of the quality of starter cultures and fermented milk products. They will understand the role of microbes in food spoilage and how different factors affect this process.

Unit 2: Gathers information regarding microbes causing food intoxications and food-borne infections. The students will learn different diagnostics methods and preventive measures.

Unit 3: Knows traditional food preservation techniques including drying, salting, refrigeration, vacuum packaging, canning/bottling, chemical preservation and irradiation. The students will also learn use of modern techniques viz. high-pressure processing (HHP), bacteriocins, manosonication (MS), etc. They will be aware of fermentation protocols of different food products and understands the use and production of probiotics, prebiotics and nutraceuticals.

Unit 4: Gains 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. Understands the relevance of microbial standards for food safety, quality assurance programs that revolutionize food safety.

Contents:

Unit I: Microbiology of food

Microorganisms important in food microbiology

- a) Food associated bacteria, yeasts and molds. Microbiome of food material.
- b) Factors influencing microbial growth in food.

Microbial spoilage of foods

- a) Spoilage of cereals and its products, vegetables, fruits, meat and meat products, milk and milk products, canned and sugar products, fish, seafood and poultry
- b) Study of microbes responsible for spoilage and brief insights into chemical and physical spoilage of foods.

Unit II:

Food borne infections

- a) Bacterial food borne infections and intoxications-*Brucella*, *Campylobacter*, *Clostridium*, *Escherichia* (ETEC/EHEC/EPEC/EAEC), *Salmonella*, *Shigella*, *Listeria*, *Vibrio*, and *Yersinia*.
- b) Non- bacterial food borne infections and intoxications- Nematodes, protozoa, algae, fungi, and viruses.
- c) General methods for diagnosis of infections, intoxications and preventive measures.

Unit III:

Food preservation

General principles of food preservation – Classical, Physical, chemical and biological food preservation methods

Fermentative food products

Starter cultures for fermented foods, Fermented milk products: Yogurt, Cheese, Kefir, etc

Oriental fermented foods: Shoyu, Temph, Kimchi, etc, Fermented vegetables – Sauerkraut

Food beverages: Malt beverages, wines, vinegar.

Role of Probiotics, prebiotics and nutraceuticals

Unit IV:

Molecular techniques in detection of food pathogens and GM foods. Biosensors in food

Food research organizations/institutes in India

Food sanitation – Microbiology of food plant sanitation, water and milk testing

Food laws and quality control – HACCP, Codex alimentarius, PFA, FPO, MFPO, BIS, AGMARK.

References:

1. Food Microbiology by W.C. Frazier, D.C. Westhoff, K.N. Vanitha. 5th edition. McGraw Hill Education. 2013.
2. Fundamental Food Microbiology by B. Ray and A. Bhunia. 5th edition. CRC press. 2013.
3. Food Microbiology by M. R. Adams, M. O. Moss, P. McClure. 4th edition. Royal Society of Chemistry. 2015.
4. Food Microbiology: Fundamentals and Frontiers by M. P. Doyle, L. R. Beuchat. 3rd Edition
5. Dairy Microbiology by Robinson. Volume II and I

CHARUTAR VIDYAMANDAL UNIVERSITY
VALLABH VIDHANAGAR
SEMESTER I
M.Sc MICROBIOLOGY
SYLLABUS EFFECTIVE FROM: JUNE-2020-21

PG04EMIC03: Microbial Physiology

Course Objectives:

The major objective of this paper is to develop clear understanding of various aspects of microbial physiology along with diverse metabolic pathways existing in bacteria in relation to its survival and propagation. This course enables the students to understand stress responses, intracellular signaling mechanism and acquainted with different protective resistance microbial responses.

Course Learning Outcomes:

Unit 1: The students will understand the structural details of cell wall, cell membrane and other outer appendages. Will have gained an in-depth knowledge of primary, secondary and group translocation transport systems existing in bacteria, simultaneously learning membrane transport proteins and kinetics of solute transport. Moreover, it will be useful to understand motility types and mechanism of microbial cells.

Unit 2: Will be acquainted with methods of measuring microbial growth, calculating growth kinetic parameters with understanding of steady state and continuous growth. Understands various microbial stress mechanisms and responses

Unit 3: Is conversant with intracellular signalling in bacteria in response to various nutritional and physiological stresses. Helpful to understand role of signalling compounds, its regulation and its response for quorum sensing, biofilm formation and bioluminescence process.

Unit 4: Will be acquainted with different mechanisms of drug resistance, various protective mechanisms of microbes upon infections. Moreover, make the students well aware with bacterial immune system, CRISPR/Cas

Contents:

Unit-I

Bacterial Cell Structure and its type, Bacterial Cell wall structure function and synthesis Membrane transport in procaryotes-simple, group translocation, ABC transporters, Protein export in bacteria– Type 1,2,3,4, Protein export pathways. 2. Permeation- Primary Active transport, secondary active transport, Co transport

Transport of ions across the membrane V-type, F-type and p-type ATPases

Bacterial organs for locomotion: Flagella: structure, synthesis, function and mechanism of locomotion, Swarming motility, Motility in spirochetes, Gliding motility, Twitching.

