# SAVITRIBAI PHULE PUNE UNIVERSITY

# (Formerly University of Pune)

# B. Sc. Degree Course in MICROBIOLOGY

# **Choice Based Credit System [CBCS]2019 Pattern**

Syllabus for Third Year

(To be implemented from Academic Year 2021-22)

Board of Studies (Microbiology)

Savitribai Phule Pune University [SPPU] Pune-411007

# **GENERAL INFORMATION**

## **Eligibility at third year B. Sc. Microbiology:**

Student shall clear all First Year B. Sc. Microbiology courses and satisfactorily keep terms of Second Year of B. Sc. with Microbiology as one of the subjects.

**Course Structure**: T. Y. B. Sc. Microbiology course includes 12 theory papers (DSEC-Discipline Specific Elective Course), 06 practical courses and 04 skill enhanced courses (SEC). The 06 theory papers, 03 practical courses and 02 skill enhanced courses (SEC) will be taught in semester V and the remaining 06 theory papers, 03 practical courses and 02 skill enhanced courses (SEC) will be taught in semester VI. The examination will be held semester-wise for theory and practical papers.

## Note:

- i. Each lecture (L) will be of 50 minutes.
- ii. Each practical of 4 hours 20 minutes and 12 practical sessions per semester
- iii. 12 weeks for teaching 03 weeks for evaluation of students (theory as well as practical).
- iv. For details refer UG rules and regulations (CBCS for Science program under Science andTechnology) published on SPPU website.

## **Evaluation Pattern (As per CBCS rules, SPPU 2019 Pattern)**

- 1. Each theory and practical course carry 50 marks equivalent to 2 credits.
- 2. Each course will be evaluated with Continuous Assessment (CA) and University Assessment (UA) mechanism.
- 3. Continuous assessment shall be of 15 marks (30%) while university Evaluation shall be of 35 marks (70%).
- 4. To pass each course, a student has to secure 40% mark in continuous assessment as well as university assessment i.e. 6 marks in continuous assessment and 14 marks in university assessment for the respective course.
- 5. For Continuous Assessment (internal assessment) minimum two tests per paper must be organized, of which one must be written test of 10 marks. 6. Method of assessment for internal exams: Viva-Voce, Project, survey, field visits, tutorials, assignments, group discussion, etc.

2.2 Mandatory Credit courses for award of B.Sc. Degree:

**In addition to the compulsory credits of 132, the student has to earn additional 8 credits** from following groups by taking/participating/conducting respective activities.

Courses in Group I are compulsory.

The student can earn maximum 04 credits from an individual group from Group 2 to Group -9.

These extra credits will not be considered for GPA calculation, however these are mandatory for the completion and award of B. Sc. Degree.

Group 1:	Physical Education (at F. Y. B. Sc. Sem. I) -01 credit
	Physical Education (at F. Y. B. Sc. Sem. II) - 01 credit
(Note: Grou	p I is compulsory for all the students as stated above.)
Group 2:	Sport representation at College level - 01 credit
	Sport representation at University/Statelevel - 02 credits
Group 3:	National Social Service Scheme (participation in Camp): 01 credits
	N.C.C.(with participation in annual camp) -01 credit
	N. C. C. (with B certificate/C certificate award)- 02 credits
cenve courses ;	N.S.S./N.C.C. Republic day parade participation - 04 credits
Group 4:	Avishkar participation; Extension activity participation, Cultural
	activity participation -01 credit
	Avishkar selection at University level - 02 credits
	Avishkar winner at state level - 04 credits
Group 5:	Research paper presentation at State/National level - 01credits
	Research paper presentation at International level - 02 credits
Group 6:	Participation in Summer school/programme; Short term course (not
i vitersvinU dos	less than 1-week duration) - 03 credit.
Group 7:	Scientific Survey, Societal survey, - 02 credits.
Group 8:	Field Visits; Study Tours; Industrial Visits; Participation in curricular/
	cocurricular competitions- 01 Credit.
Group 9:	Online certificate Courses /MOOC Courses/ Career Advancement
	Course up to 04 credits (Minimum 10 Hrs. / credit)

#### **T. Y. B. Sc.**

#### Equivalences for the New Courses (w. e. f. from 2021-22) with Old Courses (2013 Pattern) in Microbiology T. Y. B. Sc. Microbiology Semester - V

Theory/ Practical/ Skill Enhancement		Old Course Semester-III		New Course Semester-V (CBCS 2019 Pattern)
	Course Number	Course Title	Course Number	Course Title
Discipline	MB 331	Medical Microbiology-I	MB 351	Medical Microbiology-I
Specific Elective	MB 334	Immunology-I	MB 352	Immunology-I
Course (DSEC)	MB 333	Enzymology	MB 353	Enzymology
Theory	MB 332	Genetics and Molecular Biology-I	MB 354	Genetics
	MB 335	Fermentation Technology -I	MB 355	Fermentation Technology-I
	MB 346	Agricultural and Environmental Microbiology	MB 356	Agricultural Microbiology
Discipline Specific Elective Course (DSEC)	MB 349	Practical Course-III Diagnostic Microbiology and Immunology	MB 357	Practical course-I based on: MB 351 Medical Microbiology-I MB 352 Immunology I
Practical	MB 348	Practical Course-II Biochemistry and Genetics	MB 358	Practical course-II based on MB 353 Enzymology MB 354 Genetics
	MB 347	Practical Course I Applied Microbiology	MB 359	Practical course-III based on: MB 355 Fermentation Technology-I MB 356Agricultural Microbiology
Skill Enhancement course	-	-	MB 3510	Marine Microbiology
	-	-	MB 3511	Dairy Microbiology

#### **T. Y. B. Sc.**

## Equivalences for the New Courses (w. e. f. from 2021-22)

## With old Courses (2013 Pattern) in Microbiology

## T. Y. B. Sc. MicrobiologySemester-VI

Theory/ Practical/ Skill Enhancement		Old Course Semester-III	New Course Semester-VI (CBCS 2019 Pattern)		
	Course Number	Course Title	Course Number	Course Title	
Discipline Specific	MB 341	Medical Microbiology-II	MB 361	Medical Microbiology II	
Elective Course	MB 344	Immunology-II	MB 362	Immunology II	
(DSEC) Theory	MB 343	Metabolism	MB 363	Metabolism	
	MB 342	Genetics and Molecular Biology-II	MB 364	Molecular Biology	
	MB 345	Fermentation Technology-II	MB 365	Fermentation Technology II	
	MB 346	Food and Dairy Microbiology	MB 366	Food Microbiology	
Discipline Specific Elective Course	MB 349	Practical course-III Diagnostic Microbiology and Immunology	MB 367	Practical course-I. Based on: MB 361 Medical Microbiology II and MB 362 Immunology II	
(DSEC) Practical	MB 348	Practical course-II Biochemistry and Genetics	MB 368	Practical course-II. Based on: MB 363 Metabolism and MB 364 Molecular Biology	
	MB 347	Practical course-I Applied Microbiology	MB 369	Practical course III. Based on: MB 365 Fermentation technology-II and MB 366 Food Microbiology	
Skill Enhancement	-	-	MB 3610	Waste management	
course	-	-	MB 3611	Nano biotechnology	

## **Evaluation Pattern**

# T. Y. B. Sc. Microbiology

	Courses						
	Seme	ster-V			Semeste	er-VI	
Paper	Course Title	Internal examination Marks	University examination Marks	Paper	Course Title	Internal Exam Marks	University examination Marks
MB 351	Medical Microbiology I	15	35	MB 361	Medical Microbiology II	15	35
MB 352	Immunology I	15	35	MB 362	Immunology II	15	35
MB 353	Enzymology	15	35	MB 363	Metabolism	15	35
MB 354	Genetics	15	35	MB 364	Molecular Biology	15	35
MB 355	Fermentation technology I	15	35	MB 365	Fermentation Technology II	15	35
MB 356	Agricultural Microbiology	15	35	MB 366	Food Microbiology	15	35
MB 357	Practical course-I Based on: MB351 and MB 352	15	35	MB 367	Practical course I Based on: MB 361 and MB 362	15	35
MB 358	Practical course- II Based on MB 353 and MB 354	15	35	MB 368	Practical course II Based on: MB 363 and MB 364	15	35
MB 359	Practical course-III Based on:MB 355 and MB 356	15	35	MB 369	Practical course III Based on: MB 365 Fermentation technology II, MB 366 Food Microbiology	15	35
MB 3510	Marine Microbiology	15	35	MB 3510	Waste Management	15	35
MB 3511	Dairy Microbiology	15	35	MB 3511	Nano biotechnology	15	35

## DSEC-MB 351: Medical Microbiology- I

## [2 Credits; 36 Lectures]

## [1 credit=15hrs x 60 mins = 900mins/50mins=18 lectures]

## **Course Outcomes:**

- Understand the human anatomy, pathogens associated with diseases.
- Acquire knowledge of principles underlying establishment of pathogens in human body.
- Comprehend of pathogenesis of specific pathogens causing microbial diseases.
- Assess epidemiological patterns of microbial disease transmission as various modes, intensity at local and global level.
- Gain Knowledge principles of chemotherapy of microbial diseases and development of drug resistance among pathogens and strategies to mitigate.
- Develop identification systems for microbial disease diagnosis, disease treatment and prevention measures.

