# UNIVERSITY VISION AND MISSION

#### VISION

B.S. Abdur Rahman Institute of Science and Technology aspires to be a leader in Education, Training and Research in Engineering, Science, Technology and Management and to play a vital role in the socio-Economic progress of the Country.

#### MISSION

- To blossom into an internationally renowned University
- To empower the youth through quality education and to provide professional leadership
- To achieve excellence in all its endeavors to face global challenges
- To provide excellent teaching and research ambience
- To network with global institutions of excellence, Business, Industry and Research Organizations
- To contribute to the knowledge base through scientific enquiry, Applied research and Innovation

# VISION AND MISSION OF THE SCHOOL OF LIFE SCIENCES

#### VISION

To attain new heights in biotechnology research, shaping life sciences into a premier precision tool for the future for creation of wealth and ensuring social justice-specially for the welfare of the poor.

#### MISSION

The mission of the School of Life Sciences is to maximize the benefits of biotechnology to the University, the nation and the globe by being an excellent quality, comprehensive, multidisciplinary school that supports, coordinates, disseminates and advances biotechnology in the areas of social welfare and entrepreneurship.

# PROGRAMME EDUCATIONAL OBJECTIVES AND OUTCOMES M.Sc (Biotechnology)

#### **PROGRAMME EDUCATIONAL OBJECTIVES**

The course aims to provide an advanced understanding of the core principles and topics of Modern day Biotechnology, and to enable students to acquire a specialized knowledge and understanding of selected aspects by means of a lecture series and a research project. Hence, the main objectives of the program are:

- To provide strong fundamentals of biotechnology and its industrial application.
- To discover in depth knowledge of animal and plant biotechnology, and also broad area of biochemistry, Immunology and molecular biology.
- It will provide the students to develop independent learning skills all biochemical and biotechnology studies.
- This course will provide the students to apply their knowledge and skills in their future professional areas.
- This course will help in contributing to the education of academics which impart its effect for university to play an active role in other advanced studies.

#### **PROGRAMME OUTCOMES**

#### After successfully completing this course, the student should be able to:

- Understand the basic knowledge and concepts of biotechnology and other related areas.
- Understand the ability to apply their knowledge for practical which they can conduct independently.
- Apply their knowledge in other advanced subject area like nanobiotechnology, immunotechnology, and animal and plant biotechnology for the betterment and advancement of their professional career.
- Learn the theoretical and practical exposure to the basic and the advanced fields of biotechnology.



# REGULATIONS 2013 FOR M.TECH. DEGREE PROGRAMMES (WITH AMENDMENTS INCORPORATED TILL JUNE 2015)

### B.S. ABDUR RAHMAN UNIVERSITY, CHENNAI 48. REGULATIONS -2013 FOR M.TECH / MCA / M.Sc. DEGREE PROGRAMMES

(With amendments incorporated till June 2015)

#### 1.0 PRELIMINARY DEFINITIONS AND NOMENCLATURE

In these Regulations, unless the context otherwise requires

- i) **"Programme"** means Post Graduate Degree Programme (M.Tech./ MCA / M.Sc.)
- ii) **"Course"** means a theory or practical subject that is normally studied in a semester, like Applied Mathematics, Structural Dynamics, Computer Aided Design, etc.
- iii) "University" means B.S.Abdur Rahman University, Chennai, 600048.
- iv) **"Institution"** unless otherwise specifically mentioned as an autonomous or off campus institution means B.S.Abdur Rahman University.
- v) "Academic Council" means the Academic Council of this University.
- vi) **"Dean (Academic Affairs)"** means Dean (Academic Affairs) of B.S.Abdur Rahman University.
- vii) **"Dean (Student Affairs)"** means Dean(Student Affairs) of B.S.Abdur Rahman University.
- viii) **"Controller of Examinations"** means the Controller of Examinations of B.S.Abdur Rahman University who is responsible for conduct of examinations and declaration of results.

# 2.0 PROGRAMMES OFFERED, MODE OF STUDY AND ADMISSION REQUIREMENTS

#### 2.1 P.G. Programmes Offered

The various P.G. Programmes and their modes of study are as follows:

Degree	Mode of Study
M.Tech.	Full Time
M.Tech.	Part Time – Day / Evening
M.C.A.	Full Time
M. Sc.	Full Time
M. Sc.	Full Time

# 2.2 MODES OF STUDY

### 2.2.1 Full-time

Students admitted under "Full-Time" shall be available in the Institution during the complete working hours for curricular, co-curricular and extra-curricular activities assigned to them.

**2.2.2** A full time student, who has completed all non-project courses desiring to do the Projectwork in part-time mode for valid reasons, shall apply to the Dean (Academic Affairs) through the Head of the Department, if the student satisfies the clause 2.3.4 of this Regulation. Permission may be granted based on merits of the case. Such conversion is not permitted in the middle of a semester.

# 2.2.3 Part time - Day time

In this mode of study, the students are required to attend classes for the courses registered along with full time students.

### 2.2.4 Part time - Evening

In this mode of study, the students are required to attend normally classes in the evening and on Saturdays, if necessary.

**2.2.5** A part time student is not permitted to convert to full time mode of study.

# 2.3 ADMISSION REQUIREMENTS

- **2.3.1** Students for admission to the first semester of the Master's Degree Programme shall be required to have passed the appropriate degree examination of this University as specified in the Table shown for eligible entry qualifications for admission to P.G. programmes or any other degree examination of any University or authority accepted by this University as equivalent thereto.
- **2.3.2** Eligibility conditions for admission such as class obtained, number of attempts in the qualifying examination and physical fitness will be as prescribed by this Institution from time to time.
- **2.3.3** All part-time students should satisfy other conditions regarding experience, sponsorship etc., which may be prescribed by this Institution from time to time.

- **2.3.4** A student eligible for admission to M.Tech. Part Time / Day Time programme shall have his/her permanent place of work within a distance of 65km from the campus of this Institution.
- 2.3.5 Student eligible for admission to M.C.A under lateral entry scheme shall be required to have passed three year degree in B.Sc (Computer Science) / B.C.A / B.Sc (Information Technology)

#### 3.0 DURATION AND STRUCTURE OF THE P.G. PROGRAMME

**3.1** The minimum and maximum period for completion of the P.G. Programmes are given below:

Programme	Min.No.of Semesters	Max.No.of Semesters
M.Tech. (Full Time)	4	8
M.Tech.(Part Time)	6	12
M.C.A. (Full Time)	6	12
M.C.A. (Full Time) – (Lateral Entry)	4	8
M.Sc. (Full Time)	4	8

- 3.2 The PG. programmes consist of the following components as prescribed in the respective curriculum
  - i. Core courses
  - ii. Elective courses
  - iii. Project work / thesis / dissertation
  - iv. Laboratory Courses
  - v. Case studies
  - vi. Seminars
  - vii. Industrial Internship
- **3.3** The curriculum and syllabi of all PG. programmes shall be approved by the Academic Council of this University.
- **3.4** The minimum number of credits to be earned for the successful completion of the programme shall be specified in the curriculum of the respective specialization of the P.G. programme.
- **3.5** Each academic semester shall normally comprise of 80 working days. Semester-end examinations will follow immediately after the last working day.

SI. No.	Name of the Department	P.G. Programmes offered	Qualifications for admission
01.	Civil Engineering	M.Tech. (Structural Engineering)	B.E / B.Tech. (Civil Engineering) / (Structural Engineering)
01.	Civil Engineering	M.Tech. (Construction Engineering and Project Management)	B.E. / B.Tech. (Civil Engineering) / (Structural Engineering)
02.	Mechanical Engineering	M.Tech. (Manufacturing Engineering)	B.E. / B.Tech. (Mechanical / Auto / Manufacturing / Production / Industrial Mechatronics / Metallurgy / Aerospace /Aeronautical / Material Science / Marine Engineering)
03.	Polymer Technology	M.Tech. (Polymer Technology)	B.E./ B.Tech. degree Mech./Production. Polymer Science or Engg or Tech / Rubber Tech / M.Sc (Polymer Sc./ Chemistry Appl. Chemistry)
0.4	Electrical and	M.Tech. (Power Systems Engg)	B.E / B.Tech (EEE / ECE / E&I / I&C / Electronics / Instrumentation)
04.	Electronics Engineering	M.Tech. (Power Electronics & Drives)	B.E / B.Tech (EEE / ECE / E&I / I&C / Electronics / Instrumentation)
05.	Electronics and Communication	M.Tech. (Communication Systems)	B.E / B.Tech (EEE/ ECE / E&I / I&C / Electronics / Instrumentation)
	Engineering	M.Tech.(VLSI and Embedded Systems)	B.E./ B.Tech. in ECE / Electronics / EIE
06.	ECE Department jointly with Physics Dept	M.Tech. (Optoelectronics and Laser Technology)	B.E./B.Tech. (ECE / EEE / Electronics / EIE / ICE) M.Sc (Physics / Materials Science / Electronics / Photonics)
07.	Electronics and Instrumentation Engineering	M.Tech. (Electronics and Instrumentation Engineering)	B.E./ B.Tech. (EIE/ICE/Electronics/ECE/ EEE)
		M.Tech. (Computer Science and Engineering)	B.E. /B.Tech. (CSE/IT/ECE/EEE/EIE/ICE/ Electronics / MCA)
08.		M.Tech. (Software Engineering)	B.E. / B.Tech. (CSE / IT) MCA
	Engineering	M.Tech (Network Security)	B.E. /B.Tech. (CSE/IT/ECE/EEE/EIE/ICE Electronics / MCA)
		M.Tech (Computer and Predictive Analytics)	B.E. /B.Tech. (CSE/IT/ECE/EEE/EIE/ICE Electronics / MCA)
09	InformationTechnology	M.Tech. (Information Technology)	B.E /B.Tech. (IT/CSE/ECE/EEE/EIE/ICE/ Electronics) MCA
		M.Tech. (Information Security & Digital Forensics)	B.E /B.Tech. (IT/CSE/ECE/EEE/EIE/ICE/ Electronics) MCA
10		M.C.A.	Bachelor Degree in any discipline with Mathematics as one of the subjects (or) Mathematics at +2 level
	Computer Applications	M.Tech. (Systems Engineering and Operations Research)	BE / B.Tech. (Any Branch) or M.Sc., (Maths / Physics / Statistics / CS / IT / SE) or M.C.A.
		M.Tech. (Data & Storage Management)	BE / B.Tech. (Any Branch) or M.Sc., (Maths / Physics / Statistics / CS / IT / SE) or M.C.A.

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ELIG	ELIGIBLE ENTRY QUALIFICATIONS FOR ADMISSION TO P.G. PROGRAMMES					
SI. No.	Name of the Department	P.G. Programmes offered	Qualifications for admission			
		M.C.A.	Bachelor Degree in any discipline with Mathematics as one of the subjects (or) Mathematics at +2 level			
10	Computer Applications	M.C.A. (Full Time) – (Lateral Entry)	B.Sc Computer Science / B.Sc Information Technology / B.C.A			
		M.Tech. (Systems Engineering and Operations Research)	BE / B.Tech. (Any Branch) or M.Sc., (Maths / Physics / Statistics / CS / IT /			
		M.Tech. (Data & Storage Management)	SE) or M.C.A.			
11	Mathematics	M.Sc. (Actuarial Science)	Any Degree with Mathematics / Statistics as one of the Subjects of Study.			
		M.Sc. Mathematics	B.Sc. (Mathematics)			
12	Physics	M.Sc.(Physics)	B.Sc.(Physics / Applied Science / Electronics / Electronics Science /			
		M.Sc. (Material Science)	Electronics / Electronics Science / Electronics & Instrumentation)			
13	Chemistry	M.Sc.(Chemistry)	B.Sc (Chemistry) of B.Sc. (Applied Science)			
		M.Sc. Molecular Biology & Biochemistry	B.Sc. in any branch of Life Sciences			
		M.Sc. Genetics	B.Sc. in any branch of Life Sciences			
14	Life Sciences	M.Sc. Biotechnology	B.Sc. in any branch of Life Sciences			
		M.Sc. Microbiology	B.Sc. in any branch of Life Sciences			
		M.Sc. Bioscience	B.Sc. in any branch of Life Sciences			
		M.Tech. Biotechnology	B.Tech. (Biotechnology / Chemical Engineering) / M.Sc. in any branch of Life Sciences			

**3.6** The curriculum of PG programmes shall be so designed that the minimum prescribed credits required for the award of the degree shall be within the limits specified below:

Programme	Minimum prescribed credit range	
M.Tech.	75 to 85	
M.C.A.	120 to 130	
M.Sc.	75 to 85	

- **3.7** Credits will be assigned to the courses for all P.G. programmes as given below:
  - \* One credit for one lecture period per week
  - \* One credit for one tutorial period per week
  - \* One credit each for seminar/practical session/project of two or three periods per week
  - \* One credit for two weeks of industrial internship.
- **3.8** The number of credits registered by a student in non-project semester and project semester should be within the range specified below:

P.G. Programme	Non-project Semester	Project semester		
M.Tech. (Full Time)	15 to 29	12 to 20		
M.Tech. (Part Time)	6 to 18	12 to 16		
M.C.A. (Full Time)	15 to 29	12 to 20		
M.Sc. (Full Time)	15 to 25	12 to 20		

- **3.9** The electives from the curriculum are to be chosen with the approval of the Head of the Department.
- **3.10** A student may be permitted by the Head of the Department to choose electives offered from other PG programmes either within the Department or from other Departments up to a maximum of three courses during the period of his/her study, provided the Heads of the Departments offering such courses also agree.
- **3.11** To help the students to take up special research areas in their project work and to enable the department to introduce courses in latest/emerging areas in the curriculum, "Special Electives" may be offered. A student may be permitted to register for a "Special Elective" up to a maximum of three credits during the period of his/her study, provided the syllabus of this course is recommended by the Head of the Department and approved by the Chairman, Academic Council before the commencement of the semester, in which the special elective course is offered. Subsequently, such course shall be ratified by the Board of Studies and Academic Council.

