

## KARNATAK UNIVERSITY, DHARWAD ACADEMIC (S&T) SECTION

ಕರ್ನಾಟಕ ವಿಶ್ವವಿದ್ಯಾಲಯ, ಧಾರವಾಡ ವಿದ್ಯಾಮಂಡಳ (ಎಸ್&ಟಿ) ವಿಭಾಗ



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NAAC Accredited

website: kud.ac.in

No. KU/Aca(S&T)/JS/MGJ(Gen)/2023-24/59

Date: 04 09 2023

### ಅಧಿಸೂಚನೆ

ವಿಷಯ: 2023–24ನೇ ಶೈಕ್ಷಣಿಕ ಸಾಲಿನಿಂದ ಎಲ್ಲ ಸ್ನಾತಕ ಪದವಿಗಳಿಗೆ 5 ಮತ್ತು 6ನೇ ಸೆಮೆಸ್ಟರ್ NEP-2020 ಪಠ್ಯಕ್ರಮವನ್ನು ಅಳವಡಿಸಿರುವ ಕುರಿತು.

ಉಲ್ಲೇಖ: 1. ಸರ್ಕಾರದ ಅಧೀನ ಕಾರ್ಯದರ್ಶಿಗಳು(ವಿಶ್ವವಿದ್ಯಾಲಯ 1) ಉನ್ನತ ಶಿಕ್ಷಣ ಇಲಾಖೆ ಇವರ ಆದೇಶ ಸಂಖ್ಯೆ: ಇಡಿ 104 ಯುಎನ್ಇ 2023, ದಿ: 20.07.2023.

2. ವಿದ್ಯಾವಿಷಯಕ ಪರಿಷತ್ ಸಭೆಯ ನಿರ್ಣಯ ಸಂಖ್ಯೆ: 2 ರಿಂದ 7, ದಿ: 31.08.2023.

3. ಮಾನ್ಯ ಕುಲಪತಿಗಳ ಆದೇಶ ದಿನಾಂಕ: ೦५ (೦೨ (2023

ಮೇಲ್ಕಾಣಿಸಿದ ವಿಷಯ ಹಾಗೂ ಉಲ್ಲೇಖಗಳನ್ವಯ ಮಾನ್ಯ ಕುಲಪತಿಗಳ ಆದೇಶದ ಮೇರೆಗೆ, 2023–24ನೇ ಶೈಕ್ಷಣಿಕ ಸಾಲಿನಿಂದ ಅನ್ವಯವಾಗುವಂತೆ, ಎಲ್ಲ B.A./ BPA (Music) /BVA / BTTM / BSW/ B.Sc./B.Sc. Pulp & Paper Science/ B.Sc. (H.M)/ BCA/ B.A.S.L.P./ B.Com/ B.Com (CS) / BBA & BA ILRD ಸ್ನಾತಕ ಪದವಿಗಳ 5 ಮತ್ತು 6ನೇ ಸೆಮೆಸ್ಟರ್ಗಳಿಗೆ NEP-2020ರ ಮುಂದುವರೆದ ಭಾಗವಾಗಿ ವಿದ್ಯಾವಿಷಯಕ ಪರಿಷತ್ ಸಭೆಯ ಅನುಮೊದಿತ ಕೋರ್ಸಿನ ಪಠ್ಯಕ್ರಮಗಳನ್ನು ಕ.ವಿ.ವಿ. ಅಂತರ್ಜಾಲ www.kud.ac.in ದಲ್ಲಿ ಭಿತ್ತರಿಸಲಾಗಿದೆ. ಸದರ ಪಠ್ಯಕ್ರಮಗಳನ್ನು ಕ.ವಿ.ವಿ. ಅಂತರ್ಜಾಲದಿಂದ ಡೌನಲೋಡ ಮಾಡಿಕೊಳ್ಳಲು ಸೂಚಿಸುತ್ತ ವಿದ್ಯಾರ್ಥಿಗಳ ಹಾಗೂ ಸಂಬಂಧಿಸಿದ ಎಲ್ಲ ಬೋಧಕರ ಗಮನಕ್ಕೆ ತಂದು ಅದರಂತೆ ಕಾರ್ಯಪ್ರವೃತ್ತರಾಗಲು ಕವಿವಿ ಅಧೀನದ/ಸಂಲಗ್ನ ಮಹಾವಿದ್ಯಾಲಯಗಳ ಪ್ರಾಚಾರ್ಯರುಗಳಿಗೆ ಸೂಚಿಸಲಾಗಿದೆ.

ಅಡಕ: ಮೇಲಿನಂತೆ

ಖಲ್ಗಳ ಇತ್ತಿತಿಕೆ ಕೆಲಸಚಿವರು.

... ಕರ್ನಾಟಕ ವಿಶ್ವವಿದ್ಯಾಲಯದ ವ್ಯಾಪ್ತಿಯಲ್ಲಿ ಬರುವ ಎಲ್ಲ ಅಧೀನ ಹಾಗೂ ಸಂಲಗ್ನ ಮಹಾವಿದ್ಯಾಲಯಗಳ ಪ್ರಾಚಾರ್ಯರುಗಳಿಗೆ. (ಕ.ವಿ.ವಿ. ಅಂರ್ತಜಾಲ ಹಾಗೂ ಮಿಂಚಂಚೆ ಮೂಲಕ ಬಿತ್ತರಿಸಲಾಗುವುದು)

#### ಪ್ರತಿ:

- 1. ಕುಲಪತಿಗಳ ಆಪ್ತ ಕಾರ್ಯದರ್ಶಿಗಳು, ಕ.ವಿ.ವಿ. ಧಾರವಾಡ.
- 2. ಕುಲಸಚಿವರ ಆಪ್ತ ಕಾರ್ಯದರ್ಶಿಗಳು, ಕ.ವಿ.ವಿ. ಧಾರವಾಡ.
- 3. ಕುಲಸಚಿವರು (ಮೌಲ್ಯಮಾಪನ) ಆಪ್ತ ಕಾರ್ಯದರ್ಶಿಗಳು, ಕ.ವಿ.ವಿ. ಧಾರವಾಡ.
- 4. ಅಧೀಕ್ಷಕರು, ಪ್ರಶ್ನೆ ಪತ್ರಿಕೆ / ಗೌಪ್ಕ / ಜಿ.ಎ.ಡಿ. / ವಿದ್ಯಾಂಡಳ (ಪಿ.ಜಿ.ಪಿಎಚ್.ಡಿ) ವಿಭಾಗ, ಸಂಬಂಧಿಸಿದ ಕೋರ್ಸುಗಳ ವಿಭಾಗಗಳು ಪರೀಕ್ಷಾ ವಿಭಾಗ, ಕ.ವಿ.ವಿ. ಧಾರವಾಡ.
- 5. ನಿರ್ದೇಶಕರು, ಕಾಲೇಜು ಅಭಿವೃದ್ಧಿ / ವಿದ್ಯಾರ್ಥಿ ಕಲ್ಯಾಣ ವಿಭಾಗ, ಕ.ವಿ.ವಿ. ಧಾರವಾಡ.



