# Savitribai Phule Pune University Faculty of Science & Technology



Curriculum

For Second Year (S.E.) B.Tech. Biotechnology (Choice Based Credit System)

(2019 Course)

(With Effect from Academic Year 2020-21)

	(*****		Se	emes	ter-I	II								
Course Code	Course Name	Tea Sc (Hour	achin hemo rs/Wo	e eek)	E	xamin	ation Ma	Schei rks	ne a	ind	5	Cred	it	
		Theory	Practical	Tutorial	IN-Sem	End-Sem	ΜL	PR	OR	Total	ΗT	PR	TUT	Total
215461	Biochemistry I	3		-	30	70		-	-	100	3		-	3
207004	Engineering Mathematics III	3	-	01	30	70	25	-	-	125	3	-	1	4
215462	Fluid Flow & Unit Operations	3	-	-	30	70	-	-	-	100	3	-	-	3
215463	Heat transfer	3	-	-	30	70	-	-	-	100	3	-	-	3
215464	Microbiology	3	-		30	70	-	-	-	100	3	-	-	3
215465	Biochemistry I Lab	-	4	-	Ē	-	50	-		50	-	2	-	2
215466	Fluid Flow & Unit Operations Lab		2	-	-	-	25	-	-	25	-	1	-	1
215467	Heat transfer Lab	C	2	-	-	-	-	-	50	50	-	1	-	1
215468	Microbiology Lab		4	-	-	-	-	50	-	50	-	2	-	2
215469	Mandatory Audit Course 3	-	-	-	-	-	-	-	-	-	-	-	-	-
	Total	15	12	01	150	350	100	50	50	700	15	06	1	22

#### Abbreviations:

TH: Theory OR : Oral

TW : Term Work TUT : Tutorial

Note: Interested students of S.E. (Biotechnology) can opt any one of the audit course from the list of audit courses prescribed by BoS (Biotechnology Engineering)

PR : Practical

	Savit (Wi	ribai P SE(Bi th effect	hule io-Te from Se	Pur ech) Aca emest	ne U 201 demi er-IV	nive 9 Co <u>c Yea</u>	rsity ourse or 202	, Pur 0-21)	ne					
Course Code	Course Name	Teachin (Houn	ng Sch rs/Weo	eme ek)	ŀ	Exami	nation Ma	ı Sche arks	me a	nd		Cre	dit	
		Theory	Practical	Tutorial	IN-Sem	End-Sem	TW	PR	OR	Total	HL •	PR	TUT	Total
215470	Biochemistry II	3	-	1	30	70	-			100	3	-	1	4
215471	Cell Biology & Tissue Culture	3	-	1	30	70	-	- (	7	100	3	-	1	4
215472	Thermodynamics	3	-	1	30	70	25	-	-	125	3	-	1	4
215473	Genetics &Molecular Biology	3	-	1	30	70	2	X	-	100	3		1	4
215474	Biochemistry II Lab	-	2	-			25	50	-	75		1	-	1
215475	Cell Biology & Tissue Culture Lab	-	2	-		-	25	-	50	75		1	-	1
215476	Genetics & Molecular Biology Lab	-	4			<u> </u>	25	-	50	75		2	-	2
215477	Project Based Leaning	-	4	-	-	-	50	-		50	-	2	-	2
215478	Mandatory Audit Course 4	_		D	-	-	-	-	_	-	-	-	-	-
	Total	12	12	04	120	280	150	50	100	700	12	06	04	22
Abbreviatio	ns: TH : Theory TW	: Term V	Vork	PR	: Pra	ctical	1	1	1			1		

OR : Oral TUT : Tutorial

Note: Interested students of S.E. (Biotechnology) can opt any one of the audit course from the list of audit courses prescribed by BoS (Biotechnology Engineering)

Semest	ter I
Savitribai Phule Pr	une University, Pune
Second Year of B.Tech. B	Siotechnology (2019 Course)
215461	:Biochemestry I
Сг	redit
TH: 03	
Teaching Scheme:	ExaminationScheme:
TH: 03hrs/week	TH In Sem: 30
	TH End sem :70
	Total :100
Prerequisites: - Basic knowledge of Biology and	l chemistry
Course Objectives:	<b>N'0'</b>
1. To make students acquainted with the fur	nctioning of the bufferingsystem
2. To understand the working of basicbiomo	lecules
3. To recognize the clinical manifestations o	f vitamins and mineraldeficiency
Course Outcomes:	
On completion of the course, the learner will be a	ble to-
A. Understand the functioning of various but	ffering system existed in the humanbody
B.Recognize the structure and function of va	ariousbiomolecules
C.Correlate various diseases associated with	vitamin and mineraldeficiency
Course Contents	
UnitI	(07Hrs)
Water and buffer: Weak interactions in aqueous s	systems, Ionization of water, weak acid weak bases,
Ion product of water, acids and bases, buffers, b	uffering against pH changes in biological systems,
the fitness of the aqueous environment for 1	iving organisms, Problems using theHenderson-
Hasselbalchequation, Blood, Lungs, and buffer: 7	The bicarbonate buffer system, Water as a reactant
UnitII	(08Hrs)
Carbohydrate:Monosaccharidesanddisaccharides,	Polysaccharides, Homopolysaccharides in the
role of fuel and structural, Heteropolysaccharides	, Glycoconjugates- Proteoglycans, Glycoproteins,
and glycolipids Carbohydrate as informational m	polecules- Lectin

#### (07Hrs)

Proteins:CommonStructuralfeaturesofaminoacids,classificationofaminoacids(onthebasisofR groups, functions of uncommon amino acids, acid base properties of amino acids and titration curve.Peptidesandproteins,Separation,Purificationandcharacterizationofproteinsby

electrophoresis. Covalent structure of proteins, The Lambert-Beer law, Ramachandran plot.

#### UnitIV

UnitIII

(08Hrs)

Nucleotides and Nucleic acids: Characteristic bases and pentoses, phosphodiester bond, properties of nucleotidebases and three-dimensional structure of nucleic acids, Nucleic acid structure, Nucleic acid chemistry, Determination of sequences of DNA, Functions of nucleotides.

#### UnitV

#### (08Hrs)

Lipids:Storagelipids,Fattyacidsashydrocarbonderivatives,Triacylglycerol,Waxesasenergystores and water repellents, Structural lipids in membranes, Glycerophospholipids, Galactolipids, Sulfolipids, Sphingolipids, Composition and architecture of membranes. Inherited human diseases resultingfromabnormalaccumulationsofmembranelipids,Lipidsassignals,cofactorsandpigments,

Lipid extraction, Adsorption chromatography and Gas-liquid chromatography in the separation of lipid,

Composition and architecture of membranes.

#### UnitVI

#### (07Hrs)

Vitamins and Minerals: classification and functions of vitamins, (vit B1, B2, B6, B12, vit C), fat soluble vitamins (vit A, D, E, K), Vitamins deficiencies (night blind ness, keratomalacia, rickets, osteomalacia prolonged clotting time etc.) clinical manifestations of mineral deficiency (termatitis, dementia, diarrhea, pernicious anemia, survey etc)

#### Books Text:

1. DJVoet,JGVoet,CWPratt,"PrinciplesofBiochemistry",3<sup>rd</sup>ed.,JohnWiley&Sons,Inc.2008

2. D T. Plummer, "An Introduction to practical biochemistry", Tata McGraw Publishing Company Ltd, 1988

#### Reference:

1. J H Weil, "General Biochemistry", New Ages International (P) Ltd.1997.

2. JMBerg, JLTymoczko, LStryer, "Biochemistry", 6thed., FreemanWH&Company, NewYork, 2007

3. DLNelson,MMCox"PrinciplesofBiochemistry",4<sup>th</sup>ed.,W.H.Freemanandcompany,New

York, 2007

### Savitribai Phule Pune University, Pune Second Year of B.Tech. Biotechnology (2019 Course) 207004:Engineering Mathematics III Credit TH: 03 **TUT:01 Teaching Scheme: ExaminationScheme:** TH: 03 hrs/week TH TUT:01hr/week InSem:30TH End Sem :70 **TW :25 Total :125 Prerequisites: -**Differential & amp; Integral calculus, Linear Differential equations of first order and first degree, Collection, classification & amp; representation of data, Permutations & amp; combinationsFourier series and Vector algebra. Course Objectives: To make the students familiarize with concepts and techniques in Ordinary and Partialdifferential equations, Fourier transform, Laplace transform and Vector calculus. The aim is toequip them with the techniques to understand advanced level mathematics and its applications that would enhance analytical thinking power, useful in their disciplines. **Course Outcomes:** At the end of this course, students will be able to A. Solve higher order linear differential equations and its applications to engineering problems in their disciplines. B. Apply Integral transform techniques such as Fourier transform & amp; Laplace transform to solve differential equations involved in Vibration theory, Heat transfer, Liquid level systems and related engineering applications. C. Apply Statistical methods like correlation & amp; regression and probability theory as applicable to analyzing and interpreting experimental data in testing and quality control. D. Perform vector differentiation & amp; integration, analyze the vector fields and apply to fluid flow problems. E. Solve Partial differential equations such as wave equation, one and two dimensional heat flow equations.

#### **Course Contents**

Unit I: Linear Differential Equations (LDE) and Applications (08 Hours)

LDE of n th order with constant coefficients, Complementary Function, Particular Integral,

Method of Variation of parameters, Cauchy's and Legendre's DE, Simultaneous and

Symmetricsimultaneous DE. Applications of LDE to engineering problems and Mass spring system.

Unit II: Laplace Transform (LT) and Applications (08 Hours)

Definition of LT, Inverse LT, Properties & amp; theorems, LT of standard functions, LT of some

special functions viz. Periodic, Unit Step, Unit Impulse, Error, Si(t) and Ei(t), first order

Bessel's.

Applications of LT for solving ordinary differential equations, liquid level systems consisting of

single tank and two tanks in series (interacting and non-interacting systems), Second order

systems (Damped vibrator).

Unit III: Fourier Transform (FT) (07 Hours)

Fourier integral theorem. Fourier Sine & amp; Cosine integrals. Fourier transform, Fourier Cosine

transform, Fourier Sine transforms and their inverses. Finite FT, Application of FT to problems

on one and two dimensional heat flow problems.

Unit IV: Statistics and Probability (07 Hours)

Measures of central tendency, Measures of dispersion, Coefficient of variation, Moments,

Skewness and Kurtosis, Correlation and Regression, Reliability of Regression estimates.

Probability, Probability density function, Probability distributions: Binomial, Poisson, Normal,

Test of hypothesis: Chi-square test.

Unit V: Vector Calculus (08 Hours) Vector differentiation, Gradient, Divergence and Curl, Directional derivative, Solenoidal and Irrotational fields, Vector identities. Line, Surface and Volume integrals, Green's Lemma, Gauss's Divergence theorem and Stoke's theorem.

**Unit VI:**Applications of Partial Differential Equations (PDE)

(08Hours)

Basic concepts, modeling of Vibrating string, Wave equation, one and two dimensional Heat flow equations, method of Separation of variables, use of Fourier series, Applications of PDE to problems

of Chemical and allied engineering.

#### Text Books:

1. Higher Engineering Mathematics by B.V. Ramana (Tata McGraw-Hill).

2. Higher Engineering Mathematics by B. S. Grewal (Khanna Publication, Delhi).

#### **Reference Books:**

- 1. Advanced Engineering Mathematics, 10e, by Erwin Kreyszig (Wiley India).
- 2. Advanced Engineering Mathematics, 2e, by M. D. Greenberg (Pearson Education).
- B. Advanced Engineering Mathematics, 7e, by Peter V. O'Neil (Cengage Learning).
- 4. Differential Equations, 3e by S. L. Ross (Wiley India).
- 5. Introduction to Probability and Statistics for Engineers and Scientists, 5e, by Sheldon M. Ross (Elsevier Academic Press)

6.Partial Differential Equations for Scientists and Engineers by S. J. Farlow (Dover Publications, 1993)

#### Guidelines for Tutorial and Term Work:

i) Tutorial shall be engaged in four batches (batch size of 20 students maximum) per division.

ii) Term work shall be based on continuous assessment of six assignments (one per each unit) and

performance in internal tests.

