

Affiliated to the University of Mumbai

Programme: Sciences M.Sc I Microbiology

Syllabus for the Academic Year 2024-2025 based on the National Education Policy 2020



DEPARTMENT OF MICROBIOLOGY

COURSE DETAILS:

		SEMES	STER 1		SEMESTER 2			
TITLE	Virolog y and cell biology -I	Genetic s-I	Microb ial Bioche mistry	Resear ch Method ology	Virology And Cell Biology-I I	Genetics- II	Food Microbiol ogy	Field Project/O n Job Training (OJT)
TYPE OF COURSE - M.Sc	Manda tory I	Manda tory 2	Electiv e	Resear ch Metho dology	Mandato ry I	Mandato ry 2	Elective	On Job Training (OJT)
CREDITS	6	6	4	4	6	6	4	4

Preamble:

The M.Sc program at Sophia College (Autonomous) is open to both female and male students. The M.Sc course is an extension of the undergraduate curriculum dealing with all the branches of Microbiology at a considerable depth and blends the upcoming fields as well as advances in the subject. Research is an integral aspect of the curriculum and includes planning and execution of a dissertation. The outcomes of a number of the dissertations have been published in peer reviewed journals.Participation and presentations - both oral and posters in conferences, workshops and research meets is encouraged. Field projects, Educational visits and short-term internships are also included. The students who complete the postgraduate programme in Microbiology are well trained in the subject and find employment in areas like Quality control, Research and Development, Clinical Research, Teaching etc.



PROGRAMME OBJECTIVES

PO 1	To provide a comprehensive understanding of fundamental and advanced theoretical concepts and practical skills in microbiology.
PO 2	Inculcate research skills and develop ability to create hypotheses, design experiments, and interpret and document scientific results effectively.
PO 3	Prepare students for employment opportunities in research and industry based on their fields of interest.
PO 4	Enable students to apply microbiological knowledge and skills to real-world problems in environmental and industrial contexts.

PROGRAMME SPECIFIC OUTCOMES

PSO 1	Students will gain theoretical and practical knowledge about general microbiology, molecular biology, genetics, cell biology, microbial biochemistry, medical microbiology and immunology.
PSO 2	Students will learn to formulate hypotheses, design experiments, analyze and articulate results using various microbiological methods and consequently be capable of conducting a scientific enquiry.
PSO 3	Students will develop skills essential for employability in the academia, research and industry in Microbiology and Life Science sector.
PSO 4	Foster a sense of responsibility towards societal issues related to microbiology, including ethical considerations in research.



Programme: Sciences MICROBIOLOGY M	SC-I	Semeste	r – 1
Course Title: Virology an	nd cell biology-I	Course C	ode: SMCB511MJ
 COURSE OBJECTIVES: 1. To explain and describe 2. To discuss the life cycle Viroids. 3. To develop an underse eukaryotes. 4. To explain cell biology and animal viruses. 	the replication and regulation e and other details of plant vir standing of cell biology of of humans and animals in or	of transcription uses and agen eukaryotic m der to understa	on of bacteriophages. ts that infect plants such as icroorganisms and higher and the life cycle of human
 <u>COURSE OUTCOMES</u>: The learner will be able to : explain and compare rep explain the structure, recontrol of plant viral infe describe the role of menchloroplast in eukaryotes explain and discuss eukand vesicle transport. elaborate vacuoles of eular 	lication and regulation of gene plication and life cycle of spections. mbrane proteins and transport, s. aryotic nuclear pore complex, karyotic microorganisms such	e expression of pecific plant v , mitochondria , Endoplasmic as algae and a	f various bacteriophages. viruses and prevention and al ETC, ATP synthesis and reticulum, Golgi complex umoeba.
Lectures per week (1 Lecture	is 60 minutes)		4
Total number of Hours in a S	emester		60
Credits			4
Evaluation System	Semester End Examination	2 Hours	50 marks
	Internal Assessment		50 marks



UNIT 1 Bacteriophages	1.1	<i>E.coli</i> Phage T7: Genetic organization, regulation of transcription, DNA replication and maturation	15 hours
	1.2	 a. <i>E.coli</i> Phage φX174: Replication, transcription, packaging b. Filamentous DNA phages- M13: Attachment and entry, replication, assembly and release 	
	1.3	Single stranded RNA phages MS-2 and Qβ: Genetic organization and life cycle	
	1.4	a. Lambda phage: lytic and lysogenic cycleb. Bacteriophage Mu: Properties, Genetic organization and replication	
UNIT 2 Plant Viruses (1 Credit)	2.1	Viruses causing plant diseases: history, structure, transmission, symptoms, detection, prevention and control	15 hours
	2.2	Life cycles- overview Tobacco Mosaic Virus and Brome Mosaic Virus- Life cycle, host range, transmission, symptoms, diagnosis and control	
	2.3	Antiviral plant defense mechanisms: physical factors and RNA interference	
	2.4	Plant satellites and Viroids	
UNIT 3 Plasma		Students to revise basic properties of cells, different classes of cells and functions of plasma membrane	15 hours
membrane, Mitochondria and Chloroplast (1 Credit)	3.1	 Plasma membrane a. Chemical composition of membranes- (in brief) - Membrane lipids (phosphoglycerides, sphingolipids, cholesterol), carbohydrates b. Structure and functions of membrane proteins - Integral membrane proteins, 	



		 peripheral membrane proteins, lipid anchored membrane proteins c. Movement of substances across cell membranes - Diffusion of substances through membranes (Voltage-gated channels, Ligand-gated channels, Mechano-gated channels), Facilitated diffusion, Active transport 	
	3.2	 Mitochondria a. Mitochondrial structure and function- membrane and matrix b. Oxidative metabolism in the mitochondrion c. Role of mitochondria in the formation of ATP - Electron transport, types of electron carriers, Establishment of proton motive force d. Machinery for ATP formation - Structure of ATP synthase, basis of ATP formation, Rotational catalysis 	
	3.3	 Chloroplast a. Chloroplast structure and function b. Photosynthetic metabolism c. Photosynthetic pigments, Photosynthetic units and reaction centers - PSII operations, PSI operations, and Photophosphorylation 	
UNIT 4 Endomembra ne system	4.1	Nuclear envelope, Structure of the Nuclear Pore Complex and its role in Nucleocytoplasmic exchange	15 hours
(1 Credit)	4.2	The endoplasmic reticulum, The smooth endoplasmic reticulum , Functions of the rough endoplasmic reticulum- synthesis and processing of proteins	



4.3	The Golgi complex, Types of vesicle transport and their functions- Cop II- coated vesicles, Cop I-coated vesicles and Endocytic pathway	
4.4	Lysosomes, Contractile Vacuoles in algae and amoeba	
4.5	Exosomes	

PRACTICAL	1	
Course Title:	Virology And Cell	Biology-I

Course Code: SMCB511MJP

COURSE OUTCOMES:

The learner will be able to :

- 1. use the plaque assay to enumerate bacteriophages and calculate plaque forming units/ml
- 2. perform one step growth curve experiment
- 3. apply the fundamentals and concepts of lysogeny for other bacteriophages
- 4. assess the integrity of cell membrane using neutral red uptake method
- 5. perform the extraction of mitochondria and chloroplast from eukaryotic cells

Lectures per week (1 Lecture is 120 minutes)		2		
Total number of Hours in a Semester		60		
Credits		2		
Evaluation System	Semester End Examination	2 Hours	50 marks	
	Internal Assessment			

1	Enumeration of coliphages by plaque assay.	
2	Study of one step growth curve of a bacteriophage.	60 hours
3	Study of lysogeny in <i>E. coli</i> .	