# **B.Sc.in MICROBIOLOGY**

## **SYLLABUS**

WithEffectfrom2023-24

DISCIPLINE SPECIFIC CORE COURSE (DSCC) FOR SEM V &VI,

SKILL ENHANCEMENT COURSE(SEC) FOR SEMV

INTERNSHIP FOR SEM VI

**ASPER NEP-2020** 

## **Karnatak University, Dharwad** B.Sc.in **MICROBIOLOGY**

Effective from 2023-24

					Instruc	Total	Dunation		lts		
Sem.	Type of Course	Theory/P ractical	Course Code	Course Title	tion hour/w eek	hours /sem	Duration Of Exam	Format ive	Sum mativ e	Total	Credits
	DSCC-9	Theory	035MCB011	Microbial genetics and molecular biology	04hrs	56	02hrs	40	60	100	04
	DSCC-10	Practical	035MCB012	Microbial genetics and molecular biology	04hrs	56	03hrs	25	25	50	02
	DSCC-11	Theory	035MCB013	Food and dairy microbiology	04hrs	56	02hrs	40	60	100	04
V	DSCC-12	Practical	035MCB014	Food and dairy microbiology	04hrs	56	03hrs	25	25	50	02
	Other Subject										04
	Other Subject										04
	Other Subject										04
	SEC-3	Practical	035MCB061	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	04hrs	56	03hrs	25	25	50	02
				industries(Practical)  Total				275	375	650	24
	DCCC 12	TP1	02CMCD011	1 0001	04hrs	56	0.21	40		100	04
VI	DSCC-13	Theory	036MCB011	Immunology and medical microbiology	04nrs	36	02hrs	40	60	100	04
	DSCC-14	Practical	036MCB012	Immunology and medical microbiology	04hrs	56	03hrs	25	25	50	02
	DSCC-15	Theory	036MCB013	Genetic engineering and industrial microbiology	04hrs	56	02hrs	40	60	100	04
	DSCC-16	Practical	036MCB014	Genetic engineering and industrial microbiology	04hrs	56	03hrs	25	25	50	02
	Other Subject										04
	Other Subject										04
	Other Subject										04
	Internship-1		036MCB091		04hrs	56	03hrs	50	0	50	02
		I	I .	Total		I		300	350	650	26

## DisciplineSpecific Core CourseDSCC-9

Course Title: Microbial genetics and molecular biology

### Course Code:035 MCB 011

DSCC-9	Theory	04	04	56hrs.	2hrs.	40	60	100
				/Semester		arks	Marks	
ofCourse	/Practical	Credits	rperweek	res/Hours	Exam	ssessmentM	assessment	rks
Type	Theory		Instructionhou	TotalNo.ofLectu	Durationof	FormativeA	Summative	TotalMa

## CourseObjectives

- 1. UnderstandthefundamentalprinciplesandtechniquesofMolecular Biology.
- 2. StayupdatedonemergingtrendsandadvancementsinMolecular Biology.

## CourseOutcomes (COs): Afterthesuccessful completion of the course, the student will be ableto:

- 1. Understandconceptsinvolvedinreplication,transcription,translation,regulationofgeneexpressionin bacteriaandEukaryotes.
- 2. Differentiatetheprocessofreplication,transcription,translation,regulationofgeneexpressioninbacteria and Eukaryotes.
- 3. Understandthegenetic switchin bacteriophages.
- 4. Compareandcontrasthousekeeping,constitutive,inducibleandrepressiblegenesCO5.Outlineregulatory mechanisms in bacteriato control cellularprocesses.

UNIT I: Mendelian and Classical Genetics	14hr
Mendel's principles of inheritance: Special features of pea plants as an ideal system to study genetics and Mendel's cross breeding experimental approach to prove genetic	
principles. Principles of dominance and Segregation; phenotype, genotype, traits	
controlled by genes, existence of alleles (dominant and recessive).  Principle of segregation of alleles during the fertilization: Monohybrid (single	
character) cross, F1 and F2 generation, heterozygous, homozygous, test cross and back	
cross to test genotype of F1 plants.  Principle of independent assortment; Dihybrid (two characters) cross, pattern of	
assortment of alleles. Chromosomal basis of inheritance; chromosome number,	
haploid (n), diploid (2n). Chromosomal theory of Heredity; Experimental evidence linking the inheritance of	
genes to chromosomes, Chromosomes as arrays of genes, Chromosomal basis of Mendel's principles of segregation and independent assortment.	
UNIT II: Molecular basis of Life and Genetic recombination	14hr
Molecular basis of Life:	
Historical developments of DNA as a genetic material; Griffith experiment of	
Transformation, Proof that genetic information stored in DNA, Enzymatic approach to prove DNA mediates transformation by A very, MacLeod and McCarty, Hershey and	
Chase experiment to prove DNA carries the genetic information in T2 bacteriophage.	
RNA stores the genetic information in some viruses, viroids and prions. Structure of Watson Crick model of DNA, Plasmid DNA.	
Organization of genes in mitochondria and chloroplast.	
Genetic Recombination:	
Bacterial Transformation: Types of transformation mechanisms found in prokaryotes,	
Natural and artificial methods of transformation. Bacterial Conjugation: U-tube experiment to prove physical contact between bacteria is essential for gene transfer,	
properties of the F plasmid, F <sup>+</sup> x F <sup>-</sup> conjugation, sexductionF'x F <sup>-</sup> conjugation, Hfr x F <sup>-</sup>	
conjugation, Gene mapping in bacteria by conjugation. Transduction: Generalized and specialized transduction, plasmids and episomes. Genetics of Fungi: life cycle of Yeast	
and Neurospora, Terad analysis, unordered tetrad analysis in yeast, ordered tetrad	
analysis in Neurospora, two point and three-point test cross, detecting linkage and	
mapping genes in yeast and neurospora.  UNIT III: DNA Replication and protein synthesis	14hrs
<b>DNA Replication:</b> IntroductionandMechanism of DNA replication, enzymes involved	
in replication and regulation.	
<b>Prokaryotic transcription:</b> Transcription bubble, Stages of transcription, Bacterial RNA polymerase - structure and mechanism, recognition of promoters and DNA	
melting, abortive initiation. Elongation, Termination, antitermination.	
<b>Eukaryotic Transcription</b> : Eukaryotic RNA polymerases - RNA polymerase I, II, III.	
Mechanism of RNA polymerase in detail. Promoters, Transcription factors, basal apparatus, promoter clearance, elongation. Enhancers, silencers, termination.	
RNA splicing and Processing: mRNA capping, pre-mRNA splicing, lariat, snRNPs,	
spliceosome, autocatalytic splicing, alternative splicing, polyadenylation, tRNA splicing and maturation, production of rRNA, Catalytic RNAs - auto splicing,	
ribozymes, rinonuclease, viroids and virusoids, RNA editing	

