

मराठवाडा विद्यापीठ, नांदेड स्वामी रामानंद ती

'ज्ञानतीर्थ', विष्णुपुरी, नांदेड - ४३१ ६०६ (महाराष्ट्र राज्य) भारत

SWAMI RAMANAND TEERTH MARATHWADA UNIVERSITY, NANDED

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विज्ञान व तंत्रज्ञान विद्याशाखे अंतर्गत राष्ट्रीय शैक्षणिक धोरण २०२० नुसार पदव्यूत्तर द्वितीय वर्षाचे अभ्यासकम (Syllabus) शैक्षणिक वर्ष २०२४-२५ पासून लागू करण्याबाबत.

सहा.कुलसचिव

शैक्षणिक (१-अभ्यासमंडळ) विभाग

परिपत्रक

या परिपत्रकान्वये सर्व संबंधितांना कळविण्यात येते की, या विद्यापीठा अंतर्गत येणा-या सर्व संलग्नित महाविद्यालयामध्ये शैक्षणिक वर्ष २०२४-२५ पासून राष्ट्रीय शैक्षणिक धोरणानुसार पदव्यूत्तर द्वितीय वर्षाचे अभ्यासकम लागू करण्याच्या दृष्टीकोनातून विज्ञान व तंत्रज्ञान विद्याशाखे अंतर्गत येणा—या अभ्यासमंडळांनी तयार केलेल्या पटव्यूत्तर द्वितीय वर्षाच्या अभ्यासक्रमांना मा. विद्यापरिषदेने दिनांक १५ मे २०२४ रोजी संपन्न झालेल्या बैठकीतील विषय कमांक १५/५९-२०२४ च्या ठरावाअन्वये मान्यता प्रदान केली आहे. त्यानुसार विज्ञान व तंत्रज्ञान विद्याशाखेतील खालील एम. एस्सी द्वितीय वर्षाचे अभ्यासक्रम (Syllabus) लागू करण्यात येत आहेत.

- 1) M. Sc. II year Biotechnology (Affiliated College)
- 2) M. Sc. II year Biotechnology (Campus)
- 3) M. Sc. II year Bioinformatics (Sub Campus Latur)
- 4) M. Sc. II year Bioinformatics (Affiliated College)
- 5) M. Sc. II year Clinical Research (Affiliated College)
- 6) M. Sc. II year Botany (Campus)
- 7) M. Sc. II year Herbal Medicine
- 8) M. Sc. II year Boany (Affiliated College)
- M. Sc. II year Geology (Campus)
- 10) M. Sc. II year Dairy Science
- 11) M. Sc. II year Electronics
- 12) M. Sc. II year Environmental Science
- 13) M. Sc. II year Environmental Science (Campus)
- 14) M. Sc. II year Geography (Campus)
- M. Sc. II year Applied Mathematics
- M. Sc. II year Mathematics
- 17) M. Sc. II year Mathematics (Campus)
- 18) M. Sc. II year Microbiology
- M. Sc. II year Microbiology (Campus)
- 20) M. Sc. II year Statistics
- 21) M. Sc. II year Statistics (Campus)

सदरील परिपत्रक व अभ्यासक्रम प्रस्तुत विद्यापीठाच्या www.srtmun.ac.in या संकेतस्थळावर उपलब्ध आहेत. तरी सदरील बाब ही सर्व संबंधितांच्या निदर्शनास आणून द्यावी, ही विनंती.

'ज्ञानतीर्थ' परिसर,

विष्णुप्री, नांदेड - ४३१ ६०६.

जा.क्र.:शै-१/एनइपी/विवत्रंविपदवी/२०२४-२५/९० €

दिनांक १२.०६.२०२४

प्रत : १) मा. आधिष्ठाता, विज्ञान व तंत्रज्ञान विद्याशाखा, प्रस्तुत विद्यापीठ.

- २) मा. संचालक, परीक्षा व मुंल्यमापन मंडळ, प्रस्तुत विद्यापीठ.
- ३) मा. प्राचार्य, सर्व संबंधित संलग्नित महाविद्यालये, प्रस्तृत विद्यापीठ.
- ४) मा. संचालक, सर्व संकुले परिसर व उपपरिसर, प्रस्तुत विद्यापीठ
- ५) सिस्टीम एक्सपर्ट, शैक्षणिक विभाग, प्रस्तृत विद्यापीठ. याना देवून कळविण्यात येते की, सदर परिपत्रक संकेतस्थळावर प्रसिध्द करण्यात यावे.



SWAMI RAMANAND TEERTH MARATHWADA UNIVERSITY, NANDED

STRUCTURE AND SYLLABUS OF TWO-YEAR MASTERS PROGRAM IN SCIENCE

UNDER

NATIONAL EDUCATION POLICY (NEP 2020)

In

SUBJECT: BIOTECHNOLOGY

FACULTY OF SCIENCE AND TECHNOLOGY

M. Sc. Second Year

SCHOOL OF LIFE SCIENCES

SWAMI RAMANAND TEERTH MARATHWADA UNIVERSITY, NANDED

With Effect From June 2024

From the Desk of the Dean, Faculty of Science and Technology:

Swami Ramanand Teerth Marathwada University, Nanded, enduring to its vision statement "Enlightened Student: A Source of Immense Power", is trying hard consistently to enrich the quality of science education in its jurisdiction by implementing several quality initiatives. Revision and updating curriculum to meet the standard of the courses at national and international level, implementing innovative methods of teaching-learning, improvisation in the examination and evaluation processes are some of the important measures that enabled the University to achieve the 3Es, the equity, the efficiency and the excellence in higher education of this region. To overcome the difficulty of comparing the performances of the graduating students and also to provide mobility to them to join other institutions the University has adopted the cumulative grade point average (CGPA) system in the year 2014-2015. Further, following the suggestions by the UGC and looking at the better employability, entrepreneurship possibilities and to enhance the latent skills of the stakeholders the University has adopted the Choice Based Credit System (CBCS) in the year 2018-2019 at graduate and post-graduate level. This provided flexibility to the students to choose courses of their own interests. To encourage the students to opt the world-class courses offered on the online platforms like, NPTEL, SWAYM, and other MOOCS platforms the University has implemented the credit transfer policy approved by its Academic Council and also has made a provision of reimbursing registration fees of the successful students completing such courses.

SRTM University has been producing a good number of high calibre graduates; however, it is necessary to ensure that our aspiring students are able to pursue the right education. Like the engineering students, the youngsters pursuing science education need to be equipped and trained as per the requirements of the R&D institutes and industries. This would become possible only when the students undergo studies with an updated and evolving curriculum to match global scenario.

Higher education is a dynamic process and in the present era the stakeholders need to be educated and trained in view of the self-employment and self-sustaining skills like start-ups. Revision of the curriculum alone is not the measure for bringing reforms in the higher education, but invite several other initiatives. Establishing industry-institute linkages and initiating internship, on job training for the graduates in reputed industries are some of the important steps that the University would like to take in the coming time. As a result, revision of the curriculum was the need of the hour and such an opportunity was provided by the New Education Policy 2020. National Education Policy 2020 (NEP 2020) aims at equipping students with knowledge, skills, values, leadership qualities and initiates them for lifelong learning. As a result the students will acquire expertise in specialized areas

of interest, kindle their intellectual curiosity and scientific temper, and create imaginative individuals.

The curriculum given in this document has been developed following the guidelines of NEP-2020 and is crucial as well as challenging due to the reason that it is a transition from general science based to the discipline-specific-based curriculum. All the recommendations of the *Sukanu Samiti* given in the NEP Curriculum Framework-2023 have been followed, keeping the disciplinary approach with rigor and depth, appropriate to the comprehension level of learners. All the Board of Studies (BoS) under the Faculty of Science and Technology of this university have put in their tremendous efforts in making this curriculum of international standard. They have taken care of maintaining logical sequencing of the subject matter with proper placement of concepts with their linkages for better understanding of the students. We take this opportunity to congratulate the Chairman(s) and all the members of various Boards of Studies for their immense contributions in preparing the revised curriculum for the benefits of the stakeholders in line with the guidelines of the Government of Maharashtra regarding NEP-2020. We also acknowledge the suggestions and contributions of the academic and industry experts of various disciplines.

We are sure that the adoption of the revised curriculum will be advantageous for the students to enhance their skills and employability. Introduction of the mandatory *On Job Training, Internship program* for science background students is praise worthy and certainly help the students to imbibe first hand work experience, team work management. These initiatives will also help the students to inculcate the workmanship spirit and explore the possibilities of setting up of their own enterprises.

Dr. M. K. Patil
Dean

Faculty of Science and Technology

Preamble:

The National Education Policy 2020 (NEP 2020) is formulated to revamp education system and lay down road map for new India. This policy is framed based on the fundamental pillars of access, equity, quality, affordability, and accountability and seeks to transform India into a thriving knowledge society and a global knowledge superpower.

Some of the important features of National Education Policy are increasing gross enrolment ratio in higher education, holistic and multidisciplinary education with multiple entry/exit options, establishment of academic bank of credit, setting up of multidisciplinary education and research Universities and National Research Foundation, expansion of open and distance learning to increase gross enrolment ratio, internationalization of education, motivated / energized and capable faculty, online and digital education and effective governance and leadership.

As per the National Education Policy, the Government of Maharashtra has proposed a model curriculum framework and an implementation plan for the State of Maharashtra. It is to suggest and facilitate the implementation of schemes and programs, which improve not only the level of academic excellence but also improve the academic and research environment in the state. The proposed curriculum framework endeavors to empower the students and help them in their pursuit for achieving overall excellence.

In view of NEP priority and in-keeping with its vision and mission, process of updating the curriculum is initiated and implemented in SRTM University at UG and PG level from the academic year 2023-2024.

Biotechnology is often considered as the technology of hope for meeting future challenges like feeding our increasing population, cleaning dangerously polluted environments and potentiating healthcare sector etc. Establishment of new IISERs, Central Universities and IITs indicate that we are already on the track of developing infrastructure and human resource. Our dream of becoming future 'superpower' will not be possible without Biotechnology and inclusive efforts. Therefore, it is necessary to attract young and bright students and train them in the field of Biotechnology.