Credit No.	Topics	No. of Lectures
	Introduction to infectious diseases and Epidemiology	18
	1. Introduction to infectious diseases of following human body	
	systems: (Brief anatomy and Physiology, Diseases, Pathogens, common	
	symptoms)	
	a. Respiratory system	2
	b. Gastrointestinal system and liver	2
	c. Urogenital system	2
Credit I	d. Central nervous system	2
Creuit I	2. Epidemiology:	
	a. Case control and cohort studies – Study design and application	2
	b. Principle and methods – Clinical trials of drugs and vaccines	3
	(Randomized control trials Concurrent parallel and cross-over trials)	
	c. Epidemiology of infectious diseases	
	i. Sources and Reservoirs of Infection	1
	ii. Modes of Transmission of Infections	1
	iii. Disease Prevention and Control Measures, Vaccine-	3
	preventablebacterial diseases and nonvaccine-preventable	
	bacterial diseases	

	Study of bacterial pathogens:	18
	<b>3. Study of following groups of bacterial pathogens:</b> (With respect	
	to- Classification and Biochemical characters, Antigenic structure,	
	Viability characteristics, Pathogenicity, Pathogenesis, Symptoms,	
Credit	Laboratory diagnosis, Epidemiology, Prophylaxis and Chemotherapy):	
II	a. Salmonella, Vibrio	2
	b. Streptococcus pneumoniae, Streptococcus pyogenes,	4
	Neisseria meningitidis and Neisseria gonorrhoeae	
	c. Pseudomonas aeruginosa	2
	d. Treponema, Leptospira	2
	e. Clostridium tetani	2
	f. Mycobacterium tuberculosis and Mycobacterium leprae	4
	g. Orientia tsutsugamushi and Rickettsia rickettsii	2

## DSEC-MB 361: Medical Microbiology II

## [2 Credits; 36 Lectures]

## [1 credit=15hrs x 60 mins = 900mins/50mins=18 lectures]

Credit No.	Topics	No. of lectures
	Chemotherapy	18
	1. Routes of drug administration.	1
	2. Mode of action of antimicrobial agents on:	
	a. Bacteria:	
	i. Cell wall: Beta lactams:1 <sup>st</sup> to 6 <sup>th</sup> Generation- e.g. Meropenem,	2
	Imipenem, Piperacillin, Tazobactam	
	ii. Cell membrane: Polymyxin	1
Credit I	iii.Protein synthesis: Streptomycin, Tetracycline	1
	iv.Nucleic acids: Fluroquinolones, Rifamycin	1
	v. Enzyme inhibitors: Trimethoprim, Sulfomethaxazole	1
	b.Fungi: Griseofulvin, Amphotericin B, Anidulafungin, Vericonazole	3
	c. Viruses: Acyclovir, Oseltamivir, Remdecivir	1
	d.Protozoa: Metronidazole, Chloroquine	1

	3. Mechanisms of drug resistance on:	
	a. Genetic basis:	3
	i. Mutations in gene(s)	
	ii. Acquisition of foreign DNA coding for resistance	
	determinantsthrough horizontal gene transfer.	
	b. Mechanisms of drug resistance by:	3
	i. Limiting uptake of a drug.	
	ii. Modification of a drug target.	
	iii. Inactivation of a drug.	
	iv. Active efflux of a drug.	
	Human and Animal Viruses, Fungal and Protozoal Pathogens	18
	4. Introduction to cultivation of viruses	2
	5. Study of following groups of viral pathogens:	
	a. Human viruses (with respect to – Virion, Characteristics, Viability	
	characteristics, Pathogenicity, Pathogenesis, Symptoms, Laboratory	
	diagnosis including serological diagnosis, Epidemiology, Prophylaxis	
	and Chemotherapy):	
	i. Respiratory Viruses: Influenza Virus, Corona Virus	2
	ii. Hemorrhagic Virus: Dengue	2
	iii. Hepatic Virus: Hepatitis A Virus	1
	iv. Gastrointestinal Virus: Rotavirus	1
Credit	v. Cutaneous Viruses: Human papillomavirus	1
II	vi. Neurological Viruses: Japanese Encephalitis Virus	1
	b. Animal Viruses: FMD Virus and Rinderpest Virus	2
	6. Study of following groups of parasites (with respect to -	
	Classification, Lifecycle, Morphological characteristics, Viability	
	characteristics, Pathogenicity, Pathogenesis, Symptoms, Laboratory	
	diagnosis (Serological diagnosis wherever applicable), Epidemiology,	
	Prophylaxis and Chemotherapy):	
	a. Plasmodium	2
	b. Entamoeba	1
	7. Study of following groups of yeast and fungal pathogens (With	
	respect to - Morphological and cultural characteristics, Classification,	
	Pathogenicity, Pathogenesis, Symptoms, Laboratory diagnosis,	

1

1

1

Epidemiology, Prophylaxis and Chemotherapy)

- a. Aspergillus species (Pathogenic)
- b. Cryptococcus neoformans
- c. Histoplasma capsulatum

## References: MB 351 Medical Microbiology-I and MB 361 Medical Microbiology- II

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- Mayers D. L., Sobel J.D., Ouellette M., Kaye K.S. and Marchaim D. (Eds.) (2017). Antimicrobial Drug Resistance: Mechanisms of Drug Resistance. Volume 1. Edition 2. Springer. ISBN 978-3-319-46718-4
- 13. Mayers D. L., Sobel J.D., Ouellette M., Kaye K.S. and Marchaim D. (Eds.) (2017).

Antimicrobial Drug Resistance: Clinical and Epidemiological Aspects. Volume 2. Edition 2. Springer. ISBN 978-3-319-47266-9

- Mukherjee K. L. and Ghosh S. (2010). Medical Laboratory Technology. Volume I: Procedure Manual for Routine Diagnostic Tests. 2nd edition. McGraw Hill Education (India) Private Limited. ISBN-13: 978-1259061233
- Mukherjee K. L. and Ghosh S. (2010). Medical Laboratory Technology. Volume II: Procedure Manual for Routine Diagnostic Tests. 2nd edition. McGraw Hill Education (India) Private Limited. ISBN-13: 978-1259061240
- Mukherjee K. L. and Ghosh S. (2010). Medical Laboratory Technology. Volume III: Procedure Manual for Routine Diagnostic Tests. 2nd edition. McGraw Hill Education (India) Private Limited. ISBN-13: 978-1259061257
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#### Links:

- 1. https://www.who.int/travel-advice/disease-information
- 2. https://Microbenotes.Com/Remdesivir/#Mechanism-Of-Action-Of-Remdesivir
- 3. Aspergillus https://www.cdc.gov/fungal/diseases/aspergillosis/index.html
- 4. Histoplasma capsulatum https://www.cdc.gov/fungal/diseases/histoplasmosis/
- 5. Cryptococcus neoformans www.cdc.gov/fungal/diseases/cryptococcosis-neoformans/

## DSEC-MB-352 Immunology- I

## [2 Credits; 36 Lectures]

#### [1 credit=15hrs x 60 mins = 900mins/50mins=18 lectures]

#### **Course Outcomes**

- Understand immune system structure, composition, function and comparison of different types of immunity.
- Acquire knowledge about antigens, Recognition of pathogens; antigen processing and presentation; Immunity to infection and pathological consequences of immunodeficiencies.
- To learn the applications of Immunology in monoclonal antibodies, vaccines production and Immunotherapy.
- Understand abnormal working of Immune system in hypersensitivity, auto immune diseases, immune tolerance and transplantation immunology.
- To develop strategies for Diagnosis of diseases based on antigen and antibody reactions with emphasis on prevailing communicable diseases.