**Translation:** Genetic code, tRNA structure, charging of tRNA, differences between initiator tRNA and elongator tRNA, ribosome structure. Accuracy of translation. Stages of translation. Role of IFs in initiation of bacterial translation, Formation of initiation complex. Initiation of eukaryotic translation - Scanning model of mRNA, IRES, Role of eIFs. Elongation of polypeptide - EF-Tu, EF-G, peptide bond formation, peptidyl transferase activity, translocation, eEFs. Termination.

## UNIT IV: Protein folding and control of regulation of gene expression

14hrs

**Post translational modifications of proteins:**Protein maturation and secretion - protein splicing, molecular chaperones. Protein translocation and secretion in bacteriaRegulation of translation.

**Regulatory mechanisms in bacteria:**Positive and negative transcriptional control in bacteria. Operon concept, polycistronic mRNA. *lac* operon - negative inducible, allolactose, mutants of *lac* operon structure of *lac* repressor, mechanism of binding of repressor to operator. Catabolite repression of *lac* operon. Regulation by lac repressor and CAP. *trp* operon regulation - repressor control& attenuator control.

**Regulation through modification of gene structure:** DNase I hypersensitivity, histone modifications, chromatin remodelling, DNA methylation. Regulation through transcriptional activators, Co-activators and repressors, enhancers and insulators. Regulation through RNA processing and degradation. Regulation through RNA interference

## Pedagogy: Lectures, Seminars, Industry Visits, Debates, Quiz and Assignments

FormativeAssessmentforTheory						
AssessmentOccasion/type	Marks					
InternalAssessmentTest1	10					
InternalAssessmentTest2	10					
Quiz/Assignment/SmallProject	10					
Seminar	10					
Total	40Marks					
FormativeAssessmentasperguid	lelines.					

## **B.Sc. Semester–V**

**DisciplineSpecific Core Course (DSCC)-10** 

# Course Title: Microbial genetics and molecular biology Course Code: 035MCB012

Type	Theory		Instructionhou	TotalNo.ofLectu	Durationof	FormativeA	Summative	TotalMa
ofCourse	/Practical	Credits	rperweek	res/Hours	Exam	ssessmentM	assessment	rks
				/Semester		arks	Marks	
DSCC-10	Practical	02	04	56hrs.	3hrs.	25	25	50

#### **PracticalContent**

- 1. Good Laboratory Practices and Safety Measures of Biohazard materials.
- 2. Standard operating procedure for molecular biology tools/equipment's.
- 3. Extraction of crude DNA from bacteria and yeast by phenol/chloroform method.
- 4. Demonstration of estimation of DNA by spectrophotometric method.
- 5. Extraction and visualization of genomic DNA from bacterial cultures
- 6. Determination of DNA melting point and GC content
- 7. Extraction and visualization of plasmids from bacterial cultures
- 8. Characterization of DNA by SpectrophotometricAssay and Melting Temperature (Tm).
- 9. Study of semi-conservative replication of DNA through micrographs / schematic representations
- 10. DNA fingerprinting technique through micrographs / schematic representations
- 11. Determination of linkage and cross-over analysis
- 12. Experiments on probability and Chi-square test.
- 13. Demonstration of monohybrid and Dihybrid cross by Punnett squares.

## PracticalAssessment

FormativeAssessment		SummativeAssessment	TotalMarks
AssessmentOccasion/type Weightage		PracticalExams	
	inMarks		
Scheme of Practical Examination (marks): 25 marks for Semester end Major Question Minor QuestionIdentify and comment Viva03Marks	examination 10 Marks -06 Marks	25	50
Practical Records	03Marks		
Total	25	25	

	same shall be used for semester the Examination
Refe	rences
1	Karp'sCellandMolecularBiologybyGeraldKarp,JanetIwasa,Wallace Marshall.NinthEdition. 2020
2	Lewin's Genes XII. Jocelyn EKrebs, Elliott SGoldstein, Stephen TKilpatrick. Jones and Bartlett Learning. 2017
3	James D. Watson, Tania A. Baker, Stephen P. Bell, Alexander Gann, Michael Levine, Richard Losick. Molecular Biology of the Gene, 7th edition. 2017
4	Freifelder's Essentials of MOLECULAR BIOLOGY. George MMalacinski, 4 <sup>th</sup> ed. 2015
5	FreifelderD (2012).MolecularBiology,5th edition.NarosaPublishingHouse,India
6	BergJM, TymoczkoJL, GattoGJ and StryerL
	(2015)Biochemistry,8thEdition,WHFreeman&Co.,New York
7	AlbertsBruce,JohnsonA, LewisJ, RaffM, RobertsK, WalterP (2014) MolecularBiology of
	theCell.5th Edition,Taylorand Francis.New York,USA.
8	TroppBE(2012)MolecularBiology: GenestoProteins.4rdEdition, Jones&Bartlett,Learning,
	Burlington,MA
9	AllisonA.Elizabeth (2012)FundamentalMolecularBiology,2nd Edition.JWilley andSons,
	Hoboken, New Jersey

