

DEPARTMENT OF BIOMEDICAL SCIENCES M.SC., BIOMEDICAL SCIENCES

REGULATIONS AND SYLLABUS [For the candidates admitted from the Academic Year 2022 – 2023 onwards]



ALAGAPPA UNIVERSITY (A State University Accredited with "A+" grade by NAAC (CGPA: 3.64) in the Third Cycle and Graded as Category-I University by MHRD-UGC) Karaikudi -630003, Tamil Nadu.

The panel of Members-Broad Based Board of Studies

Chairperson: Name: Dr. S. Ravikumar, Designation: Professor and Head, Department of Biomedical Sciences, Alagappa University, Teaching Experience:28 years, Research Experience:28 years, Area of research: Drugs from the Ocean	
Foreign Expert: Name: Dr. K. Anand, D. Tech, PrChemSA, MRSC , Designation: Research Scientist: NRF-DSI Innovation Fellow, Department: Department of Chemical Pathology, University of the Free State: Teaching Experience: Nil , Research Experience: 8 years, Area of Research: Medical Biochemistry	
Indian Expert: Name: Dr. K.Ruckmani, Designation: Professor Department: Department of Pharmaceutical Technology: Anna University, Teaching Experience:28 years, Research Experience:28 years, Area of Research: Nanobio translational research	
Indian Expert: Name: Dr. N. Thajuddin, Designation: Professor Department: Dept. of Microbiology, Bharathidasan University, Teaching Experience: 30 years, Research Experience: 30 years, Area of Research: Micro algal technology	
Industry Expert: Name: Dr. S. Jacob Inbaneson, Designation: General Manager, Athenese-DX. Pvt. Ltd, Module No. 407 & 408, 4 th Floor, TICEL Bio Park II No.5, CSIR Road, Taramani, Chennai 600113, Teaching Experience:4 years; Research experience: 12 years, Area: Biomedical device	P
Members (All Department faculty) Name: Dr. S. Ravikumar, Designation: Professor and Head, Department, Department of Biomedical Sciences, Alagappa University, Teaching Experience: 28 years, Research Experience: 28 years, Area of Research: Drugs from the Ocean	
Members (Special Invitee) Name: DR. R. Aananthi, Designation: Assistant Professor & Medical Officer, Department: Alagappa University College of physical education, Alagappa University, Teaching Experience:5 years, Research Experience:5 years, Area of Research: General physician	
Alumnus/Alumna: Name: Mr. R. Jayasathya, Current position-Nil Type of Profession, Nil, Professional address: Nil	

ALAGAPPA UNIVERSITY DEPARTMENT OF BIOMEDICAL SCIENCES

Karaikudi -630003, Tamil Nadu.

REGULATIONS AND SYLLABUS-(CBCS-University Department)

[For the candidates admitted from the Academic Year 2022 - 2023 onwards]

Name of the Department: Biomedical SciencesName of the Programme: Biomedical SciencesDuration of the Programme: Full Time (Two Years)

Choice-Based Credit System

A choice-Based Credit System is a flexible system of learning. This system allows students to gain knowledge at their own tempo. Students shall decide on electives from a wide range of elective courses offered by the University Departments in consultation with the Department committee. Students undergo additional courses and acquire more than the required number of credits. They can also adopt an inter-disciplinary and intra-disciplinary approach to learning, and make the best use of the expertise of available faculty.

Programme

"Programme" means a course of study leading to the award of a degree in a discipline.

Courses

'Course' is a component (a paper) of a programme. Each course offered by the Department is identified by a unique course code. A course contains lecture s/tutorials/laboratory /seminar/project / practical training/report writing /Viva-voce, etc or a combination of these, to meet effectively the teaching and learning needs.

Credits

The term "Credit" refers to the weightage given to a course, usually in relation to the instructional hours assigned to it. Normally in each of the courses credits will be assigned on the basis of the number of lectures/tutorial/laboratory and other forms of learning required to complete the course contents in a 15-week schedule. One credit is equal to one hour of lecture per week. For laboratory/field work one credit is equal to two hours.

Semesters

An Academic year is divided into two **Semesters.** In each semester, courses are offered in 15 teaching weeks and the remaining 5 weeks are to be utilized for conduct of examination and evaluation purposes. Each week has 30 working hours spread over 5 days a week.

Medium of Instruction:

English only

Departmental committee

The Departmental Committee consists of the faculty of the Department. The Departmental Committee shall be responsible for admission to all the programmes offered by the Department including the conduct of entrance tests, verification of records, admission, and evaluation. The Departmental Committee determines the deliberation of courses and specifies the allocation of credits semester-wise and course-wise. For each course, it will also identify the number of credits for lectures, tutorials, practical, seminars etc. The courses (Core/Discipline Specific Elective/Non-Major Elective) are designed by teachers and approved by the Departmental Committees. Courses approved by the Departmental Committees shall be approved by the Board of Studies/Broad Based Board of Studies. A teacher offering a course will also be responsible for maintaining attendance and performance sheets (CIA -I, CIA-II, assignments and seminar) of all the students registered for the course. The Non-major elective programme, MOOCs coordinator and Internship Mentor are responsible for submitting the performance sheets to the Head of the department. The Head of the Department consolidates all such performance sheets of courses pertaining to the programmes offered by the department. Then forward the same to be Controller of Examinations.

PEO-1	The program is to foster high-quality innovative research and teaching and interdisciplinary knowledge to develop specialist academicians and intellectual leaders with excellent professional skills in biomedical sciences.	
PEO-2	The programme will provide students with a firm grounding in current understanding of the structure and function of the human body in health and disease	
PEO-3	To develop into highly-skilled and knowledgeable scientists whom we expect to flourish in the new era of biomedicine	
PEO-4	To learn current developments in the field of biomedical sciences and further pursue to do advanced research into the underlying causes of these disorders, diseases, diagnosis and treatments	
PEO-5	To orient the students to solve laboratory skills such as planning of experiments, data acquisition, management and analysis to a selected research problem	
PEO-6	To create a passion for research while inculcating a scientific temperament and a knowledge inquisitive mind with the main aim of contributing towards human health through basic cum applied research	
PEO-7	To engage the students with the fundamental concepts in the field of human anatomy & Physiology, biochemistry, medical oncology and toxicology to a succinct research problem in the chosen specialty area with modern techniques with infrastructure and equipment facility	
PEO-8	To contributed in the field of research and development, involved in the development of molecular diagnostic devices and technologies, pharmaceuticals, veterinary biomedicine and medical interventional strategies	
PEO-9	To expose the students to various advanced molecular, immunological, genetics, Bioinformatics techniques sequence analysis and related services to the needs of academics, industries, hospitals, and other service sectors	
PEO-10	To expose and make the students orientation with solving the research problems, development of the technologies and products for further applications and translational investigation	

Programme Educational Objectives- (PEO)

Programme Specific Objectives-(PSO)

PSO-1	Enable students to acquire laboratory skills in biomedical science
PSO-2	Unique courses on forensic science, bioengineering and artificial organs
PSO-3	Learn courses from other departments and MOOCS open platform
PSO-4	Knowledge on the exploration of newer drugs from marine origin
PSO-5	Training in diagnostic laboratory/hospital during two months summer holidays

Programme Outcome-(PO)

PO-1	Thorough knowledge in handling the human disease diagnosis kits	
PO-2	Understanding the principles and applications of the forensic science, bio engineering and artificial organs	
PO-3	Gain knowledge on the courses other than the core and elective courses	
PO-4	Trained with additional practical skills through hands on training	
PO-5	Recognize the contribution of the basic biomedical sciences to advancing public health sciences.	
PO-6	Apply fundamental concepts of the biomedical sciences to explore public health issues, in particular, prevention, diagnosis, and treatment of disease.	
PO-7	Students would be oriented for undertaking a research career in a top most Institution across the globe or to take a jobs in Biomedical, pharmaceutical, diagnostic and clinical-related industries.	
PO-8	Students would be ready to solve and biomedical problems to develop and formulate appropriate questions, organize and test hypotheses, and apply research results to improve healthcare sectors by molecular diagnostics and process and products	
PO-9	Highly competent human resources could be generated with knowledge in one or more of the following areas: Human Genetics, Medical oncology, pharmacology, Clinical Biochemistry etc., with various advanced techniques.	
PO-10	 Pharmacology, Clinical Biochemistry etc., with various advanced techniques. With the extensive hands-on training in the program, after completin this course, students would be highly knowledgeable in the biomedical sciences and their implication to the advancement of human health and biomedical developments. 	

Programme Specific Outcome-(PSO)

PSO-1	Gain knowledge about contribution of each organ system to the maintenanc	
	of homeostasis.	
PSO-2	Acquire knowledge on the biomolecules and their importance in normal functioning of living organisms.	
PSO-3	Empathize the cascade mechanisms underlying the process over identification of pharmaceutically active compound	
PSO-4	Understand the mode of transmission of diseases and its diagnosis.	
PSO-5	Understanding the kinds of naturally occurring and synthetic toxic substances with our different biological systems	

Eligibility for admission

A candidate who has passed Bachelor's Degree in Biological Sciences (Anatomy, Physiology, Genetics, Medical Biochemistry, Pathology, Physiology, Pharmacology and Environmental toxicology, Endocrinology, Microbiology, Biochemistry, Biotechnology, Biomedical Science, Botany, Zoology, Bioinformatics, Marine Biology, Computational Biology, B. Pharm, B.Sc., Nursing (3or4years), Pharmacology) with at least 55% of marks and 50% marks for SC/ST candidates as main course of study of any university accepted by the syndicate as equivalent thereto, subject to such condition as may be prescribed therefore shall be permitted to appear and qualify for the M.Sc. Degree in Biomedical Science of this university after a course of study of two academic years.

Minimum Duration of programme

The programme is for a period of two years. Each year shall consist of two semesters viz. Odd and Even semesters. Odd semesters shall be from June / July to October / November and even semesters shall be from November / December to April / May. Each semester there shall be 90 working days consisting of 6 teaching hours per working day (5 days/week).

220-

Components

A PG programme consists of a number of courses. The term "course" is applied to indicate a logical part of the subject matter of the programme and is invariably equivalent to the subject matter of a "paper" in the conventional sense. The following are the various categories of the courses suggested for the PG programmes:

- A. Core courses (CC)- "Core Papers" means "the core courses" related to the programme concerned including practical's and project work offered under the programme and shall cover core competency, critical thinking, analytical reasoning, and research skill.
- B. Discipline-Specific Electives (DSE) means the courses offered under the programme related to the major but are to be selected by the students, shall cover additional academic knowledge, critical thinking, and analytical reasoning.
- C. Non-Major Electives (NME)- Exposure beyond the discipline
- Students have to undergo a total of two Non Major Elective courses with 2 credits offered by other departments (one in II Semester another in III Semester).
- A uniform time frame of 3 hours on a common day (Tuesday) shall be allocated for the Non-Major Electives.
- Non Major Elective courses offered by the departments pertaining to a semester should be announced before the end of previous semester
- Registration process: Students have to register for the Non-Major Elective course within 15 days from the commencement of the semester either in the department or NME portal (University website).

D. Self Learning Courses from MOOCs platforms.

- > MOOCs shall be on voluntary for the students.
- > All PG programmes students have to undergo a total of 2 Self Learning Courses (MOOCs) one in II semester and another in III semester.
- > The actual credits earned through MOOCs shall be transferred to the credit plan of programmes as extra credits. Otherwise 2 credits/course be given if the Self Learning Course (MOOCs) is without credit.
- > While selecting the MOOCs, preference shall be given to the course related to employability skills.

E. Projects / Dissertation /Internships (Maximum Marks: 200)

The student shall undertake the dissertation/Project work during the fourth Semester.

Plan of work \succ

Project/Dissertation

The candidate shall undergo Project/Dissertation Work during the final semester. The candidate should prepare a scheme of work for the dissertation/project and should get approval from the guide. The candidate, after completing the dissertation /project work, shall be allowed to submit it to the university departments at the end of the final semester. If the candidate is desirous of availing the facility from other departments/universities/laboratories/organizations they will be permitted only after getting approval from the guide and HOD. In such a case, the candidate shall acknowledge the same in their dissertation/project work.

Format to be followed for dissertation/project report \triangleright

The format /certificate for thesis to be followed by the student are given below

- Title page \triangleright
 - Certificate
 - Acknowledgment
 - Content as follows:

Chapter No	Title	Page number
1	Introduction	
2	Aim and objectives	
3	Review of literature	
4	Materials and methods	
5	Result	
6	Discussion	
7	Summary	
8	References	

Format of the title page Title of Dissertation/Project work

Dissertation/Project submitted in partial fulfillment of the requirement for the degree of Master of Science to the Alagappa University, Karaikudi -630003.

By (Student Name) (Register Number) University Logo

Department of -----

Alagappa University

(A State University Accredited with "A+" grade by NAAC (CGPA: 3.64) in the Third Cycle and Graded as Category-I University by MHRD-UGC, 2019: QS ASIA Rank-216, QS BRICS Rank-104,QS India Rank-20)

Karaikudi - 630003 (Year)

Format of certificates

Certificate -Guide

Place: Karaikudi

Date:

Research Supervisor

Certificate - (HOD)

This is to certify that the thesis entitled "______" submitted by Mr/Mis ------(Reg No: ------) to the Alagappa University, in partial fulfilment for the award of the degree of Master of ------ in ------- is a bonafide record of research work done under the supervision of Dr.-----, Assistant Professor, Department of -------, Alagappa University. This is to further certify that the thesis or any part thereof has not formed the basis of the award to the student of any degree, diploma, fellowship, or any other similar title of any University or Institution.

Place: Karaikudi Date:_____

Head of the Department

Declaration (student)

I hereby declare that the dissertation entitled "-------" submitted to the Alagappa University for the award of the degree of Master of Science in ------" has been carried out by me under the guidance of Dr. ------, Assistant Professor, Department of -------, Alagappa University, Karaikudi – 630 003. This is my original and independent work and has not previously formed the basis of the award of any degree, diploma, associateship, fellowship, or any other similar title of any University or Institution.

Place: Karaikudi

Date:

<u>Internship</u>

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The students who have opted for an Internship must undergo industry/hospital training in the reputed organizations to accrue industrial knowledge in the second semester vacation holidays or other than the regular class hours. The student has to find industry/hospital related to their discipline (Public limited/Private Limited/owner/NGOs etc.,) in consultation with the faculty in charge/Mentor and get approval from the head of the department and Departmental Committee before going for an internship.

Format to be followed for Internship report

 The format /certificate for internship report to be followed by the student are given below

 >
 Title page -Format of the title pageTitle of internship report

Internship report submitted in partial fulfilment of the requirement for the Master of degree into the Alagappa University, Karaikudi -630003. By (Student Name) (Register Number) University Logo

Department of -----

Alagappa University

(A State University Accredited with "A+" grade by NAAC (CGPA: 3.64) in the Third Cycle and Graded as Category-I University by MHRD-UGC, 2019: QS ASIA Rank-216, QS BRICS Rank-104,QS India Rank-20) Karaikudi - 630003 (Year)

Certificate-(Format of certificate – faculty in-charge)

This is to certify that the report entitled "______" submitted to Alagappa University, Karaikudi-630 003 in partial fulfilment for the Master of Science in ------by Mr/Mis------ (Reg No ------) under my supervision. This is based on the work carried out by him/her in the organization M/S ------. This Internship report or any part of this work has not been submitted elsewhere for any other degree, diploma, fellowship, or any other similar record of any University or Institution.

Place	:		
Date:			

Research Supervisor

Certificate (HOD)

This is to certify that the Internship report entitled ""
submitted by Mr/Mis (Reg No) to the Alagappa University, in
partial fulfilment for the award of the Master of Science in is a bonafide record of Internship
report done under the supervision of, Assistant Professor, Department
of, Alagappa University and the work carried out by him/her in the organization
M/S This is to further certify that the internship report or any part thereof has
not formed the basis of the award to the student of any degree, diploma, fellowship, or any other
similar title of any University or Institution.

Place: Karaikudi Date:

Head of the Department

Certificate-(Format of certificate - Company supervisor or Head of the Organization)

This is to certify that the Internship report entitled "-----

----" submitted to Alagappa University, Karaikudi-630 003 in partial fulfilment for the Master of Science in ------by Mr/Mis------ (Reg No ------) under my supervision. This is based on the work carried out by him/her in our organization M/S ------

----- for the period of three months or -----. This Internship report or any part of this work has not been submitted elsewhere for any other degree, diploma, fellowship, or any other similar record of any University or Institution.

