

**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 and Technology) 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: Group 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  
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 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)

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

**Equivalences for the New Courses (w. e. f. from 2021-22)****With old Courses (2013 Pattern) in Microbiology****T. Y. B. Sc. Microbiology Semester-VI**

<b>Theory/ Practical/ Skill Enhancement</b>	<b>Old Course Semester-III</b>		<b>New Course Semester-VI (CBCS 2019 Pattern)</b>	
	<b>Course Number</b>	<b>Course Title</b>	<b>Course Number</b>	<b>Course Title</b>
<b>Discipline Specific Elective Course (DSEC) Theory</b>	<b>MB 341</b>	Medical Microbiology-II	<b>MB 361</b>	Medical Microbiology II
	<b>MB 344</b>	Immunology-II	<b>MB 362</b>	Immunology II
	<b>MB 343</b>	Metabolism	<b>MB 363</b>	Metabolism
	<b>MB 342</b>	Genetics and Molecular Biology-II	<b>MB 364</b>	Molecular Biology
	<b>MB 345</b>	Fermentation Technology-II	<b>MB 365</b>	Fermentation Technology II
	<b>MB 336</b>	Food and Dairy Microbiology	<b>MB 366</b>	Food Microbiology
<b>Discipline Specific Elective Course (DSEC) Practical</b>	<b>MB 349</b>	Practical course-III Diagnostic Microbiology and Immunology	<b>MB 367</b>	Practical course-I. Based on: MB 361 Medical Microbiology II and MB 362 Immunology II
	<b>MB 348</b>	Practical course-II Biochemistry and Genetics	<b>MB 368</b>	Practical course-II. Based on: MB 363 Metabolism and MB 364 Molecular Biology
	<b>MB 347</b>	Practical course-I Applied Microbiology	<b>MB 369</b>	Practical course III. Based on: MB 365 Fermentation technology-II and MB 366 Food Microbiology
<b>Skill Enhancement course</b>	-	-	<b>MB 3610</b>	Waste management
	-	-	<b>MB 3611</b>	Nano biotechnology

**Evaluation Pattern****T. Y. B. Sc. Microbiology**

Courses							
Semester-V				Semester-VI			
Paper	Course Title	Internal examination Marks	University examination Marks	Paper	Course Title	Internal Exam Marks	University examination Marks
<b>MB 351</b>	Medical Microbiology I	15	35	<b>MB 361</b>	Medical Microbiology II	15	35
<b>MB 352</b>	Immunology I	15	35	<b>MB 362</b>	Immunology II	15	35
<b>MB 353</b>	Enzymology	15	35	<b>MB 363</b>	Metabolism	15	35
<b>MB 354</b>	Genetics	15	35	<b>MB 364</b>	Molecular Biology	15	35
<b>MB 355</b>	Fermentation technology I	15	35	<b>MB 365</b>	Fermentation Technology II	15	35
<b>MB 356</b>	Agricultural Microbiology	15	35	<b>MB 366</b>	Food Microbiology	15	35
<b>MB 357</b>	Practical course-I Based on: MB351 and MB 352	15	35	<b>MB 367</b>	Practical course I Based on: MB 361 and MB 362	15	35
<b>MB 358</b>	Practical course-II Based on MB 353 and MB 354	15	35	<b>MB 368</b>	Practical course II Based on: MB 363 and MB 364	15	35
<b>MB 359</b>	Practical course-III Based on:MB 355 and MB 356	15	35	<b>MB 369</b>	Practical course III Based on: MB 365 Fermentation technology II, MB 366 Food Microbiology	15	35
<b>MB 3510</b>	Marine Microbiology	15	35	<b>MB 3610</b>	Waste Management	15	35
<b>MB 3511</b>	Dairy Microbiology	15	35	<b>MB 3611</b>	Nano biotechnology	15	35

**Semester V****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
	<b>Introduction to infectious diseases and Epidemiology</b>	<b>18</b>
Credit I	<b>1. Introduction to infectious diseases of following human body systems:</b> (Brief anatomy and Physiology, Diseases, Pathogens, common symptoms) <ul style="list-style-type: none"> <li>a. Respiratory system</li> <li>b. Gastrointestinal system and liver</li> <li>c. Urogenital system</li> <li>d. Central nervous system</li> </ul>	2 2 2 2
	<b>2. Epidemiology:</b> <ul style="list-style-type: none"> <li>a. Case control and cohort studies – Study design and application</li> <li>b. Principle and methods – Clinical trials of drugs and vaccines (Randomized control trials Concurrent parallel and cross-over trials)</li> <li>c. Epidemiology of infectious diseases               <ul style="list-style-type: none"> <li>i. Sources and Reservoirs of Infection</li> <li>ii. Modes of Transmission of Infections</li> <li>iii. Disease Prevention and Control Measures, Vaccine-preventable bacterial diseases and nonvaccine-preventable bacterial diseases</li> </ul> </li> </ul>	2 3 1 1 3

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

**References: MB 351 Medical Microbiology-I**

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- Champoux J. J., Neidhardt F. C., Drew W. L. and Plorde J. J. (2004). Sherris Medical Microbiology: An Introduction to infectious diseases. 4th edition. Ryan K. J. and Ray C. G. (editors). McGraw-Hill Companies. DOI: 10.1036/0838585299
- Dey N. C., Dey T. K. and Sinha D. (2013). Medical Bacteriology Including Medical Mycology and AIDS. 17<sup>th</sup> Edition. New Central Book Agency (P) Ltd (Publisher). India
- Dulbecco R., Eisen H. N. and Davis B. D. (1990). Microbiology. United States: Publisher -Lippincott. ISBN: 9780608072432
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14. 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
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22. Tortora G. J., Funke B. R. and Case C. L. (2016). Microbiology: An introduction 12th Edition, Pearson. ISBN-13: 9780321929150

**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/](https://www.cdc.gov/fungal/diseases/cryptococcosis-neoformans/)