Chemotaxis: Molecular mechanism and physiological significance.

Two component signal transduction in prokaryotes

Unit-II

Bacterial cell division: molecular mechanisms involved in formation of Z-ring, Cell division machinery.

Bacterial differentiation: endospore formation, physiological and genetic aspects of sporulation, Sporulation inducing signals & events in sporulation

Microbial stress responses: Heat shock, pH, aerobic-anaerobic shifts- Arc and Fnr system, Oxygen toxicity: Mechanism of oxygen toxicity and its mechanism to overcome toxicity-catalase, peroxidase and superoxide dismutase, Osmotic pressure, Osmolarity regulation in *E.coli* (Omp system), Phosphate assimilation in *E.coli* (Pho system).

Unit-III

Quorum sensing process in microorganisms

Bioluminescence process, biochemistry, genetics and significance.

Bacterial biofilms formation steps, dispersion and control strategies

Siderophores; structure, function and significance

Microbial fuel cells: Energy generation principle and application. Production of Hydrogen.

Unit-IV

Mechanism of drug resistance.

Bacteriocins: Structure, Classification and physiological significance of it.

Host Parasite interactions: Structures and functions involved in Host-parasite interactions, Bacterial damages to host upon infection. Structure and Mechanism of Endotoxin, Exotoxin and Exoenzymes formed by bacteria.

The prokaryotic “immune system”, CRISPR/Cas

References:

- Microbial Physiology by A.G. Moat, J. W. Foster, M. P. Spector. 3rd Edition. John Wiley & Sons. 2002
- The Physiology and Biochemistry of prokaryotes, David White
- Bacterial signalling, Kramar and Jung
- Bacterial physiology: A molecular approach, W. E. Sharoud
- Topic related review articles

CHARUTAR VIDYAMANDAL UNIVERSITY
VALLABH VIDHANAGAR
SEMESTER I
M.Sc MICROBIOLOGY
SYLLABUS EFFECTIVE FROM: JUNE-2020-21

PG01EMIC04: Virology

Course Objectives:

The main objective of the course is to make students to understand the taxonomical classification, phenotypic and biochemical identification of various viruses. The course will teach the strategies to develop isolation and preservation of viruses and their life cycle for development of antiviral compounds for the viral infections and the role of various viruses in the development of vaccines etc.

Course Learning Outcomes:

Unit 1: Student will be able to describe the defining viral attributes, the general properties of viruses, and steps in virus infection cycle. The principle of virus classification, list the virus families, and describe methods of study virus infection. General overview of viral genomes and their types as well as isolation & preservation of viruses.

Unit 2: Student will be able to receive information regarding various bacteriophages life cycles, which are model viruses for the study. Applications of phages in therapy; Concern over phage contamination in food & fermentation industry.

Unit 3: Students will be able to know various plant and animal viruses and their isolation, preservation and classifications. Student will be able to describe host defense against virus infection and able to describe general characteristics of acute viral infections, pathogenesis of Influenza virus, Polio virus, Measles virus, and Rotavirus infection. Student will be able to describe general characteristics of chronic, persistent, latent infections

Unit 4: Student will be able to describe how different antiviral drugs and their mode of action of viruses, student knows how live viral vaccines are made, how inactivated viral vaccines are made, Polio vaccine and story of polio eradication.

Student is able to describe antiviral drug discovery process, mechanism of drug resistance and use of interferons for viral infections.

Contents:

Unit – I: Prokaryotic Viruses

Discovery of bacteriophages, Structure and composition of bacteriophages, Classification system of Baltimore & ICTV

Phage biodiversity, Genome diversity and host- specific interactions

Isolation and purification by filtration, ultracentrifugation and affinity chromatography

Plaque assays

One step growth, single burst and eclipse experiments

Unit – II:

Life cycle of model bacteriophages infecting *E coli* – λ (lytic lysogenic)

Life cycle of model bacteriophages: ϕ X 174, M13

Life cycle of model bacteriophages: T4, T7

Life cycle of model bacteriophages: Q β , Mu

Applications of phages - therapy; Concern over phage contamination in industry (dairy)

Unit – III : Eukaryotic Viruses

Discovery and classification of plant and animal viruses, structure of viruses, viroids, virusoids

Classification of viruses – ICTV and Baltimore classifications

Host – viruses interactions, permissive/non – permissive hosts; Cytopathic effects

Isolation and purification of viruses, Cultivation and propagation

Assay methods – pock assay, hemagglutination assay, transformation assay.

Structure, Life cycle and Pathogenicity of Gemini virus

Structure, Life cycle and Pathogenicity of TMV

Structure, Life cycle and Pathogenicity of Adenovirus

Structure, Life cycle and Pathogenicity of Rotavirus

Structure, Life cycle and Pathogenicity of Rubella, Influenza and Measles viruses

Structure, Life cycle and Pathogenicity of HIV and Hepatitis B Virus

Unit – IV: Prevention & control of viral diseases

Antiviral compounds and their mode of action

Interferon and their mode of action.