Place: Date:_____ Supervisor or in charge

Declaration (student)

I hereby declare that the internship report entitled "-------" submitted to the Alagappa University for the award of the **Master of Science in** --------has been carried out by me under the supervision of ------, Assistant Professor, Department of------, Alagappa University, Karaikudi – 630 003. This is my original and independent work carried out by me in the organization M/S ------ for the period of three months or ------ and has not previously formed the basis of the award of any degree, diploma, associateship, fellowship, or any other similar title of any University or Institution.

Place: Karaikudi Date:_____

- Acknowledgment
- ➢ Content as follows:

Chapter No	Title	Page number
1	Introduction	Par Shi
2	Aim and objectives	IVERSITY 8
3	Organisation profile /details	- V.
4	Methods / Work	
5	Observation and knowledge gained	
6	Summary and outcome of the Internship study	
7	References	

No. of copies of the dissertation/project report/internship report

The candidate should prepare three copies of the dissertation/project/report and submit the same for the evaluation of examiners. After evaluation, one copy will be retained in the department library, one copy will be retained by the guide and the student shall hold one copy.

Teaching methods

The class room teaching would be through conventional lecture, use of OHP, power point presentation and novel innovative teaching ideas like television and computer aided instruction. Periodic field visit enable the student for gathering the practical experience and up to date industrial scenario. Student seminars would be arranged to improve their awareness and communicative skill. In the laboratory, instruction would be given for the safe handling of chemicals and instruments. The practical experiments shall be conducted withspecial efforts to inculcate scientific knowledge among students. The students shall be trained to handle advanced instrumental facilities and shall be allowed to do experiments individually. Periodic test would be conducted to students to assess their knowledge. Slow learners would be identified and will be given special attention.

Attendance

Students must have earned 75% of attendance in each course for appearing for the examination. Students who have earned 74% to 70% of attendance need to apply for condonation in the prescribed form with the prescribed fee. Students who have earned 69% to 60% of attendance need to apply for condonation in the prescribed form with the prescribed fee along with the Medical Certificate. Students who have below 60% of attendance are not eligible to appear for the End Semester Examination (ESE). They shall redo the semester(s) after completion of the programme.

Examination

The examinations shall be conducted separately for theory and practical's to assess (remembering, understanding, applying, analysing, evaluating, and creating) the knowledge required during the study. There shall be two systems of examinations viz., internal and external examinations. The internal examinations shall be conducted as Continuous Internal Assessment tests I and II (CIA Test I & II).

A. Internal Assessment

The internal assessment shall comprise a maximum of 25 marks for each subject. The following procedure shall be followed for awarding internal marks.

Theory -25 marks

Sr.No	Content	Marks
1.	Average marks of two CIA test	15
2.	Seminar/group discussion/quiz	5
3.	Assignment/field trip report/case study report	5
	Total	25

Practical -25 Marks

1	Major Experiment	10 marks
2	Minor Experiment	5 marks
3	Spotter $(2x 5/4 x4)$ or any other mode	10 marks
	Total	25 Marks

Project/Dissertation/internship-50 Marks (assess by Guide/incharge/HOD/supervisor)

1	Two presentations (mid-term)	30 Marks
2	Progress report	20 Marks
	Total	50 Marks

B. External Examination

There shall be examinations at the end of each semester, for odd semesters in the month of October / November; for even semesters in April / May.

- A candidate who does not pass the examination in any course(s) may be permitted to appear in such failed course(s) in the subsequent examinations to be held in October / November or April / May. However candidates who have arrears in Practical shall be permitted to take their arrear Practical examination only along with Regular Practical examination in the respective semester.
- A candidate should get registered for the first semester examination. If registration is not possible owing to shortage of attendance beyond condonation limit / regulation prescribed OR belated joining OR on medical grounds, the candidates are permitted to move to the next semester. Such candidates shall re-do the missed semester after completion of the programme.
- For the Project Report/ Dissertation Work / internship the maximum marks will be 100 marks for project report evaluation and for the Viva-Voce it is 50 marks (if in some programmes, if the project is equivalent to more than one course, the project marks would be in proportion to the number of equivalent courses).
- Viva-Voce: Each candidate shall be required to appear for Viva-Voce Examination (in defense of the Dissertation Work /Project/ internship).

C. Scheme of External Examination (Question Paper Pattern)

Theory - Maximum 75 Marks

Section A			10 questions – 2 each from every unit
Section B	5 questions Either / or type like 1.a (or) b. All questions carry equal marks	The second se	5 questions – 1 each from every unit
Section C	5 questions Either / or type like 1.a (or) b. All questions carry equal marks		5 question –Should cover all units

Practical - Maximum 75 Marks

Section A	Major experiment	15 Marks
Section B	Minor experiment	10 Marks
Section C	Experimental setup	5 Marks
Section D	Spotters (5 x 5 marks)	25 Marks
Section E	Record note	10 Marks
Section F	Vivo voce	10 Marks

Dissertation /Project report/Internship report Scheme of evaluation

Dissertation /Project report/Internship report	100 Marks
Vivo voce	50 Marks

Results

The results of all the examinations will be published through the Department where the student underwent the course as well as through University Website

Passing minimum

- A candidate shall be declared to have passed in each course if he/she secures not less than 40% marks in the End Semester Examinations and 40% marks in the Internal Assessment and not less than 50% in the aggregate, taking Continuous assessment and End Semester Examinations marks together.
- The candidates not obtained 50% in the Internal Assessment are permitted to improve their Internal Assessment marks in the subsequent semesters (2 chances will be given) by writing the CIA tests and by submitting assignments.
- Candidates, who have secured the pass marks in the End-Semester Examination and in the CIA but failed to secure the aggregate minimum pass mark (E.S.E + C I.A), are permitted to improve their Internal Assessment mark in the following semester and/or in University examinations.
- A candidate shall be declared to have passed in the Project / Dissertation / Internship if he /she gets not less than 40% in each of the Project / Dissertation / Internship Report and Viva-Voce and not less than 50% in the aggregate of both the marks for Project Report and Viva-Voce.
- A candidate who gets less than 50% in the Project / Dissertation / Internship Report must resubmit the thesis. Such candidates need to take again the Viva-Voce on the resubmitted Project report.

Grading of the Courses

The following table gives the marks, Grade points, Letter Grades and classifications meant to indicate the overall academic performance of the candidate.

RANGE MARKS	OF	GRADE POINTS	LETTER GRADE	DESCRIPTION
90 - 100		9.0 - 10.0	0	Outstanding
80 - 89		8.0 - 8.9	D +	Excellent
75 - 79		7.5 – 7.9	D	Distinction
70 - 74		7.0 – 7.4	A+	Very Good
60 - 69		6.0 - 6.9	Α	Good
50 - 59		5.0 - 5.9	В	Average
00 - 49		0.0	U	Re-appear
ABSENT		0.0	AAA	ABSENT

Conversion of Marks to Grade Points and Letter Grade (Performance in Paper / Course)

- a) Successful candidates passing the examinations and earning GPA between 9.0 and 10.0 and marks from 90 100 shall be declared to have Outstanding (O).
- b) Successful candidates passing the examinations and earning GPA between 8.0 and 8.9 and marks from 80
 89 shall be declared to have Excellent (D+).
- c) Successful candidates passing the examinations and earning GPA between 7.5 7.9 and marks from 75 79 shall be declared to have Distinction (D).
- d) Successful candidates passing the examinations and earning GPA between 7.0 7.4 and marks from 70 74 shall be declared to have Very Good (A+).
- e) Successful candidates passing the examinations and earning GPA between 6.0 6.9 and marks from 60 69 shall be declared to have Good (A).
- f) Successful candidates passing the examinations and earning GPA between 5.0 5.9 and marks from 50 59 shall be declared to have Average (B).
- g) Candidates earning GPA between 0.0 and marks from 00 49 shall be declared to have Re-appear (U).
- h) Absence from an examination shall not be taken as an attempt.

From the second semester onwards the total performance within a semester and continuous performance starting from the first semester are indicated respectively by Grade Point Average (GPA) and Cumulative Grade Point Average (CGPA). These two are calculated by the following formulate

GRADE POINT AVERAGE (GPA) = $\Sigma_i C_i G_i / \Sigma_i C_i$

GPA = <u>Sum of the multiplication of Grade Points by the credits of the courses</u> Sum of the credits of the courses in a Semester

Classification of the final result

CGPA	Grade	Classification of Final
		Result
9.5 - 10.0	0+	First Class – Exemplary*
9.0 and above but below 9.5	0	
8.5 and above but below 9.0	D++	First Class with Distinction*
8.0 and above but below 8.5	D+	
7.5 and above but below 8.0	D	
7.0 and above but below 7.5	A++	First Class
6.5 and above but below 7.0	A+	
6.0 and above but below 6.5	Α	
5.5 and above but below 6.0	B+	Second Class
5.0 and above but below 5.5	В	
0.0 and above but below 5.0	U	Re-appear

The final result of the candidate shall be based only on the CGPA earned by the candidate.

- a) Successful candidates passing the examinations and earning CGPA between 9.5 and 10.0 shall be given Letter Grade (O+), those who earned CGPA between 9.0 and 9.4 shall be given Letter Grade (O) and declared to have First Class –Exemplary*.
- b) Successful candidates passing the examinations and earning CGPA between 7.5 and 7.9 shall be given Letter Grade (D), those who earned CGPA between 8.0 and 8.4 shall be given Letter Grade (D+), those who earned CGPA between 8.5 and 8.9 shall be given Letter Grade (D++) and declared to have First Class with Distinction*.
- c) Successful candidates passing the examinations and earning CGPA between 6.0 and 6.4 shall be given Letter Grade (A), those who earned CGPA between 6.5 and 6.9 shall be given Letter Grade (A+), those who earned CGPA between 7.0 and 7.4 shall be given Letter Grade (A++) and declared to have First Class.
- d) Successful candidates passing the examinations and earning CGPA between 5.0 and 5.4 shall be given Letter Grade (B), those who earned CGPA between 5.5 and 5.9 shall be given Letter Grade (B+) and declared to have passed in Second Class.
- i) Candidates those who earned CGPA between 0.0 and 4.9 shall be given Letter Grade (U) and declared to have Re-appear.
- e) Absence from an examination shall not be taken as an attempt.

CUMULATIVE GRADE POINT AVERAGE (CGPA) = $\Sigma_n \Sigma_i C_{ni} G_{ni} / \Sigma_n \Sigma_i C_{ni}$

CGPA = <u>Sum of the multiplication of Grade Points by the credits of the entire Programme</u> Sum of the credits of the courses for the entire Programme

Where 'Ci' is the Credit earned for Course i in any semester; 'Gi' is the Grade Point obtained by the student for Course i and 'n' refers to the semester in which such courses were credited.

CGPA (Cumulative Grade Point Average) = Average Grade Point of all the Courses passed starting from the first semester to the current semester.

Note: * The candidates who have passed in the first appearance and within the prescribed Semesters of the PG Programme are alone eligible for this classification.

Maximum duration of the completion of the programme

The maximum period for completion of **M.Sc.**, in Biomedical Sciences shall not exceed eight semesters continuing from the first semester.

Conferment of the Master's Degree

A candidate shall be eligible for the conferment of the Degree only after he/ she has earned the minimum required credits for the Programme prescribed therefor (i.e. 90 credits). Programme).

Village Extension Programme

The Sivaganga and Ramnad districts are very backward districts where a majority of people Lives in poverty. The rural mass is economically and educationally backward. Thus the aim of the introduction of this Village Extension Programme is to extend out to reach environmental awareness, social activities, hygiene, and health to the rural people of this region. The students in their third semester have to visit any one of the adopted villages within the jurisdiction of Alagappa University and can arrange various programs to educate the rural mass in the following areas for three day based on the theme.1. Environmental awareness 2. Hygiene and Health. A minimum of two faculty members can accompany the students and guide them.



M.Sc., Biomedical Sciences

			Programme structure						
S. No	Paper Code	T	Title of the paper	T/P	Credits	Hours/ Week	M	arks	
]	I Semester				Ι	Ε	Total
1	508101	Core 1	Human Anatomy and Physiology	Т	5	5	25	75	100
2	508102	Core 2	Medical Biochemistry	Т	4	4	25	75	100
3	508103	Core 3	Clinical Pathology	Т	4	4	25	75	100
4	508104	Core 4	Lab-I : Human Anatomy and Physiology Medical Biochemistry Clinical Pathology	Р	4	8	25	75	100
6	508105	Core 5	Lab-II : Techniques in Biomedical Science-I	Р	4	6	25	75	100
7	508501	DSE*-1	Bioinformatics and IPR /	Т	3	3	25	75	100
	508502		Marine Pharmaceuticals						
					24	30	150	450	600
			II Semester						
8	508201	Core 6	Medical Genetics	Т	5	5	25	75	100
9	508202	Core 7	Pharmacology	Т	5	5	25	75	100
10	508203	Core 8	Lab-III: Medical Genetics, Pharmacology	Р	4	8	25	75	100
11	508204	Core 9	Lab-IV: Techniques in Biomedical Sciences-II	b-IV: Techniques in Biomedical P 4 6 iences-II		6	25	75	100
12	508503	DSE*2	Forensic Science /	Т	3	3	25	75	100
	508504		Artificial organs						
13		Non-Majo	or Elective **	Т	2	3	25	75	100
14		Self-learn	ing course (SLC) –MOOCs***	Т	E	xtra crec	lit		
	•				23	30	150	450	600
			III Semester						
15	508301	Core 10	Toxicology	Т	5	5	25	75	100
16	508302	Core 11	Medical Oncology	Т	5	5	25	75	100
17	508303	Core 12	Lab-V : Toxicology, Medical Oncology	Р	4	8	25	75	100
18	508304	Core 13	Lab-VI-Techniques in Biomedical Sciences-III	Р	4	6	25	75	100
19	508505 508506	DSE*3	Biomaterials and Tissue Engineering / Bio-imaging Technology	Т	3	3	25	75	100
20		Non-Majo	or Elective **	Т	2	3	25	75	100
21		Self-learn	ing course (SLC) –MOOCs***	Т	E	xtra crec			
					23	30	150	450	600
			IV Semester			1		1	
22	508999	Core 14	****Dissertation Work	Р	15	20	50	150	200
23	508777	Core 15	Hospital training	Р	5	10	50	150	200
					20	30	100	300	400
			Total		90 +		550	1650	2200

DSE – Student Choice and it may be conducted by parallel sections.

** NME -Student have to select courses offered by other (Faculty) departments.

*** SLC- Voluntary basis

*** Dissertation / internship report -Marks -Viva-voce (50) + thesis (100) + internal (50) = 200

T-Theory P-Practical

					Sen	nester -	- I				
Core Cou	rse code:]	Humar	n Anator	my an	d Physi	iology	Т	Cred	its:5	Hours: 5
50	08101										
Objective 1	loorn tha	a gra	ss mor	rnhology	Uni [®]		and fun	ctions	of vari	0116 0	rgans of the
	human bo		55 1101	photogy	, suu	icture i	inu run	cuons	or vari	045 0	igans of the
Structure of Ce generation and groups, estimation tissue, connective Functions of ski regulation by ski	conduction on of RBC e tissue, m n temperat	n, ele C, W nuscle	ectrical BC and e tissue	stimulat d platele e, nervou	ion. I et. Tiss is tissu	Blood C sues and ie and ti	Cell – co d histol issue me	omposit ogy, er mbrane	ion, ori nbryoni s. Anato	gin o c tiss omy o	f RBC, blood sue, epithelial f human skin.
Outcome 1	Acquire l	know	vledge o	on the co	ells an	nd tissue	es				K2
					Unit						
Objective 2	Describe t	the va	arious 1	homeost	tatic n	nechani	sms and	l their i	imbalar	ices	
	s and recep and function us system. Understan	eptors, on of	, brain, Ènervou	brainste 1s tissue, ture and	em, ve , reflex d func	ntricles x action	and spin , velocit	nal cord	l, periph nduction	neral a	and automatic
	systems		ŝ	04/0540		NIVERS	1	8			
Objective 3	Understan	1.4		<u></u>	Unit		6 1.66	_		61	
	piration, di	listurt absorj owlec	oance o ption, n dge abo	f respira novemer	atory f nt of C	function. GI tract.	, pulmo Accesso	nary fui ory orga	nction te ins-liver	est. C , sple	rganization of
P			I HOINE	0514515.	Unit	IV	10	-			
Objective 4	Know-how	w the	specia	l senses			ts				
General Charact mechanism of hormones. Struc regulation of ov Spermatogenesis	eristic and action of cture and f ulation, fer	d clas pitui funct ertiliza	ssificati itary, h ion of ation, ir	on of he hypothala reprodu mplantati	ormon amus, ctive (ion, ge	e, synth thyroid organs, estation,	nesis, seo l, parath hormon parturit	yroid, es of to ion and	adrenal, estes ar l lactatio	pano nd ov on, oc	creas, thymus ary, hormonal ogenesis,
	Have an u relevant so			minolog	y	ological	l proces	ses acci	urately	with	K2
				Unit `							
Objective 5	Acquire th	he co	ordinat	ted worl	king p	attern	of differ	ent org	gans		
	l, determin of nephro ble of thirs faction, tas It attempt	nants on., H st. Re ste, vi ts to	of glor Regulat enal reg isual syst highlig	merular ion: reg gulation stem, he sht the r	filtration gulation of pot aring a	on rate n of ex tassium, and bala	(GFR)., xtracellu calcium nce.	Reabso lar flui a, phosp	orption d osmo hate and	and s blarity d mag	ecretion along and sodium
	sciences an	nd m	edicine	e							

Suggested Readings:

Cinnamon L. VanPutte, Jennifer L. Regan and Andrew F. Russo, (2014). Seeley's Anatomy and Physiology, 10th edition Publisher: McGraw Hill International.