**Semester V****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
	<b>Organs of immune system, Innate immunity, Antigen and Immunoglobulins</b>	<b>18</b>
Credit I	<p><b>1. Organs of immune system:</b></p> <p>a. Primary lymphoid organs (Thymus and Bone Marrow): Thymus – structure, thymic education (positive and negative selection) Bone marrow –Structure and Negative selection</p> <p>b. Secondary lymphoid organs – structure and functions of spleen and lymph node, mucous associated lymphoid tissue, lymphatic system and lymph circulation</p>	2 2
	<p><b>2. Innate immunity: Non-specific mechanisms of defense: Second line of defense:</b></p> <p>a. Humoral components: Defensins, pattern recognition proteins (PRP) and pathogen associated molecular patterns (PAMPs), complement, kinins, and acute phase reactants.</p> <p>b. Cellular components: Phagocytic cells – PMNL, macrophages (reticulo-endothelial cell system) and dendritic cells</p> <p>c. Phagocytosis (oxygen dependent and independent systems), Complement activation (Classical, Alternative and lectin pathway), Inflammation (cardinal signs, mediators, vascular and cellular changes, role of Toll-like receptors)</p>	1 1 5

Credit II	<b>3. Antigen:</b>	
	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, Autoantigens, Isoantigens	1
	<b>4. Immunoglobulins:</b>	
	a. Characteristic of domain structure, functions of light and heavy chain domains and antigenic nature of immunoglobulin molecules	2
	b. Molecular basis of antibody diversity (kappa, lambda and heavy chain)	2
	<b>Antigen- Antibody Interactions, Major Histocompatibility Complex, Transplantation and Immunity and Hybridoma Technology and Monoclonal Antibodies</b>	<b>18</b>
<b>5. Antigen- Antibody Interactions:</b>		
A. <b>Principles of interactions:</b> Antibody affinity and avidity, ratio of antigen antibody, lattice hypothesis and two stage theory, antigen-antibody reaction kinetics (dialysis equilibrium experiment)		2
B. <b>Visualization of antigen antibody complexes:</b>		
a. Precipitation reactions: in fluid and in gel, immunoelectrophoresis		1
b. Agglutination reactions: hemagglutination, bacterial agglutination, passive agglutination and agglutination-inhibition		1
c. Immunofluorescence techniques: direct and indirect, fluorescence-activated cell sorting (FACS)		2
d. Enzyme-linked immunosorbent assay (ELISA), biotin-avidin system and enzyme-linked immune absorbent spot (ELISpot) assay		2
e. Radioimmunoassay RIA		1
<b>6. Major Histocompatibility Complex:</b>		
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. Transplantation and Immunity;</b>		
a. Types of Grafts, Allograft rejection mechanisms		2
b. Prevention of allograft rejection		1

	<b>8. Hybridoma Technology and Monoclonal Antibodies;</b> a. Preparation, HAT selection and propagation of hybridomas secreting monoclonal antibodies b. Applications of monoclonal antibodies	2 1
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**References: MB-352 Immunology- I**

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24. 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 No.	Topics	No. of lectures
	<b>Enzymes:</b>	<b>18</b>
	<b>1. Structure of enzymes:</b> a. Methods to determine amino acid residues at active site (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: Occurrence, Structure and Biochemical functions of the following: i. Thiamine (Vitamin B1) and Thiamine Pyrophosphate ii. Vitamin D	3 2
<b>Credit I</b>	<b>2. Enzyme assays:</b> a. Principles of enzyme assays and calculation of enzyme unit, specific activity b. Enzymes assays with examples by: i. Spectrophotometric methods ii. Radioisotope assay	1 2

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

**References: MB 353 Enzymology**

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**CBCS: 2019 Pattern****T. Y. B. Sc.****Microbiology**

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6. Nelson D. L. and Cox M. M. (2021). Lehninger's Principles of Biochemistry. 8<sup>th</sup> Edition. Mac Millan Worth Pub. Co. New Delhi. ISBN:9781319228002
7. Palmer T. (2001) Enzymes: Biochemistry, Biotechnology and Clinical chemistry. Horwood Pub. Co. Chichester, England. ISBN-9781898563785
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**Semester V****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 No.	Topics	No. of lectures
	<b>DNA Replication and Gene Expression</b>	<b>18</b>
I	<b>1. Process of prokaryotic DNA replication</b> <ul style="list-style-type: none"> <li>a. Single replicon</li> <li>b. Bidirectional movement of replication fork</li> <li>c. Ori C</li> <li>d. Pre-priming and Priming reaction.</li> <li>e. DNA polymerases, DNA synthesis of leading, lagging strand Okazaki fragments.</li> <li>f. Termination- Ter sequence, Tusprotein</li> </ul> <b>2. Prokaryotic and Eukaryotic Transcription</b> <ul style="list-style-type: none"> <li><b>i. Transcription in Prokaryotes</b> <ul style="list-style-type: none"> <li>a. Structure of promoter</li> <li>b. Structure and function of RNA polymerase</li> <li>c. Steps of transcription: Initiation, Elongation and termination</li> </ul> </li> <li><b>ii. Transcription in eukaryotes with respect to protein coding Gene:</b> <ul style="list-style-type: none"> <li>a. Promoter, promoter proximal elements and enhancers</li> <li>b. Transcription regulatory proteins</li> <li>c. RNA polymerases</li> <li>d. Steps in transcription: Initiation, Elongation, Termination</li> <li>e. Post transcriptional modifications: 5' capping, 3' polyadenylation and introduction to RNA splicing</li> </ul> </li> </ul>	4
		3