Second Year of B.Tech. Big	otechnology (2019 Course)
215462: Fluid Fl	ow and Unit Operations
Cre	edit
TH: 03	
Teaching Scheme:	ExaminationScheme:
TH: 03hrs/week	ТН
	InSem:30TH
	End Sem :70
	Total :100
Prerequisites: -	0
Basic Knowledge of Physics and Mathematics.	
Problem Solving ability, Information manipulation	and Processing skills.
Course Objectives:	0.0.
1. To provide familiarity with the nature and particular the nature an	roperties of fluids and to understand the basic
equations of fluid flow along with their resp	ectiveapplications
2. To provide concept of pressure drop and energy	ergy losses during fluidflow
3. To familiarize students with unit operations equipment's	based on solid liquid systems and the related
4. To introduce students to particle technology	- basic concepts, laws and unitoperations
involved in pretreatment for different bioprocesses	
Course Outcomes:	
On completion of the course, learner will be able to	0—
A. Characterize fluids encountered in Biotech	nology industries and predict their flowbehavior.
B. Select and operate equipment based on the	properties of the material beinghandled.
C. Select, design and operate systems based of	nfluidization.
D. Select and operate equipment based on the	properties of the material beinghandled.
UnitI	(07Hrs)
System of units and conversions; Basics of Unit O	perations & Fluid – Definition and important
properties, viscosity, temperature and pressure dependent	dence,Newton'slaw,Classificationoffluids;

Laminar and turbulent flow – Concept of Reynold's number; Formation and separation of boundary layer.

#### UnitII

#### (08Hrs)

(**07Hrs**)

Laws of incompressible potential flow: Mass balances - Continuity equation and its applications to fluid dynamics, Energy balances in fluid dynamics: Euler's equation, Bernoulli's equation and its applications, Flow measurement using venturimeter, orificemeter and pitot tube; Hagen Poiseuelle equation, turbulent flow in pipes, effect of roughness, friction in flowing fluid, Moody, sdiagram; Minor losses in pipe flow, effect of fittings and valves.

#### UnitIII

Introduction to the dynamics of suspended particles: Lift and drag forces, drag coefficients; Flow of solids through fluids: Gravity settling of particles, Terminal velocity, Stoke's law and Newton'slaw, Freeandhinderedsettling,Sinkandfloatmethod,Differentialsettlingmethod;Sedimentation:Batch andcontinuous,equipmentsforsedimentation,Centrifugalsettling:Advantagesandequipments– cyclones and hydrocyclones.

#### UnitIV

Flow of fluid through solids: Characteristics of flow through packed beds - Darcy's equation, Equations for laminar flow (Kozeny Carmen) and turbulent flow (Burke Plummer), Ergun equation; Introductionoffluidization, minimumfluidization velocity, characteristics of fluidized systems, types of fluidization and their applications, Introduction to computational fluid dynamics (CFD)

#### UnitV

Fluid moving machinery-pumps, Types of pumps: positive displacement pump and centrifugal pumps, Characteristics of Centrifugal Pumps, NPSH; Valves and their types; Mixing and Agitation - Necessityofmixingandagitation, TypesofImpellers–Radialandaxialflow, Differentflowpatterns in mixing, Agitator selection, Calculation of power requirement, Mixing equipment; Mixing equipment for pastes and viscous material

#### UnitVI

Particle Technology: Properties of solids - Particle size and shape, Mixtures of particles, Determination of particle size; Screening - Standard screen series, screen analysis, Screen effectiveness and capacity, Industrial screening equipment; Size reduction: Crushing efficiency, energyrequirementscalculationsbyusingdifferentcrushinglaws,Sizereductionequipment:Primary crushers,secondarycrushers,Intermediateandfinegrinders,OpencircuitandClosedcircuitgrinding.

#### (07Hrs)

(08Hrs)

(08Hrs)

#### Books

#### Text:

1. R K Bansal, "A Textbook of Fluid Mechanics and Hydraulic Machines", 9th ed., Laxmi Publications, New Delhi,2004

2. McCabe, Smith, Harriot, "Unit Operations in Chemical Engineering", 7th ed., Tata McGraw Hill

#### Publications

#### **Reference:**

- 1. R K Rajput, "A Textbook of Fluid Mechanics", S. Chand Ltd., 2008
- 2. George Granger Brown, "Unit Operation"; Asia Publishing House', FirstEdition
- 3. Bird R.B., Stewart W.E., Lightfoot E.N. "Transport phenomena" 2ed., Wiley Publications, 2002.

	Savitribai Phule Pune University, Pune
Seco	ond Year of B. Tech. Biotechnology (2019 Course)
	215463: Heat Transfer
	Credit
TH: 03	
<b>Teaching Scheme:</b>	ExaminationScheme:
TH: 03hrs/week	THINSem:
	30TH Endsem
	:70
	Total:100
Prerequisites: -	
Basic Knowledge of Ph	ysics, Mathematics and fluid mechanics.
Problem Solving ability	with concept understanding and applications
<b>Course Objectives:</b>	<u> </u>
1. To make studen	ts aware of basic principles and mechanism of heat transferprocess.
2. To develop unde	erstanding of heat transfer systems and heat balanceequations.
3. To study various	s heat transfer equipments and theirapplication.
4. To provide infor	rmation about the scope and applications of heat transfer in the fieldof
biotechnology	
<b>Course Outcomes:</b>	
On completion of the co	burse, learner will be able to-
A. Understand and	apply knowledge of heat transferprinciples
B. Understand and	write heat balances of thesystem.
C. Choose suitable	heat transfer equipment for the requiredprocess.
D. Design heat tran	sferequipment.
<b>Course Contents</b>	
UnitI	(07Hrs)
Introduction:Modesofhe	eattransfer, conduction, convection, and radiation, Applications of heat
transfer in biotechnolog	gy;Conduction: Fourier's law of heat conduction, thermal conductivity
of liquids, gases and so	olids, Concept of thermal resistance, thermal conductance and contact
resistance,Introductiont	osteadyandunsteadystateconduction, Steadystateconduction in
infinitely long slab, hol	low cylinder and hollow spheres.

#### UnitII

#### (08Hrs)

Conduction: Heat transfer through Composite Materials, Thermal resistance in composite slab andcylinder,Heatlossesthroughpipe,thermalinsulationandoptimumthicknessofinsulation, properties of insulator, Heat transfer from extended surfaces with uniform crosssection, classification of extended surfaces, efficiency of longitudinal fin.

#### UnitIII

#### (07Hrs)

Convection: Types, Newton's law of cooling, Individual and overall heat transfer coefficient, Natural and forced convection in laminar and turbulent flow, Heat transfer with phasechanges: Condensation and Boiling liquids Modes and features, Dimensional Analysis, units of various quantities used in heat transfer, Importance of dimensional analysis in experimental design and datareduction,laminarandforcedconvectiongoverningequationsbyAlgebraicRayleigh's method

#### UnitIV

Convection: Analogy between heat, mass and momentum transfer. Principal and heat balance equation in laminar flow and empirical equations for turbulent flow through tube, through annulus, over the plate, Concept of thermal boundary layer and its significance Radiation:Fundamentalfactsanddefinitionofterms:Emissivity,absorptivity,blackbody,gray body,opaquebody,StefanBoltzmanlaw,Kirchoffslaw,Plankslaw,Wien'slaw,Theshape factor

#### UnitV

#### (07Hrs)

(08Hrs)

Heat exchange equipment: Types of heat exchangers including compact heat exchangers, parallelflowarrangement,foulingfactor,LMTDinparallelandcounterflow,Effectiveness NTU method, shell and tube heat exchanger.

#### UnitVI

#### (08Hrs)

Evaporation: Types of evaporators, performance, capacity and economy, Boiling point elevation, heat transfer coefficients, Material balance calculations, Multiple effect evaporators: Feed Forward and Feed Backward evaporator, Methods of feeding, capacity and economy, effect of liquid head and boiling point elevation

#### Books

#### Text:

- 1. S. P. Sukhatme, "A Textbook on Heat Transfer", 4th ed, Universities Press (India),2005
- 2. K. A. Gavhane, "Heat Transfer", 10 th edition, NiraliPrakashan, 2010
- 3. R.C. Sachdeva "Fundamentals of Engineering *Heat* and Mass *Transfer*" 5<sup>th</sup>edition, New age international Publisher,2010

#### **Reference:**

- 1. Frank Kreith, Mark Bohn, "Principles of Heat Transfer" 5th edition, PWS Publishing company,Boston(1997)
- 2. D.Q.Kern, "ProcessHeatTransfer", 11thed., TataMcGrawHillPublication, NewDelhi
- 3. Böckh, Peter, Wetzel, Thomas, Heat Transfer, Basics and Practice, Springer, 2012
- 4. Sinnout R.K. "Coulson Richardson's chemical engineering vol.6" Pergamon Press, 1993

Second Year of B.7	Fech. Biotechnology (2019 Course)
	215464: Microbiology
	Credits
TH: 03	
Teaching Scheme:	Examination Scheme:
TH: 03 hrs/week	TH Insem:30
	TH Endsem:70
	Total:100
Prerequisites: Basic knowledge of Bic	ology and Chemistry
Course Objectives:	
1. To introduce the concept, dev	velopment and significance of microbiology in day to
day life.	
2. Totrainthestudentstoasepticha	andlingtechniquesalongwith,culturing,utilizationand
elimination of microorganism	as from differentenvironments.
3. To make the students aware of	of the ubiquitous nature, diversity and growth of
different microorganisms and	I their significance inbiotechnology
4. To introduce the concept of d	liseases and role of microorganism in differentdiseases
Course Outcomes:	
<b>Course Outcomes:</b> On completion of the course, learner w	ill be able to-
<b>Course Outcomes:</b> On completion of the course, learner w A. Graduates are made aware of sig	ill be able to– gnificance ofmicrobiology
Course Outcomes: On completion of the course, learner w A. Graduates are made aware of sig B. Graduates would be trained to id	ill be able to– gnificance ofmicrobiology dentify, handle, cultivate and eliminate microorganisms
Course Outcomes: On completion of the course, learner w A. Graduates are made aware of sig B. Graduates would be trained to it from different environments	ill be able to– gnificance ofmicrobiology dentify, handle, cultivate and eliminate microorganisms
Course Outcomes: On completion of the course, learner w A. Graduates are made aware of sig B. Graduates would be trained to it from different environments C. Graduates are trained to underst	ill be able to– gnificance ofmicrobiology dentify, handle, cultivate and eliminate microorganisms tand growth requirements of differentmicroorganisms
Course Outcomes: On completion of the course, learner w A. Graduates are made aware of sig B. Graduates would be trained to in from different environments C. Graduates are trained to underst D. Graduates are able to understand	ill be able to– gnificance ofmicrobiology dentify, handle, cultivate and eliminate microorganisms tand growth requirements of differentmicroorganisms d significance of clean and aseptictechniques.
Course Outcomes: On completion of the course, learner w A. Graduates are made aware of sig B. Graduates would be trained to in from different environments C. Graduates are trained to underst D. Graduates are able to understand E. Graduates are made aware of di	ill be able to– gnificance ofmicrobiology dentify, handle, cultivate and eliminate microorganisms tand growth requirements of differentmicroorganisms d significance of clean and aseptictechniques. fferent microbiology related processes in the fields of

F. Graduates can identify general principles of microbiology, underlying cause, treatment and prevention of diseases.

#### **CourseContents**

#### **Unit I: Introduction to Microbiology:**

The History and scope of microbiology, Types of Microorganisms: Bacteria, fungi, algae, protozoa, Actinomycetes, viruses. Ultrastructure of Prokaryotic and Eukaryotic cell with functions. Microscopy and Staining methods, Bacterial classification

#### Unit II: Cultivation and Isolationofmicroorganisms:

Basic Nutritional and environmental Requirements- macro and micronutrient requirements, growth factors, Nutritional types of microorganisms: Photoautotrophs, Photo heterotrophs, Chemoautotrophs, Chemoheterotrophs, Environmental Effects on Bacterial Growth: Temperature, Oxygen, pH, Osmotic pressure classes, Extremophiles.

Design, types, composition and use of Media: Solid, liquid, semi-solid media, general purpose, selective media, differential media, enrichment media, enriched media. Microbial cultivation-Concept of pure culture, co-culture, mixed culture and colony characteristics. Isolation of microorganism & Pure culture techniques: Streak, Spread and Pour plate method. Culture Techniques for Anaerobes, Preservation and maintenance methods.

#### Unit III: Microbial Growth and Enumeration Techniques

Reproduction in microorganisms: Binary fission, asexual methods, Growth curve, Logarithmic representation of bacterial population, calculation of generation time and specific growth rate, Enumerationofbacteria-Directcount,StandardPlateCount,Microscopiccount,Turbidometry, Flow cytometer, Biomass determination, other microscopic and nucleic acid basedmethods.

Batch, Continuous and Synchronous cultures.

Unit IV: Controlof Microorganisms:

Concept and definition-Sterilization and disinfection. Control of Microorganisms by Physical and Chemical Agents: Concept, Heat (Dry, Moist),Pasteurization, Radiation, filtration, Chemical agents-Alcohols, Quaternary ammonium compounds, phenoliccompounds,

Disinfectants, Antimicrobial agents, Antibiotics, Concept of MIC and MBC, Drug resistance.