Frederic H. Martini, Judi L. Nath and Edwin F. Bartholomew, (2015). Fundamentals of Anatomy and Physiology, 10th edition Publisher: Pearson Education Limited.

Gerard J. Tortora and Bryan Derrickson., (2015) Anatomy and Physiology-Workbook, 1st edition Publisher: Wiley India Pvt. Ltd.

Gerard, T. J., & Bryan, D. (2015). Anatomy & physiology. *Indian edition, Wiley India pvt. Ltd., New Delhi*, 603-623.

Martini, F. H., Nath, J. L. & Bartholomew, E. F. (2015). Fundamentals of Anatomy and Physiology. 2001. *Pentice Hall: New Jersey*, 538-557.

VanPutte, C. (2016). Seeley's anatomy & physiology. McGraw-Hill Higher Education.

Bruce, J. Colbert, Jeff, J. ankney and Karen, T, Lee (2020). Anatomy, Physiology and Diseases.

Tobin C.E., "Basic Human Anatomy", McGraw-Hill Publishing Co., Ltd., Delhi 1997

Online resources

Text Book Of Human Anatomy & Physiology- https://www.kobo.com/in/en/ebook/text-book-of-human-anatomy-physiology

Anatomy and Physiology Books- https://www.bioexplorer.net/best-anatomy-and-physiology-books.html/

K1-Remember, K2-Understand, K3- Apply K4- Analyze, K5-Evaluate, K6- Create

	Course Outcome VS Programme Outcomes											
CO	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10		
CO-1	S(3)	M(2)										
CO-2	S(3)	M(2)	M(2)	L(1)	L(1)	S(3)	M(2)	M(2)	L(1)	L(1)		
CO-3	S(3)	S(3)	L(1)	M(2)	M(2)	M(2)	S(3)	L(1)	M(2)	M(2)		
CO-4	S(3)	M(2)	L(1)	L(1)	L(1)	L(1)	M(2)	L(1)	L(1)	L(1)		
CO-5	S(3)	L(1)	S(3)	L(1)	M(2)	L(1)	L(1)	S(3)	L(1)	M(2)		
W.AV	3	2	1.8	1.4	1.6	1.8	2)	1.8	1.4	1.6		

1. Low, 2. Medium, 3. Strong

Cours	Course Outcome VS Programme Specific Outcomes										
CO	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5						
CO-1	S (3)	M(2)	M(2)	M(2)	M(2)						
CO-2	S (3)	M(2)	M(2)	L(1)	L(1)						
CO-3	S (3)	S (3)	L(1)	M(2)	M(2)						
CO-4	S (3)	M(2)	L(1)	L(1)	L(1)						
CO-5	S (3)	L(1)	S (3)	L(1)	M(2)						
W.AV	3	2	1.8	1.4	1.6						

			Semester -	Ι				
Core	Course code:	Me	dical Biochemi	stry	Т	Credits:4	Hours:4	
	508102							
			Unit-I					
Objective	1 Understand Biomolecules	the struct		,	nctions		nportance of	
	of carbohydrate m							
-	ypo and Hypergly st. Diabetes mellitu				-	and Galactos	semia, Glucose	
Outcome 1						tance in	K2	
Outcome 1	normal function			and then	mpor		<u>K2</u>	
		-	Unit-II					
Objective 2	Learn the eler catalytic activi		yme structure	that explai	ns the	ir substrate s	pecificity and	
	lipid metabolism,							
	des, Phospholipid	s and Chol	esterol metabol	ism. Stea	atorrhe	a. Inborn er	rrors of lipid	
metabolism.		4 41					TZ 4	
Outcome 2	Conditions	tion the m	etabolic pathw	ays iinked	i with	pathological	K4	
	C C C C C C C C C C C C C C C C C C C	39	Unit III	100				
Objective 3	Outline the s	equence of r	eactions in ana	erobic met	abolisi	m.		
acid metabol	amino acid and pr ism in starvation, nmonia. Uremia, U	Disorders of	plasma protein	 γ-globuli 	inemia	, proteinuria.	urea, uric acid	
	anaemia, cushings							
Outcome 3	Outcome 3Get comprehensive and concise overview of the metabolic disordersK4							
		S	Unit IV	918				
Objective 4	Describe the immune respo		role of horn	ones and	basis	of innate	and adoptive	
	nucleic acid metab ome, orotic aciduri					abolism, Gout	, Lesch-	
Outcome 4	Understand th	e role of pla	telets in hemos			oosis and	K2	
	basis of immu	ne response.	TI •4 T/					
			Unit V					
Objective 5	-	e	e allergic react					
Immunologic autoimmune	cal disorders-diso diseases-SLE, rheu		mmunoglobulin tis, psoriasis, m	2		ergy and hy	persensitivity,	
Outcome 5	Learn the auto	o immune di	seases				K5	
Suggested R					4.			
•	W. & Dominiczak			•			b	
2	. A., & Ferrier, D.	R. (2011). L	ippincott's illus	rated revie	ws: Bio	ochemistry (7	^m ed.). Wolters	
	India Pvt. Ltd.							
	& Voet, J. G. (201		•			•	Inc, 492.	
	Ferrier (2017). Lip	•			•		000) 11	
	Murray and Daryl istry, 25 th edition	K. Granner	and Peter A. M	ayes and V	ictor W	v. Kodwell. (2	000). Harper's	
	oet and Judith G. V	V_{OPT}	Riochemistry ?	rd edition				
	Nelson and Micha		•		ininger	principles of	biochemistry	

Online resources Biochemistry Books, Ebooks And Journalshttps://www.us.elsevierhealth.com/medicine/biochemistry Medical Biochemistry - An Essential Textbook- https://www.thieme.in/Biochemistry-Medical-Biochemistry-An-Essential-Textbook-Panini Textbook of Medical Biochemistryhttps://books.google.co.in/books/about/Textbook_of_Medical_Biochemistry.html?id=BVpDI7n2M9gC& redir_esc=y

K1-Remember, K2-Understand, K3- Apply K4- Analyze, K5-Evaluate, K6- Create

	Course Outcome VS Programme Outcomes											
CO	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10		
CO-1	S(3)	M(2)	S(3)	M(2)	S(3)	M(2)	M(2)	M(2)	M(2)	M(2)		
CO-2	M(2)	S(3)	S(3)	M(2)	S(3)	L(1)	L(1)	M(2)	M(2)	L(1)		
CO-3	L(1)	M(2)	M(2)	S(3)	S(3)	M(2)	M(2)	S(3)	L(1)	M(2)		
CO-4	L(1)	L(1)	S(3)	M(2)	S(3)	L(1)	L(1)	M(2)	L(1)	L(1)		
CO-5	M(2)	S(3)	M(2)	L(1)	S(3)	L(1)	M(2)	L(1)	S(3)	L(1)		
W.AV	1.8	2.2	2.6	2	3	1.4	1.6	2	1.8	1.4		

1. Low, 2. Medium, 3. Strong

Cours	Course Outcome VS Programme Specific Outcomes										
CO	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5						
CO-1	S(3)	S(3)	S(3)	S(3)	S(3)						
CO-2	M(2)	S(3)	S(3)	M(2)	S(3)						
CO-3	L(1)	S (3)	M(2)	S(3)	S(3)						
CO-4	L(1)	S(3)	S(3)	M(2)	S(3)						
CO-5	M(2)	S(3)	M(2)	L(1)	S(3)						
W.AV	1.8	3	2.6	2	3						

Alla V

		Semester – I								
Core	Course code:	Clinical Pathology	Т	Credits:4	Hours:4					
	508103									
		Unit-I			1					
Objective	1 Understand the common infection	concepts of cell injury, clinico-path ous and non-infectious diseases.	ologic	cal correlati	on of					
•••		rculatory Disturbances: Edema, chror		•						
		k, fluid and electrolyte imbalance. G			^					
•	carcinogenesis, tumor; Laboratory diagnosis: cytological techniques including FNAC, Biopsy. Organ transplantation: Immunologic basis of rejection and graft versus host reaction.									
^										
Outcome	1 Acquire knowl host disease.	edge on the cytological techniques a	nd Gra	aft-versus-	K2					
	Unit-II									
Objective 2		al and altered morphology of diffe								
		extent needed for understanding of	diseas	se processes	and their					
	clinical significa		- 1							
	•	rial diseases: tuberculosis and lepr	•		• •					
		s, measles; rickettsial; dengue, Chic								
		pportunistic infections; Parasitic dise		-						
		smission, diagnostic procedures and l		•						
		r Pathology: Rheumatic heart diseas								
	-	tion, Hypertensive heart disease, Co	ngenit	al neart dise	ase,					
• •		rdiovascular diseases.		1**	TZ A					
Outcome 2	Understand the	e mode of transmission of diseases a	na its	diagnosis	K4					
Objective 3	Loorn the com	Unit III mon immunological disorders and	thoir	rocultant	ffoots on the					
Objective 5	human body.	mon minunological disorders and	unen	resultant	inclusion inc					
Inflammatory		ni; pneumonia; pulmonary tuberculo	sis; o	ccupational	lung disorders					
-		l renal function, urine analysis; neph		-	-					
Ų		olycystic kidneys, diagnosis of urina	•							
		esophagus; salivary gland tumors; p								
	•	ntestine, appendix and large intestine;	-							
intestinal tract				C	C					
Outcome 3	Decipher the p	athogenesis of renal and gastrointes	tinal	tract	K4					
	diseases									
	· · · · · · · · · · · · · · · · · · ·	Unit IV								
Objective 4	Have an und	lerstanding of the common haemato	logica	l disorders						
Regulation of	hematopoiesis;	nutritional anaemias: Iron deficienc	y ana	emia, folic	Acid/Vit. B12					
•		ernicious anaemia, hemolytic anaemi								
-		lofibrosis, multiple myeloma; Liver								
Jaundice, hep	atitis, cirrhosis, he	epatocellular and metastatic carcinon	na; Di	seases of th	e gall bladder:					
Cholecystitis,	cholelithiasis. lyn	phoreticular System: Lymphadenitis	, Hod	gkin's and	Non-Hodgkin's					
lymphoma; D	iseases of spleen .	Splenomegaly & Thymus -myasthen	ia gra	vis. Diagnosi	is of liver and					
biliary tract di	seases.		-	_						
Outcome 4	Lean the neces	sity of hemostatic disorders and abn	ormal	lities	K2					
	associated with	menstrual cycle.								
		Unit V								
Objective 5	5	f necessity to diagnose them and to			5					
-	-	s of cervix, Hormonal influences a		-	~ ~					
		cycle and the abnormalities associa								
-		of the breast; prostate; ovarian and			-					
-	reproductive system diseases. Osteopathology: Osteomyelitis; Metabolic									
liseases Rickets/osteomalacia, osteoporosis, Endocrine Pathology: Diagnosis of Diabetes Mellitus; goiter, tumors of thyroid, adrenal diseases; pituitary tumors. Neuropathology: Diagnosis of pyogenic										
	and tuberculous meningitis, brain abscess, tuberculoma; CNS tumors; CSF and its disturbances.Outcome 5Familiar with the knowledge on pyogenic and tuberculousK5									
Outcome 5	Familiar with meningitis.	the knowledge on pyogenic	and	tuberculous	К5					

Suggested Readings: Goodman, C. C., & Fuller, K. S. (2016). Pathology for the Physical Therapist Assistant. Larson, M.T & Donna, D.L.M. (2016). Clinical chemistry: Fundamentals and Laboratory Techniques (1st ed.). Mete, O., & Asa, S. L. (2016). Endocrine Pathology with Online Resource. Cambridge University Press.(ed.,) Rubin, R., Strayer, D. S., & Rubin, E (2008). Rubin's pathology: clinicopathologic foundations of medicine. (ed.,) Salvo, S. G. (2017). Mosby's Pathology for Massage Therapists Raphael Rubin, David S. Strayer. (2011). Pathology: Clinical pathologic foundations of Medicine(6th ed.,) Ozgur Mete, Sylvia L.Asa (2016). Endocrine Pathology (1st ed.,). Donna Larson (2016). Clinical Chemistry: Fundamentals and Laboratory Techniques (1st ed.,) **Online resources** Oxford Handbook of Clinical Pathology- https://www.kobo.com/in/en/ebook/oxford-handbook-ofclinical-pathology-2e Books, Pathology **Ebooks** And https://www.us.elsevierhealth.com/medical-Journalsstudents/pathology K1-Remember, K2-Understand, K3- Apply K4- Analyze, K5-Evaluate, K6- Create

		С	ourse O	utcome	VS Prog	ramme	Outcome	5		
СО	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10
CO-1	S(3)	S(3)	S(3)	M(2)	M(2)	S (3)	M(2)	M(2)	M(2)	M(2)
CO-2	M(2)	S(3)	S(3)	M(2)	M(2)	S(3)	M(2)	M(2)	L(1)	L(1)
CO-3	L(1)	M(2)	M(2)	S(3)	L(1)	S(3)	S(3)	L(1)	M(2)	M(2)

CO-4

CO-5

W.AV

M(2)

L(1)

1.8

S(3)

M(2)

2.6

S(3)

S(3)

2.8

M(2)

L(1)

2

1. Low, 2. Medium, 3. Strong

L(1)

S(3)

1.8

S(3)

S(3)

3

M(2)

L(1)

2

L(1)

S(3)

1.8

L(1)

L(1)

1.4

L(1)

M(2)

1.6

Course Outcome VS Programme Specific Outcomes										
CO	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5					
CO-1	S(3)	S(3)	S(3)	M(2)	S(3)					
CO-2	M(2)	S(3)	S(3)	M(2)	S(3)					
CO-3	L(1)	M(2)	M(2)	S(3)	S(3)					
CO-4	M(2)	S(3)	S(3)	M(2)	S(3)					
CO-5	L(1)	M(2)	S(3)	L(1)	S(3)					
W.AV	1.8	2.6	2.8	2	3					
	1 T	ow 2 Modiu	m 3 Stro							