	<b>3. Regulation of transcription:</b> Concept and components of operon: Lac operon: Inducible operon	2
	<b>4. Translation in prokaryotes and eukaryotes</b> a. Structure and role of m-RNA, t-RNA and Ribosomes in 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	5
	<b>Gene transfer and mapping techniques</b>	<b>18</b>
	<b>5. Gene transfer by Transformation</b> a. Discovery of Transformation b. Natural transformation Systems- <i>Streptococcus pneumoniae</i> and <i>Haemophilus influenzae</i> . c. Factors affecting transformation i. Competence development ii. Size of DNA iii. Concentration of DNA	4
Credit II	<b>6. Gene transfer by Conjugation</b> a. Discovery of Conjugation, b. Properties of F plasmid, F <sup>+</sup> , F <sup>-</sup> , Hfr and F' strains c. Process of conjugation between F <sup>+</sup> and F <sup>-</sup> , Hfr and F <sup>-</sup> , F' and F <sup>-</sup>	4
	<b>7. Gene transfer by Transduction</b> a. Discovery of Transduction b. Generalized transduction mediated by P22 c. Specialized transduction mediated by lambda phage	4
	<b>8. An introduction to Gene mapping</b> 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-transduction and interrupted mating	6

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**Reference-Links:**

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<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC388511/?page=1>
2. National Academies Press: Introduction of Recombinant DNA-Engineered Organisms Into the Environment: Key Issues: <https://www.nap.edu/download/18907#>
3. Guidelines and Handbook for Institutional Biosafety Committees (DBT, Govt. of India and BCIL):<https://thsti.res.in/pdf/IBG.pdf>
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>

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

<b>Credit</b>  <b>II</b>	<b>3. Sterilization of Media:</b>	
	a. Methods of sterilization	1
	b. Batch sterilization and Continuous sterilization (direct and indirect methods)	1
	c. Concept and derivation of Del factor	1
	d. Filter sterilization of liquid media	1
	<b>4. Scale-up and Scale-down:</b>	
	a. Objectives of scale-up	1
<b>Credit</b>  <b>II</b>	b. Levels of fermentation (laboratory, pilot-plant and production level – flow sheet to explain scale up)	1
	c. Criteria of scale-up for critical parameters [Aeration (kLa 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)]	1
	d. Scale-down (example of anyone commercial fermentation)	1
	<b>Downstream processing and Quality assurance of fermentation products</b>	<b>18</b>
	<b>5. Downstream processing of fermentation products: (method, principle, types, examples of fermentations, factors affecting, merits and demerits at large scale operation)</b>	
	a. Cell disruption methods	1
	b. Filtration	1
<b>Credit</b>  <b>II</b>	c. Centrifugation	1
	d. Liquid-liquid extraction	1
	e. Distillation	1
	f. Drying	1
	<b>6. Quality assurance of fermentation products (as per IP, USP)</b>	
	a. Methods of detection and Quantification of the fermentation product: physicochemical, biological and enzymatic methods	2
	b. Sterility testing (direct inoculation method, membrane filtration method)	1
	c. Bioburden test	1

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

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**Reference links:**

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[https://www.who.int/biologicals/vaccines/Tetanus\\_Recommendations\\_TRS\\_980\\_Annex\\_5](https://www.who.int/biologicals/vaccines/Tetanus_Recommendations_TRS_980_Annex_5)

<https://academic.oup.com/jimb/article-pdf/18/5/340/34773995/jimb0340.pdf>.

25. Large scale production of rabies vaccine:

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**Semester V****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 No.	Topics	No. of lectures
Credit I	<b>Plant Pathology</b>	18
	1. Plant growth improvement and Stages in development of a disease <ol style="list-style-type: none"> <li>a. Plant growth improvement with respect to disease resistance</li> <li>b. Stages in development of a disease: Infection, invasion, colonization, dissemination of pathogens and perennation</li> </ol>	3
	2. Classification of disease based on symptoms (with one example of the following): Canker, Downy mildew, Mosaic	3
	3. Plant disease epidemiology Concepts of monocyclic, polycyclic and polyetic diseases with one example of each, disease triangle and forecasting of plant diseases.	6
	4. Methods of plant disease control <ol style="list-style-type: none"> <li>i. Eradication</li> <li>ii. Chemical control</li> <li>iii. Biological control (employing bacterial and fungal cultures)</li> <li>iv. Integrated pest management</li> <li>v. Genetic engineering for disease resistant plants</li> </ol>	6

	<b>Microorganisms in sustainable Agriculture and tools in plant genetic engineering</b>	<b>18</b>
<b>Credit</b>	<b>5. Microorganisms in sustainable Agriculture</b>	
<b>II</b>	a. Soil Micro biome (plant Micro biome)	2
	b. Concept, Composition, functioning and methods to study:	
	i. Conservation of soil health: Role of microorganisms in soil health	1
	ii. Phytonutrient availability by soil microorganisms Mechanism of diazotrophy, Phosphate solubilization, Potassium mobilization, micronutrient availability	4
	iii. Biofilm in plant surfaces, Biofilm formation; Biofilm in Phyllosphere and rhizosphere, Examples of plant- microbe interactions in biofilms, Biotechnological applications of plant biofilms	3
	<b>6 Microorganisms in plant genetic engineering:</b>	
	a Concept of GM crops (Transgenic crops) w.r.t. to edible vaccines, insecticide resistance, herbicide resistance, improved varieties, new variants, disease resistance	2
	b. Tools and techniques:	
	i. Microorganisms as tools in plant genetic engineering (Shuttle vectors)	1
	ii Technology of BT resistant crops	1
	iii. Concept of edible vaccines	1
	iv Technique of use of plant viruses in genetic engineering	1
	c. RNAi Technology and antisense RNA technology in disease resistant plant varieties	2

**References: MB 356 Agricultural Microbiology**

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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**

<b>Sr. No</b>	<b>Title of the Practical</b>	<b>No. of Practicals</b>
1.	<b>Clinical microbiology:</b>  <b>Physical, Chemical and Microscopic examination of Clinical samples -</b> Urine, stool and pus	2
2.	<b>Isolation, identification of following pathogens from clinical samples:</b>  i. <i>Klebsiella</i> spp. ii. <i>Salmonella</i> spp. iii. <i>Pseudomonas</i> spp iv. <i>Streptococcus</i> spp  <b>and <i>Enterococcus</i> spp</b>  (for identification use of keys as well as Bergey's Manual is recommended)	4
3.	<b>Agglutination tests:</b>  Widal test (Slide test and Tube Test) and  Rapid Plasma Reagins (RPR) test	1
4.	<b>Epidemiological survey:</b>  Development of hypothesis, Data collection, organization, statistical analysis, graphical representation using computers and interpretation, Preparation of report	2
5.	<b>Hemogram:</b>  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	3