10	ArandaPS,LaJoieDM,Jorcyk CL(2012).BleachGel: ASimpleAgaroseGelforAnalyzingRNA
	Quality.Electrophoresis.33(2):366–369.Doi:10.1002/elps.201100335.
11	BlochKD;GrossmannB(1995).DigestionofDNAwithRestrictionEndonucleases.
	https://doi.org/10.1002/0471142727.mb0301s31
12	ChomczynskiP,Sacchi N(2006). "Thesingle-stepmethodofRNAisolationby acidguanidinium
	thiocyanate-phenol-chloroform extraction: twenty-something years on". Nat Protoc. 1 (2): 581–
	5.doi:10.1038/nprot.2006.83.
13	ElkinsKM(2013). DNAExtractionForensic DNABiology.
14	FrederickM.Ausubel,RogerBrent,RobertE.Kingston,DavidD.Moore,
	J.G.Seidman, John A. Smith, Kevin Struhl (2003). Current Protocols in Molecular Biology. John Wiley
	&Sons, New
	York, United States.
15	JohnsonM(2019).RNA extraction,SynatomResearch,Princeton,NewJersey,UnitedStates.
	DOI//dx.doi.org/10.13070/mm.en.2.201.
16	LewisM.Agarosegelelectrophoresis (basicmethod).Department ofPathology, Universityof
	Liverpool. <a href="http://diyhpl.us/~bryan/irc/protocol-online/protocolcache/agarogel.html">http://diyhpl.us/~bryan/irc/protocol-online/protocolcache/agarogel.html</a>
17	RandallDR.(2009).MolecularBiologyLaboratorymanual.
18	SambrookJF,Russell DW(2001).MolecularCloning:a LaboratoryManual.3rd edition.Cold
	SpringHarbor,N.Y.ColdSpringHarborLaboratoryPress
19	Struhl K, Seidman J G, Moore D D, Kingston RE, Brent R, Ausubel FM, Smith JA. (2002).
	hortProtocolsinMolecularBiology:ACompendiumofMethodsfromCurrentProtocolsinMolecular
	Biology.JohnWiley&SonsInc.,New York,UnitedStates
20	SurzyckiS (2000).Basic techniquesinmolecularbiology.Springer.
21	Yılmaz M, Ozic C, Gok İ (2012). Principles of Nucleic Acid Separation by Agarose
	GelElectrophoresis. Gel Electrophoresis - Principles and Basics, Dr. Magdeldin S (Ed.), ISBN:
	978-953-51-0458-2, InTech.http://www.intechopen.com/books/gel-electrophoresis-principles-
	Andbasics

## DisciplineSpecific Core CourseDSCC-11

Course Title: Food and dairy microbiology (Theory)

Course Code: 035MCB013

Type	Theory		Instructionhou	TotalNo.ofLectu	Durationof	FormativeA	Summative	TotalMa
ofCourse	/Practical	Credits	rperweek	res/Hours	Exam	ssessmentM	assessment	rks
				/Semester		arks	Marks	
DSCC-11	Theory	04	04	56hrs.	2hrs.	40	60	100

## **Course Pre-requisite(s):**

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

- CO1. To understand the association of microbes in food and dairy, quality testing of food and dairy products
- CO2. To understand the preservation and food safety protocols
- CO3. To understand the methods of spoilage of food and the diseases associated with it
- CO4. To learn the properties of milk and the types of preservation of milk.
- CO5. To learn the human microbiota and its significance in Diet.

CONTENTS	56 Hrs
Unit I-Microbes and Food	14
Food as a substrate for microorganisms: Intrinsic and extrinsic parameters affecting the	hrs
growth of microbes. Microorganisms in food and their sources (molds, yeast and bacteria).	
Food borne infections and Intoxication: Causative agents, foods involved, symptoms and	
preventive measures for Salmonella, Shigella, Yersinia enterocolitica, Staphylococcus,	
Clostridium. Salmonella, Bacillus cereus, Brucella, Listeria monocytogens, Mycotoxin,	
Phycotoxins.	
<b>Fermented Food:</b> Fermented vegetable-sauerkraut, pickles. Meat- sausage. Beverages	
kombucha. Sourdough. Microbes as food- SCP, SCO. Neutrceuticals and Synbiotics.	
Unit II-Spoilage of Food, Preservation and Food safety	14hr
<b>Spoilage of Food :</b> Principles of food spoilage. Sources of food contamination, Types of spoilage. Spoilage of meat and poultry, Fish and sea foods. Spoilage cereals, fruits and vegetables.	
Spoilage of canned food.	
<b>Food Preservation</b> : Principles of food Preservation. Methods of preservation Physical (temperature, drying, irradiation), chemical (Class I and Class II). Bio preservation. Canning, Food Packaging-Types of packaging materials, properties and benefits.	
Quality control in Food-Food Sampling, preparation and handling, Surface and environmental monitoring in food industry, basic physical and chemical analysis of food,	
Microbiological analysis of food and food products, Rapid microbiological and molecular	
methods to detect food pathogens.  Food Sopitation and Sofaty: Good Hygiana practices: GLP GMP (Weste treatment)	
<b>Food Sanitation and Safety:</b> Good Hygiene practices, GLP, GMP (Waste treatment disposal methods), Food Safety HACCP, FSSAI and Food safety and Standard act 2006,	
Food control agencies and their regulation.	

### **Unit III: Dairy Microbiology**

**History:** Introduction and Significance of Dairy microbiology, Hygienic milk production, Dairy associated microorganisms. Properties of milk. Types of milk- dried, liquid, condensed, Microorganisms associated with milk (beneficial and harmful)

**Microbiology of milk:** Sources of contamination and spoilage of milk. Microbiological methods for milk testing in Diary Industry: Rapid platform tests (Organoleptic, COB, alcohol test, Phosphatase test, DMC, sedimentation test.). Reductase tests. SPC. Biochemical changes of milk-souring, gassy fermentation, proteolysis, lipolysis, ropiness. Effect of processing on microorganisms in milk.

**Preservation of milk:** Pasteurization. Dehydration, sterilization, Packing of milk and dairy products. Effect of processing on microorganisms in milk.

**Dairy products:** Therapeutic value of Yoghurt and Butter milk, Cheese (Types and production), Tofu, Yoghurt, Acidophilus milk. Prebiotics, Probiotics. Starter culture types and role (single, mixed). Antimicrobial substances in milk.

### **Unit IV: Human Microbiome and Diet**

Human Microbiome: Definition, origin, formation, development, structure and functions of human microbiome and its evolution. Factors affecting microbial diversity and functions of microbiome: -age, genetics, environment, diet, anatomy, physiology, immunity, and psychology of host(human). Dynamics microbiome changes from birth to death; pregnancy and the microbiome; personnel microbiome concepts. Geography, Ethnicity -Specific Variations in Human microbiome. "diseases of civilization" -allergies, diabetes, asthma, obesity, inflammatory bowel disease. Debate on "nature" vs. "nurture". Biodiversity and major genera of human microbiome, human microbiome system as a "holobiont" or "superorganism" microbiome distributions in healthy Individuals- hands, neck, scalp, axilla, groin, toes, ear; anterior nares, oral cavity, throat, stomach, small intestine and large intestine and birth canal.