		Semester – I								
Core	Course code:	Practical –I	P	Credits:4	Hours:8					
	508104	Human Anatomy and Physiology	΄,							
		Medical Biochemistry,								
		Clinical Pathology								
Course		ds on experience on specimen collect	ion, label	ing and docu	imentation					
Objectives		earn the techniques involved in the bl								
	System State	erstand the identification of bones and	1 mechan	isins of vario	us inner					
		in the quantitative and qualitative ana	lysis of r	nicro and m	nacro					
	mol	ecules and cells in blood	-							
	> Lear	n the diagnostics procedures in impo	rtant mici	robial infection	ons					
	1. Specimen	collection and Processing: Coll	action	of specime	en, labeling,					
	documentat	•	ection	of specific	in, labening,					
		nt of blood pressure by using sphygm	omanom	eter and digit	tal meter					
		ion of bones identification and side								
		merus, radius, ulna, lower limb-femu								
	column, rib	s, sternum, sacrum.,	_							
	 Demonstration of major muscles of the body-limbs, head & neck. Demonstration of heart-major vessels of the body-Aorta, subclavian, carotid, 									
			ody-Aorta	, subclavian,	carotid,					
		lial, ulna, femoral, renal.,	stem							
	6. Demonstration of different parts of respiratory system7. Demonstration of the part of digestive system									
	8. Demonstration of other organs- spleen, testis, uterus.									
	9. Protein estimation by Lowry's & Bradford methods,									
		arity of protein & subunit structure by								
		genomic & plasmid DNA by using A	garose g	el electropho	resis					
		on of Blood Groups	na and m	anosting of	maana DDC					
		termination of blood counts - Staini ential leucocytes count using Leishma		eporting of	sinears- KBC,					
		on of packed cell Volume,	ii staiii,							
		on of Erythrocyte sedimentation rate	[ESR]							
		on of clotting time, bleeding time								
		of blood glucose, cholesterol, serum t								
	2	d chemical examinations of urine in	cluding s	ugar, protein	, ketone, bile					
C	salts, bile pi	gments.								
Suggested	Readings:	iczak, M.H. (2019). Medical Biocher	nistm. (5 th	ad)						
		er, D. R. (2011). Lippincott's illustrate			try (7 th ed)					
		(2011). Biochemistry, 4-th Edition. A								
	SonsInc, 492.	(
		a, G. (2012). Update: anatomy & phy								
		Derrickson, B. (2014). Anatomy of	and Physi	iology-WorkE	Book. CBS					
	publication.	P (2010) Tout Pool of Provet 1	Dhua: -1	an (2 rd - 1) I Inivanciti					
Pal,	, G. K., & Pravati, Press (India) Priva	P., (2010). <i>Text Book of Practical</i> te Limited	r nysiolo	gy, (5 ean.	<i>i</i> . Universities					
Pal		nda. N. & Amudharaj. D. (2015). A	tlas of F	Iuman Anata	pmy , $(1^{st} ed)$					
, i ai,	Jordi Vigue. Cham		oj 1							
Rim		R. E., & Korf, B. (Eds.). (2013). En	nery and	Rimoin's esse	ential medical					
	genetics. Elsevier.									
Online	Decourage Harris	Anotomy And Dhysiols and LL at M	mucl II.	mat//\/	o In/We					
		Anatomy And Physiology – I Lab Ma Iuman-Anatomy-Physiology-I-Lab-M			J.m/ w p-					
Essentia			hysiolog		Manual-					
		tbooks/essentials-of-human-anatomy-								
	5373400-08053734			•						

Course Outcomes (CO)	CO1- Learn specimen collecting, labeling and processing. CO2- Understanding the arrangement of various types of bones and inner
	systems CO3- Gain knowledge on the measurements of micro and macro molecules in human blood
	CO4- Acquire knowledge on the laboratory diagnosis of microbial infections. CO5- Learn the various tests involved in the assessment of blood cell counts

	-	C	ourse Ou	utcome	VS Prog	ramme	Outcome	S		-
СО	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10
CO-1	S(3)	S(3)	S(3)	S(3)	M(2)	M(2)	M(2)	M(2)	S(3)	M(2)
CO-2	S(3)	S(3)	M(2)	S(3)	M(2)	M(2)	M(2)	M(2)	M(2)	M(2)
CO-3	S(3)	L(1)	L(1)	S(3)	L(1)	S(3)	L(1)	L(1)	L(1)	S(3)
CO-4	S(3)	L(1)	M(2)	S(3)	L(1)	M(2)	L(1)	M(2)	S(3)	M(2)
CO-5	S(3)	M(2)	L(1)	S(3)	S(3)	L(1)	S(3)	L(1)	M(2)	L(1)
W.AV	3	2	1.8	3	1.8	2	1.8	1.6	2.2	2

1. Low , 2. Medium , 3. Strong

Cour	se Outcom	e VS Progra	mme Spec	ific Outco	mes
CO	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5
CO-1	S(3)	S(3)	S (3)	S(3)	M(2)
CO-2	S(3)	S(3)	S(3)	S(3)	M(2)
CO-3	S(3)	L(1)	S(3)	S(3)	L(1)
CO-4	S(3)	L(1)	S(3)	S(3)	L(1)
CO-5	S(3)	M(2)	S(3)	S(3)	S(3)
W.AV	3	2	3	3	1.8

			Semester -	- I					
Core	Course code: 508105		Techniques in sciences-I		Р	Credits: 4	Hours: 6		
Course Objectives	Get of meth ➤ Gain	comprehensive ods in microbi	nciples and appli- e and concise over al identifications in the laboratory di- zation techniques	erview of the liagnosis of 1	e appli nicrot	ication of sta	C		
	DeciFami	iliar with the p	principles and pra	ctice of bion	nedica	l wastes			
	ow/high pow pes of micros en, autoclave	scopes.							
	 Demonstration of commonly used culture media: LB broth, LB agar, nutrien broth, nutrient agar, blood agar, chocolate agar, MacConkey medium Lowenstein Jensen (LJ) media, Robertson cooked meat media, sabouraud dextrose agar 								
		n staining, Al	bert's staining, a	cid fast stain	ning, 1	lactophenol of	cotton blue		
	5. Moti voge catal	lity tests and s proskauer t	biochemical tests est, citrate utiliz se test for bacteria	ation test, t	riple				
			the disposal of bi	omedical wa	stes				
Aneja Arora Baren In Harv Bayn Harv ed Naiga Lt Perry Voet, Sc	a, D.R., & Aron c, M. R., & Irv. fections: Patho ealth Sciences. es, J.W. & Don ey, R. A., & F l.). Wolters Khu aonkar, M. A. d c, J. J., Staley, J onsInc, 49.	ra, B. (2007). ing, W. L. (20 ogenesis, Imn miniczak, M.H. errier, D. R. (uwer India Pv (2008). A man I. T., & Lory, G. (2011). Bio	nu <mark>al of medical l</mark> S. (2002). Microl ochemistry, (4 th e	ology (2 nd ea crobiology E ry Investigat al Biochemiss tt's illustrate aboratory tec bial life. Sina d.). NewYor	l.). CE E-Book tion d try (5 th d revi chnold tuer A tk: Jol	BS Publicatio :: A Guide to and Control ^h ed.). ews: Bioche pgy. Pragati 1 ssociates Inc hn Wiley&	n. 9 <i>Microbial</i> 9. Elsevier emistry (7 th Books Pvt. orporated.		
ed Micro		ntificpubonline 16/0. Laboratory	e.com/bookdetail/		/-labo				
Course (CO)	Outcomes	CO2- Aware CO3- Under CO4- Comp	n the microscopy the various mether the various the various rehend the methor the principles and	nods of steril culture medi ds of identif	ization a icatior	ns of microbe	es		

	Course Outcome VS Programme Outcomes										
CO	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10	
CO-1	S(3)	S(3)	S(3)	S(3)	M(2)	M(2)	M(2)	M(2)	M(2)	M(2)	
CO-2	S(3)	S(3)	M(2)	S(3)	M(2)	M(2)	M(2)	L(1)	L(1)	M(2)	
CO-3	S(3)	M(2)	S(3)	S(3)	L(1)	S(3)	L(1)	M(2)	M(2)	S(3)	
CO-4	S(3)	M(2)	M(2)	S(3)	L(1)	M(2)	L(1)	L(1)	L(1)	M(2)	
CO-5	S(3)	M(2)	L(1)	S(3)	S(3)	L(1)	S(3)	L(1)	M(2)	L(1)	
W.AV	3	2.4	2.2	3	1.8	2	1.8	1.4	1.6	2	

1.	Low	,	2.	Medium,	3.	Strong
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Co	Course Outcome VS Programme Specific Outcomes											
CO	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5							
CO-1	S(3)	S(3)	S(3)	S(3)	M(2)							
CO-2	S(3)	S(3)	M(2)	S(3)	M(2)							
CO-3	S(3)	M(2)	S(3)	S(3)	L(1)							
CO-4	S(3)	M(2)	M(2)	S(3)	L(1)							
CO-5	S(3)	M(2)	L(1)	S(3)	S(3)							
W.AV	3	2.4	2.2	3	1.8							



			Semester – I			
DSE-1	Co	ourse code:	Bioinformatics and IPR	Т	Credits:3	Hours:3
		508501				
			Unit-I			
Object	ive 1	To explain	the technical know – of information that	nt ca	n be collecte	ed from web
		sources Inv	olved in patenting of novel drugs.			
			bioinformatics web servers. Computer sys			
		_	logy (LAN, WAN, &MAN)Internet basics:	Inter	net connectio	n, web
browsir	-					
Outcon	ne 1		of biological data, submission and ret		al it from	K2
		databases a	nd design databases to store the informat Unit-II	ion		
Ohiaatir		Understand			field of action	as for hotton
Objectiv	ve Z		the essential features of the interdiscipling biological data	lary	field of scien	ice for better
Role of	bioint		numan genome project databases, nucleic a	cid s	equence data	bases (NCBI,
EMBL, I	DDJB), protein seq	uence database (SWISS-PORT); data base s	earch	ning: BLAST	
Outcom	e 2	Demonstrat	e the most important bioinformatics dat	abas	es, perform	K4
		text-and seq	uence-based searches, and analyze the re-	esult	s in light of	
		molecular bi	ological knowledge			
			Unit III			
Objectiv			udents opportunity to interact with algor	ithm	s, tools and	current
		scenario	5			
•		÷	pair wise and multiple sequences. analys	-		
			gene finding and gene scan, protein pre		-	
and func		condary struc	ture, alpha- beta structure, motifs, tertiary	struc	speciali	zeu structure
Outcom	e 3	Experiment	pair wise and multiple sequence alig	nme	nt and will	K4
		analyze the	s <mark>econ</mark> dary and tertiary structures of prote	<mark>ein s</mark> e	equences.	
			Unit IV		L	
Objectiv			e students look <mark>at a</mark> biologic <mark>al</mark> problem fro	om a	computation	al point
		of view				
			rotein confirmation and visualization tool			
	-	-	istory, analog & structural, ligand designing	& op	otimization, de	ocking,
<u> </u>			deling in drug discovery.	. 1.	· · · · · · · · · · · · · · · · · · ·	V.A
Outcom	e 4		the data structure (databases) used in t the information (especially: find genes;			K2
		_	understand and be aware of current			
			ating to this area.	10	scarch and	
		problems re	Unit V			
Objectiv	ve 5	To find out	the methods for analyzing the expressi	on. s	structure and	l function of
- ~J		DNA, RNA	• • •	, ~		
WTO-G.	ATT o	& TRIPS.Dif	ferent types of intellectual property rights	(IPR)).Intellectual	property laws
And the	Intern	net. Trade Re	lated Aspects of Intellectual Property Right	s.Pate	ents-patent ap	plications and
-		g patent- Sele	ected examples of patent in biotechnology. I	licens	sing and com	oulsory
licensing						
Outcom	e 5		out gene and protein expression pattern	is an	d modelling	К5
		cellular i	nteractions and processes.			

Suggested Readings:

Lewis PO., and Zaykin D., (1997). Genetic data analysis: Computer program for the analysis of Allelic data. Available at http://chee.unm.edu/gda.

Amitava chakraborti, (2000). Patenting Biotechnology certain aspects.Pp.69-75.Subbaram, N.R., (2004). Basics of IPR.Pp. 5-13.

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- De Robertis, E. D., & De Robertis, E. M. (1987). Cell and molecular biology. Lea & Febiger.
- Lehninger, A. L., Nelson, D. L., & Cox, M. M. (2004). Overhead Transparency Set for Lehninger Principles of Biochemistry (4th ed.).
- WH Freeman. Murray, R. K., Granner, D. K., Mayes, P. A.,& Rodwell, V. W., (2006). Harper's Biochemistry (27th ed.). McGraw Hill.

Shaik, N.,Halid Rahman, H., Babajan, B., Elango, R. (2019). Essentials of Bioinformatics Vol.II.Springer edition., pp.328.

Online resources

Bioinformatics An Introductory Textbook- https://link.springer.com/book/10.1007/978-3-662-65036-3 **Bioinformatics, 4th Edition-** https://www.wiley.com/en-ca/Bioinformatics%2C+4th+Edition-p-9781119335580

K1-Remember, K2-Understand, K3- Apply K4- Analyze, K5-Evaluate, K6- Create

	Course Outcome VS Programme Outcomes									
СО	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10
CO-1	M(2)	S(3)	S(3)	M(2)	M(2)	M(2)	M(2)	M(2)	M(2)	M(2)
CO-2	L(1)	L(1)	S (3)	L(1)	M(2)	M(2)	L(1)	L(1)	M(2)	M(2)
CO-3	M(2)	S(3)	S(3)	M(2)	S (3)	L(1)	M(2)	M(2)	S(3)	L(1)
CO-4	M(2)	L(1)	S(3)	L(1)	M(2)	L(1)	L(1)	L(1)	M(2)	L(1)
CO-5	M(2)	L(1)	S(3)	L(1)	L(1)	S(3)	L(1)	M(2)	L(1)	S(3)
W.AV	1.8	1.8	3	1.4	2	1.8	1.4	1.6	2	1.8

Course Outcome VS Programme Specific Outcomes									
CO	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5				
CO-1	M(2)	S(3)	S(3)	M(2)	M(2)				
CO-2	L(1)	S(3)	S(3)	L(1)	M(2)				
CO-3	M(2)	S(3)	S(3)	M(2)	S(3)				
CO-4	M(2)	S(3)	S(3)	L(1)	M(2)				
CO-5	M(2)	S(3)	S(3)	L(1)	L(1)				
W.AV	1.8	3	3	1.4	2				

1. Low, 2. Medium, 3. Strong

		Semester – I			
DSE-1	Course code: 508502	Marine pharmaceuticals	Т	Credits:3	Hours:3
		Unit-I	1 0		
-	_	hensive knowledge on the various kin			-
		of natural origin- Complex pol ins, flavonoids, tannins, terpenoids. Ster			
		and peptides, antibiotics, biologics and			,
Outcome	1 Understand marine sourc	an idea about the therapeutic lead a ses	nolecule	es from	K2
	1	Unit-II			
Objective		the essential features of the interdisc ng biological data	iplinary	field of scien	ce for better
actinomycete	s), algae, seag	from marine flora and fauna-m rasses, mangroves, invertebrates –spo hinoderms, tunicates, bryozoans, pisce	nges, cn		ls, bryozoans,
Outcome 2	Recognise t Production	he importance of marine fauna	and flo	ora in drug	K4
Objective	3 Logra the te	Unit III chniques involved in the evaluation o	forndo	drugs	
•		environment-Development of novel for		0	Low
		al enrichment - food supplements. Food		÷	
Outcome 3	Gain theoret	ical knowledge on neutraceuticals			K4
		Unit IV			
Objective 4	Versed in the	e design an <mark>d</mark> formulation of drug dos	age forn	ıs	
		vascular drugs, cytotoxic compounds odic agents and other therapeutically va			
Outcome 4	Recognise th	e marine toxins as drugs	1		K2
	·	Unit V			
Objective 5	Familiarize v	vith the types of toxins and poisons ir	n marine	e organisms	
the biosynthe the major cla	esis of marine r sses of the met		netabolit	es on the proc	•
		plems encountered with the types of o	losage fo	orms	K5
Suggested R Fingerma	0	Recent advances in Marine Biotechnolog	gy, (ed.,))	
Morries I	H. Baslow, (196	9). Marine Pharmacology. The William	ns & Wil	kins Co., Balti	more.
Hall., S a	and G.Strichartz	, (1990). Marine toxins: Origin, Structu	ire and M	Molecular Phar	macology
Paul Sing	gleton (1999).Ba	acteria in Biology, Biotech and Medicin	e, 5th ed	••	
Treves B	rown, K.M., (2	000). Applied Fish Pharmacology. (ed.,)	1		
Dennis W	V. Ross, (2002)	Introduction to molecular medicine 3r	d (ed.,)		
Rodney J	.Y. (2003). Bio	tech and Biopharmaceuticals (ed.,).			
Trivedi, I	P.C. (2004). He	rbal Drugs and Biotechnology (ed.,).			