Keeping in mind, BOS in Biotechnology and Bioinformatics prepared the curriculum to ensure up-to-date level of understanding of Biotechnology. Studying Biotechnology prepares the students for their career working either in educational institutions or industries in which they can be directly involved in the teaching, research and development. Also, to ensure uniform curriculum and its quality at UG/PG level, curriculum of different Indian Universities, syllabus of NET, SET, MPSC, UPSC and the UGC model curriculum are referred to serve as a base in updating the same.

The comments or suggestions from all teachers, students and other stakeholders are welcome for upbringing this curriculum.

Salient Features:

The syllabus of M Sc Biotechnology has been framed to meet the requirement of Choice Based Credit System under NEP 2020. The courses offered here in will train and orient the students in the specific fields of Biotechnology.

The Core Courses deal with Biochemistry, Cell and Developmental Biology, Immunology and Virology, Genetics and Molecular Biology, Bioanalytical Techniques, Bioprocess Engineering and Technology, r-DNA Technology, Bioinformatics, Pharmaceutical Biotechnology, Plant Biotechnology and Genomics and Proteomics.

Apart from the core courses, the Department Specific Elective Courses deal with Microbial and Enzyme Technology, Environmental Biotechnology, Diagnostic Biology, Animal Biotechnology, Nanobiotechnology, Techniques in Microbiology and Food Biotechnology. These courses offered during this program are designed with the aim of imparting specific skills to the students which will lead to the employability of the students. There are also two Research Projects in third and fourth semester respectively.

This would help students to lay a strong foundation in the field of Biotechnology.

Overall, after completion of this program, students will also acquire fundamental knowledge of applications of Biotechnology.

Program Educational Objectives:

The Objectives of this program are:

PEO1: To offer postgraduate program in Biotechnology based on the needs of industries, academic and research institutions worldwide.

PEO2: To promote and popularize Biotechnology at grass root level and attract young and budding talents.

PEO3: To expose the students to the different emerging fields of Biotechnology.

PEO4: To update curriculum by introducing recent advances in the subject that enable the students to face NET, SET, MPSC, UPSC and other competitive examinations successfully.

PEO5: To train and orient the students so as to develop human resource for the educational institutes and other organizations.

PEO6: To inculcate analytical and application-oriented abilities to create active and frontline researchers and human resource for the industries.

PEO7: To develop specific skills amongst students for self-employability and also for the development of their own enterprises.

Program Outcomes:

The Outcomes of this program are:

PO1: This Biotechnology program shall promote and popularize Biotechnology at grass root level and attract young and budding talents.

PO2: This program will expose the students to the different emerging fields of Biotechnology.

PO3: This will provide updated curriculum with recent advances in the subject that enable the students to face NET, SET, MPSC, UPSC and other competitive examinations successfully.

PO4: This program shall train and orient the students so as to develop human resource for the educational institutes and other organizations.

PO5: This program shall train and orient the students so as to develop active and frontline researchers and human resource for the industries.

PO6: This will also develop specific skills amongst students for self-employability and also for the development of their own enterprises.

Prerequisite:

Basic knowledge of Science at B.Sc. level. The optional courses of this program are offered to the students registered for post-graduate programs. Such students should have the basic knowledge of Biotechnology and willing to gain additional knowledge in the field of Biotechnology.

The students seeking admission to this program should have cleared any B Sc or B Pharm or B Sc Agri from any statutory University.

Dr. Sunita D. Lohare

Chairman, BOS in Biotechnology and Bioinformatics Swami Ramanand Teerth Marathwada University, Nanded 431606.

Details of the Board of Studies Members in the subject Biotechnology and Bioinformatics under the Faculty of Science & Technology, S.R.T.M. University, Nanded.

Sr No	Name of the Member	Designation	Sr No	Name of the Member	Designation
1	Dr Sunita Dhundiraj Lohare Shri Havgiswami Mahavidyalaya, Udgir, Dist. Latur Mob 9284161504	Chairman	2	Dr Babasaheb S Surwase School of Life Sciences SRTM, University, Nanded 431606. Mob 9075829767	Member
3	Dr Pratap V. Deshmukh Nagnath Arts, Commerce and Science College, Aundha Nagnath, Dist. Hingoli Mob 9637202024	Member	4	Dr Komal S. Gomare Dept of Biotechnology, Dayanand Science College, Latur Mob 9284238413	Member
5	Dr Vaibhav D. Deshpande General Manager, Quality Corporate Office, Wockhardt, Mumbai Mob 9100988260	Member			
		Invit	ee Memb	pers	
6	Dr Laxmikant Kamble School of Life Sciences, SRTM University, Nanded 431606. Mob 8669695555	Member	7	Dr M M V Baig Dept of Biotechnology, Yeshwant Mahavidyalaya, Nanded. Mob 9422170641	Member
8	Dr Arun Ingale School of Life Sciences, North Maharashtra University, Umavinagar, Jalgaon Mob 9822708707	Member	9	Dr Prashant Thakare Department of Biotechnology, SGB Amravati University, Amravati. Mob 9822222822	Member
10	Dr A B Gulwe School of Technology, SRTM University Sub Campus, Latur. Mob 7387120874	Member	11	Dr Sanjog T. Thul Environmental Biotechnology and Genomics Division, National Environmental and Engineering Research Institute (CSIR-NEERI). Nagpur. Mob 9881877072	Member
12	Dr Shivraj Hariram Nile Department of Food Science and Agriculture, National Agri- Food Biotechnology Institute (NABI), Mohali, Punjab Mob 9561740707	Member	13	Dr Sunil Hajare Department of Biotechnology, New Model Degree College, Hingoli . Mob 8378878817	Member



Swami Ramanand Teerth Marathwada University, Nanded-431606

Faculty of Science & Technology

Credit Framework and Structure of Two Year PG Program (NEP 2020)
Subject: M Sc Biotechnology (Campus School of Life Sciences) (R-2023)

Year & Level	Sem	Major Subject	t	VEC	OJT / FP/CS	Research Project	Practicals (1-Cr)	Credits	Total Credits
Levei		(DSC-4 Cr)	(DSE-3 Cr)		(3-Cr)	rroject	(I-Cr)		Credit
1	1	SBTTC-401 Biochemistry SBTTC-402 Cell and Developmental Biology SBTTC-403 Immunology and Virology	SBTTE-401 Biostatistics and Basic Computer OR SBTTE-403 Techniques in Microbiology	SVECR-401 Research Methodology (3-Cr)			SBTTP-401 Lab Course in Biochemistry SBTTP-402 Lab Course in Cell and Developmental Biology SBTTP-403 Lab Course in Immunology and Virology SBTTE-402 Lab Course in Biostatistics and Basic Computer OR SBTTE-404 Lab Course in Techniques in Microbiology	22	44
	2 SBTTC-451 Genetics and Molecular Biology OR SBTTC-452 Bioanalytical Techniques SBTTE-453 Diagnostic Biology SBTTC-453 Bioprocess Engineering and Technology OR SBTTC-453 Bioprocess Engineering and Technology OR SBTTC-454 Lab Course in Bioprocess Engineering and Technology SBTTC-455 Lab Course in Environmental Biotechnology SBTTC-456 Lab Course in Bioprocess Engineering and Technology SBTTC-456 Lab Course in Environmental Biotechnology SBTTC-457 Lab Course in Environmental Biotechnology SBTTC-458 Lab Course in Environmental Biotechnology OR SBTTC-454 Lab Course in Diagnostic Biology OR SBTTC-454 Lab Course in Diagnostic Biology						22		
		Exi	t option: Exit Option with PG I	Diploma in Basic I	Biotechnology	(After 2024-2	5)		
SE		SBTTC-501 r DNA Technology SBTTC-502 Bioinformatics SBTTC-503 Pharmaceutical Biotechnology	Biotechnology OR SBTTE-503 Microbial and Enzyme Technology Ology			Research Project SBTTR- 501 (4-Cr)	SBTTP-501 Lab Course in rDNA Technology and Bioinformatics SBTTP-502 Lab Course in Pharmaceutical Biotechnology SBTTE-502 Lab Course in Animal Biotechnology OR SBTTE-504 Lab Course in Microbial and Enzyme Technology	22	44
	4	SBTTC-551-Plant Biotechnology SBTTC-552-Genomics and Proteomics	SBTTE-551 Nanobiotechnology OR SBTTE-553 Food Biotechnology	SVECP- 551 Publication Ethics (2-Cr)		Research Project SBTTR- 551	SBTTP-551 Lab Course in Plant Biotechnology SBTTP-552 Lab Course in Genomics and Proteomics SBTTE-552 Lab Course in Nanobiotechnology	22	
						(6-Cr)	OR SBTTE-554 Lab Course in Food Biotechnology		

DSE indicates Department Specific Elective Course. Biotechnology student, in a particular semester, can opt either of these courses OR a course offered by other programs of the school. DSC- Department Specific Core, OJT- On Job Training, FP- Field Project, CS- Case Study, R- Research Methodology, P- Publication Ethics, Cr- Credit, VEC- Value Education Course, R- Revision, Credits of four semesters = 88, Total Marks of All Four Semesters = 2200



M. Sc. Second Year Semester III (Level 7.0)

Teaching Scheme

Subject	Course	Course Name	Cı	edits Assign	ed	Teac	hing Scheme
	Code		Theory	Practical	Total	Theory (Hrs/Week)	Practical (Hrs/Week/Batch)
Major	SBTTC-501	rDNA Technology	04		04	04	
(DSC)	SBTTC-502	Bioinformatics	04		04	04	
	SBTTC-503	Pharmaceutical Biotechnology	04		04	04	
Elective (DSE)	SBTTE-501 SBTTE-503	Animal Biotechnology OR Microbial and Enzyme Technology	03		03	03	
Research Project	SBTTR-501	Research Project		04	04		08
DSC	SBTTP-501	Lab Course in rDNA Technology and Bioinformatics		01	01		02
Practical	SBTTP-502	Lab Course in Pharmaceutical Biotechnology		01	01		02
DSE Practical	SBTTE-502	Lab Course in Animal Biotechnology OR		01	01		02
	SBTTE-504	Lab Course in Microbial and Enzyme Technology					
	Total C	redits	15	07	22	15	14



M. Sc. Second Year Semester III (Level 7.0)

Examination Scheme

[20% Continuous Assessment (CA) and 80% End Semester Assessment (ESA)]