#### Unit V: Microbial interactionsinenvironment

Microbial interactions in environment, commensalisms, antagonism, symbiosis.Microbiology of air, soil, food, milk, water and waste water, Potability of water.

#### (07 Hrs)

(08 Hrs)

(08 Hrs)

(07Hrs)

(07 Hrs)

Unit VI:Medicalmicrobiology

The Epidemiology of Infectious Diseases, Human diseases caused by bacteria –Typhoid, Cholera, Tuberculosis. Viral diseases- Rabies, HIV, Influenza, fungal diseases – Candidiasis

#### **Books:**

#### Text:

1. Prescott Harley Klein, Microbiology, Fifth Edition, "Microbiology", 5th Edition, The McGraw Hill Companies,2002

2. Michael Pelczar, "Microbiology", 5th Edition, Tata McGraw-Hill Education, 1993

3. Michael T., Madigan, John M. Martinko, Jack Parker, "Brock biology of microorganisms", Prentice Hall,2000.

#### **Reference:**

1. A. J. Salle, "Fundamental Principles of Bacteriology", 7th edition, Tata McGraw-Hill education.

2. Roger Y. Stainier et al. "General Microbiology", 5th edition., PHIPublication.

3. Tortora, "Microbiology: An Introduction", 9th edition, Pearson Education India,2008.

4. Schlegel H.G. – "General Microbiology", 8th edition, Cambridge University Press, 1995.

5. Robert Cruikshank, "Medical Microbiology", Churchill Livingstone, 1975.

6. Thomas Jones Mackie, et al, "Mackie & McCartney medical microbiology: a guide to the

laboratory diagnosis and control of infection", Churchill Livingstone, 1989

	Savitribai Phule Pu	ne University, Pune
	Second Year of B.Tech. Bi	otechnology (2019 Course)
	215465	Biochemistry I Lab
	Cre	edit
		PR : 02
Teaching	Scheme:	ExaminationScheme:
PR: 04hr	s/week	TW :50
Prerequi	sites: - Basic knowledge of Biology and	chemistry
Course C	Objectives:	
1. To	develop basic lab skills	
2. То	e learn methods for the isolation of biomolect	ules
Course C	<b>Dutcomes:</b>	<u> </u>
	etion of this course, students will be able to	and a state and intermentation of neurlin
A. P.	repare laboratory reagents, learn to extract bi	omolecules, and interpretation of results.
B.Ju	stify the use of buffers in studying biological	systems
C.Id	entify biomolecules by suitable tests	
Suggeste	d List of Laboratory Assignments (Any	7 <b>8</b> )
Sr. No.	Group A	
1.	Preparation of molar and normal solution	ons
2.	Calibration of pH meter	
3.	Calibration of pipette	
4.	Preparation of Buffer solution	
Group B		
1.	Extraction of protein	
2	Qualitative analysis of protein	
2.	<b>C</b>	

4.	Iodine test for polysaccharide
Group	C
1.	The determination of pKa
2.	To check the resistance of buffer for pH change
3.	To check the reproducibility of the experiment
4.	Quantitative estimation of protein
5.	Qualitative Testing of Lipid
Guidel	ines for Instructor's Manual
1. Stud	ents should be briefed with Risk Assessment and Biosafety Levels
2. All t	he instruments to be validated before use
3. All ť	he experiments should be standardized
<b>4.</b> The	instructor is responsible for seeing that the consequences of student are rectified, including
the cor	rection of damages and violations and take-down of experiments.
Guidel	ines for Student's Lab Journal
1. Use a	a bound notebook.
2. Lab	notebooks should be done in pen and no erasing or white-out is allowed
3. Num	ber the pages
4. Title	and underline each lab exercise at the top of the page and date it. Each lab writeup should be
done se	eparately even if more than one exercise is performed in a lab period. Leave enough room in
the lab	notebook to complete the entire lab including results and discussions.
5. Briet	fly explain the lab exercise objectives in a few sentences.
6. Reco	ord observations, diagrams, and results from the exercise.
7. Cond	clude the report with a brief discussion in the essay form.
8. Wri	te neatly, be organized, and follow a standard format.
Note: T	The purpose of the lab notebook is to encourage students to compile and organize their
laborat	ory notes and to understand the purpose of the laboratory exercises and the meaning of their
results.	

#### Guidelines for Lab /TW Assessment Lab Assessment will be based on the following points

- 1. Present/Absent
- 2. A completion date of the journal
- 3. Regularity
- 4. Understanding

**5.** Presentation

#### Guidelines for Laboratory Conduction The following rules must be observed during laboratory conduction

- 1. Lab coat should be worn by students before entering the laboratory
- 2. Students shall keep their belongings on storage rack
- 3. Loose hair and flowing parts of apparel shall be properly tied before commencing of work
- 4. Enter the usage of chemicals and equipment's in a logbook
- 5. The instruction manual should be read before operating any instrument
- 6. Students should make aware of hazard warning symbols on reagent bottle
- 7. Protective devices must be worn when it is necessary to protect the eyes and face from splashes
- 8. All chemicals, glassware, reagents and plastic wares should be kept on their appropriate place after use
- 9. Reagents to be stored should be labeled with due discarding date

10.Instructions for proper disposal of waste material should be followed

- 10. Report accidental cuts or burns to the instructor immediately.
- 11. Perform the experiment. Collect data in a clear and organized fashion. Be sure to note the units for each measurement. Also, be sure to participate in the experiment rather than just recording data for your group

#### **General Guidelines:**

Before starting any experiment, clearly define the goals. What question are you answering or what principle are you trying to demonstrate? What data is needed to solve the problem? Identify the methods of measurement and instrumentation to be used.

Savitribai Phule I	rune University, rune
Second Year of B.Tech. Biotechnology (2019	course)
215466 :Fluid Flow	and Unit Operations Lab
Credit	
	PR : 01
Teaching Scheme	Examination Scheme:
PR: 02hrs/week	Tw: 25
Prerequisites: - Basic Knowledge of Physics ar	nd Mathematics.
Problem Solving ability, Information manipulation	on and Processing skills.
Course Objectives:	
1. To learn the fluid properties and fundamentals	of fluid flow.
2. To introduce the flow measuring devices.	
3. To impart knowledge in measuring pressure, d	lischarge and velocity of fluid flow.
4. To provide practical knowledge in verification	of principles of fluid flow.
Course Outcomes:	
On successful completion of the course students	will be able to
A.Understand basic physics of fluid.	
B. Calculate and design engineering applications	involving fluid.
C.Analyse flow in terms of mass, momentum and	1 energy balance.
D. Predict the coefficient of discharge for flow th Guidelines for 1	rough Venturimeter and Orifice. Instructor's Manual
The Fluid Flow Laboratory is a substantial par	t of the course "Fluid Flow and Unit Operations"
and is constructed to complement the lecture	portion of the course. The labs are designed to
provide the student with a physical understa	anding of the fundamental principles and basic
equations of fluid mechanics. This understandir	ng is gained through the application of "text book"
concepts and equations to real problems.	
The student is to read the lab manual chapter as tothe lab. Some labs contain thought questions of proceeding.	ssigned to each laboratory period BEFORE coming or require that you perform some derivations before

#### Guidelines for Lab /TW Assessment

Attendance is required for all of the lab sessions. Each session, except one demonstration activity, requires the completion of a formal labreport. These reports are the basis of your final lab grade. Each assignment represents a substantial fraction of your total score.

#### **Guidelines for Laboratory Conduction**

#### Safety Guidelines:

Beverycarefulandawareofthevariousexperimentcontrols(startbutton,stopbutton,speedcontrol) for each labsession.

Ask lab instructor, if you are not sure about what to

do. Make sure all spilled liquids are wiped

upimmediately. Do not leave experimentsunattended.

Any injuries should be reported immediately for proper care.

#### **General Guidelines:**

Before starting any experiment, clearly define the goals. What question are you answering or what principle are you trying to demonstrate? What data is needed to solve the problem?

Identify the methods of measurement and instrumentation to be used. At the research stations, "play around" with the equipment so that you understand how the instruments work, what you are measuring, and how what you are measuring connects with the physics of the problems at hand. Perform the experiment. Collect data in a clear and organized fashion. Be sure to note the units for each measurement. Also, be sure to participate in the experiment rather than just recording data for your group.

Suggested List of Laboratory Assignments (Any 8)

 Sr. No.
 Determination of viscosity

 1.
 Determination of viscosity

 2.
 Flow through venturimeter

 3.
 Flow through orifice meter.

 Group B

 1.
 Friction during flow through pipe

 2.
 Verification of Bernoulli's theorem

3.	Verification of Stoke's law		
Group	C		
1.	Verification of Darcy's law		
2.	Flow through pipe fittings		$\sim$
3.	Flow through packed bed		
		G	
		S.	
		<b>~</b> ' <b>~</b> `	

	Savitribai Phule Pu	ne University, Pune
	Second Year of B.Tech. Bi	otechnology (2019 Course)
	215467:	Heat Transfer Lab
	Cre	edit
		PR : 01
Teaching	Scheme:	Examination Scheme:
PR: 02hrs	s/week	UK :50
Prereauis	ites: - Basics of mathematics and physic	cs. Fundamental principles of heat transfer and
theory con	itent related to experiment	
Course O	bjectives:	
1. To 1	nake students aware of application of ba	asic principles and mechanism of heat transfer
proc	cess	
2. To a	levelop understanding of various heat tr	ansfer systems and heat balance equations.
3. To s	study various heat transfer equipments, o	lesigns and their application.
4. To s	study the industrial scope and application	ns of heat transfer in the field of biotechnology
Course O	utcomes:	
In complet	tion of the course lab, learner will be abl	e to-
A. Unc	lerstand and apply knowledge of basic h	eat transfer principles
B. Unc	lerstand and write heat balances around	the system.
C. Und	lerstand design aspects and choose suita	ble heat transfer equipment for the required
proc	cess/application	
D. Calo	culate the rate and area of heat transfer, i	individual and overall co-efficient of heat transfer,
effic	ciency etc.	
Suggested	List of Laboratory Assignments (An	y 8)
Sr. No.	Group A – Conduct	tion, Convection and Radiation
1	Heat transfer from fin in a natural convec	tion
2	Heat transfer in forced convection	
	Composite wall apparatus	
3		

	Co- current and countercurrent heat exchanger
7	Plate type heat exchanger
8	Shell and tube heat Exchanger
	Group C –Study Experiments
9	Double effect evaporator
10	Agitated Vessel
Guid 1. St 2. A	elines for Instructor's Manual udents should be briefed with risk assessment and safety levels. Il the instruments to be validated before use.
3. A	Il the experiments should be standardized.
4. T	ne instructor is responsible for seeing that the consequences of student are rectified, including
co	rrection of damages, violations and take down of experiments.
Guid 1.	elines for Student's Lab Journal Use a bound notebook.
2.	Lab notebooks should be done in pen and no erasing or white-out is allowed
3.	Number the pages
3. 4.	Number the pages Title and underline each lab exercise at the top of the page and date it. Each lab write-up
3. 4.	Number the pages Title and underline each lab exercise at the top of the page and date it. Each lab write-up should be done separately even if more than one exercise is performed in a lab period. Leave
3. 4.	Number the pages Title and underline each lab exercise at the top of the page and date it. Each lab write-up should be done separately even if more than one exercise is performed in a lab period. Leave enough room in the lab notebook to complete the entire lab including observations,
3. 4.	Number the pages Title and underline each lab exercise at the top of the page and date it. Each lab write-up should be done separately even if more than one exercise is performed in a lab period. Leave enough room in the lab notebook to complete the entire lab including observations, calculations, results and discussions.
3. 4. 5.	Number the pages Title and underline each lab exercise at the top of the page and date it. Each lab write-up should be done separately even if more than one exercise is performed in a lab period. Leave enough room in the lab notebook to complete the entire lab including observations, calculations, results and discussions. Briefly explain the lab exercise objectives in a few sentences.
<ol> <li>3.</li> <li>4.</li> <li>5.</li> <li>6.</li> </ol>	Number the pages Title and underline each lab exercise at the top of the page and date it. Each lab write-up should be done separately even if more than one exercise is performed in a lab period. Leave enough room in the lab notebook to complete the entire lab including observations, calculations, results and discussions. Briefly explain the lab exercise objectives in a few sentences. Record observations, diagrams and results from the exercise.
<ol> <li>3.</li> <li>4.</li> <li>5.</li> <li>6.</li> <li>7.</li> </ol>	Number the pages Title and underline each lab exercise at the top of the page and date it. Each lab write-up should be done separately even if more than one exercise is performed in a lab period. Leave enough room in the lab notebook to complete the entire lab including observations, calculations, results and discussions. Briefly explain the lab exercise objectives in a few sentences. Record observations, diagrams and results from the exercise. Conclude the report with a brief discussion in essay form.
<ol> <li>3.</li> <li>4.</li> <li>5.</li> <li>6.</li> <li>7.</li> <li>8.</li> </ol>	Number the pages Title and underline each lab exercise at the top of the page and date it. Each lab write-up should be done separately even if more than one exercise is performed in a lab period. Leave enough room in the lab notebook to complete the entire lab including observations, calculations, results and discussions. Briefly explain the lab exercise objectives in a few sentences. Record observations, diagrams and results from the exercise. Conclude the report with a brief discussion in essay form. Write neatly, be organized and follow a standard format.