Online resources Marine Pharmacy Guide- https://www.kobo.com/in/en/ebook/marine-pharmacy-guide Encyclopedia of Marine Biotechnologyhttps://onlinelibrary.wiley.com/doi/book/10.1002/9781119143802 Marine Biomaterials- https://link.springer.com/book/10.1007/978-981-16-4787-1

K1-Remember, K2-Understand, K3- Apply K4- Analyze, K5-Evaluate, K6- Create

	Course Outcome VS Programme Outcomes									
CO	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10
CO-1	M(2)	S(3)	S(3)	M(2)	M(2)	M(2)	M(2)	M(2)	M(2)	S(3)
CO-2	L(1)	M(2)	S(3)	L(1)	M(2)	M(2)	L(1)	L(1)	M(2)	M(2)
CO-3	M(2)	L(1)	S(3)	M(2)	S(3)	L(1)	M(2)	M(2)	S(3)	L(1)
CO-4	L(1)	L(1)	S(3)	L(1)	M(2)	L(1)	L(1)	L(1)	M(2)	L(1)
CO-5	M(2)	M(2)	S(3)	L(1)	L(1)	S(3)	L(1)	M(2)	L(1)	M(2)
W.AV	1.6	1.8	3	1.4	2	1.8	1.4	1.6	2	1.8

1. Low, 2. Medium, 3. Strong

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No al agappa inversity (20									
Course Outcome VS Programme Specific Outcomes									
СО	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5				
CO-1	M(2)	S(3)	S(3)	M(2)	M(2)				
CO-2	L(1)	M(2)	S(3)	L(1)	M(2)				
CO-3	M(2)	L(1)	S (3)	M(2)	S(3)				
CO-4	L(1)	L(1)	S(3)	L(1)	M(2)				
CO-5 🥌	M(2)	M(2)	S(3)	L(1)	L(1)				
W.AV	1.6	1.8	3	1.4	2				

						Sem	nester-II						
Core	Course 5082				N		l genetics	5		Т	Cred	its :5	Hours:5
							Unit-I						
Objectiv							tance an	<u> </u>		•			
	of Human												
	tial mode											, ped	igree
÷	, construct		-	-	-				etic da	ta in pe			
Outcom	e 1Aco	quire ki	now	vledge	on the		ly history	y			K	2	
	- 1					-	nit-II						
Objective		scribe t nan chr				omosoi	mes ban	ding, no	omencl	ature a	nd pat	holog	y of
Human cl chromoso aberration Genetics dystrophy	me band Numeri of fetal w	ing, ka cal and vastage.	aryot Id St . Sez	otype Structur x-linke	analys ral ab ed inho	is; Mo erration eritance	olecular ns. Com e: colour	cytogen mon ch blindne	etics: romoso ess, hae	FISH, ome abi emophili	CGH., normalit a and	Chro ies i nuscu	omosomal n cancer,
Outcome		ow-how nophilia		ne sex-	linked	l inher	ritance s	uch as	colour	blindn	ess and	K	4
					S.	-	nit III	18					
Objective Inborn erracid me		etabolisi	sm, o	disord	ers of	etic tec amino	hnique s acid me	tabolism	n, disor	ders of	branch		ain amino etabolism,
Inborn err acid me Mucopoly variation Pharmaco	rors of me tabolism, ysaccharid by the ef genomics	etabolisi disorc oses an fect of : Anima	sm, o ders ind drug al m	disord s of Albini lgs, He nodels	ers of carbo sm., l eredita in pha	etic tec amino ohydrat Pharma ry disc armacog	hnique s acid me e metal codynam orders wi genomics	tabolism bolism, hics: De th altere , Ecogen	n, disor disor efinitior ed drug	ders of ders o n, drug	branch f lipic metab	mo maco maco	etabolism, , Genetic ogenetics,
Inborn err acid me Mucopoly variation	rors of me tabolism, ysaccharid by the ef genomics	etabolisi disorc oses an fect of : Anima	sm, o ders ind drug al m	disord s of Albini lgs, He nodels	ers of carbo sm., l eredita in pha	etic tec amino ohydrat Pharma ry disc armacog	hnique s acid me e metal codynam orders wi	tabolism bolism, hics: De th altere , Ecogen	n, disor disor efinitior ed drug	ders of ders o n, drug	branch f lipic metab	me me	etabolism, , Genetic ogenetics,
Inborn err acid me Mucopoly variation Pharmaco	rors of me tabolism, ysaccharid by the ef genomics	etabolisi disorc oses an fect of : Anima	sm, o ders ind drug al m	disord s of Albini lgs, He nodels	ers of carbo sm., l eredita in pha ponse a	etic tec amino ohydrat Pharma ry disc armacog	hnique s acid me e metal codynam orders wi genomics	tabolism bolism, hics: De th altere , Ecogen	n, disor disor efinitior ed drug	ders of ders o n, drug	branch f lipic metab	mo maco maco	etabolism, , Genetic ogenetics,
Inborn err acid me Mucopoly variation Pharmaco	rors of me tabolism, vsaccharid by the ef genomics 3 Lea e 4 Des	etabolisi disorc oses an fect of : Anima arn the	sm, o rders and drug al m e dru the t	disord of Albini ags, He nodels 1g res p molec	ers of carbc sm., l eredita in pha ponse	etic tec amino ohydrat Pharma ry disc armacog and m	hnique s acid me e metal codynam orders wi genomics	tabolism bolism, nics: De th altere , Ecoger n	n, disor disord efinitior ed drug netics	ders of ders o n, drug ; respon	branch f lipic metab se, Pha	molism macc K	etabolism, , Genetic ogenetics,
Inborn err acid me Mucopoly variation Pharmaco Outcome Objective	rors of me tabolism, vsaccharid by the ef genomics 3 Lea e 4 Des of 1	etabolism disorci oses an fect of : Anima arn the scribe the netabol	sm, o rders ind drug al m e dru blism	disord s of Albini nodels ng resp molec	ers of carbo sm., l eredita in pha ponse Un ular a	etic tec amino ohydrat Pharma ry disc armacog and m ait IV and bio	hnique s acid me e metal codynam orders wi genomics etabolism	tabolism bolism, hics: De th altere , Ecogen n l pathw	n, disor disorc finitior d drug netics /ays of	ders of ders o n, drug ; respon	branch f lipic metab se, Pha	mo olism macc Ka	etabolism, , Genetic ogenetics, 4
Inborn err acid me Mucopoly variation Pharmaco Outcome Objective Inherited	rors of me tabolism, vsaccharid by the ef genomics 3 Lea 2 4 Des of 1 immunod	etabolisi disorc oses an fect of Anima arn the scribe th netabol eficienc	sm, o rders ind drug al m e dru e dru the i blism	disord s of Albini ugs, Ha nodels ug resp molec n disorde	ers of carbo sm., l eredita in pha ponse Un ular a	etic tec amino ohydrat Pharma ry disc armacog and m ait IV and bio ood gr	hnique s acid me e metal codynam orders wir genomics etabolism ochemica oups., G	tabolism bolism, nics: De th altere , Ecoger n l pathw enetic fa	n, disor disora finitior d drug netics vays of actors	ders of ders o n, drug respon	branche f lipic metab se, Pha	molism maco Ka Seeases	etabolism, , Genetic ogenetics, 4 s: genetic
Inborn err acid me Mucopoly variation Pharmaco Outcome Objective	rors of me tabolism, vsaccharid by the ef genomics 3 Lea 2 4 Des of r immunod lity to co	etabolisi disorc oses an fect of : Anima orn the cribe t netabol eficienc ommon	sm, o rders ind i drug al m e dru the f blism cy d	disord s of Albini ags, He nodels 1g res molec 1 disorde seases,	ers of carbo sm., l eredita in pha ponse Un ular a types	etic tec amino ohydrat Pharma ry disc urmacog and m ait IV and bic ood gr and n	hnique s acid me e metal codynam orders wi genomics etabolism ochemica oups., G nechanis	tabolism bolism, hics: De th altere , Ecogen n l pathw enetic fa ms of g	n, disor disor disor d drug netics vays of actors genetic	ders of ders o n, drug respon inborn in com suscept	branche f lipic metab se, Pha errors mon di tibility,	molism rmacco Ka seases appro	etabolism, , Genetic ogenetics, 4 s: genetic oaches to
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Inborn err acid me Mucopoly variation Pharmaco Outcome Objective Inherited susceptibi demonstra	rors of me tabolism, vsaccharid by the ef genomics 3 Lea 2 4 Des of r immunod lity to co ating gene	etabolisi disorc oses an fect of Anima arn the scribe the metabol eficienc ommon etic susc ease, Ali	sm, o rders ind drug al m e dru e dru the the lism cy d disc scept lzhei	disord s of Albini ags, He nodels ag resp molec n disorde seases, tibility imer d	ers of carbo sm., l eredita in pha conse Un ular a ers, blo types , type isease	etic tec amino ohydrat Pharma ry disc armacog and m and bio ood gr and 1 1 diabo	hnique s acid me e metal codynam orders wi genomics etabolism ochemica oups., G nechanis	tabolism bolism, hics: De th altere , Ecogen n l pathw enetic fa ms of g 2 diabo	n, disor disore efinition d drug netics /ays of actors genetic etes, C	ders of ders o n, drug respon inborn in com suscept	branche f lipic metab se, Pha errors mon di tibility,	molism rmacco Ka seases appro	etabolism, , Genetic ogenetics, 4 s: genetic paches to ension,
Inborn err acid me Mucopoly variation Pharmaco Outcome Objective Inherited susceptibi demonstra coronary a	rors of me tabolism, vsaccharid by the ef genomics 3 Lea 2 4 Des of r immunod lity to co ating gene	etabolisi disorc oses an fect of Anima arn the scribe the metabol eficienc ommon etic susc ease, Ali	sm, o rders ind drug al m e dru e dru the the lism cy d disc scept lzhei	disord s of Albini ags, He nodels ag resp molec n disorde seases, tibility imer d	ers of carbo sm., l eredita in pha oonse Un ular a ers, blo types , type isease n the g	etic tec amino ohydrat Pharma ry disc armacog and m and bio ood gr and 1 1 diabo	hnique s acid me e metal codynam orders wi genomics etabolism ochemica oups., G nechaniss etes, type	tabolism bolism, hics: De th altere , Ecogen n l pathw enetic fa ms of g 2 diabo	n, disor disore efinition d drug netics /ays of actors genetic etes, C	ders of ders o n, drug respon inborn in com suscept	branche f lipic metab se, Pha errors mon di tibility,	k maco maco K k seases appro ypert	etabolism, , Genetic ogenetics, 4 s: genetic paches to ension,
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Inborn err acid me Mucopoly variation Pharmaco Outcome Objective Inherited susceptibi demonstra coronary a Outcome Objective Haematolo	rors of me tabolism, vsaccharid by the ef genomics 3 Lea	etabolisi disorci oses an fect of Anima arn the scribe the netabol eficience ommon etic susce ase, Ali uip kno sorders of gene	sm, c ders al m drug al m drug drug drug drug drug scept lzhei owle netic like mutike mutike disc	disord s of Albini ags, He nodels ag resj molec n disorde seases, tibility imer d edge on e facto ce tha utation	ers of carbo sm., l eredita in pha oonse Un ular a ers, blo types , type isease n the <u>g</u> U rs in c lassem s in hu	etic tec amino ohydrat Pharma ry disc armacog and m ait IV and bio ood gr and 1 diabo gene m nit V commo iia, sic umans,	hnique s acid me e metal codynam orders wir genomics etabolism ochemica oups., G mechanisr etes, type utations n disease ckle cell Human	tabolism bolism, hics: De th altere , Ecogen n l pathw enetic fa ms of g 2 diabe in hum es anemia	n, disor disore efinition d drug netics /ays of actors genetic etes, C an a, hae ndrial o	ders of ders o n, drug respon in com suscept rohn di moglobi diseases	brancho f lipic metab se, Pha errors mon di tibility, sease, h nopathi Loss c	Kalendaria (Construction) Kalendaria (Construct	etabolism, , Genetic ogenetics, 4 s: genetic paches to ension, 2 Molecular action and
Inborn err acid me Mucopoly variation Pharmaco Outcome Objective Inherited susceptibi demonstra coronary a Outcome Objective Haematolo pathology Gain of t	rors of me tabolism, vsaccharid by the ef genomics 3 Lea 6 4 Des of r immunod lity to co ating gene artery dise 4 Equ c 5 Lea ogical dis classes functional	etabolisi disorci oses an fect of : Anima arn the cribe th netabol eficience ommon etic susce ase, Ali ip kno arn gene sorders of gene mutati	sm, c ders al m drug al m drug drug drug drug drug scept lzhei owle metic lik e mu ions	disord s of Albini ags, He nodels 1g resp molec 1 disorde seases, tibility <u>imer d</u> edge on c facto ac tha utation s in hu	ers of carbc sm., l eredita in pha oonse Un ular a ers, blo types , type isease n the g U usrs in c lassem s in hu umans,	etic tec amino ohydrat Pharma ry disc armacog and m ait IV and bic ood gr and r 1 diabo gene m nit V commo aia, sic umans, , Agan	hnique s acid me e metal codynam orders wi genomics etabolism ochemica oups., G nechanist etes, type nutations n disease ckle cell Human maglobi	tabolism bolism, hics: De th altere , Ecoger n l pathw enetic fa ms of g 2 diabe in hum es anemia	n, disor disore disore dinition d drug netics vays of actors genetic etes, C an a, hae ndrial c Disease	ders of ders o n, drug respon in com suscept crohn di moglobi diseases es of co	brancho f lipic metab se, Pha errors mon di tibility, sease, h nopathi Loss c pollagens	seases approvement seas approvement seases approvement seases approvement seases approvement seases approvement seases approvement seases approvement seas approvement seas approvement seas approvement seas approvement seas approvement seas approvement seas approvement seas approvement seas approvement seas approvement seas approvement seas approvement seas approvement seas approvement seas approvement seas approvement seas approvement seas approvement seas approvement s approvement s approvement s approvement s approveme	etabolism, , Genetic ogenetics, 4 s: genetic oaches to ension, 2 Molecular nction and ngle gene
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Suggested Readings:

Chattopadhyay, I. (2018). Fundamentals of Genetiics (1st ed.). Vinod Kumar Jain, Scientific International (Pvt.) Ltd.

Rimoin, D. L., Pyeritz, R. E., & Korf, B. (Eds.). (2013). Emery and Rimoin's essential medical genetics. Elsevier.

Turnpenny, P. D., & Ellard, S. (2016). Emery's Elements of Medical Genetics E-Book. Elsevier Health Sciences.

Maloy, S.R. Cronan, J.E and Freifelder (D.J.B). (2002) Modern Genetic Analysis, 2 nd (ed.,)

Freifielder, J.B. (1993). Essentials of Molecular Biology, 2 nd (ed.,)

Amita Sarkar (2011). A Textbook of Human Genetics (ed.,)

Watson et al.,(2014). Molecular Biology of the Gene, 7 th (ed.,)

Online resources

Medical Genetics- https://shop.elsevier.com/books/medical-genetics/jorde/978-0-323-59737-1 Essential Medical Genetics, Includes Desktop Edition, 6th Edition- <u>https://www.wiley.com/en-fr/Essential+Medical+Genetics,+Includes+Desktop+Edition,+6th+Edition-p-9781405169745</u> K1-Remember, K2-Understand, K3- Apply K4- Analyze, K5-Evaluate, K6- Create

USUB BOD

	Course Outcome VS Programme Outcomes										
СО	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10	
CO-1	M(2)	S(3)	S(3)	S(3)	M(2)	S(3)	M(2)	S(3)	M(2)	S(3)	
CO-2	S(3)	L(1)	S(3)	S(3)	M(2)	S(3)	L(1)	M(2)	M(2)	M(2)	
CO-3	S(3)	S(3)	M(2)	S(3)	L(1)	S(3)	M(2)	L(1)	L(1)	L(1)	
CO-4	S(3)	L(1)	M(2)	S (3)	L(1)	S (3)	L(1)	L(1)	M(2)	S(3)	
CO-5	S(3)	L(1)	M(2)	S (3)	S(3)	S (3)	M(2)	M(2)	L(1)	M(2)	
W.AV	2.8	1.8	2.4	3	1.8	3	1.6	1.8	1.6	2.2	

1. Low, 2. Medium, 3. S	Strong
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Course Outcome VS Programme Specific Outcomes									
CO	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5				
CO-1	M(2)	S(3)	S(3)	S(3)	M(2)				
CO-2	S(3)	L(1)	S(3)	S(3)	M(2)				
CO-3	S(3)	S(3)	M(2)	S(3)	L(1)				
CO-4	S(3)	L(1)	M(2)	S(3)	L(1)				
CO-5	S(3)	L(1)	M(2)	S(3)	S(3)				
W.AV	2.8	1.8	2.4	3	1.8				