Credit No.	Topics	No. of Lectures
	Organs of immune system, Innate immunity, Antigen and Immunoglobulins	18
	1. Organs of immune system:	
	a. Primary lymphoid organs (Thymus and Bone Marrow):	2
	Thymus – structure, thymic education (positive and negative selection)	
	Bone marrow –Structure and Negative selection	
	b. Secondary lymphoid organs – structure and function of spleen and	2
	lymph node, mucous associated lymphoid tissue, lymphatic system and lymph	
Credit I	circulation	
Crean I	2. Innate immunity: Non-specific mechanisms of defense: Second line of	
	defense:	
	a. Humoral components: Defensins, pattern recognition proteins (PRP) and	1
	pathogen associated molecular patterns (PAMPs), complement, kinins,	
	andacute phase reactants.	
	b. Cellular components: Phagocytic cells - PMNL, macrophages (reticulo-	1
	endothelial cell system) and dendritic cells	
	c. Phagocytosis (oxygen dependent and independent systems), Complement	5
	activation (Classical, Alternative and lectin pathway), Inflammation	
	(cardinal signs, mediators, vascular and cellular changes, role of Toll-like	
	receptors)	

	3. Anti	igen:		
	a.	Factors affecting immunogenecity	1	
	b.	Antigenic determinants, haptens and cross-reactivity, Carrier, Adjuvants	1	
	c.	Types of antigens: Thymus-dependent and thymus-independent		
		antigens, Synthetic antigens, Soluble and particulate antigens,	1	
		Autoantigens, Isoantigens		
	4. Imn	nunoglobulins:	2	
	a.	Characteristic of domain structure, functions of light and heavy chain	2	
		domainsand antigenic nature of immunoglobulin molecules	2	
	b.	Molecular basis of antibody diversity (kappa, lambda and heavy chain)	2	
	Antige	n- Antibody Interactions, Major Histocompatibility Complex,	18	
	Transj	plantation and Immunity and Hybridoma Technology and		
	Monoc	clonal Antibodies		
	5. Anti	gen- Antibody Interactions:		
	A. Pri	nciples of interactions: Antibody affinity and avidity, ratio of antigen	2	
	an	tibody, lattice hypothesis and two stage theory, antigen-antibody		
	rea	actionkinetics (dialysis equilibrium experiment)		
	B. Visualization of antigen antibody complexes:			
	a.	Precipitation reactions: in fluid and in gel, immunoelectrophoresis	1	
	b.	Agglutination reactions: hemagglutination, bacterial agglutination,	1	
		passive agglutination and agglutination-inhibition		
Credit	c.	Immunofluorescence techniques: direct and indirect, fluorescence-	2	
Π		activated cell sorting (FACS)		
п	d.	Enzyme-linked immunosorbent assay (ELISA), biotin-avidin system and	2	
		enzyme-linked immune absorbent spot (ELISpot) assay		
	e.	Radioimmunoassay RIA	1	
	6. Maj	or Histocompatibility Complex:		
	a.	Structure of MHC in man and mouse	1	
	b.	Structure and functions of MHC class-I and class-II molecules	1	
	c.	MHC antigen typing (microcytotoxicity and mixed lymphocyte reaction)	1	
	<b>7.</b> Tra	nsplantation and Immunity;		
	a.	Types of Grafts, Allograft rejection mechanisms	2	
	b.	Prevention of allograft rejection	1	

#### **T. Y. B. Sc.**

8. Hybridoma Technology and Monoclonal Antibodies;	
a. Preparation, HAT selection and propagation of hybridomas	2
secretingmonoclonal antibodies	
b. Applications of monoclonal antibodies	1

## Semester VI

## DSEC-MB 362 Immunology- II

## [2 Credits; 36 Lectures]

## [1 credit=15hrs x 60 mins = 900mins/50mins=18 lectures]

Credits	Topics	No. of
		Lectures
	Cytokines, Adaptive / Acquired Immunity, Hypersensitivity,	18
	Autoimmunity and Autoimmune diseases and Immunodeficiency	
	1. Cytokines:	
	a. Concept- Cytokines, lymphokines, monokines, interleukines,	1
	chemokines, interferons and tumor necrosis factor	
	b. Properties, Attributes and biological functions of cytokines	2
	2. Adaptive / Acquired Immunity (Third line of defense):	
	A. Humoral Immune Response	
	i. Primary and secondary response kinetics, significance in	2
Credit	vaccination programs	
Ι	ii. Response of secondary lymphoid organs to antigen	1
	iii. Antigen processing and presentation (Major	5
	Histocompability class I and class II restriction pathways),	
	cell-cell interactions and adhesion molecules, response to	
	super-antigens, role of cytokines in activation and	
	differentiation of B-cells	
	B. Cell Mediated Immune Response	
	i. Activation and differentiation of T cells, role of cytokines in	2
	activation	
	ii. Mechanism of Cytotoxic T lymphocytes (CTL) mediated	3
	cytotoxicity, Antibody-dependent cellular cytotoxicity (ADCC)	
	iii.Significance of Cell Mediated Immune Response (CMI)	1
	iv. Immune response against tumors and foreign transplanted cells	1

	Hypersensitivity, Autoimmunity and Autoimmune diseases and	18
	Immunodeficiency	
	3. Hypersensitivity	
Credit	a. General principles of different types of hypersensitivity reactions	2
II	b. Gell and Coomb's classification of hypersensitivity -	
	mechanism with examples for type I (Immediate), II, III and IV	5
	(delayed)	
	4. Autoimmunity and Autoimmune diseases:	
	a. Immunological tolerance	1
	b. Types of autoimmune diseases	1
	c. Factors contributing development of autoimmune diseases	1
	d. Immunopathological mechanisms	1
	e. Diagnosis and treatment of autoimmune diseases: Myasthenia	2
	gravisand Rheumatoid arthritis	
	f. Therapeutic immunosuppression for autoimmunity	1
	5.Immunodeficiency:	
	i. Complement deficiencies	2
	ii. Introduction to congenital immunodeficiency disorders:	2
	Common Variable Immune Deficiency (CVID) and acquired	
	immunodeficiency: Immune mechanisms in AIDS	

## References: MB-352 Immunology- I and MB 362- Immunology-II

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#### **CBCS: 2019 Pattern**

#### **T. Y. B. Sc.**

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- Zanetti M. (2005). The role of cathelicidins in the innate host defense of mammals.Curr. Issues Mol. Biol. 7:179–196.

## **DSEC-MB 353: Enzymology**

## [2 Credits; 36 Lectures]

#### [1 credit=15hrs x 60 mins = 900mins/50mins=18 lectures]

#### **Course Outcomes**

- To understand methods of active site determination, role of enzymes and its cofactors in microbial physiology.
- To learn to perform enzyme assay, purification and quantification of enzymes activity, enzyme kinetics in terms of initial, final velocity, mathematical expression of enzyme kinetic parameters.
- To correlate regulation of metabolism at enzymatic levels and apply, methodology for commercial applications of enzymes
- To learn mechanisms of transport of solutes across the membrane
- To get acquainted with mechanism of biosynthesis and degradation of bio molecules
- To comprehend basic concept of autotrophic mode of metabolism of prokaryotes

Credit	Topics	No. of
No.		lectures
	Enzymes:	18
	1. Structure of enzymes:	
	a. Methods to determine amino acid residues at active site	3
	(Physical method e.g. x-ray crystallography and chemical	
	methods such as trapping of ES complex, use of inhibitors,	
	use of pseudo-substrate, change of pH)	
	b. Role of vitamins in metabolism:	2
	Occurrence, Structure and Biochemical functions of the following:	
Credit	i. Thiamine (Vitamin B1) and Thiamine Pyrophosphate	
Ι	ii. Vitamin D	
	2. Enzyme assays:	
	a. Principles of enzyme assays and calculation of	1
	enzyme unit, specific activity	
	b. Enzymes assays with examples by:	2
	i. Spectrophotometric methods	
	ii. Radioisotope assay	

3. Principles and Methods of Enzyme purification:	
a. Methods of cell fractionation	2
b. Principles and methods of enzyme purification:	
i. Based on molecular size	2
ii. Based on charge	2
iii. Based on solubility differences	2
iv. Based on specific binding property and selective adsorption	1
c. Construction of enzyme purification chart	1
Enzyme Kinetics, metabolic regulation and Immobilized Enzymes:	18
4. Enzyme Kinetics:	
a. Concept and use of initial velocity	2
b. Michaelis Menton equation for the initial velocity of single	5
substrate enzyme catalyzed reaction. Brigg's Haldane	
modification of Michaelis Menton equation. Michaelis Menton	
plot, Lineweaver and Burk plot. Definition with significance of	
Km, Ks, Vmax	
5. Metabolic Regulations:	
a. Enzyme compartmentalization at cellular level	1
b. Allosteric enzymes	1
c. Feedback mechanisms	2
d. Covalently modified regulatory enzymes (Glycogenphosphorylase)	1
e. Proteolytic activation of zymogens	1
f. Isozymes - concept and examples	1
g. Multienzyme complex e.g. Pyruvate dehydrogenasecomplex(PDH)	1
6. Immobilization of enzymes:	3
Concept, methods of immobilization and applications	
	<ul> <li>a. Methods of cell fractionation</li> <li>b. Principles and methods of enzyme purification: <ol> <li>Based on molecular size</li> <li>Based on charge</li> <li>Based on solubility differences</li> <li>Based on specific binding property and selective adsorption</li> <li>c. Construction of enzyme purification chart</li> </ol> </li> <li>Enzyme Kinetics, metabolic regulation and Immobilized Enzymes: <ol> <li>Concept and use of initial velocity</li> <li>Michaelis Menton equation for the initial velocity of single substrate enzyme catalyzed reaction. Brigg's Haldane modification of Michaelis Menton equation. Michaelis Menton plot, Lineweaver and Burk plot. Definition with significance of Km, Ks, Vmax</li> </ol> </li> <li>5. Metabolic Regulations: <ol> <li>Enzyme compartmentalization at cellular level</li> <li>Allosteric enzymes</li> <li>Feedback mechanisms</li> <li>Covalently modified regulatory enzymes (Glycogenphosphorylase)</li> <li>Proteolytic activation of zymogens</li> <li>Isozymes - concept and examples</li> <li>Multienzyme complex e.g. Pyruvate dehydrogenasecomplex(PDH)</li> </ol> </li> </ul>