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4	Assignment on any plant virus (other than TMV and BMV).	
5	Study of cell membrane integrity using uptake of neutral red.	
6	Isolation of mitochondria	
7	Isolation of chloroplasts.	

Programme: Sciences MICROBIOLOGY MSC-I	Semester – 1	
Course Title: Genetics-I	Course Code:SMCB512MJ	

COURSE OBJECTIVES:

- 1. To explain coordination of DNA replication, septum formation and chromosome partitioning in bacteria.
- 2. To describe the molecular details of gene expression and its regulation in bacteria and eukaryotes.
- 3. To discuss recombination at the molecular level in bacteria and eukaryotic microorganisms such as yeast.
- 4. To explain the complementation test and its significance in mapping of genes.
- 5. To understand the lac operon and develop critical thinking skills
- 6. To gain knowledge of epigenetic modifications of genes in eukaryotes.

COURSE OUTCOMES:

The learner will be able to

- 1. explain the role of bacterial proteins in septum formation and segregation of chromosomes and also in partitioning of plasmids.
- 2. describe molecular details of transcription, RNA processing, splicing and translation.
- 3. explain the DSB repair model of recombination, role of proteins in bacterial and eukaryotic recombination, mating type switching in *Saccharomyces cerevisiae* and compare homologous recombination in bacteria and eukaryotes.
- 4. explain the complementation test and fine structure mapping and their significance.
- 5. distinguish between different mechanisms of regulation of bacterial operons



6. compare different mechanisms of eukaryotic gene regulation.				
Lectures per week (1 Lecture	is 60 minutes)	4		
Total number of Hours in a Semester		60		
Credits		4		
Evaluation System	Semester End Examination	2 Hours	50 marks	
	Internal Assessment		50 marks	

UNIT 1 Bacterial Cell Division, Chromosome partitioning and Gene expression	1.1	 Cell division and chromosome partitioning in bacteria a. Replication and cell cycle b. Septum formation in bacteria, Function of FtsZ, MinCD and MinE c. Partitioning of Chromosomes d. Partitioning of single copy plasmids 	15 hours
(1 Credit)	1.2	Gene expression- Transcriptiona. Bacterial Transcriptionb. Eukaryotic Transcription	
	1.3	 RNA molecules and processing - a. Messenger RNA- Structure, processing, addition of the 5' Cap, addition of the Poly (A) tail, RNA splicing, self splicing introns, Alternative processing pathways, RNA editing b. Transfer RNA- Structure of transfer RNA, tRNA gene structure and processing c. Ribosomal RNA- Structure of the ribosome, rRNA gene structure and processing. 	
	1.4	Gene expression - Translation	



		a. The process of translation- The binding of amino acids to transfer RNAsb. Initiation, elongation and termination of translationc. Posttranslational modifications of proteins	
UNIT 2 Recombinatio n, Mutation and Genetic Complementa tion (1 Credit)	2.1	 Recombination a. DSB repair model - steps b. Proteins involved in Homologous recombination in prokaryotes - RecBCD, RecA, RuvA, RuvB and RuvC c. Homologous recombination in eukaryotes and proteins involved in the same d. Mating type switching in Saccharomyces cerevisiae (Gene conversion) e. Concept of linkage 	15 hours
	2.2	Mutation a. Somatic mutation and germline mutation b. Study of mutants	
	2.3	Genetic Complementation - Complementation test and fine structure mapping	
UNIT 3 Regulation of gene expression in bacteria (1 Credit)	3.1	 Operons a. The <i>lac</i> operon of <i>E. coli</i> - Experimental evidence for the regulation of <i>lac</i> genes, mutations in the protein-coding and regulatory genes, and positive control of the <i>lac</i> operon b. The <i>ara</i> operon of <i>E. coli</i>: Positive and negative control c. The <i>trp</i> operon of <i>E. coli</i>- Attenuation 	15 hours
	3.2	Other regulatory mechanisms - Antisense RNA, Riboswitches, Sigma factor switching- Sporulation in <i>Bacillus subtilis</i>	



UNIT 4	4.1	Gene regulation in Eukaryotes-	
Eukaryotes gene expression – Regulation and epigenetic modification s		 a. Changes in chromatin structure and histone modifications b. Regulation of transcription factors and activators c. RNA Processing- Examples- SV40, sex differentiation in Drosophila, Degradation of RNA, RNA interference (in brief) d. Processes that affect translation and modification of proteins. 	
(1 Credit)	4.2	 Epigenetic modifications that alter gene expression a. Dosage compensation of genes on X chromosomes- mechanism of X chromosome inactivation b. Gene Imprinting – Mechanisms and imprinting disorders c. Noncoding RNAs d. DNA Methylation 	15 hours

PRACTICAL
Course Title: Genetics-ICourse Code: SMCB512MJP

COURSE OUTCOMES:

The learner will be able to :

- 1. prepare agarose gels, load DNA samples, run electrophoresis, visualize the separated DNA bands and interpret the gel image to understand the plasmid topology and size.
- 2. perform experimental procedures to study bacterial conjugation and analyze the order of the gene transfer.
- 3. perform the necessary steps to expose microorganisms to UV radiation for mutagenesis and isolate streptomycin-resistant mutants using selective culturing techniques.



- 4. enrich and isolate auxotrophic mutants using penicillin enrichment and replica plate techniques and determine the proportion of auxotrophic mutants
- 5. perform the β -galactosidase assay and acquire skills in quantifying and analyzing β -galactosidase activity.
- 6. explain the regulation of Lac operon and apply critical as well problem-solving skills to lac operon-related analytical questions.

Lectures per week (1 Lecture is 120 minutes)		2	
Total number of Hours in a Semester		60	
Credits		2	
Evaluation SystemSemester End Examination		2 Hours	50 marks
Internal Assessment			

1	Separation of plasmid or genomic DNA using agarose gel electrophoresis	60 hours
2	Bacterial conjugation	00 110410
3	UV mutagenesis	
4	Penicillin enrichment technique	
5	β- galactosidase assay	
6	Problems on <i>lac</i> operon	

Programme: Sciences MICROBIOLOGY MSC-I	Semester – 1	
Course Title: Microbial Biochemistry	Course Code:SMCB511E	



COURSE OBJECTIVES:

- 1. To revise the structure, properties and functions of important macromolecules.
- 2. To develop understanding of different analytical methods for studying macromolecules
- 3. To assimilate the principles behind common methods of extraction , purification and study of proteins.