Subject	Course Code	Course Name		The	eory		Pr	actical	Total
			Contin	nuous Assessi	ment (CA)	ESA			
			Test I	Test II	Avg of (T1+T2)/2	Total	CA	ESA	
Major (DSC)	SBTTC-501	rDNA Technology	20	20	20	80			100
(DSC)	SBTTC-502	Bioinformatics	20	20	20	80			100
	SBTTC-503	Pharmaceutical Biotechnology	20	20	20	80			100
Elective (DSE)	SBTTE-501	Animal Biotechnology OR	15	15	15	60			75
(DSE)	SBTTE-503	Microbial and Enzyme Technology							
Research Project	SBTTR-501	Research Project					20	80	100
DSC Practical	SBTTP-501	Lab Course in rDNA Technology and Bioinformatics					05	20	25
	SBTTP-502	Lab Course in Pharmaceutical Biotechnology					05	20	25
DSE Practical	SBTTE-502	Lab Course in Animal Biotechnology					05	20	25
	SBTTE-504	OR Microbial and Enzyme Technology							



M. Sc. Second Year Semester IV (Level 7.0)

Teaching Scheme

Subject	Course Code	Course Name	Cı	redits Assigne	d	Teacl	ning Scheme
			Theory	Practical	Total	Theory (Hrs/Week)	Practical (Hrs/Week/Batch)
Major (DSC)	SBTTC-551	Plant Biotechnology	04		04	04	
Major (DSC)	SBTTC-552	Genomics and Proteomics	04		04	04	
Elective (DSE)	UR		03		03	03	
Value Education Course SVECP-551 Publication Ethics		02		02	02		
Research Project	SBTTR-551	Research Project		06	06		12
DSC Practical	SBTTP-551	Lab Course in Plant Biotechnology		01	01		02
DSC Practical	SBTTP-552	Lab Course in Genomics and Proteomics		01	01		02
DSE Practical SBTTE-552 SBTTE-554 Lab Course in Nanobiotechnology OR Lab Course in Food Biotechnology			01	01		02	
	Total Credits			09	22	13	18



M. Sc. Second Year Semester IV (Level 7.0)

Examination Scheme

[20% Continuous Assessment (CA) and 80% End Semester Assessment (ESA)]

Subject	Course Code	Course Name		The	ory		Pr	actical	Total
			Conti	nuous Assessn	nent (CA)	ESA			
			Test I	Test II	Avg of (T1+T2)/2	Total	CA	ESA	
Maior (DSC)	SBTTC-551	Plant Biotechnology	20	20	20	80			100
Major (DSC)	SBTTC-552	Genomics and Proteomics	20	20	20	80			100
Elective	SBTTE-551	Nanobiotechnology OR	15	15	15	60			75
(DSE)	SBTTE-553	Food Biotechnology	13	15	13				73
Value Education Course	SVECP-551	Publication Ethics					10	40	50
Research Project	SBTTR-551	Research Project					30	120	150
DSC Practical	SBTTP-551	Lab Course in Plant Biotechnology					05	20	25
	SBTTP-552	Lab Course in Genomics and Proteomics					05	20	25
DSE Practical	SBTTE-552 SBTTE-554	Lab Course in Nanobiotechnology OR Lab Course in Food					05	20	25
	5B11L 554	Biotechnology							

SBTTC-501: rDNA Technology Teaching Scheme

Course	Course Name	Teaching S	Scheme (Hrs.)	Credits Assigned			
Code		Theory	Practical	Theory	Practical	Total	
SBTTC-501	rDNA Technology	04		04		04	

Assessment Scheme

Course Code	Course		Th	eory		Practical		Total
	Name	CA						
		Test I	Test II	Avg of (T1+T2)/2	ESA	CA	ESA	
SBTTC-501	rDNA Technology	20	20	20	80			100

Course pre-requisite:

> Students should be aware of the basics of different processes of Genetics and Molecular Biology.

Course objectives:

➤ The objective of this course is to familiarize the students with concept of rDNA, clone and gene cloning, cloning strategies, tools and techniques, applications and advantages and alternatives to transgenics etc.

Course outcomes: Students will be able to be

> GM literate i.e. aware about rDNA technology, its advantages and disadvantages in addition to tools and techniques. It will help in avoiding spread of misconception about GMO in society.

Curriculum Details:

Module	Unit	Topic	Hrs.					
No.	No.							
1.0		Fundamentals of Genetic Engineering	15					
	1.1	Introduction to concept of r-DNA, clone and gene cloning. Scope and						
		lestones in Genetic Engineering.						
	1.2	Strategies and Molecular Tools: Restriction and modifying enzymes. DNA and RNA markers.						
	1.3	Vectors: Cloning and expression vectors; vector components: Promoters, selectable markers, reporter genre, ori, URRs, codon optimization, Properties and Applications.						
	1.4	Commonly used vectors: Plasmids, bacteriophages, Phagemids and						

		cosmids. Artificial chromosomes.	
2.0		Gene Cloning strategies and tools	15
	2.1	Isolation and purification of chromosomal and plasmid DNA, Yield analysis, Nucleic acid amplification and its applications. Genomic and c-DNA library preparation and application.	
	2.2	Cloning Methods: Blunt end cloning, Sticky end and sticky end PCR cloning, TA cloning, PCR recombination, Integration PCR, In-Fusion TM Cloning, TOPO Cloning, Gateway cloning etc.	
	2.3	Methods of screening: Selection by complementation, antibiotic resistance, colony PCR etc.	
	2.4	Expression analysis: Phenotype, RNA and Protein level. Northern blot, Primer extension, S1 mapping, RNase protection assay, Reporter assays, RT-PCR and Real time q-PCR, Nucleic acid microarray, Transcriptome sequencing, Western blotting.	
3.0		Applications of r DNA Technology I	15
	3.1	Heterologous expression of proteins. Vector engineering and codon optimization, host engineering	
	3.2	Expression in bacteria, expression in mammalian and plant cells,	
	3.3	Processing of Recombinant proteins: Purification and refolding, characterization of recombinant proteins, stabilization of proteins.	
	3.4	Process and applications of Phage Display. GMO (Microorganisms, Plants and animals) with traits having applications in different sectors: A. Health, B. Agriculture, C. Environment and D. Industrial	
4.0		Applications of r DNA Technology II	15
	4.1	Gene silencing: Strategies, applications and advantages. Genome editing: Strategies, applications and advantages.	
	4.2	Gene therapy: Principles of Gene therapy: Vector engineering.	
	4.3	Strategies of gene delivery. Gene replacement/augmentation therapy, success and limitations of gene therapy.	
	4.4	Genetic engineering guidelines, Regulatory bodies, GEAC, RCGM and IBSC.	
		Total	60

References:

- 1. Sambrook, J., Fritisch E. F. and Maniatis, T. Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory Press, New York, 2000.
- 2. Glover, D. M. and Hames, B. D. DNA Cloning: a Practical Approach, IRL, Press, Oxford, 1995.
- 3. Kaufman, P. B., Wu, W., Kim, D. and Cseke, L. J. Molecular and Cellular Methods in Biology and Medicine, CRC Press, Florida, 1995.
- 4. Berger, S. L. and Kimmel, A. R. Methods in Enzymology ,Vol. 152, Guide to Molecular Cloning Techniques, Academic Press, Inc. San Diego, 1998.
- 5. Goeddel, D. V. Methods in Enzymology, Vol 185, Gene Expression Technology, Academic Press, Inc., San Diegoo, 1990.

- 6. Mickloss, D. A. and Freyer, G. A. DNA Science. A First Course in Recombinant Technology, Cold Spring Harbor Laboratory Press, New York, 1990.
- 7. Primrose, S. B. DNA Science: A First Course in Recombinant Technology, Blackwell Scientific Publishers, Oxford, 1994.
- 8. Davies, J. A. and Raznikoff,, W. S. Milestones in Biotechnology. Classic papers on Genetic Engineering, Butterworth-Heinemann, Boston, 1992.
- 9. Walker, M. R. and Rapley, R. Route Maps in Gene Technology, Blackwell Science Ltd., Oxford, 1997.
- 10. Kingsman, S. M. and Kingsman, A. J. Genetic Engineering. An Introduction to gene analysis and exploitation in eukaryotes, Blackwell Scientific Publications, Oxford, 1998.
- 11. Glick, B. R., Pasternak, J. J. and Patten, C. L. Molecular Biotechnology: Principles and Applications of Recombinant DNA, 6th Edition, American Society for Microbiology, 1994.

SBTTP-501 Lab Course in rDNA Technology

Part A

- 1. Genetic recombination (conjugation, transformation, transduction) in bacteria.
- 2. Gene cloning: Restriction, digestion and ligation, DNA Cloning in plasmid vectors and analysis of gene products.
- 3. Preparation of competent cells and transformation by CaCl2 method.
- 4. DNA amplification.
- 5. DNA fingerprinting: RFLP, RAPD
- 6. Blotting and hybridization techniques: Western, Southern & Northern hybridization.
- 7. Gene expression in *E. coli*
- 8. Agarose gel electrophoresis by using DNA markers for molecular weight determination

SBTTC-502: Bioinformatics

Teaching Scheme

Course	Course Name	Teaching	Scheme (Hrs.)	Credits Assigned			
Code		Theory	Practical	Theory	Practical	Total	
SBTTC-502	Bioinformatics	04		04		04	

Assessment Scheme

Course Code C	Course Name		Th	eory		Pra	ctical	Total
		CA						
		Test I	Test II	Avg of (T1+T2)/2	ESA	CA	ESA	
SBTTC-502 B	sioinformatics	20	20	20	80			100

Course pre-requisite:

The students should be aware of the basics of computers and Molecular Biology.

Course objectives:

The objective of this course is to familiarize students with basic concepts of sequences, structural alignment, database searching and protein structure prediction.