General principles of viral vaccination

Applications of Virology:

Use of viral vectors in cloning and expression, Gene therapy and Phage display

References:

1. Principles of Virology, (Vol I & II) Flint SJ, Enquist LW, Racaniello VR, Skalka AM, Pub ASN Press
2. Introduction to Modern Virology – Dimmock
3. Basic Virology – Wagner
4. Virology – Saravanan

5. Virology – Maharajan
6. Molecular Virology – A. J. Cann
7. An introduction to Viruses – Biswas

PG01CMIC04: Practicals based on PG01CMIC01and PG01CMIC02

List of Practicals

1. Amino acid titration curve
2. DNA estimation by DPA method and UV absorption
3. RNA estimation by orcinol method
4. Isolation of chromosomal DNA
5. Separation of proteins by PAGE
6. Introduction to pH, buffer preparation, molar, normal and % solutions.
7. Calculations for making stock solution
8. Separation of amino acids by TLC
9. Separation of cells by density gradient centrifugation
10. Determination of partition coefficient

PG01CMIC05: Practicals based on PG01CMIC03and PG01EMIC01

List of Practicals

1. Estimation of Reducing Sugar in Jaggery by Cole's Method
2. Estimation of Protein by Folin-Lowry Method
3. Estimation of Reducing Sugar by DNS Method
4. Total Sugar Estimation by Phenol Sulphuric acid estimation
5. Estimation of RNA by Orcinol Method
6. Localization of Cell Organelle and Determination of Chlorophyll and Carotenoids
7. Estimation of Amino Acid (Proline)
8. Estimation of Amino Acid (Methionine from Food Grains)
9. Study of Cell structure (Eukaryotic & Prokaryotic)
10. Study of Meiosis and Mitosis

CHARUTAR VIDYAMANDAL UNIVERSITY
VALLABH VIDHANAGAR
SEMESTER II
M.Sc MICROBIOLOGY
SYLLABUS EFFECTIVE FROM: JUNE-2020-21

PG02CMIC01: Fermentation Technology

Course Objectives:

The major objective of this paper is to impart knowledge about fermentation processes and its relevant aspects. The course will teach strain improvement strategies, preservation methods, sterilization of media and air. It will be useful to understand various bioreactors and their applications. This course will enable the students to understand aeration-agitation concept, its importance in fermentation process and growth kinetics. Lastly, useful to understand downstream processes of the fermentation process.

Course Learning Outcomes:

Unit 1. Is able to describe the role of microbes in fermentation processes. The students will understand different strategies of strain improvement. It will also be useful to understand the role of medium components on product formation.

Unit 2. Understands aseptic environment, sterilization and its various methods. Will know fermenter design, its components and its variable control parameters.

Unit 3. Understands microbial growth, its kinetics and association of product formation with growth. The students will understand the concept of mass transfer and various methods to determine $K_L a$.

Unit 4. Is able to describe various methods of product recovery. Will know the role of various chromatography in product purification. Moreover, makes the student aware of desalting, drying and crystallization processes.

Contents:

Unit I

Isolation, Screening: Primary and Secondary, Preservation and maintenance of Industrially important microorganisms

Strain Improvement of industrially important microbes: Isolation of mutant producing primary and secondary metabolites, isolation and use of auxotrophic mutants, isolation and use of revertant mutants and use of recombination systems

Media for industrial fermentation processes: Energy sources, antifoam agents and medium optimization

Unit II

Sterilization methods and principles: Media sterilization, mathematical modelling of sterilization processes, Arrhenius equation, Del factor, effect of sterilization on media quality and yield coefficients, batch and continuous sterilization, filter and steam sterilization at industrial scale

Design of fermenter and reactors: Basic components of a fermenter, laboratory and industrial scale fermenters, mechanical, Types of fermenter like stirred tank, bubble column, airlift, packed beds, fluidized beds, perfusion cultures, photo-bioreactors and animal cell culture bioreactor, Plug flow reactors, Immobilized enzyme reactors.

Bioprocess Control parameters: Instrumentation for monitoring bioreactor and fermentation processes, Sensors, Controllers, fermentation control systems and architecture, Incubation and sequence control, advanced control Scale up and Scale down and containment

Unit III

Microbial Growth kinetics: Kinetics of growth and substrate utilization in batch, fed batch and continuous systems. Inoculum development, aseptic inoculation and sampling.

Agitation and aeration: Mass transfer of oxygen, Determination of K_{La} , factors affecting K_{La} , fluid rheology, newtonian and non-newtonian fluids, bingham plastic, pseudo plastic, power number, Reynolds number.

Unit IV

Recovery and Purification of fermentation Products: Bio separation: filtration, centrifugation, sedimentation, flocculation, cell disruption, liquid-liquid extraction.

Purification by chromatographic techniques, Membrane Processes, drying, crystallization, storage and packaging.

Fermentation Economics

References:

- Principles of Fermentation Technology : Whitekar & Stanbury
- Comprehensive Biotechnology : Murray Moo Young
- Methods in Industrial Microbiology : Sikyta
- Fermentation Microbiology and Biotechnology, El Mansi and Bryc

CHARUTAR VIDYAMANDAL UNIVERSITY
VALLABH VIDHANAGAR
SEMESTER II
M.Sc MICROBIOLOGY
SYLLABUS EFFECTIVE FROM: JUNE-2020-21

PG02CMIC02: Basics of Microbial Genetics

Course Objectives:

The objectives of this course are to take students through basics of microbial genetics covering different types of mutations, plasmid biology, prokaryotic genetics and agrobacterium genetics. On covering the course the student will be exposed to concepts of mutation, DNA damage and repair, plasmid biology, microbial and phage genetics.