COURSE OUTCOMES:

The learner will be able to

- 1. describe the correlation between structure and functions of cellular macromolecules like proteins, lipids, carbohydrates.
- 2. explain the details of extraction & purification of proteins by salt precipitation and dialysis and separating mixture of proteins by chromatography and electrophoresis.
- 3. elaborate on protein folding mechanism in cells
- 4. discuss the principle and applications of spectroscopic techniques and X ray diffraction analysis done to characterize proteins.
- 5. outline use of radioisotopes in biology experiments

Lectures per week (1 Lecture is 60 minutes)		2	
Total number of Hours in a Semester		30	
Credits		2	
Evaluation System	Semester End Examination	2 Hours	50 marks
	Internal Assessment		50 marks

1.1	Structure and function of Proteins: Peptide bond and its stability, Ramachandran plot. Factors determining primary, secondary, tertiary and quaternary structure of proteins, thermodynamics of folding, role of disulfide bonds, dynamics of globular protein folding, chaperonins. Motifs and domains, protein families, protein stability, protein-protein interactions.	15 hours
	protein-protein interactions.	
	1.1	1.1 Structure and function of Proteins: Peptide bond and its stability, Ramachandran plot. Factors determining primary, secondary, tertiary and quaternary structure of proteins, thermodynamics of folding, role of disulfide bonds, dynamics of globular protein folding, chaperonins. Motifs and domains, protein families, protein stability, protein-protein interactions.



	1.2	Glycobiology: Types of carbohydrates, glycosidic bond and its stability, Structure and functions of glycoconjugates, proteoglycans, glycoproteins, glycolipids and homopolysaccharides.	
	1.3	Lipids: Classification of lipids, structure and functions of glycerolipids, ether lipids, galactolipids, sulfolipids, lipids in archaebacteria, sphingolipids, terpenes, isoprenoids.	
UNIT 2 Analytical Biochemistry	2.1	General methods of purification of proteins: Use of salting out / salting in, organic solvents, column chromatography, electrophoresis.	15 hours
(1 Credit)	2.2	Spectroscopic methods: Principle, Instrumentation and applications of Raman spectroscopy, IR spectroscopy, FTIR, Circular dichroism, NMR, ESR, X ray diffraction and mass spectroscopy	
	2.3	Radiolabeling techniques: Different types of radioisotopes, their detection, measurement and clinical applications.	

PRACTICAL

Course Title: Microbial Biochemistry

Course Code: SMCB511EP

COURSE OUTCOMES:

The learner will be able to :

- 1. extract cholesterol, separate fats by chromatography and determine iodine number of oils
- 2. isolate lactose and detect it using osazone test as well as estimate total sugar content by phenol sulphuric acid method
- 3. estimate polyphenol concentration in food stuff.
- 4. become competent in extracting, purifying and performing assay of enzyme amylase.

Lectures per week (1 Lecture is 120 minutes)	2
Total number of Hours in a Semester	60



Credits		2	
Evaluation System	Semester End Examination	2 Hours	50 marks
	Internal Assessment		

1	Extraction of total lipids.	60 hours
2	Isolation of cholesterol and lecithin from egg yolk.	00 110013
3	Identification of fatty acids and other lipids by TLC.	
4	Determination of degree of unsaturation of fats and oils.	
5	Isolation of lactose from bovine milk.	
6	Estimation of total sugars by phenol-sulphuric acid method.	
7	Isolation of glutamic acid from gluten.	
8	Isolation, Purification of Beta amylase from fungi using salting out and dialysis	
9	Estimation of beta amylase activity using the DNSA method.	
10	Estimation of protein content of beta amylase and calculation of specific activity.	
11	Estimation of polyphenols/tannins by Folin-Denis method	



12	Visit to an Instrumentation facility	

Programme: Sciences MICROBIOLOGY MSC-I	Semester – 1		
Course Title: Research Methodology	Course Code:SMCB511RM		
 COURSE OBJECTIVES: 1. To learn about the process of research, types of research and research design. 2. To learn about different types of sampling methods, sampling designs and variables. 3. To learn about methods of data collection, interpretation and report writing. 4. To learn about scientific writing and ethics in research and publication. 5. To use ICT as a tool to assist in writing research proposals and research outcomes. 6. To learn about the use of biostatistics software in interpretation of data. 			
 <u>COURSE OUTCOMES</u>: The learner will be able to design a research proposal. use appropriate methods of sample collection, methods of carrying out the research and write a report on the same. use anti plagiarism software to check if the proposal is acceptable, prepare a manuscript present research in a written / oral format using ICT. learn the use of biostatistics software so that it can be applied to the data collected for validity and interpretation. 			
Lectures per week (1 Lecture is 60 minutes)	4		
Total number of Hours in a Semester	60		
Credits 4			

Semester End Examination

Evaluation System



	Internal Assessment		50 marks
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UNIT 1 Basics of Research (1 Credit)	1.1	Meaning and objectives of research, research and scientific method, research process, research methods vs methodology. Criteria of good research, Problems encountered by researchers in India.	15 hours
	1.2	Types of research: conceptual vs empirical, applied vs fundamental, descriptive vs analytical, qualitative vs quantitative.	
	1.3	Research designs: Features of a good research design, different research designs. Case study, cross over study, case control design, cohort study design, multifactorial design, ex post facto	
UNIT 2 Sampling, data collection, interpretation and report writing. (1 Credit)	2.1	Sampling and sampling design: Steps and different types of sample design. Methods of sampling: non probability, simple random, systematic, stratified, quota, cluster and area sampling, multistage and sequential sampling. Problems due to unintended sampling, ecological and statistical population in the laboratory. Variables: Nominal, ordinal, discontinuous and continuous.	15 hours
	2.2	Collection of data: Methods and techniques of data collection. Types of data collection: Primary and Secondary. Methods of primary data collection: Observation, Experimentation, Questionnaire, Interview, Schedules, Case pilot study etc. Methods of secondary data collection- Internal and External.	
	2.3	Interpretation and report writing:	



		Techniques of interpretation and different steps involved in report writing, types of report, mechanics of writing a research report.	
3.1 UNIT 3 Scientific writing and Ethics in research and publication (1 Credit)		Abstract, Writing of Literature review, Aim and Objectives Methodology, References/ Bibliography and Preparation of manuscript for publication of research/ review paper.15 hourPeer reviewed, UGC CARE listed, indexed journals, citation index and role of citation, impact factor of a journal.15 hour	
	3.2	Use of computer in research: Computer technology, computer and researchers, software tools in the structure, design and preparation of thesis, layout, labeling of figures, legends, preparation of tables, layout, etc. Preparation of oral presentation and posters.	
	3.3	Ethics in research and publication: Citations, acknowledgement, conflict of interest, plagiarism, plagiarism checking tools. Overview of ethics in research: Overview of legislation and regulation, ethical guidelines in animal and clinical research. IPR and patent law.	
UNIT 4 Biostatistics (1 Credit)	4.1	Basics of Biostatistics: Measure of central tendencies, mean, mode, median. Measure of dispersion, Standard deviation, Standard error of means, P value concept.	15 hours



	Use of appropriate software for computation of statistical data.
4.2	Types of hypothesis: Basics concepts, types of hypothesis - Null and Alternate hypothesis, levels of hypothesis and testing of hypothesis. Parametric test: Z test, t test (1 tailed and 2 tailed test) of hypothesis. Different types of ANOVA test Non parametric test
4.3	Correlation analysis & Regression analysis: interpolation and extrapolation, nonlinear data fitting, probit analysis etc. Software used for all of the above.
	Student activity: A hands-on workshop will be organized to help students learn about the various biostatistics softwares. A talk will be organized to inform students on how to go about writing scientific articles to promote science journalism as a career choice.