Course outcomes: On completion of this course, the students shall:

- Explain the theoretical knowledge of database system and algorithms.
- Analyze and discuss the results in light of molecular biological knowledge (sequence alignment and Phylogenetic tree plot)
- Collect the proficient knowledge to solve biological system- a multi-disciplinary problem
- ➤ Develop the key skills of molecular modeling techniques currently practiced in any pharmaceutical research and development unit

Curriculum Details:

Module	Unit	Topic	Hrs.					
No.	No.							
1.0		Introduction to Bioinformatics	15					
	1.1	Introduction to Bioinformatics Resources Bioinformatics Resources:						
		NCBI, EBI, ExPASy						
	1.2	RCSB, And DDBJ: The knowledge of databases and bioinformatics tools						
		available at these resources						
	1.3	Organization of databases: data contents, purpose and utility.						
	1.4	Open access bibliographic resources and literature databases: PubMed,						
		BioMed Central, Public Library of Sciences (PloS), Cite Xplore						
2.0		Sequence databases	15					

	2.1	Sequence databases: Nucleic acid sequence databases: GenBank, EMBL,	
		DDBJ; Protein sequence databases: Uniprot-KB: SWISS-PROT,	
		TrEMBL, UniParc	
	2.2	Structure Databases: PDB, NDB, PubChem, ChemBank.	
	2.3	Sequence file formats: Various file formats for bio-molecular sequences:	
		GenBank, FASTA, GCG, MSF etc.	
	2.4	Protein and nucleic acid properties: Proteomics tools at the ExPASy server, GCG utilities and EMBOSS, Computation of various parameters	
3.0		Sequence analysis	15
	3.1	Sequence Analysis: Basic concepts of sequence similarity, identity and	
		homology, definitions of homologues, orthologues, paralogues and	
		xenologues	
	3.2	Scoring matrices: basic concept of a scoring matrix, Matrices for nucleic	
		acid and proteins sequences	
	3.3	PAM and BLOSUM series	
	3.4	Matrix derivation methods and principles	
4.0		Sequence alignment	15
	4.1	Sequence alignment: Measurement of sequence similarity; Similarity and	
		homology.	
	4.2	Pairwise sequence alignment: Basic concepts of sequence alignment	
	4.3	Needleman and Wunsch, Smith and Waterman algorithms for pairwise	
		alignments, gap penalties	
	4.4	Use of pairwise alignments for analysis of Nucleic acid and protein	
		sequences and interpretation of results.	
		Total	60

References

- 1. Mount, D., Bioinformatics: Sequence and Genome Analysis, Cold Spring Harbor Laboratory Press, New York, 2004.
- 2. Baxevanis, A. D. and Francis Ouellellette, B. F. Bioinformatics- a Practical Guide to the Analysis of Genes and Proteins, Wiley India Pvt Ltd, 2009.
- 3. David, P. S., Attwood, T. K. Introduction to bioinformatics, 1st Edition, Pearson Education, 1999.

SBTTP-501 Lab Course in Bioinformatics

Part B

- 1. Proteomics tools
- 2. Structural and functional predictions
- 3. Phylogenetic Analysis, Phylogenetic tree construction
- 4. DNA and protein sequence and PDB file formats
- 5. Local and global sequence alignments of protein and DNA sequences
- 6. Needleman Wunsch and Smith-Waterman algorithm
- 7. BLAST, Multiple sequence alignment

SBTTC-503: Pharmaceutical Biotechnology

Teaching Scheme

Course Code	Course Name	Teaching Scheme (Hrs.)		(Credits Ass	igned
		Theory	Practical	Theory	Practical	Total
SBTTC-503	Pharmaceutical Biotechnology	04		04		04

Assessment Scheme

Course Code	Course Name		Theory				ctical	Total
			CA					
		Test I	Test II	Avg of (T1+T2)/2	ESA	CA	ESA	
SBTTC-503	Pharmaceutical Biotechnology	20	20	20	80			100

Course pre-requisite:

- Basic understanding of Molecular Biology, Genetics, and Microbiology.
- ➤ Knowledge of basic concepts in pharmaceutical sciences.

Course objectives:

- ➤ The objective of this course is to apply the basic concepts in the specific field of pharmaceutical biotechnology.
- ➤ To expose students to gain insights into identification and design of drugs that could be potentially useful in the identification of candidate drug which have efficacy in cell culture or animal models and thus the most effective compound could be employed based on the above results to put into clinical trials.

Course outcomes: Students will be able to

- Explain the strategies and various steps of new drug discovery process.
- Explain the concept of pharmacodynamics and pharmacokinetics
- Apply the knowledge of pharmaceutical manufacturing in the production of biopharmaceuticals like antibiotics, vaccines, proteins and hormones
- Earry out the quality control procedures in the production of various biopharmaceuticals
- Explain the regulatory aspects in the development of pharmaceuticals.

Curriculum Details:

Module No.	Unit No.	Topic	Hrs.
1.0	110.	Antibiotics and Synthetic antimicrobial agents	15
	1.1	Concept of bioassay, therapeutic index, MIC and LD50 Mechanism of action, microbial resistance	13
	1.2	Therapeutic, prophylactic usage and adverse reactions of antibiotic and	
	1.2	synthetic antimicrobial agents: β -lactam, aminoglycosides, tetracyclines, ansamycins, macrolides, peptide antibiotics	
	1.3	Synthetic antibiotics: Sulphonamides, Chloramphenicol, Quinolone. Antifungal antibiotics: Amphotericin B, Griseofulvin and Fluconazole	
	1.4	Antiviral drugs: Acyclovir, Zidovudine, Amantadine. Antitumor drugs: Bleomycin, Dactinomycin	
2.0		Drug discovery methods	15
	2.1	Drug Discovery Process, biological activity directed and other types of screening, natural products	
	2.2	Combinatorial chemistry; General overview of validation techniques, Methods of Drug Discovery and development, QSAR and SAR.	
	2.3	Concepts of Bioavailability, Process of drug absorption, Pharmacokinetic processes, Timing for optimal therapy	
	2.4	Drug delivery considerations for the new biotherapeutics.	
3.0		Pharmacology of drugs	15
	3.1	Physicochemical properties in relation to biological action, Effects of route of administration, Drug Targets, Validation techniques of Pharmaceutical targets	
	3.2	Pharmacokinetics and pharmacodynamics of drugs, Drug Toxicity	
	3.3	Basic terminologies in drug delivery and drug targeting, Doses forms, Various routes of administration of drugs, Strategies for enhanced therapeutic efficacies	
	3.4	DNA vaccines, Vaccines & Monoclonal antibody-based pharmaceuticals and other pharmaceutical products: Streptokinase, streptodornase	
4.0		Formulations and Regulations of drug	15
	4.1	Formulation of Biotechnological Products, Drug Delivery,	
	4.2	Examples of some Biotechnological products in clinical development.	
	4.3	Role of FDA, ICH Guidelines	
	4.4	The Regulation of Pharmaceutical Biotechnological Products and Ethical Issues.	
		Total	60

Text Books and Reference Books

1. Hugo, W. B. & Russel, A. R., Pharmaceutical Microbiology, Sixth Edition. Blackwell Scientific Publications, 1998.

- 2. Kadam, S. S., Mahadik, K. R. and Bothra, K. G. Principles of medicinal chemistry, Vol. 1, Edition: 17, Nirali Publication, 2017.
- 3. Gokhale, S. D., KoKate, C. K. Pharmacognosy, Edition:18, Nirali Prakashan Educational Publication, 2017.
- 4. Singh., B. D. Biotechnology Expanding Horizon, First Edition, Kalyani Publication, Delhi, 2004.
- 5. Vyas, S. P., & Dixit, V. K. Pharmaceutical Biotechnology, 12th edition, CBS publishers & distributors, New Delhi, 2009.
- 6. Hooper, D. C., Wolfson. J. S. Quinolone antimicrobial agents, Second edition, Washington, D.C. : American Society for Microbiology, 1993.
- 7. Murray, S., Cooper Quality control in the Pharmaceutical industry, Vol. 2, Academic Press Inc. U. S., New York, 1974.
- 8. Rhem, H. J. & Reed G. Biotechnology: Measuring, Modelling and Control, vol 4, Wiley India Pvt Ltd publications, 2010.
- 9. Gregory Gregoriadis, Drug carriers in biology & medicine, Academic Press New York London, 1979.
- 10. Hillery, A. M., Lloyd, A. W. and Swarbrick, J. Drug Delivery and Targeting For Pharmacists and Pharmaceutical Scientists, Harwood Academic Publisher, 2001.
- 11. Ansel, H. C., Allen, L. V. and Popovich, N. G. Pharmaceutical Dosage Forms and Drug Delivery Systems, 8th Edition, Lippincott Williams and Wilkins Publisher, 2005.
- 12. Shyam Mohapatra, Shivendu Ranjan, Nandita Dasgupta, Raghvendra Kumar and Sabu Thomas. Applications of Targeted Nano Drugs and Delivery Systems, 1st Edition, Elsevier, 2018.

SBTTP-502 Laboratory Course in Pharmaceutical Biotechnology

- 1. Quality control of antibiotic and non-antibiotic formulations using titrimetric, spectrophotometric, chromatographic methods as per IP/US Pharmacopoeia.
- 2. Microbiological assays of antibiotics.
- 3. Sterility testing and stability testing of parenteral formulations.
- 4. Sterility testing of pharmaceutical products (intra-venous injections, antibiotics and vitamins)
- 5. Assays for screening antimicrobial/antifungal agents from plants and other natural sources.
- 6. Test for disinfectants (phenol coefficient / RWC method).
- 7. Determination of antibacterial spectrum for drugs/antibiotics.
- 8. Testing for antibiotic / drug sensitivity /resistance.
- 9. Determination of MIC value for antimicrobial chemicals.

SBTTE-501 Animal Biotechnology

Teaching Scheme

Course Code	Course	Teaching Sc	heme (Hrs.)	Credits Assigned			
	Name	Theory	Practical	Theory	Practical	Total	
SBTTE-501	Animal Biotechnology	03		03		03	

Assessment Scheme

Course Code	Course	Theory				Pra	ctical	Total
	Name	CA						
		Test I	Test II	Avg of (T1+T2)/2	ESA	CA	ESA	
SBTTE-501	Animal Biotechnology	15	15	15	60		1	75

Course pre-requisite:

➤ Basic understanding of Cell and developmental Biology.

Course objectives:

➤ The objective of this course is to enable students to develop basic skills for vertebrate cell culture, maintenance of cell lines and in vitro application of cell and molecular techniques and also to understand the principles of animal cloning and its applications.