**References: MB 357: Diagnostic Microbiology and Immunology**

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6. Bergey's Manual of Systematic Bacteriology. (2009). Volume Three: The Firmicutes. Part C: The Proteobacteria. Vos, P., Garrity, G., Jones, D., Krieg, N.R., Ludwig, W., Rainey, F.A., Schleifer, K.-H., Whitman, W. (Eds.). Springer. ISBN 978-0-387-95041-9
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**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. No.	Title of the Practical	No. of Practical
1.	Determination of absorption spectra and molar extinction co-efficient of two different dyes (by colorimetry /spectrophotometry)	1
2.	Qualitative analytical tests using flow charts for <ul style="list-style-type: none"> <li>i. Proteins (tests for aromatic amino acids, sulfur containing amino acids, different amino acids)</li> <li>ii. Carbohydrates (tests for monosaccharides, disaccharides, and polysaccharides)</li> </ul>	2
3.	Preparation of buffers and calibration of pH meter	1
4.	Paper Chromatography <ul style="list-style-type: none"> <li>i. Separation and Identification of amino acids from mixture by paper chromatography</li> <li>ii. Separation and Identification of sugars from mixture by paper chromatography</li> </ul>	1
5.	Extraction and quantitative estimation of total carbohydrate /proteins from natural sample: <ul style="list-style-type: none"> <li>i. Estimation of total carbohydrates from natural sources by Phenol Sulphuric acid method</li> <li>ii. Estimation of reducing sugar from natural sources by DNSA method</li> <li>iii. Estimation of proteins from natural sources by Folin Lowry method</li> </ul>	3
6.	Isolation of genomic DNA from bacteria	1
7.	Determination purity of DNA and its quantification: <ul style="list-style-type: none"> <li>a. Estimation of DNA by UV- spectrophotometric method, 260/280 ratio</li> <li>b. Estimation of DNA by the diphenylamine</li> </ul>	1
8.	Bacterial Conjugation	1
9.	Chromosome Staining (G-banding) Giemsa staining of chromosome from eukaryotic cell extract	1

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

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- Indian Pharmacopeia. (2018 Addendum 2021). <https://www.indianpharmacopoeia.in/index.php>
- USA Clinical Laboratory Standards Institute(CLSI) Guidelines 2021 on <https://clsi.org/>
- Sterility Testing:  
[https://www.who.int/medicines/publications/pharmacopoeia/TestForSterility-RevGenMethod\\_QAS11-413FINALMarch2012.pdf](https://www.who.int/medicines/publications/pharmacopoeia/TestForSterility-RevGenMethod_QAS11-413FINALMarch2012.pdf)
- Microbiological assay of antibiotics: <https://apps.who.int/phtml/pdf/b/7.3.1.3.1-Microbiological-assay-of-antibiotics.pdf>

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<https://lib.dr.iastate.edu/cgi/viewcontent.cgi?article=14117&context=rtd>.

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7. Isolation of *Xanthomonas citri* from citrus canker:—

<https://www.plantbiosecuritydiagnostics.net.au/app/uploads/2018/11/NDP-9-Asiatic-citrus-canker-Xanthomonas-V1.2.pdf>.

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- Dube H. C. and Bilgrami K.S.1976 Text book of modern pathology. Vikas Publishing House. New Delhi.
- Mehrotra R. S. (1994). Plant Pathology. Tata McGraw-Hill Limited.
- Rangaswami G. (2005). Diseases of Crop Plants in India. 4th edition. Prentice Hall of India Pvt. Ltd., New Delhi.

9. Isolation of *Plasmopara viticola* from grapes (Downy Mildew):

M. A. Mane, S. S. Bodke and R. N. Dhawale (2018). Isolation and Identification of *Plasmopara viticola* associated with Grapevine from Marathwada Region. International Journal of Current Microbiology and Applied Sciences ISSN: 2319-7706 Special Issue-6 pp. 714-728

10. Validation of standards of biofertilizers:

- Manual - <https://law.resource.org/pub/in/bis/S06/is.6092.3.2.2004.pdf>
- Rhizobial and azotobacterial biofertilizers: <https://bio-fit.eu/q8/lo6-quality-control-of-biofertilizers?start=4>.

- Organic Farming: Organic Inputs and Techniques:

[http://agritech.tnau.ac.in/org\\_farm/orgfarm\\_biofertilizertechnology.html](http://agritech.tnau.ac.in/org_farm/orgfarm_biofertilizertechnology.html).

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- Yadav A. K. and Chandra K. (2014). Mass Production and Quality Control of Microbial Inoculants. Proc Indian Natn Sci Acad. 80 (2): 483-489.

11. Isolation of PGPR with PSB:

- <https://www.ijnpnd.com/article.asp?issn=2231-0738;year=2013;volume=3;issue=1;spage=29;epage=33;aulast=Ranjan>.
- <https://scielo.conicyt.cl/pdf/jsspn/v16n2/aop4316.pdf>

**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.

**Theory Total Lectures: 21**

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

**Skilled Based Elective MB 3510:****Marine Microbiology Practical****Total Lectures: 15 Practical 03 x 05 lectures=15 lectures**

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

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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 to local and global market demand.