#### **Guidelines for Lab /TW Assessment**

Term work marks distribution should be carried out based on following points:

- 1. Regularity and sincerity of students during lab practicals
- 2. Journal presentation
- 3. Understanding of the experiment
- 4. Performance in unit tests
- 5. Attendance during theory lectures

#### **Guidelines for Laboratory Conduction**

- 1. Please do not touch open wires.
- 2. Make the connection as per the circuit diagram.
- 3. Check permissible range heat input written on the setup before setting parameters current and voltage.
- 4. Ensure that the connections are properly tightened then Start the power supply.
- 5. After the experiment is over reduce the dimmerstat to zero.
- 6. Switch off the power supply and remove the connections.
- 7. Turn off all valves and water supply completely.

#### Savitribai Phule Pune University, Pune Second Year of B.Tech. Biotechnology (2019 Course) 215468:Microbiology Lab

Credit

	PR : 02
Teaching Scheme: PR: 04hrs/week	Examination Scheme: PR :50 Total :50
Course Objectives:	
1. To train the students to aseptic	handling techniques and elimination of
microorganisms from different enviro	onments.
2. To train the students different method	ls of isolation and culturing of microorganisms
from different sources.	
3. To make the students aware of the	e ubiquitous nature, diversity and growth of
different microorganisms	
4 To aware students for the significance	e of microorganism in Biotechnology
+. To aware students for the significance	e of interoorganism in Diotechnology
Course Outcomes:	
A Graduates are able to understand sign	able to-
A. Graduates are able to understand sign	intelace of clean and aseptic techniques.
B. Graduates would be trained to	identify, handle, cultivate and eliminate
microorganisms from different enviro	onments
C. Graduates are trained to unde	rstand growth requirements of different
microorganisms	
D. Graduates are made aware of differe	nt microbiology related processes in the fields
of agriculture, medicine and food.	
Suggested List of <mark>L</mark> aboratory Assignments Group A Study of laboratory equipment's	5
1. Introduction and working of basic lab	poratory instruments.
Aseptic Handling Techniques-Laminar	Flow System, Autoclaving,
Group B Culture media preparation and a	aseptic techniques
Cotton Plug preparation. Plate and pipe	ettes wrapping & autoclaving

2. Nutrient Media preparation and sterilization for Bacteria and Fungi-- Broth, Plate and Slant Culture transfer and preservation techniques- subuturing, Glycerol stocks.

#### **Group C Bacterial Morphology and Staining**

- 3. Smear Preparation and Simple Staining
- 4. Differential Gram's staining.

#### Group D Isolation of microorganisms:

- 5. Serial dilution
- 6. Spread plate method
- 7. Pour plate method
- 8. Streak plate method
- 9. Colony characteristics- Size, Shape, color, margin, Consistency. Opacity, Gram's Character

Group E Microbial Growth Characteristics

- 10. Determination of generation time of E.coli
- 11. Effect of temperature/pH/atmospheric oxygen on growth of *E.coli*

Group F Study of Molds and yeast (any one)

- 12. Standard plate count/Total Viable Count from food/milk
- 13. Morphology of fungi on direct microscopic count/media.
- 14. Antibiotic sensitivity test

#### **Guidelines for Instructor's Manual**

- 1. Discuss the syllabus for the course
- 2. Go through the general procedures for lab safety
- 3. Review the guidelines for laboratory reports
- 4. Practice some of the scientific calculations that will be used throughout the semester
- 5. Practice some basic laboratory methods such as the use of balances, pipets, and Micropipettes
- 6. Students should be briefed with Risk Assessment and Biosafety Levels
- 7. All the instruments to be validated before use
- 8. All the experiments should be standardized

The instructor is responsible for seeing that the consequences of student are rectified, including correction of damages and violations and take-down of experiments

#### **Guidelines for Student's Lab Journal**

Please read these instructions now and refer to them regularly! These instructions must be followed carefully.

- 1. Use a bound notebook for notes and lab observations.
- 2. Lab notebooks should be done in pen and no erasing or white-out is allowed
- 3. For the Journal use Journal Pages and ensure you draw neat diagrams wherever necessary
- 4. Number the pages
- 5. Title and underline each lab exercise at the top of the page and date it.
- 6. Each lab write-up should be done separately even if more than one exercise is performed in a lab period.
- 7. Briefly explain the lab exercise objectives in a few sentences.
- 8. Record observations, diagrams and results from the exercise.
- 9. Conclude the report with a brief discussion in essay form.
- 10. Write neatly, be organized and follow a standard format.

Note: The purpose of the lab notebook is to encourage students to compile and organize their laboratory notes and to understand the purpose of the laboratory exercises and the meaning of their results.

#### Guidelines for Lab /TW Assessment

Lab assessment will be based on following points

- 1. Present / Absent
- 2. Completion date of journal
- 3. Regularity
- 4. Understanding
- 5. Presentation

#### **Guidelines for Laboratory Conduction**

Basic Principles for Students Working in Cell Biology Laboratories :

The following rules must be observed at all times to prevent accidental injury to and

infection of yourself and others and to minimize contamination of the lab environment:

- 1. Never place books, backpacks, purses, etc., on bench tops. Always place these in the assigned cubicles. Keep manuals and pens on pull-out desks.
- 2. Clean your work area with dilute bleach solution at the beginning AND end of each

```
lab.
```

- 3. Wash your hands with soap and dry with paper towels when entering and leaving the lab.
- 4. Wear a lab coat at all times while working in the lab to prevent contamination or accidental staining of your clothing.
- 5. Closed-toe shoes (no sandals) are to be worn in the lab.
- 6. Long hair must be tied back to prevent exposure to flame and contamination of cultures.
- 7. Do not place anything in your mouth or eyes while in the lab. This includes pencils, food, and fingers. Keep your hands away from your mouth and eyes.
- 8. Eating and drinking are prohibited in the lab at all times. This includes gum, cough drops, and candy.
- 9. Never pipet by mouth. Use a mechanical pipetting device.
- 10. Do not remove media, equipment, or bacterial cultures from the laboratory. This is absolutely prohibited and unnecessary.
- 11. Do not place contaminated instruments such as inoculating loops, needles, and pipettes on bench tops. Loops and needles should be sterilized by incineration, and pipettes should be disposed of in designated receptacles of bleach solution.
- 12. Carry cultures in a test tube rack when moving around the lab or when keeping cultures on bench tops for use. This prevents accidents and contamination of your person or belongings.
- 13. Immediately covers spilled cultures or broken culture tubes with paper towels and then saturate them with disinfectant solution. Notify your instructor that there has been a spill. After 15 minutes, dispose of the towels and broken items as indicated by your instructor.
- 14. Report accidental cuts or burns to the instructor immediately.
- 15. At the end of each lab session, place all cultures and materials in the proper disposal area.
- 16. Electronic devices should not be brought into the lab. This includes, but is not limited to iPods, MP3 players, radios, cell phones, and calculators.

#### **General Guidelines:**

Before starting any experiment, clearly define the goals. What question are you answering or what principle are you trying to demonstrate? What data is needed to solve the problem?Identify the methods of measurement and instrumentation to be used.

**Reference Books:** Laboratory Exercises in Microbiology-Harley Prescott 5<sup>th</sup> Edition

#### 215469:Mandatory Audit Course 3

In addition to credits courses, it is recommended that there should be audit course (non-credit course). Audit course is for the purpose of self-enrichment and academic exploration. Audit course carry no academic credit. Selection of audit courses helps the learner to explore the subject of interest in greater details resulting in achieving objective of audit course's inclusion. Evaluation of audit course will be done at institute level. Method of conduction and method of assessment for audit courses is suggested.

#### **Criteria:**

The student registered for audit course shall be awarded the grade PP and shall be included such grade inthesemestergradereportforthatcourse, provided students has the minimum attendance as prescribed by the Savitribai Phule Pune university and satisfactory in-semester performance and secured a passing grade in that audit course. No grade point is associated with this "PP" grade and performance in these courses is not accounted in the calculation of the performance indices SGPA and CGPA.

#### Guidelines for Conduction and Assessment (Any one or more of following but not limited to)

- 1. Lecture/Guestlecture
- 2. Visit (Social/field) and reports
- 3. Demonstrations
- 4. Surveys
- 5. Mini project
- 6. Hands on experience on specific focusedtopic.
- 7. Seminar/Workshop

#### Guidelines for Assessment (Any one or more of following but not limited to)

- 1. Writtentest
- 2. Quiz
- 3. Demonstrations/practicaltest
- 4. Presentations
- 5. IPR/publication
- 6. Report

#### Audit course 2 Options (Anyone)

- 1. Bio safety inBiotechnology
- 2. Leadership and PersonalityDevelopment

# Semester II

Savitribai Phule Pr	une University, Pune
Second Year of B. Tech. B	Biotechnology (2019 Course)
215470	:Biochemistry II
Cr	edits
TH: 03	TUT:01
Teaching Scheme:	ExaminationScheme:
TH: 03 hrs/week	TH Insem : 30
TUT: 01 hrs/week	TH Ensem :70
	Total:100
Prerequisites: - Basic Knowledge of structure and	nd function of biomolecules
Course Objectives:	0
1. To examine the main pathways of metabolic	blism and how they are integrated withother
Pathways within thecell.	
2. To provide an understanding about disord	ders occur due to defects inmetabolism
3. To understand the basic functioning ofen:	zymes
Course Outcomes:	
On completion of the course, the learner will be	able to-
A. Understand the metabolism of major clas	ses of biomolecules like carbohydrates,
lipidsand proteins in living systems and the	heir relation inbetween
B. Apply knowledge of biochemistry to und	erstand genetic diseases involving disordersof
metabolism	
C. Understand the functioning of enzymes	
Course Contents	
UnitI Principles of Bioenergetics: Bioenergetics and	(07Hrs) thermodynamics, Phosphoryl group transfer and
ATP, The free energy of hydrolysis of ATP	within cells, Energy required to the assembly of
informational macromolecules, Transphosphoryl	ations between nucleotides, Potential phosphoryl
group donor	
UnitII	(08Hrs)
Primary, secondary (alpha helix, beta sheets, turn	ns, and loops), tertiary (myoglobin) and quaternary
structure(hemoglobin)ofprotein,Namingandclass	ificationofenzymes,enzymecofactors,the kinetics
ofenzyme-catalyzedreactions,MichaelisMentene	quation,the
effectofpH,temperatureonenzymeactivity.	

#### UnitIII

Glycolysis,PreparatoryphaseandPayoffphase,Canceroustissuesandglucosemetabolism,feeder pathwaysforGlycolysis,Fatesofpyruvateunderanaerobiccondition,gluconeogenesisPentose phosphate pathway of glucose oxidation, Glucose 6-Phosphate dehydrogenase deficiency

### UnitIV

# Glycogens break down, Synthesis of glycogen and Starch and the role of nucleoside diphosphate sugars, TCAcycle, discovery of TCAcycle, intracellular location of the enzymes of the TCAcycle,

reactions of the TCA cycle, regulation of the TCA cycle, electron transport chain, oxidative

phosphorylation, Cori cycle, Disorders of carbohydrate metabolism E.g., glycogen storage disease

#### UnitV

Protein purification (molecular size, solubility difference, electric charge, selective adsorption, affinity chromatography), digestion and absorption of proteins, removal of nitrogen in amino acid degradation, ammoniatoxicity, pathways of a mino acid degradation, inherited defects of the urea

#### cycle

#### UnitVI

# Digestionandabsorptionoflipidsbeta-oxidationoffattyacid,ketonebodies,ketoacidosis,oxidation of fatty acid in peroxisomes, degradation of odd chain fatty acid, oxidation of PUFA. Synthesis of fattyacid:FormationofmalonylCoA,the roleoffattyacidsynthaseinthe synthesisoffattyacids,transferof

acetyl CoA to the cytoplasm, sources of NADPH for fatty acid synthesis.