- **3.12** The medium of instruction, examination, seminar and project/thesis/ dissertation reports will be English.
- **3.13** Industrial internship, if specified in the curriculum shall be of not less than two weeks duration and shall be organized by the Head of the Department.

#### 3.14 PROJECT WORK/THESIS/DISSERTATION

- **3.14.1** Project work / Thesis / Dissertation shall be carried out under the supervision of a qualified teacher in the concerned Department.
- **3.14.2** A student may however, in certain cases, be permitted to work for the project in an Industrial/Research Organization, on the recommendation of the Head of the Department. In such cases, the project work shall be jointly supervised by a faculty of the Department and an Engineer / Scientist from the organization and the student shall be instructed to meet the faculty periodically and to attend the review committee meetings for evaluating the progress.
- **3.14.3** Project work / Thesis / Dissertation (Phase II in the case of M.Tech.) shall be pursued for a minimum of 16 weeks during the final semester, following the preliminary work carried out in Phase-1 during the previous semester.
- **3.14.4** The Project Report/Thesis / Dissertation report / Drawings prepared according to approved guidelines and duly signed by the supervisor(s) and the Head of the Department shall be submitted to the concerned department.
- **3.14.5** The deadline for submission of final Project Report / Thesis / Dissertation is within 30 calendar days from the last working day of the semester in which Project / Thesis / Dissertation is done.
- **3.14.6** If a student fails to submit the Project Report / Thesis / Dissertation on or before the specified deadline he / she is deemed to have not completed the Project Work / Thesis / dissertation and shall re-register the same in a subsequent semester.
- **3.14.7** A student who has acquired the minimum number of total credits prescribed in the Curriculum for the award of Masters Degree will not be permitted to enroll for more courses to improve his/her cumulative grade point average (CGPA).
- 4.0 CLASS ADVISOR AND FACULTY ADVISOR
- 4.1 CLASS ADVISOR

A faculty member will be nominated by the HOD as Class Advisor for the whole class.

He/she is responsible for maintaining the academic, curricular and cocurricular records of all students throughout their period of study.

#### 4.2 FACULTY ADVISOR

To help the students in planning their courses of study and for general counseling on the academic programme, the Head of the Department of the students will attach a certain number of students to a faculty member of the department who shall function as Faculty Advisor for the students throughout their period of study. Such Faculty Advisor shall offer advice to the students on academic and personal matters, and guide the students in taking up courses for registration and enrolment every semester.

#### 5.0 CLASS COMMITTEE

- **5.1** Every class of the PG Programme will have a Class Committee constituted by the Head of the Department as follows:
  - i. Teachers of all courses of the programme
  - ii. One senior faculty preferably not offering courses for the class, as Chairperson.
  - iii. Minimum two students of the class, nominated by the Head of the Department.
  - iv. Class Advisor / Faculty Advisor of the class Ex-Officio Member
  - v. Professor in-charge of the PG Programme Ex-Officio Member.
- **5.2** The Class Committee shall be constituted by the respective Head of the Department of the students.
- **5.3** The basic responsibilities of the Class Committee are to review periodically the progress of the classes to discuss problems concerning curriculum and syllabi and the conduct of classes. The type of assessment for the course will be decided by the teacher in consultation with the Class Committee and will be announced to the students at the beginning of the semester. Each Class Committee will communicate its recommendations to the Head of the Department and Dean (Academic Affairs). The class committee, without the student members, will also be responsible for finalization of the semester results and award of grades.

**5.4** The Class Committee is required to meet at least thrice in a semester, first within a week of the commencement of the semester, second, after the first assessment and the third, after the semester-end examination to finalize the grades.

# 6.0 COURSE COMMITTEE

Each common theory course offered to more than one group of students shall have a "Course Committee" comprising all the teachers teaching the common course with one of them nominated as Course coordinator. The nomination of the Course coordinator shall be made by the Head of the Department / Dean (Academic Affairs) depending upon whether all the teachers teaching the common course belong to a single department or to several departments. The Course Committee shall meet as often as possible and ensure uniform evaluation of the tests and arrive at a common scheme of evaluation for the tests. Wherever it is feasible, the Course Committee may also prepare a common question paper for the test(s).

# 7.0 REGISTRATION AND ENROLMENT

- **7.1** For the first semester every student has to register and enroll for all the courses.
- **7.2** For the subsequent semesters registration for the courses will be done by the student during a specified week before the semester-end examination of the previous semester. The curriculum gives details of the core and elective courses, project and seminar to be taken in different semester with the number of credits. The student should consult his/her Faculty Adviser for the choice of courses. The Registration form shall be filled in and signed by the student and the Faculty Adviser.
- **7.3** From the second semester onwards all students shall pay the prescribed fees and enroll on a specified day at the beginning of a semester.
- 7.4 A student will become eligible for enrolment only if he/she satisfies clause 9 and in addition he/she is not debarred from enrolment by a disciplinary action of the Institution. At the time of enrolment a student can drop a course registered earlier and also substitute it by another course for valid reasons with the consent of the Faculty Adviser. Late enrolment will be permitted on payment of a prescribed fine up to two weeks from the date of commencement of the semester.

- **7.5** Withdrawal from a course registered is permitted up to one week from the date of the completion of the first assessment test.
- **7.6** Change of a course within a period of 15 days from the commencement of the course, with the approval of Dean (Academic Affairs), on the recommendation of the HOD, is permitted.
- **7.7** Courses withdrawn will have to be taken when they are offered next if they belong to the list of core courses.
- 7.8 A student should have registered for all preceding semesters before registering for a particular semester.

#### 8.0 TEMPORARY BREAK OF STUDY FROM THE PROGRAMME

A student may be permitted by the Dean (Academic Affairs) to avail temporary break of study from the programme up to a maximum of two semesters for reasons of ill health or other valid grounds. Such student has to rejoin only in the same semester from where he left. However the total duration for completion of the programme shall not exceed the prescribed maximum number of semesters (vide clause 3.1).

# 9.0 MINIMUM REQUIREMENTS TO REGISTER FOR PROJECT / THESIS / DISSERTATION

**9.1** A student is permitted to register for project semester, if he/she has earned the minimum number of credits specified below:

Programme	Minimum No. of credits to be earned to enroll for project semester
M.Tech. (Full time)	18 (III semester)
M.Tech. (Part time)	18 (V semester)
M.C.A. (Full time)	45 (V semester)
M.C.A. (Full time) – (Lateral Entry)	22 (V semester)
M.Sc.(Full time)	30 (IV semester) if project is in IV semester 18 (III semester) if project is in III semester
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**9.2** If the student has not earned minimum number of credits specified, he/she has to earn the required credits, at least to the extent of minimum credits specified in clause 9.1 and then register for the project semester.

#### 10.0 DISCIPLINE

- **10.1** Every student is required to observe discipline and decorous behavior both inside and outside the campus and not to indulge in any activity, which will tend to bring down the prestige of the Institution.
- **10.2** Any act of indiscipline of a student reported to the Head of the Institution will be referred to a Discipline and Welfare Committee for taking appropriate action.
- **10.3** Every student should have been certified by the HOD that his / her conduct and discipline have been satisfactory.

#### 11.0 ATTENDANCE

- **11.1** Attendance rules for all Full Time Programme and Part time day Time Programmes are given in the following sub-clause.
- **11.2** Ideally every student is expected to attend all classes and earn 100% attendance in the contact periods of every course, subject to a maximum relaxation of 25% for genuine reasons like on medical grounds, representing the University in approved events etc., to become eligible to appear for the semester-end examination in that course, failing which the student shall be awarded "I" grade in that course. If the course is a core course, the student should register for and repeat the course when it is offered next. If the course is an elective, either he/she can register and repeat the same elective or can register for a new elective.
- **11.3** The students who have not attended a single hour in all courses in a semester and awarded 'l' grade are not permitted to write the examination and also not permitted move to next higher semester. Such students should repeat all the courses of the semester in the next Academic year.

#### 12.0 SUMMER TERM COURSES

**12.1** Summer term courses may be offered by a department on the recommendation of the Departmental Consultative Committee and approved by the Dean (Academic Affairs). No student should register for more than three courses during a summer term.

- **12.2** Summer term courses will be announced by the Head of the department at the end of the even semester before the commencement of the end semester examinations. A student will have to register within the time stipulated in the announcement. A student has to pay the fees as stipulated in the announcement.
- **12.3** The number of contact hours and the assessment procedure for any course during summer term will be the same as those during regular semesters.

Students with U grades will have the option either to write semester end arrears exam or to redo the courses during summer / regular semesters, if they wish to improve their continuous assessment marks subject to the approval of the Head of the department.

**12.4** Withdrawal from a summer term course is not permitted. No substitute examination will be conducted for the summer term courses.

#### 13.0 ASSESSMENTS AND EXAMINATIONS

**13.1** The following rule shall apply to the full-time and part-time PG programmes (M.Tech./ M.C.A. / M.Sc.)

For lecture-based courses, normally a minimum of two assessments will be made during the semester. The assessments may be combination of tests and assignments. The assessment procedure as decided in the Class Committee will be announced to the students right from the beginning of the semester by the course teacher.

- **13.2** There shall be one examination of three hours duration, at the end of the semester, in each lecture based course.
- **13.3** The evaluation of the Project work will be based on the project report and a Viva-Voce Examination by a team consisting of the supervisor concerned, an Internal Examiner and External Examiner to be appointed by the Controller of Examinations.
- **13.4** At the end of industrial internship, the student shall submit a certificate from the organization and also a brief report. The evaluation will be made based on this report and a Viva-Voce Examination, conducted internally by a Departmental Committee constituted by the Head of the Department.

#### 14.0 WEIGHTAGES

**14.1** The following shall be the weightages for different courses:

<ul> <li>(i) Lecture based course</li> <li>Two continuous assessments</li> <li>Semester-end examination</li> </ul>	- 50% - 50%
<ul> <li>(ii) Laboratory based courses</li> <li>Laboratory work assessment</li> <li>Semester-end examination</li> </ul>	- 75% - 25%
(iii) Project work Periodic reviews Evaluation of Project Report by External Examiner Viva-Voce Examination	- 50% - 20% - 30%

- **14.2** Appearing for semester end examination for each course (Theory and Practical) is mandatory and a student should secure a minimum of 40% marks in semester end examination for the successful completion of the course.
- **14.3** The markings for all tests, tutorial, assignments (if any), laboratory work and examinations will be on absolute basis. The final percentage of marks is calculated in each course as per the weightages given in clause 13.1.

#### 15.0 SUBSTITUTE EXAMINATION

- **15.1** A student who has missed for genuine reasons any one of the three assessments including semester-end examination of a course may be permitted to write a substitute examination. However, permission to take up a substitute examination will be given under exceptional circumstances, such as accident or admissions to a hospital due to illness, etc.
- **15.2** A student who misses any assessment in a course shall apply in a prescribed form to the Dean (Academic Affairs) through the Head of the department within a week from the date of missed assessment. However the substitute tests and examination for a course will be conducted within two weeks after the last day of the semester-end examinations.

#### 16.0 COURSEWISE GRADING OF STUDENTS AND LETTER GRADES

**16.1** Based on the semester performance, each student is awarded a final letter grade at the end of the semester in each course. The letter grades and the corresponding grade points are as follows, but grading has to be relative grading

Letter grade	Grade points
S	10
A	9
В	8
С	7
D	6
E	5
U	0
W	-
I	-
AB	-

Flexible range grading system will be adopted

- "W" denotes withdrawal from the course.
- "I" denotes inadequate attendance and hence prevention from semesterend examination
- "U" denotes unsuccessful performance in a course.

"AB" denotes absent for the semester end examination

- **16.2** A student is considered to have completed a course successfully if he / she secure five grade points or higher. A letter grade 'U' in any course implies unsuccessful performance in that course.
- **16.3** A course successfully completed cannot be repeated for any reason.

### 17.0 AWARD OF LETTER GRADE

- **17.1** A final meeting of the Class Committee without the student member(s) will be convened within ten days after the last day of the semester end examination. The letter grades to be awarded to the students for different courses will be finalized at the meeting.
- **17.2** After finalization of the grades at the class committee meeting the Chairman will forward the results to the Controller of Examinations, with copies to Head of the Department and Dean (Academic Affairs).

### 18.0 DECLARATION OF RESULTS

- **18.1** After finalization by the Class Committee as per clause 16.1 the Letter grades awarded to the students in the each course shall be announced on the departmental notice board after duly approved by the Controller of Examinations.
- **18.2** In case any student feels aggrieved about the results, he/she can apply for revaluation after paying the prescribed fee for the purpose, within one week from the announcement of results.

A committee will be constituted by the concerned Head of the Department comprising of the Chairperson of the concerned Class Committee (Convener), the teacher concerned and a teacher of the department who is knowledgeable in the concerned course. If the Committee finds that the case is genuine, it may jointly revalue the answer script and forward the revised marks to the Controller of Examinations with full justification for the revision, if any.