## DSEC-MB 363: Metabolism

## [2 Credits; 36 Lectures]

## [1 credit=15hrs x 60 mins = 900mins/50mins=18 lectures]

Credit	Topics	No of
No.	Membrane transport and Bioenergetics	lectures 18
	1. Membrane transport mechanisms:	6
	i. Passive transport - Diffusion, Osmosis, Facilitated transport	
	ii. Active transport - Active transport systems in bacteria	
	iii. Group translocation of sugars in bacteria	
	iv. Ionophores: Mechanism and examples	
	2. Bioenergetics:	
	i. Laws of thermodynamics- first and second law	1
	ii. Concepts of free energy, entropy, high energy compounds:	4
Credit I	Pyrophosphate, enolic phosphates, acyl phosphates, thioester	
	compounds, and guanidinium compounds	
	iii. Mitochondrial electron transport chain: components, arrangement	7
	of different components in the inner membrane, structure and	
	function of ATP synthatase, inhibitors and uncouplers of ETC and	
	oxidative phosphorylation, energetics of mitochondrial electron	
	transport chain	
	Metabolic pathways and Autotrophy	18
	3. Biosynthesis and Degradation:	
	a. Chemistry, concept of polymerization of macromolecules:	6
	Polysaccharides. (Starch, and peptidoglycan) and Lipids (Fatty acids,	
	triglycerides and phospholipids)	
	b. Degradation of macromolecules - Polysaccharides (starch), Lipids	6
Credit	(fatty acids oxidation e.g. $\beta$ oxidation), Proteins (urea cycle)	
II	4. Bacterial Photosynthesis: Photosynthetic bacteria with reference to	
	photosynthetic apparatus, energy generation, and CO <sub>2</sub> fixation	
	a. Cyanobacteria,	2
	b. Purple bacteria	2
	5 Chemolithotrophy:	2
	Concept and one example, Iron oxidizing bacteria	
		10100

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## **DSEC -MB 354: Genetics**

## [2 Credits; 36 Lectures]

#### [1 credit=15hrs x 60 mins = 900mins/50mins=18 lectures]

#### **Course Outcomes**

- To exhibit a knowledge base in Genetics and Molecular Biology
- To understand the central dogma of Molecular Biology
- To construct genetic map of bacteria and fungi
- To get introduced to concept of recombination and bacteriophage Genetics
- To understand the concept cloning in bacteria
- To demonstrate the knowledge of common and advanced laboratory practices in Molecular Biology

Credit	Topics	No. of
No.		lectures
	DNA Replication and Gene Expression	18
	1. Process of prokaryotic DNA replication	4
	a. Single replicon	
	b. Bidirectional movement of replication fork	
	c. Ori C	
	d. Pre-priming and Priming reaction.	
	e. DNA polymerases, DNA synthesis of leading, lagging strand	
	Okazaki fragments.	
	f. Termination- Ter sequence, Tus protein	
Credit	2.Prokaryotic and Eukaryotic Transcription	3
I	Transcription in Prokaryotes	
	a. Structure of promoter	
	b. Structure and function of RNA polymerase	
	c. Steps of transcription: Initiation, Elongation and termination	
	Transcription in eukaryotes with respect to protein coding Gene:	4
	a. Promoter, promoter proximal elements and enhancers	
	b. Transcription regulatory proteins	
	c. RNA polymerases	
	d. Steps in transcription: Initiation, Elongation, Termination	
	e. Post transcriptional modifications: 5' capping, 3'	
	polyadenylation and introduction to RNA splicing	

	3. Regulation of transcription:	2
	Concept and components of operon:	_
	Lac operon: Inducible operon	
	4. Translation in prokaryotes and eukaryotes	5
	a. Structure and role of m-RNA, t-RNA and Ribosomes in	C
	Translation	
	b. Role of Aminoacyl t-RNA synthetase in translation	
	c. Steps in translation: Initiation, elongation, translocation and	
	termination of protein synthesis	
	d. Salient features of Eukaryotic translation	
	Gene transfer and mapping techniques	18
	5. Gene transfer by Transformation	4
	a. Discovery of Transformation	
	<ul><li>b. Natural transformation Systems-</li></ul>	
	Streptococcus pneumoniae and Haemophilus influenzae.	
	c. Factors affecting transformation	
	i. Competence development	
	ii. Size of DNA	
	iii. Concentration of DNA	
Credit	6. Gene transfer by Conjugation	4
	a. Discovery of Conjugation,	+
II	<ul> <li>b. Properties of F plasmid, F<sup>+</sup>, F<sup>-</sup>, Hfr and F' strains</li> </ul>	
	c. Process of conjugation between $F^+$ and $F^-$ , Hfr and $F^-$ , F 'and F-	
	7. Gene transfer by Transduction	4
	a. Discovery of Transduction	
	b. Generalized transduction mediated by P22	
	c. Specialized transduction mediated by lambda phage	
	8. An introduction to Gene mapping	6
	a. Gene linkage and concept of genetic recombination	
	b. Recombination mapping: Map unit, recombination frequency	
	c. Mapping of genes by co-transformation	
	d. Mapping of genes by co-transduction	
	e. Mapping by interrupted mating experiment	
	f. Numerical problems based on co-transformation, co-	
	transductionand interrupted mating	
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## **DSEC -MB-364: Molecular Biology**

## [2 Credits; 36 Lectures]

## [1 credit=15hrs x 60 mins = 900mins/50mins=18 lectures]

Credit No	Topics	No. of lectures
	Genetic Recombination and Bacteriophage Genetics.	18
	1. Gene linkage and crossing over	9
	a. Mendel's laws: Eukaryotic Cell cycle, Mitosis, Meiosis	
	b. Holliday model for Homologous recombination, Role of Rec	
	and Ruvproteins	
Credit	c. Genetic mapping by Tetrad analysis in N. crassa (Numerical	
I	Calculationsusing PD, TT and NPD)	
	d. Genetic Mapping by Parasexual cycle in A. nidulans	
	2. Bacteriophage Genetics	9
	a. Lytic cycle: Virulent phages, T-series phages, Concept and	
	formation ofplaque, Lysogenic cycle: Temperate phage (λphage)	
	b. Bacteriophage mutants: Plaque morphology (r type), Host	
	range, Conditional lethal mutants (Ts and Am)	
	c. Concept of Genetic Complementation and Cis-trans test of genetic	
	function.(Intergenic- rII locus of T4 phage, Mechanism of Intragenic	
	complementation.)	
	d. Fine structure mapping of rII locus of T4 phage using Benzer's spot	
	testsand deletion mapping	
	DNA damage and repair mechanisms, Recombinant DNA technology	18
	3. DNA damage and Repair mechanisms	5
	a. DNA damage by hydrolysis, deamination, alkylation,	
Credit	oxidation, Radiation (X rays and UV rays)	
II	b. DNA repair by Photo reactivation	
	c. DNA repair by Mismatch repair mechanism	
	d. DNA repair by Excision repair mechanisms (BER/NER)	

4. Rec	ombinant DNA Technology Tools and basics of recombinant DNA	10
techn	ology	
a.	Introduction to recombinant DNA technology	
b.	Restriction enzymes: Concept, Nomenclature, properties and types	
	withexamples (Eco R1, Sma I, Pst I).	
c.	Vectors: Features of an ideal vector	
	i. Plasmids: pBR322	
	ii. Bacteriophage vectors: Lambda	
	iii. Cosmids	
	iv. High capacity vectors: YACs, BACs	
	v. Expression vectors	
d.	Joining of DNA molecules- DNA Ligases (E. coliand T4 phage), Use	
	of Linker / Adaptor / Homopolymer tailing	
e.	Methods to transfer recombinant DNA into bacterial host cells	
	(Physical – Electroporation, Gene gun, Chemical –CaCl2 mediated,	
	liposome mediated)	
f.	Methods of screening recombinants using selective markers and Blue-	
	whitescreening	
5. Mol	lecular techniques used in RDT	3
a.	Isolation of genomic DNA	
b.	Principle and methodology of Agarose gel electrophoresis and its	
	applications	
c.	Concept, Methodology and applications of Southern, Northern and	
	Western blotting	

## **References: MB 354 Genetics and MB 364 Molecular Biology**

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- National Academies Press: Introduction of Recombinant DNA-Engineered Organisms Into theEnvironment: Key Issues: <u>https://www.nap.edu/download/18907#</u>
- Guidelines and Handbook for Institutional Biosafety Committees (DBT, Govt. of India and BCIL):<u>https://thsti.res.in/pdf/IBG.pdf</u>
- 4. University of North Carolina's Biosafety Guidelines (Principles, Risk assessment, Biosafety levels, Guidelines):

https://ehs.unca.edu/laboratory-safety/biological-safety/ http://www.informatics.jax.org/silver/chapters/7-1.shtml

## DSEC -MB 355 Fermentation Technology-I

## [2 Credits; 36 Lectures]

#### [1 credit=15hrs x 60 mins = 900mins/50mins=18 lectures]

#### **Course Outcomes**

- To impart technical understanding of commercial fermentations.
- To apply classical, advanced strain improvement and isolation techniques for fermentationprocesses.
- To optimize and sterilize media used in fermentation industry for commercially economical and efficient fermentations.
- To recover the product using suitable methods and ensuring quality of the finished productby quality assurance tests.
- To acquaint fermentation economics, process patentability, process validation.
- To comprehend the large scale productions of commercially significant fermentation products of classical and recent significance.