ASSESSMENT PATTERN AND EVALUATION For NEP PG

A. Evaluation of Mandatory, Elective Courses and Common Course (Research Methodology) for MSc Part -1:

Assessment and evaluation pattern would be 50:50. There will be two subheads, namely, Summative Assessment (SA) and Continuous Assessment (CA) of 50 marks each for Mandatory courses.

- 1. Mandatory, elective and practical will have separate heads of passing.
- 2. A student needs to secure 40% marks for passing individually in SA and CA.
- 3. If a student fails, he/she will have to appear for an ATKT examination.
- **4.** Students who have missed the SA for a genuine reason (supported with a document subject to approval by the authorities) will appear for an Additional SA of 50 marks. This Additional/ATKT SA will be held after the declaration of the respective semester results and at the discretion of the PG exam committee.
- 5. Students will be declared FAIL if she scores less than 20 marks out of 50 marks.
- 6. Staff will show assessed answer papers of SA to students and discuss the rubric of assessment with them on a day fixed by the PG Exam Committee.
- 7. Grievance Redressal Mechanism for addressing grievances related to SA:

Students may apply for Reassessment, Photocopying and Revaluation of the SA answer books after the declaration of results in response to the notice posted by the College Office for the same.

8. Students with learning disabilities (LD) will be given extra time for SA as per the University rules.

B. Continuous Assessment (CA) for Mandatory Courses:

1. CA activities will be planned and conducted by the respective departments.

The departments are required to share the details of the CA activities with the Deputy Controller of PG exam and PG Co-ordinator (VP- Science).

- 2. Students' CA activity-related scores with assessed papers and feedback on their work (tests, other activities, assignments etc.) must be shared with students.
- 3. Format of CA for Mandatory courses: Two CA activities of 25 marks each.

CA 1: Test - 25 marks (Duration for answering the Test: Max. 60 Minutes)



CA 2: Any Activity - 25 marks

- 4. The minimum score to pass the Course will be 20 marks out of 50 marks.
- 5. If a student fails to pass (scores less than 20) then, the student will have to appear for

ATKT – one IA Test of 25 marks and one assignment of 25 Marks.

C. Evaluation for Elective and Common Courses (Research Methodology) under NEP:

1. Format of CA for Elective Courses: Two tests of 25 marks each of subjective type.

Only CA is to be conducted with 50 marks.

CA 1: Test - 25 marks (Duration for answering the Test: Max. 60 Minutes)

CA 2: Test- 25 marks (Duration for answering the Test: Max. 60 Minutes)

2. Format of CA for Common Courses:

CA 1: Test - 25 marks (Duration for answering the Test: Max. 60 Minutes)

CA 2: Any Activity - 25 marks

- **3.** If a student fails to pass (scores less than 20) then, the student will have to appear for 50 marks ATKT one **IA** Test of 25 marks and **one assignment** of 25 Marks.
- **4.** The minimum score to pass the Course will be 20 marks out of 50 marks. Students' CA activity-related scores with assessed papers and feedback (tests, other activities, assignments etc.) will be shared individually with students.
- 5. Grievance Redressal Mechanism for addressing grievances related to CAS:

Students will apply in a prescribed format to the respective Vice Principals. The grievance will be addressed by involving the concerned faculty and the other Exam Committee member/s deputed by the Principal.

REFERENCES

Mandatory Paper 1 SMCB511MJ

- 1. Becker, William M., Kleinsmith, Lewis J., & Hardin, Jeff. (2019). Becker World of Biology, 11th edn, *Pearson*.
- 2. Cann, Alan. (2015). Principles of Molecular Virology, 6th edn. Academic Press.



- 3. Fields, Bernard N., Knipe, David, M., Howley, Peter.M., & Griffin, Diane E. (2001). Fields Virology 4th edn, *Lippincott Williams and Wilkins*
- 4. Freifelder, David. (2004). Molecular Biology, 2nd edn. *Narosa Publishing House*.
- 5. Karp, Gerald. (2010). Cell and Molecular Biology, 6th edn. John Wiley & Sons, Inc.
- 6. Lodish, Harvey., Berk, Arnold., & Kaiser, Chris A. (2007). Molecular Cell Biology, 6th edn. *W.H. Freeman & Co Ltd.*
- Madigan, M., Martinko, J., Bender, K., Buckley, D., & Stahl, D. (2015). Brock Biology of Microorganisms 14th edn. *Pearson*.
- 8. Mahy, Brian WJ., & Regenmortel, Marc HV Van. (2010). Desk Encyclopedia of General Virology. *Elsevier*.
- 9. Shors, Teri. (2009). Understanding viruses, 1st edn. Jones and Bartlett Publishers.
- 10. Shors, Teri. (2016). Understanding viruses, 3rd edn. Jones and Bartlett Publishers.
- 11. Willey, Joanne M., Sherwood Linda M., & Woolverton Christopher J. (2014) Prescott's Microbiology, 9th edn, *McGraw-Hill Higher Education*.

Mandatory Paper 2 SMCB512MJ

- 1. Brooker, Robert. (2017). Genetics: Analysis and Principles, 6th edn. McGraw-Hill Higher Education
- 2. Lewin, Benjamin. (2004). Genes VIII. Pearson.
- 3. Lewin, Benjamin. (2007). Genes IX. Jones and Bartlett publishers.
- 4. Pierce, Benjamin A. (2003). Genetics- A Conceptual approach, Worth Publishers Inc., US.
- 5. Pierce, Benjamin A. (2013). Genetics- A Conceptual approach, 5th edn, W.H. Freeman
- 6. Russell, Peter J. (2010). iGenetics: A Molecular Approach, 3rd edn. *Pearson*.
- 7. Stanier, Roger Y., Adelberg, Edward A., & Ingraham, John L. (1976). General Microbiology, 4th edn. *Macmillan*.
- 8. Tamarin, Robert H. (2002). Principles of Genetics, 7th edn. McGraw-Hill.
- 9. Watson, James D., Baker, Tania A., Bell, Stephen P., Gann A., Levine, M., & Losick., R. (2003). Molecular Biology of the Gene, 5th edn. *Cold Spring Harbor Laboratory Press*.
- 10. Watson, James D., Baker, Tania A., Bell, Stephen P., Gann A., Levine, M., & Losick., R. (2013). Molecular Biology of the Gene, 7th edn. *Pearson*.
- 11. Watson, James D., Caudy Amy A., Myers, Richard M., & Witkowski Jan A. (2007) Recombinant DNA, Genes and Genomics A short course, 3rd edn. *W.H. Freeman and Company*.
- 12. Weaver, Robert F. (2012). Molecular Biology, 5th edn. *McGraw-Hill*.

ELECTIVE SMCB511E



- 1. Beedu, Rao., & Deshpande, S. Experimental biochemistry –A student companion. *IK international Pvt. Ltd.*
- Conn, Eric E., Stumpf, Paul K., Bruening, George., & Doi, Roy H. (2006). Outlines of Biochemistry, 5th edn, *Wiley India Edition*.
- 3. Jayaraman. Laboratory manual in biochemistry. New Age International Publishers.
- 4. Nelson, D., & Cox, M., (2005) Lehninger: Principles of Biochemistry, 4th edn., *New York, W.H. Freeman & Co.*
- 5. Pratt-Cornley. (2013). Essential Biochemistry (illustrated), 3rd edn. Wiley.
- 6. Price, Nicholas C., & Nairn, Jacqueline. (2009). Exploring proteins: Student's guide to experimental skills and methods. *Oxford University Press*.
- 7. Segel, I.R., (2004). Biochemical calculations, 2nd edn. John Wiley and Sons.
- 8. White, David. (2011). The physiology and biochemistry of prokaryotes, 4th edn, *Oxford University Press.*
- 9. Wilson, K., & Walker, J. (1994). Principles and techniques of practical biochemistry, 4th P edn, *Cambridge University Press*.