			Semester-II					
Core		rse code:	Pharmacology	Т	Credits:	5 Hours:5		
	5	08202						
	. 1	A	Unit-I					
	Objective 1Acquire knowledge about the principles of pharmacologyDrug Nomenclature, Generic name, Fixed dose combinations, dosage forms and Posology of Drugs.							
			gs. Routes of administration of drugs (
Ŭ			ages. Absorption, Distribution, Biotransforr t, Factors modifying dose of a drug. Mec					
			c principles, mechanism of action and type					
		-	ial agonist, inverse agonist, antagonist).	S OI DI	ug receptor	is and		
Outcom			basic and clear understanding over the co	meants	and			
Outcom	6 1		ntal principles of pharmacology	ncepts	anu	K2		
		Tunuunit	Unit-II			IX 2		
Objective	2	I oorn th	e mechanism of action of several drugs	over th	o various	nhysiological		
Objective	: _	systems	e mechanism of action of several drugs	over th	e various	physiological		
Pharmaco	logical	C C	on of drugs: Pharmacology of drugs acting	y on aut	tonomic (cl	holinergic and		
			eral and central nervous systems (anesthet					
•	•	· • •	agents). Drugs acting on the respiratory	•				
			ints and anticoagulants. hematonics, Ca					
			ertensive agents, peripheral vasodilators an					
Ŭ	•	• •	d applications of drugs used as immuno	•				
			sants, narcotic analgesics, antagonists, n					
			adrenergic receptor blockers, neuron blocke					
neuromus				10, guile				
Outcome		Acquire	knowledge over the basic principl	es of	systemic	K4		
0	-	pharmaco		••• •••	5,5000000			
			Unit III					
Objective	3	Describe	the basics of Chemotherapeutic drug	s and	steps inv	olved in the		
N			tion of new drugs	<u> </u>				
			of infection, infestation and neoplastic dis					
	-	÷	Drug addiction and abuse. Drug discovery	-		.		
			ug Discovery process, Early Drug Discove	-		-		
<u> </u>			cal assays, cell cultures and various anin					
· ·			or alternative models.) and Clinical Phases			, · ·		
			A). Gene Therapy $-$ An introduction and					
		s in drug di	ransgenic animal models in the developme	III OI IIE	ew drugs. v	Leff filles and		
Outcome		-	the cascade mechanisms underlying the	nrocos	s ovor	K4		
Outcome	5	-	tion of pharmaceutically active compound	-	S UVEI	174		
		aunina	Unit IV					
Objective	4	Know the	e principle of identification, continuous m	onitorir	ng and han	dling of		
Sojective	•		vents that arouses owing to the drug		-5 and nan	wing vi		
Pharmaco	vigilan		detection systems and uses advanced data	analyti	cs. Importa	ance of safety		
	-	-	HO international drug monitoring program	-	-	-		
	-		and PSUR. Investigation of ADR/ADE ar		-	-		
			Definitions and classification of adverse			-		
			sality assessment, Severity and seriousness	-				
			Aanagement of adverse drug reactions. Po			-		
-	•		ilization studies, therapeutic audit, essential		-			
Outcome		Realize	the necessity for continuous detection,	-	-	K2		
			ding and prevention of adverse effects of					
L				<u> </u>				

Unit V								
Objective 5 Highlights the standards to meet the regulatory standards and prohibition of Drugs								
Commonly used animals in pharmacological research, limitations of animal tests, Animal handling and animal care Methods (anaesthetizing, restraining, blood collection and euthanasia.). Institutional Animal Ethics Committee, CPCSEA guidelines for breeding and stocking of animals, performance of experiments, transfer and acquisition of animals for experiment. Drugs and Cosmetics Act, 1940 and its rules 1945, Legal definitions of schedules to the Act and Rules. Classes of drugs and cosmetics prohibited from import Manufacture and sale of certain drugs with Offences and penalties., narcotic drugs and psychotropic substances Act-1985 and Rules. Manufacture of drugs for test, examination, analysis and commercial use and requirements to adopt and follow Regulatory commitments (ICH,								
GCP, PICS, WHO GMP, GxP) Outcome 5 Understand the key concepts over necessary of ethical clearance to carry out animal studies								
 Suggested Readings: August, J.T., Anders, M.W., Murad, F., & Coyle, J.C (eds.) (1994). Advances in Pharmacology (1st ed.). Academic Press Katzung, B. G. (2017). Basic and clinical pharmacology. McGraw-Hill Education. Derasari&Brahmankar (2015). Elements of Pharmacology.(ed.,) Bose, B.C.(2015) Text Book Pharmacology (ed.) Goodman and Gilman's,(2012). The Pharmacological Basis of Therapeutics (ed.) Tripathi, K.D. (2020). Essentials of Medical Pharmacology, JAYPEE Brothers Medical Publishers (P) Ltd, New Delhi. (ed.) Sharma H. L., Sharma K. K., (2018). Principles of Pharmacology, Paras medical publisher (ed.) Online resources Pharmacology Books, Ebooks And Journals- https://www.uk.elsevierhealth.com/medicine-and- 								
<u>surgery/pharmacology</u> K1-Remember, K2-Understand, K3- Apply K4- Analyze, K5-Evaluate, K6- Create								

Course Outcome VS Programme Outcomes PO-2 PO-3 PO-4 PO-5 CO PO-1 PO-6 PO-7 PO-8 **PO-9** PO-10 M(2) M(2) M(2) S(3) CO-1 S(3) S(3) M(2) M(2)M(2) M(2) M(2) CO-2 S(3) S(3) L(1) L(1) M(2) M(2) M(2) M(2) L(1) M(2) CO-3 S(3) M(2) M(2) M(2) S(3) L(1) L(1) L(1) M(2) CO-4 S(3) L(1) L(1) L(1) L(1) M(2) L(1) M(2) S(3) L(1) CO-5 L(1)S(3) S(3) M(2) S(3) M(2) L(1) L(1) L(1) L(1) W.AV S(3) 2.2 1.8 2.2 1.8 1.4 1.6 2 1.6 1.4

CARDIN EXCELLENC

1. Low, 2. Medium, 3. Strong

Cours	Course Outcome VS Programme Specific Outcomes								
CO	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5				
CO-1	S(3)	S(3)	M(2)	M(2)	M(2)				
CO-2	S(3)	M(2)	S(3)	L(1)	L(1)				
CO-3	S(3)	M(2)	M(2)	M(2)	M(2)				
CO-4	S(3)	L(1)	L(1)	L(1)	L(1)				
CO-5	S(3)	S(3)	L(1)	L(1)	M(2)				
W.AV	3	2.2	1.8	1.4	1.6				

1. Low, 2. Medium, 3. Strong



		Semester-II								
Core	Course code: 508203	Practical –III Medical Genetics, Pharmacology	Р	Credits:4	Hours:8					
Course Objectives	> Ph	ntification of drugs by morphological chara ysical and chemical tests for evaluation of d section and analysis of source organs flora	rug v	wherever ap						
	dru > Ide > Iso	 drug Identification of fibres and surgical dressings Isolation of nature products such as caffeine, starch, emetine, fixed oils 								
	 Constructi Mounting Mitosis in Identificat Diagnosis Demonstra Study of h Animal ha Routes of Topical ap Analgesic Determina 	on of Barr body in buccal epithelial cell, of biochemical disorder- Alkaptonuria tion on structure and molecular organization ereditary disorder with the aid of chromoson ndling and precautions drug administration plication of atropine on rabbit eye effect of diclofenac in mice using hot plate tion and calculation of lethal dose (LD50) v	n of me k meth alue	chromosom aryotyping od						
	 14. Collection, processing, identification and extraction of bioactive crude extraction from marine fauna and flora 15. Quantitative and Qualitative analysis of phytochemicals from marine fauna andflora 16. Extraction of neutraceuticals from marine sources 									
Course Outcomes (CO)	CO1- To knov CO2- Familar CO3- Underst CO4- Learn th	w the family history se the mutations on chromosomes and the scientific validation and quality eval e routes of administration of drugs on of safe and efficacy of drugs	luatio	on of drugs						
Laboratory-Ma	ces: nd Genetics Lab Ma nual-17-18.pdf.Gen	nual. https://sjce.ac.in/wp-content/uploads/2018/0 etics Laboratory Manual. i/viewcontent.cgi?article=1008&context=ny_o		ll-Biology-G	enetics-					

https://academicworks.cuny.edu/cgi/viewcontent.cgi?article=1008&context=ny_oers. Molecular Genetics Laboratory Procedures. https://bpb-us_e1.wpmucdn.com/wordpressua.uark.edu/dist/1/802/files/2020/07/MANUAL.pdf

	Course Outcome VS Programme Outcomes									
СО	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10
CO-1	S(3)	M(2)	S(3)	S(3)	M(2)	M(2)	M(2)	S(3)	M(2)	M(2)
CO-2	S(3)	M(2)	S(3)	S(3)	M(2)	M(2)	M(2)	M(2)	M(2)	L(1)
CO-3	S(3)	M(2)	M(2)	S(3)	S(3)	L(1)	L(1)	L(1)	S(3)	M(2)
CO-4	S(3)	L(1)	M(2)	S(3)	M(2)	L(1)	M(2)	S(3)	M(2)	L(1)
CO-5	S(3)	S(3)	L(1)	S(3)	L(1)	S(3)	L(1)	M(2)	L(1)	L(1)
W.AV	3	2	2.2	3	2	1.8	1.6	2.2	2	1.4

Course Outcome VS Programme Specific Outcomes								
CO	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5			
CO-1	S(3)	M(2)	S(3)	S(3)	M(2)			
CO-2	S(3)	M(2)	S(3)	S(3)	M(2)			
CO-3	S(3)	M(2)	M(2)	S(3)	S(3)			
CO-4	S(3)	L(1)	M(2)	S(3)	M(2)			
CO-5	S(3)	S(3)	L(1)	S(3)	L(1)			
W.AV	3	2	2.2	3	2			

1. Low, 2. Medium, 3. Strong



		Semester-II									
Core	Course code:	Practical –IV Techniques in	Р	Credits:4	Hours:6						
	508204	Biomedical sciences-II									
Course	➢ Learr	the basic concepts and applications of	of inst	ruments app	lied in						
Objectives	bioch	emical analysis.	minati	on of al volu	as of amino						
	acids	arn the preparation of buffers and detern	mnau	on of pr valu	les of amino						
		 Recognize the principles and procedures behind the chromatographic 									
		techniques									
		Clear understanding the principles of spectrometric analysis and their									
	practi	cal application explored.	· · · · ·	1 (1	1.1.1						
		ire the features of chromatography tech cations	inique	s and their	biological						
		nd applications of pH meter, Col	orime	ter Spectro	nhotometer						
	Centrifuge	nd applications of pri meter, con		ier, speeno	photometer,						
	Preparation of	of buffers									
		f Henderson-Hasselbach equation									
		f pKa values in acid-base titrations									
		n of pI value of amino acids		_							
		f amino acids and sugars by Paper chron	natog	raphy							
		of plant pigments by TLC									
		f organic compounds by column chroma on of GC, HPLC, AAS,NMR,FTIR and									
	• Demonstratio	on of GC, HPLC, AAS, NMR, FTIR and	wass	spectroscopy	/						
Suggested R	eadings:	ALADAPEA DRIVERSITY									
Baynes, J.	W., & Dominiczak,	M. H. (2014). Medical Biochemistry (e	d.,)								
		81). Introduction to biomedical equipme									
		& Schenken, J. R. (1994). Laboratory ir									
		(2001). Introduction to biomedical equi		t technology.	(ed.,)						
		ook of biomedical instrumentation. (ed.,		ad Laboustor							
	ues (1st ed.).	(2016). Clinical chemistry: Fundament	lais ai	nd Laborator	y						
Technic	lues (Ist ed.).										
		Manual of Biochemistry Joy P P, Sur									
	prsvkm.kau.in/sites/d	efault/files/documents/prsvkm_laborator	y_ma	nual_of_bioc	hemistry.						
pdf.											
		stry. Https://Jru.Edu.In/Studentcorner/L	ab-Ma	inual/Bpharm	1/2nd-						
	b%20manual%20-%		of	I waatan							
Course C (CO)		 Learn the principles and applications Understand the role of buffers in bio 									
		3- Hands on experience on the biochem			nniques						
		4- Know the principle, instrumentation a			inques						
	0	chromatography	up	prioution of							
	CO5	- Gain knowledge on the principle, inst	rumer	ntation and a	pplication						
		of spectroscopy									

	Course Outcome VS Programme Outcomes									
CO	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10
CO-1	S(3)	S(3)	M(2)	S(3)						
CO-2	S(3)	M(2)	S(3)	L(1)	L(1)	M(2)	M(2)	L(1)	M(2)	M(2)
CO-3	S(3)	L(1)	L(1)	M(2)	M(2)	S(3)	L(1)	M(2)	L(1)	L(1)
CO-4	S(3)	M(2)	M(2)	L(1)	L(1)	M(2)	L(1)	L(1)	M(2)	S(3)
CO-5	S(3)	S(3)	L(1)	L(1)	M(2)	L(1)	S(3)	L(1)	L(1)	M(2)
W.AV	3	2.2	1.8	1.4	1.6	2	1.8	1.4	1.6	2.2

Cours	Course Outcome VS Programme Specific Outcomes								
CO	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5				
CO-1	S(3)	S(3)	M(2)	S(3)	M(2)				
CO-2	S(3)	M(2)	S(3)	S(3)	L(1)				
CO-3	S(3)	L(1)	L(1)	S(3)	M(2)				
CO-4	S(3)	M(2)	M(2)	S(3)	L(1)				
CO-5	S(3)	S(3)	L(1)	S(3)	M(2)				
W.AV	3	2.2	1.8	3	1.6				



			Semester-II						
DSE-2	Course code: 508503		Forensic Science	Т	Credits:3	Hours:3			
			Unit-I	I	-	4			
Objective 1 Understand the basic concepts of forensic science and their scope.									
Definition and scope of forensic science, history and development of forensic science. Scope and									
development of forensic science in India, growth of core laboratories, set up in country									
Outcome	1 Acquire kno	wledge on th	e forensic laboratories a	nd developm	ient. F	K2			
			Unit-II						
Objective 2	2 Learn crime	scene procee	lures and types of crime	s.					
Introductior	to crime, soci	ological aspe	ct in society, types of c	rimes, crimes	in India, c	rime scene			
managemen		v	, protection of crime sc						
collection o			ene management in manm						
Outcome			forensic scientists.			K4			
	I		Unit III						
Objective	3 Describe the	various divis	sions of crime investigat	ion					
Duties of forensic scientist, various divisions of crime investigation – toxicology, biology, serology, chemistry, physics ballistics prohibition document and other divisions.									
			research methods an		ssues in 1	K4			
		S	Unit IV	2					
Objective	4 Gain knowle	dge on the b	asic concepts of psychological	ogy					
analysis, b		scillation sign	science laboratory – DNA nature proficiency (BEOS 2.						
Outcome 4	Know-how	the methods	of <mark>cyb</mark> er foren <mark>sic</mark> s		1	K2			
			Unit V						
Objective	5 Familiariz	<mark>e the</mark> special	ized facilities in national	fo <mark>rensic</mark> lab	oratory				
professional psychology.	ls psychology; T	The science a	psychology, modern pe nd research methods, pro	fessional and					
Outcome 5		e various div	isions in forensic labora	tories	k	K5			
Basu, R Bertino, Leela D Heath, Y	F. A. (2007). Pri . (2009). Fundan A. J. (2012). Fo pubey (2018). Fo W. (2018). Psych Sharma, R.K. (2) purces	mentals of Fo orensic Scienco prensic Scienco hology Resea 008). Practica	rch Methods: Connecting and Viva in Forensic Me	estigations (e Research to	d.,) Students' Liv				
Forensic Se	Forensic Science: A Multidisciplinary Approach- https://onlinelibrary.wiley.com/doi/book/10.1002/9783527693535								
Forensic S	cience Resource	es: Books/eb	ooks- https://guides.uflib.u	ufl.edu/c.php?	<u>g=147333&</u>	<u>p=968747</u>			
K1-Remen	iber, K2-Under	stand, K3- A	pply K4- Analyze, K5-E	Evaluate, K6-	Create				

	Course Outcome VS Programme Outcomes									
CO	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10
CO-1	S(3)	S(3)	S(3)	M(2)	M(2)	M(2)	M(2)	M(2)	M(2)	S(3)
CO-2	M(2)	S(3)	M(2)	M(2)	M(2)	M(2)	L(1)	L(1)	M(2)	M(2)
CO-3	M(2)	S(3)	L(1)	L(1)	S(3)	L(1)	M(2)	M(2)	L(1)	L(1)
CO-4	M(2)	S(3)	S(3)	M(2)	M(2)	L(1)	L(1)	L(1)	M(2)	S(3)
CO-5	L(1)	S(3)	M(2)	L(1)	L(1)	S(3)	L(1)	M(2)	L(1)	M(2)
W.AV	2	3	2.2	1.6	2	1.8	1.4	1.6	1.6	2.2