**Human Diet and Microbiota**: Microbiome vs microbiota; microbiota development and functions in early life; Microbiota transmission-pregnancy, birth and postnatal. Microbiota perturbations: medical practices, hygiene and antibacterials. Nutritional modulation of the gut microbiome for metabolic health- animal models, human obesity, human type 2 diabetes, life longevity.

## Pedagogy: Lectures, Seminars, Industry Visits, Debates, Quiz and Assignments

Assessment Occasion/type	Marks	
Internal Assessment Test 1	10	
Internal Assessment Test 2	10	
Quiz/Assignment/Small Project	10	
Seminar	10	
Total	40 Marks	

15 hrs

## **Discipline Specific Core Course**DSCC-12

Course Title: Food and dairy microbiology

Course Code: 035MCB014

Type of	Theory		Instruction	Total No. of	Duration	Formative	Summative	Total
Course	/Practical	Credits	hour per week	Lectures/Hours	of Exam	Assessment	assessment	Marks
				/Semester		Marks	Marks	
DSCC-12	Practical	02	04	56hrs.	3hrs.	25	25	50

### **ContentsofPractical**

- 1. Standard procedures for food sampling, preparation and handling in food industry.
- 2. Standard procedures surface and environmental monitoring in food industry.
- 3. Isolation and enumeration of Aerobic Plate count and yeast and moulds from infected fruits and vegetables, ready to eat and cooked foods and fermented foods.
- 4. Enumeration of *E.coli, S.aureus*, *Salmonella, Shigella* and *Bacillus cereus* form ready to eat/cooked food using selective culture medias.
- 5. Reductase tests-MBRT, Resazurin and Litmus milk test.
- 6. Estimation of Titrable acidity in milk.
- 7. Estimation of Fat Gerber's method
- 8. Bacterial examination of milk by SPC, DMC
- 9. Estimation of lactose in milk
- 10. Production of fermented foods
- 11. Detection of Mastitic milk.
- 12. Visit to Food and Milk processing, alcoholic beverage Industries and report should be written and submitted along with the practical record.

FormativeAssessment		SummativeAssessment	TotalMarks		
AssessmentOccasion/type	Weightage	PracticalExams			
	inMarks				
Scheme of Practical Examination (	listribution of				
marks): 25 marks for Semester end	examination				
Major Question		25	50		
Minor Question	06 Marks	23	50		
Identify and comment	-3x1 = 03 Marks				
Viva03Marks					
Practical Records	03Marks				
Total	25	25			

Re	eferences
1	Adams, M.Rand Moss, MO. 1995. Food Microbiology. The Royal Society
	ofChemistry,Cambridge.
2	James.M.Jay, 1992,Modernfood microbiology4ed.
3	Frazier W.C. and Westhoff C.D. 2008 Food Microbiology. Tata McGraw Hill Publishing
	CompanyLimited,New Delhi,India.
4	DoyleM.P.andBeuchatL.R.(2007).FoodMicrobiology-Fundamentals.Frontiers,ASMPress.
5	Garbutt J. (1997). Essentials of Food Microbiology, Armold- International Students
	edition,London.8. Marriott N. G. and Gravani R. B. (2006).
6	PrinciplesofFoodSanitation,FoodSciencetextSeries,Springer International,NewYork,USA.
7	ThomasJ., Matthews, Karl; Kniel, Kalmia E(2017), Food Microbiology: An Introduction, American Soci
	etyfor (ASM).
8	DeakT.andBeuchatL.R.(1996).HandBookof FoodSpoilageYeasts,CRCPress,NewYork.

## **Skill Enhancement Course: SEC-3**

Course Title: Microbial quality control in food and pharma industries

Course Type: SEC-3
Course Code:035MCB061

	Type ofCourse	Theory /Practical		Instructionhou r/week	TotalNo.ofLectu res/Hours		FormativeA ssessmentM		
	orcourse	/I ractical	Credits	1/ WCCK	/Semester	Lam	arks	Marks	11.5
•	SEC-3	Practical	02	04	56hrs.	3hrs.	25	25	50

### CourseOutcomes(COs): Attheendofthecoursethestudentshouldbeableto:

- 1. Demonstrate skills as per National Occupational Standards (NOS) of the "LabTechnician/Assistant" QualificationPack issued by the Life Sciences SectorSkillDevelopmentCouncil-LFS/Q0509.
- 2. Develop knowledge of laboratory safety procedures and protocols and acquireskillsinhandlingandmaintaininglaboratoryequipmentandinstruments.
- $3. \quad Operate analytical equipment and instruments as per standard operating procedures (SOP)\\$
- 4. Knowledgeaboutmajoractivitiesofthebiotechindustry,regulationsandcompliance, environment, health and safety (EHS), good laboratory practices(GLP),andGoodManufacturingPractices(GMP)aspertheindustrystandar ds.
- 5. Demonstrate soft skills, such as decision-making, planning, organizing, problem-solving, analyticalthinking, critical thinking, and documentation.

#### **Contents**

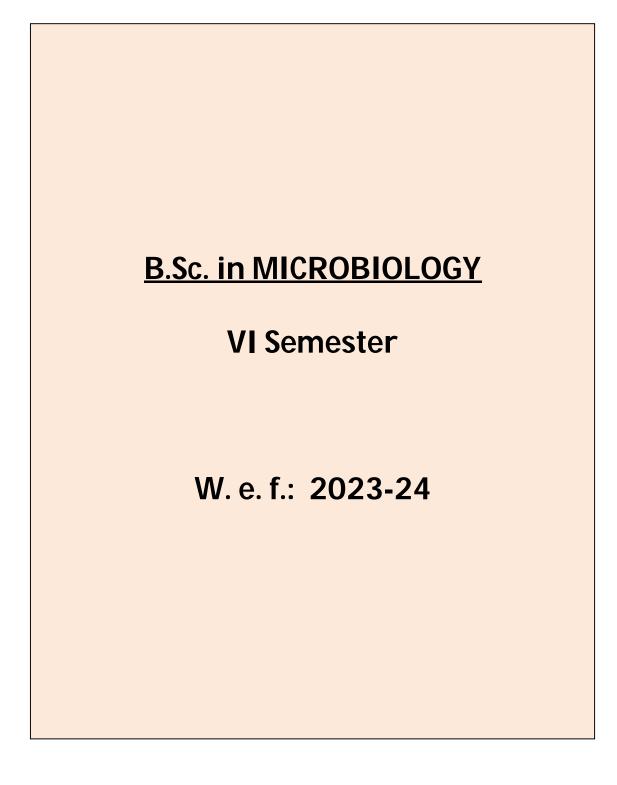
- 1. SOP for Swab and food sampling, handling and homogenization in food industry.
- 2. Guidelines and procedure for GLP, GMP and GDP in quality control food and pharma industries.
- 3. Procedure for cleaning, disposal, decontamination, sanitation, fumigation and sterility in Microbiology laboratory.
- 4. Monitoring and validation of autoclave process by chemical and biological indicators in quality control microbiology.
- 5. Media preparation and its importance of media in pharmaceutical and food industries.
- 6. Pure cultures maintenance techniques in quality control microbiology.
- 7. Growth Promotion Test (GPT) to verify the fertility of culture media.
- 8. Physical and chemical analysis of raw water used in food and pharma industries.
- 9. Enumeration of Total Viable Count (TVC), Total Yeast and Mould Count (TYMC) and specified pathogens by membrane filtration techniques in pharma industries.
- 10. Enumeration of Total Coliforms and E.coli form drinking, raw and DM water by membrane filtration techniques.
- 11. Environmental monitoring of Surface and personal hygiene swabs in industries.
- 12. Perform BET and MLT sterility tests.
- 13. Visit to Pharma, Food and food processing, alcoholic beverage industries. Tour/Project Report should be submitted.