**18.3** The "U" and "AB" grade once awarded stays in the grade sheet of the students and is not deleted when he/she completes the course successfully later. The grade acquired by the student later will be indicated in the grade sheet of the appropriate semester.

# 19.0 COURSE REPETITION AND ARREARS EXAMINATION

**19.1** A student should register to re-do a core course wherein "I" or "W" grade is awarded. If the student is awarded "I" or "W" grade in an elective course either the same elective course may be repeated or a new elective course may be taken.

- **19.2** A student who is awarded "U" or "AB" grade in a course shall write the semester-end examination as arrear examination, at the end of the next semester, along with the regular examinations of next semester courses.
- **19.3** A student who is awarded "U" or "AB" grade in a course will have the option of either to write semester end arrear examination at the end of the subsequent semesters, or to redo the course whenever the course is offered. Marks earned during the redo period in the continuous assessment for the course, will be used for grading along with the marks earned in the end-semester (re-do) examination.
- **19.4** If any student obtained "U" or "AB" grade, the marks earned during the redo period for the continuous assessment for that course will be considered for further appearance as arrears.
- **19.5** If a student with "U" or "AB" grade prefers to redo any particular course fails to earn the minimum 75% attendance while doing that course, then he/she will not be permitted to write the semester end examination and his / her earlier 'U' grade and continuous assessment marks shall continue.

#### 20.0 GRADE SHEET

- **20.1** The grade sheet issued at the end of the semester to each student will contain the following:
  - (i) the credits for each course registered for that semester.
  - (ii) the performance in each course by the letter grade obtained.
  - (iii) the total credits earned in that semester.
  - (iv) the Grade Point Average (GPA) of all the courses registered for that semester and the Cumulative Grade Point Average (CGPA) of all the courses taken up to that semester.
- **20.2** The GPA will be calculated according to the formula

$$GPA = \frac{\sum_{i=1}^{n} (C_i) (GPi)}{\sum_{i=1}^{n} C_i} \quad Where \ n = number \ of \ courses$$

where Ci is the number of credits assigned for ith course

GP<sub>i</sub> - Grade point obtained in the i<sup>th</sup> course

For the cumulative grade point average (CGPA) a similar formula is used except that the sum is over all the courses taken in all the semesters completed up to the point of time.

#### 'I' and 'W' grades will be excluded for GPA calculations.

#### 'U', 'AB' 'I' and 'W' grades will be excluded for CGPA calculations.

**20.3** Classification of the award of degree will be as follows:

CGPA	Classification
8.50 and above, having completed all courses in first appearance	First class with Distinction
6.50 and above, having completed within a period of 2 semesters beyond the programme period	First Class
All others	Second Class

However, to be eligible for First Class with Distinction, a student should not have obtained U or I grade in any course during his/her study and should have completed the PG Programme within a minimum period covered by the minimum duration (clause 3.1) plus authorized break of study, if any (clause 8). To be eligible for First Class, a student should have passed the examination in all courses within the specified minimum number of semesters reckoned from his/her commencement of study plus two semesters. For this purpose, the authorized break of study will not be counted. The students who do not satisfy the above two conditions will be classified as second class. For the purpose of classification, the CGPA will be rounded to two decimal places. For the purpose of comparison of performance of students and ranking, CGPA will be considered up to three decimal places.

# 21.0 ELIGIBILITY FOR THE AWARD OF THE MASTERS DEGREE

- **21.1** A student shall be declared to be eligible for the award of the Masters Degree, if he/she has:
  - i) successfully acquired the required credits as specified in the Curriculum corresponding to his/her programme within the stipulated time,
  - ii) no disciplinary action is pending against him/her.

**21.2** The award of the degree must be approved by the University.

### 22.0 POWER TO MODIFY

Notwithstanding all that have been stated above, the Academic Council has the right to modify any of the above regulations from time to time.

# CURRICULUM & SYLLABI FOR M.Sc. BIOTECHNOLOGY (FOUR SEMESTERS / FULL TIME)

#### CURRICULUM

#### **SEMESTER I**

SI. No.	Course Code	Course Title	L	т	Ρ	С
1	LSB6101	Advanced Biochemistry	4	0	0	4
2	LSB6102	Cell & Molecular Biology	3	0	0	3
3	LSB6103	Biostatistics	3	0	0	3
4	LSB6104	Immunology	3	0	0	3
5	LSB6105	Biomedical Instrumentation	3	0	0	3
6	LSB6106	Microbiology	3	0	0	3
7	LSB6107	Biochemistry Lab	0	0	3	1
8	LSB6108	Cell Biology Lab	0	0	3	1
9	LSB6109	Immunology Lab	0	0	3	1
				Credits 2		22
		SEMESTER II				
SI. No.	Course Code	Course Title	L	т	Ρ	С
1	LSB6221	Stem cells in Health care	3	0	0	3
2	LSB6222	Drug Design and Development	3	0	0	3
3	LSB6203	Genetic Engineering	3	0	0	3
4	LSB6204	Computational Biology	3	0	0	3
5		Elective I	3	0	0	3
6	LSB6205	Computational Biology Lab	0	0	3	1
7	LSB6206	Genetic Engineering Lab	0	0	3	1
8		Mini Dusis st	0	~	~	1
	LSB6223	Mini Project	0	0	3	I

SEMESTER III							
SI. No.	Course Code	Course Title		L	т	Ρ	С
1	LSB7121	Protein Engineering		3	0	0	3
2	LSB7102	Plant & Medical Biotechnology		3	0	0	3
3		Elective II		3	0	0	3
4		Elective III		3	0	0	3
5	LSB7122	Mini Project		3	0	0	6
				C	Cred	lits	18
		SEMESTER IV					
SI. No.	Course Code	Course Title		L	т	Ρ	С
1	LSB7221	Project work		0	0	20	18
				C	red	its	18
			TOTAL	CR	EDI	ΤS	76

ELECTIVES										
Course Code	Electives I	L	Т	Ρ	С					
LSBY021	Bioenterpreneurship	3	0	0	3					
LSBY022	IPR and Patent Law	3	0	0	3					
LSBY023	Biosafety & Bioethics	3	0	0	3					
Electives II										
LSBY024	Molecular Diagnostics	3	0	0	3					
LSBY025	Food Process technology	3	0	0	3					
LSBY026	Animal Biotechnology	3	0	0	3					
Electives III										
LSBY027	Environmental Biotechnology	3	0	0	3					
LSBY028	Bionanotechnology	3	0	0	3					
LSBY029	Industrial Biotechnology	3	0	0	3					

#### SYLLABUS

#### SEMESTER I

LSB6101

ADVANCED BIOCHEMISTRY

L T P C 3 0 0 3

#### **OBJECTIVES:**

This course aims to develop in the students' mind a concept regarding

- The diversity of metabolic processes occurring in biological system.
- The effect of the structural and functional role of the enzymes governing the metabolic processes.
- Importance of the metabolic pathways in maintaining homeostasis in biological system.
- The clinical implications of the metabolic pathway.

# MODULE I AMINO ACIDS & PROTEIN: STRUCTURE AND FUNCTIONS

Amino acids- Classification, structure and function, proteins- primary, secondary, tertiary and quaternary structure, Ramachandran plot, super secondary structures and helix loop.

#### MODULE II ENZYMOLOGY

Classification of enzymes. How do enzymes work: activation energy, substrate specificity. Enzyme-substrate interaction: Lock and Key mechanism and Induced Fit mechanism. Effect of temperature and pH on enzyme action. Enzyme Kinetics: Michaelis-Menten Equation, Km, Measurement of Km and Vmax (Lineweaver-Burk equation). Kinetics of multisubstrate reaction: Sequential reactions and ping-pong reactions. Enzyme inhibition: reversible (competitive, uncompetitive and mixed) and irreversible. Allosteric regulation of enzyme activity.Multienzyme complex and multifunctional enzymes.

# MODULE III ENERGY PRODUCTION AND OXIDATIVE PHOSPHORYLATION

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Introduction to metabolism: Anabolism, catabolism, metabolic pathways. Characteristics of metabolic pathways

Glycolysis: glycolytic pathway. Molecular mechanism of action of the glycolytic enzymes. Energetic of glycolysis. Glycolysis and cancer biology—Warburg Hypothesis and PET scanning. Fates of Pyruvate under anaerobic conditions: alcohol and lactic acid fermentation. Importance of lactic acid fermentation.

TCA Cycle: Formation of Acetyl CoA and reactions of citric acid cycle. Molecular mechanism of pyruvate dehydrogenase complex and enzymes involved in Kreb's cycle.Energetic of TCA cycle and substrate level phosphorylation.

Lipid metabolism: Hormonal regulation of the mobilization of triglycerides from adiposities. Transport of fatty acid into mitochondria.Beta oxidation of saturated fatty acid (both even and odd).Regulation.Energetic.

Electron Transport Chain: structure and function of Electron carriers: Complex I—V. Passage of electrons from complex I to IV. Mitchell's chemiosmotic hypothesis and proton gradient.Structure of complex V or ATP synthase, Catalytic sites of ATP synthesis. Mechanism of ATP generation by Boyer's binding change mechanism—rotational catalysis. Energetic of ATP synthesis and efficiency of ATP synthase.

#### MODULE IV METABOLIC INTERRELATIONSHIP

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Starve-Fed cycle.Glucose homeostasis. Switching of metabolism of liver between starve and fed cycle. Metabolic relationship of tissues in various nutritional and hormonal states—insulin resistance, diabetes, exercise, pregnancy, lactation, stress, liver and renal diseases, alcohol consumption.

#### MODULE V REGULATORY MECHANISMS OF METABOLIC PATHWAYS 7

Feed back inhibition by allosteric modulation of enzymes. Covalent modifications of enzymes. Isozymes. Propetolytic cleavage. Regulationg the amount of enzyme—regulation gene expression in prokaryotes and eukaryotes.

#### Total Hours: 45

#### **REFERENCE:**

- 1. Nelson D.L, Cox M. M. Lehninger's Principle of Biochemistry. 5<sup>th</sup> Ed.,W. H. Freeman, 2008.
- 2. Biochemistry by Lubert Stryer 7<sup>th</sup> ed. W. H. Freeman & Company.

 Textbook of Biochemistry with Clinical Correlations. 4<sup>th</sup> Ed. Thomas M. Devlin. Wiley-Liss publication. 1997.

#### OUTCOMES:

At the completion of the course the student will develop an understanding about the

- Various metabolic processes occurring in biological system and their role in governing homeostasis and normal physiology.
- The importance of enzymes as a regulatory molecule in metabolism.
- The interrelationship of metabolic pathways different physiological conditions.
- The role of liver in regulating metabolism.

LSB6102	CELL & MOLECULAR BIOLOGY	L	т	Ρ	С
		3	0	0	3

#### **OBJECTIVES:**

- To get overview of classes of cells and structural and function aspects of plasma membrane and cell organelle.
- To develop skill to understand molecular aspects of cell cycle and cell division.
- To get familiar with transcription and translation in details.
- To understand the signaling pathways in cell functioning

#### MODULE I **INTRODUCTION TO CELL**

Basic properties of cell, Different classes of cell: Prokaryotic, animal and plant cell.Plasma membrane- structure and function, Chemical composition of membranes, membrane lipids and proteins,fluid mosaic model, Transport across the membranes- diffusion, osmosis, facilitated diffusion, passive and active transport; membrane potential and nerve impulses.

#### MODULE II MEMBRANE TRANSPORT

Endoplasmic Reticulum, Golgi complex-glycosylation, Vesicle transport-COPI and COPII; Lysosomes-autophagy;Endocytic pathway- endocytosis and phagocytosis, transport of proteins into peroxisomes, mitochondria and chloroplast;

#### MODULE III ENERGY CONVERSION

Structure of mitochondria and organization of respiratory chain; Proton Pump and ATP generation in mitochondria; Structure of chloroplast and Photosynthesis, photorespiration; Genetic system of mitochondria and chloroplast.

#### MODULE IV BASIC GENETIC MECHANISMS

The structure and function of DNA, DNA packaging and Chromosomes, chromatin structure and function, DNA replication mechanisms, DNA damage and repair and homologous recombination and transposable elements, Telomeres, telomerase and end replication. Role of telomerase in aging and cancer.

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#### MODULE V TRANSCRIPTION AND TRANSLATION

Transcription- Prokaryotic and eukaryotic Transcription- RNA polymerasesgeneral and specific transcription factors- regulatory elements- mechanism of transcription, Transcription termination Post transcriptional modificationsplicing- editing- nuclear export of mRNA- mRNA stability; Translation- Genetic code, Mechanism of initiation-elongation and termination- Regulation of translation.

#### **Total Hours:45**

#### REFERENCES

- 1. Molecular Biology of Cell by Alberts et.al. John Wiley & Sons, 6<sup>th</sup> Ed, 2015
- 2. The Cell by Cooper. ASM Press, 4<sup>th</sup> Ed, 2007
- 3. Cell and Molecular Biology by Karp. John Wiley & Sons, 7<sup>th</sup> Ed, 2013
- 4. Lodish H. F.Cell and Molecular Biology. W.H. Freeman & Co Ltd, 7<sup>th</sup> Ed, 2000.

#### OUTCOMES:

• On the completion of the above objectives student will be able to get the overview of classes of cells and structural and function aspects of plasma membrane and cell organelle. They can develop skill to understand molecular aspects of cell cycle, cell division, transcription and translation.