Credit	Topics	No. of
No.		lectures
	Upstream processes of fermentations	18
	1. Strain Improvement:	
	a. Objectives of strain improvement	1
	b. Methods for strain improvement:	
	i. Types of mutants used in strain improvement (altered cell	1
Credit I	permeability mutants, auxotrophs, analogue resistant mutants, revertants)	
	ii. Selection of different types of mutants (replica plate method,	2
	filtration enrichment, penicillin enrichment method, gradient	
	platetechnique)	
	iii. Application of rDNA technology (significance, technique for	1
	commercial recombinant products like insulin)	
	2. Media optimization	
	a. Objectives of media optimization	1
	b. Methods of media optimization:	
	i. Classical approach – One factor at a time, Full factorial design	1
	ii. Plackett and Burman Design (with example) (Numerical	2
	problems of PBD can be discussed using software)	
	iii. Response Surface Methodology (RSM)	1

	3. Ste	rilization of Media:	
	a.	Methods of sterilization	1
	b.	Batch sterilization and Continuous sterilization (direct and	1
		indirectmethods)	
	с.	Concept and derivation of Del factor	1
	d.	Filter sterilization of liquid media	1
	4. Sca	le-up and Scale-down:	
	a.	Objectives of scale-up	1
	b.	Levels of fermentation (laboratory, pilot-plant and	1
		productionlevel – flowsheet to explain scale up)	
	с.	Criteria of scale-up for critical parameters [Aeration (kLa	1
		Volumetric Mass transfer coefficient), Agitation (P/V	
		ratio, $N_{Re}$ Reynolds number, $N_p$ Power number, $N_{Fr}$	
		Froudes number), Sterilization and broth rheology	
		(Newtonian and non Newtonian fluids - bacterial and	
		mycelia fungal fermentations)]	
	d.	Scale-down (example of any one commercial fermentation)	1
	Down	stream processing and Quality assurance of fermentation	18
	produ	icts	
	5. Dov	wnstream processing of fermentation products:	
	(meth	od, principle, types, examples of fermentations, factors	
	affect	ing, merits and demerits at large scale operation)	
	a.	Cell disruption methods	1
	b.	Filtration	1
Credit	с.	Centrifugation	1
II	d.	Liquid-liquid extraction	1
	e.	Distillation	1
	f.	Drying	1
	6. Qu	ality assurance of fermentation products (as per IP, USP)	
	a.	Methods of detection and Quantification of the fermentation	2
		product: physicochemical, biological and enzymatic methods	
	b.	Sterility testing (direct inoculation method, membrane	1
	b.	Sterility testing (direct inoculation method, membrane filtration method)	1

d.	Microbial limit test	1
e.	Pyrogen testing: Endotoxin detection (LAL test)	1
f.	Ames test and modified Ames test	1
g.	Toxicity testing (Acute toxicity)	1
h.	Shelf life determination	1
<b>7.</b> F	ermentation economics:	
a.	Contribution of various expense heads to a process (Recurring	1
	and nonrecurring expenditures) citing any suitable example.	
b.	Introduction to Intellectual Property Rights -	1
	Types of IPR (patenting in fermentation industry)	
c.	Concept of validation( significance of SOPs)	1

## DSEC - MB 365 Fermentation Technology – II

## [2 Credits; 36 Lectures]

## [1 credit=15hrs x 60 mins = 900mins/50mins=18 lectures]

Credit	Topics	No. of
No.		lectures
	Solid state and Submerged state fermentations and Large scale	18
	fermentations	
	1. Introduction to Solid State Fermentation and Submerged	1
Credit	Fermentation:	
I	Process, production strains, media, fermentor design, fermentation	
	conditions, applications, merits and demerits	
	2. Large scale production of (process with flow sheet, nature of the	
	product, production pathway, applications, production strains, media,	
	fermentation process, parameters, product recovery)	
	a. Primary Metabolites:	
	i. Vitamins (B12 and B2)	3
	ii. Amino acids - Glutamic acid, Lysine	3
	iii. Organic acids (Citric acid, Vinegar and Lactic acid)	4
	b. Secondary metabolites:	
	i. Bioethanol	1
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	ii. Alcoholic Beverages -	3
	a. Beer (Lagering, Maturation, Types of beer)	
	b. Wine (Aging, Malo-lactic acid fermentation, types of wine, wine	
	defects, comparison of white and red wine)	
	iii. Antibiotics [Penicillin (natural and semi synthetic) and	3
	Streptomycin]	
	Large scale production of enzymes, steroids, biomass based products,	18
	milk products, vaccines, immune sera and Modern trends in microbial	
	production	
	3. Enzymes	
	i. Amylase	1
	ii. Esterases	1
	iii. Proteases	1
	4. Microbial transformation of steroids	2
	5. Biomass based products:	
	i. Yeast: Baker's and Distiller's yeast	2
	ii. Probiotics: Lactobacillus sporogenes	1
	6. Milk products:	
	i. Cheese (Processed, soft, semi-hard, hard ripened types- bacterial and	2
	mold)	
	ii. Yogurt (plain, flavoured, fruit, sundae style. Stirred type, set type,	2
	probiotic yoghurt)	
Credit	7. Vaccines	
II	i. Polio – Inactivated Polio Vaccine, Oral Polio Vaccine	1
	ii. Tetanus – Tetanus toxoid (TT)	1
	iii. Rabies – HDCC, Chick embryo cell line, Vero cell line	1
	8. Immune sera	
	i. Anti tetanus serum (ATS)	1
l	ii. Anti rabitic serum (ARS)	1
	9. Modern trends in microbial production:	
	Biosurfactant and bioemulsifier	1

## References: MB 355 Fermentation Technology- I and MB 365 Fermentation Technology- II

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#### **Reference links:**

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- Large scale production of rabies vaccine: https://academic.oup.com/jimb/article-pdf/18/5/340/34773995/jimb0340.pdf.
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## DSEC - MB 356: Agricultural Microbiology

## [2 Credits; 36 Lectures]

#### [1 credit=15hrs x 60 mins = 900mins/50mins=18 lectures]

#### **Course Outcomes**

- To understand plant growth improvement with respect to disease resistance, environment tolerance.
- To correlate stages of plant disease development, epidemiology, symptom based classification, control methods.
- To understand the importance of microorganisms in sustainable agriculture, biotechnological application of bio films, edible vaccines.
- To correlate Soil Micro biome and Role of microorganisms in soil health
- To determine the use of Microorganisms as tools in plant genetic engineering.

Credit	Topics	No. of
No.		lectures
	Plant Pathology	18
	<b>1.</b> Plant growth improvement and Stages in development of a disease	3
	a. Plant growth improvement with respect to disease resistance	
	b. Stages in development of a disease: Infection, invasion,	
	colonization, dissemination of pathogens and perennation	
	2. Classification of disease based on symptoms (with one example of	3
Credit	the following):	
Ι	Canker, Downy mildew, Mosaic	
	3. Plant disease epidemiology	6
	Concepts of monocyclic, polycyclic and polyetic diseases withone	
	example of each, disease triangle and forecasting of plant diseases.	
	4. Methods of plant disease control	6
	i. Eradication	
	ii. Chemical control	
	iii. Biological control (employing bacterial and fungal cultures)	
	iv. Integrated pest management	
	v. Genetic engineering for disease resistant plants	

	Microorganisms in sustainable Agriculture and tools in plant	18
	genetic engineering	
	5. Microorganisms in sustainable Agriculture	
	a. Soil Micro biome (plant Micro biome): Concept,	3
Credit	Composition, functioning and methods to study plant Micro	
II	biome	
	b. Conservation of soil health: Role of microorganisms in soil	1
	health	
	c. Phytonutrient availability by soil microorganisms	2
	Mechanism of diazotrophy, Phosphate solubilization,	
	Potassium mobilization, micronutrient availability	
	d. Biofilm in plant surfaces, Biofilm formation; Biofilm in	3
	Phyllosphere and rhizosphere, Examples of plant- microbe	
	interactions in biofilms, Biotechnological applications of	
	plant biofilms	
	6 Microorganisms in plant genetic engineering:	
	a. Concept of GM crops (Transgenic crops) w.r.t. to edible	2
	vaccines, insecticide resistance, herbicide resistance,	
	improved varieties, new variants, disease resistance	
	b. Tools and techniques:	
	i. Microorganisms as tools in plant genetic engineering	2
	(Shuttlevectors)	
	ii Technology of BT resistant crops	1
	iii. Concept of edible vaccines	1
	iv Technique of use of plant viruses in genetic	1
	engineering	
	c. RNAi Technology and antisense RNA technology in	2
	disease resistant plant varieties	

## **References: MB 356 Agricultural Microbiology**

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**CBCS: 2019 Pattern** 

#### **T. Y. B. Sc.**

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## **DSEC - MB 366: Food Microbiology**

#### [2 Credits; 36 Lectures]

#### [1 credit=15hrs x 60 mins = 900mins/50mins=18 lectures]

#### **Course Outcome**

- To describe food safety problems and solutions in India and global scale.
- Identify and classify types of microorganisms in food processing and compare their Characteristics and behavior
- To learn food classification based on their perishability, intrinsic and extrinsic factors affecting the growth of microbes in foods, role of microorganisms in food fermentation.
- To acquire knowledge about food spoilage, food borne diseases, predisposition and preventive and control measures.
- To apply principles of sanitation, heat treatment, irradiation, modified atmosphere, antimicrobial preservatives and combination of method (hurdle concept) to control microbial growth with emphasis on HACCP guidelines.