Cours	Course Outcome VS Programme Specific Outcomes								
CO	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5				
CO-1	S(3)	S(3)	S(3)	M(2)	M(2)				
CO-2	M(2)	S(3)	M(2)	M(2)	M(2)				
CO-3	M(2)	S(3)	L(1)	L(1)	S(3)				
CO-4	M(2)	S(3)	S(3)	M(2)	M(2)				
CO-5	L(1)	S(3)	M(2)	L(1)	L(1)				
W.AV	2	3	2.2	1.6	2				



			Semester-II			
DSE 2	Course code: 508504		Artificial Organs	Т	Credits:3	Hours:3
		-	Unit-I			
Objective			tive medicine and organ	-		2
•	U U		medicine, biomaterial co ation of artificial organs.	oncentra	tion, outlook	for organ
Outcome	1 Understand	d the artificial	l organs and their mech	anisms	5	K2
			Unit-II			
			art and circulatory assi			
of valve pro categories,	ostheses, thromb intra- aortic ball	ous deposition, oon pump, per	ces- design of artificial h durability, mechanical ci ccutaneous cardiopulmona	irculato ary byp	ry assistance, ass.	two main
Outcome	2 Acquire kr devices	iowledge abou	t artificial lungs and blo	od gas	exchange	K4
			Unit III			
Objective	-	0	nd cardio pulmonary by	-		-
cardio puli carbon-di-	monary bypass, oxide transport.	ECMO, comp	e devices- artificial lung arison of artificial lungs	and nat	tural lungs, o	xygen transport,
Outcome	-		ng concepts that charac	eterize	the quality	of K4
	imaging te	chniques	Unit IV			
Objective	1 Describe t	a ranal trans	plantation and dialysis.			
					4-4:	
			Artificial kidney: renal tra itoneal dialysis equipme			
therapy, th			insulin administration sys			
Outcome	4 Versed in ultrasound		of image formation, ca	pture a	and display o	of K2
		1	Unit V			
Objective	5 Know-how	the artificial	blood and liver			
classificatio		on of substitu	icial blood: plasmaphere tes. Artificial liver: liver n replacement.			
Outcome	5 Learn abo	ut the artificia	l lung and cardio pulm	onary	bypass	K5
Miller, E Ong, J. I enginee Poole, D. agents. Cromwel and mea Drexler, Y (ed.,) Online reso Artificial (Tissue Eng	(2017). Biomat. . G. (2006). Arti L., Appleford, I ring application . L., & Mackwo (ed.,) 1, L., Weibell, H asurements (ed., W., & Fujimoto, ources Drgans- https:// gineering for A	ficial Organs. M. R., & Man s. (ed.,) rth, A. K. (201 F. J., Pfeiffer, 1 J. G. (2008). (link.springer.co Artificial Orga	and tissue engineering: pr (ed.,) i, G. (2014). Introduction 0). Artificial Intelligence: E. A., & Usselman, L. B. Optical coherence tomogra om/book/10.1007/978-3-0 ans: Regenerative Media ary.wiley.com/doi/book/1	n to bid founda (1973) aphy: te 031-016 icine, S	omaterials: ba ations of com). Biomedical echnology and (11-0 Smart Diagne	asic theory with putational instrumentation applications.
			n/us/en/ebook/artificial-on pply K4- Analyze, K5-E	-	K6 Cmart	
K1-Veinen	iber, K2-Ullder	stallu, NJ- A	ppiy R4- Allalyze, R5-E	vaiual	e, NU- Creat	¢

		Co	ourse Ou	itcome	VS Prog	ramme	Outcome	s		
CO	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10
CO-1	M(2)	S(3)	M(2)	M(2)	M(2)	S(3)	M(2)	M(2)	M(2)	M(2)
CO-2	S(3)	M(2)	L(1)	L(1)	M(2)	S(3)	M(2)	M(2)	L(1)	L(1)
CO-3	S(3)	L(1)	M(2)	M(2)	L(1)	S(3)	S(3)	L(1)	M(2)	M(2)
CO-4	S(3)	L(1)	L(1)	L(1)	L(1)	S(3)	M(2)	L(1)	L(1)	L(1)
CO-5	S(3)	M(2)	M(2)	M(2)	S(3)	S(3)	L(1)	S(3)	L(1)	M(2)
W.AV	2.8	1.8	1.6	1.6	1.8	3	2	1.8	1.4	1.6

1.	Low,	2.	Medium,	3.	Strong
	,			-	

Cour	se Outcom	e VS Progra	amme Spec	cific Outco	omes
СО	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5
CO-1	M(2)	S(3)	M(2)	M(2)	S(3)
CO-2	S(3)	M(2)	L(1)	L(1)	S(3)
CO-3	S(3)	L(1)	M(2)	M(2)	S(3)
CO-4	S(3)	L(1)	L(1)	L(1)	S(3)
CO-5	S(3)	M(2)	M(2)	M(2)	S(3)
W.AV	2.8	1.8	1.6	1.6	3



		Semester-III			
Core	Course code: 508301	Toxicology	T	Credits:	5 Hours:5
		Unit-I			
-		basics underlying toxicology			
		rinciples of toxicology, mechanisms an			
-	• •	toxicity and its measurement: acu			
^	•	s of exposure of toxicants (oral, inl			· ·
-	-	of absorption, distribution, excretion; ma			
		ase 2 reactions; control of metabolism schemical basis of toxicity: mechanism of		-	-
		actors influencing toxic effects.	toxicity	. Distuita	
Outcome 1	Imparts descript	tive idea over the fundamentals of to	kic subs	tances	K2
	and mechanism of	Unit-II			
Objective	2 Describe the cate	egory of toxins and its effects			
-		ial toxins and animal toxins. Synthetic	organic	compound	ls: chemical
		in the work place, solvents, vapo	-	-	
		xicology, food toxicology. Toxic respo			
	-	ystem, endocrine system, heart and vascu			
-	-	ty: Disturbance of excitable membrane fu	-		
•		o cellular macromolecules.	,		
Outcome 2	8	he kinds of naturally occurring and our different biological systems	syntheti	c toxic	K4
		Unit III			
Objective	3 Elicit the threats	posses <mark>sed by the carcin</mark> ogenic and xen	obiotic	agents	
Classificatio	on of carcinogens	(known and suspected carcinogens), typ	pes of c	arcinogen	s (chemical,
physical an	nd onco-genic). Ca	ategorization of carcinogens (Category	1A, 1I	3 and 2).	Multistage
		of action of chemical carcinogens			
		organic carcinogens (arsenic, beryllium,			
		, types, mutagenic agents and effects. C tagenicity. Xenobiotics: Introduction, cla			
		reign compounds: Direct toxic action: t			
		armacological, physiological and biochem			
		lge over the several carcinogenic substa			K4
	mankind				
		Unit IV			
Objective4	Acquire knowled	lge over the several poisons and its elin	nination	methods	
Types of p	oison, clinical sign	s and symptoms, diagnosis, managemer	nt and n	nedicolega	l aspects of
		oisons; neural poisons; somniferous;			
		sphyxiants; drug abuse. Treatment of po			
	-	addict forming drugs. Incidence of act	-		vention and
	· ·	l symptoms and management of barbitura		-	_
U 1		und and lead, mercury and arsenic pois	soning. 1	Biomarker	s criteria of
÷		ation of toxicity interactions		. 1	
Outcome 4		cal status of variety of poisons and th	nerapeut	tic	K2
	approaches to ex	posed poisoning			

	Unit V
Objective 5	Learn about the multiple toxicity evaluation methodologies
Evaluation of	toxicity, median effective dose, median toxic dose, median lethal dose, therapeutic index
and therapeut	ic window. Toxicity testing : Short-term tests for mutagenicity, genetic toxicity and
mutagenesis a	ssay: bacterial mutation tests-reversion test, ames test, fluctuation test, and eukaryotic
mutation test.	Biochemical mechanisms of tissue toxicity, organ, neurotoxicity; gastro-intestinal
toxicity, skin	toxicity/ photosensitivity, genetic toxicology, reproduction toxicity, carcinogenicity
studies ,exagg	erated and unwanted toxicological effects, single dose, repeat dose toxicity studies, safety
pharmacology	, studies (including segment I, II, and III).
Outcome 5	Learn the key concept's in testing methodologies of toxicants and K5 mutagens
Suggested Re	adings:
Parikh C.k	K. Parikh 's (2007) Textbook of Medical Jurisprudence and Toxicology, 6th ed.,
Franklin, C	C.A (2020).Modi's medical Jurisprudence and Toxicology .21st ed.,.
H.P. Rang	, M.M. Dale, J.M. Ritter & P.K. Moore (2012). Pharmacology. (ed.,)
Bertram G	. Katzung. (2008). Basic and Clinical Pharmacology. (ed.,)
W.	C. Bowman, M. J. Rand.(2016). Text book of Pharmacology (ed.,)
Online resour	
	Of Modern Toxicology-
	unp.ac.id/file/abstrak_kki/EBOOKS/A%20textbook%20of%20Modern%20Toxicology.
	um=email&utm_source=transaction
	of Modern Toxicology- https://onlinelibrary.wiley.com/doi/book/10.1002/0471646776
KI-Remembe	r, K2-Understand, K3- Apply K4- Analyze, K5-Evaluate, K6- Create

	Course Outcome VS Programme Outcomes											
CO	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10		
CO-1	S(3)	M(2)	M(2)	M(2)	M(2)	S (3)	M(2)	M(2)	M(2)	M(2)		
CO-2	M(2)	L(1)	L(1)	M(2)	L(1)	S (3)	M(2)	L(1)	L(1)	M(2)		
CO-3	L(1)	M(2)	S(3)	L(1)	M(2)	S(3)	L(1)	M(2)	M(2)	L(1)		
CO-4	M(2)	L(1)	S(3)	L(1)	L(1)	S(3)	L(1)	L(1)	L(1)	L(1)		
CO-5	S(3)	S(3)	S(3)	S(3)	L(1)	S(3)	S(3)	L(1)	M(2)	S(3)		
W.AV	2.2	1.8	2.4	1.8	1.4	3	1.8	1.4	1.6	1.8		

Cours	se Outcom	e VS Progra	mme Spee	cific Outc	omes
CO	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5
CO-1	S(3)	S(3)	M(2)	M(2)	M(2)
CO-2	M(2)	S(3)	L(1)	M(2)	L(1)
CO-3	L(1)	S(3)	S(3)	L(1)	M(2)
CO-4	M(2)	S(3)	S(3)	L(1)	L(1)
CO-5	S(3)	S(3)	S(3)	S(3)	L(1)
W.AV	2.2	3	2.4	1.8	1.4

^{1.} Low, 2. Medium, 3. Strong

					Sem	ester-III			
Core		urse code: 508302		Me	dical On	cology	T	Credits: 5	Hours:5
						Unit-I			
								– cell interacti	
			•	•		•		•	ns, invasions by
		s, angiogene	esis, m	orphog	ens, mec	chanism of de	regulation	n of cell cycle	during cancer,
Apoptosi Outcon	ne 1	Understand cancer.	the m	nechani	ism of d	eregulation o	f cell cyo	cle during	K2
	r				τ	J nit-II			
Objectiv	ve 2	Know the d	egree	of mali	ignancy a	and types of c	chromoso	mal translocat	tions
WHO	classif	fication,	stagir	ig and g	grading, o	degree of mali	gnancy, t	disease, tumo ypes of chromo tors- Src, Wnt,	
Outcome 2 To learn the relationship between oncogene products and growth factors								K4	
					U	nit III			
Objectiv	ve 3	Learn the o	ncoger	nic mut		n growth prom	noting p	roteins	
cycle co genes),	ontrol, necros	evasion of sis.	growt	h inhib	oitory sig	nals, cancer g	genes (on	-	loss of cell mor suppressor
Outcom	e 3	Know-how t	the mu	itations		g loss of cell c	ycle cont	rol	K4
						Init IV			
•				0	0	ed delivery of		0	
	d deliv	very of antication						cancer deliver tems for the de	
Outcom	e 4	Gain knowle	edge tl	he criti	ical anal	y <mark>sis</mark> of cancer	therapy	and vaccines	K2
				3	Unit V		1.7		
Objectiv	ve 5	Fo learn the	e vario	us type	es of the	rapy in cance	r treatmo	ent	
						immunotherap of cancer ther		entional chemot	herapy,
Outcom	e 5	Acquire the	cance	r diagr	nosis and	l treatment			K5
textb Leon Paul Cava Cava of me Rudle	oook of idas C M. (20 lli, F., Prir lli, F., edical oph K biol	f oncology (2. Platanias, (222). Silvern Kaye, B.K., nciples and p Kaye, S. B oncology. (e Lenhard, Na ogy. (ed.,)	(3 rd ed. (2022) nan,Or , Hanso practice 3., Han ed.,) anoma	,) Advar acologic en, H.H e of onc sen, H. terials f	nces in or c Imaging I., Armita cology (4 H., Arm for cance	ncology, 1 st ed g: A Multidisc age, O.J., Picca th ed.). nitage, J. O., & r diagnosis W	iplinary A art, J.M. d & Piccart ang, E. (2	Approach, (2nd & Gebhart. (20	09). Cancer- 2009). Textbook ystems
Anto	nio Ru	isso, <u>Marc P</u>	<u>eters</u> ,	<u>Lorena</u>	Incorvai	<u>a, Christian Ro</u>	<u>lfo</u> , (202	1). Practical Me	edical
		cology Text						,	
		0,		. 77					

Online resou	irces									
Textbook	of	Medical	Oncology-	https://www.routledge.com/Textbook-of-Medical-						
Oncology/Cavalli-Kaye-Hansen-Armitage-Piccart-Gebhart/p/book/9780415477482										
Oncology/Cancer: Books- https://uscmed.sc.libguides.com/c.php?g=377964&p=2558699										
K1-Rememb	K1-Remember, K2-Understand, K3- Apply K4- Analyze, K5-Evaluate, K6- Create									

	Mapping Course Outcome VS Programme Outcomes											
СО	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10		
CO-1	S(3)	S(3)	M(2)									
CO-2	S(3)	M(2)	L(1)	M(2)	L(1)	M(2)	M(2)	L(1)	L(1)	M(2)		
CO-3	S(3)	S(3)	S(3)	L(1)	M(2)	S(3)	L(1)	M(2)	M(2)	L(1)		
CO-4	S(3)	M(2)	S(3)	L(1)	L(1)	M(2)	L(1)	L(1)	L(1)	L(1)		
CO-5	S(3)	L(1)	S(3)	S(3)	L(1)	L(1)	S(3)	L(1)	M(2)	S(3)		
W.AV	3	2.2	2.4	1.8	1.4	2	1.8	1.4	1.6	1.8		

Cours	e Outcome	e VS Progra	amme Spe	cific Outc	omes
CO	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5
CO-1	S(3)	S(3)	M(2)	M(2)	M(2)
CO-2	S(3)	M(2)	L(1)	M(2)	L(1)
CO-3	S(3)	<mark>S(3</mark>)	S(3)	L(1)	M(2)
CO-4	S(3)	M (2)	S (3)	L(1)	L(1)
CO-5	S(3)	L(1)	S(3)	S(3)	L(1)
W.AV	3	2.2	2.4	1.8	1.4