PracticalAssessment					
FormativeAssessment		SummativeAssessment	TotalMarks		
AssessmentOccasion/type	Weightage	PracticalExams			
	inMarks				
Scheme of Practical Examination (	listribution of				
marks): 25 marks for Semester end	examination				
Major Question	10 Marks	35	50		
Minor Question	06 Marks	25	50		
Identify and comment	-3x1 = 03 Marks				
Viva03Marks					
Practical Records	03Marks				
Total	25	25			

The same shall be used for semester end Examination

### References

- 1. Baird, R. M., Hodges, N. A. and Denyer, S. P. (2005). Handbook of Microbiological Quality control in Pharmaceutical and Medical Devices, Taylor and Francis Inc.
- 2. Garg, N., Garg, K. L. and Mukerji, K. G. (2010). Laboratory Manual of Food Microbiology I K International Publishing House Pvt. Ltd.
- Harrigan, W. F. (1998). Laboratory Methods in Food Microbiology, 3<sup>rd</sup> ed. Academic Press.
   Jay, J. M., Loessner, M. J., Golden, D. A. (2005). Modern Food Microbiology, 7<sup>th</sup> edition. Springer.



## DisciplineSpecific Core Course(DSCC)-13

# Course Title:Immunology and medical microbiology Course Code:036MCB011

-	DSCC-13	Theory	04	04	56hrs.	2hrs.	40	60	100
					/Semester		arks	Marks	
	ofCourse	/Practical	Credits	rperweek	res/Hours	Exam	ssessmentM	assessment	rks
	Type	Theory		Instructionhou	TotalNo.ofLectu	Durationof	FormativeA	Summative	TotalMa

## CourseObjectives

- 1. To understand the various aspects of immunity, elicitation of immuneresponses, factors determining the outcome of immune responses andmajorplayersofimmunity, relevance betweennutritional supportand immunity, and immunological techniques.
- 2. To provide knowledge on essential features of antigens and antibodies and their types and different theories of Antibody formation.
- 3. Toacquireknowledgeontypesofimmunity,phagocytosis,interferons,and thecomplementsystem.
- 4. To explain the concept of hypersensitivity, autoimmunity, and transplantation.
- 5. Toprovideknowledgeonimmunedeficienciesandseveralimmunologicaltechniques

## **CourseOutcomes**

#### Attheendofthecourse, the students hould be able to:

- 1. Demonstratecomprehensionoftheunderlyingstructureandfunct ionoftheimmunesystemandrelateddisorders.
- 2. Demonstrate an understanding of the role of cells and molecules in immuner eactions and responses
- 3. Demonstratetechnicalskillsinimmunologicaltoolsandtechniques
- 4. Applythedomainspecificknowledgeandskillsacquiredinimmunologyforinnovati vetherapiesandImmunotechnologies
- 5. Understandthefundamentalconceptsofimmunity, and the contributions of the organisand cells in immuner esponses.
- 6. RealizehowtheMHCmolecule's function and hosten counters an immune in sult.
- 7. Understandtheantibodiesandcomplementsystem
- 8. Understandthemechanismsinvolvedintheinitiationofspecificimmuneresponses
- 9. Differentiatethehumoralandcell-mediatedimmunemechanisms
- 10. Comprehendtheoverreactionbyourimmunesystemleadingtohypersensitivecon ditions and its consequences
- 11. Understandunique properties of cancercells, immune recognition of tum ors, immune evasion of cancers

56Hrs
14hrs
14Hrs
14Hrs

UNIT-IV 14Hrs

#### Antigen:

Immunogenicity and antigenicity, epitopes, haptens. Properties of antigen contribute to immunogenicity; Chemical nature (proteins, carbohydrates, lipids and nucleic acids), degree of foreignness, molecular weight, chemical composition and complexity, degradability. Adjuvants (alum, freunds incomplete and complete) and their importance. B and T cell epitopes.

## **Antibody**:

Basic structure of antibody, light and heavy chain, variable and constant region, hinge region, Fab and Fc. Structure and functions of different types of antibodies (IgM, IgG, IgA, IgE, and IgD). Antigenic determinants on immunoglobulins:

Isotype, allotype and idiotype. Monoclonal antibody production by hybridoma technology

## **Principles and applications of antigen-antibody interactions:**

Definition of affinity and avidity. Immunoprecipitation; Radial (Mancini) and double (Ouchterlony) immunodiffusion.

**Agglutination reactions:** Hemagglutination, Bacterial agglutination, passive agglutination, and agglutination inhibition. Enzyme linked immune-sorbent assay (ELISA): Direct, indirect, sandwich and competitive ELISA. Radioimmunoassay (RIA). Immunofluorescence. Complement mediated opsonization, complement fixation test.

## **Hypersensitive reactions:**

Classification, Humoral Immunity mediated hypersensitivity; Type I (IgE), Type II (IgG and IgM-ADCC), Type III (Antigen-antibody complex), and Cell mediated hypersensitivity TypeIV (DTH).

Pedagogy: Lectures, Seminars, Industry Visits, Debates, Quiz and Assignments

Formative Assessment for Theory					
Assessment Occasion/type	Marks				
InternalAssessmentTest1	10				
InternalAssessmentTest2	10				
Quiz/Assignment/Small Project	10				
Seminar	10				
Total	40 Marks				
Formative Assessment as per guidelines.					