Credit No	Topics	No. of lectures
	Introduction to properties of food and spoilage of food	18
Credit	1. Classification of food- Perishable, non-perishable, and stable.	4
Ι	Sensory characters of food-	
	a. Definition of food	
	b. Sensory or organoleptic factors- appearance factors-(size,shape,	
	color, gloss, consistency, wholeness)	
	c. Textural factors-texture changes	
	d. Flavor factors (taste, smell, mouthfeel, temperature	
	2. Factors affecting Microbial growth in food	5
	a. Intrinsic factors- pH, water activity, O-R potential, nutrient content,	
	biological structure of food, inhibitory substances in food.	
	b. Extrinsic factors-Temperature of storage, Relative humidity,	
	concentration of gases.	
	3. Sources of food spoilage microorganisms	9
	a. Contamination and spoilage of perishable foods- vegetables and	
	fruits, Meat and meat products, Fish and other sea food, Egg and	
	poultry products.	
	b. Contamination and spoilage of canned foods	
	c. Contamination and spoilage of- cereals and cereal products, sugar	
	and sugar products, salad dressings, spices and condiments.	

	Food Preservation and food in relation to disease	18
	4. Principles of food preservation	10
	a. Importance of TDP, TDT, D, F, Z values	
	b. Use of low and high temperature for food preservation.	
	c. Use of chemicals and antibiotics in food preservation,	
Credit	d. Canning	
II	e. Dehydration	
	f. Use of radiation	
	g. Tetra pack technology	
	h. Food grade bio preservatives	
	5. Microbial food poisoning and food infection	4
	a. Food poisoning - Clostridium botulinum, Aspergillus flavus	
	b. Food infection-Salmonella typhimurium, Vibrio parahaemolyticus	
	6. Concept of Prebiotic and Probiotic and fermented food- definition,	2
	Health effects, Quality assurance, Safety, side effects and risk.	
	Potential applications of Prebiotic, Probiotic and fermented food	
	7. Food sanitation and regulatory authorities (ISO, FDA, WHO)	2

# **References: MB 366 Food Microbiology**

- Alias A K., Paliyath G. and Bhat R. (2012). Progress in Food Preservation. United Kingdom: Wiley. ISBN: 9780470655856
- 2. Banwart G. J. (1989). Basic Food Microbiology. 2nd edition. Chapman and Hall. International Thompson Publishing.
- Bozoglu T. F. and Erkmen O. (2016). Food Microbiology. 2 Volume Set: Principles into Practice. United Kingdom: Wiley. ISBN: 9781119237761
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- 5. Early R. (2012). Guide to quality management for the food Industry. Blackie Academic and Professional. ISBN-13: 978-1461358879
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# Semester V

# **Practical Course-I**

# DSEC-MB – 357: Diagnostic Microbiology and Immunology

# [2 Credits; 78 Lectures]

### [1 credit=15hrs x 130 mins = 1950 mins/50 mins=39 lectures]

78 L distributed as 60 L for performing practicals and 18 L for internal evaluation

# **12 Practicals x 5 lectures = 60 Lectures**

Sr.	Practical	No. of
No		Practicals
1.	Clinical microbiology:	2
	Physical, Chemical and Microscopic examination of Clinical samples -	
	Urine, stool and pus	
2.	Isolation, identification of following pathogens from clinicalsamples:	5
	i. <i>Klebsiella</i> spp.	
	ii. Salmonella spp.	
	iii. Pseudomonas spp	
	iv. Streptococcus spp.	
	v. Enterococcus spp.	
	(for identification use of keys as well as Bergey's Manual is	
	recommended)	
3.	Agglutination tests: Widal test (Slide test and Tube Test) and	1
	Rapid Plasma Reagin (RPR) test	
4.	Epidemiological survey:	2
	Development of hypothesis, Data collection, organization, statistical	
	analysis, graphical representation using computers and interpretation,	
	Preparation of report	
5.	Hemogram:	2
	a. Estimation of hemoglobin (Acid hematin and Cyan-methemoglobin	
	method)	
	b. ESR and PCV determination,	
	c. White blood cell differential count from peripheral blood	
	d. Counting of RBCs and WBCs using counting chamber	
	e. Calculation of hematological indices	

# Semester VI

# **Practical Course-I**

# DSEC-MB – 367: Diagnostic Microbiology and Immunology

# [2 Credits: 78 Lectures]

### [1 credit=15hrs x 130 mins = 1950 mins/50 mins=39 lectures]

78 L distributed as 60 L for performing practicals and 18 L for internal evaluation

# **12 Practicals x 5 lectures = 60 Lectures**

Sr.	Practicals	No. of
No.		Practicals
1.	Study of permanent slides of following microbial pathogens:	3
	a. Entamoeba histolytica	
	b. <i>Giardia</i> spp.	
	c. <i>Plasmodium</i> spp.	
	d. Mycobacterium (tuberculosis and leprae)	
	e. Trichophyton spp.	
	f. Epidermophyton spp.	
	g. Microsporum spp.	
2.	Isolation and identification of following yeast and fungalpathogens:	1
	Cryptococcus neoformans / Histoplasma capsulatum	
3.	Antibiotic sensitivity testing of the bacterial pathogens (for Gram	1
	negative and Gram Positive)	
4.	Immunohematology:	2
	Cross-matching (Major and Minor) and Coomb's test (Direct andIndirect )	
5.	Immuno chromatographic test:	1
	The qualitative differential detection of IgM and IgG antibodies to	
	dengue virus in Human serum /Plasma	
	Or	
	Advantage Mal Card visual immunoassay:	
	The qualitative diagnosis of <i>Plasmodium</i> spps	
6.	Immunoprecipitation:	1
	Double diffusion (Ouchterlony) technique	
7.	Demonstrations of:	2
	a. Serum protein separation by electrophoresis	
		1

### References: MB 357 and MB 367: Diagnostic Microbiology and Immunology

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- 3. Bergey's Manual of Systematic Bacteriology. (2005). Volume Two: The Proteobacteria, Part A: Introductory Essays. Garrity G. editor. Springer. ISBN 978-0-387-24143-2
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- Mukherjee K. L. and Ghosh S. (2010). Medical Laboratory Technology, Volume III: Procedure Manual for Routine Diagnostic Tests. 2nd edition. McGraw Hill Education (India) Private Limited. ISBN-13: 978-1259061257
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# Links:

- 1. *Candida*:https://www.slideshare.net/Sciblack/candidiasis-clinical-manifestationsand-lab-diagnosis-of-oral-candidiasis
- 2. Cryptococcus neoformans:
  - i) (https://www.cdc.gov/fungal/diseases/cryptococcosis-neoformans/symptoms.html)
  - ii) Rev. Inst. Med. Trop. Sao Paulo, 57(4):295-298, July-August, 2015, http://dx.doi.org/10.1590/S0036-46652015000400004, isolation of cryptococcus neoformansfrom environmental samples collected in southeastern Nigeria, emeka i. nweze(1), fred a. kechia(2,3), uju e. dibua(1), charles eze(4) and uwakwe s. onoja(5)
  - iii) https://academic.oup.com/mmy/article/43/6/565/1008990
- 3. *Histoplasma*: https://www.slideshare.net/shahmanthan24/histoplasmoismycology-epidemiology-laboratory-diagnosis?from\_action=save