#### **Text Book:**

 D J Voet, J G Voet, C W Pratt, "Principles of Biochemistry", 3rd ed., John Wiley & Sons, Inc.
 2008 2. D T. Plummer, "An Introduction to practical biochemistry", Tata McGraw Publishing Company Ltd, 1988

#### Reference Book:

1. J H Weil, "General Biochemistry", New Ages International (P) Ltd.1997.

2.JMBerg, JLTymoczko, LStryer, "Biochemistry", 6thed., Freeman WH&Company, New York, 2007
3. D L Nelson, M M Cox "Principles of Biochemistry", 4th ed., W.H. Freeman and company,

New York, 2007

#### (07Hrs)

(08Hrs)

# (07Hrs)

(08Hrs)

Savitribai Phu	ıle Pune University, Pune
Second Year of B. Te	ch. Biotechnology (2019 Course)
215471: C	ell Biology & Tissue Culture
	Credits
TH :03	TUT:01
Teaching Scheme:	ExaminationScheme:
TH: 03 hrs/week	TH Insem : 30
TUT: 01 hrs/week	TH Endsem :70
December 2014 - Decis have a labor of Distance	Total :100
Prerequisites: - Basic knowledge of Biolog	gy and Chemistry
1 To study call structure and functions	of organallafunctions
<ol> <li>To study cell structure and functions</li> <li>Exposure on transportations through</li> </ol>	
<ol> <li>Exposure on transportations through</li> <li>To focus on different receptors and t</li> </ol>	model of signaling
4 To introduce the concept of cellsion	aling
5 Different cell types offer differentfu	nctionality
6 Application of cell biology inbiotech	netonality
Course Outcomose	niology
On completion of the course learner will be	a able to
A To understand the structure and func	tion of eukervoticcells
B. To introduce basic techniques in cell	lbiology
C. Understand the application of cell bi	ology in disease and development of the rapy
D. Give hands-on experience in cell bio	blogy techniques such as in-vitro cellculture
2. Sive hunds on experience in cell of	No5, comiques such as in vitro concutture
Course Contents	
Lourse Contents	(0711)
Structure and function of Call I	(0/Hrs)
Structure and function of Cell –I	tuents of the call, sub callular comparison stations and
organalles such as puclous, perovisiones ar	doplasmia retignium. Colai apparatus, hussoares
	CODIASHIIC TEHCHIIIII. CODI ADDATAIIIS. IVSOSOMES.

#### UnitII

#### (08Hrs)

#### Structure and function of Cell -II

Biomembranes, cytoskeleton structure and function, cell movement, ion channels transportof small molecules, Membrane proteins: Carrier proteins and active membrane transport, Endocytosis/Exocytosis/Pinocytosis, extracellular vesicles, cell–cell interactions, Junctions–

Adhesion, Tight, Gap, Plasmodesmata.

#### UnitIII

#### (07Hrs)

#### Intracellular signaling and Cell Cycle

Signaling molecules - receptors, functions, Pathways of intracellular signal transduction. General principles of communication, morphogen. Types of receptors GPCR, etc. Cell cycle, Cell cycle control system, Karyokinesis, Cytokinesis, Control of cell division and growth, Mitosis, Meiosis, Cell death and cell renewal - Programmed cell death Apoptosis.

#### UnitIV

#### **Tissues and Cancer**

Epithelial tissue, connective tissue, muscle tissue, nervous tissue, blood. Stem cells: Hematopoietic stem cells & embryonic stem cells. Cancer: Development of Cancer and properties of Cancer.

#### UnitV

#### Animal Cell and Tissue culture

Animal tissue culture: tissue culture media, Media types, Types of culture- adherent cell lines and suspension cell cultures, Passaging, Cell separation, characterization, cryopreservation of animal cells, contamination and cytotoxicity.

#### UnitVI

#### **Plant Tissue culture**

Plant tissue culture: Totipotency, Requirements, Plant growth hormones. Types of culture:Callus culture, Pollen culture, Anther culture, Protoplast fusion. Application: Production of secondary metabolites, transgenic plants.

#### Books:

#### **Text Book:**

- 1. Karp, "Cell and Molecular Biology" John Wiley and Sons Pvt.Ltd
- 2. Sudha Gangal, 'Animal tissue culture', Orient Longman, 2006
- 3. The Cell: A molecular approach by Geoffrey M.Cooper. ASM Press, Pages: 673

(08Hrs)

(08Hrs)

#### **Reference Book:**

- 1. Cooper G.M. & Hausman, "The Cell", fifth edition, ASMPress.
- Bruce Alberts, Alexander Johnson, Julian Lewis, Martin Raff, Keith Roberts, Peter Walter.
   "MolecularBiologyoftheCell",4thedition,GarlandPublishing,NewYork,London,(2002)
- 3. Harvey Lodishetal "Molecular Cell Biology", 4th edition, NewYork
- 4. M.K. Razdan, "Introduction to Plant Tissueculture".
- 5. Freshney Ian ,"Animal tissueculture"
- 6. Tortora and Grabowasky "Human anatomy and physiology", Wileypublication

	21547	2: Thermodynamics
	Cı	redit
	TH :03	TUT:01
Teachi	ing Scheme:	ExaminationScheme:
TH:	03hrs/week	ТН
TUT:	01hr/week	Insem:30TH
		Endsem :70
		TW:25
Duouso		Total :125
Prereg	Decision and the formula and the i	
•	Basic concepts of fundamental and deri	ved properties like mass, acceleration, kinetic
	Desig his chamical activities and common	
•	Basic biochemical pathways and genera	i cenmetabolism.
Course	e Objectives:	
1. 2	To introduce students to basic concepts	and first law othermodynamics
2.	To develop an understanding of neat end	ergy and its application to industrial and
3	Tomakastudantsawarashoutfunctioning	Sindustrialsystems by introducing the multiple
5.	second law of Thermodynamics	sindusu taisystemsoy nu odue nightem with
4	Provideanunderstandingoftheconceptofe	auilibriumanditsrelevanceto solution
	thermodynamics	
5.	To introduce students to concepts related	d to thermodynamics of chemically reacting
	systems	
6.	To demonstrate the applications of them	nodynamic principles in context of biological
	systems	

#### **Course Outcomes:**

On completion of the course, learner will be able to-

- A. Analyze simple systems from thermodynamic aspect as also to apply this knowledge to evaluate the efficiency and feasibility of a physical or biochemicalprocess
- B. Characterize systems comprising of solutions and mixtures
- C. Predict equilibrium, conversion and efficiency of achemicalor biochemicalprocesses
- D. Examine biological systems and biochemical reactions from a thermodynamic view and understand the various applications in thiscontext

#### **Course Contents**

#### UnitI

**Introduction to engineering thermodynamics**: Scope and importance, fundamental and derived quantities, Joule's experiment, Internal Energy, The First law of Thermodynamics, Energy Balance for Closed Systems, Thermodynamic State and State Functions, equilibrium, The Phase Rule, The Reversible Process, Constant-V and Constant-P Processes, Enthalpy,Heat Capacity, Mass and Energy Balances for open systems, Processes Involving Ideal Gases

#### UnitII

#### (08Hrs)

(07Hrs)

(07Hrs)

**HeatEffects:**Sensibleandlatentheateffects,temperaturedependenceofheatcapacity,standard heat of reaction, standard heat of formation, standard heat of combustion, Hess's law; Temperature dependence of standard heat of reaction, heat effects of industrial reactions

#### UnitIII

**Second law of Thermodynamics:** Limitations of First Law, Heat Engines, Heat Pump and Refrigerator, Statement of second law, Criterion for irreversibility, Carnot cycle and theorems, Clausius inequality, Concept of entropy and its calculation, mathematical statement of 2<sup>nd</sup>law, Principle of entropy increase

#### UnitIV

#### (08Hrs)

**SolutionThermodynamics:**Fundamentalpropertyrelations,TheChemicalPotentialandPhase Equilibria, Partial Properties, Ideal-Gas Mixtures, Fugacity and Fugacity Coefficient: Pure species,FugacityandFugacityCoefficient:SpeciesinSolution,GeneralizedCorrelationsforthe Fugacity Coefficient, The Ideal Solution, Excess Properties

#### UnitV

#### (07Hrs)

**Chemical Reaction Equilibria:** Application of the criteria for equilibrium to chemical reactions, the standard Gibbs free energy change and the equilibrium constant; effect of temperature on equilibrium constant, evaluation of the equilibrium constant, relation of equilibrium constant to composition, calculation of equilibrium conversion for single reaction; The phase rule and Duhem's theorem for reacting systems

#### UnitVI

(08Hrs)

Application of thermodynamics to biological systems: Energy transformations in biological systems, Examples of applications of laws of thermodynamics to bio-systems, Gibb's free energy concept for bio-changes and its applications; Thermodynamics of biochemicalchanges - Energy Yielding and Energy Requiring Reactions, feasibility of individual steps and overall reactions

#### **Books:**

#### **Reference:**

1. J M Smith, H C Van Ness, "Introduction to Chemical Engineering Thermodynamics", 7<sup>th</sup> ed., McGraw-Hill Education, 2005

1. T E Daubert, "Chemical Engineering Thermodynamics", McGraw-Hill Inc., 1985

Text:

- 1. K V Narayanan, "A Textbook of Chemical Engineering Thermodynamics", PHI Learning Pvt. Ltd.,2004
- 2. Y V C Rao, "Chemical Engineering Thermodynamics", University Press, 1997
- 3. D T Hayne, "Biological Thermodynamics", Cambridge UniversityPress

Second Year of B.Tech. I	Biotechnology (2019 Course)
215473:Genetic	cs and Molecular Biology
Cı	redits
TH: 03	TUT:01
Teaching Scheme:	Examination Scheme
TH: 03 hrs/week	THInsen
TUT: 01 hrs/week	30THEndsen
	Total:10
Prerequisites: Basic knowledge of Biology	
Lourse Objectives:	a consting & tomela similation of DNA
1. To understand the process of DNA rep	liestion
2. To understand the process of DNATep	tion
4 To understand the process of translatio	
Course Outcomes:	
On completion of the course, learner will be a	able to-
A. Understand the concept of Mendelian	genetics & nucleic acidstructure.
B. Comprehend the replicationmechanism	n.
C. Emphasize on role of different types o	of RNA & transcriptionprocess
D. Deduce the process of translation.	
Course Contents	
Unit I:MendelianGenetics	(07Hrs)
Introduction, Mendelianinheritancepatternstud	lyandlawsofheredity,Co-dominance,linkage,
linkage maps, Hardy-Weinberg Equation, Mo	odel systems like Drosophila, C. Elegans, Zebra
fish, Arabidopsis	
Unit II:DNATopology	(08Hrs)
Watson-Crick'sdiscoveryofstructureofDNA,M	MacLeodandMcCarty'sexperiment,Hershey and
Chase's experiment, Structure of DNA: A,	B (Watson-crick model), and Zstructure,

Chargaff's rule, Physicochemical properties of	DNA, DNA supercoiling, DNA packaging:
Chromosome, Chromatin, Chromatid, Euchrom	natin and Heterochromatin.
Unit III:DNAReplication	(07Hrs)
Introduction to replication of DNA, Chemis	try of DNA synthesis, Mechanism of DNA
polymerase, Replication Fork, DNA synthesi	is at the replication fork, Initiation of DNA
replication, Elongation of DNA replication, Fin	ishing replication, Telomere replication.
Mutation and Repair	C
UnitIV:RNA	(08Hrs)
TypesofRNA,Codingandnon-codingRNAs,tRN	A,mRNA,rRNA,andsmallRNAs,introns and
exons, chemistry of RNA splicing, splicing	pathways, alternative splicing, ribozyme,
importance of RNA, RNA world theory.	
UnitV:Transcription	(07Hrs)
Transcription, RNA polymerase, Transcription	cycle in bacteria, concept of Operon,
Transcription cycle in eukaryotes, Reverse Tran	nscriptase, .Regulation.
UnitVI:Translation	(08Hrs)
Genetic code, Protein biosynthesis, Initiation of	f translation, Translation elongation,
Termination of Translation, regulation, posttrar	islational modifications, protein synthesis in
prokaryotes and eukaryotes, chaperones, heat sl	hock proteins.
Books:         Text:         1. JamesD.Watson,TaniaA.Baker,Stephend         th         5 edition, Dorling Kindersley (India) Pv         2. Freifelder D., "Molecular Biology", Jon	P.Bell,"MolecularBiologyoftheGene" /t.Ltd. nes and Bartlett Publishers,(1987)
Reference:	
1. Benjamin Lewin, "Gene IX", Oxford Unive	rsity Press, Oxford, New York,(2000)
2. Bruce Alberts, Alexander Johnson, Julian L	ewis, Martin Raff, Keith Roberts, PeterWalter
"Molecular Biology of the Cell", 4th edit	ion, Garland Publishing, New York, Londor
(2002)	
3. T.A. Brown, "Genomes" John Wiley and Sc	ons Pvt.Ltd
/ 4. Ansumbel F.M, Brent R, Kingston R.E, M	Moore D.D., 'Curernt protocols in Molecula
Biology' Green Publishing Associates,(1988	8)
5. Old R W and Primrose SB, "Principles of G	ene manipulations: An introduction toGenetic