Course outcomes: Students will be able to

- Explain the fundamental scientific principles that underlie cell culture
- Acquire knowledge for isolation, maintaince and growth of cells.
- > Develop proficiency in establishing and maintaining of cell lines.
- Acquire knowledge in animal cloning and its applications

Curriculum Details:

Module	Unit	Topic	Hrs.
No.	No.		
1.0		Introduction to Animal Cell Science	11
	1.1	Structure and organization of animal cell. Equipments and materials for animal	
		cell culture technology.	
	1.2	Primary and established cell line cultures. Introduction to the balanced salt	
		solutions and simple growth medium.	
	1.3	Brief discussion on the chemical, physical and metabolic functions of	

		different constituents of culture medium.	
	1.4	Role of carbon dioxide. Role of serum and supplements. Serum and protein	
		free defined media and their applications. Application of animal cell culture.	
2.0		Cell lines and its applications	11
	2.1	Primary and secondary culture of animal cells: types of primary cell	
		cultures, isolation of tissue and primary culture.	
	2.2	Subcultured propagation, criteria for subculture and propagation, split ratio, subculture in suspension.	
	2.3	Basic techniques of mammalian cell culture in vitro, disaggregation of	
		tissue and primary culture; maintenance of cell culture; cell separation.	
	2.4	Scaling- up of animal cell culture.	
3.0		Cell lines and its applications	12
	3.1	Culture and maintenance of cell lines, Biology and characterization of the	
		cultured cells, Measurement of viability,	
	3.2	Model animals in animal biotechnology	
	3.3	DNA transfer techniques in to mammalian cells, Microinjection,	
		electroporation, Stem cell etc.	
	3.4	Artificial insemination, IVF, somatic cell nuclear transfer and stem cell	
4.0		technology.	
4.0		Animal Cloning	11
	4.1	Principle, Concept and application of animal cloning	
	4.2	Embryonic stem cells and their applications.	
	4.3	Cell culture based vaccines, viral vectors, Gene therapy	
	4.4	Animal ethics and bio safety, Principle of tissue engineering	
		Total	45

References

- 1. Ian, R. Freshney Culture of Animal Cells: A Manual of Basic Technique and Specialized Applications, 4th Edition, 2000.
- 2. Ranga, M. M. Animal Biotechnology, 2nd edition, Agrobios, 2007.
- 3. Masters, J. R. W. Animal Cell Culture, 3rd Edition, Oxford University Press, 2000.
- 4. Marshak, D. R., Gardner, R. L. and Gottlieb, D. Stem Cell Biology, Volume 40, Cold Spring Harbor Publication, 2001.

SBTTE- 502 Laboratory course in Animal Biotechnology

- 1. Enumeration of Red Blood cells (RBCs) from sheep and human blood
- 2. Isolation of Cells from Liver tissue and its Quantification
- 3. Cell Viability assay using dye-exclusion method
- 4. Micronucleus Assay
- 5. Hemolytic test for Staphylococcus aureus
- 6. Con-A induced Hemagglutination Assay
- 7. Anchorage independent cell culture(Lymphocyte culture)
- 8. Anchorage dependent cell culture (chick embryo cell culture)

- 9. Epithelial cell culture
- 10. Anti-angiogenic activity using chick chorio-allantoic membrane(CAM) assay
- 11. Preparation of culture media.

BTTE-503 Microbial and Enzyme Technology Teaching Scheme

Course	Course Name	Teaching S	Scheme (Hrs.)		Credits Ass	igned
Code		Theory	Practical	Theory	Practical	Total
	Microbial and Enzyme Technology	03		03		03

Assessment Scheme

Course Code	Course Name	Theory			Pra	ctical	Total	
			CA					
		Test I	Test II	Avg of (T1+T2)/2	ESA	CA	ESA	
SBTTE-503	Microbial and Enzyme Technology	15	15	15	60			75

Course pre-requisite: Students should be aware of basics in Microbiology and Enzymology Course objectives:

- > To make Students learn structural and functional relationships in enzymes and altering their structure in order to function 'better'.
- To provide basic knowledge of microbes and enzyme technology and use of enzymes as tools in industry, agriculture and medicine.
- > To exploit microbes and enzymes for the production of human beneficial products

Course outcomes: Students will be able to

- Learn kinetics of enzyme catalyzed reactions & enzyme inhibitory and regulatory process.
- > Perform immobilization of enzymes
- > Understanding of microbial pathways
- > Get exposure of wide applications of microbes and enzymes and their future potential

Curriculum Details:

Module No.	Unit No.	Topic	Hrs.
1.0	110.	Fundamentals of Microbial Metabolism	12
	1.1	Isolation, development, presentation and improvement of industrially	
		important micro-organism; isolation of auxotrophic mutants	

	1.2	Isolation of revertant mutants and use of recombinant systems for improvement of industrial microorganisms; metabolic pathways:	
		Regulatory mechanism of metabolic pathways in industrial strains	
	1.3	Bioenergetics-basic principles; equilibria and concept of free energy;	
		coupled processes; glycolysis and glycolytic enzymes regulation	
	1.4	TCA-cycle and enzyme regulation; oxidative phosphorylation and	
		enzyme regulation; fatty acid metabolism; principles of metabolic	
		regulation; regulatory steps	
2.0		Enzyme catalysis and kinetics	11
	2.1	Source of enzymes; production, isolation and purification of enzymes;	
		characterization in terms of pH, temperature, ionic strength, substrate and	
		product tolerance, effect of metal ions etc.	
	2.2	Enzyme kinetics: Enzymes as Biological catalysts; enzyme action:	
		Active site, functional group, enzyme substrate complex, cofactors	
	2.3	Michaelis-Menten equation; enzyme inhibition; methods of plotting	
		enzyme kinetics data	
	2.4	Enzyme turnover; enzyme induction, repression, covalent modification,	
		isoenzymes, allosteric effect	
		F	
3.0		Enzyme Engineering and Immobilization	11
3.0	3.1	Rationales for enzyme engineering, steps in enzyme engineering,	11
3.0		Rationales for enzyme engineering, steps in enzyme engineering, chemical methods for modifying enzyme activity	11
3.0	3.1	Rationales for enzyme engineering, steps in enzyme engineering, chemical methods for modifying enzyme activity Site directed mutagenesis, protein engineering, mechanism of protein	11
3.0	3.2	Rationales for enzyme engineering, steps in enzyme engineering, chemical methods for modifying enzyme activity Site directed mutagenesis, protein engineering, mechanism of protein folding and its pathological effects	11
3.0		Rationales for enzyme engineering, steps in enzyme engineering, chemical methods for modifying enzyme activity Site directed mutagenesis, protein engineering, mechanism of protein folding and its pathological effects Immobilized enzyme technology: different techniques of immobilization	11
3.0	3.2	Rationales for enzyme engineering, steps in enzyme engineering, chemical methods for modifying enzyme activity Site directed mutagenesis, protein engineering, mechanism of protein folding and its pathological effects Immobilized enzyme technology: different techniques of immobilization of enzymes and whole cells	11
3.0	3.2	Rationales for enzyme engineering, steps in enzyme engineering, chemical methods for modifying enzyme activity Site directed mutagenesis, protein engineering, mechanism of protein folding and its pathological effects Immobilized enzyme technology: different techniques of immobilization of enzymes and whole cells Advantages and disadvantages of immobilization; application of cell and	11
	3.2	Rationales for enzyme engineering, steps in enzyme engineering, chemical methods for modifying enzyme activity Site directed mutagenesis, protein engineering, mechanism of protein folding and its pathological effects Immobilized enzyme technology: different techniques of immobilization of enzymes and whole cells Advantages and disadvantages of immobilization; application of cell and enzyme immobilization and biosensors	
4.0	3.2	Rationales for enzyme engineering, steps in enzyme engineering, chemical methods for modifying enzyme activity Site directed mutagenesis, protein engineering, mechanism of protein folding and its pathological effects Immobilized enzyme technology: different techniques of immobilization of enzymes and whole cells Advantages and disadvantages of immobilization; application of cell and enzyme immobilization and biosensors Clinical and Industrial Enzymes	11
	3.2	Rationales for enzyme engineering, steps in enzyme engineering, chemical methods for modifying enzyme activity Site directed mutagenesis, protein engineering, mechanism of protein folding and its pathological effects Immobilized enzyme technology: different techniques of immobilization of enzymes and whole cells Advantages and disadvantages of immobilization; application of cell and enzyme immobilization and biosensors Clinical and Industrial Enzymes Clinical and diagnostics enzymes and their applications, enzymes as	
	3.2 3.3 3.4 4.1	Rationales for enzyme engineering, steps in enzyme engineering, chemical methods for modifying enzyme activity Site directed mutagenesis, protein engineering, mechanism of protein folding and its pathological effects Immobilized enzyme technology: different techniques of immobilization of enzymes and whole cells Advantages and disadvantages of immobilization; application of cell and enzyme immobilization and biosensors Clinical and Industrial Enzymes Clinical and diagnostics enzymes and their applications, enzymes as therapeutic agents	
	3.2	Rationales for enzyme engineering, steps in enzyme engineering, chemical methods for modifying enzyme activity Site directed mutagenesis, protein engineering, mechanism of protein folding and its pathological effects Immobilized enzyme technology: different techniques of immobilization of enzymes and whole cells Advantages and disadvantages of immobilization; application of cell and enzyme immobilization and biosensors Clinical and Industrial Enzymes Clinical and diagnostics enzymes and their applications, enzymes as therapeutic agents Enzymes in food processing, leather, textile, detergent and	
	3.2 3.3 3.4 4.1 4.2	Rationales for enzyme engineering, steps in enzyme engineering, chemical methods for modifying enzyme activity Site directed mutagenesis, protein engineering, mechanism of protein folding and its pathological effects Immobilized enzyme technology: different techniques of immobilization of enzymes and whole cells Advantages and disadvantages of immobilization; application of cell and enzyme immobilization and biosensors Clinical and Industrial Enzymes Clinical and diagnostics enzymes and their applications, enzymes as therapeutic agents Enzymes in food processing, leather, textile, detergent and pharmaceuticals and fine chemical industries	
	3.2 3.3 3.4 4.1	Rationales for enzyme engineering, steps in enzyme engineering, chemical methods for modifying enzyme activity Site directed mutagenesis, protein engineering, mechanism of protein folding and its pathological effects Immobilized enzyme technology: different techniques of immobilization of enzymes and whole cells Advantages and disadvantages of immobilization; application of cell and enzyme immobilization and biosensors Clinical and Industrial Enzymes Clinical and diagnostics enzymes and their applications, enzymes as therapeutic agents Enzymes in food processing, leather, textile, detergent and pharmaceuticals and fine chemical industries Enzymes in organic solvents and ionic liquids: various organic solvents	
	3.2 3.3 3.4 4.1 4.2 4.3	Rationales for enzyme engineering, steps in enzyme engineering, chemical methods for modifying enzyme activity Site directed mutagenesis, protein engineering, mechanism of protein folding and its pathological effects Immobilized enzyme technology: different techniques of immobilization of enzymes and whole cells Advantages and disadvantages of immobilization; application of cell and enzyme immobilization and biosensors Clinical and Industrial Enzymes Clinical and diagnostics enzymes and their applications, enzymes as therapeutic agents Enzymes in food processing, leather, textile, detergent and pharmaceuticals and fine chemical industries Enzymes in organic solvents and ionic liquids: various organic solvents and ionic liquids used in bio catalysis	
	3.2 3.3 3.4 4.1 4.2	Rationales for enzyme engineering, steps in enzyme engineering, chemical methods for modifying enzyme activity Site directed mutagenesis, protein engineering, mechanism of protein folding and its pathological effects Immobilized enzyme technology: different techniques of immobilization of enzymes and whole cells Advantages and disadvantages of immobilization; application of cell and enzyme immobilization and biosensors Clinical and Industrial Enzymes Clinical and diagnostics enzymes and their applications, enzymes as therapeutic agents Enzymes in food processing, leather, textile, detergent and pharmaceuticals and fine chemical industries Enzymes in organic solvents and ionic liquids: various organic solvents	