Course Learning Outcomes:

Unit 1: Students will know that genome is transient and mutation keeps on happening. They will know the types mutation and different mechanisms involved in their repair.

Unit 2: Students should be clear about types of plasmids, their compatibility regulation of copy number and segregation. Students will also learn about phage genetics and recombination.

Unit 3: Students should have understood the types and process of transformation, conjugation and transduction at the end of this unit

Unit 4: Here student should have learnt Agrobacterium genetics, types of restriction modification systems and different types of transposable elements.

Contents:

UNIT -1

Mutation, DNA damage and Repair

Spontaneous mutations (Random v/s Adaptive nature of mutation; Mutation rate and its determination, Types of DNA damage and their consequences (spontaneous and chemical induced deamination, radiation induced DNA damage, loss of nitrogen bases, alkylation, intra and inter strand cross linking) , DNA repair pathways (Mis-match repair in prokaryotes and eukaryotes, Nucleotide excision repair in prokaryotes and in eukaryotes, base excision repair, recombinational repair, SOS pathway, specific repair of oxidative DNA damage, repair of pyrimidine dimers, repair of alkylation induced damage and adaptive response and other specific repair mechanisms).

UNIT –II

Plasmid Biology, Phage Genetics & Recombination

Types of plasmids, compatibility, regulation of plasmid copy number & plasmid segregation

T-series, complementation and Fine structure analysis, biology of lambda phages.

Types of recombination, Different models of recombination, Molecular mechanism of homologous recombination in eukaryotes, Mating type switching, Site specific recombination and its biological significance.

UNIT -III

Genetic exchange in prokaryotes

Natural transformation in *Bacillus subtilis*, Transformation by inducing artificial competence, Gene linkage and mapping by transformation.

Generalized transduction in T4 bacteriophage, Specialized transduction, homologous recombination with recipient's chromosome, measuring transduction (co-transduction of markers, marker effects, abortive transduction, transduction of plasmids). Applications of transduction.

F-factor mediated Conjugation in *E. coli*, Hfr conjugation and chromosomal transfer, F-prime conjugation and merodiploids, Conjugation of fertility inhibited F-like plasmids, Non conjugative mobilizable plasmids, chromosomal mobilization of non-F plasmids, Interrupted mating and conjugational mapping.

UNIT-IV

Agrobacterium genetics, Restriction Modification Systems, Transposable Elements

Ti plasmid, Interkingdom gene transfer (Key early experiments, vir regulon, protein secretion apparatus, conjugation model of T-DNA transfer, Integration products)

Types of RM systems, Role of RM systems, salient features and insights into evolution of diverse types of Restriction endonucleases and Methyl transferases, Regulation of RM systems.

Types of bacterial transposable elements; Structure, genetic organization and mechanism of transposition of Tn5, Tn3, phage Mu, Tn7, IS911, Integrons, Retrotransposons, conjugative and mobilizable transposons, Assays of transposition.

References:

1. Lewin's Genes X: Jocelyn E. Krebs
2. Molecular Biology of the Gene 6th Edition-Watson et al.
3. Modern Microbial Genetics 2nd Edition-Uldis Streips and Ronald Yasbin
4. Microbial genetics 2nd Edition-Stanley Molay, John Cronan and David Freifelder.
5. Molecular Genetics of Bacteria 3rd Edition-Snyder and Champness.
6. Molecular Genetics: An Introductory Narrative 2nd Edition-Stent and Calender
7. Principles of Genetics 6th Edition- Snustad and Simmons
8. Molecular Biology of the Cell 5th Edition-Alberts et al.

CHARUTAR VIDYAMANDAL UNIVERSITY
VALLABH VIDHANAGAR
SEMESTER II
M.Sc MICROBIOLOGY
SYLLABUS EFFECTIVE FROM: JUNE-2020-21

PG02CMIC03: Immunology

Course Objectives:

The objective of this course is to understand various components of the host immune system; their structure, organization and role in defence mechanism. The student will gain knowledge to understand the operational mechanisms which underlie the host defence system. It would make them clear to understand genetic organization and expression of receptors to show immune response. They will also learn the role of immune system in health and diseases.

Course Learning Outcomes:

Upon successful completion of the course, the student will learn:

Unit 1: Will be able to understand the types of immunity and basic components of immune system; the role played by complement system as an interface between innate and adaptive immunity.

Unit 2: Will be able to understand the genetic organization of the genes meant for expression of immune cell receptors and the basis of the generation of their diversity. The principle of antigen-antibody interactions and methods to measure them will become clear to students.

Unit 3: Will be able to understand the importance of MHC molecule in an individual's immunity to various antigens, the mechanism of antigen processing and presentation. They will be able to understand the mechanism of B and T cell activation and memory generation.