LSB6103

#### BIOSTATISTICS

### L T P C 3 0 0 3

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#### **OBJECTIVES:**

- To learn about the concepts of biostatistics.
- To handle and analyses bulk data.
- To explore the application and scope of biostatistics

#### MODULE I INTRODUCTION TO STATISTICS

Definition of statistics-population and universe-the sample and populationstatistical inference-parameter and statistics. Construction a histograminterpretation of histogram the normal distribution- the mean-mode-and standard deviation- representing the normal curve as straight line-uncertainties in estimating a mean.

#### MODULE II PROBABILITY DISTRIBUTION

The binomial and poisson distributions, Probability, Random sampling and hypothesis testing, The binomial distribution, The poisson distribution, the normal probability distribution, Frequency distribution of continoius variables, Derivation of the normal distribution, Properties of normal distribution, Applications of the normal distribution, Departures from normality, Graphic methods.

#### MODULE III PROPORTION DATA AND ANALYSIS

Examples of Proportion data- MPM-sterility testing of medicines- animal toxicityinfection and immunization studies e.g., LD50, ED50, PD50 statistical treatment to proportion data- Chi-square test-goodness of fit to normal distribution.

#### MODULE IV COUNT DATA

Introduction to Count data, Examples of count data (bacterial cell count, radioactivity count, colony and plaque count, etc.). Statistical treatment to count datapossion distribution-standard error-confidence limits of counts.

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#### MODULE V ANALYSIS OF VARIANCE

Introduction procedure-F and t test. Correlation regression and line fitting through graph points-standard curves- correlation-linear regression (fitting the best straight line through series of points) - standards curves and interpolations of unknown values thereon.

#### Total Hours : 45

#### REFERENCES

- 1. Green. R. H. Sampling Design and Statistical Methods for Environmental Biologists .John Wiley & Sons, 1979.
- 2. Snedecor G. W. and Cochran W. G. Statistical methods. 8<sup>th</sup> Ed., Iowa State Press, 1989.
- Glover T, Mitchell K. Introduction to Biostatistics. 1<sup>st</sup> Ed., Mcgraw Hill Science, 2001.
- 4. Matthews. Sucessful Scientific writing: A step-by-step Guide for Biomedical Scientists. 2<sup>nd</sup> ed. Cambridge University Press, 2001.
- 5. Jerrold H. Z. Biostatistical Analysis. 4<sup>th</sup> ed. Pearson Education, 2006.

#### OUTCOMES:

• On completion of the course the students will possess knowledge of biostatistics to handle and analysis of bulk data in a significant way.

LSB6104

## L T P C 3 0 0 3

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## **OBJECTIVES:**

• The course is aimed at introducing the science of immunology and detailed study of various types of immune systems and their classification structure and mechanism of immune activation.

#### MODULE I OVERVIEW OF IMMUNE SYSTEM

Innate, adaptive and Comparative Immunity, Immune dysfunction and its consequences, Cells & Tissues of Immune System:Hematopoisesis, Apoptosis and Necrosis, systemic function of Immune system, organs of immune systems, Lymphoid cells and organs Evolutionary comparision. Cytokines-Properties of Cytokines, Cytokine Receptors, Cytokine Antagonists, Cytokine Secretion by TH1 and TH2 Subsets, Cytokine-Related Diseases, Therapeutic Uses of Cytokines and Their Receptors, Cytokines in Hematopoiesis.

## MODULE II MOLECULAR IMMUNOLOGY

Immunogenicity Versus Antigenicity, Factors that influence immunogenicity, Epitopes, Haptens and the Study of Antigenicity, Pattern-Recognition Receptors, drugs allegies-when medicine become immunogens, Molecular structure of antibody, Obstacles to Antibody Sequencing, Immunoglobulin Fine Structure, Antibody-Mediated Effector Functions, Antibody Classes and Biological Activities, Antigenic Determinants on Immunoglobulins, The B-Cell Receptor, The Immunoglobulin Superfamily, Monoclonal Antibodies.

## MODULE III ORGANIZATION AND EXPRESSION OF IMMUNOGLOBULIN GENES 9

Genetic Model Compatible with Ig Structure, Multigene Organization of Ig Genes, Variable-Region Gene Rearrangements, Mechanism of Variable-Region DNA Rearrangements, Generation of Antibody Diversity, Class Switching among Constant-Region Genes, Expression of Ig Genes, Synthesis, Assembly, and Secretion of Immunoglobulins, Regulation of Ig-Gene Transcription, Antibody Genes and Antibody Engineering

## MODULE IV ANTIGEN PROCESSING AND PRESENTATION

General organization and inheritance of the major histocompatibility complex (MHC), MHC molecules and genes, detailed genomic map of MHC genes, cellular distribution of MHC molecules, regulation of MHC expression, MHC and immune responsiveness, MHC and disease susceptibility self-MHC restriction of T cells, role of antigen-presenting cells, evidence for two processing and presentation pathways, endogenous antigens: the cytosolic pathway,exogenous antigens: the endocytic pathway presentation of nonpeptide antigens.

## MODULE V GENERATION OF T AND B CELL RESPONSE

T-Cell Receptor, Early Studies of the T-Cell Receptor and  $\alpha\beta$  and T-Cell Receptors: Structure and Roles, Organization and Rearrangement of TCR Genes, T-Cell Receptor Complex: TCR-CD3,T-Cell Accessory Membrane Molecules, Three-Dimensional Structures of TCR-Peptide-

MHC Complexes, Alloreactivity of T Cells, T-Cell Maturation and the Thymus, Thymic Selection of the T-Cell Repertoire,  $T_H$ -Cell Activation, T-Cell Differentiation, Cell Death and T-Cell Populations Peripheral T-Cells, B-Cell Maturation, B-Cell Activation and Proliferation, The Humoral Response, In Vivo Sites for Induction of Humoral Responses, Germinal Centers and Antigen-Induced B-Cell Differentiation, Regulation of B-Cell Development, Regulation of the Immune Effector Response.

## MODULE VI MMUNOTECHNIQUES

Strength of antigen and antibody reaction- cross reactivity, precipitation and agglutination reactions, Radioimmunoassay and ELISA, Markers of immunocompetant cells, separation and purification of immunocompetant cells. Functional tests for immunocompetant cells and histocompatibility testing.Immunological assays- Complementfixation tests, In-vivo tests/ neutralization tests, immunodiffusion, immunoblotting, immunohistochemistry and immunofluorescenece techniques. Biosenor assays for assessing ligand –receptor interaction, CMI techniques- lymphoproliferation assay, Mixed lymphocyte reaction, Cell Cytotoxicity assays, Apoptosis, Microarrays

## Total Hours : 45

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## REFERENCES

- 1. Kuby, RA Goldsby, Thomas J. Kindt, Barbara, A. Osborne Immunology, 6<sup>th</sup> Edition, Freeman, 2002.
- 2. Brostoff J, Seaddin JK, Male D, Roitt IM., Clinical Immunology, 6<sup>th</sup> Edition, Gower Medical Publishing, 2002.
- Janeway et al., Immunobiology, 4<sup>th</sup> Edition, Current Biology publications., 1999.
- 4. Paul, Fundamental of Immunology, 4<sup>th</sup> edition, Lippenco

## OUTCOMES:

After completing the course students will:

- have a detailed understanding of Component of immunity
- know antigen presentation on a detailed molecular level
- understand the concept immunology and the immune system .
- have a in depth knowledge of the cellular and molecular basis for autoimmune disease and allergies.
- have basic knowledge of tumor immunology and the development of novel recombinant antibodies for treatment of cancer and autoimmune disease

LSB6105

## L T P C 3 0 0 3

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#### **OBJECTIVES:**

- To understand the application of Biomedical instrumentation.
- To introduce the student to the various devices of electrical origin and non electrical origin.
- To provide awareness of electrical safety of medical equipments.
- To know the important and modern methods of imaging techniques.

## MODULE I FUNDAMENTALS OF MEDICAL INSTRUMENTATION 10

Role of technology in medicine, landmark developments in biomedical instrumentation, physiological systems of the body, sources of biomedical signals, basic medical instrumentation system, performance requirements of medical instrumentation systems, intelligent medical instrumentation systems, consumer and portable medical equipment, implantable medical devices, Basic components of a biomedical system, Transducers, Piezoelectric, ultrasonic transducers, Temperature measurements, Fibre optic temperature sensors. Amplifiers: Preamplifiers, differential amplifiers, chopper amplifiers Isolation amplifier.

## MODULE II BIOELECTRIC SIGNALS AND ELECTRODES

Origin of bioelectric signals, recording electrodes, silver-silver chloride electrodes, Electrodes, Limb electrodes, floating electrodes, pregelled disposable electrodes, electrodes for ECG, electrodes for EEG, electrodes for EMG, electrical conductivity of electrode jellies and creams, microelectrodes, Micro, needle and surface electrodes, Typical waveforms, Electrical safety in medical environment: shock hazards, leakage current-Instruments for checking safety parameters.

## MODULE III BIOMEDICAL RECORDER

Measurement of blood pressure, Heart rate, Pulmonary function measurements, spirometer, Photo Plethysmography, Body Plethysmography, Blood Gas analysers : pH of blood measurement of blood pCO2, pO2, finger-tip oxymeter - ESR, GSR measurements, Electrocardiograph, vector cardiograph (VCG), phonocardiograph (PCG),digital stethoscope,

electroencephalograph (EEG), electromyography, other biomedical recorders, biofeedback instrumentation.

## MODULE IV CLINICAL INSTRUMENTS AND PATIENT MONITORING SYSTEMS

Medical diagnosis with chemical tests, spectrophotometry, spectrophotometer type instruments, colorimeters, spectrophotometers, clinical flame photometers, selective-ion electrodes based electrolytes analyser, automated biochemical analysis systems, Radio graphic and fluoroscopic techniques, Computer tomography, MRI, Ultrasonography, X-ray Machines and Digital Radiography, Blood cell counter.

## MODULE V THERAPEUTIC EQUIPMENTS AND PATIENT SAFETY 9

Audiometers and Hearing Aids, Pacemakers, Defibrillators, Ventilators, Nerve and muscle stimulators, Diathermy, Heart – Lung machine, Dialysers, Lithotripsy, electric shock hazards, leakage currents, safety codes for electromedical equipment, electrical safety analyzer, testing of biomedical equipment.

## Total Hours: 45

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## OUTCOMES:

After the completion of the course

- The student acquires an adequate knowledge and could co relates the human body to the parameters that have clinical importance.
- The student learn the fundamental principles of medical equipment and patient safety.

## **REFERENCES:**

- 1. R.S.Khandpur, 'Hand Book of Bio-Medical instrumentation', McGraw Hill Publishing Co Ltd. 2003.
- 2. M.Arumugam, 'Bio-Medical Instrumentation', Anuradha Agencies, 2003.
- 3. L.A. Geddes and L.E.Baker, 'Principles of Applied Bio-Medical Instrumentation', John Wiley & Sons, 1975.
- 4. J.Webster, 'Medical Instrumentation', John Wiley & Sons, 1995.
- 5. C.Rajarao and S.K. Guha, 'Principles of Medical Electronics and Bio-medical Instrumentation', Universities press (India)

LSB6106

#### MICROBIOLOGY

## L T P C 3 0 0 3

#### **OBJECTIVES:**

- To expose the fundamentals and scope of microbiology.
- To learn the classification of bacteria, Virus, Fungi & Protozoa
- Learn microbial culture and disease related to microbes.

#### MODULE I INTRODUCTION TO MICROBIOLOGY

History and scope of microbiology; Classification of microorganisms-bacteria, fungi, virus, alga, protozoa; sterilization techniques, disinfectant and antiseptic agents. Microscopy - types of microscopes and their applications-simple and compound, bright field, dark field, fluorescence, phase-contrast and electron microscopes.

## MODULE II BACTERIOLOGY

Major groups of bacteria- Archaebacteria, Actinomycetes, chemoautotrophs, eubacteria, Pseudomonads, cyanobacteria, rickettsias, chlamydias and spirochetes; Bacterial cell- structure and functions of cellular components-cell wall composition of Gram positive and Gram negative bacteria, sub-cellular organizations, flagella, capsule and spores; bacterial staining; antimicrobial agents-antibiotics, chemotherapeutic drugs-antibacterial agents-mode of action; antibiotic resistance.

## MODULE III VIROLOGY

Classification, morphology and characteristics of virus, fungi and Protozoa.Structure of DNA and RNA viruses, viral replication, Bacteriophageslysogeny and lytic cycle; virus like agents-satellites, viroids and prions, antiviral and antifungal drugs.Classification of Helminthic parasites; Life cycle of malarial and filarial parasites.

## MODULE IV CULTURING OF MICROORGANISMS

Microbial culture, continuous and synchronous culture; composition of culture media -solid and liquid media, chemically defined media, complex and differential media; Effect of PH, temperature and radiation on microbial growth.

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## MODULE V MICROBES AND DISEASES

Major human diseases caused by bacterial, viral and fungal pathogens Diseases of the respiratory tract-diphtheria, tuberculosis, pneumonia, influenza, mumps; Diseases of the skin-systemic mycoses, candidiasis; herpes viral infections, chicken pox, zoster and small pox; Genito-urinary infections-Gonorrhea, syphilis, leptospirosis, and AIDS; trichomoniasis; Diseases of GIT-Cholera, ETEC and EIEC infections; shigellosis; Typhoid; Hepatitis, gasteroenteritis. Major human protozoan diseases-Malaria, Amebiasis, Toxoplasmiases.