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

Credit No.	Theory	No of Lectures
Credit 1.5	<p><b>1. Definition, types, microflora and pathogens:</b></p> <ul style="list-style-type: none"> <li>i. Definition of milk, Composition and physicochemical properties of Milk of different animals. Difference between colostrum and milk.</li> <li>ii. Types of milk: whole, toned, double toned, homogenized, and skimmed milk, dehydrated milk</li> <li>iii. Microflora associated with milk and its importance.</li> <li>iv. Sources of contamination of raw milk and relative importance in influencing quality of milk during production, collection, transportation, and storage, milk borne diseases.</li> </ul>	8
	<p><b>2. Processing Techniques and naturally occurring preservatives</b></p> <ul style="list-style-type: none"> <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>	4
	<p><b>3. Spoilage of Milk</b></p> <ul style="list-style-type: none"> <li>i. Spoilage of Milk</li> <li>ii. Succession of microorganisms in milk leading to spoilage</li> <li>iii. Stormy fermentation, ropiness, sweet curdling</li> <li>iv. Color and flavor defects</li> <li>v. Preservation of Milk and Milk products by physical (irradiation) and Chemical agents, food grade bio preservatives (GRAS), Bacteriocins of LAB</li> </ul>	5

	<b>4. Microbiological aspects of quality control and quality assurance in production of milk and milk products.</b> i. Good Manufacturing Practices, ii. Sanitary standard operating procedures, iii. Total quality management and application of HACCP program in dairy industry. iv. Safety concern of biofilm formation on equipment surfaces and their control measures	4
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**Skilled Base Elective MB 3511****Dairy Microbiology Practical****Total Lectures: 15Total Practical 05 x 05 lectures=15 Lectures**

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

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**Semester VI****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
Credit I	<b>Chemotherapy</b>	<b>18</b>
	<b>1. Routes of drug administration.</b>	1
	<b>2. Mode of action of antimicrobial agents on:</b>	
	<b>a. Bacteria:</b>	
	i. Cell wall: Beta lactams: 1 <sup>st</sup> to 6 <sup>th</sup> Generation- e.g. Meropenem, Imipenem, Piperacillin, Tazobactam	2
	ii. Cell membrane: Polymyxin	1
	iii. Protein synthesis: Streptomycin, Tetracycline	1
	iv. Nucleic acids: Fluroquinolones, Rifamycin	1
	v. Enzyme inhibitors: Trimethoprim, Sulfomethaxazole	1
	<b>b. Fungi:</b> Griseofulvin, Amphotericin B, Anidulafungin, Vericonazole	3
	<b>c. Viruses:</b> Acyclovir, Oseltamivir, Remdecivir	1
	<b>d. Protozoa:</b> Metronidazole, Chloroquine	1
	<b>3. Mechanisms of drug resistance on:</b>	
	<b>a. Genetic basis:</b>	3
	i. Mutations in gene(s)	
	ii. Acquisition of foreign DNA coding for resistance determinants through horizontal gene transfer.	
	<b>b. Mechanisms of drug resistance by:</b>	3
	i. Limiting uptake of a drug.	
ii. Modification of a drug target.		
iii. Inactivation of a drug.		
iv. Active efflux of a drug.		

Credit II	<b>Human and Animal Viruses, Fungal and Protozoal Pathogens</b>	<b>18</b>
	<b>4. Introduction to cultivation of viruses</b>	2
	<b>5. Study of following groups of viral pathogens:</b>	
	a. <b>Human viruses</b> (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
	v. Cutaneous Viruses: Human papillomavirus	1
	vi. Neurological Viruses: Japanese Encephalitis Virus	1
	b. <b>Animal Viruses:</b> FMD Virus and Rinderpest Virus	2
	<b>6. Study of following groups of parasites</b> (with respect to Classification, Lifecycle, Morphological characteristics, Viability characteristics, Pathogenicity, Pathogenesis, Symptoms, Laboratory diagnosis (Serological diagnosis wherever applicable), Epidemiology, Prophylaxis and Chemotherapy):	
	a. <i>Plasmodium</i>	2
	b. <i>Entamoeba</i>	1
	<b>7. Study of following groups of yeast and fungal pathogens</b> (With respect to – Morphological and cultural characteristics, Classification, Pathogenicity, Pathogenesis, Symptoms, Laboratory diagnosis, Epidemiology, Prophylaxis and Chemotherapy)	
	a) <i>Aspergillus</i> species (Pathogenic)	1
	b) <i>Cryptococcus neoformans</i>	1
	c) <i>Histoplasma capsulatum</i>	1

**References: MB 361 Medical Microbiology- II**

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**Links:**

6. <https://www.who.int/travel-advice/disease-information>
7. <https://Microbenotes.Com/Remdesivir/#Mechanism-Of-Action-Of-Remdesivir>
8. *Aspergillus* <https://www.cdc.gov/fungal/diseases/aspergillosis/index.html>
9. *Histoplasma capsulatum* <https://www.cdc.gov/fungal/diseases/histoplasmosis/>
10. *Cryptococcus neoformans* [www.cdc.gov/fungal/diseases/cryptococcosis-neoformans/](https://www.cdc.gov/fungal/diseases/cryptococcosis-neoformans/)

**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
	<b>Cytokines, Adaptive / Acquired Immunity, Hypersensitivity, Autoimmunity and Autoimmune diseases and Immunodeficiency</b>	<b>18</b>
	<b>1. Cytokines:</b> <ul style="list-style-type: none"> <li>a. Concept- Cytokines, lymphokines, monokines, interleukines, chemokines, interferons and tumor necrosis factor</li> <li>b. Properties, Attributes and biological functions of cytokines</li> </ul>	1 2
	<b>2 Adaptive / Acquired Immunity (Third line of defense):</b> <ul style="list-style-type: none"> <li><b>A. Humoral Immune Response</b> <ul style="list-style-type: none"> <li>i. Primary and secondary response kinetics, significance in vaccination programs</li> <li>ii. Response of secondary lymphoid organs to antigen</li> <li>iii. Antigen processing and presentation (Major Histocompatibility 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</li> </ul> </li> <li><b>B. Cell Mediated Immune Response</b> <ul style="list-style-type: none"> <li>i. Activation and differentiation of T cells, role of cytokines in activation</li> <li>ii. Mechanism of Cytotoxic T lymphocytes (CTL) mediated cytotoxicity, Antibody-dependent cellular cytotoxicity (ADCC)</li> <li>iii. Significance of Cell Mediated Immune Response (CMI)</li> <li>iv. Immune response against tumors and foreign transplanted cells</li> </ul> </li> </ul>	2 1 5 2 3 1 1
<b>Credit</b>		
<b>I</b>		