Unit 4: The students will gain knowledge about the mechanism of cell mediated immunity. They will learn about the cytokines, important biopharmaceuticals and their role in modulation of immune response. The students will also learn how body shows different kinds of immune response to different infections.

Contents:

Unit I

Immunity: Innate and Adaptive, Cells of the Immune system: Haematopoiesis and its regulation

Cells and organs of the immune system: Primary and secondary lymphoid organs

Induced Innate immunity: receptors of the innate immunity (TLR and sensing of PAMPs, CLR,RLR and CLR); Inflammatory responses, Natural Killer cells

Antigens: Immunogenicity versus antigenicity, Epitopes, Haptens.

Complement system: The Major Pathways of Complement Activation: Classical, alternative and lectin complement pathways, functions of complement, regulation of complement, complement deficiencies, microbial complement evasion strategies

Unit II

Antibody: Structure of immunoglobulin; classes of immunoglobulins, Signal transduction pathways emanating from the BCR

The Organization and Expression of Lymphocyte Receptor Genes: Hozumi and Tonegawa's Experiment, Multigene organization of Ig Gene, Mechanism of VDJ recombination, B cell receptor expression: Allelic exclusion, B cell isotype switching and somatic hypermutation; expression of membrane bound and soluble immunoglobulin; T cell receptor genes and expression

Basics of Antigen-antibody interactions: Immunoprecipitation and agglutination based techniques, Methods to determine affinity of antigen-antibody interactions, Immunofluorescence, FACS

Unit III

The Major Histocompatibility Complex and Antigen Presentation: The structure and function of MHC molecules, general organization and inheritance of MHC genes, The role and expression Pattern of MHC, Endogenous and exogenous pathway of antigen processing and presentation; presentation of non-peptide antigens.

B Cell activation: T dependent and T independent B cell responses and memory generation

T Cell activation: Two signal hypothesis, superantigens, activation and differentiation of T cell into effector and memory cells. T_H1 and T_H2 responses.

Unit IV

Cell mediated effector response (Generation of effector CTL's, Granzyme and Perforin Mediated Cytolysis, Fas-FasL Mediated Cytolysis, NK cell mediated cytotoxicity)

Cytokines: properties, receptors, associated diseases, therapeutic applications, cytokine signalling pathways: JAK-STAT and FAS-FASL signalling pathways

Immune response to infection by viruses, bacteria, fungi and parasite: Mechanism of Immune response and evasion by pathogen

References

- 1 Owen, J. A., Punt, J., & Stranford, S. A. (2013). *Kuby immunology* (7thEdn). New York: WH Freeman.
- 2 Murphy, K., & Weaver, C. (2016). *Janeway's immunobiology* (9thEdn) Garland Science.
- 3 Male, D., Brostoff, J., Roth, D., & Roitt, I. (2012). *Immunology* (8thEdn) *With STUDENT CONSULT Online Access*. Elsevier Health Sciences.
- 4 Abbas, A. K., Lichtman, A. H., & Pillai, S. (2014). *Cellular and molecular immunology* (6thEdn) Elsevier Health Sciences.
- 5 Relevant review articles / research papers / handouts of latest development in the subject.

CHARUTAR VIDYAMANDAL UNIVERSITY
VALLABH VIDHANAGAR
SEMESTER II
M.Sc MICROBIOLOGY
SYLLABUS EFFECTIVE FROM: JUNE-2020-21

PG02EMIC01: BIOSTATISTICS

Course Objectives:

The course aims to develop competency and expertise in the application of statistical methods applied to biological data obtained in experimental techniques.

Course Learning Outcomes:

Unit 1: Student will be able to know about basic terms and use in biostatistics. They will understand types of data, their organisation and various graphical representation methods to represent data, and will enable students to understand the basic statistics and its importance in research.

Unit 2: Student will be able to calculate various measures of central tendencies, measure of dispersion and measure of kurtosis and skewness and its importance.

Unit 3: To understand the exact method of data analysis for the problem under investigation. Student will be able to perform various hypothesis testing like T-test, F-test, and chi square tests and its application in biological sciences.

Unit 4: Understanding for drawing valid inferences and to plan for future investigations. Student will be able to perform Correlation & regression calculations and its application in Biological sciences. Student will able to perform ANOVA testing.

Contents:

Unit I:

Data Collection and Presentation

Types of Biological Data: Qualitative Data -Nominal, Ordinal, Ranked; Quantitative Data: Discrete and Continuous.

Understanding of Population and sample

Methods of Collection of Data: (i) Experimental Data and (ii) Survey Data- Simple random Sample (with and without replacement), stratified sampling and cluster sampling.

Tables: Frequency Distributions, Relative Frequencies.

Graphical Presentation: Bar charts, Histograms, Frequency Polygons, One way scatter plots, Box plots, two-way scatter plots, line graphs.

Unit II:

Descriptive Statistics

Measures of Central Tendency: Mean, Median and Mode, quartiles, deciles and percentiles (both for raw data and grouped data)

Measures of Dispersion: Range, Interquartile Range, Variance, Standard Deviation and Coefficient of Variation.

Measures of Skewness and Kurtosis.

Unit III:

Statistical hypotheses: Null and Alternative hypotheses.