## Total Hours : 45

## REFERENCES

- 1. Prescott, Harley and Klein- Microbiology-5<sup>th</sup> edition; Publisher: mcgraw Hill science 2002.
- 2. Gerard J. T, Berdell R. F, Christine L. C. Microbiology: An Introduction. 8<sup>th</sup> Ed., Benjamin Cummings, 2004.
- 3. Kenneth J. R,George R, John C. S. Medical Microbiology: An Introduction to Infectious Diseases, mcgraw-Hill Professional, 2003.

## OUTCOMES:

• On the completion of the above objectives student will be able to learn the fundamentals and scope of microbiology, classification of Protozoa, bacteria, Fungi Virus, & microbial culture and diseases.

**OBJECTIVES:** 

LSB6107	BIOCHEMISTRY LAB	LΤ	Ρ

С

0 0 3 1

- To learn the preliminary methods in biochemistry by preparing buffer and different solutions.
- To learn about the factors affecting enzymatic activity.
- learn about several techniques of separations of sugar and amino acids.

## EXPERIMENTS

- 1. Laboratory safety guidelines.
- 2. To prepare an Acetic-Na Acetate Buffer system and validate the Henderson-Hasselbach equation.
- To determine an unknown protein concentration by plotting a standard graph of BSA using UV-Vis Spectrophotometer and validating the Beer- Lambert's Law.
- 4. Determination of pH optima of an enzyme.
- 5. Determination of Km and Kcat of a particular enzyme.
- 6. Effect of temperature on enzyme activity.
- 7. Separation techniques for amino acids and sugar:
  - (a) paper chromatography
  - (b) thin layer chromatography.
- 8. Separation of proteins by native and SDS-PAGE.
- 9. Quantification of reducing sugar in different food material.
- 10. Estimation of different biochemical parameters of blood(a) sugar (b) cholesterol (c) urea.

## **REFERENCES:**

- Wilson K and Walker J, Principles and Techniques in Practical Biochemistry, 5<sup>th</sup> Ed., Cambridge University Press, 2000.
- 2. Holtzhauer M, Basic Methods for the Biochemical Lab, Springer, 2006.

3. Nigam, Lab Manual in Biochemistry: Immunology and Biotechnology, Tata McGraw-Hill Education, 2007.

## OUTCOMES:

On performing the above experiments students will be able to:

- quantify different biomolecules from unknown samples.
- develop an idea about the separation of different biomolecules like proteins and carbohydrate.
- develop an idea about the factors regulating enzyme activity.
- determine the various parameters defining enzyme activity.
- estimate the concentration of various biomolecules in a wide range of samples.

LSB6108 CELL BIOLOGY LAB L T I	с с
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0 0 3 1

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## **OBJECTIVES:**

- To learn basic techniques in molecular biology
- To study and differentiate the electrochemical properties of nucleic acids

## EXPERIMENTS

- 1. Preparation of competent cell by calcium chloride method and checking its efficiency
- 2. Preparation of slides from onion root tip for mitosis
- 3. Isolation & Purification of genomic DNA from bacteria
- 4. Isolation & Purification of plasmid DNA
- 5. Isolation of RNA
- 6. Agarose gel electrophoresis of chromosomal & plasmid DNA
- 7. Restriction Digestion of chromosomal & plasmid DNA
- 8. Isolation of DNA fragment from agarose gel

## REFERENCES

1. Michel R. G and Sambrook J. Molecular Coning- A laboratory manual. Cold spring harbor laboratory press, 2012.

## OUTCOMES:

• On the completion of the above experiments students will be able to handle DNA samples and also to isolate, purify and visualize nucleic acid.

LTB6104

## IMMUNOLOGY LAB

## L T P C 0 0 3 1

## **OBJECTIVES:**

- To acquire knowledge on immunological techniques
- To train in various techniques involving antigen and antibody reactions

## EXPERIMENTS

- 1. Double diffusion, Immuno-electrophoresis and Radial Immuno diffusion.
- 2. Rocket electrophoresis
- 3. Antibody titre by ELISA method.
- 4. ELISA for detection of antigens and antibodies-DOT ELISA
- 5. Sandwich ELISA
- 6. Blood group mapping
- 7. Separation of leucocytes by dextran method
- 8. Separation of mononuclear cells by Ficoll-Hypaque
- 9. Preparation of antigens from pathogens and parasites
- 10. Slide and tube agglutination reaction

## **REFERENCES:**

- 1. Rose et al., Manual of Clinical laboratory Immunology, 6<sup>th</sup> Ed ASM Publications, 2002.
- 2. Lefkovis and Pernis. Immunological methods. Academic Press, 1978.
- 3. Hudson L. and Hay F.C. Practical Immunology. Black Well publishers, 1989

## OUTCOMES:

• Students could independently perform diagnostics assays involving antigenantibody reaction. They also learn to perform the qualitative and quantitative analysis using antibody.

#### SEMESTER II

LSB6221

STEM CELLS IN HEALTH CARE

# LTPC

3 0 0 3

#### **OBJECTIVES**:

- To obtain knowledge on wide-ranging topics related to stem cell
- To learn about the application of stem cells in health care
- To get familiar with the issues and challenges of stem cell political and ethical issues surrounding the stem cell debate.

## MODULE I GENESIS OF CELLS

Concept of stem cells: types, self-renewal and pluripotency, isolation and characterization, Niche and its role on differentiation of stem cells, Stem cells and restorative biology, Reprogramming of genome function through epigenetic inheritance.

## MODULE II STEM CELLS

Embryonic stem cells, Stem Cells from adults. Pluripotency necessary, or is unipotency enough? What are the mechanisms? Stem-cell plasticity, Regulators of pluripotency and differentiation of stem cell. The isolation, expansion, genetic manipulation, genomic reprogramming, and cloning of stem cells. The problem of differentiation of stem cells. Stem Cells and imprinted genes. Differences between adult and embryonic stem cells, what types of cells adult stem cells can become.

## MODULE III CELL & TISSUES

From single to multicellular components - Regulation of cell division and cytoskeleton, Stem cells in regeneration, Cell specification and early signaling events during morphogenesis, Development of cell adhension and motility, Cellular imprinting.

## MODULE IV CELL GROWTH & DEVELOPMENT

Factors controlling cell development - Environmental factors like temperature, oxeygen, location, time, cell number, Chemical factors like growth factors, hormones, cytokines, microRNAs, Genetic factors.

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## MODULE V STEM CELLS AND THERAPEUTICS

Cancer stem cells, Stem cells treatment to diseases, Current stem cell therapies, how we can use stem cells for studying cancer and finding cures to other diseases, Correlation between stem cells and cancer, Stem cells and aging. Clinical applications of hematopoietic stem cells from cord blood first successful transplantation of cord blood in a child with Fanconi's anemia. Treatment of neural diseases such as Parkinson's disease, Huntington's disease and Alzheimer's disease. Repair of damaged organs such as the liver and pancreas. Ethical issues associated with stem cells.

## Total Hours: 45

## REFERENCES

- 1. Kiessling A. A, Human Embryonic Stem Cells: An Introduction to the Science and Therapeutic Potential, Jones and Bartett, 2003.
- 2. Quesenberry P.J. Stem Cell Biology and Gene therapy, 1<sup>st</sup> Edition, Willy-Less, 1998.
- 3. Lanja L, Essential of stem cell Biology, 2<sup>nd</sup> Edition, Academic Press, 2006.
- 4. Ho A.D. and Hoffiman R. Stem Cell Transplantation Biology Processes Therapy, Willy-VCH, 2006.
- 5. Potten C. S. Stem Cells, Elsevier, 2006.

## OUTCOMES:

• After the completion of the course the student will have overall knowledge of scientific research, management, implications and exploitation in Stem Cells in Health care.

LSB6222 DRUG DESIGN AND DEVELOPMENT		L	т	Ρ	С
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#### **OBJECTIVEs:**

- Provide students with an understanding of the process of drug discovery.
- Learn new aspects in Drug Discovery and Design

#### MODULE I INTRODUCTION TO DRUG

Properties of drug, Classification of drug, Inter and Intramolecular: Weak interactions in drug molecules; Optical activity & Biological effect, Types of bonding-Covalent, ion, ion-dipole, hydrogen bonding, C-H hydrogen bonding, dihydrogen bonding, van der Waals interactions.

## MODULE II DRUG TARGETS

Enzymes and receptors. Competitive, non-competitive and allosteric inhibitors, transition-state analogs and suicide substrates. Nucleic acids as drug targets: Intercalating & Non intercalating drugs, Alkylating and metallating agents, Chain cutters and terminators.

#### MODULE III HIGH-THROUGHPUT SCREENING (HTS)

Targets in HTS, Biochemical assays- FRET, FP, HTRF, FCS, Cell Based assays- Second Messenger assays, Reporter gene assays, cell proliferation assays.

## MODULE IV OPTIMIZING TARGET INTERACTIONS

Structure activity relationship (SAR), and QSAR, Binding role of alcohols and Phenols, aromatic rings, alkenes, ketones and aldehydes, amines, amides, Carboxylic acids and quarternary ammonium salts, Identification of a pharmacophore, Drug optimization: strategies in drug design, Variation of substituents. Optimization of ADME properties.

## MODULE V COMPUTATIONAL METHODS IN DRUG DISCOVERY 9

Protein Data Bank, Relationship between sequence and 3D structure, Databases, Binding site prediction, Homology Modeling, Docking tools, Ligand design and Target prediction.

Total Hours: 45

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## REFERENCES

- 1. Drug Discovery and Development- Technology in Transition. Second edition. Raymond G Hill
- 2. The Organic Chemistry of Drug Design and Drug Action, Second Edition Richard B Silverman.

## OUTCOMES:

• This course will train students encompassing different aspects of drug design and development up to date topics in pharmaceutical sciences which is associated with contemporary drug discovery.

#### **OBJECTIVES:**

- To learn about genetic engineering, principles involved in manipulating genes and DNA.
- To know about cloning strategies and expression systems.
- To acquire basic understanding of techniques in genetic engineering.

## MODULE I BASICS CONCEPTS

DNA Structure and properties; Restriction Enzymes; DNA ligase, Klenow enzyme, T4 DNA polymerase, Polynucleotide kinase, Alkaline phosphatase; Cohesive and blunt end ligation; Linkers; Adaptors; Homopolymeric tailing; Labeling of DNA: Nick translation, Random priming, Radioactive and non-radioactive probes, Hybridization techniques: Northern, Southern and Colony hybridization, Fluorescence in situ hybridization; Chromatin Immuno precipitation; DNA-Protein Interactions-Electromobility shift assay; DNaselfootprinting

#### MODULE II CLONING VECTORS

Plasmids; Bacteriophages; M13 mp vectors; PUC19 and Bluescript vectors, Phagemids; Lambda vectors; Insertion and Replacement vectors; Cosmids; Artificial chromosome vectors (YACs; BACs); Animal Virus derived vectors-SV-40; vaccinia/bacculo& retroviral vectors; Expression vectors; pMal; GST; pET-can be omitted vectors; Protein purification; His-tag; GST-tag; MBP-tag etc.; Intein-based vectors; Inclusion bodies; Methodologies to reduce formation of inclusion bodies; Baculovirus and pichia vectors system, Plant based vectors, Ti and Ri as vectors, Yeast vectors, Shuttle vectors. Criteria for selection of vectors.

#### MODULE III CLONING METHODOLOGIES

Insertion of Foreign DNA into Host Cells; Transformation; Transfection, Transduction, Construction of libraries; Isolation of mRNA and total RNA; cDNA and genomic libraries; cDNA and genomic cloning; Expression cloning; Jumping and hopping libraries; Southwestern and Far-western cloning; Protein-protein interactive cloning and Yeast two hybrid system; Phage display; Principles in

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maximizing gene expression. Methods to confirm cloning and reporter genes and proteins.

## MODULE IV PCR AND ITS APPLICATIONS

Primer design; Fidelity of thermostable enzymes; DNA polymerases; Types of PCR – multiplex, nested, reverse transcriptase, real time PCR, touchdown PCR, hot start PCR, colony PCR, cloning of PCR products; Tvectors; Proof reading enzymes; PCR in gene recombination; Deletion; addition; Overlap extension; and SOEing; Site specific mutagenesis; PCR in molecular diagnostics; Viral and bacterial detection; PCR based mutagenesis detection. Sequencing methods; Enzymatic DNA sequencing; Chemical sequencing of DNA; Automated DNA sequencing; RNA sequencing; Chemical Synthesis of oligonucleotides.

## MODULE V APPLICATION OF GENETIC ENGINEERING

Gene silencing techniques; Introduction to siRNA; siRNA technology; Micro RNA; Construction of siRNA vectors; Principle and application of gene silencing; Gene knockouts and Gene Therapy; Creation of knock out mice; Disease model; Somatic and germ-line therapy- in vivo and ex-vivo; Suicide gene therapy; Gene replacement; Gene targeting; Transgenics; cDNA and intragenic arrays; Differential gene expression and protein array. Ethics in genetic engineering and global policy.