SMCB511RM

- 1. Daniel, W. W., & Cross, C. L. (2019). Biostatistics: A Foundation For Analysis In The Health Sciences, 10th edn, *Wiley Publications*.
- 2. Kothari, C.R. (2004). Research Methodology Methods and Techniques, 2nd Revised edn, *New Age International Publishers*.
- 3. Kumar, Ranjit.(2014). Research Methodology: A Step-by-Step guide for beginners, 5th edn, *SAGE Publications*
- 4. Le, Chap T. (2003). Introductory Biostatistics. John Wiley and Sons Ltd.
- 5. Pandey, P., & Pandey, M. Mishra. (2015) Research Methodology: Tools And Techniques. *Bridge Center*.
- Yip, C., Han, N.L.R., & Sng, B. L. (2016). Legal and ethical issues in research. *Indian J Anaesth*. 60(9): 684–688.



Programme: Sciences MICROBIOLOGY MSC-I	Semester – 2
Course Title: Virology And Cell Biology-II	Course Code:SMCB523MJ

COURSE OBJECTIVES:

- 1. To explain, discuss and analyze molecular biology and the life cycle of human viruses
- 2. To discuss the role of viruses in cancer and working with them in the research laboratory.
- 3. To develop an understanding of Prions and genetic experiments performed.
- 4. To describe cytoskeletal elements and their functions.
- 5. To summarize the development of multicellular organisms such as Drosophila melanogaster.
- 6. To explain eukaryotic cell cycle, mitosis and meiosis.
- 7. To explain signalling and communication in eukaryotic microorganisms including the yeast *Candida albicans*.
- 8. To discuss programmed cell death in eukaryotes, bacteria and yeasts.

COURSE OUTCOMES:

The learner will be able to

- 1. explain and analyze the replication and life cycle of different viruses, mechanism of retroviruses induce tumors, DNA tumor viruses, oncolytic viruses and Prion only hypothesis.
- 2. explain the structure and functions of Microtubules, Intermediate filaments and Microfilaments.
- 3. recall the development of model organism *Drosophila melanogaster* and role of different genes in its development.
- 4. explain the cell cycle and checkpoints and their significance, stages of mitosis and meiosis and connect the topics with the mandatory paper 2 topics such as Mendelian Genetics, Extensions of the same and Cancer.
- 5. explain and discuss cell signaling and signal transduction, MAP kinase pathway, and Ras signaling.
- 6. explain and compare programmed cell death in eukaryotes, bacteria and yeast

Lectures per week (1 Lecture is 60 minutes)			4	
Total number of Hours in a Semester			60	
Credits			4	
Evaluation System	Semester End Examination	2 Hours	50 marks	

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Internal Assessment

50 marks

UNIT 1 Human Viruses (1 Credit)		Structure, replication, life cycle and current affairs of the following viruses	
	1.1	dsDNA viruses - (04L) a. Poxviruses (Variola major and Vaccinia) b. Herpesviruses	15 hours
	1.2	dsRNA viruses- Rotavirus .	
	1.3	 Positive ssRNA viruses (05L) a. Rhinovirus b. Severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) c. Dengue virus 	
	1.4	Negative ssRNA viruses (04L) a. Rabies virus b. Measles virus	
UNIT 2 Tumor viruses and Prions (1 Credit)	2.1	 Tumor viruses (09L) Students to revise important definitions related to Cancer and characteristics of transformed cells a. Molecular mechanisms of virally induced tumor formation by RNA tumor viruses (Retroviruses) b. DNA tumor viruses - Hepatitis B virus, Human Papillomavirus, Adenoviruses, Simian Virus- 40 c. Oncolytic viruses 	15 hours
	2.2	 Prions (03L) a. History, case studies b. PRNP gene, Prion only hypothesis c. Biochemical analysis of the prion amino acid sequence 	



		d. Genetic Research and experiments with knockout mice.	
	2.3	Working with viruses in the research laboratory (03L)	
UNIT 3 Cytoskeleton and Development of multicellular organisms (1 Credit)	3.1	Cytoskeleton (11L) a. Microtubules i. Structure and composition ii. Microtubule-associated proteins iii. Motor proteins - kinesins, cytoplasmic dynein iv. Microtubule-organizing centers (MTOCs) v. The dynamic properties of microtubules b. Intermediate filaments i. Intermediate filament assembly and disassembly ii. Types and functions c. Microfilaments i. Microfilament assembly and disassembly ii. Myosin: the molecular motor of actin filaments e. Cytoskeletal elements in bacteria Development of Multicellular Organisms (04L) a. Genetics of Pattern formation in <i>Drosophila</i> i. Egg-polarity genes ii. Segmentation genes b. Homeobox genes in other organisms	15 hours
UNIT 4 Cellular reproduction, Signaling,	4.1	Cellular Reproduction (05L) a. The cell cycle b. Control of the cell cycle c. Mitosis	15 hours

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Communicati		d. Meiosis	
on and Programmed cell death (1 Credit)	4.2	 Signaling, communication and programmed cell death (10L) a. The basic elements of cell signaling systems b. G protein-coupled receptors and signal transduction by them c. Protein tyrosine phosphorylation as a mechanism for signal transduction d. Ras-MAP Kinase pathway e. Ras signaling in pathogenic yeast <i>Candida albicans</i> f. Apoptosis g. Programmed cell death in <i>E.coli</i> and <i>Saccharomyces cerevisiae</i> h. Lysis of the mother cell during sporulation of <i>Bacillus subtilis</i> 	

PRACTICAL Course Title: Virology And Cell Biology-II

Course Code: SMCB523MJP

COURSE OUTCOMES:

The learner will be able to :

- 1. correlate and recall the experiments done at the Virology Laboratories in the Research Institutes
- 2. recall the inoculation of an embryonated egg and cultivation of an animal virus observed during the visit
- 3. construct an assignment on any of the following viruses:Ebola virus, Nipah virus, West Nile virus, Mumps virus , Hepatitis C virus etc.
- 4. identify and distinguish between the different steps of Mitosis and Meiosis
- 5. detect sporulation and germination in *Bacillus species*, and use Haemocytometer to determine the spore count

Lectures per week (1 Lecture is 120 minutes)	2
Total number of Hours in a Semester	60
Credits	2



Evaluation System	Semester End Examination	2 Hours	50 marks
	Internal Assessment		

1	Visit to NIRRH or Haffkine research institute or Animal tissue culture laboratory.	60 hours
2	Demonstration - Egg inoculation and cultivating animal virus in embryonated eggs.	
3	Assignment on any one of the following viruses - Ebola virus, Nipah virus, West Nile virus, Mumps virus, Hepatitis C virus.	
4	Study of Mitosis.	
5	Study of Meiosis.	
6	Sporulation and germination in Bacillus subtilis.	