## Discipline Specific Core CourseDSCC-14

Course Title: Immunology and medical microbiology

Course Code: 036MCB012

Ty	pe of	Theory		Instruction	Total No. of	Duration	Formative	Summative	Total
C	ourse	/Practical	Credits	hour per week	Lectures/Hours	of Exam	Assessment	assessment	Marks
					/Semester		Marks	Marks	
DS	CC-12	Practical	02	04	56hrs.	3hrs.	25	25	50
ContentofPractical									
1	Identify pathogenic bacteria (any three of <i>E. coli, Salmonella, Pseudomonas, Staphylococcus</i> and <i>Bacillus</i> ) on the basis of cultural, morphological and biochemical characteristics: IMViC, TSI, H <sub>2</sub> S production, Nirate reduction, Urease production and catalase tests.								
2	Study of composition and use of important differential media for identification of pathogenic bacteria: EMB Agar, MacConkey agar, Mannitol salt agar, Deoxycholate citrate agar, TCBS, Cetrimide agar.								
3	Study	of bacterial	flora of sl	in by swab met	thod				
4	Perfori	m antibacte	rial sensiti	vity by Kirby-E	Bauer method (Dis	sk/well and	MIC)		
5	•	• 1			help of photograps), dermatomycos			oes,	
6	Study	of various s	tages of M	Ialarial parasite	in RBCs using pe	ermanent m	ounts.		
7	Identif	ication of h	uman bloc	od group and ca	lculation of allelic	c frequency			
9	Perfori	m Different	ial Leukoc	cyte Count of th	e given blood san	nple.			
10	Separa	tion of seru	m from th	e blood sample	(demonstration).				
11	Perfori	m immunod	liffusion b	y Ouchterlony 1	method.				
12	Perfori	m DOT ELI	SA.						
13	Perfori	m immune e	electropho	resis.					

PracticalAssessment							
FormativeAssessment		SummativeAssessment	TotalMarks				
AssessmentOccasion/type Weightage		PracticalExams					
	inMarks						
Scheme of Practical Examination (							
marks): 25 marks for Semester end							
Major Question		25	50				
Minor Question	-06 Marks	23	50				
Identify and comment	-3x1 = 03 Marks						
Viva03Marks							
Practical Records	03Marks						
Total	25	25					

RE	FERENCES
1	Ananthanarayan R and Paniker C.K.J (2009) Textbook of Microbiology, 8 <sup>th</sup> 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. 26th edition. McGraw Hill Publication
3	Goering R., Dockrell H., Zuckerman M. and Wakelin D. (2007) Mims' Medical Microbiology. 4th edition. Elsevier
4	Willey JM, Sherwood LM, and Woolverton CJ. (2013) Prescott, Harley and Klein's Microbiology. 9th edition. McGraw Hill Higher Education
5	Madigan MT, Martinko JM, Dunlap PV and Clark DP. (2014). Brock Biology of Microorganisms. 14th edition. Pearson International Edition
6	Abbas AK, Lichtman AH, Pillai S. (2007). Cellular and Molecular Immunology. 6th edition Saunders Publication, Philadelphia.
7	Delves P, Martin S, Burton D, Roitt IM. (2006). Roitt's Essential Immunology.11th edition Wiley- Blackwell Scientific Publication, Oxford.
8	Goldsby RA, Kindt TJ, Osborne BA. (2007). Kuby's Immunology. 6th edition W.H. Freeman and Company, New York.
9	Murphy K, Travers.P,Walport M. (2008).Janeway's Immunobiology.7 <sup>th</sup> edition Garland Science,Publishers,New York.
10	Peakman.M.and Vergani D. (2009).Basic and Clinical Immunology,2nd edition Churchill ,Livingstone Publishers, Edinberg.
11	Richard C and Geiffrey S. (2009). Immunology. 6th edition. Wiley Blackwell Publication.

## Discipline Specific Core Course(DSCC)-15

Course Title: Genetic engineering and industrial microbiology Course Code:036MCB013

Type of	Theory		Instruction	Total No. of	Duration	Formative	Summative	Total
Course	/Practical	Credits	hour per week	Lectures/Hours	of Exam	Assessment	assessment	Marks
				/Semester		Marks	Marks	
DSCC-15	Theory	04	04	56hrs.	2hrs.	40	60	100

## **CourseObjectives:**

- 1. Performsimulationsofmicrobialgrowthandmetabolism
- 2. Designbioreactorsfortheproductionofvarious products.
- 3. Presentknowledgeaboutmajormetabolicpathwaysandthoserelatedtobiofuelproducti onfrommicrobes.
- $4. \quad Understand the fundamental concepts and principles of environmental MICROBIOLOGY and Explore the interrelationship between MICROBIOLOGY and the environment.\\$
- 5. Gainknowledge of the various applications of MICROBIOLOGY inenvironmental conservation, pollution control, and sustainability.
- 6. Learnaboutmicrobialprocessesandtheirroleinenvironmental MICROBIOLOGY.
- 7. Understand the principles of bioremediation and its application in the clean-up of environmental pollutants.
- 8. Explorethepotentialofbioenergyproductionandwastemanagementthroughbiotech nologicalapproaches.
- 9. IdentifyandcharacterizethemostimportantcontaminantsintheBioprocessandotherindus trialwastes.
- 10. Reuse/recyclethebiologicalwastetocleantechnologysuchasenergy,biofuel,biofertili zerthrough bioremediation

### Courseoutcomes:

- 1. Exploitation of microorganisms for industrial use and their improvement, and formulation of media for efficient growth and production of microbial or cell-based products.
- 2. The design, operation, and specific applications of various bioreactors.
- 3. DemonstrateacomprehensiveunderstandingofthefundamentalconceptsandprinciplesofenvironmentalMICROBIOLOGY.
- 4. Applyknowledgeofbiotechnologicaltechniques to addressenvironmentalchallenges, such as pollution control and was temanage ment.
- 5. Analyzeandevaluateenvironmental MICROBIOLOGY cases tudies, research findings, and real-world applications.
- 6. Designandimplementbiotechnological approaches for environ mental remediation, utilizing microbial processes and biodegradat ion principles.
- 7. Evaluate the ethical and sustainable aspects of environmental MICROBIOLOGY practices and make informed decisions regarding their application inenvironmental conservation.
- 8. Communicate scientific concepts and research findings related