## **Total Hours: 45**

## TEXT/REFERENCES

- 1. S.B. Primrose, R.M. Twyman and R.W.Old; Principles of Gene Manipulation. 6<sup>th</sup> Edition, S.B.University Press, 2001.
- 2. J. Sambrook and D.W. Russel; Molecular Cloning: A Laboratory Manual, Vols 1-3, CSHL, 2001.
- 3. Brown TA, Genomes, 3<sup>rd</sup> ed. Garland Science 2006
- 4. Selected papers from scientific journals.
- 5. Desmond S.T. Nicholl An Introduction to Genetic Engineering Cambridge University Press 2008

6. Technical Literature from Stratagene, Promega, Novagen, New England Biolab etc.

## OUTCOMES:

- On completion of the course the scholars will acquire knowledge on the concepts and terminology in genetic engineering.
- Students will be familiar with various cloning strategies in prokaryotes as well as in eukaryotes.
- Students will learn various techniques in genetic engineering.
- They will also get awareness about the social and ethical issues concerning cloning by genetic engineering

LSB6204	COMPUTATIONAL BIOLOGY	LΤ	Ρ	С
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#### **OBJECTIVES**:

- To understand the programming languages applied in computational biology.
- To understand the methods and applications for sequence analysis, Phylogenetics and Protein modelling.

#### MODULE I INTRODUCTION TO PROGRAMMINB LANGUAGE 9

Introduction – Programming languages – Problem solving Technique: Algorithm, Flowchart, Compiling, Testing and Debugging - Basic Perl Data Types, File handle and File Tests – Perl Modules – SQL.

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#### MODULE II PROGRAMMING IN C, C++ AND OOPS

C language Introduction – Tokens – Keywords, Identifier, Variables, Constants, Operators – Structure of a 'C' program - Expression – Data types – Control Statement - C++programming–Object Oriented Concept: Encapsulation, Inheritance, Polymorphism.

## MODULE III COMPUTATIONAL BIOLOGY AND SEQUENCE ANALYSIS 9

Molecular sequences, Genome sequencing: pipeline and data, Next generation sequencing data, Biological databases: Protein and Nucleotide databases, Sequence Alignment, Dynamic Programming for computing edit distance and string similarity, Local and Global Alignment, Needleman Wunsch Algorithm, Smith Waterman Algorithm, BLAST family of programs, FASTA algorithm, Functional Annotation, Progressive and Iterative Methods for Multiple sequence alignment, Applications.

## MODULE IV PHYLOGENETICS

Introduction to Phylogenetics, Distance and Character based methods for phylogenetic tree construction: UPGMA, Neighbour joining, Ultrametric and Min ultrametric trees, Parsimonous trees, Additive trees, Bootstrapping.

## MODULE V PROTEIN STRUCTURE, MODELLING AND SIMULATIONS 9

Protein Structure Basics, Visualization, Prediction of Secondary Structure and Tertiary Structure, Homology Modeling, Structural Genomics, Molecular Docking principles and applications, Molecular dynamics simulations

## **Total Hours: 45**

## **REFERENCES:**

- 1. Dan Gusfield. Algorithms on Strings Trees and Sequences, Cambridge University Press.
- 2. David W. Mount Bioinformatics: Sequence and Genome Analysis, Cold Spring Harbor Laboratory Press, Second Edition, 2004.
- 3. Arthur M. Lesk, Introduction to Bioinformatics by Oxford University Press, 2008.
- 4. Tisdall, James, Beginning PERL for Bioinformatics, O'Reilley Publications, 2001.
- 5. Andrew R. Leach, Molecular Modeling Principles and Applications, Second Edition, Prentice Hall.
- Baldi, P., Brunak, S. Bioinformatics: The Machine Learning Approach, 2<sup>nd</sup> ed., East West Press, 2003
- 7. Baxevanis A.D. and Oullette, B.F.F. A Practical Guide to the Analysis of Genes and Proteins, 2<sup>nd</sup> ed., John Wiley, 2002

## OUTCOMES:

• At the end of this course, students will have been familiarized with language skills and their applications in analyzing Protein structure, sequence analysis which can be used in analyzing the binding effect of drugs on proteins.

LSB6205	COMPUTATIONAL BIOLOGY LAB	LTPC
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#### **OBJECTIVES:**

- To get hands on experience on plasmid construction, mappings and analysis.
- To explore to various tools in bioinformatics.

## EXPERIMENTS

- 1. Plasmid Construction
- 2. Restriction Mapping
- 3. PCR Primer Designing
- 4. Sequence Retrieval and Format Conversion
- 5. ORF Finding
- 6. Homology Search
- 7. Multiple Sequence Alignment
- 8. Gene Prediction in prokaryotes
- 9. Motif finding in DNA and Protein Sequences
- 10. Structure Visualization
- 11. Phylogenetic Analysis
- 12. Protein Secondary Structure Prediction

## REFERENCES

- Rashidi H, Buehler L. K. Bioinformatics Basics: Applications in Biological Science and Medicine. 2<sup>nd</sup> Ed., CRC Press, 2005.
- 2. Baxevanis A. D, Ouellette B.F. F. Bioinformatics: A Practical Guide to the Analysis of Genes and Proteins. 2<sup>nd</sup> edition Wiley, John & Sons, Incorporated, 2004.
- 3. Krawetz S. A, Womble D. D. Introduction to Bioinformatics: A Theoretical and Practical Approach. Humana press, 2003

## OUTCOMES:

• Students will be familiar with various soft skills/tool used in understating modern biology. They will also be able to analyze and interpolate data starting from PCR primer designing to structure predictions.

LSB6206	GENETIC ENGINEERING LABORATORY	LTPC
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## **OBJECTIVES**:

- To practice the earned theoretical knowledge in genetic engineering techniques
- To get acquainted with DNA/gene products know about cloning strategies and expression systems.
- To get familiarize with the sequential processes in genetic engineering.

## LIST OF EXPERIMENTS:

- 1. Isolation of desired DNA/gene by PCR or restriction enzymes
- 2. Gel elution and purification of inserts
- 3. Ligation
- 4. Transformation
- 5. Verification of cloning by PCR or reporter gene or by patching the positive colonies
- 6. Plasmid isolation from PCR positive colonies
- 7. Confirmation of cloning by restriction digestion
- 8. Set up DNA sequencing reaction
- 9. Cleaning the sequencing reaction product
- 10. Automated DNA sequencing
- 11. Sequence Editing
- 12. Sequence analysis by BLAST

## **REFERENCES:**

Laboratory Manual

## OUTCOMES:

- Students will be familiar with various techniques globally used in engineering DNA and gene.
- Students will also be able to analyze the successfully cloned products
- Students will be able to independently plan and execute the cloning of desired gene.

## SEMESTER III

LSB7121

PROTEIN ENGINEERING

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## **OBJECTIVES:**

• This course aims to develop in the students' mind a concept regarding the structure and function of proteins of particular importance, the student will know the production of recombinant protein and in general how to engineer protein to be used as therapeutics.

## MODULE I INTRODUCTION TO PROTEIN ENGINEERING

Forces stabilizing proteins – Van der waals, electrostatic, hydrogen bonding and weakly polar interactions, hydrophobic effects; Entropy – enthalpy compensation, Protein engineering and its applications, features or characteristics of proteins that can be engineered (definition and methods of study) – affinity and specificity; Spectroscopic properties; Stability to changes in parameters as pH, temperature and amino acid sequence, aggregation propensities, etc.

## MODULE II TECHNIQUES FOR PROTEIN ENGINEERING

Methods of measuring the stability of a protein; Spectroscopic methods to study physicochemical properties of proteins: far-UV and near-UV CD; Fluorescence; UV absorbance; ORD; Hydrodynamic properties–viscosity, hydrogen-deuterium exchange.

## MODULE III SITE DIRECTED MUTAGENESIS AND PROTEIN ENGINEERING

Altering Proteins by mutagenesis methods, techniques for Oligonucleotide directed mutagenesis by using single stranded DNA as template, Denatured double stranded DNA as template, PCR based mutagenesis, Engineering proteins by chemical modifications, Genetic fusion of domains, alteration of function by selection and screens, deletion mutagenesis. Introduction of selected mutagenesis by Oligonucleotide directed mutagenesis, Scanning mutagenesis, Insertion of unnatural mutagenesis.

## MODULE IV ENGINEERING PROTEINS FOR PURIFICATION

Introducing cleavage sites, engineering Proteins for Chromatography, Immunoaffinity chromatography, Ion exchange chromatography, Metal affinity chromatography.

#### MODULE V STABILIZATION AND MODIFICATION OF PROTEINS

Principles of structure stabilization by solvent components, sources of exclusion, Balance between cosolvent exclusion and binding, cosolventintertactions in the denaturation reaction, Practical considerations, Post translational modifications- Involving peptide bond, C-terminal, side chain. Modification methods- Enzymatic, Non enzymatic, Specificity, chaperones mediated. Applications of Post translational modifications.

**Total Hours:** 

LSB7102 PLANT AND MEDICAL BIOTECHNOLOGY L T P C

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## **OBJECTIVES:**

- to learn about embryogenesis and other type of hybridization techniques.
- to know about genetic transformation and techniques about gene delivery.
- to have an idea about gene mapping and cloning and different type of biotic and abiotic stress.
- to know about protein engineering and different type of bioinformatics analysis.

## MODULE I PLANT TISSUE CULTURE

Totipotency, organogenesis, somatic embryogenesis, artificial seed production, Micropropagation, somaclonal variation, Germplasm conservation and cryopreservation.Protoplast Culture and Somatic Hybridization Protoplast isolation- its culture and usage, Somatic hybridization and its applications.

## MODULE II AGROBIOLOGY

Agrobacterium-plant interaction; Virulence; Ti and Ri plasmids; Opines and their significance; T-DNA transfer, Genetic Transformation Agrobacteriummediated gene delivery, Direct gene transfer - PEG-mediated, electroporation, particle bombardment and alternative methods; Screenable and selectable markers, Characterization of transgenics, Gene targeting.

## MODULE III MOLECULAR MAPPING & MARKER ASSISTED SELECTION (MAS) 9

Marker assisted selection for genes of agronomic importance, e.g. insect resistance, grain quality and grain yield, Molecular polymorphism, RFLP, RAPD, STS, AFLP, SNP markers; Construction of genetic and physical map, Gene mapping and cloning, strategies for Introducing Biotic and Abiotic Stress Resistance/Tolerance Bacterial resistance; Viral resistance; Fungal resistance; Insects and pathogens resistance; Herbicide resistance; Drought, salinity, thermal stress, flooding and submergence tolerance.

## MODULE IV MOLECULAR THERAPEUTICS

Basic concept of stem cell therapy, neutraceuticals, nanotechnology and clinical trials, revolution in diagnosis, changing approaches of therapy, FDA Organization chart and regulatory measures for drug discovery. Drug discovery - Overview, rational drug design, combinatorial chemistry in drug development, computer assisted drug design, role of bioinformatics in genome based therapy, antisense DNA technology for drug designing. Stem cells in therapy.

## MODULE V VACCINES

Biotechnological approaches to obtain blood products: Tissue plasminogen activator and erythropoietin, Vaccine technology: Subunit vaccines, drawbacks of existing vaccines, criteria for successful vaccine, peptide vaccine, minicells as vaccines, impact of genetic engineering on vaccine production, viral vector vaccines and AIDS vaccine chiral technology.

## Total Hours: 45

## **REFERENCES**:

- 1. Adrian Slater, Nigel Scott and Mark Fowler, Plant Biotechnology: The genetic manipulation of plants, 1<sup>st</sup> Edition, Oxford University Press, 2003
- 2. Edited by BR Jordan, 2<sup>nd</sup> Edition, The Molecular Biology and Biotechnology of Flowering, CABI, 2006.
- 3. Neil Wille, Phytoremediation: Methods and Reviews, 1<sup>st</sup> Edition, Humana Press, 2007.
- 4. Denis Murphy, Plant Breeding and Biotechnology: Societal Context and the Future of Agriculture, Cambridge University Press, 2007.

## OUTCOMES:

On the completion of course student will be able to understand

- different hybridization techniques and basics of embrogenesis.
- they will be able to learn about different gene delivery techniques.
- they will learn about genomics, protein engineering and other bioinformatics tools.

#### Electives-I

LSBY021

## **BIO-ENTREPRENEURSHIP**

# LTPC

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#### **OBJECTIVES**:

The objective of the course is to

- To understand concepts and process involved with bio-entrepreneurship
- To make the students aware of the importance of entrepreneurship opportunities available in the society for the entrepreneur.
- Acquaint them with the challenges faced by the entrepreneur

## MODULE I ACCOUNTING AND FINANCE

Taking decision on starting a venture; Assessment of feasibility of a given venture/new venture; Approach a bank for a loan; Sources of financial assistance; Making a business proposal/Plan for seeking loans from financial institution and Banks; Funds from bank for capital expenditure and for working; Statutory and legal requirements for starting a company/venture; Budget planning and cash flow management; Basics in accounting practices: concepts of balance sheet, P&L account, and double entry bookkeeping; Estimation of income, expenditure, profit, income tax etc.

## MODULE II MARKETING

Assessment of market demand for potential product(s) of interest; Market conditions, segments; Prediction of market changes; Identifying needs of customers including gaps in the market, packaging the product; Market linkages, branding issues; Developing distribution channels; Pricing/Policies/ Competition; Promotion/Advertising; Services Marketing

## MODULE III NEGOTIATIONS/STRATEGY

With financiers, bankers etc.; with government/law enforcement authorities; with companies/Institutions for technology transfer; Dispute resolution skills; External environment/changes; Crisis/ Avoiding/Managing; Broader vision–Global thinking

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## MODULE IV INFORMATION TECHNOLOGY & HUMAN RESOURCE DEVELOPMENT

How to use IT for business administration; Use of IT in improving business performance; Available software for better financial management; E-business setup, management. Human Resource Development (HRD)- Leadership skills; Managerial skills; Organization structure, pros & cons of different structures; Team building, teamwork; Appraisal; Rewards in small scale set up.