Programme: Sciences MICROBIOLOGY MSC-I	Semester – 2
Course Title: GENETICS-II	Course Code:SMCB524MJ
COUDSE OD IECTIVES.	

COURSE OBJECTIVES:

- 1. To discuss Mendelian genetics, principles of inheritance and extensions of and deviations from Mendelian genetics and solve problems related to the topics.
- 2. To develop an understanding of concepts and principles associated with population genetics
- 3. To explain the genetic basis of cancer.
- 4. To describe the Transposable genetic elements in prokaryotes and eukaryotes.



- 5. To explain the techniques used for study of genetics.
- 6. To discuss basics and applications of bioinformatics.

COURSE OUTCOMES:

The learner will be able to

- 1. explain the Mendelian principles and acquire knowledge of its extensions and deviations.
- 2. solve the problems on Mendelian Genetics and develop critical thinking.
- 3. discuss the principles of population genetics.
- 4. explain the genetic basis of cancer.
- 5. describe the Transposable genetic elements in prokaryotes and eukaryotes.
- 6. compare the techniques used for study of genetics.
- 7. explain the basics of computational biology and apply the knowledge to solve practical problems

Lectures per week (1 Lecture	is 60 minutes)	4	
Total number of Hours in a S	emester	60	
Credits		4	
Evaluation System	Semester End Examination	2 Hours	50 marks
	Internal Assessment		50 marks

UNIT 1	1.1	Mendelian Genetics (05 L)	15 hours
Mendelian		a. Mendel's experimental design	
Genetics,		b. Monohybrid crosses and Mendel's principle of	
Extensions of		Segregation	
Mendelian		i. Branch diagram of monohybrid crosses	
Genetics and		ii. Use of testcrosses	
Extranuclear		c. Dihybrid crosses and Mendel's principle of	
Inheritance		Independent Assortment	
(1 Credit)		i. Branch diagram of dihybrid crosses	
		d. Trihybrid crosses	
		e. Mendelian genetics in Humans- Pedigree analysis	
		f. Problems on Mendelian Genetics	



	1.2	Extensions of and Deviations from Mendelian Genetic Principles (10 L) a. Multiple Alleles b. Modification of dominance relationships i. Incomplete dominance ii. Codominance iii. Molecular explanations c. Essential genes and lethal alleles d. Gene expression and environment e. Epistasis i. Recessive epistasis ii. Dominant epistasis f. Problems on extensions of Mendelian Genetics g. Extranuclear Inheritance (non-Mendelian) i. Extranuclear genomes ii. Rules of extranuclear inheritance iii. Examples of extranuclear inheritance	
UNIT 2 Population Genetics, Transposable genetic elements and Cancer (1 Credit)	2.1	 Population Genetics (06 L) a. Genotypic and allelic frequencies b. Calculation of genotypic and allelic frequencies for autosomal and X linked loci c. Hardy-Weinberg Law and calculation of genotypic frequency at Hardy Weinberg equilibrium d. Factors affecting genotypic and allelic frequencies e. Changes in genetics structure of populations (mutation, migration & gene flow, genetic drift and natural selection) f. Measuring genetic variation 	15 hours
	2.2	 a. Transposable genetic clements (05 L) a. Transposable elements in prokaryotes: An overview b. The medical significance of bacterial transposons c. Transposable elements in eukaryotes i. Ac and Ds elements in Maize 	



		 ii. P elements and hybrid dysgenesis in Drosophila iii. Mariner, an ancient and widespread transposon d. Retrotransposons Retrovirus like elements Retroposons e. The genetic and evolutionary significance of transposable elements Transposons as mutagens Transposons and genome organization 	
	2.3	 Genetic basis of cancer (04 L) a. Cancer- Introduction b. Mutations in different types of genes c. Change in chromosome number and structure, d. Changes in DNA methylation e. Sequential mutations 	
UNIT 3 Genomics (1 Credit)	3.1	Identifying genes (03) a. Techniques i. Positional Cloning ii. Exon trapping and CpG Islands b. Mutated genes associated with human disease- Huntington disease	15 hours
	3.2	 Techniques in genomic sequencing (05) a. Traditional DNA sequencing methods- Sangers sequencing (Dideoxynucleotide method) and Maxam-Gilbert Sequencing b. Next Generation Sequencing (NGS) methods - Primer Walking, Pyrosequencing, Sequencing Using Reversible Chain Terminators and Sequencing by Ligation 	



	3.3	Strategies for large-scale sequencing of genomes (02)a. Shotgun Cloning Strategyb. Cyclic Array Sequencing	
	3.4	Functional Genomics / Transcriptomics (02)a. Gene Expression profiling- DNA microarraysb. Serial Analysis of Gene Expression (SAGE)	
	3.5	 Comparative Genomics (03) a. Bacteria Minimal/ small genome Horizontal gene transfer- significance in comparative genomics Significance of comparative genomics in studying microbial evolution. 	
UNIT 4 Bioinformati cs (1 Credit)	4.1	 a. Introduction to bioinformatics, scope and applications b. Databases and tools/software Nucleotide sequence databases Protein sequence databases Protein sequence databases c. Sequence alignment and alignment scores Pairwise – Global and local sequence alignment Algorithms - Needleman–Wunsch, Smith–Waterman, BLAST and FASTA Multiple sequence alignment- ClustalW d. Identification of genes on prokaryotic DNA Prediction of genes in genome sequences- insilico methods 	15 hours



 f. Phylogenetic analysis Distance based methods Maximum likelihood method Bayesian phylogenetics Parsimony-based methods g. Protein classification and structure prediction Domain identification and annotation (e.g., Pfam) Protein structure databases- PDB_CATH and 	
i. Domain identification and annotation (e.g.,Pfam)ii. Protein structure databases- PDB, CATH and	
SCOP e. Structure visualization f. Packages for genomic analysis	
g. Introduction to Linux, Python and K programming	

PRACTICAL
Course Title: Genetics-IICourse Code: SMCB524MJP

COURSE OUTCOMES:

The learner will be able to :

- 1. analyze, classify and solve problems on Mendelian Genetics, Population Genetics and Restriction mapping
- 2. perform DNA transformation and plasmid curing experiments and apply these experiments in molecular biology research in future
- 3. isolate and purify genomic DNA from bacteria and lymphocytes and confirm its presence using UV-visible spectrophotometry
- 4. design primers to carry out the amplification of genes using Polymerase chain reaction
- 5. apply the tools and softwares of Bioinformatics in computational biology research
- 6. do a workshop or an online course in Molecular Biology or Genetics and apply the knowledge.

Lectures per week (1 Lecture is 120 minutes)	2
Total number of Hours in a Semester	60
Credits	2



Evaluation System	Semester End Examination	2 Hours	50 marks
	Internal Assessment		

1	Problems on Mendelian genetics.	
2	Problems on Population genetics.	60 hours
3	DNA Transformation.	
4	Curing of plasmids.	
5	Isolation of genomic DNA from bacteria and lymphocytes.	
6	Problems on restriction mapping	
7	Design of primer & PCR.	
8	Bioinformatics practicals - i. Exploring DNA databases and analysis of gene record ii. Introduction to LINUX, R and Python commands, iii. Construction of Phylogeny Tree using Clustal omega	
9	Online course related to any aspect of Genetics OR Workshop on Molecular Biology/Genetics in an institute.	