# Savitribai Phule Pune University, Pune Second Year of B.Tech. Biotechnology (2019 Course) 215474: Biochemistry II Lab Credit **PR**:01 **ExaminationScheme: Teaching Scheme:** TW :25 PR: 02hrs/week **PR:50** Total :75 Prerequisites: - Basic knowledge of Biochemistry I **Course Objectives:** 1. To learn fundamental approaches for conduction of the experiment 2. To give hands-on experience to the students on the isolation of biomolecules 3. To make students acquainted with the kinetics of biocatalyst **Course Outcomes:** On completion of this course, students will be able to A. Conduct experiments with satisfactory analysis of data and interpretation of results

B.To apply methods for extraction of biomolecules

C.Understand the functioning of enzymes (biocatalysts)

#### Suggested List of Laboratory Assignments (Any 8)

Sr. No.	Group A
1.	Determination of Amax
2.	
	Quantitative estimation of reducingsugar
3.	
	Protein estimations by Lowry/ Biuret/ Bradfordmethod

#### **Group B**

1.	Estimation of protein by spectrophotometer at 280nm.
2.	Extraction of cholesterol/lipids
3.	Estimation of cholesterolconcentration

1.	Isolation of Enzyme
2.	Varying enzymeassay
3.	To determine Km forenzyme
Guide	lines for Instructor's Manual
1. Stud	dents should be briefed with Risk Assessment and Biosafety Levels
2. All	the instruments to be validated before use
3. All	the experiments should be standardized
<b>4.</b> The	instructor is responsible for seeing that the consequences of student are rectified, including
the	correction of damages and violations and take-down of experiments.
Guide	lines for Student's Lab Journal
1. Use	a bound notebook.
2. Lab	notebooks should be done in pen and no erasing or white-out is allowed
3. Nun	nber the pages
4. Title	e and underline each lab exercise at the top of the page and date it. Each lab writeup should be
don	e separately even if more than one exercise is performed in a lab period. Leave enough room
in th	he lab notebook to complete the entire lab including results and discussions.
5. Brie	fly explain the lab exercise objectives in a few sentences.
6. Rec	ord observations, diagrams, and results from the exercise.
7. Con	clude the report with a brief discussion in the essay form.
8. Wri	te neatly, be organized, and follow a standard format.
Note: ' labora results	The purpose of the lab notebook is to encourage students to compile and organize their tory notes and to understand the purpose of the laboratory exercises and the meaning of their .
Guide Lab A	lines for Lab /TW Assessment ssessment will be based on the following points
1. Pres	sent/Absent
2. A c	ompletion date of the journal
3. Reg	ularity

#### Guidelines for Laboratory Conduction The following rules must be observed during laboratory conduction

- 1. Lab coat should be worn by students before entering the laboratory
- 2. Students shall keep their belongings on storage rack
- 3. Loose hair and flowing parts of apparel shall be properly tied before commencing of work
- 4. Enter the usage of chemicals and equipment's in a logbook
- 5. The instruction manual should be read before operating any instrument
- 6. Students should make aware of hazard warning symbols on reagent bottle
- 7. Protective devices must be worn when it is necessary to protect the eyes and face from splashes
- 8. All chemicals, glassware, reagents and plastic wares should be kept on their appropriate place after use 8. Reagents to be stored should be labeled with due discarding date
- 9. Instructions for proper disposal of waste material should be followed
- 10. Report accidental cuts or burns to the instructor immediately.
- 11. Perform the experiment. Collect data in a clear and organized fashion. Be sure to note the units for each measurement. Also, be sure to participate in the experiment rather than just recording data for your group

#### **General Guidelines:**

Before starting any experiment, clearly define the goals. What question are you answering or what principle are you trying to demonstrate? What data is needed to solve the problem? Identify the methods of measurement and instrumentation to be used.