#### MODULE V ROLE OF KNOWLEDGE CENTRE AND R&D

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Support mechanism for entrepreneurship in India; Knowledge centres like universities and research institutions; Role of technology and upgradation; Assessment of scale of development of Technology; Managing Technology Transfer; Regulations for transfer of foreign technologies; Technology transfer agencies.

## **Total Hours: 45**

## **REFERENCES:**

- 1. Roy Rajeev, Entrepreneurship Oxford Latest Edition
- 2. E. Gordon & K. Natarajan Entrepreneurship Development Himalaya 2008
- 3. Coulter Entrepreneurship in Action PHI 2<sup>nd</sup> Edition
- 4. P. C. Jain Handbook For New Entrepreneur .Oxford Latest Edition
- 5. S. S. Khanka Entrepreneurial Development S. Chand, Latest Edition
- 6. Thomas W. Zimmerer& Norman M. Scarborough Essentials of Entrepreneurship and small business management, PHI 4th Edition
- 7. Dr. VidyaHattangadi Entrepreneurship, Himalaya 2007
- 8. Vasant Desai Small Scale Industries and Entrepreneurship, Himalaya 2008
- 9. Dr. v. B. Angadi, Dr. H. S. Cheema & Dr. M. R. Das Entrepreneurship, Growth, and Economic Integration A linkage, Himalaya 2009

## OUTCOMES:

On successful completion of this module, learners will be able to have:

- An understanding of accounting and finance related to bio-entrepreneurship
- The capability to apply advanced assessment strategies on marketing of product
- The capability to apply knowledge on negotiations strategy related to bioentrepreneurship
- The ability to apply and handle information technology & human resource development
- Addressing the problems associated with role of knowledge centre and R&D

#### INTELLECTUAL PROPERTY RIGHTS LTPC LSBY022 **& PATENT LAW** 3 0 0 3

## **OBJECTIVES:**

- To learn about the Intellectual Property Rights
- To understand about criteria in applying and maintaining patents.
- To be familiarized with the law and enforcement in Intellectual Property Rights

#### MODULE I INTRODUCTION TO IPR

General regime of intellectual property rights and law. Theories of Intellectual Property Rights, Kinds of Intellectual Property.Intellectual Property as an Instrument of Development, Economic importance of Intellectual Property.Need for Protecting Intellectual Property.National and international perspectives.

## MODULE II TRADE MARK

Introduction to Trade mark, Trade mark registration and maintenance Process, Transfer of Rights, Inter parties Proceeding, Infringement, Dilution Ownership of Trade mark, Likelihood of confusion, Trademarks claims, Trademarks Litigations and International Trade mark Law. Trade Secret, Employee Limitation, Unfair Competition and Trade Secret Litigations.

## MODULE III COPYRIGHTS

Introduction to Copyrights, Principles of Copyright, Copyright Law, Copy right Ownership, Transfer and duration, Right to prepare Derivative works, Rights of Distribution, Rights of Perform the work Publicity Copyright Formalities and Registrations, Limitions, Copyright disputes and International Copyright Law.

## MODULE IV GEOGRAPHICAL INDICATIONS

Registration, Duration of Protection and Renewal; Infringement, Penalties and Remedies.Layout designs of Integrated Circuits- Semiconductor Integrated Circuits Layout-Design Act, 2000, Registration and Effect of Registration, Assignment and Transmission.Protection of Plant Varieties and Farmers' Rights - Authority and Registry, Duration, Effect of Registration and Benefit Sharing, Farmers' Rights, Plant Varieties Protection Appellate Tribunal, Infringement, Offences and Penalties.

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## MODULE V IPR LEGISTLATION AND PATENTING

World Intellectual property organization WIPO – establishment, role, membership, etc., Indian IPR legislation, Indian patent act, national intellectual property policy. Rationale for Intellectual Property Protection in Biotechnology, Patenting Biotechnology Inventions-Objective, Concept of Novelty, Concept of inventive step, Microorganisms, Moral Issues in Patenting Biotechnological inventions. Protection of Plant Varieties.Protection of Traditional Knowledge. Case studies on Basmati rice, turmeric, neem and also current cases.

## Total Hours: 45

## **REFERENCES:**

- 1. Debirag E. B. Intellectual Property. Cengage learning, New Delhi
- 2. Prabhuddha G. Intellectual Property Rights. Tata Mc-Graw–Hill, New Delhi
- 3. Gopalakrishnan N. S. and Agitha, T. G. Principles of Intellectual Property, Eastern Book Company, Lucknow 2009.
- 4. Subbaram N. R. Handbook of Indian patent law and practice, S. Viswanathan printers and publishers Pvt Ltd, 1998.

## OUTCOMES:

• On the completion of the above objectives student will be able to know about IPR and also the importance of protecting their innovation. They will be familiar with international and national law practiced and also recent issues on it.

LSBY023	<b>BIOSAFETY AND BIOETHICS</b>	L	т	Ρ	С

## **OBJECTIVES:**

- Developing a good work ethics and laboratory working condition
- Understanding the importance of following and maintaining laboratory safety guidelines

## MODULE I ETHICS IN BIOLOGY

Principles and purpose of studying bioethics, legal, moral and ethical issues in biological research, human rights, privacy and justice, IPR and technology transfer.

## MODULE II BIOSAFETY

Biosafety in laboratory practices, laboratory associated infections and other hazards, assessment of biological hazards and levels of biosafety, biosafety regulations in handling of recombinant DNA processes and products.

## MODULE III GENETICALLY MODIFIED CROPS AND FOOD

Genetically modified food and biosafety assessment procedures for GM foods and related consumables, including transgenic food crops, ecological safety assessment of recombinant organisms and transgenic crops, case studies of relevance (e.g. BT cotton).

## MODULE IV ETHICAL ISSUES IN LABORATORY RESEARCH

Ethical issues and guidelines for research with laboratory animals, current uses of laboratory animals in biomedical research, animal experimentation using hazardous chemicals, animal care and maintenance, CPSEA guidelines for laboratory animals.

## MODULE V ETHICAL ISSUES IN CLINICAL RESEARCH

Ethical issues and guidelines for research with clinical samples and humans studies, Role of Institutional Human ethical board, ICMR's ethical guidelines and clinical trials registration in India and challenges in clinical trials.

## Total Hours: 45

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## **REFERENCES:**

- 1. Thomas, J.A., Fuch, R.L. Biotechnology and Safety Assessment (3rd Ed). Academic Press, 2002
- 2. Fleming, D.A., Hunt, D.L. Biological safety Principles and practices (3rd Ed). ASM Press, Washington, 2000.
- 3. H.-J. Rehm and G. Reed, Biotechnology A comprehensive treatise (Vol. 12). Legal economic and ethical dimensions VCH.

## OUTCOMES:

• At the end of the course student will develop an idea about the importance of good laboratory practice in high quality research. They will also develop an awareness about the basic fundamental safety measures that a researcher should follow in laboratory.

## ELECTIVES II

LSBY024

MOLECULAR DIAGNOSTICS

# LTPC

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## **OBJECTIVES:**

- Developing the basic concept of molecular diagnostics
- Understanding the common procedures and which are used in disease diagnosis
- To be familiar with various types of diseases diagnosis methods and progression of diagnosed disease.

## MODULE I INTRODUCTION TO MOLECULAR DIAGNOSTICS

Collection, preservation and storage of clinical samples, biopsy, Principles, application and limitations of Biological assays used in diagnosis- PCR, ELISA, FISH, gene sequencing, microarrays, protein arrays. GLP, SOP and ethics in molecular diagnostics.

## MODULE II INFECTIONS

Infection and mode of transmission, types of infectious diseases- bacterial and fungal infections, diagnosis of infections caused by Streptococcus, Coliforms, Salmonella, Shigella, Vibrio, and Mycobacterium- diagnosis of fungal infections, major fungal diseases, Dermetophytoses, Candidiosis and Aspergillosis. Diagnosis of DNA and RNA viruses- pox virus, rhabdo virus, hepatitis; virus diagnosis of protozoan diseases- amoebiosis, malaria, trypanosomiosis, leishmaniasis- study of helminthic diseases- Fasciola hepatica and Ascarislumbricoides. Filariasis and Schistosomiasis.Diagnosis of chicken guinea and swine flu.

## MODULE III CLINICAL GENETICS

Chromosomes chemistry and packaging, Cytogenetic, Structural and numerical abnormalities of chromosomes, Chromosome bands, banding techniques, mutation and polymorphism analysis, human genome project, cancer genetics- oncogenes, tumor suppressor genes- gene therapy, genetic counseling, nucleic acid hybridization techniques, Disease linked with mitochondrial DNA Genetic linkage and chromosome and genetic mapping in human diseases, Prenatal

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## MODULE IV IMMUNODIAGNOSTICS

Introduction to immunodiagnostics, antigen-antibody reactions, antibody production, antibody markers, CD markers, FACS, Human Leukocyte Antigen (HLA) typing, agglutination (ABO/Bacterial), immunoprecipitation, immunodiffusion, flocytometer.

## MODULE V FORENSIC SCIENCE

Introduction to Forensic Science, DNA fingerprinting / DNA Profiling / DNA Testing in Forensic Science.; Ethics, Rules and Procedures in DNA analysis.Autopsy and toxicological diagnosis.Determination of Paternity-Human identification and sex determination.semen analysis, Case study.

#### Total Hours: 45

## **REFERENCES:**

- 1. Tietz Textbook of Clinical Chemistry, Carl A. Burtis, Edward R. Ashwood,
- 2. Harcourt Brace & Company Aisa Pvt. Ltd.
- 3. Essentials of Diagnostic Microbiology, Lisa Anne Shimeld
- 4. The Science of Laboratory Diagnosis, Crocker Burnett

## OUTCOMES:

- Learners will be able to define basic terminology and describes basic concepts in molecular diagnostics
- The students will know the importance and the relevance of molecular diagnostic techniques and applications of molecular diagnostics in various field including medical, forescenic, etc..

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LSBY025	FOOD PROCESS TECHNOLOGY	LTPC
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#### **OBJECTIVES:**

- To explore about food process and technology.
- To get overview of processing of various types of food
- To expose themselves to storage and handling of food and food products.

#### MODULE I STORAGE & HANDLING OF CEREALS

Infestation control; Drying of grains, Processing of rice and rice products.Milling of wheat and production of wheat products, including flour and semolina.Milling of corn, barley, oat, coarse grains including sorghum, ragi and millets; Processing of tea, coffee and cocoa.

#### MODULE II FRESH FRUITS AND VEGETABLES

Preservation of fruits and vegetable by heat treatment.Production and preservation of fruits and vegetable juices, preservation of fruit juice by hurdle technology.Non-alcoholic beverages; Food Laws, food rules and standards, Statistical Quality Control; Various types of packaging.

## MODULE III SEA FOOD

Commercial handling, storage and transport of raw fish; Average composition of fish; Freshness criteria and quality assessment of fish; Spoilage of Fish; Methods of Preservation of fish: Canning, Freezing, Drying, Salting, Smoking and Curing. Quality control of processed fish; Fish processing industries in India.

# MODULE IV ANIMAL PRODUCT

Slaughtering technique of animal; Meat cuts and portions of meat, muscle; Color of meat; Post mortem changes of meat; Meat processing - curing and smoking; fermented meat products (meat sausages & sauces); Frozen meat & meat storage. Classification of poultry meat; Composition and nutritional value of poultry meat & eggs; Processing of poultry meat and eggs; Spoilage and control; Byproduct utilization and future prospects; Poultry farms in India.

# MODULE V DIARY PRODUCT

Composition of milk; Varieties of milk; Checks for purity of milk; Handling of fresh milk.Pasteurization of milk; HTST and UHT techniques; Packaging of milk; Fermentation of milk and fermented milk products. Manufacture of milk products like evaporated milk, powder milk, condensed milk, cream butter, cheese, yogurt, ice cream, ghee, baby food and sweet meat. Quality control of milk and milk products; Milk plant hygiene and sanitation.

# Total Hours: 45

# REFERENCES

- 1. Principles of Food Science, Vol-I by FennmaKarrel
- 2. Modern Dairy Products, Lampert LH; 1970, Chemical Publishing Company.
- 3. Developments in Dairy Chemistry Vol 1 & 2;
- 4. Processed Meats; Pearson AM & Gillett TA; 1996, CBS Publishers.
- 5. Meat; Cole DJA & Lawrie RA; 1975, AVI Pub.
- 6. Post Harvest Technology of cereal pulse and oil seeds by Chakraborty, AC
- Egg and poultry meat processing; Stadelman WJ, Olson VM, Shemwell GA & Pasch S; 1988, Elliswood Ltd.
- 8. Preservation of Fruits & Vegetables by Girdhari Lal, Sidhapa and Tandon
- 9. Developments in Meat Science I & II, Lawrie R; Applied Science Pub. Ltd.
- 10. Egg Science & Technology; Stadelman WJ & Cotterill OJ; 1973, AVI Pub.
- 11. Technology of Food Preservation by Desrosier Fish as Food; Vol 1 & 2; Bremner HA; 2002, CRC Press.
- 12. Fish & Fisheries of India; Jhingram VG; 1983, Hindustan Pub Corp.
- 13. Robinson RK; 1996; Modern Dairy Technology, Vol 1 & 2; Elsevier Applied Science Pub.
- 14. Milk & Milk Processing; Herrington BL; 1948, McGraw-Hill Book Company.
- 15. Fox PF; Applied Science Pub Ltd. Outlines of Dairy Chemistry, De S; Oxford.