# Semesters V

# **Practical Course – II**

### MB 358: Enzymology and Genetics

# [2 Credits; 78 Lectures]

# [1 credit=15hrs x 130 mins = 1950 mins/50 mins=39 lectures]

78 L distributed as 60 L for performing practicals and 18 L for internal evaluation

## **12 Practicals x 5 lectures = 60 Lectures**

Sr.	Practical	No. of
No.		Practical
1.	Determination of absorption spectra and molar extinction co-efficient of two	1
	different dyes(by colorimetry /spectrophotometry)	
2.	Qualitative analytical tests using flow charts for Proteins (tests for aromatic	2
	amino acids, sulfur containing amino acids, different amino acids)	
	i. Carbohydrates (tests for monosaccharides, disaccharides, and	
	polysaccharides)	
3.	Preparation of buffers and calibration of pH meter	1
4.	Paper Chromatography	1
	i. Separation and Identification of amino acids from mixture by paper	
	chromatography	
	ii. Separation and Identification of sugars from mixture by paper	
	chromatography	
5.	Extraction and quantitative estimation of total carbohydrate /proteins from	3
	natural sample:	
	i. Estimation of total carbohydrates from natural sources by Lane-Eynon	
	method	
	ii. Estimation of reducing sugar from natural sources by DNSA method	
	iii.Estimation of proteins from natural sources by Folin Lowry method	
6.	Isolation of genomic DNA from bacteria	1
7.	Determination purity of DNA and its quantification:	1
	a. Estimation of DNA by UV- spectrophotometric method, 260/280 ratio	
	b. Estimation of DNA by the diphenylamine	
8.	Bacterial Conjugation	1
9.	Chromosome Staining (G-banding)	1
	Giemsa staining of chromosome from eukaryotic cell extract	

# **References: MB 358 Enzymology and Genetics**

- Ausubel F. M., Brent R., Kingston R. E., Moore D. D., Seidman J.G., Smith J. A. and Struhl K. (Editors.). (2003). Current Protocols in Molecular Biology. Copyright © John Wiley and Sons, Inc. ISBN: 047150338X
- Bollet C., Gevaudan M.J., de Lamballerie X., Zandotti C. and de Micco P. (1991). A simple method for the isolation of chromosomal DNA from Gram positive or acid-fast bacteria. Nucleic Acids Research. 19(8): 1955. https://doi.org/10.1093/nar/19.8.1955
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- Wilson K. and Walker J. (Editors). (2010). Principles and Techniques of Biochemistry and Molecular Biology. 7<sup>th</sup> edition. Cambridge University Press, New York. ISBN-13: 978-0521731676
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### Semester VI

## **Practical Course – II**

### DSEC-MB 368: Metabolism and Molecular Biology

### [2 Credits: 78 Lectures]

### [1 credit=15hrs x 130 mins = 1950 mins/50 mins=39 lectures]

78 L distributed as 60 L for performing practicals and 18 L for internal evaluation

#### **12 Practicals x 5 lectures = 60 Lectures**

Sr. No.	Practicals	No. of Practical
1.	Clinical Biochemistry - Estimations of	3
	i. Blood sugar	
	ii. Blood urea	
	iii. Serum cholesterol	
	iv. Serum proteins and albumin	
2.	Enzyme production, purification, quantification and Immobilization:	4
	i. Lab scale production of amylase using isolates	
	ii. Precipitation of amylase from fermentation broth (salt/solvent)	
	iii. determination of specific activity of crude and purified amylase	
	iv. Immobilization of Amylase using calcium alginate	
3.	Enrichment, Isolation and Enumeration of Bacteriophages (Principle,	2
	Methodology and Calculations of phage titer in PFU/ml)	
4.	Isolation of Plasmid DNA and Agarose Gel Electrophoresis	1
	(Demonstration/hands on as per infrastructure availability)	
5.	Study of Mitotic cell division from onion root tips	1
6.	Visit to a Biotechnology/ Biochemistry institute	1

### **References: MB 368 Metabolism and Molecular Biology**

- Ausubel F. M., Brent R., Kingston R. E., Moore D. D., Seidman J.G., Smith J. A. and Struhl K. (Editors.). (2003). Current Protocols in Molecular Biology. Copyright © John Wiley and Sons, Inc. ISBN: 047150338X
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- Wilson K. and Walker J. (Editors). (2010). Principles and Techniques of Biochemistryand Molecular Biology. 7<sup>th</sup> edition. Cambridge University Press, New York. ISBN-13: 978-0521731676

# Semester V

# Practical course-III

### DSEC-MB 359 Fermentation Technology- I and Agricultural Microbiology

### [2 Credits; 78 Lectures]

### [1 credit=15hrs x 130 mins = 1950 mins/50 mins=39 lectures]

78 L distributed as 60 L for performing practicals and 18 L for internal evaluation

### **12 Practicals x 5 lectures = 60 Lectures**

Sr.	Title of the Practical	No. of
No		Practical
1.	Sterility Testing of pharmaceuticals (non-biocidal injectables): Direct	2
	inoculation method, membrane filtration method, using control test cultures as	
	per IP guidelines (availability at the center).	
2.	Minimum inhibitory concentration and minimum bactericidal concentration of	2
	antibacterial compounds (MIC and MBC)	
3.	Antibiotic and growth factor assay (agar gel diffusion technique)	2
4.	Isolation and identification of Xanthomonas spp. from citrus canker	1
5.	Isolation of Aspergillus niger from black rot of onion	1
6.	Collection of plant disease specimens and study of symptoms/ Project based on	1
	digital record of plant diseases(Group Activity)	
7.	Isolation of PGPR with phosphate solubilization potential/Vesicular-	2
	Arbuscular Mycorrhiza (VAM), Preparation of liquid bioinoculants	
8.	Validation of commercial formulations of bioinoculants based on BIS	1
	standards, Pot studies to check effect of bioinoculants on plant growth	

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### Semester VI

### **Practical Course-III**

### DSEC-MB 369 Fermentation Technology- II and Food Microbiology

### [2 Credits; 78 Lectures]

### [1 credit=15hrs x 130 mins = 1950 mins/50 mins=39 lectures]

78 L distributed as 60 L for performing practicals and 18 L for internal evaluation

#### **12** Practicals x 5 lectures = 60 Lectures

Sr.	Title of the practical	No. of
No		practicals
1.	Lab Scale production of the fermentation products:	2
	a. Ethanol (fermentation, recovery by simple distillation,	
	estimation of end product by CAN method and fermentation	
	efficiency)	
	or	
	b. Citric acid (fermentation, recovery by acid base precipitation	
	and estimation of product by titrometry)	
2.	Solid state fermentation for production of any one fermentation	1
	product (Trichoderma sp. / mushrooms / enzymes)	
3.	Isolation and identification of Probiotic microflora from natural	2
	sources or any commercial formulation.	
4.	Study of SOPs for pharmaceutical industry	1
	a. disinfectant efficacy testing	
	b. Physical monitoring of microbiology section	
	c. Handling of biological indicators	
	d. Microbiological testing of vials	
	e. Identification of contaminant in sterile area	
5.	Detection of aflatoxin	1
6.	Determination of TDP and TDT value	2
7.	Determination of TDR and D value	1
8.	HACCP guidelines for food industry (activity based)	1
9.	Visit to any food industry or a fermentation industry	1

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#### T. Y. B.Sc.

### SEM V

# Skilled Base Elective MB 3510 Marine Microbiology

2 Credit Course: 1.5 credit theory+0.5 credit Practical

### **Course Outcome:**

- To impart the awareness of unseen and unexplored niche of marine ecosystem of microbes.
- To acquire advances in the knowledge of marine microbes and marine ecology.
- To learn the field research on marine processes and laboratory research on microorganisms.
- To comprehend the role of marine microbes in bioremediation and bioprospecting.
- To avail career opportunities in marine education, industry and research.

Credit	Theory	No. of
		lectures
	1. Marine ecology and sampling	
	a. Marine Habitats – estuaries, mangroves, coral reefs, salt	3
	marshes, coastal ecosystems, deep sea, hydrothermal vents,	
	Polar habitat –Arctic, Antarctica, Southern Ocean	
	b. Physiology of marine microorganisms – metabolic diversity,	4
	marineloop, marine snow, Role of marine microorganisms in	
	biogeochemical cycles, nutrient cycling and hydrocarbon	
	degradation	
	c. Sampling methods- water sampling (Niskin sampler) and	4
	sedimentsampling (Grab sampler, box corer, gravity corer),	
Credit	Culturing methods – VBNC, biofilm, mats from vents and	
1.5	estuarine sample.	
	2. Marine microbes, role in bioremediation and bioprospecting	
	a. Extremophilic microorganisms – econiches, different types	2
	withexamples and significance	
	b. Archaea –biodiversity, stress response, adaptation and	3
	significance	2
	c. Marine mycology – econiche, types of marine fungi	
	andsignificance	3
	d. Bioremediation – heavy metals, hydrocarbon pollutants – tar ball	
	and oil spills	

#### **Theory Total Lectures: 21**

# Skilled Based Elective MB 3510:

# **Marine Microbiology Practical**

### Total Lectures: 15 Practical 03 x 05 lectures=15 lectures

Credit	Practical	No. of Practicals
Credit	<b>1.</b> Physico-chemical analysis of sea water	1 1
		1
0.5	<b>2.</b> Isolation of marine bacteria/ fungi from different econiches –	1
	coastal waters, deep sea, estuarine waters, sediments	
	<b>3.</b> Isolation of extremophilic bacteria – halophiles, thermophiles,	1
	acidophiles, alkalophiles, psychrophiles, osmophiles (any two	
	of these)	

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# Semester V

# Skilled Base Elective MB 3511 Dairy Microbiology

# 2 Credit Course: Total lectures: 36: Theory-21 L; Practical-15L

# **Course Outcome:**

- To understand prospects of dairying at commercial marketing.
- To acquire skills of processing of milk and dairy products.
- To assess quality control in dairy industry.
- To comprehend production of dairy products of commercial significance with emphasis tolocal and global market demand.