Programme: Sciences MICROBIOLOGY MSC-I	Semester – 2	
Course Title: Food Microbiology	Course Code:SMCB522E	



COURSE OBJECTIVES:

- 1. To list microorganisms that are commonly associated with fermented foods
- 2. To outline the process for making fermented foods & understand the benefits of using fermentation as a food processing method, also appreciate the similarities and difference among fermentations of dairy and vegetable products.
- 3. To evaluate claims about health benefits of probiotic bacteria.
- 4. To recognize the difference between methods available for microbiological analysis of food and compare the methods in terms of advantages and disadvantages.
- 5. To discuss the importance of HACCP system with respect to food safety and quality

COURSE OUTCOMES:

The learner will be able to

- 1. relate the steps of bread, cheese, idli & sauerkraut making to microbial fermentation and final characteristics.
- 2. prepare food samples for determination of microbial load, understand why some sampling plans are more stringent than others and choose appropriate sampling plans as per case number.
- 3. differentiate among conventional and rapid methods of detection of pathogens.
- 4. explain the basis of immunological, nucleic acid, and biochemical methods and recognize appropriate rapid method suitable for specific use
- 5. differentiate among the various microbiological criteria
- 6. recognize how indicator organisms are used in microbiological criteria
- 7. identify and list steps required to manage microbiological hazards in foods
- 8. outline the basic concepts of GMPs and recognize its limitations
- 9. understand the process for development of a HACCP program
- 10. identify role of national and international agencies involved in food safety and quality

Lectures per week (1 Lecture is 60 minutes)		2	
Total number of Hours in a Semester		30	
Credits		2	
Evaluation SystemSemester End ExaminationInternal Assessment			
			50 marks



UNIT 1 Application s of Microorga nisms in food industry (1 Credit)	1.1	 Microbiology of fermented foods (08L): a. Starter cultures: Bacterial, Yeast and molds. Concentrated cultures, Problems in starter cultures and control methods b. General method of production Bread Idli Cheese – Types, Production of Cheddar, Swiss and Blue cheese Fermented vegetable products – Sauerkraut Popular Oriental fermented foods: Kimchi, Soy sauce, Tempeh 	15 hours
	1.2	 Microbial products used in food industry (07L): a. Enzymes in food processing, b. Food grade pigments, Flavour compounds, Exopolysaccharides c. Microbes used as Probiotics (Examples, properties and benefits) 	
UNIT 2 Microbiologic al quality of Food (1 Credit)	2.1	 Detection and enumeration of microbes in food (08L): a. Conventional Methods i. Direct Enumeration :Microscopic Counts, Count using nonselective , selective, differential chromogenic media ii. Indirect count : Dilution to extinction, MPN, Dye Reduction test b. Detection of Microbial Toxins c. Rapid and automated methods for detection of Pathogens: Metabolic Fingerprinting, Immunomagnetic Separation, Reverse Passive Latex Agglutination (RPLA) Method, Immunochromatographic Lateral Flow Assay, Hybridization Method , Microarrays and Mass-Spectrometry d. Bacteriophage for detection of pathogens 	15 hours



	e. Biosensors for detection of microbes in food.	
2.2	 Food Quality and Safety (07L) a. New emerging food borne pathogens of concern. b. Control at source. c. Indicator microorganisms : Characteristics, Coliform and enterococci. d. Microbiological Criteria. e. Sampling plan, Types (2 class and 3 class) and sampling procedures f. HACCP system g. Regulations and agencies monitoring microbiological safety of food: ICMSF, CDC, Food net, Codex Alimentarius, ISO22000, FSSAI 	

PRACTICAL Course Title: Food Microbiology

Course Code: SMCB522EP

COURSE OUTCOMES:

The learner will be able to :

- 1. determine the microbial load and changes in the population of lactic acid bacteria during Sauerkraut fermentation.
- 2. isolate probiotic bacterium from fermented dairy products using Rogosa agar and check its ability to produce bacteriocin.
- 3. detect microorganisms with lipase / amylase/ protease activity on Gorodkowa's, starch agar and milk agar respectively.
- 4. design and conduct an experiment to comment on the effect of any one parameter (Temperature, time, ratio of ingredients, type of ingredients) on the leavening of bread by *Saccharomyces cerevisiae*.
- 5. prepare food samples for determination of microbial load and determine the APC and coliform count in carrot and apple juice, salad, mayonnaise to comment on hygienic quality and shelf life
- 6. carry out Quality Assessment and Analysis of Milk (Raw, Packed) by performing DMC, RPT and SPC / LPC, Thermophilic/Psychrophilic, yeast-mold counts
- 7. detect and identify pathogens associated with frozen fish/poultry/meat using specific selective / differential chromogenic media.
- 8. conduct literature survey on latest novel detection methods for food borne pathogens/ toxins.



Lectures per week (1 Lecture is 120 minutes)		2	
Total number of Hours in	n a Semester	60	
Credits		2	
Evaluation System Semester End Examination		2 Hours	50 marks
Internal Assessment			

1 Microbiological study of fermented foods (Sauerkraut)		
2	Isolation of Probiotic bacteria and checking the antimicrobial effect of the bacteriocin produced by probiotic organisms.	60 hours
3	Production of Microbial Enzymes of commercial importance.	
4	Student activity: Bread Production and studying the effect of various factors affecting Bread Production (5 marks)	
5	Microbiological load in juices, salad, mayonn- aise.	
6	Quality Assessment and Analysis of food :Milk (Raw, Packed)	
7	Detection of pathogens in frozen fish/poultry/ meat	
8	Report to be written in journal on Novel detection methor food borne pathogens/ toxins.	



Programme: Sciences MICROBIOLOGY MS	Semester – 2			
Course Title: Field Proje (OJT)	Course Code: SMCB521OJT			
 <u>COURSE OBJECTIVES:</u> 1. To develop and establish practical skills during the internship/on job training at an indust hospital, pathology laboratory etc. 2. To prepare a report on the same and present the experiments and skills learnt during t internship in the form of a Powerpoint presentation 				
 <u>COURSE OUTCOMES</u>: The learner will be able to 1. develop skills and apply the knowledge in the future 2. write a report on the internship/On job training and present in the form of a Powerpoint presentation. 				
Hours per week		8		
Total number of Hours in a Semester		120		
Credits			4	
Evaluation System	Semester End Examination	-	-	
	Internal Assessment	-	-	



ASSESSMENT PATTERN AND EVALUATION For NEP PG

A. Evaluation of Mandatory, Elective Courses and Common Course (Research Methodology) for MSc Part -1:

Assessment and evaluation pattern would be 50:50. There will be two subheads, namely, Summative Assessment (SA) and Continuous Assessment (CA) of 50 marks each for Mandatory courses.

- 1. Mandatory, elective and practical will have separate heads of passing.
- 2. A student needs to secure 40% marks for passing individually in SA and CA.
- **3.** If a student fails, he/she will have to appear for an ATKT examination.
- **4.** Students who have missed the SA for a genuine reason (supported with a document subject to approval by the authorities) will appear for an Additional SA of 50 marks. This Additional/ATKT SA will be held after the declaration of the respective semester results and at the discretion of the PG exam committe.
- 5. Students will be declared FAIL if she scores less than 20 marks out of 50 marks.
- 6. Staff will show assessed answer papers of SA to students and discuss the rubric of assessment with them on a day fixed by the PG Exam Committee.
- 7. Grievance Redressal Mechanism for addressing grievances related to SA:

Students may apply for Reassessment, Photocopying and Revaluation of the SA answer books after the declaration of results in response to the notice posted by the College Office for the same.