# OUTCOMES:

• On the completion of the above objectives student will have a sound knowledge on the various techniques involved in food processing, storage and handling of food and food products.

LSBY026

## ANIMAL BIOTECHNOLOGY

# L T P C 3 0 0 3

# **OBJECTIVES:**

The student will learn about

- The basic idea about animal cell culture, drug toxicity and its application.
- Vaccines production and other technology related to antibody production.
- The basic concept of cloning and several issues related with that.
- Genomics and the role of DNA forensics.

# MODULE I ANIMAL CELL CULTURE

Morphology and ultra-structure of animal cell, requirements for animal cell culture, media and reagents, Primary culture, secondary culture, maintaining cell line, Suspension cultures, Somatic cell cloning and hybridization, transfection and transformation, Stem cells and their application, Animal cell culture application for in vitro testing of drugs, toxicity of environmental pollutants in cell culture.

# MODULE II ANIMAL HEALTH BIOTECHNOLOGY

Introduction to immune system, History of vaccines development, concept of vaccines, vaccine production, conventional methods and recombinant approaches, technology for antibody production, Phage display technology, radio immunoassays and enzyme immunoassays, Immunoblotting, Nucleic acid based diagnostic methods including nucleic acid probe hybridization.

# MODULE III ANIMAL REPRODUCTIVE BIOTECHNOLOGY

Culture of embryos, Cryopreservation of embryos, Embryo transfer, Micromanipulation of animal embryos, Artificial insemination, in vitro fertilization, Transgenic animal technology and its different applications, Animal cloningbasic concepts, Cloning from embryonic cells and adult cells; Cloning of farm animals; Cloning for conservation of endangered species, Ethical, social and moral issues related to cloning, Human Cloning.

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#### MODULE IV ANIMAL GENOMICS

Introduction to animal genomics; Different methods for characterization of animal genomes, SNP, STR, QTLS, RFLP, RAPD, proteomics, metobolomics, different breeds of cattle, buffalo, sheep, goats, pigs, camels, horses, canines and poultry, characterization of livestock breeds; Marker assisted breeding of livestock and poultry, Genetic basis for disease resistance; Gene knock out technology and animal models for human genetic disorders.

## MODULE V DNA FORENSICS

Nucleic acid based methods for identification of animal species, Detection of food/feed adulteration with animal protein, adulteration detection in meat using DNA based methods, Identification of wild animal species using DNA based methods using different parts including bones, hair, blood, skin and other parts confiscated by anti-poaching agencies; Human forensics; Microbial forensics; Bioterror agents; Biocrimes and Bioterrorism.

#### **Total Hours: 45**

#### **REFERENCES:**

- 1. Animal Cell Culture Practical Approach, 3<sup>rd</sup> Edition, Oxford University, Ed. John R.W. Masters, Press, 2000.
- 2. Ed. Martin, Clynes Animal Cell Culture Techniques, Springer, 1998.
- 3. Animal Cell Biotechnology. Portner, 2<sup>nd</sup> Edition, Humana Press, 2007.
- 4. A. Puller (ed), Genetic engineering in Animals, VCH Publishers.
- 5. Gordon, Reproductive Technologies in Farm Animals, CAB Intl., 2005.
- 6. Pinkert, Transgenic animal technology, Academic Press, 2006.

## OUTCOMES:

on the completion of the above course student will learn about

- basic of animal cell culture and the production of antibodies.
- how vaccines is being produced and its importance in several aspects
- the social and moral issues related to cloning.
- the importance of studying Forensic science.

# ELECTIVES III

#### LSBY027 ENVIRONMENTAL BIOTECHNOLOGY LTPC

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# **OBJECTIVES:**

- To learn the environment protection Act and Law related to environmental biotechnology
- To give basic idea on environmental sample analysis
- To understand the basic principles involved in waste water management
- To get the information on usage of Bioremediation-biotechnology
- To inform students about Biooxidation& microbial leaching

#### INTRODUCTION TO ENVIRONMENTAL MODULE I BIOTECHNOLOGY

Water, Soil and Air: their sources and effects. Removal of Specific Pollutants: Sources of Heavy Metal Pollution, Microbial Systems for Heavy Metal Accumulation, Biosorption& detoxification mechanisms, Environment protection Act: Environmental laws, Environmental policies, Environmental ethics. UN declaration. Environmental protection and conservation. Environmental Impact Assessment, Ecoplanning and Sustainable Development

# MODULE II ENVIRONMENTAL SAMPLE ANALYSIS

Physicochemical and bacteriological analysis of soil and water, Problems associated with soil alkali soils, sodic soils, and solid waste, Fate of insecticides fungicides, pesticides in soil, use of genetically modified (insect-, pest- and pathogen resistant) plants. Ecotoxicology of soil pollutants, Municipal solid waste treatment strategies.

# MODULE III WASTE WATER MANAGEMENT

Waste water constituents, Analysis and selection of flow rates and loadings, Process Selection, Physical unit operations, Chemical unit operations, Fundamentals of biological treatment, Role of biotechnology in water purification systems. Types and kinetics of biological treatment, Advanced waste water treatment, Biological Processes for Industrial and domestic

effluent, Treatment, Aerobic Biological Treatment, Anaerobic Biological Treatment.

# MODULE IV BIOREMEDIATION-BIOTECHNOLOGY

Bioremediation-Biotechnology for clean environment, Biomaterials as substitutes for non-degradable materials, Metal microbe interactions: Heavy Metal Pollution and impact on environment, Microbial Systems for Heavy Metal Accumulation, Biosorption, molecular mechanisms of heavy metal tolerance Bioindicators and biosensors for detection of pollution. Biotechnology for Hazardous Waste Management, Persistent organic pollutants, Xenobiotics, Biological Detoxification of PAH, Biotechniques for Air Pollution Control. Solid Waste Management.

# MODULE V BIOOXIDATION & MICROBIAL LEACHING

Biooxidation – Direct and Indirect Mechanisms – Biooxidation Kinetics; Bacterial oxidation of Sphalerite, Chalcopyrite and Pyrite.; Extraction of metals from ores; Recovery of metals from solutions; Microbes in petroleum extraction; Microbial desulfurization of coal.

# Total Hours:45

# **REFERENCES:**

- 1. Amann, R.I. Stromley, J. Stahl : Applied & Environmental Microbiology
- 2. Environmental Microbiology, W.D. Grant & P.E. Long, Blakie, Glassgow and London.
- 3. Microbial Gene Technology, H. Polasa (ED.) South Asian Publishers, New Delhi.
- 4. Biotreatment Systems, Vol. 22, D. L. Wise (Ed.), CRC Press, INC.
- 5. Standard Methods for the Examination of Water and Waste Water (14<sup>th</sup> Education), 1985. American Public health Association

# OUTCOMES:

On successful completion of this module, learners will be able to have:

• An understanding of environment protection regulations and source of environmental pollutions.

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- The capability to apply advanced knowledge on environmental sample analysis
- The capability to apply avalanched discipline in waste water management
- The ability to formulate technique for bioremediation process
- An understanding of how biooxidation& microbial leaching helping in the industries.

LSBY028	BIONANOTECHNOLOGY	L	Т	Ρ	С
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#### **OBJECTIVES:**

- To provide an introduction to nanobiotechnology.
- To make the students understand about the functional principles of nanobiotechnology

#### MODULE I FUNDAMENTALS OF NANOSCIENCE

Introduction, the nanoscale dimension and paradigm, definitions and historical evolution (colloids etc.) and current practice, types of nanomaterials and their classifications (1D, 2D and 3D etc. nanocrystal, Nanoparticle, Quantum dot, Quantum Wire and Quantum Well etc), Polymer, Carbon, Inorganic, Organic and Biomaterials –Structures and characteristics.

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# MODULE II CHARACTERIZATIONS IN BIONANOTECHNOLOGY 9

Optical (UV-Vis/Fluorescence), X-ray diffraction, Imaging and size (Electron microscopy, light scattering, Zeta potential), Surface and composition (ECSA, EDAX, AFM/STM etc), Vibration (FT-IR and RAMAN), SERS -3, Magnetic, Electrical and Electrochemical.

## MODULE III APPLICATIONS OF BIONANOTECHNOLOGY

Materials in Biosystems: Proteins - Lipids - RNA and DNA, Protein Targeting – Small Molecule/Nanomaterial - Protein Interactions Nanomaterial-Cell interactions-Manifestations of Surface Modification (Polyvalency),Drugs-Photodynamic therapy, molecular motors, neuroelecronic interphases, development of nanoluminiscent tags.

## MODULE IV NANOMATERIALS AND DIAGNOSTICS

Drug Delivery and Therapeutics, MRI, Imaging, Surface Modified Nanoparticles, MEMS/NEMS, based on Nanomaterials, Peptide/DNA Coupled Nanoparticles, Lipid Nanoparticles For Drug Delivery, Inorganic Nanoparticles For Drug Delivery, Metal/Metal Oxide Nanoparticles (antibacterial/anti fungal/anti viral), Anisotropic and Magnetic Particles (Hyperthermia).

# MODULE V NANOMATERIALS AND TOXICITY EVALUATION

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Designer biopolymers, Procollagen, DNA Polynode, RNA topoisomerase, Protein – magnetic materials, Cyto-toxicity, Geno-toxicity, In vivo tests/assays.

#### **Total Hours: 45**

# **REFERENCES**:

- 1. C. M. Niemeyer, C. A. Mirkin, Nanobiotechnology: Concepts, Applications and Perspective, Wiley VCH, 2004.
- 2. 2 T. Pradeep, Nano: The Essentials, McGraw Hill education, 2007.
- 3. Nicholas A. Kotov, Nanoparticle Assemblies and Superstructures, CRC, 2006.
- 4. David S Goodsell, "Bionanotechnology", John Wiley & Sons, 2004.

# OUTCOMES:

• After the completion of the course the student will have the basic knowledge of nanotechnology in biotechnology. In detail understanding of the application of Nanomaterials in biotechnology and acquire the knowledge about the DNA, proteins, amino acids, drug delivery, biomedicine etc.

LSBY029	INDUSTRIAL BIOTECHNOLOGY	L	Т	Ρ	С
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#### **OBJECTIVES:**

- To obtain knowledge on wide-ranging topics related to applications of biotechnology in industries.
- To learn about bioprocess technology and its applications
- To get familiar with enzymes and microbes used for industrial purposes.

# MODULE I FERMENTATION & PROCESSING

Introduction to fermentation technology: Upstream and downstream processing of biomolecules. Isolation, Preservation and Improvement of Industrial Micro-Organisms; Medium requirements for fermentation process; Criteria for good medium; Sterilization - batch and continuous heat sterilization of liquid media, filter sterilization of liquid media and Air. Design of sterilization equipment

## MODULE II KINETICS OF SUBSTRATE UTILIZATION, PRODUCT FORMATION AND BIOMASS PRODUCTION

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Phases of cell growth in batch cultures - transient growth kinetics, Simple unstructured kinetic Models for microbial growth, Growth of filamentous organisms; Environmental conditions affecting growth kinetics, substrate and product inhibition on cell growth and product formation; structured kinetic Models, segregated kinetic Models of growth. Production of primary and secondary metabolites. The production of some commercially important Organic acids, amino acids and alcohols, study of production processes for various classes of low molecular weight secondary metabolites: Antibiotics, quinones, aromatics, Vitamins and Steroid.

## MODULE III BIOPROCESSING

Industrial use of micro organisms; Microbes exploited commercially-Saccharomyces, Lactobacillus, Penicillium, Acetobactor, Bifidobacterium, Lactococcus, Streptococcus etc; Fermentation-process, media and systems; Upstream and downstream processing; Product development; Dairy fermentation and fermented products

#### MODULE IV BIOREACTORS

Animal Cells as bioreactors, characteristics of bioreactors, expression and over production of targeted proteins –human growth hormones – production of and interferon's. Good manufacturing practice bio safety issues bioethics, Intellectual Property patenting issues.

#### MODULE V INDUSTRIAL APPLICATION OF ENZYMES

Immobilized enzymes - principles & techniques of immobilization - commercial production of enzymes; amylases, proteases, cellulose, artificial enzymes, industrial applications, fermentation, enzymes Modification, site directed mutagenesis; immobilized enzyme in industrial processes.Structure and function of coenzyme - reactions involving TPP, pyrodoxal phosphate, nicotinamide, flavin nucleotide, coenzyme A and biotin. Industrial utilization of enzymes, food, detergents, energy, waste treatment, pharmaceuticals and medicine.

#### Total Hours: 45

#### REFERENCES

- 1. Maheshwari, D. K. et. al., Biotechnological applications of microorganisms, IK . International, New Delhi, 2006
- 2. Stanbury, P. F. et. al., Principles of Fermentation Technology, 2<sup>nd</sup> Edition, Elsevier, UK, 1995.
- 3. Waites, M. J. et.al., Industrial Biotechnology: An Introduction, Blackwell publishing, UK, 2007

## OUTCOMES:

 After the completion of the course the student will have overall knowledge of scientific industrial biotechnology and applications of microbes and enzymes used in industry.