Statistical Tests: Acceptance region and Rejection Region. Types of errors and power of the test.

Goodness of fit tests.

Random Variables: Discrete and Continuous. Some examples from biological sciences.

Probability Distributions: General Normal Distribution, Standard Normal Distribution ; Sampling Distributions- t, chi-square and F distributions.

Significance Tests for Normal Distribution: One sample tests for mean – z test and t-test.

Two sample tests for normal distributions: Tests for means (i) when variances are known (ii) when variances are unknown. Tests for equality of variances.

Paired t-test for equality of means.

Confidence Intervals

Unit IV:

Correlation: Covariance, Calculation of covariance, correlation analysis and correlation Coefficient calculated from ungrouped data.

Regression: Simple linear regressions analysis, regression coefficients, Linear regression line or equation

Analysis of Variance: Completely Randomized Design, Randomized Block Design

References:

- Fundamentals of statistics by S.C. Gupta
- Principles of Biostatistics by Marcello Pagano and Kimberlee Gaurea
- Biostatistics : A Foundation For Analysis in the Health Sciences by Daniel, Wayne (Seventh Edition), Wiley India Pub.

CHARUTAR VIDYAMANDAL UNIVERSITY
VALLABH VIDHANAGAR
SEMESTER II
M.Sc MICROBIOLOGY
SYLLABUS EFFECTIVE FROM: JUNE-2020-21

PG02EMIC02- Microtechniques

Course Objectives:

The course will enable the students to understand the principle of microscopy types of microscopy used to explore the knowledge of microtechniques. The measurement of size of microorganisms, sanctioning of the bigger specimens by using microtome, preparation of temporary and permanent slides of the specimen will be known.

Course Learning Outcomes:

Unit 1: Deals with the concept and principle of microscopy. It provides the understanding of different optical components of microscopy,

Unit 2: Enrich the knowledge of different types of microscopes such as Light microscope, Compound microscope, Dark field, Bright field, Stereo microscope, Confocal, Phase contrast microscope, Fluorescent microscope, Transmission Electron Microscopy (TEM) and Scanning Electron Microscopy (SEM).

Unit 3: Explore the need and methods of measurement of microorganisms by micrometry. The calibration and working with the stage and ocular micrometer. Illustrations and concept of photomicrography will also be known.

Unit 4: The concept of killing and fixation agents, dehydration of the specimens, embedding of specimens in paraffin wax, free hand sanctioning, mounting of sanctioned specimen on slide, staining of specimens and different types of staining will be known.

Contents

Unit 1

Principles of microscopy – eyepiece lens and objective lenses; Magnification, Resolving power, numerical aperture. Mechanical components: base, pillar, stage, sub stage, body tube, focusing knobs, nose pieces. Optical components: mirror, objectives, ocular lens, condenser, Focussing slides under low/ high power and oil immersion.

Unit 2

Types of microscopes: Light microscope, Compound microscope, Dark field, Bright field, Interference microscope (Stereo microscope), Confocal, Inverted microscope, Phase contrast

microscope, Fluorescent microscope, Electron microscope: Transmission Electron Microscopy (TEM) and Scanning Electron Microscopy (SEM)

Unit 3

Measurement of Microorganisms- Micrometry – Stage micrometer, Ocular micrometer, Calibration and working. Preparation of illustrations using camera lucida, digital camera and photomicrography.

Unit 4

Killing and fixation agents - carnoy's formula, F. A. A.

Dehydration– general account of dehydration (Ethanol, Isopropyl alcohol, Acetone, Glycerine). Ethanol – Xylene series and Tertiary Butyl Alcohol Series

Infiltration – paraffin wax method, Embedding

Free hand sectioning- Microtome (Rotary and sledge) serial sectioning and its significance.

Mounting- A brief account on whole mounting, maceration, smears and squash preparation, application of permanent whole mounts, permanent sections.

Staining- Classification: natural dyes, coal tar dyes, double staining, vital staining; simple, Gram staining, negative staining, capsule staining, spore staining, flagellar staining, nuclear staining and acid-fast staining, stains: saffranin, hematoxylin, acetocarmine.

References

1. Plant Microtechnique, Johansen D.A. 1940, Mc Graw – Hill Book Company, Inc. New York.
2. Manual of Microbiology – Tools and Techniques, Kanika S. 2007, Ane's student edition.
3. Botanical Microtechnique; principles and Practice, Khasim S.K., 2002, Capital Publishing Company New Delhi.
4. Essentials of botanical microtechnique, Toji T. 2004, Apex Infotec Publ.
5. Murphy D. B. Fundamental of Light Microscopy & Electron Imaging. 1st ed. Wiley-Liss, 2001.
6. R. Marimuthu – Microscopy and Microtechnique, MJP Publishers, 2015.

CHARUTAR VIDYAMANDAL UNIVERSITY
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SEMESTER II
M.Sc MICROBIOLOGY
SYLLABUS EFFECTIVE FROM: JUNE-2020-21

PG02EMIC03: Omics and Computational Biology

Course Objectives:

The course will enable the students to understand the concept of genome mapping, genome sequencing, functional genomics, basic concepts of proteomics tools, data mining, basic concepts and tools of lipidomics, glycomics and phosphoproteomics. Storage and retrieval of various types of databases collection and storing of sequence data will be understood by the students. Students will also be able to know the local and global alignment through scoring matrices, gene prediction methods, RNA fold analysis, splice site identification.

