

Model curriculum for VI semester



Government of Karnataka Model Curriculum

Program Name	BSc in Microbiology	Semester	VI
Course Title	IMMUNOLOGY AND MEDICAL MICROBIOLOGY (Theory)		
Course Code:	MIC C13-T	No. of Credits	4
Contact hours	60 Hours(4 hours per week)	Duration of SEA/Exam	2 hours
Formative Assessment Marks	40	Summative Assessment Marks	60

Course Pre-requisite(s): Common to the Course Programme at Entry Level

Course Outcomes (COs): After the successful completion of the course, the student will be able to:

CO1: To gain a preliminary understanding about various immune mechanisms.

CO2: To familiarize with Immunological techniques and serodiagnosis of infectious diseases

CO3: To understand pathogenic bacterial infections, symptoms, diagnosis and treatment process.

CO4: To understand pathogenic bacterial infections, symptoms, diagnosis and To understand pathogenic bacterial infections, symptoms, diagnosis and treatment process treatment process

Contents	60 Hrs
<p>UNIT-I</p> <p>Normal microflora of the human body and host pathogen interaction</p> <p>Normal microflora of the human body: Importance of normal microflora, normal microflora of skin,throat, gastrointestinal tract, urogenital tract Host pathogen interaction: Definitions - Infection, Invasion, Pathogen, Pathogenicity, Virulence, Toxigenicity, Carriers and their types, Opportunistic infections, Nosocomial infections. Transmission of infection, Pathophysiologic effects of LPS. Sample collection, transport and diagnosis.</p> <p>Clinical Microbiology</p> <p>Medical Bacteriology</p> <p>The following diseases in detail with Symptoms, mode of transmission, prophylaxis and control</p> <p>respiratory diseases: <i>Streptococcus pyogenes</i>, <i>Haemophilus influenzae</i>, <i>Mycobacterium tuberculosis</i></p> <p>Gastrointestinal Diseases: <i>Escherichia coli</i>, <i>Salmonella typhi</i>, <i>Vibrio cholerae</i>, Others: <i>Staphylococcus aureus</i>, <i>Bacillus anthracis</i>, <i>Clostridium tetani</i>, (10 hrs)</p>	15 hrs.
<p>UNIT-II</p> <p>Medical Virology, parasitology and Mycology</p> <p>The following diseases in detail with Symptoms, mode of transmission, prophylaxis and control</p> <p>Polio, Herpes, Hepatitis, Rabies, Dengue, AIDS, Corona, Influenza, swine flu, Ebola, Chikungunya, Japanese Encephalitis</p> <p>Protozoan diseases: Malaria, Kala-azar, Entamoeba</p> <p>Fungal infections- Cutaneous mycoses: Tinea, pedis (Athlete's foot) Systemic mycoses: Histoplasmosis Opportunistic mycoses: Candidiasis (10 Hrs)</p> <p>Antimicrobial agents: General characteristics and mode of action Antibacterial agents: Inhibitor of nucleic acid synthesis; Inhibitor of cell wall synthesis; Inhibitor of cell membrane function; Inhibitor of protein synthesis; Inhibitor of metabolism</p> <p>Antifungal agents: Mechanism of action of Amphotericin B, Griseofulvin Antiviral agents: Mechanism of action of Amantadine, Acyclovir, Azidothymidine . Antibiotic resistance, MDR, XDR, MRSA, NDM-1 5hrs</p>	15 Hrs



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Course Title	IMMUNOLOGY AND MEDICAL MICROBIOLOGY (Practical)	Practical Credits	2
Course Code	MIC C14-P	Contact Hours	4Hours/week
Formative Assessment	25 Marks	Summative Assessment	25 Marks
Practical Content			

1	Identify pathogenic bacteria (any three of <i>E. coli</i> , <i>Salmonella</i> , <i>Pseudomonas</i> , <i>Staphylococcus Bacillus</i>) on the basis of cultural, morphological and biochemical characteristics: IMViC, TSI, nitrate reduction, urease production and catalase tests
2	Study of composition and use of important differential media for identification of pathogenic bacteria: EMB Agar, McConkey agar, Mannitol salt agar, Deoxycholate citrate agar, TCBS
3	Study of bacterial flora of skin by swab method
4	Perform antibacterial sensitivity by Kirby-Bauer method
5	Study symptoms of the diseases with the help of photographs: Polio, anthrax, herpes, chicken pox, HPV warts, AIDS (candidiasis), dermatomycoses (ring worms)
6	Study of various stages of Malarial parasite in RBCs using permanent mounts.
7	Identification of human blood groups.
8	Perform Total Leukocyte Count of the given blood sample.
9	Perform Differential Leukocyte Count of the given blood sample.
10	Separate serum from the blood sample (demonstration).
11	Perform immunodiffusion by Ouchterlony method.
12	Perform DOT ELISA.
13	Perform immunoelectrophoresis.