Reference Books:

- 1. Dixon, M. and Webb, E. C. Enzyme Inhibition and Activation in: Enzymes, 3rd Edition, Pp. 126–136, 332–399, Academic Press, New York, 1979.
- 2. Palmer, T. Understanding enzymes, 4th edition, Prentice Halls/Ellis Horwood Ltd. Publication, London, 1995.
- 3. Price, N. C. and Stevens, L. Fundamentals of Enzymology: Cell and Molecular Biology of Catalytic Proteins, 3rd Edition, Oxford science publications, New York, 1999.
- 4. Buchholz, K., Kasche, V. and Bornscheuer, U. T. Biocatalysts and enzyme technology, 2nd Edition, Wiley-Blackwell Publication, 2012.

- 5. Copeland, Robert, A. Enzymes: a practical introduction to structure, mechanism and data analysis, John Wiley & Sons, 2004.
- 6. Balasubramanian, D., Bryce, C. F. A., Dharmalingam, K., Green, J. and Jayaraman, K. Concepts in Biotechnology, 2nd Edition, Universities Press, 2004.
- 7. Rastogi, S. C., Mendiratta, N. and Rastogi, P., Bioinformatics Methods and Applications :(Genomics, Proteomics and Drug Discovery), 4th Edition, PHI Learning, 2006.
- 8. Satyanarayana, U. and Chakrapani, U. Biotechnology, 1st Edition, Books and Allied Ltd. 2005.
- 9. Smith, J. E. Biotechnology, 5th Edition, Cambridge University Press, 2009.
- 10. Berg, J. M., Tymoczko, J. L., Gatto G. J. and Stryer, L. Biochemistry, 8th Edition, WH Freeman and Company, 2015.
- 11. Creighton, T. E. Protein Structure and Molecular Properties, 2nd Edition, W.H. Freeman and Co. Ltd, 1993.
- 12. Primrose, S. B. and Twyman, R. M., Principles of Gene Manipulation and Genomics, 7th Edition, Blackwell Publishing 2006.
- 13. Sambrook, J., Fritsch, E. R. and Maniatis, T. Molecular Cloning: A Laboratory Manual, 2nd Edition, Cold Spring Harbo Laboratory Press, 1989.

SBTTE-504 Laboratory Course in Microbial and Enzyme Technology

- 1. Isolation of microbial strains for the production of commercially important enzymes.
- 2. Production of commercially important enzymes from microbial source.
- 3. Standardization of medium, composition for the optimum production of enzymes
- 4. Partial purification of isolated enzymes
- 5. Determination of enzyme activity and specific activity of α -amylase/lipase.
- 6. Study of Maltose calibration curve
- 7. Characterization of enzymes effect of pH, temperature metal ions, substrate and inhibitors on enzyme activity.
- 8. Study of kinetic parameters Km, Vmax and Kcat
- 9. Molecular weight determination of enzyme by gel filtration method
- 10. Method of checking the purity of the enzymes SDS PAGE
- 11. Immobilization of enzymes
- 12. Characterization of clinical and industrially important enzymes

SEMESTER IV

SBTTC-551: Plant Biotechnology

Teaching Scheme

Course	Course Name	Teaching S	cheme (Hrs.)		Credits	Assigned
Code		Theory	Practical	Theory	Practical	Total
SBTTC-551	Plant Biotechnology	04		04		04

Assessment Scheme

Course Code	Course Name	Theory				Pra	ctical	Total
		CA						
		Test I	Test II	Avg of (T1+T2)/2	ESA	CA	ESA	
SBTTC-551	Plant Biotechnology	20	20	20	80			100

Course pre-requisite:

The students should be familiar with the fundamentals of Plant Sciences and Biotechnology

Course objectives:

- To acquaint the students with basic principles and various methods of Tissue Culture.
- To impart knowledge about varied methods of gene transfer and transgenic plant development.
- > To underhand basics of secondary metabolites and their engineering.
- > To acquire knowledge about molecular markers and their use in plant breeding.

Course outcomes: Students will

- ➤ Demonstrate the knowledge about the techniques of Plant Tissue Culture and acquire comprehensive knowledge on GM technology for quality characteristics and their role in crop improvement.
- Acquire knowledge in metabolic engineering and industrial products.
- > Develop skills in molecular marker studies and their use in plant breeding.
- > Shall develop scientific skills to work in Plant tissue culture, Pharmaceutical and Research laboratories.

Curriculum Details:

Module	Unit	Topic	Hrs.			
No.	No.					
1.0		UNIT I	15			
	1.1	History: Important events in the history of plant tissue culture; Laboratory				
		Requirements and General Techniques; Cellular Totipotency				
	1.2	Tissue Culture Media: Introduction, media constituents, media selection,				
		media preparation; Callus Culture; Micropropagation: Introduction,				
		techniques, applications, production of pathogen free plants				
	1.3	Somatic Embryogenesis; Haploid Production: Introduction, techniques,				
		factor affecting androgenesis, ontogeny of androgenic haploids				

	1.4	Plant regeneration from pollen embryos, homozygous diploids, applications, limitations; Triploid production.	
2.0		UNIT II	15
	2.1	Somaclonal & gametoclonal variations; Protoplast Culture: Protoplast isolation, fusion and regeneration,	
	2.2	Cybrids; Embryo Culture and embryo rescue: Introduction, techniques; Synthetic Seeds	
	2.3	Cell and Suspension Culture: Introduction, isolation of single cells, suspension cultures, culture of single cells, plant cell reactors, applications of cell culture	
	2.4	Production of secondary metabolites: Introduction, strategies used to optimize product yield, commercial aspects	
3.0		UNIT III	15
	3.1	Introduction to transgenic technology: Conventional breeding versus Transgenesis; Introduction to <i>Agrobacterium tumefaciens</i> and <i>A. rhizogenes</i> , Features of Ti and Ri Plasmids and their use as vectors, Binary and co-integrate vectors	
	3.2	Agrobacterium mediated transformation, Direct DNA transfer to plants, Detection, characterization and expression of Transformants	
	3.3	Applications of plant transformation for productivity and performance: GM technology for Conferring resistance to biotic stresses (pests, viruses and fungi) and abiotic stresses (draught and salt), Herbicide resistance	
	3.4	Increasing shelf life of fruits and flowers, Enhancing the nutritional quality (pro-vitamin A), Chloroplast Transformation	
4.0		UNIT IV	15
	4.1	Metabolic engineering and industrial products: Plant secondary metabolites: alkaloids, industrial enzymes	
	4.2	Biodegradable plastic: polyhydroxybutyrate, therapeutic proteins, lysosomal enzymes, antibodies, edible vaccines, oleosin partitioning technology etc	
	4.3	Aspects related to commercial release of transgenic crops.	
	4.4	Molecular marker aided breeding: RFLP, RAPD, Microsatelites, AFLP etc.	
		Total	60

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- 1. Razdan, M. K. Introduction to Plant Tissue Culture, 2nd Edition, Oxford and IHB publishing Co. Pvt. Ltd. 2003.
- 2. Kumar, U. Methods in Plant Tissue Culture, 1st Edition, Bikaner: Agro Botanica, 1999.
- 3. Misawa M. Plant tissue culture: An Alternative for Production of Useful Metabolites, Daya Publishing House, New Delhi, 1994.
- 4. Bhojwani, S. S., and Razdan, M. K. Plant Tissue Culture, Theory and Practice a Revised Edition, Volume 5, Elsevier India Ltd, 1996.
- 5. S. Ignacimuthu, S. J. Applied Plant Biotechnology, Science Publishers, U.S., 1997.
- 6. Young, M. M., Flower, M. W. and Warren, G. S. Plant Biotechnology: Comprehensive Biotechnology, Second Supplement, 1st Edition, Oxford Pergaman press, 1992.

- 7. Hammond, J.; McGarvey, P., and Yusibov, V. Plant Biotechnology: New Products and Applications: 240 (Current Topics in Microbiology and Immunology), Springer-Verlag Berlin and Heidelberg GmbH & Co, New Delhi, 2000.
- 8. Mantell, J. H., Matthews, J. A. and Mckee, R. A. Principles of Plant Biotechnology: An Introduction to Genetic Engineering in Plants, Oxford, Blackwell Scientific Publication, 1985.
- 9. Singh, C. H. Biotechnology in Crop Improvement, International Book Distribution Company, Lucknow, 1998.
- 10. Gupta, P. K. Elements of Biotechnology, Rastogi and Company Meerut, 1996.
- 11. Pareek, L. K. and Swarnakar, P. L. Trends in Plant Tissue Culture and Biotechnology, Agro Botanica Publishers, 1997.
- 12. Esra Galun, Adina Breiman, Transgenic Plants, Imperial College Press, 1997.
- 13. Singh, B. D. Biotechnology, Kalyani Publishers, Ludhiana, India, 1998.
- 14. Narayanaswamy, S. Plant Cell and Tissue Culture, Tata McGraw Hill Education Publication, 1992.
- 15. Hammond, J., McGarvey, P. and Yusibo, V. Plant Biotechnology: New Products and Applications: 240 (Current Topics in Microbiology and Immunology), Springer-Verlag Berlin and Heidelberg GmbH & Co., 1999.
- 16. Fu, T. J., Singh, G. and Curist, W. R. Plant Cell and Tissue Culture for the production of Food Ingredients, Springer Boston MA press, 1999.
- 17. Chawla, H. S. Biotechnology in Crop Improvement, International Book Distribution Company, 1998.