# toenvironmental MICROBIOLOGY effectively, both in written and oral forms, to diverse audiences.

and oralforms,todiverseaudiences.	
GENETIC ENGINEERING AND INDUSTRIAL MICROBIOLOGY Contents	
Unit I: Introduction and tools of Microbial Genetic Engineering	14Hrs
Historical prospective: Definition of genetic engineering, milestones in genetic engineering, prospects and problems of genetic engineering.  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.  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.  Cloning host- Cloning in Escherichia coli, cloning in Saccharomyces cerevisiae,  Unit II: Techniques and applications in Microbial Genetic Engineering	14 Hrs
	14 Hrs
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 hybridization, blue white selection, antibiotic selection, colony and plaque hybridization.  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.  Recombinant microorganisms: Application of recombinant microorganisms in basic research, industry, medicine, agriculture, environment.  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.	
Unit III: Introduction to Industrial Microbiology	14 Hrs
Introduction to Industrial microbiology: History, scope and development of industrial microbiology. Isolation and screening (Primary and Secondary) of industrially important microorganisms, Strain improvement methods. Preservation of industrially important microbes. Basic features; design and components of a bioreactor, Specialized bioreactors and their applications: tubular bio reactors, fluidized bed reactor, packed bed reactors, membrane bioreactors, Photo-bioreactors and anaerobic bioreactors.  Role of industrial microorganisms for ecovery of Minerals (Bioleaching) and Petroleum (Microbial Enhanced Oil Recovery-MOER). Role of microbes in production of biofuel by bacteria, algae and fungi.	
Unit IV: Fermentation Process and Scale up Fermentation Process:Types of fermentation process: Submerged fermentation, Solid state fermentation (Koji), batch fermentation, continuous fermentation, kinetics of fermentation process. Inoculum preparation. Media components and formulation (Crude media components, Anti-foam agents, Precursors, Inducers, Inhibitors and Buffering agents). Sterilization of media and raw materials and maintenance of Sterility at critical points during fermentation. Scale up of Fermentation: Upstream and Downstream processing, Objectives and significance of downstream processing: Overview of steps in extraction and purification ofproducts (Antibiotic, Enzyme, Hormones, Anti-cancerous compounds); Precipitation Filtration and centrifugation; cell disruption- Physical, chemical and biological methods; Product extraction; product purification, recovery drying, crystallization and product testing. Merits and demerits. Immobilization of cells and enzymes –Types, advantages and applicationsin fermentation industry.	

Pedagogy: Lectures, Seminars, Industry Visits, Debates, Quizand Assignments

Formative Assessment for Theory			
Assessment Occasion/type	Marks		
InternalAssessmentTest1	10		
InternalAssessmentTest2	10		
Quiz/Assignment/Small Project	10		
Seminar	10		
Total	40 Marks		
Formative Assessment as per g	uidelines.		

## **Discipline Specific Core Course**DSCC-16

Course Title: Genetic engineering and industrial microbiology

Course Code: 036MCB014

DSCC-16	Practical	02	04	56hrs.	3hrs.	25	25	50
				/Semester		Marks	Marks	
Course	/Practical	Credits	hour per week	Lectures/Hours	of Exam	Assessment	assessment	Marks
Type of	Theory		Instruction	Total No. of	Duration	Formative	Summative	Total

### **ContentofPractical**

- 1. Preparation of buffers-TE, TAE and Lysisbuffer.
- 2. Isolation of genomic DNA from *Escherichia coli*.
- 3. Preparation of master and replica plates.
- 4. Designing of primers for DNA amplification.
- 5. Demonstration of amplification of DNA by PCR(By chart).
- 6. Demonstration of southern, western and northern blotting techniques (By chart).
- 7. Preparation of wine from different fruits an Estimation of Alcohol by Specific gravity method.
- 8. Production and estimation of citric acid by Aspergillus brasilensis
- 9. Production of enzyme (amylase/protease/cellulose /invertase by submerged fermentation).
- 10. Production and estimation of any one secondary metabolite.
- 11. Immobilization of cells and enzymes by solid entrapment.
- 12. Preservation of microbes with glycerol/soil/oil/sand.
- 13. Visit to Molecular biology laboratories, Research institutes, Sugar Distillery, Alcoholic beverages industry and report should be written and submitted along with the practical record.

1.

PracticalAssessment						
FormativeAssessment		SummativeAssessment	TotalMarks			
AssessmentOccasion/type	Weightage	PracticalExams				
	inMarks					
Scheme of Practical Examination (	listribution of					
marks): 25 marks for Semester end	examination					
Major Question	10 Marks	25	50			
Minor Question	06 Marks	23	50			
Identify and comment	3x1 = 03 Marks					
Viva03Marks						
Practical Records	03Marks					
Total	25	25				

R	References					
1	Brown TA. (2010). Gene Cloning and DNA Analysis. 6th edition. Blackwell Publishing, Oxford, U. Clark DP and Pazdernik NJ. (2009). Biotechnology: Applying the Genetic Revolution. Elsevier Academic Press, USA	K.				

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.

## Microbiology InternshipforgraduateProgramme

Coursetitle	InternshipDisciplinespecific
Course code	036MCB091
Noofcontacthours	56
Nocredits	2
Methodofevaluation	Presentations/Reportsubmission/Both

**ProjectAssessment** 

Type ofCourse	Theory		Instructionhou	TotalNo.ofLectu	Durationof	FormativeA	Summative	TotalMa
	/Practical	Credits	r/week	res/Hours	Exam	ssessmentM	assessment	rks
				/Semester		arks	Marks	
Internship-1	Practical	02	04	56 hrs.	3hrs.	50	0	50

- Internship shall be Discipline Specific of 56 hours (2 credits) with duration 1-2 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.
- Incase Internship in a company or institute not possible or college did not permit then mini projects on Microbiology topics may be given. Viz., Wine production, Human microbiome etc,

UG programme: 2023-24

## GENERAL PATTERN OF THEORY QUESTION COURSE FOR DSCC/OEC

## (60 marks for semester end Examination with 2 hrs duration)

#### Part-A

1. Question number 1-06 carries 2 marks each. Answer any 05 questions : 10 marks

#### Part-B

**2.** Question number 07-11 carries 05Marks each. Answer any 04 questions : 20 marks

### Part-C

**3.** Question number 12-15 carries 10 Marks each. Answer any 03 questions : 30 marks (Minimum 1 question from each unit and 10 marks question may have sub questions for 7+3 or 6+4 or 5+5 if necessary)

### **Total: 60 Marks**

Note: Proportionate weight age shall be given to each unit based on number of hours

Prescribed