Pedagogy: Experiential learning, Problem solving, Project

Formative Assessment for Practical	
Assessment Occasion/ type	Marks
Attendance	05 Marks
Records	05 Marks
Performance	05 Marks
Test	10 Marks
Total	25 Marks

Formative Assessment as per guidelines are compulsory

REFERENCES

1	Ananthanarayan R and Paniker C.K.J (2009) Textbook of Microbiology, 8 th Edition, University Press, Publication.
2	Brooks G.F., Carroll K.C., Butel J.S., Morse S.A. and Mietzner, T.A. (2013) Jawetz, Melnick and Adelberg's Medical Microbiology. 26 th edition. McGraw Hill Publication
3	Goering R., Dockrell H., Zuckerman M. and Wakelin D. (2007) Mims' Medical Microbiology. 4 th edition. Elsevier
4	Willey JM, Sherwood LM, and Woolverton CJ. (2013) Prescott, Harley and Klein's Microbiology. 9 th edition. McGraw Hill Higher Education
5	Madigan MT, Martinko JM, Dunlap PV and Clark DP. (2014). Brock Biology of Microorganisms. 14 th edition. Pearson International Edition
6	Abbas AK, Lichtman AH, Pillai S. (2007). Cellular and Molecular Immunology. 6 th edition Saunders Publication, Philadelphia.
7	Delves P, Martin S, Burton D, Roitt IM. (2006). Roitt's Essential Immunology. 11 th edition Wiley-Blackwell Scientific Publication, Oxford.
8	Goldsby RA, Kindt TJ, Osborne BA. (2007). Kuby's Immunology. 6 th edition W.H. Freeman and Company, New York.
9	Murphy K, Travers.P, Walport M. (2008). Janeway's Immunobiology. 7 th edition Garland Science, Publishers, New York.
10	Peakman.M.and Vergani D. (2009). Basic and Clinical Immunology, 2 nd edition Churchill, Livingstone Publishers, Edinberg.
11	Richard C and Geiffrey S. (2009). Immunology. 6 th edition. Wiley Blackwell Publication.



**Government of Karnataka
Model Curriculum**

Program Name	BSc in Microbiology	Semester	VI
Course Title	MICROBIAL GENETIC ENGINEERING (Theory)		
Course Code:	MIC C15-T	No. of Credits	3
Contact hours	45 Hours(3 Hours per week)	Duration of SEA/Exam	2 hours
Formative Assessment Marks	40	Summative Assessment Marks	60

Course Pre-requisite(s): Common to the Course Programme at Entry Level

- CO1 : To acquire knowledge on the concepts and terminology in genetic engineering
 CO2 : To learn about principles involved in manipulating genes and DNA
 CO3 : Familiar with various cloning strategies in prokaryotes
 CO4 : Learn techniques in genetic engineering
 CO5 : To have awareness of IPR, the social and the ethical issues concerning cloning by genetic engineering

MICROBIAL GENETIC ENGINEERING	45Hrs
Unit 1: Introduction to Microbial Genetic Engineering	15 Hrs
<p>Historical prospectives: Definition of genetic engineering, milestones in genetic engineering, prospects and problems of genetic engineering.</p> <p>Tools in Microbial Genetic Engineering: Restriction modification systems- Types, Mode of action, nomenclature, applications of restriction enzymes in genetic engineering. DNA modifying enzymes and their applications: DNA polymerases, methylases, Terminal deoxynucleotidyl transferase, kinases and phosphatases and DNA ligases.</p>	
Unit 2: Cloning vectors, DNA transfer methods and identification of recombinants	15 Hrs
<p>Cloning Vectors: Definition and Properties. Characteristics of cloning vectors. Plasmid vectors: pBR and pUC series. Bacteriophage lambda, cosmids, BACs, YACs. Use of linkers and adaptors. Expression vectors: Baculovirus based vectors, mammalian SV40-based expression vectors.</p> <p>Cloning host- Cloning in <i>Escherichia coli</i>, cloning in <i>Saccharomyces cerevisiae</i>, cloning in GRAS microorganism. Gene Library: Construction of cDNA library, genomic library. DNA transfer methods: Microinjection, Biolistic, Electroporation, Calcium phosphate and Liposome mediated DNA transfer. Identification and selection of recombinants: DNA hybridisation, blue white selection, antibiotic selection, colony and plaque hybridization.</p>	
Unit 3: Techniques and applications in Microbial Genetic Engineering	15 Hrs
<p>Isolation and Detection of DNA: Isolation of DNA, restriction digestion and ligation of DNA, Agarose gel electrophoresis, Blotting techniques- Southern blotting, Northern blotting, dot blot, DNA microarray analysis, Western blotting. DNA sequencing- Sanger's method. PCR techniques and applications.</p> <p>Recombinant microorganisms: Application of recombinant microorganisms in basic research, industry, medicine, agriculture, environment.</p> <p>Products of recombinant DNA technology: Products of human therapeutic interest - insulin, hGH, antisense molecules. Bt transgenic - cotton, brinjal, Gene therapy, recombinant vaccines. Biological, ethical and social issues of gene cloning and IPR.</p>	