SBTTP-551 Laboratory Course in Plant Biotechnology

- 1. Preparation of MS medium
- 2. Surface sterilization
- 3. Micro propagation of plant through multiplication of pre-existing meristems.
- 4. Hardening of *in vitro* raised plants
- 5. Encapsulation of somatic embryos
- 6. Embryo culture and embryo rescue.
- 7. Protoplast isolation, fusion and culture.
- 8. In vitro production of fast growing normal root culture for production of secondary metabolites
- 9. Elicitation of plant cells for secondary metabolites
- 10. Agrobacterium Ti plasmid based vector mediated transformation, selection of transformants, reporter gene assay.
- 11. Transformation of plant tissues using Agrobacterium rhizogenes for hairy root production
- 12. Transformation and expression of GFP gene in suitable host.
- 13. Developing RFLP maps, Developing RAPD maps

SBTTC-552: Genomics and Proteomics

Teaching Scheme

Course Code	Course Name	Teaching So	cheme (Hrs.)	Credits Assigned			
Code		Theory	Practical	Theory	Practical	Total	
SBTTC-552	Genomics and Proteomics	04		04		04	

Assessment Scheme

Course Code	Course Name	Theory				Pra	ctical	Total
		CA						
SDTTC 552	Genomics and	Test I	Test II	Avg of (T1+T2)/2	ESA	CA	ESA	
SBTTC-552	Proteomics	20	20	20	80			100

Course pre-requisite:

> The students should be familiar with the Computers and Basic concepts of Bioinformatics.

Course objectives:

- > To understand the concept of proteomics and their applications.
- > To expose the students to methods of studying genetic materials obtained from various environmental samples.
- ➤ To provide students the understanding of basic concepts of sequences, structural alignment, database searching, protein structure prediction and computer-based drug designing.

Course outcomes: On completion of this course, the students shall be able to

- > Perform alignment of sequences and construct the matrix for alignment based on dynamic programming
- > Construct the phylogenetics of different sequences.
- Analyze sequence and structure of bio-macromolecule data
- > Edit the three-dimensional structure of protein using structural bioinformatics tools
- Explain the properties of genetic materials and storage and processing of genetic information.
- Analyze genomic data.
- Explain biological phenomena based on comparative genomics
- > Design transcriptomics and proteomics experiments for studying differential gene expression and related analysis
- > Use metagenomics approach for studying phenomena associated with microbial communities.

Curriculum Details:

Module	Unit	Торіс	Hrs.
No. 1.0	No.	Committee	1.5
1.0	1.1	Genomics Litraduction Methods of granging DNA (isolation and in situation)	15
		Introduction, Methods of preparing DNA (isolation and <i>in-vitro</i> chemical synthesis)	
	1.2	DNA separation techniques and DNA sequence analysis: Introduction, methods, sanger and dideoxy method	
	1.3	Automated sequencing, fluorescence method	
	1.4	Genome mapping: Introduction, methods, construction, use of online tools for genome map construction	
2.0		Genome Engineering	15
	2.1	Microarray technology: Introduction, methods and application,	
	2.2	Combinational genomics: Introduction and application,	
	2.3	Synthetic chromosome pharmacogenomics and proteomics	
	2.4	Synthetic life pharmacogenomics and proteomics.	
3.0		Proteomics	15
	3.1	Introduction and scope	
	3.2	Polyacrylamide gel electrophoresis, Isoelectric focusing	
	3.3	Two dimensional PAGE for proteome analysis and image analysis of 2D gel	
	3.4	Mass spectrometry for protein identification	
4.0		Protein Structure Prediction	15
	4.1	Introduction, principle, tools, application for structure prediction of primary to quaternary structure of proteins	
	4.2	Protein engineering: Introduction, protein chips and functional proteomics,	
	4.3	Protein modeling, introduction, methods and tools	
	4.4	Assigning secondary structure	
		Total	60

References:

- 1. Primrose, S. B. and Twyman, R. M. Principles of Gene Manipulation and Genomics, 7th edition, Wiley Blackwell Publishing, 2006.
- 2. Lesk, A. M. Introduction to Genomics, 7th Edition, Oxford University Press, 2007.
- 3. Lesk, A. M. Introduction to Bioinformatics, 3rd Edition, Oxford University Press, 2017.
- 4. Ghosh, Z. and Mallick, B. Bioinformatics: Principles and Applications, Oxford University Press India, 2008.
- 5. Lewin, B. Genes VIII: International Edition, Pearson Education International, 2003.

SBTTP-552 Laboratory course in Genomics and Proteomics

- 1. Proteomics tools,
- 2. Structural and functional predictions,

- 3. Phylogenetic Analysis, Phylogenetic tree construction
- 4. DNA and protein sequence and retrieval and submission
- 5. Local and global sequence alignment of protein and DNA sequences,
- 6. Needleman Wunsch and Smith-Waterman algorithm,
- 7. BLAST, Multiple sequence alignment

SBTTE-551: Nanobiotechnology

Teaching Scheme

Course	Course Name Teaching Scheme (I		Scheme (Hrs.)		Credits Assigned		
Code		Theory	Practical	Theory	Practical	Total	
SBTTE-551	Nanobiotechnology	03		03		03	

Assessment Scheme

Course Code	Course Name	Theory				Pra	ctical	Total
		CA						
		Test I	Test II	Avg of (T1+T2)/2	ESA	CA	ESA	
SBTTE-551	Nanobiotechnology	15	15	15	60			75

Course pre-requisite: General awareness regarding Nanomaterials and Biotechnological techniques Course objectives:

> To introduce to the students various opportunities in the emerging field of Bioscience & nano-bioscience through nanotechnology and to make the students familiar with the important concepts applicable to bioscience and nano biotechnology and its applications.

Course outcomes: On completion of this course, the students will able to

- ➤ Comprehend the concept of "nanotechnology" and its interdisciplinary aspects.
- Learn various approaches of synthesizing nanomaterials, their advantages and limitations.
- > Gain knowledge about various techniques used for characterizing nanomaterials.
- ➤ Comprehend the importance of engineered nanomaterials for biomedical, therapeutic and environmental applications.
- > Evaluate the potential toxic effects of nanotechnology on living organisms and the environment.

Curriculum Details:

Module	Unit	Topic	Hrs.
No.	No.		
1.0	1	Introduction to Nano Science	11
	1.1	Basic concepts of quantum mechanics and quantum theory	
	1.2	mechanical properties of nanomaterials, structural properties, optical properties, melting and electrical conductivity of nanoparticles	
	1.3	Analytical techniques used in Nanotechnology: Diffraction, UV-visible spectroscopy, optical microscope	
	1.4	Analytical techniques used in Nanotechnology: Atomic force microscope, electron microscope SEM and TEM	
2.0	2	Methods of Synthesis of Nanoparticles	12

	2.1	Use of bacteria, Fungi, Actinomycetes	
	2.2	Magnetotactic bacteria for synthesis of magnetic nanoparticles,	
		mechanism of formulation	
	2.3	Viruses as components for formulation of nanostructured materials,	
		Synthesis process and applications	
	2.4	Role of plants in nanoparticles synthesis.	
3.0	3	Nano-biomaterials	11
	3.1	Biocompatibility, anti-bacterial activity	
	3.2	Principles involved applications. Biomaterial	
	3.3	Nanocircuitry, protein based nanocircuitry	
	3.4	DNA nanostructures for mechanics and computing, DNA based	
		nanomechanical devices	
4.0	4	Nanomedicines	11
	4.1	Development of nanomedicine, nanodrug administration	
	4.2	Diagnostic application, therapeutic applications	
	4.3	Nano-tribology, drug and gene delivery for human health	
	4.4	Nano-biosensors, Nanomatechnology of cleaning environment for	
		heavy metals and bioremediation.	
		Total	45

References:

- **1.** Sulbha Kulkarni, Nanotechnology Principles and Practices, 3rd Edition, Springer Nature Publication, 2015.
- **2.** Mark Ratner and Donier Ratner Nanotechnology A Gentle Introduction to Next Big Idea, 1st Edition, Pearson Publication, 2002.
- 3. Satya Prakash, Quantum Mechanics, Pragati Prakashan Publication Limited, Meerut, 2016.
- **4.** Charles P. P. (Jr.), Frank J. O. Introduction to Nanotechnology, 1st Edition, Wiley-Interscience Publication, 2003.
- **5.** G. Cao, Nanostructures and Nanomaterials: Synthesis, Properties and Applications, Imperial College Press, 2004.
- **6.** Niemeyer, C. M. and Mirkin, C. A. Nanobiotechnology: Concepts, Applications and Perspectives, Wiley, VCH, 2004.
- **7.** Leggett, G. J. and Jones, R. A. L. Bionanotechnology: In Nanoscale Science and Technology, John Wiley & Sons, 2005.
- **8.** Murthy, B. S., Shankar, P., Raj, B., Rath, B. B. and Murday, J. Textbook of Nanoscience and Nanotechnology (Universities Press-IIM Series in Metallurgy and Materials Science), Springer Publication, 2012.
- **9.** Pradeep, T. Nano: The Essentials Understanding Nanoscience and Nanotechnology, McGraw Hill Publishing Company Ltd. 2007.
- 10. Goodsell, D. S. Bionanotechnology: Lessons from Nature, John Wiley & Sons, 2004.
- 11. Bhushan, B. Springer Handbook of Nanotechnology, 2nd Edition, Springer Publications, 2007.
- **12.** Nalwa, H. S. Encyclopedia of Nanoscience and Nanotechnology, Volume1-10, American Scientific Publishers, 2004.