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

**References: MB 362- Immunology-II**

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**Semester VI****DSEC-MB 363: Metabolism****[2 Credits; 36 Lectures]****[1 credit=15hrs x 60 mins = 900mins/50mins=18 lectures]**

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

**References: MB 363 Metabolism**

11. Berg J. M., Stryer L., Tymoczko J. and Gatto G. (2019). Biochemistry. 9<sup>th</sup> Edition. Palgrave Macmillan. ISBN-978-1319114657.
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**Semester VI****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
	<b>Genetic Recombination and Bacteriophage Genetics.</b>	<b>18</b>
Credit I	<p><b>1. Gene linkage and crossing over</b></p> <ul style="list-style-type: none"> <li>a. Mendel's laws: Eukaryotic Cell cycle, Mitosis, Meiosis</li> <li>b. Holliday model for Homologous recombination, Role of Rec and Ruvproteins</li> <li>c. Genetic mapping by Tetrad analysis in <i>N. crassa</i> (Numerical Calculations using PD, TT and NPD)</li> <li>d. Genetic Mapping by Parasexual cycle in <i>A. nidulans</i></li> </ul> <p><b>2. Bacteriophage Genetics</b></p> <ul style="list-style-type: none"> <li>a. Lytic cycle: Virulent phages, T-series phages, Concept and formation of plaque, Lysogenic cycle: Temperate phage (<math>\lambda</math>phage)</li> <li>b. Bacteriophage mutants: Plaque morphology (r type), Host range, Conditional lethal mutants (Ts and Am)</li> <li>c. Concept of Genetic Complementation and Cis-trans test of genetic function. (Intergenic- rII locus of T4 phage, Mechanism of Intragenic complementation.)</li> <li>d. Fine structure mapping of rII locus of T4 phage using Benzer's spot tests and deletion mapping</li> </ul>	9
	<b>DNA damage and repair mechanisms, Recombinant DNA technology</b>	<b>18</b>
Credit II	<p><b>3. DNA damage and Repair mechanisms</b></p> <ul style="list-style-type: none"> <li>a. DNA damage by hydrolysis, deamination, alkylation, oxidation, Radiation (X rays and UV rays)</li> <li>b. DNA repair by Photo reactivation</li> <li>c. DNA repair by Mismatch repair mechanism</li> <li>d. DNA repair by Excision repair mechanisms (BER/NER)</li> </ul>	5

	<b>4. Recombinant DNA Technology Tools and basics of recombinant DNA technology</b> a. Introduction to recombinant DNA technology b. Restriction enzymes: Concept, Nomenclature, properties and types with examples (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 ( <i>E. coli</i> and T4 phage), Use of Linker / Adaptor / Homopolymer tailing e. Methods to transfer recombinant DNA into bacterial host cells (Physical – Electroporation, Gene gun, Chemical –CaCl <sub>2</sub> mediated, liposome mediated) f. Methods of screening recombinants using selective markers and Blue-White screening	10
	<b>5. Molecular techniques used in RDT</b> 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	3

**References: MB 364 Molecular Biology**

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**Reference-Links:**

5. Potential biohazards of recombinant DNA molecules:  
<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC388511/?page=1>
6. National Academies Press: Introduction of Recombinant DNA-Engineered Organisms Into the Environment: Key Issues: <https://www.nap.edu/download/18907#>
7. Guidelines and Handbook for Institutional Biosafety Committees (DBT, Govt. of India and BCIL):<https://thsti.res.in/pdf/IBG.pdf>
8. 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>

**Semester VI****DSEC - MB 365 Fermentation Technology – II****[2 Credits; 36 Lectures]****[1 credit=15hrs x 60 mins = 900mins/50mins=18 lectures]**

Credit No.	Topics	No. of lectures
	<b>Solid state and Submerged state fermentations and Large scale fermentations</b>	<b>18</b>
<b>Credit I</b>	<b>1. Introduction to Solid State Fermentation and Submerged Fermentation:</b> Process, production strains, media, fermentor design, fermentation conditions, applications, merits and demerits	<b>1</b>
	<b>2 Large scale production of</b> (process with flow sheet, nature of the product, production pathway, applications, production strains, media, fermentation process, parameters, product recovery)	<b>3</b>
	<b>a. Primary Metabolites:</b> i. Vitamins (B12 and B2) ii. Amino acids - Glutamic acid, Lysine iii. Organic acids (Citric acid, Vinegar and Lactic acid)	<b>1</b>
	<b>b. Secondary metabolites:</b> i. Bioethanol ii. Alcoholic Beverages - 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 Streptomycin]	<b>3</b>

Credit <b>II</b>	<b>Large scale production of enzymes, steroids, biomass-based products, milk products, vaccines, immune sera and Modern trends in microbial production</b>	<b>18</b>
	<b>3. Enzymes</b>	
	i. Amylase	1
	ii. Esterases	1
	iii. Proteases	1
	<b>4. Microbial transformation of steroids</b>	<b>2</b>
	<b>5. Biomass based products:</b>	
	i. Yeast: Baker's and Distiller's yeast	2
	ii. Probiotics: <i>Lactobacillus sporogenes</i>	1
	<b>6. Milk products:</b>	
	i. Cheese (Processed, soft, semi-hard, hard ripened types- bacterial and mold)	2
	ii. Yogurt (plain, flavoured, fruit, sundae style. Stirred type, set type, probiotic yoghurt)	2
	<b>7. Vaccines</b>	
	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
	<b>8. Immune sera</b>	
	i. Anti tetanus serum (ATS)	1
	ii. Anti rabitic serum (ARS)	1
	<b>9. Modern trends in microbial production:</b>	
	Biosurfactant and bioemulsifier	1