Course Learning Outcomes:

Unit 1: Deals with genome, genomics and transcriptomics. The concept and application of physical map, genetic map, genome sequencing, functional genomics, small or large regulatory RNAs and dark matter will be known.

Unit 2: Gathers information regarding concept of proteomics, metabolomics and lipidomics. The the basic tools of proteomics, metabolomics, lipidomics and their applications will be learnt by the students.

Unit 3: Deals with the primary and secondary databases, collection, storage and retrieval of databases, knowledge of freeware, software and hardware. The sequence databases, sequence format, annotation and archival of databases will be understood.

Unit 4: Accords the sequence alignment and applications. The choice of alignment, local alignment, global alignment scoring matrices, codon usages analysis, RNA fold analysis, splice site identification will also be studied by the students.

Contents

Unit 1

Genome, Genomics & Transcriptomics:

Genome mapping: Physical and Genetic Map, Genome Sequencing, Next generation sequencing methods, Genome Annotation, Functional Genomics. Transcription factor binding sites, RNA-Seq, Microarrays, Regulatory RNAs: small or large, Computational prediction of miRNA target genes, RNA Dark matter

Unit 2

Proteomics, Metabolomics & Lipidomics:

Basic concepts, Tools of proteomics- SDS PAGE, 2D PAGE, Liquid chromatography, Mass Spectrometry (ESI and MALDI), Protein identification by peptide mass fingerprinting, Applications of proteomics.

Fundamental concept, data integration and data mining; Tools of metabolomics-Capillary electrophoresis, Gas chromatography, Electrochemical detectors.

Basic concepts and tools of lipidomics, glycomics and phosphoproteomics.

Unit 3

Biological Literature Information access, storage and retrieval systems- Primary and secondary databases of genomics, transcriptomics, proteomics and metabolomics. Knowledge on freeware and commercial software. Importance of hardware and software creations.

Collecting and Storing Sequence Data: Sequence assembly; Submission of Sequences; Sequence accuracy; Sequence databases; Sequence formats; Annotation and Archival.

Unit 4

Sequence alignment and applications: Uses: Choice to be made for alignment; Scoring matrices; Homology and related concepts; Dot Matrix methods; Dynamic programming methods for global and local alignments tools- FASTA, BLAST, statistical and Biological significance.

Nucleic acid sequence analysis: Reading frames; Codon Usage analysis; Translational and transcriptional signals; Splice site identification; Gene prediction methods; RNA fold analysis

References:

1. Introduction to Proteomics -Tools for the New Biology by Daniel C. Liebler, Humana Press.
2. Mass Spectrometry for Biotechnology by Gary Siuzdak, Academic Press.
3. Proteomics for Biological Discovery by Timothy Veenstra and John Yates, Wiley.
4. Metabolomics- Methods and Protocols by Wolfram Weckwerth, Humana Press.
5. Lipidomics- Technologies and Applications by Kim Ekroos, Wiley-VCH.
6. Web/Journal Resources.
7. Transcriptomics: Expression Pattern Analysis, Virendra Gomase, Somnath Tagore; VDM Publishing, 2009 – Science.
8. Current Protocols in Bioinformatics, Edited by A.D. Baxevanis et al, Wiley Publishers. 2005.

9. Bioinformatics by David W. Mount, Cold Spring Harbor Laboratory Press. 2001.
10. Fundamental concepts of Bioinformatics by D.E. Krane and M.L Raymer, Pearson Education. 2003.
11. Bioinformatics and Functional Genomics by Pevsner, J., John Wiley and Sons, New Jersey, USA. 2003
12. Principles of Genome Analysis and Genomics (3rd Ed.) by Primrose, S.B. and Twyman, R.M., Blackwell Publishing Company, Oxford, UK. 2003.
13. Introduction to proteomics – Tools for the new biology (1st Ed.) by Liebler, D.C., 2002, Human Press Inc., New Jersey, USA.
14. Bioinformatics: Sequence and Genome Analysis by Mount, D., Cold Spring Harbor Laboratory Press, New York. 2004.

CHARUTAR VIDYAMANDAL UNIVERSITY
VALLABH VIDHANAGAR
SEMESTER II
M.Sc MICROBIOLOGY
SYLLABUS EFFECTIVE FROM: JUNE-2020-21

PG02EMIC04: Medical Microbiology

Course Objectives:

The objective of this course is to make the students understand various attributes which make the microbes pathogenic or disease-causing, the emergence of newer pathogens with relevance to India and the various tools for their local or global spread. The students would also learn the mechanisms of resistance of bacteria to antibiotics and role of newer vaccines in controlling infectious diseases. The course would also enable students to describe the diagnostic methods and automated equipment which may be used for diagnosis of diseases caused by microorganisms.