SBTTE-452 Lab Course in Nanobiotechnology

- 1. Synthesis of nanomaterials by using microbial sources (bacteria, Fungi, Actinomycetes)
- 2. Synthesis of nano-magnetic particles
- 3. Detection and Analysis of Nanomaterials using UV-Spectrophotometer
- 4. Study of topography of Nanomaterials by Atomic Force Microscopy
- 5. Characterizations and Analysis of Nanomaterials by SEM/TEM
- 6. Antimicrobial activity of Nanomaterials
- 7. Evaluation of efficacy of drug conjugated with nanomaterials.
- 8. Bioremediation of heavy metals by Nanotechnology
- 9. Pollutant removal using Nanomaterials from industrial effluent.

SBTTE-553: Food Biotechnology

Teaching Scheme

Course	Course Name	Teaching S	Scheme (Hrs.)	Credits Assigned			
Code		Theory	Practical	Theory	Practical	Total	
SBTTE-553	Food Biotechnology	03		03		03	

Assessment Scheme

Course Code	Course Name	Theory				Practical		Total
		CA						
		Test I	Test II	Avg of (T1+T2)/2	ESA	CA	ESA	
SBTTE-553	Food Biotechnology	15	15	15	60			75

Course pre-requisite:

> General awareness of food items

Course objectives:

- ➤ To provide basic knowledge about food biotechnology and preliminary preparation of food before actual processing steps.
- > To know the effect of microorganisms on food and to make the clear understanding about chemical and microbiological properties of food etc.
- > To understand the concept of food born infections and to aware about laws and standards in food biotechnology.

Course outcomes: On completion of this course, students shall be able to:

- > Comprehend the different microorganisms roles involved in food biotechnology with different food items.
- ➤ Define and explain different preliminary steps before and after food fermentation.
- Comprehend phenomenon of food degradation and spoilage by microorganisms with change in the properties of food.
- ➤ To create awareness about different laws and standards in food biotechnology.

Curriculum Details:

Module	Unit	Topic	Hrs.
No.	No.		
1.0		Microbes and Food	12
	1.1	Characteristics of Microorganisms in Food, Types of microorganisms	
		associated with food	

	1.2	Factors affecting the growth of micro-organisms in food, Microbial Food	
		Spoilage, Sources of Microorganisms in foods	
	1.3	Some important food spoilage microorganisms, Spoilage of specific food	
		groups- Milk and dairy products, Meat, poultry and seafood's, Cereal	
		and cereal products	
	1.4	Fruits, vegetables and Canned food products.	
2.0		Food fermentations	11
	2.1	Food Fermentations - Definition and types of fermented food,	
		Microorganisms used in food fermentations	
	2.2	Dairy Fermentations- starter cultures and their types	
	2.3	Concept of probiotics, biotechnological process for food fortification	
	2.4	Prebiotics and oligosaccharides, fermented foods-types, methods of	
		manufacture for vinegar, sauerkraut, tempeh, miso, soya sauce, beer,	
		wine and traditional Indian fermented foods.	
3.0		Food-borne Diseases	11
	3.1	Foodborne Diseases, Types - Food borne infections, food borne	
		intoxications and toxin infections, with common and recent examples.	
	3.2	Control of microorganisms in foods, Principles and methods of	
		preservation, Physical Methods of Food Preservation- Dehydration,	
		Freezing, Cold storage, Heat Treatment, Irradiation	
	3.3	Bio-preservatives i.e Bacteriocins and other natural products	
	3.4	Role of lactic acid bacteria in preservation of food items	
4.0		Food safety laws and Standards	11
	4.1	Food safety Laws and Standards: Food quality & analysis: Pre and Post-	
		harvest factors in food quality	
	4.2	Physical, Chemical and Microbiological factors of quality, proximate	
		analysis of foods, sample and sample preparation in foods.	
	4.3	Food laws: voluntary and mandatory food laws in India.	
	4.4	Food certification agencies.	
		Total	45

References:

- 1. Hotchkiss, N. N., Potter and Joseph, H. Food Science, 5th Edition, Science Technology Publication, 2007.
- 2. M. Shafiur Rahman, Handbook of Food Processing, 2nd Edition, CRC Press Taylor and Francis group, 2007.
- 3. Frazier, W. C., Westhoff, D. C. and Vanitha N. M. Food Microbiology, 5th Edition, McGraw Hill Education Publication, 2004.
- 4. Sivasankar, B. Food Processing and Preservation, Prentice Hall India Learning Private Limited Publication, 2002.
- 5. Avantina Sharma, Textbook of Food Science & Technology, Vol-I & II, 1st Edition, International Book Distributing Company, 2006.

- 6. Cheung, P. C. K., Mehta, B. M. Handbook of Food Chemistry, Ist Edition, Springer-Verlag Berlin Heidelberg, 2015.
- 7. Jay, J. M., Golden, D. A. and Loessner M. J. Modern Food Microbiology (Food Science Text Series), Springer-Verlag New York Inc. Publication, New Delhi, 2000.
- 8. Garbutt, John. Essentials of Food Microbiology, Hodder Arnold H&S Publication, London, 1997.
- 9. Pelczar, M. J., Chan, E. C. S. and Krieg, N. R. Microbiology, 5th Edition, Affiliated East-West Press Private Limited Publication, 1993.
- 10. Lawley, R., Curtis L. and Davis, J. The Food Safety Hazard Guidebook, Royal Society of Chemistry publication, 2004.

SBTTE-454 Laboratory course in Food Biotechnology

- 1. Isolation of microorganism from spoiled food (Dairy products, Meat products, Vegetables & Fruits)
- 2. Isolation of microorganism from traditional Indian fermented foods
- 3. Production and characterization of bacteriocin from probiotic microorganisms.
- 4. Antimicrobial activity of spices and oils against food spoilage causing microorganism
- 5. Microbiological Analysis of Milk: Raw and Pasteurized Milk MBRT, SPC, Coliform, Sterilized Milk / LHT Milk Spore Count.
- 6. Microbiological analysis of Milk Products: Analysis of butter, ice cream, Paneer, Standard plate count, coliform count, yeast and mold count.
- 7. Proximate analysis of dairy & fermented foods

SVECP-551: Publication Ethics

Teaching Scheme

Course	Course Name	Teaching So	cheme (Hrs.)	Credits Assigned			
Code		Theory	Practical	Theory	Practical	Total	
SVECP-551	Publication Ethics	02		02		02	

Assessment Scheme

Course Code	Course Name	Theory				Practical		Total
			CA					
		Test I	Test II	Avg of (T1+T2)/2	ESA	CA	ESA	
SVECP-551	Publication Ethics	10	10	10	40			50

Course pre-requisite:

> General awareness regarding publication basics

Course objectives:

- To know rules, issues, options, and resources for research ethics.
- > To familiarize with various institutional ethics review boards/academic integrity guidelines.
- > To understand the purpose and value of ethical decision-making.
- > To have a positive disposition towards continued learning about research ethics

Course outcomes:

- To have a positive disposition towards continued learning about research philosophy & ethics.
- To know Rules, Regulations, Issues, Options, and Scientific Resources of Research Ethics.
- > To learn the culture of fairness, honesty and integrity in academic communications and to understand the purpose and value of ethical decision-making.
- Avoid wasteful and duplicate publications & encourage original contributions to advance Academic Research and Scholarship.
- Acquiring knowledge & professional competence and expertise about Patents, Copyrights, and other forms of Intellectual Property Rights.
- To promote social good and prevent or mitigate societal hazards through innovative ideas, creativity and research advocacy

Curriculum Details:

Curriculum Details:

Module No.	Unit No.	Торіс	Hrs.
1.0		Publication ethics	
	1.1	Publication ethics: definition, introduction and importance, Best practices/standards setting initiatives and guidelines: COPE, WAME, etc. Conflicts of interest	
	1.2	Publication misconduct: definition, concept, problems that lead to unethical behavior and vice verse, types	08
	1.3	Violation of publication ethics, authorship and contributor ship	
	1.4	Identification of publication misconduct, complaints and appeals. Predatory publishers and journals	
2.0		Open access publishing	
	2.1	Open access publications and initiatives.	
	2.2	SHERPA/RoMEO online resource to check publisher copyright and self- archiving policies	07
	2.3	Software tool to identify predatory publications developed by SPPU	
	2.4	Journal finder/ journal suggestion tools viz. JANE	
3.0		Publication misconduct	
	3.1	Subject specific ethical issues, FFP, authorship	
	3.2	Conflicts of interest	07
	3.3	Complaints and appeals: examples and fraud from India and abroad	U /
	3.4	Use of plagiarism software like Turnitin, Urkund and other open source software tools.	
4.0		Databases and research metrics	
	4.1	Databases: Indexing databases	
	4.2	Citation databases: Web of Science, Scopus, etc.	08
	4.3	Research Metrics: Impact Factor of journal as per journal citation report, SNIP, SJR, IPP, Cite Score.	VO
4.4		Metrics: h-index, g index, i10 index, altmetrics	
		Total	30

References:

- 1. Donna M. Mertens, Pauline E. Ginsberg The Handbook of Social Research Ethics, SAGE (2009).
- 2. Rose Wiles, Bloomsbury What are Qualitative Research Ethics? (2013).
- 3. Robin Levin Penslar, eds, Research Ethics: Cases and Materials, Indiana University Press (1995).
- 4. Gary Comstock, Research Ethics: A Philosophical Guide to the Responsible Conduct of Research, Cambridge University Press (2013)
- 5. Bird, A. Philosophy of Science. Routledge, 2006.

- 6. MacIntyre, Alasdair A Short History of Ethics London, 1967
- 7. P. Chaddah Ethics in Competitive Research: Do not get scooped; do not get plagiarized, 2018
- 8. National Academy of Sciences, National Academy of Engineering and Institute of Medicine. (2009).
- 9. On being a Scientist: A Guide to Responsible Conduct in Research. Third Edition. National Academies Press.
- 10. Resnik, D. B. What is ethics in research & why is it important. National Institute of Environmental Health Sciences, 2018.
 - Retrieved from https www.nichs.nih.gov/research/resources/bioethics/whatis/index.cfm
- 11. Beall, J. Predatory publishers are corrupting open access. Nature, 2012. https://doi.org/10.1038/489179a
- 12. Indian National Science Academy (INSA), Ethics in Science Education, Research and Governance(2019), SBN:978-81-939482-1-7. http://www.insaindia.res.in/pdf/Ethics Book.pdf