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**Reference links:**

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[https://www.who.int/biologicals/vaccines/Tetanus\\_Recommendations\\_TRS\\_980\\_Annex\\_5](https://www.who.int/biologicals/vaccines/Tetanus_Recommendations_TRS_980_Annex_5)  
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52. Large scale production of rabies vaccine:
53. Large scale production of tetanus vaccine:  
<http://nopr.niscair.res.in/bitstream/123456789/26533/1/JSIR%2060%2810%29%20773-778.pdf>.
54. USA Clinical Laboratory Standards Institute(CLSI) Guidelines 2021: <https://clsi.org/>

**Semester VI****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 behaviour
- 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
	<b>Introduction to properties of food and spoilage of food</b>	<b>18</b>
<b>Credit I</b>	<b>1. Classification of food- Perishable, non-perishable, and stable.</b> <b>Sensory characters of food-</b> <ol style="list-style-type: none"> <li>a. Definition of food</li> <li>b. Sensory or organoleptic factors- appearance factors (size, shape, color, gloss, consistency, wholeness)</li> <li>c. Textural factors-texture changes</li> <li>d. Flavor factors (taste, smell, mouthfeel, temperature)</li> </ol>	4
	<b>2. Factors affecting Microbial growth in food</b> <ol style="list-style-type: none"> <li>a. Intrinsic factors- pH, water activity, O-R potential, nutrient content, biological structure of food, inhibitory substances in food.</li> <li>b. Extrinsic factors-Temperature of storage, Relative humidity, concentration of gases.</li> </ol>	5
	<b>3. Sources of food spoilage microorganisms</b> <ol style="list-style-type: none"> <li>a. Contamination and spoilage of perishable foods- vegetables and fruits, Meat and meat products, Fish and other sea food, Egg and poultry products.</li> <li>b. Contamination and spoilage of canned foods</li> <li>c. Contamination and spoilage of- cereals and cereal products, sugar and sugar products, salad dressings, spices and condiments.</li> </ol>	9

<b>Credit II</b>	<b>Food Preservation and food in relation to disease</b>	<b>18</b>
	<b>a. Principles of food preservation</b> <ul style="list-style-type: none"> <li>a. Importance of TDP, TDT, D, F, Z values</li> <li>b. Use of low and high temperature for food preservation.</li> <li>c. Use of chemicals and antibiotics in food preservation,</li> <li>d. Canning</li> <li>e. Dehydration</li> <li>f. Use of radiation</li> <li>g. Tetra pack technology</li> <li>h. Food grade bio preservatives</li> </ul>	10
	<b>5. Microbial food poisoning and food infection</b> <ul style="list-style-type: none"> <li>a. Food poisoning -<i>Clostridium botulinum, Aspergillus flavus</i></li> <li>b. Food infection-<i>Salmonella typhimurium, Vibrio parahaemolyticus</i></li> </ul>	4
	<b>6. Concept of Prebiotic and Probiotic and fermented food-</b> definition, Health effects, Quality assurance, Safety, side effects and risk. Potential applications of Prebiotic, Probiotic and fermented food	2
	<b>7. Food sanitation and regulatory authorities (ISO, FDA, WHO)</b>	2

### References: MB 366 Food Microbiology

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**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. No.	Title of the Practical	No. of Practicals
1.	<b>Study of permanent slides/ of following microbial pathogens:</b> a) <i>Entamoeba histolytica</i> b) <i>Giardia</i> spp. c) <i>Plasmodium</i> spp. d) <i>Mycobacterium</i> (tuberculosis and leprosy) e) <i>Epidermophyton</i> spp.	1
2.	<b>Isolation and identification of following:</b> Isolation and identification of <i>Candida</i> from skin/mouth. (Slide Culture Technique) a. i. Isolation and identification of <i>Aspergillus niger</i> ii. Determination of Koch's Postulates using <i>Aspergillus niger</i> . iii. Total fungal spore count by Neubauer's chamber	4
3.	<b>Antibiotic sensitivity testing of the bacterial pathogens (for Gram negative and Gram Positive)</b>	1
4.	<b>Immunohematology:</b> a. Determination of titre of Anti-A and Anti-B in human serum b. Cross-matching (Major and Minor) and Coomb's test (Direct and Indirect)	2
5.	Qualitative detection of Rheumatoid factor (RA factor) and Streptolysin O using Slide test.	1
6.	<b>Immunoprecipitation:</b> Double diffusion (Ouchterlony) technique	1
7.	<b>Demonstrations of:</b> a. ELISA (Antigen/ Antibody detection) b. Egg inoculation technique	1
8.	<b>Visit to blood bank and preparation of visit report</b>	1

**References: MB 367: Diagnostic Microbiology and Immunology**

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28. Talib V. H. (2019). Handbook Medical Laboratory Technology. 2nd edition. CBS Publishers and Distributors Pvt. Ltd. ISBN-13: 978-8123906775

**Practical Course – II****SEC-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.	Title of the Practical	No. of Practical
1.	Clinical Biochemistry - Estimations of i. Blood sugar ii. Blood urea iii. Serum cholesterol iv. Serum proteins and albumin	3
2.	Enzyme production, purification, quantification and Immobilization: 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	4
3.	Enrichment, Isolation and Enumeration of Bacteriophages (Principle, Methodology and Calculations of phage titer in PFU/ml)	2
4.	Isolation of Plasmid DNA and Agarose Gel Electrophoresis (Demonstration/hands on as per infrastructure availability)	1
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**

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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. No	Title of the practical	No. of Practicals
1.	Lab Scale production of the fermentation products: a. Ethanol (fermentation, recovery by simple distillation, estimation of end product by CAN method and fermentation efficiency) <b>or</b> b. Citric acid (fermentation, recovery by acid base precipitation and estimation of product by titrometry)	2
2.	Solid state fermentation for production of any one fermentation product ( <i>Trichoderma sp.</i> / mushrooms / enzymes)	1
3.	Isolation and identification of Probiotic microflora from natural sources or any commercial formulation.	2
4.	Study of SOPs for pharmaceutical industry 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	1
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