# Skilled Base Elective MB 3511 Dairy Microbiology Theory Total Lectures: 21

Credit	Theory	No of
No.	1 Definition types microflere and nethogens:	Lectures
	<ol> <li>Definition, types, microflora and pathogens:         <ol> <li>Definition of milk, Composition and physicochemical properties of Milk of different animals. Difference between colostrum and milk.</li> <li>Types of milk: whole, toned, double toned, homogenized, and skimmed milk, dehydrated milk</li> <li>Microflora associated with milk and its importance.</li> <li>Sources of contamination of raw milk and relative importance in influencing quality of milk during production, collection,</li> </ol> </li> </ol>	8
	transportation, and storage, milk borne diseases.	
	2. Processing Techniques and naturally occurring preservatives	4
Credit 1.5	<ul> <li>i. Bacteriological aspects of processing techniques like bactofugation, thermisation, pasteurization (in detail process is expected), sterilization and boiling.</li> <li>ii. Naturally occurring preservative systems in milk like LP system, immunoglobulins, Lysozyme, Lactoferrin etc.</li> </ul>	
	3. Spoilage of Milk	5
	i. Spoilage of Milk	
	ii. Succession of microorganisms in milk leading to spoilage	
	iii. Stormy fermentation, ropiness, sweet curdling	
	iv. Color and flavor defects	
	v. Preservation of Milk and Milk products by physical (irradiation)	
	and Chemical agents, food grade bio preservatives (GRAS), Bacteriocins of LAB	

4. Mi	crobiological aspects of quality control and qualityassurance in	4
pro	oduction of milk and milk products.	
i.	Good Manufacturing Practices,	
ii.	Sanitary standard operating procedures,	
iii.	Total quality management and application of HACCP programin	
	dairy industry.	
iv.	Safety concern of biofilm formation on equipment surfaces and	
	their control measures	

# Skilled Base Elective MB 3511

### **Dairy Microbiology Practical**

### **Total Lectures: 15Total Practical 05 x 05 lectures=15 Lectures**

Credit	Practicals	Number of
		Practicals
	1. Microbiological analysis of milk:	1
	Enumeration of bacteria. (Standard Plate Count (SPC) and	
	DirectMicroscopic Count) – raw milk and pasteurized milk	
	2. Microbiological quality control tests for milk:	1
	i. Dye reduction tests (MBRT/Resazurin)	
	ii. Mastitis test	
	iii. Somatic cell count	
	iv. Phosphatase test	
Credit	3. Microbiological quality of indigenous dairy products:	1
0.5	i. Khoa	
	ii. Kulfi	
	iii. Shrikhand	
	iv. Paneer	
	v. Curd/ Buttermilk	

#### **References:**

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#### T. Y. B.Sc.

### Semester VI

# Semester VI Skilled Base Elective MB 3610 Waste Management

### **2 Credit Course: Total lectures: 36:** Theory-21 L; Practical-15L

### **Course Outcome:**

- To understand waste management and it practicable applicability.
- To assess the magnitude and influence of hazardous content of waste, pollution of waters and waste water treatment technologies.
- To learn the design and working of treatment plants and methods used for liquid and solid waste treatment.
- To impart the understanding of kinetics of biological systems used in waste treatment.
- To learn the standards of waste management and competent authorities involved at National and international level.

Credit	Theory	No. of
<b>a 1</b>		Lectures
Credit	A. Liquid Waste Management	
1.5	1. Principles of Wastewater Treatment	4
	i. The need for treatment of wastewater	
	ii. General characteristics of liquid waste - pH, Color Turbidity,	
	Odor, Electrical conductivity, COD, BOD, Total Solids, Total	
	Dissolved Solids, Total Suspended Solids, Total Volatile	
	Solids, Chlorides, Sulphates, Oil and Grease.	
	2. Microbiology of Wastewater	4
	Role of microorganisms in wastewater treatment	
	i. Aerobic and Anaerobic digestion models; attached /	
	anchored and suspended growth.	
	ii. Removal of pathogenic microbes, indicator microbes,	
	enumeration of different types of microbes	
	3. Unit operations in wastewater treatment plant	4
	i. Collection system - Methods of collection, conservancy systems,	
	water carriage system, sewerage system.	
	ii. Screen chamber, Grit chamber, Oil and grease removal	
	iii. Stabilization pond, Aerated lagoon	
	iv. Activated sludge process, Trickling filter	
	v. Rotating biological contactors, anaerobic digestion processes,	
	fluidized bed reactor.	

#### Skilled Base Elective MB 3610 Waste Management Theory Total Lectures 21

	Торіс	No. of lectures
	B. Solid Waste Management and hazardous waste	
4.	Characterization of solid wastes: Dairy and e-waste	2
5.	Biomedical waste: Definition, Types, Processing	2
6.	Solid biodegradable waste processing: Composting,	2
	Vermicomposting, Biogas production	
7.	Post-processing by-products of municipal solid waste	3
	treatment:leachate refused-derived fuel (RDF)	

# Skilled Base Elective MB 3610 Waste Management Practicals Total Lectures 15

### Total Practicals 05 x 05 lectures= 15 lectures

Credit	Practicals	No. of Practicals
Credit 0.5	1. Determination of Solids in wastewater: Total Solids, Suspended Solids, Dissolved Solids, Volatile Solids, Fixed Solids, Settleable Solids	1
	2. Determination of Dissolved Oxygen, BOD and COD of waste water (before and after treatment) (MPCB Standards)	1
	<b>3.</b> Preparation of Project report based on a case study (Hotel/ Industry-Dairy, Food processing)	1
	Study of the source, generation rates and characteristics of hazardous wastes and their regulation, handling, treatment, and disposal. Special emphasis is placed on process design of waste handling, treatment and	
	emphasis is placed on process design of waste handling, treatment and disposal systems.	

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# Semester VI

# Skilled Base Elective MB 3611 Nano-biotechnology 2 Credit Course: 1.5 credit theory+0.5 credit Practical Theory-21 L; Practical-15L

### **Course Outcome**

- To understand design, development and application of Nanomaterials and their application in Nanodevices.
- To learn fundamentals of nanotechnology as to Synthesis and characterization techniques of nanoparticles.
- To acquire knowledge of applications of nanomaterials in different disciplines of humanlife.
- To compare the merits of using nanotechnology with existing technologies.

# Skilled Base Elective MB 3611 Nano-biotechnology Theory [total lectures 21]

Sr. No.	Торіс	No. of
		Lectures
	1. Introduction to Nano-biotechnology:	6
	a. Introduction to nanoscale, nanomaterials, nanoscience and nanotechnology	
	b. Nanoscale bioassemblies	
	c. Liposomes, viruses, DNA, polysaccharides and proteins (Protein	
	nanotubes, nanofibers, peptide nanoparticles).	
	d. Biomedical applications of bioassemblies	
	e. Cell targeting, drug delivery, bioimaging and vaccine development.	
	2. Microbial mediated metallic nanoparticles synthesis:	5
	a. Gold nanoparticles (AuNPs)	
	b. Silver nanoparticles (AgNPs)	
	c. Au-Ag alloy nanoparticles	
Credit	d. Oxide nanoparticles	
1.5	e. Magnetic nanoparticles	
	f. Non-magnetic oxide nanoparticles	
	g. Sulfide nanoparticles etc.	
	3. Characterization techniques for nanomaterials:	6
	UV-visual spectroscopy, Fourier transform infrared (FTIR), X-ray diffraction	
	(XRD), X-ray photoelectron spectroscopy (XPS), Scanning electron	
	microscopy (SEM), Transmission electron microscopy (TEM) and dynamic	
	light scattering (DLS).	
		4

# 4. Applications of nanoparticles:

Antibacterial agent, drug delivery, biosensor, animal industry and

nanotechnology inwastewater treatment.

# Skilled Base Elective: MB 3611 Nano-biotechnology. Practicals [total lectures 15]

Credit	Practical	No. of
		Practicals
	1. Microbial synthesis of metallic nanoparticle synthesis (any two ): silver, chromium, cobalt)	1
Credit 0.5	<ul> <li>2. Detection and Characterization of metallic nanoparticles in colloidal solutions by:</li> <li>a. UV-Spectrophotometer</li> <li>b. FTIR analysis</li> </ul>	1
	<b>3.</b> Application of nanoparticles- checking antimicrobialactivities against the microbial synthesized metallic nanoparticles (any two)	1

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