Course Articulation Matrix: Mapping of Course Outcomes (COs) with Program Outcomes (POs 1-12)

Course Outcomes (COs) / Program Outcomes (POs)	Program Outcomes (POs)														
	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15
To acquire knowledge on the concepts and terminology in genetic engineering	√					√									
To learn about principles involved in manipulating genes and DNA	√		√						√						
Familiar with various cloning strategies in prokaryotes									√	√					
Learn techniques in genetic engineering						√						√			
To have awareness of IPR, the social and the ethical issues concerning cloning by genetic engineering										√					

Pedagogy: Lectures, Seminars, Industry/Institute Visits, Debates, Quiz, Project and Assignments

Formative Assessment for Theory	
Assessment Occasion/ type	Marks
Attendance	10
Seminar	10
Debate/Quiz/Assignment	10
Class test	10
Total	40 Marks
<i>Formative Assessment as per guidelines are compulsory</i>	


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Course Title	MICROBIAL GENETIC ENGINEERING (Practical)	Practical Credits	02
Course Code	MIC C16-P	Contact Hours	4 Hours/ week
Formative Assessment	25 Marks	Summative Assessment	25 Marks
Practical Content			

Practical: Microbial Genetic Engineering

Preparation of buffers-TE, TAE and Lysis buffer.
 Isolation of plasmid DNA from *Escherichia coli*.
 Estimation of DNA by DPA method.
 Demonstration of estimation of DNA by spectrophotometric method.
 Resolution and visualization of DNA by agarose gel electrophoresis.
 Induction of mutations in bacteria by UV light.
 Preparation of competent cells and demonstration of bacterial transformation.
 Demonstration of bacterial transformation and calculation of transformation efficiency.
 Digestion of DNA with restriction enzymes.
 Demonstration of ligation of DNA fragments.
 Preparation of master and replica plates.
 Designing of primers for DNA amplification.
 Demonstration of amplification of DNA by PCR.
 Demonstration of Southern blotting.
 Study of recombinant products-as per theory syllabus.

Pedagogy: Experiential learning, Problem solving, Project

Formative Assessment for Practical	
Assessment Occasion/ type	Marks
Class Records	05
Test	10
Attendance	05
Performance	05
Total	25 Marks
<i>Formative Assessment as per guidelines are compulsory</i>	

REFERENCES :

1	Brown TA. (2010). Gene Cloning and DNA Analysis. 6th edition. Blackwell Publishing, Oxford, U.K. Clark DP and Pazdernik NJ. (2009). Biotechnology: Applying the Genetic Revolution. Elsevier Academic Press, USA
2	Krebs J, Goldstein E, Kilpatrick S (2013). Lewin's Essential Genes, 3rd Ed., Jones and Bartlett Learning Primrose SB and Twyman RM. (2006). Principles of Gene Manipulation and Genomics, 7th edition. Blackwell Publishing, Oxford, U.K.
3	Primrose SB and Twyman RM. (2008). Genomics: Applications in human biology. Blackwell Publishing, Oxford, U.K.
4	Russell PJ. (2009). i Genetics- A Molecular Approach. 3rd Ed, Benjamin Cummings
5	Sambrook J and Russell D. (2001). Molecular Cloning-A Laboratory Manual. 3rd edition. Cold Spring Harbor Laboratory Press
6	Sambrook J and Russell DW. (2001). Molecular Cloning: A Laboratory Manual. 4th Edition, Cold Spring Harbour Laboratory press.
7	Watson JD, Baker TA, Bell SP et al. (2008) Molecular Biology of the Gene, 6th Ed., Benjamin Cummings Wiley JM, Sherwood LM and Woolverton CJ. (2008). Prescott, Harley and Klein's Microbiology. McGraw Hill Higher Education.

Internship for graduate Programme (As Per UGC & AICTE)

Course title	Internship Discipline specific
No of contact hours	90
No credits	2
Method of evaluation	Presentations/Report submission/Activity etc.,

- ❖ Internship shall be Discipline Specific of 90 hours (2 credits) with a duration 4-6 weeks.
- ❖ Internship may be full-time/part-time (full-time during semester holidays and part-time in the academic session)
- ❖ Internship mentor/supervisor shall avail work allotment during 6th semester for a maximum of 20 hours.
- ❖ The student should submit the final internship report (90 hours of Internship) to the mentor for completion of the internship.
- ❖ The detailed guidelines and formats shall be formulated by the universities separately as prescribed in accordance to UGC and AICTE guidelines.


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CBCS Question Paper Pattern for UG Semester DSC

Paper Code:	Paper Title:		
Duration of Exam	2 Hours	Max Marks	60
Instruction:	Answer all the sections		

Section-A

.....	15 Marks
I. Answer any Five of the following questions (5x3=15)	
1. 2. 3. 4. 5. 6. 7.	

Section-B

.....	25 Marks
II. Answer any FIVE of the following questions (5X5=25)	
8. 9. 10. 11. 12. 13. 14.	

Section-C

.....	20 Marks
III. Answer any TWO of the following questions (2X10=20)	
15. 16. 17. 18.	