**References: MB 369 Fermentation Technology- II and Food Microbiology**

## 1. Lab scale fermentations:

- Casida L. E., Jr. (2019). Industrial Microbiology, New Age International Publishers, New Delhi. ISBN- 9788122438024

- Patel A. H. (2016). Industrial Microbiology. Trinity Press (Publisher). ISBN-13-9789385750267  
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  - <https://iopscience.iop.org/article/10.1088/1757-899X/612/2/022111/pdf>.
  - <https://www.scielo.br/j/babt/a/vDHdsFscjRYSW6jkRfKQCDM/?lang=en>.
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  - <https://www.frontiersin.org/articles/10.3389/fmicb.2019.01382/full>.
  - <https://www.hindawi.com/journals/ijmicro/2020/8865456/>.
- 4. Study of SOPs:
  - <https://www.pharmaguideline.com/2012/01/sop-for-physical-monitoring-of.html>
  - <http://biomanufacturing.org/uploads/files/989767618742858542-sop-visual-inspection-process.pdf>.
- 5. Detection of aflatoxin:
  - [https://old.fssai.gov.in/Portals/0/Pdf/Draft\\_Manuals/MYCOTOXIN.pdf](https://old.fssai.gov.in/Portals/0/Pdf/Draft_Manuals/MYCOTOXIN.pdf).
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  - Jay J. M. and Loessner M. J. (2005). Modern Food Microbiology. 7th edition. Springer. ISBN 978-0-387-23413-7.
- 7. HACCP:  
[https://www.fsai.ie/food\\_businesses/haccp/principles\\_of\\_haccp.html](https://www.fsai.ie/food_businesses/haccp/principles_of_haccp.html).

**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 its 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.

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

Credit	Theory	No. of Lectures
1.5	<p><b>A. Liquid Waste Management</b></p> <p><b>1. Principles of Wastewater Treatment</b></p> <ul style="list-style-type: none"> <li>i. The need for treatment of wastewater</li> <li>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.</li> </ul> <p><b>2. Microbiology of Wastewater</b></p> <ul style="list-style-type: none"> <li>Role of microorganisms in wastewater treatment</li> <li>i. Aerobic and Anaerobic digestion models; attached / anchored and suspended growth.</li> <li>ii. Removal of pathogenic microbes, indicator microbes, enumeration of different types of microbes</li> </ul> <p><b>3. Unit operations in wastewater treatment plant</b></p> <ul style="list-style-type: none"> <li>i. Collection system - Methods of collection, conservancy systems, water carriage system, sewerage system.</li> <li>ii. Screen chamber, Grit chamber, Oil and grease removal</li> <li>iii. Stabilization pond, Aerated lagoon</li> <li>iv. Activated sludge process, Trickling filter</li> <li>v. Rotating biological contactors, anaerobic digestion processes, fluidized bed reactor.</li> </ul>	4 4 4

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

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

Total Practicals 05 x 05 lectures= 15 lectures

Credit	Practicals	No. of Practicals
	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
0.5	3. Preparation of Project report based on a case study (Hotel/ Industry-Dairy, Food processing)  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 disposal systems.	1

### References: Skilled Base Elective MB 3610 Waste Management

- Chandrappa R. and Das D. B. (2012). Solid Waste Management-Principles and Practice. In Environmental Science and Engineering. Springer (Firm).
- Dutta S., Neela Priya D., Chakradhar B. and Sasi Jyothsna T.S. (2019) Value Added By-products Recovery from Municipal Solid Waste in Waste Valorisation and Recycling. Springer, Singapore.
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10. Standard Methods for the Examination of Water and Wastewater. (2017). 23<sup>rd</sup> Edition. American Public Health Association, American Water Works Association, and Water Environment Federation
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13. Wesley Eckenfelder W. Jr. (2000). Industrial Water Pollution Control. 3rd Edition. McGraw Hill.

**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 human life.
- To compare the merits of using nanotechnology with existing technologies.

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

Sr. No.	Topic	No. of Lectures
1.5	<p><b>1. Introduction to Nano-biotechnology:</b></p> <ul style="list-style-type: none"> <li>a. Introduction to nanoscale, nanomaterials, nanoscience and nanotechnology</li> <li>b. Nanoscale bioassemblies</li> <li>c. Liposomes, viruses, DNA, polysaccharides and proteins (Protein nanotubes, nanofibers, peptide nanoparticles).</li> <li>d. Biomedical applications of bioassemblies</li> <li>e. Cell targeting, drug delivery, bioimaging and vaccine development.</li> </ul> <p><b>2. Microbial mediated metallic nanoparticles synthesis:</b></p> <ul style="list-style-type: none"> <li>a. Gold nanoparticles (AuNPs)</li> <li>b. Silver nanoparticles (AgNPs)</li> <li>c. Au-Ag alloy nanoparticles</li> <li>d. Oxide nanoparticles</li> <li>e. Magnetic nanoparticles</li> <li>f. Non-magnetic oxide nanoparticles</li> <li>g. Sulfide nanoparticles etc.</li> </ul> <p><b>3. Characterization techniques for nanomaterials:</b></p> <p>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).</p>	6 5 6 4

	<b>4. Applications of nanoparticles:</b> Antibacterial agent, drug delivery, biosensor, animal industry and nanotechnology in wastewater treatment.	
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**Skilled Base Elective: MB 3611 Nano-biotechnology. Practicals [total lectures 15]**

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

**References: Skilled Base Elective: MB 3611 Nano-biotechnology.**

1. Bujold K. E., Lacroix A., and Sleiman H. F. (2018). DNA Nanostructures at the Interface with Biology. *Chem.* 4: 495–521. Elsevier Inc.
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<http://dx.doi.org/10.1098/rsif.2012.0740>
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