Course Learning Outcomes:

Unit 1: Understands infection, its types and various host pathogen interaction. The students will be able to know the operation and the mechanisms which underlie the immune response to understand the phenomena like host defence. Useful to study various tools available to work on epidemiology.

Unit 2: Will gain in depth knowledge of Morphology, Cultural Characteristics, Antigenic structures, Pathogenesis, Laboratory Diagnosis of certain prominent and newer disease-causing bacteria.

Unit 3: Will get the information of different significant viral diseases, their characteristics, pathogenicity, antigenic properties, diagnosis and its preventive and control measures.

Unit 4: Understands different fungal and protozoal infections, their life cycles, and pathogenesis. Also useful to study and evaluate preventive and control mechanisms.

Contents:

Unit-I Basics in Medical Microbiology

Host pathogen interaction: Definitions - Infection, Invasion, Pathogen, Pathogenicity, Virulence, Toxigenicity, Carriers and their types, Opportunistic infections, Nosocomial infections. Transmission of infection, Pathophysiological effects of LPS

Nonspecific host defences, virulence factors, normal flora and gnotobiology Epidemiology: Infectious diseases, disease cycle, epidemiological methods, diagnostic principles, control, prevention, antimicrobial therapy.

Unit-II Bacterial Diseases

Morphology, Cultural Characteristics, Antigenic structures, Pathogenesis, Laboratory Diagnosis of following bacteria: *Staphylococcus*, *Streptococcus* including *Pneumococcus*, *Corynebacterium*, *Clostridium*, *Mycobacteria*, *E. coli*, *Salmonella*, *Shigella*, *Spirochaetes*, *Neisseria*

Unit-III Viral Diseases

The Nature and classification of viruses, Morphology: virus structure and Virus replication.

General properties, diseases caused, lab diagnosis and prevention of Herpes (HSV), Hepatitis (HAV & HAB), Picorna (Polio virus), Orthomyxo (Influenza), Paramyxo (Mumps and Measles), Rabdo (Rabies), Ebola, Zika and HIV virus.

Viral vaccines and antiviral agents.

Unit-IV Fungal and Protozoal Diseases

Fungal Morphology, diseases caused and lab diagnosis of:

Opportunistic fungi – *Candida and Aspergillus*

Fungi causing Cutaneous mycoses- *Dermatophytes*

Subcutaneous mycoses - *Mycetoma*

Systemic mycoses- *Histoplasma*

Protozoal Morphology, life cycle, laboratory diagnosis of following parasites

Parasites: *Entamoeba, Giardia, Leishmania, Plasmodium*

References:

1. Textbook of Microbiology by Surinder Kumar
2. Medical Parasitology by R. Karyakarte.
3. P. B. Godkar. Text Books of Medical Laboratory Technology
4. Anathanarayana & Panikar – A Text Book of Medical Microbiology
5. P. Chakraborty- A Text Book of Microbiology
6. Chatterjee, KD – Parasitology
7. Danial Greenwood et al, Medical Microbiology, A guide to Microbial Infections, Pathogenesis, Immunity, Laboratory Diagnosis and control.
8. Jagdish Chander, Textbook of medical mycology.
9. Teri Shores- Understanding Viruses.
10. Biswas SB and Biswas A: An Introduction to Viruses.

PG02CMIC04: Practicals based on PG02CMIC01 and PG02CMIC02

List of Practicals

1. Optimization of centrifugation for separation of cells
2. Measurement of growth by various methods (Absorbance, SPC, Direct count, Wet weight, Dry weight, Indirect method)
3. Determination of KLa by sulfite oxidation method
4. Demonstration of laboratory scale fermenter
5. Production of ethanol by yeast cells
6. Production of penicillin and its recovery
7. Recovery of citric acid
8. Partial purification of proteins by precipitation
9. Conjugation in *E. coli*.
10. Transduction in *E. coli*
11. Transposon assay
12. β -galactosidase induction and assay
13. Isolation and enumeration of bacteriophage
14. Demonstration of Lysogeny

PG02CMIC05: Practicals based on PG02CMIC03 and PG02EMIC01

List of Practicals

1. To perform total WBC count using Haemocytometer
2. To Perform Differential Leukocyte count
3. To learn the technique of Ouchterlony Double Diffusion
4. To learn the technique of Radial Immunodiffusion
5. To learn the technique of Immunoelectrophoresis
6. To perform sandwich Dot ELISA test for antigen
7. To learn the technique of latex -agglutination
8. To separate lymphocytes by density gradient method
9. To convert ungrouped data in to grouped data using Sturge's formula.
10. To study representation of data by one dimensional diagram.
11. To study representation of data by two dimensional diagram.
12. To study representation of data by means of graphs. (Histogram & frequency polygon).
13. To study the data representation by graphs (Frequency polygon & frequency curve).
14. To study how to calculate descriptive statistics for the given data. (Mean mode, median, standard deviation and mean deviation).
15. To study the concept of permutation and combination in practical counting problems.
16. To study the concept of normal distribution and apply it to practical problems.

17. To study the concept of estimation (point estimation and interval estimation).
18. To apply the concept of skewness in the field of biosciences.
19. To apply the concept of F- test for biological problems.
20. To apply the concept of χ^2 – test for biological problems.