8. Students with learning disabilities (LD) will be given extra time for SA as per the University rules.

B. Continuous Assessment (CA) for Mandatory Courses:

1. CA activities will be planned and conducted by the respective departments.

The departments are required to share the details of the CA activities with the Deputy Controller of PG exam and PG Co-ordinator (VP- Science).



- 2. Students' CA activity-related scores with assessed papers and feedback on their work (tests, other activities, assignments etc.) must be shared with students.
- 3. Format of CA for Mandatory courses: Two CA activities of 25 marks each.

CA 1: Test - 25 marks (Duration for answering the Test: Max. 60 Minutes)

CA 2: Any Activity - 25 marks

- 4. The minimum score to pass the Course will be 20 marks out of 50 marks.
- 5. If a student fails to pass (scores less than 20) then, the student will have to appear for ATKT one IA Test of 25 marks and one assignment of 25 Marks.

C. Evaluation for Elective Courses under NEP:

1. Format of CA for Elective Courses: Two tests of 25 marks each of subjective type.

Only CA is to be conducted with 50 marks.

CA 1: Test - 25 marks (Duration for answering the Test: Max. 60 Minutes)

CA 2: Test- 25 marks (Duration for answering the Test: Max. 60 Minutes)

REFERENCES

SMCB523MJ Mandatory 1

- 1. Burrell, Christopher., Howard, Colin., & Murphy, Frederick. (2016). Fenner and White's Medical Virology, 5th edn. *Academic Press*.
- 2. Cann, Alan. (2015). Principles of Molecular Virology, 6th edn. Academic Press.
- 3. Farrugia, Gianluca., & Balzan, Rena. (2012). Oxidative stress and programmed cell death in yeast. *Front. Oncol.*
- 4. Graumann, Peter L. (2007). Cytoskeletal Elements in Bacteria. Annu. Rev. Microbiol. 61:589-618.
- 5. Hardin, Jeff., Bertoni, Gregory., & Kleinsmith, Lewis J. (2016). Becker's World of the Cell, 8th Global edn. *Pearson*.
- 6. Hardin, J., Bertoni, G., & Kleinsmith, L. J. (2019). Becker's World of Cell Biology (9th ed.). Pearson
- Jacobs, Samantha E., Lamson, Daryl M., St. George, Kirsten., & Walsh, Thomas J. (2013). Human Rhinoviruses. *Clinical Microbiology Reviews*. Volume 26 Number 1, p. 135–162.
- 8. Karp, Gerald. (2010). Cell and Molecular Biology, 6th edn. John Wiley & Sons, Inc.



- 9. Le Kerr, Shannic., Mathew, Cynthia., & Ghildyal, Reena. (2021). Rhinovirus and Cell Death. *Viruses*. 13, 629.
- 10. Lewis, Kim. (2000). Programmed Death in Bacteria. *Microbiology and molecular biology reviews*. 1092(2172):503-514.
- 11. McDonald, Sarah M., & Patton, John T. (2011). Assortment and packaging of the segmented rotavirus genome. *Trends Microbiol*. 19(3): 136–144.
- 12. Pentland, Daniel R., Piper-Brown, Elliot., Mühlschlegel, Fritz A., & Gourlay, Campbell W. (2017). Ras signalling in pathogenic yeasts. *Microbial Cell*. Vol 5(2):63-74.
- 13. Pierce, B. (2008). Genetics- a conceptual approach, 3rd edn, W.H. Freeman and company.
- 14. Rota, Paul A., Moss, William J., Takeda Makoto., de Swart, Rik L., Thompson, Kimberly M., & Goodson, James L. (2016). Measles. *Primer. Nature Reviews*. Volume 2, Article 16049.
- 15. Shors, Teri. (2009). Understanding viruses, 1st edn. Jones and Bartlett Publishers.
- 16. Shors, Teri. (2016). Understanding viruses, 3rd edn. Jones and Bartlett Publishers.
- 17. Stobart, Christopher C., Nosek, Jenna M., & Moore, Martin L. (2017) Rhinovirus Biology, Antigenic Diversity, and Advancements in the Design of a Human Rhinovirus Vaccine. *Frontiers in Microbiology*. Volume 8, Article 2412.

SMCB524MJ Mandatory 2

- 1. Attwood Teresa K., & Parry-Smith David J. (1999). Introduction to Bioinformatics. 1st edn. *Addison Wesley Longman Limited*.
- 2. Glick, Bernard R., Pasternak, Jack J., & Patten, Cheryl L. (2010). Molecular Biotechnology: Principles and Applications of Recombinant DNA, 4th edn. *ASM Press*.
- 3. Klug William S., & Cummings Michael R. (2000). Concepts of genetics. 6th edn. Prentice Hall.
- Krane Dan E., & Raymer Michael L. (2002). Fundamental Concepts of Bioinformatics, 1st edn. Benjamin Cummings.
- 5. Mount, David W. (2004). Bioinformatics: sequence and genome analysis, 2nd edn. CSHL Press
- 6. Pierce, B. (2008). Genetics- a conceptual approach, 3rd edn, W. H. Freeman and Company.
- 7. Primrose, S. B., & Twyman, R. M. (2006). Principles of Gene Manipulation, 7th edn. *Blackwell Publishing*
- 8. Russell, Peter J. (1998). Genetics, 5th edn, Benjamin Cummings.
- 9. Russell, Peter J. (2010). iGenetics: A Molecular Approach, 3rd edn. Pearson.
- Snustad, Peter D., & Simmons, Michael J. (2003). Principles of Genetics, 3rd edn. John Wiley & Sons, Inc.
- 11. Snustad, Peter D., & Simmons, Michael J. (2012). Principles of Genetics, 6th edn. *John Wiley & Sons, Inc.*
- 12. Weaver, Robert F. (2012). Molecular Biology, 5th edn. McGraw-Hill.



- 13. Watson, J.D., Caudy, A.A., Myers, R.M., & Witkowski, J.A. (2007). Recombinant DNA Genes and Genomes: A Short Course. 3rd edn. *W.H. Freeman and Company*.
- 14. Xiong, Jin. (2006). Essential Bioinformatics. Cambridge: Cambridge University Press

SMCB522E Elective

- 1. Adams, M R., & Moss, M O. (2007). Food Microbiology, 3rd edn. New age international publishers.
- 2. Jay, James M., Loessner, Martin J., & Golden, David A. (2005). Modern Food Microbiology 7th edn, *Springer*.
- 3. Montville, Thomas J., Matthews, Karl R., & Kniel, Kalmia E.(ed). (2012). Food Microbiology: An Introduction, 3rd edn. *ASM press*.
- 4. Ray, Bibek., & Bhunia, Arun . (2014). Fundamental Food Microbiology 5th edn. CRC Press.
- 5. Varzakas, Theodoros., & Tzia, Constantina. (2016). Handbook of Food Processing, *CRC press-Taylor* –*Francis group*.