Second Year of B.Tech. Biotechnology (2019 Course)         215475:Cell Biology and Tissue CultureLab         Credit         PR : 01         Reaching Scheme:         ExaminationSche         Two         Two         Of the Cell and its components.         Course Objectives:         1. To offer basic understanding of how eukaryotic cell functions         2. Role of lab and its equipments in helping understand the functioning and growth of the cells         3. To help understand the process of isolation and characterization of different types of cells.         4. To give hands on experience to the students on isolation, characterization, culturing, st culturing and preservation of animal cells.         Course Outcomes:         A. The students will understand the different types of microscopy techniques to observe to physiological characteristics of the cells.         C. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth         D. The students will be able to understand the media sterilization and cryopreservati techniques         auggested List of Laboratory Assignments (Any 8)         Title of the Practical         Introduction to Cell biology and tissue culture facility.       Microscope observation of adherent and suspension	Savitribai Phule I	Pune University, Pune
215475:Cell Biology and Tissue CultureLab         Credit         PR : 01         ExaminationSche         Two         Coredit         PR : 01         ExaminationSche         Two         Prerequisites: Basic Understanding of the Cell and its components.         Course Objectives:         1. To offer basic understanding of how eukaryotic cell functions         2. Role of lab and its equipments in helping understand the functioning and growth of the cells         3. To help understand the process of isolation and characterization of different types of cells.         4. To give hands on experience to the students on isolation, characterization, culturing, st culturing and preservation of animal cells.         Course Outcomes:         A. The students will understand the different types of microscopy techniques to observe to physiological characteristics of the cells.         B. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth         D. The students will be able to understand the media sterilization and cryopreservati techniques         augusted List of Laboratory Assignments (Any 8)         Title of the Practical         Introduction to Cell biology and tissue culture facility.       Micro	Second Year of B.Tech.	Biotechnology (2019 Course)
Credit         PR : 01         Creaching Scheme:       ExaminationSche         PR: 02hrs/week       TW         Ol       Tota         Prerequisites: Basic Understanding of the Cell and its components.       Tota         Course Objectives:       Image: Cell and its components.         2. Role of lab and its equipments in helping understand the functioning and growth of the cells       State         3. To help understand the process of isolation and characterization of different types of cells.       Image: Cell and preservation of animal cells.         Course Outcomes:       Image: Cell and the difference between various types of cells.       Image: Cell and will be able to use different types of microscopy techniques to observe to physiological characteristics of the cells.         C. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth       Image: Cell and will be able to understand the media sterilization and cryopreservati techniques         Maggested List of Laboratory Assignments (Any 8)       Image: Cell biology and tissue culture facility.         Introduction to Cell biology and tissue culture facility.       Image: Cell counting using hemocytometer: Animal cell count for passaging/WBC count.	215475:Cell Biology	y and Tissue CultureLab
PR : 01         Feaching Scheme:       ExaminationSche         PR: 02hrs/week       DOI         Tota       Tota         Prerequisites: Basic Understanding of the Cell and its components.       Tota         Course Objectives:       Tota         1. To offer basic understanding of how eukaryotic cell functions       Present the students of the process of isolation and characterization of different types of cells         3. To help understand the process of isolation and characterization, culturing, su culturing and preservation of animal cells.       Present types of cells.         B. The students will understand the difference between various types of cells.       B.         B. The students will understand the difference between various types of cells and will be able to use different types of microscopy techniques to observe the physiological characteristics of the cells.         C. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth         D. The students will be able to understand the media sterilization and cryopreservati techniques         suggested List of Laboratory Assignments (Any 8)         Tr. No.       Title of the Practical         O       Introduction to Cell biology and tissue culture facility.         Microscope observation of adherent and suspension culture and confluence using invertinicroscope         O       Cell counting using hemocytometer: Animal cell count for passagin	(	Credit
Catching Scheme:       ExaminationScheme         PR: 02hrs/week       ExaminationScheme         Prerequisites:       Basic Understanding of the Cell and its components.         Course Objectives:       To offer basic understanding of how eukaryotic cell functions         2.       Role of lab and its equipments in helping understand the functioning and growth of the cells         3.       To help understand the process of isolation and characterization of different types of cells         4.       To give hands on experience to the students on isolation, characterization, culturing, st culturing and preservation of animal cells.         Course Outcomes:       A.         A.       The students will be able to use different types of microscopy techniques to observe to physiological characteristics of the cells.         C.       The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth         D.       The students will be able to understand the media sterilization and cryopreservati techniques         Suggested List of Laboratory Assignments (Any 8)       Title of the Practical         I.       Introduction to Cell biology and tissue culture facility.         Microscope observation of adherent and suspension culture and confluence using invert microscope         C.       Cell counting using hemocytometer: Animal cell count for passaging/WBC count.		PR : 01
<b>R: 02hrs/week</b> O) <b>Prerequisites:</b> Basic Understanding of the Cell and its components. <b>Course Objectives:</b> 1. To offer basic understanding of how eukaryotic cell functions <b>2</b> .         2. Role of lab and its equipments in helping understand the functioning and growth of the cells <b>3</b> .         3. To help understand the process of isolation and characterization of different types of cells <b>4</b> .         4. To give hands on experience to the students on isolation, characterization, culturing, st culturing and preservation of animal cells. <b>Course Outcomes:</b> A. The students will be able to use different types of microscopy techniques to observe to physiological characteristics of the cells. <b>8</b> .         B. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth <b>D</b> .         D. The students will be able to understand the media sterilization and cryopreservati techniques <b>1</b> . <b>augested List of Laboratory Assignments (Any 8) 1</b> . <b>Title of the Practical 1</b> .         Introduction to Cell biology and tissue culture facility.       Microscope observation of adherent and suspension culture and confluence using inverting microscope         .       Cell counting using hemocytometer: Animal cell count for passaging/WBC count.	ching Scheme:	ExaminationScheme:
Tota         Prerequisites: Basic Understanding of the Cell and its components.         Course Objectives:         1. To offer basic understanding of how eukaryotic cell functions         2. Role of lab and its equipments in helping understand the functioning and growth of the cells         3. To help understand the process of isolation and characterization of different types of cells         4. To give hands on experience to the students on isolation, characterization, culturing, st culturing and preservation of animal cells.         Course Outcomes:         A. The students will understand the difference between various types of cells.         B. The students will be able to use different types of microscopy techniques to observe to physiological characteristics of the cells.         C. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth         D. The students will be able to understand the media sterilization and cryopreservati techniques         uggested List of Laboratory Assignments (Any 8)         rt No.       Title of the Practical         Introduction to Cell biology and tissue culture facility.         Microscope observation of adherent and suspension culture and confluence using invertinicroscope         .       Cell counting using hemocytometer: Animal cell count for passaging/WBC count.	02hrs/week	OR:50
<ul> <li>Prerequisites: Basic Understanding of the Cell and its components.</li> <li>Course Objectives:         <ol> <li>To offer basic understanding of how eukaryotic cell functions</li> <li>Role of lab and its equipments in helping understand the functioning and growth of the cells</li> <li>To help understand the process of isolation and characterization of different types of cells</li> <li>To give hands on experience to the students on isolation, characterization, culturing, st culturing and preservation of animal cells.</li> </ol> </li> <li>Course Outcomes:         <ol> <li>The students will understand the difference between various types of cells.</li> <li>The students will be able to use different types of microscopy techniques to observe a physiological characteristics of the cells.</li> <li>The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth</li> <li>The students will be able to understand the media sterilization and cryopreservati techniques</li> </ol> </li> <li>Introduction to Cell biology and tissue culture facility.         <ol> <li>Microscope observation of adherent and suspension culture and confluence using inverting microscope</li> <li>Cell counting using hemocytometer: Animal cell count for passaging/WBC count.</li> </ol> </li></ul>		Total :75
Course Objectives:         1. To offer basic understanding of how eukaryotic cell functions         2. Role of lab and its equipments in helping understand the functioning and growth of the cells         3. To help understand the process of isolation and characterization of different types of cells         4. To give hands on experience to the students on isolation, characterization, culturing, st culturing and preservation of animal cells.         Course Outcomes:         A. The students will understand the difference between various types of cells.         B. The students will be able to use different types of microscopy techniques to observe the physiological characteristics of the cells.         C. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth         D. The students will be able to understand the media sterilization and cryopreservati techniques         Augested List of Laboratory Assignments (Any 8)         ir. No.       Title of the Practical         Microscope observation of adherent and suspension culture and confluence using invertimicroscope         .       Cell counting using hemocytometer: Animal cell count for passaging/WBC count.	requisites: Basic Understanding of the Cell	and its components.
<ol> <li>To offer basic understanding of how eukaryotic cell functions</li> <li>Role of lab and its equipments in helping understand the functioning and growth of the cells</li> <li>To help understand the process of isolation and characterization of different types of cells</li> <li>To give hands on experience to the students on isolation, characterization, culturing, st culturing and preservation of animal cells.</li> <li>Course Outcomes:         <ul> <li>A. The students will understand the difference between various types of cells.</li> <li>B. The students will be able to use different types of microscopy techniques to observe to physiological characteristics of the cells.</li> <li>C. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth</li> <li>D. The students will be able to understand the media sterilization and cryopreservati techniques</li> <li>Suggested List of Laboratory Assignments (Any 8)</li> <li>introduction to Cell biology and tissue culture facility.</li> <li>Microscope observation of adherent and suspension culture and confluence using invertemicroscope</li> <li>Cell counting using hemocytometer: Animal cell count for passaging/WBC count.</li> </ul> </li> </ol>	ursa Objectives.	
<ol> <li>To offer basic understanding of how eukaryotic cell functions</li> <li>Role of lab and its equipments in helping understand the functioning and growth of the cells</li> <li>To help understand the process of isolation and characterization of different types of cells</li> <li>To give hands on experience to the students on isolation, characterization, culturing, su culturing and preservation of animal cells.</li> <li><b>Course Outcomes:</b></li> <li>A. The students will understand the difference between various types of cells.</li> <li>B. The students will be able to use different types of microscopy techniques to observe a physiological characteristics of the cells.</li> <li>C. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth</li> <li>D. The students will be able to understand the media sterilization and cryopreservati techniques</li> <li><b>uggested List of Laboratory Assignments (Any 8)</b></li> <li><b>ir. No.</b> Title of the Practical</li> <li>Introduction to Cell biology and tissue culture facility.</li> <li>Microscope observation of adherent and suspension culture and confluence using invertimicroscope</li> <li>Cell counting using hemocytometer: Animal cell count for passaging/WBC count.</li> </ol>		
<ol> <li>Role of lab and its equipments in helping understand the functioning and growth of the cells</li> <li>To help understand the process of isolation and characterization of different types of cells</li> <li>To give hands on experience to the students on isolation, characterization, culturing, su culturing and preservation of animal cells.</li> <li>Course Outcomes:         <ul> <li>A. The students will understand the difference between various types of cells.</li> <li>B. The students will be able to use different types of microscopy techniques to observe the physiological characteristics of the cells.</li> <li>C. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth</li> <li>D. The students will be able to understand the media sterilization and cryopreservation techniques</li> <li>Figure List of Laboratory Assignments (Any 8)</li> <li>introduction to Cell biology and tissue culture facility.</li> <li>Microscope observation of adherent and suspension culture and confluence using invertimicroscope</li> <li>Cell counting using hemocytometer: Animal cell count for passaging/WBC count.</li> </ul> </li> </ol>	To offer basic understanding of how eukary	votic cell functions
<ul> <li>3. To help understand the process of isolation and characterization of different types of cells</li> <li>4. To give hands on experience to the students on isolation, characterization, culturing, st culturing and preservation of animal cells.</li> <li>Course Outcomes:</li> <li>A. The students will understand the difference between various types of cells.</li> <li>B. The students will be able to use different types of microscopy techniques to observe the physiological characteristics of the cells.</li> <li>C. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth</li> <li>D. The students will be able to understand the media sterilization and cryopreservative techniques</li> <li>uggested List of Laboratory Assignments (Any 8)</li> <li>r. No. Title of the Practical</li> <li>Introduction to Cell biology and tissue culture facility.</li> <li>Microscope observation of adherent and suspension culture and confluence using inverted microscope</li> <li>Cell counting using hemocytometer: Animal cell count for passaging/WBC count.</li> </ul>	Role of lab and its equipments in helping un	nderstand the functioning and growth of the cells
<ul> <li>4. To give hands on experience to the students on isolation, characterization, culturing, st culturing and preservation of animal cells.</li> <li>Course Outcomes: <ul> <li>A. The students will understand the difference between various types of cells.</li> <li>B. The students will be able to use different types of microscopy techniques to observe the physiological characteristics of the cells.</li> <li>C. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth</li> <li>D. The students will be able to understand the media sterilization and cryopreservation techniques</li> </ul> </li> <li>Fuggested List of Laboratory Assignments (Any 8)</li> <li>Fr. No. Title of the Practical <ul> <li>Introduction to Cell biology and tissue culture facility.</li> <li>Microscope observation of adherent and suspension culture and confluence using inverted microscope</li> <li>Cell counting using hemocytometer: Animal cell count for passaging/WBC count.</li> </ul> </li> </ul>	To help understand the process of isolation	and characterization of different types of cells
culturing and preservation of animal cells. Course Outcomes: A. The students will understand the difference between various types of cells. B. The students will be able to use different types of microscopy techniques to observe a physiological characteristics of the cells. C. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth D. The students will be able to understand the media sterilization and cryopreservation techniques Cuggested List of Laboratory Assignments (Any 8) C. The of the Practical C. Introduction to Cell biology and tissue culture facility. C. Microscope observation of adherent and suspension culture and confluence using inverted microscope C. Cell counting using hemocytometer: Animal cell count for passaging/WBC count.	To give hands on experience to the stude	ents on isolation, characterization, culturing, sub-
<ul> <li>Course Outcomes:</li> <li>A. The students will understand the difference between various types of cells.</li> <li>B. The students will be able to use different types of microscopy techniques to observe a physiological characteristics of the cells.</li> <li>C. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth</li> <li>D. The students will be able to understand the media sterilization and cryopreservati techniques</li> <li>vuggested List of Laboratory Assignments (Any 8)</li> <li>ir. No.</li> <li>Title of the Practical</li> <li>Introduction to Cell biology and tissue culture facility.</li> <li>Microscope observation of adherent and suspension culture and confluence using invertemicroscope</li> <li>Cell counting using hemocytometer: Animal cell count for passaging/WBC count.</li> </ul>	culturing and preservation of animal cells.	
<ul> <li>A. The students will understand the difference between various types of cells.</li> <li>B. The students will be able to use different types of microscopy techniques to observe a physiological characteristics of the cells.</li> <li>C. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth</li> <li>D. The students will be able to understand the media sterilization and cryopreservati techniques</li> <li>buggested List of Laboratory Assignments (Any 8)</li> <li>F. No. Title of the Practical</li> <li>Introduction to Cell biology and tissue culture facility.</li> <li>Microscope observation of adherent and suspension culture and confluence using invertemicroscope</li> <li>Cell counting using hemocytometer: Animal cell count for passaging/WBC count.</li> </ul>	irse Outcomes:	
<ul> <li>B. The students will be able to use different types of microscopy techniques to observe a physiological characteristics of the cells.</li> <li>C. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth</li> <li>D. The students will be able to understand the media sterilization and cryopreservative techniques</li> <li>Suggested List of Laboratory Assignments (Any 8)</li> <li>Fr. No. Title of the Practical</li> <li>Introduction to Cell biology and tissue culture facility.</li> <li>Microscope observation of adherent and suspension culture and confluence using inverted microscope</li> <li>Cell counting using hemocytometer: Animal cell count for passaging/WBC count.</li> </ul>	The students will understand the difference	between various types of cells.
<ul> <li>physiological characteristics of the cells.</li> <li>C. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth</li> <li>D. The students will be able to understand the media sterilization and cryopreservati techniques</li> <li>Suggested List of Laboratory Assignments (Any 8)</li> <li>Ar. No. Title of the Practical</li> <li>Introduction to Cell biology and tissue culture facility.</li> <li>Microscope observation of adherent and suspension culture and confluence using inverted microscope</li> <li>Cell counting using hemocytometer: Animal cell count for passaging/WBC count.</li> </ul>	The students will be able to use differen	it types of microscopy techniques to observe the
<ul> <li>C. The students will be able to prepare the lab for the growth of cells and will be able characterize and understand the phenomena of growth</li> <li>D. The students will be able to understand the media sterilization and cryopreservative techniques</li> <li>Suggested List of Laboratory Assignments (Any 8)</li> <li>Fr. No. Title of the Practical</li> <li>Introduction to Cell biology and tissue culture facility.</li> <li>Microscope observation of adherent and suspension culture and confluence using inverted microscope</li> <li>Cell counting using hemocytometer: Animal cell count for passaging/WBC count.</li> </ul>	physiological characteristics of the cells.	
characterize and understand the phenomena of growth         D. The students will be able to understand the media sterilization and cryopreservative techniques <b>buggested List of Laboratory Assignments (Any 8) fr. No. Title of the Practical</b> .       Introduction to Cell biology and tissue culture facility.         .       Microscope observation of adherent and suspension culture and confluence using inverted microscope         .       Cell counting using hemocytometer: Animal cell count for passaging/WBC count.	The students will be able to prepare the	lab for the growth of cells and will be able to
<ul> <li>D. The students will be able to understand the media sterilization and cryopreservative techniques</li> <li>Suggested List of Laboratory Assignments (Any 8)</li> <li>Sr. No. Title of the Practical</li> <li>Introduction to Cell biology and tissue culture facility.</li> <li>Microscope observation of adherent and suspension culture and confluence using inverted microscope</li> <li>Cell counting using hemocytometer: Animal cell count for passaging/WBC count.</li> </ul>	characterize and understand the phenomena	a of growth
techniques         Suggested List of Laboratory Assignments (Any 8)         Sr. No.       Title of the Practical         .       Introduction to Cell biology and tissue culture facility.         .       Microscope observation of adherent and suspension culture and confluence using inverted microscope         .       Cell counting using hemocytometer: Animal cell count for passaging/WBC count.	The students will be able to understan	nd the media sterilization and cryopreservation
Suggested List of Laboratory Assignments (Any 8)         Sr. No.       Title of the Practical         .       Introduction to Cell biology and tissue culture facility.         .       Microscope observation of adherent and suspension culture and confluence using inverted microscope         .       Cell counting using hemocytometer: Animal cell count for passaging/WBC count.	techniques	
Sr. No.       Title of the Practical         .       Introduction to Cell biology and tissue culture facility.         .       Microscope observation of adherent and suspension culture and confluence using inverter microscope         .       Cell counting using hemocytometer: Animal cell count for passaging/WBC count.	gested List of Laboratory Assignments (A	Any 8)
Introduction to Cell biology and tissue culture facility.         Microscope observation of adherent and suspension culture and confluence using invertance microscope         Cell counting using hemocytometer: Animal cell count for passaging/WBC count.	No.   Title of the Practical	
<ul> <li>Microscope observation of adherent and suspension culture and confluence using invertant microscope</li> <li>Cell counting using hemocytometer: Animal cell count for passaging/WBC count.</li> </ul>	Introduction to Cell biology and tissu	e culture facility.
. Cell counting using hemocytometer: Animal cell count for passaging/WBC count.	Microscope observation of adherent a microscope	and suspension culture and confluence using inverted
	Cell counting using hemocytometer: A	Animal cell count for passaging/WBC count.
. WBC Different cell types stain and visualize	WBC Different cell types stain and vi	isualize

5.	Onion Root Tip nuclear staining and observation of stages of Mitosis
6.	Filter sterilization of media for animal cell culture
7.	Passaging of adherent animal cell cultures.
8.	Cryopreservation of animal cells.
9.	Revival of animal cells
10.	Viability staining
	Guidelines for Instructor's Manual
1.	Discuss the syllabus for the course
2.	Go through the general procedures for lab safety
3.	Review the guidelines for laboratory reports
4.	Practice some of the scientific calculations that will be used throughout the semester
5.	Practice some basic laboratory methods such as the use of balances, pipets, and Micropipettes
6.	Students should be briefed with Risk Assessment and Biosafety Levels
7.	All the instruments to be validated before use
8.	All the experiments should be standardized
9.	The instructor is responsible for seeing that the consequences of student are rectified, including
	correction of damages and violations and take-down of experiments
Guide	lines for Student's Lab Journal
Please	read these instructions now and refer to them regularly!
These 1.	instructions must be followed carefully. Use a bound notebook for notes and lab observations.
2.	Lab notebooks should be done in pen and no erasing or white-out is allowed
3.	For the Journal use Journal Pages and ensure you draw neat diagrams wherever necessary
4.	Number the pages
5.	Title and underline each lab exercise at the top of the page and date it.
5. 6.	Title and underline each lab exercise at the top of the page and date it. Each lab write-up should be done separately even if more than one exercise is performed in a
5. 6.	Title and underline each lab exercise at the top of the page and date it. Each lab write-up should be done separately even if more than one exercise is performed in a lab period.
5. 6. 7.	Title and underline each lab exercise at the top of the page and date it. Each lab write-up should be done separately even if more than one exercise is performed in a lab period. Briefly explain the lab exercise objectives in a few sentences.
5. 6. 7. 8.	Title and underline each lab exercise at the top of the page and date it. Each lab write-up should be done separately even if more than one exercise is performed in a lab period. Briefly explain the lab exercise objectives in a few sentences. Record observations, diagrams and results from the exercise.

10. Write neatly, be organized and follow a standard format.

Note: The purpose of the lab notebook is to encourage students to compile and organize their laboratory notes and to understand the purpose of the laboratory exercises and the meaning of their results.

#### Guidelines for Lab /TW Assessment

Lab assessment will be based on following points

- 1. Present / Absent
- 2. Completion date of journal
- 3. Regularity
- 4. Understanding
- 5. Presentation

#### **Guidelines for Laboratory Conduction**

Basic Principles for Students Working in Cell Biology Laboratories : The following rules must be observed at all times to prevent accidental injury to and infection of yourself and others and to minimize contamination of the lab environment:

- 1. Never place books, backpacks, purses, etc., on bench tops. Always place these in the assigned cubicles. Keep manuals and pens on pull-out desks.
- 2. Clean your work area with dilute bleach solution at the beginning AND end of each lab.
- 3. Wash your hands with soap and dry with paper towels when entering and leaving the lab.
- 4. Wear a lab coat at all times while working in the lab to prevent contamination or accidental staining of your clothing.
- 5. Closed-toe shoes (no sandals) are to be worn in the lab.
- 6. Long hair must be tied back to prevent exposure to flame and contamination of cultures.
- 7. Do not place anything in your mouth or eyes while in the lab. This includes pencils, food, and fingers. Keep your hands away from your mouth and eyes.
- 8. Eating and drinking are prohibited in the lab at all times. This includes gum, cough drops, and candy.
- 9. Never pipet by mouth. Use a mechanical pipetting device.
- 10. Do not remove media, equipment, or bacterial cultures from the laboratory. This is absolutely prohibited and unnecessary.
- 11. Do not place contaminated instruments such as inoculating loops, needles, and pipettes on bench tops. Loops and needles should be sterilized by incineration, and pipettes should be

disposed of in designated receptacles of bleach solution.

- 12. Carry cultures in a test tube rack when moving around the lab or when keeping cultures on bench tops for use. This prevents accidents and contamination of your person or belongings.
- 13. Immediately covers spilled cultures or broken culture tubes with paper towels and then saturate them with disinfectant solution. Notify your instructor that there has been a spill. After 15 minutes, dispose of the towels and broken items as indicated by your instructor.
- 14. Report accidental cuts or burns to the instructor immediately.
- 15. At the end of each lab session, place all cultures and materials in the proper disposal area.
- 16. Electronic devices should not be brought into the lab. This includes, but is not limited to iPods, MP3 players, radios, cell phones, and calculators.

#### **General Guidelines:**

Before starting any experiment, clearly define the goals. What question are you answering or what principle are you trying to demonstrate? What data is needed to solve the problem? Identify the methods of measurement and instrumentation to be used.

Teaching Scheme:         PR:04hrs/week         Course Objectives:         1. To understand the significance biology         2. To isolate and purify the DNA         3. To estimate the quantitative pu         4. To understand the process of acid.         Course Outcomes:         On completion of the course, learner w         A. Graduates are learn the princip molecular biology	Credits         PR:02         Examination Scheme:         TW:25         OR:50         Total: 75         ce of different stocks and reagents used in molecular         from prokaryotic and eukaryotic cells.         arity of isolated DNA.         gel electrophoresis for qualitative analysis of nucleic
Teaching Scheme: PR:04hrs/week         Course Objectives:         1. To understand the significance biology         2. To isolate and purify the DNA         3. To estimate the quantitative pu         4. To understand the process of acid.         Course Outcomes:         On completion of the course, learner w         A. Graduates are learn the princip molecular biology	PR:02 Examination Scheme: TW:25 OR:50 Total: 75 ce of different stocks and reagents used in molecular from prokaryotic and eukaryotic cells. rity of isolated DNA. gel electrophoresis for qualitative analysis of nucleic
<ul> <li>Teaching Scheme: PR:04hrs/week</li> <li>Course Objectives: <ol> <li>To understand the significance biology</li> <li>To isolate and purify the DNA</li> <li>To estimate the quantitative put</li> <li>To understand the process of acid.</li> </ol> </li> <li>Course Outcomes: On completion of the course, learner we A. Graduates are learn the princip molecular biology</li></ul>	Examination Scheme:         TW:25         OR:50         Total: 75         ce of different stocks and reagents used in molecular         from prokaryotic and eukaryotic cells.         urity of isolated DNA.         gel electrophoresis for qualitative analysis of nucleic
<ul> <li>PR:04hrs/week</li> <li>Course Objectives: <ol> <li>To understand the significance biology</li> <li>To isolate and purify the DNA</li> <li>To estimate the quantitative put</li> <li>To understand the process of acid.</li> </ol> </li> <li>Course Outcomes: On completion of the course, learner w A. Graduates are learn the princip molecular biology</li></ul>	TW:25 OR:50 Total: 75 ce of different stocks and reagents used in molecular from prokaryotic and eukaryotic cells. nrity of isolated DNA. gel electrophoresis for qualitative analysis of nucleic
<ul> <li>Course Objectives: <ol> <li>To understand the significance biology</li> <li>To isolate and purify the DNA</li> <li>To estimate the quantitative puther of the process of acid.</li> </ol> </li> <li>Course Outcomes: On completion of the course, learner were and the process of acid are learn the principe molecular biology</li></ul>	OR:50 Total: 75 ce of different stocks and reagents used in molecular from prokaryotic and eukaryotic cells. writy of isolated DNA. gel electrophoresis for qualitative analysis of nucleic
<ul> <li>Course Objectives: <ol> <li>To understand the significance biology</li> <li>To isolate and purify the DNA</li> <li>To estimate the quantitative pu</li> <li>To understand the process of acid.</li> </ol> </li> <li>Course Outcomes: On completion of the course, learner w A. Graduates are learn the princip molecular biology</li></ul>	Total: 75 ce of different stocks and reagents used in molecular from prokaryotic and eukaryotic cells. urity of isolated DNA. gel electrophoresis for qualitative analysis of nucleic
<ul> <li>Course Objectives: <ol> <li>To understand the significance</li> <li>biology</li> <li>To isolate and purify the DNA</li> <li>To estimate the quantitative pu</li> <li>To understand the process of acid.</li> </ol> </li> <li>Course Outcomes: On completion of the course, learner w A. Graduates are learn the princip molecular biology</li></ul>	ce of different stocks and reagents used in molecular from prokaryotic and eukaryotic cells. writy of isolated DNA. gel electrophoresis for qualitative analysis of nucleic
<ol> <li>To understand the significanc biology</li> <li>To isolate and purify the DNA</li> <li>To estimate the quantitative pu</li> <li>To understand the process of acid.</li> </ol> Course Outcomes: On completion of the course, learner w A. Graduates are learn the princip molecular biology	ce of different stocks and reagents used in molecular from prokaryotic and eukaryotic cells. writy of isolated DNA. gel electrophoresis for qualitative analysis of nucleic
<ul> <li>biology</li> <li>2. To isolate and purify the DNA</li> <li>3. To estimate the quantitative pu</li> <li>4. To understand the process of acid.</li> </ul> Course Outcomes: On completion of the course, learner w A. Graduates are learn the princip molecular biology	from prokaryotic and eukaryotic cells. writy of isolated DNA. gel electrophoresis for qualitative analysis of nucleic
<ol> <li>To isolate and purify the DNA</li> <li>To estimate the quantitative pu</li> <li>To understand the process of acid.</li> </ol> Course Outcomes: On completion of the course, learner w A. Graduates are learn the princip molecular biology	from prokaryotic and eukaryotic cells. arity of isolated DNA. gel electrophoresis for qualitative analysis of nucleic
<ul> <li>3. To estimate the quantitative pu</li> <li>4. To understand the process of acid.</li> </ul> Course Outcomes: On completion of the course, learner w A. Graduates are learn the princip molecular biology	urity of isolated DNA. E gel electrophoresis for qualitative analysis of nucleic
<ul> <li>4. To understand the process of acid.</li> <li>Course Outcomes:</li> <li>On completion of the course, learner w</li> <li>A. Graduates are learn the princip molecular biology</li> </ul>	gel electrophoresis for qualitative analysis of nucleic
acid. Course Outcomes: On completion of the course, learner w A. Graduates are learn the princip molecular biology	
Course Outcomes: On completion of the course, learner w A. Graduates are learn the princip molecular biology	
	will be able to- ple of various buffer system and reagents required in
B. Graduates are skilled to isolat organisms.	te the nucleic acids from prokaryotic and eukaryotic
C. Graduates are trained to use	agarose gel electrophoresis system for estimation of
nucleic acids	
D. Graduates are trained to examine	he the DNA by qualitative and quantitative methods
Suggested List of Laboratory Assign	nments
1. Preparation of Stock & Workin	ng solutions-Tris-Cl, EDTA, Extraction Buffers, TE
2. Isolation of plant genomic DNA	A
3. Isolation of E.coli genomic DN	NA
4. Quantification and purity che	ask of antroated DNA from Plant & E coli by a

- 4. Quantification and purity check of extracted DNA from Plant & E. coli by at 260/280,260/230 nm Spectrophotometer
  - 5. Agarose Gel Electrophoresis
  - 6. Study of effect of agarose concentration on electrophoretic mobility of DNA

- 7. Visualization of plant genomic DNA by agarose gel electrophoresis
- 8. Visualization of E. coli genomic DNA by agarose gel electrophoresis

#### **Guidelines for Instructor's Manual**

Students should be briefed with risk assessment and Biosafety levels

- All the instruments to be validated before use
- All the experiments should be standardized

• The instructor is responsible for seeing that consequences of student are rectified, including correction of damages and violations and take-down of experiments

#### **Guidelines for Student's Lab Journal**

- Use provided templates of experiment write ups
- Follow the sequence of experiments as per the index, while arranging journal file
- Draw necessary diagrams with pencil and fill other fields like observations, calculations, conclusion etc. with Pen
- Paste Images e.g. of specialized equipment, Gel pictures, isolated DNA wherever necessary
- Avoid overwriting and copying of results, conclusions etc.

#### Guidelines for Lab /TW Assessment

- Each experiment will be assessed based on following terms.
- Student should attend each practical in scheduled batch to gain full marks for that practical
- Regularity will be assessed throughout the semester for practical.
- Presentation of students in laboratory during practical will be assessed.
- Understanding and application of steps involved in practical to achieve good results will contribute in term work/lab assessment.

• For final term work assessment along with above all points, unit test marks, theory lecture attendance will also be considered

Savitribai Phule P	une University, Pune		
Second Year of B.Tech. Biotechnology (2019 Course) 215477:Project based Learning Credits			
			PR: 02
		Teaching Scheme:	Examination Scheme:
PR : 04 hrs/week	TW:50		
	Total:50		
Projectbasedlearning(PBL)requirescontinuous	mentoringbyfacultythroughoutthesemester for		
successful completion of the tasks selected l	by the students per batch. While assigning the		
teaching workload a load of 2 Hrs/week/batch	needs to be considered for the faculty involved.		
The Batch needs to be divided into sub-group	ps of 5 to 6 students. Assignments / activities /		
models/projectsetc.underprojectbasedlearning	iscarriedthroughoutsemesterandCreditfor		
PBLhastobeawardedonthebasisofinternalcontin	nuousassessmentandevaluationattheend		
ofsemester.			
	$\sim$		
Few examples of problem based learning are a	as follows		
1. Evaluation of Protein Content among Sprou	ited and Un-SproutedSeeds		
2. To check pH of different watersamples			
3. To estimate carbohydrate content in differen	nt foodsamples		
4. Diversity of microorganisms of varioussam	ples.		

#### 215478:Mandatory Audit Course 4

In addition to credits courses, it is recommended that there should be audit course (non-credit course). Audit course is for the purpose of self-enrichment and academic exploration. Audit course carry no academic credit. Selection of audit courses helps the learner to explore the subject of interest in greater details resulting in achieving objective of audit course's inclusion. Evaluation of audit course will be done at institute level. Method of conduction and method of assessment for audit courses is suggested.

#### Criteria:

The student registered for audit course shall be awarded the grade PP and shall be included such grade in the semester grade report for that course, provided students has the minimum attendance as prescribed by the Savitribai Phule Pune university and satisfactory in-semester performance and secured a passing grade in that audit course. No grade point is associated with this "PP" grade and performance in these courses is not accounted in the calculation of the performance indices SGPA and CGPA.

#### Guidelines for Conduction and Assessment (Any one or more of following but not limited to)

- 1. Lecture/Guestlecture
- 2. Visit (Social/field) and reports
- 3. Demonstrations
- 4. Surveys
- 5. Mini project
- 6. Hands-on experience on a specific focusedtopic.
- 7. Seminar/Workshop

#### Guidelines for Assessment (Any one or more of following but not limited to)

- 1. Writtentest
- 2. Quiz
- 3. Demonstrations/practicaltest
- 4. Presentations
- 5. IPR/publication
- 6. Report

#### Audit course 2 Options (Anyone)

- 1. Professional Ethics and Etiquettes
- 2. Entrepreneurship Development inBiotechnology