Semester-III											
Core	Course code:	Practical V- Taxicology, Madical	Р	Credits:4	Hours:8						
	508303	Oncology									
Course		arn the routes of administration in animal 1									
Objectives		derstand the effect of drugs action and ger	neral and	esthesia.							
		equire the acute toxicity in given drugs. in knowledge on the study of cell culture t	echniqu	60							
	\rightarrow Kr	now-how the study of the various biologica	and bi	ochemical m	arkers						
		andling and precautions									
		routes of administration									
		plication of atropine and pilocarpine on ra	bbit eye	;							
	U	effect of diclofenac on mice or rat									
		effect of general anaesthesia with ketamine	e								
		the acute toxicity of a given drug									
		tion and calculation of TD 50/ TC 50 and 1									
		ation of the detection of organophosphorou	us pestic	cides in biolo	gical						
	sample	n of modio, core for primary call culture									
	Preparation of media, sera for primary cell culturePreparation of established cell lines										
		ing and viability									
	 Preservation 										
	 Demonstra 										
		of cell lines- MCF and VERO									
		city assay- MTT assay, apoptosis assay, ne	utral red	d assav							
	•	dical scavenging assay		assaj							
	LDH (Lactate dehydrogenase assay)										
		DNA fragmentation assay									
	Ultrasound imaging										
	Principles and production of X-rays										
Suggested R	•										
		rinciples of Toxicology Testing (2nd ed.).									
Katzı	ung, B. G. (2017).	Basic and clinical pharmacology. (ed.,)									
		v, O. P. (2014). The essentials of forensic n									
		B., Hansen, H. H., Armitage, J. O., &	k Picca	rt-Gebhart,	M.						
		edical oncology.(ed.,)	(1)								
		2010). Nanomaterials for cancer diagnosis (aterials science and tissue engineering: prin		nd mathada	(ad)						
		M. R., & Mani, G. (2014). Introduction									
	engineering applic			materials. Das	sie theory						
	0 0 11	nour, E. R. (2003). Medical imaging physic	cs. (ed.))							
		mental Techniques In Cell Culture. Labo									
		com/deepweb/assets/sigmaaldrich/marketing	g/global/	documents/42	25/663/						
	ntal-techniques-in-		Cultu		Manual						
Animal	Cell	And Tissue n/web_assets/srm_mainsite/files/files/BT%2	Cultu		Manual.						
		20AND% 20TISSUE% 20CULTURE% 20L									
Surachai			Animal	Cell	Culture.						
	,	tents/Lec%20no%201(2).pdf.	IIIIIiii	con	Culture.						
-											
Course		and the routes of administration of drugs									
Outcomes		the scientific validation of drugs									
(CO)	CO3- Know - CO4- Underst	how the side of the drugs and the different kinds of bio imaging tech	nniques								
	CO5- Recogn	ize the cell culture techniques for the evalu	ation of	drugs							

	Course Outcome VS Programme Outcomes										
CO	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10	
CO-1	S(3)	M(2)	S(3)	S(3)	M(2)	M(2)	M(2)	M(2)	M(2)	M(2)	
CO-2	S(3)	M(2)	S(3)	S(3)	L(1)	M(2)	M(2)	L(1)	L(1)	M(2)	
CO-3	S(3)	M(2)	M(2)	S(3)	M(2)	S(3)	L(1)	M(2)	M(2)	L(1)	
CO-4	S(3)	L(1)	M(2)	S(3)	L(1)	M(2)	L(1)	L(1)	L(1)	L(1)	
CO-5	S(3)	S(3)	L(1)	S(3)	L(1)	L(1)	S(3)	L(1)	M(2)	S(3)	
W.AV	3	2	2.2	3	1.4	2	1.8	1.4	1.6	1.8	

1.	Low,	2.	Medium,	3.	Strong
----	------	----	---------	----	--------

Cours	Course Outcome VS Programme Specific Outcomes											
CO	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5							
CO-1	S(3)	M(2)	S(3)	S(3)	M(2)							
CO-2	S(3)	M(2)	S(3)	S(3)	L(1)							
CO-3	S(3)	M(2)	S(3)	S(3)	M(2)							
CO-4	S(3)	L(1)	S(3)	S(3)	L(1)							
CO-5	S(3)	S(3)	S(3)	S(3)	L(1)							
W.AV	3	2	3	3	1.4							



		Semester-III							
Core	Course code:	Practical VI- Techniques in Biomedical	Р	Credits:4	Hours:6				
	508304	sciences-III							
Course		and the techniques in isolation, identification, e	extrac	tion, structura	al and				
Objectives	function	nal elucidation, of medicinal compounds.							
	Compre	when the principles and applications of microsc	opes	ia taabaiguag					
	\rightarrow Know-f	now the principles and applications of chromato the principles and applications of electrophore	graph	ne techniques					
	\succ Have a	n understanding the methods of electrochemical	meth	ods of analys	is				
	• Extraction of bioactive compounds (hot and cold method)								
	 Lyophi 								
		al of solvent by rotary flash evaporator							
	Demon	stration of							
		ing Electron Microscope (SEM)							
		nission Electron Microscope(TEM)							
		cal Microscope							
		preparation and histopathological examination							
		processing and sectioning							
		ctrophoresis							
		fingerprinting							
		solation and quantification							
	• PCR, R	APD, RELP and DNA sequencing							
Suggested	Readings.	S ALAGAPPA UNIVERSITY 8							
Rich Dav cond Will Irfar Den	hard F. Venn, (id L. spector & cepts from c lard, H., Dean, h A. Khan, Atiy nis W. Ross, (2	x, M.M. W.H. (2006). Lehninger's Biochemistry 2004). Principles and practice of bioanalysis (ed. & Robert D. Goldman. (2006). Basic methods is ells: A laboratory manual (ed.,) S, Instrumental Methods of Analysis (1986) va Khanm (1998). Role of Biotech Medicinal an 2002). Introduction to molecular medicine, 3rd (e b). Herbal Drugs and Biotechnology (ed.,)	.) in mi <mark>d Ar</mark> c	croscopy: Pro					
), Herour Drugs and Drottermorogy (ear,)							
	Resources:	einstrumentation Dischargistru Missehisters (7 ₋₁₁ F						
Technolog		oinstrumentation, Biochemistry, Microbiology, C		biology and El	Izyme				
		net/publication/329390135_Laboratory_Manual	for	Rioinstrument	tation B				
		y_Cell_Biology_and_Enzyme_Technology_For_							
Laboratory					procedure,				
		ege.edu.in/download/downloads/1803201242057		Ų	1 ,				
•		-	•						
Course		erstand the extraction, isolation and identification	on of	bioactive mol	ecules				
Outcomes (CO)									
	CO2- Learn the various microscopic techniques involved in the drug discovery and								
	developme	· ·		ε.					
	-	uire the separation of phytochemicals by using i	noten	monte					
	-				. • •				
		w the separation of macro molecules for the de-	-		e peptides				
	CO5- Viev	v the toxicity of drugs through histopathological	studi	es					

	Course Outcome VS Programme Outcomes											
CO	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10		
CO-1	S(3)	S(3)	S(3)	S(3)	M(2)	S(3)	M(2)	M(2)	M(2)	M(2)		
CO-2	S(3)	S(3)	M(2)	S(3)	M(2)	S(3)	M(2)	L(1)	L(1)	M(2)		
CO-3	S(3)	L(1)	L(1)	S(3)	S(3)	S(3)	L(1)	M(2)	M(2)	L(1)		
CO-4	S(3)	L(1)	M(2)	S(3)	M(2)	S(3)	L(1)	L(1)	L(1)	L(1)		
CO-5	S(3)	M(2)	L(1)	S(3)	L(1)	S(3)	S(3)	L(1)	M(2)	S(3)		
W.AV	3	2	1.8	3	2	3	1.8	1.4	1.6	1.8		

Course	Course Outcome VS Programme Specific Outcomes											
СО	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5							
CO-1	S(3)	S(3)	S(3)	S(3)	M(2)							
CO-2	S(3)	S(3)	M(2)	S(3)	M(2)							
CO-3	S(3)	L(1)	L(1)	S(3)	S(3)							
CO-4	S(3)	L(1)	M(2)	S(3)	M(2)							
CO-5	S(3)	M(2)	L(1)	S(3)	L(1)							
W.AV	3	2	1.8	3	2							



		Semester-III			
DSE 3	Course code: 508505	Biomaterials and Tissue Engineering	P	Credits 3	Hours 3
		Unit-I			
		the concepts of biomaterial implants and ti			
	• I ·	ges and disadvantages., Bio ceramics for imperson set and other biomedical alloys, impla		0	
Outcome	1 Acquire kno engineering.	wledge on the biomaterials, implant and tiss	sue		K2
	8 8	Unit-II			
Objective	2 Learn appli	cations of nanomaterials in various body pa	arts im	plant bioma	aterials
-		ials use as implants, biological response of i		=	
		of the body with implanted materials., Materia			
	bioceramics, mo				
Outcome	2 Understand	the desirable and undesirable reactions of	the b	ody with	K4
	implanted n	naterials.			
		Unit III			
°		the advantages and disadvantages of implar			
and vascu		modes of dental implant failure, wear debris des of cartilage implant, vascular implant, im			
		t tissue engineering and bioactive scaffold.			K4
Objective	4 Gain kno	wledge on the protein interactions with imp	planted	d materials	
•		implanted materials, cellular recognition of			on material
		ion, differentiation, cellular extra cellular mati	•		
	•	n-body response, inflammatory response	1		C
Outcome		he protein interaction with the implanted n	nateria	ls	K2
		Unit V		1	
Objective	-	ne applications of natural and degrad	lable	polymers	for tissue
Ti	engineering			mus dissus le	
cellular si	gnaling, extrace	cells, morphogenesis, generation of tissue in the llular matrix as a biologic scaffold for	tissue	engineerin	g, scaffold
		old, natural polymers in tissue engineering ap	pplicati	ons, degrad	able
* *	for tissue enginee				7 =
Outcome		the role of applications of polymers in tissu	e engli	heering H	K5
	Readings:	tterials science and tissue engineering: princip	loc and	mathada (a	(be
		tificial Organs. (ed.,)	ies and	methous. (e	<i>)</i>
		M. R., & Mani, G. (2014). Introduction to b	oiomate	rials: basic t	theory with
	ering applications		10111400		
Poole,	D. L., & Mackwo	orth, A. K. (2010). Artificial Intelligence: found	dations	of computat	tional
agents.		-		-	
Donglu		omaterial and Tissue engineering,(ed.,)			
	•	er et al., (2004). Biomaterial Science: An Intro	oduction	n to material	ls in
Online -	Medicine., (ed.,))			
Online res Biomatori		Engineering- https://link.springer.com/book/1	10 100-	1/078 2 667	06104 6
		Biomaterials for Tissue Engineering-	10.1007	//9/8-3-002-	00104-0
		pook/9781681085364/			
		n Tissue Engineering- <u>https://www.wiley.cor</u>	m/en-		
		+in+Tissue+Engineering-p-9781118371213			
		stand, K3- Apply K4- Analyze, K5-Evaluat	te, K6-	Create	

	Course Outcome VS Programme Outcomes										
СО	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10	
CO-1	M(2)	S(3)	S(3)	M(2)							
CO-2	L(1)	M(2)	S(3)	M(2)	M(2)	L(1)	L(1)	L(1)	M(2)	M(2)	
CO-3	M(2)	L(1)	S(3)	S(3)	L(1)	M(2)	M(2)	M(2)	S(3)	L(1)	
CO-4	L(1)	L(1)	S(3)	M(2)	L(1)	L(1)	L(1)	M(2)	M(2)	L(1)	
CO-5	M(2)	M(2)	S(3)	L(1)	S(3)	L(1)	M(2)	S(3)	L(1)	S(3)	
W.AV	1.6	1.8	3	2	1.8	1.4	1.6	2	2	1.8	

Course	Course Outcome VS Programme Specific Outcomes											
CO	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5							
CO-1	M(2)	S(3)	S(3)	M(2)	S(3)							
CO-2	L(1)	M(2)	S(3)	M(2)	S(3)							
CO-3	M(2)	L(1)	S(3)	S(3)	S(3)							
CO-4	L(1)	L(1)	S(3)	M(2)	S(3)							
CO-5	M(2)	M(2)	S(3)	L(1)	S(3)							
W.AV	1.6	1.8	3	2	3							



		Semester-III			
DSE 3 C	ourse code: 508506	Bio-Imaging Technology	Τ	Credits 3	Hours 3
		Unit-I			
U		asic principles and applications of micros			
fluorescens m	icroscope, sca	e, principles and applications of optical m nning electron microscope, transmission el		•	· ·
dead assay wi					
Outcome 1	Learn abou	t the principles and applications of micr	oscop	y	K2
	~	Unit-II			
-		dge on the ultrasound imaging systems			
principles of	A, B M Mod	sics of ultrasound- principles of image e, scan converters- Doppler ultrasound- pul	sed ar	nd continuous.	
Outcome 2	Understand imaging tech		he qu	ality of	K4
		Unit III			
•		the principles and production of X-rays			
		of X-rays-soft and hard, radiographic an nage intensifier systems, computed and dig			
Outcome 3	Acquired k	nowledge about the principles of image	form	ation,	K4
	-	display of ultrasound and X-ray.			
		Unit IV			
Objective 4	To acqui	red knowledg <mark>e</mark> of imaging system theory	and t	their applicat	ions
		omography, mammography, transverse ton			
		cal applications, CT Angiography basic p			
		nner princi <mark>ple</mark> s, SPECT, Computer <mark>te</mark> chniq			on.
Outcome 4	Understand NMR spectr		graph	y, MRI and	K2
	•	Unit V	ľ		
		applications of image acquisition in mag			
spin-echo tech	inique and spir	etic resonance imaging MRI-T1 MRI-T2 n relaxation technique- various types of pu R spectroscopy.			
	Clear in the	role of applications various types of pu on of imaging	ilse se	equences for	K5
instrument Drexler, V application Hendee, V Khandpur Ong, J. L engineerin Po cor Online resou Bio-Imaging Principles-Teo Biomedical In	, L., Weibell, tation and mea V., & Fujimoto Is. (ed.,) V. R., & Riter , R. S. (1987) , Appleford, g applications ole, D. L., & <u>mputational ag</u> rces Principles, T chniques-and- naging: Princi	Mackworth, A. K. (2010). Artificial Intellig	mogra s. (ed. (ed.,) bion gence: w.rout	aphy: technolo ,) naterials: basic foundations c tledge.com/Bic	e theory with
K1-Remembe	er, K2-Under	stand, K3- Apply K4- Analyze, K5-Eval	uate,	K6- Create	

	Course Outcome VS Programme Outcomes										
CO	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10	
CO-1	S(3)	M(2)	S(3)	S(3)							
CO-2	M(2)	L(1)	L(1)	M(2)	M(2)	L(1)	L(1)	M(2)	M(2)	S(3)	
CO-3	S(3)	S(3)	S(3)	S(3)	L(1)	M(2)	M(2)	L(1)	L(1)	S(3)	
CO-4	M(2)	L(1)	S(3)	M(2)	L(1)	L(1)	L(1)	M(2)	S(3)	S(3)	
CO-5	S(3)	S(3)	S(3)	L(1)	S(3)	L(1)	M(2)	L(1)	M(2)	S(3)	
W.AV	2.6	2	2.4	2	1.8	1.4	1.6	1.6	2.2	3	

С	Course Outcome VS Programme Outcomes										
CO	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5						
CO-1	S(3)	S(3)	M(2)	M(2)	M(2)						
CO-2	M(2)	S(3)	L(1)	M(2)	M(2)						
CO-3	S(3)	S(3)	S(3)	S(3)	L(1)						
CO-4	M(2)	S(3)	S(3)	M(2)	L(1)						
CO-5	S(3)	S(3)	S(3)	L(1)	S(3)						
W.AV	2.6	3	2.4	2	1.8						
	11	ow 2 Medi	im 3 Strot	nα							



	Semester-IV								
Core	Course code: 508999	Dissertation Work	Credits 15	Hours20					



Semester-IV									
Core	Course code: 508777	Hospital Training	Credits 5	Hours 10					



Courses offer to other Departments

		Semester-II	L		
NME	Course code:	Hospital Management an	d Biosafety T	Credit	s 2 Hours 3
	course coue.	Unit-I	a biosarcey 1	Cituli	
Objective	Understand t	theories of management			
Ū.		volution of management,		nortance of	f management
		t thought- overall suppor			
	enance and comp				
Outcome 1		importance of manageme	ent and different l	bodies of	K2
	management	,			
		Unit-II			
Objective 2	Know Mana hospital admi	hospitals by understan istrator	ding the comple	exity, levels	s and role of
Epidemiolo	gical basis for he	hcare management, Mana	gement developm	ent-towards	development
-	-	f Indian hospitals, Manag		-	-
•		of hospital management, Op	eration concept- u	se of model	s, Health
		managerial methods			1
Outcome 2	2 Acquire know management	edge on the epidemiolog	cal basis for hea	lthcare	K4
		Unit III	N 40		·
		anagement process and in			
		nciples in planning hospit			
		nent survey of community, anagement & implementati			
engineering	, lighting etc.	DIAX			_
Outcome	3 Recognise the	organization of the hospit	al and functionar	ies	K4
	T (1	Unit IV	1	• •	
Objective 4		ent issues that have an im			
committee	and hospital fu	management structure, ty tionaries, duties and res gement of quality assured	ponsibilities of v	arious pos	itions hospital
	4 Know the hore requirements	pital infrastructure and f	actors determinin	ng legal	K2
	requirements	Unit V			
Objective 5	Recognise the	biosafety regulatory frame	work and its soc	io economi	c impact
U U	9	ospital waste management,			-
·	thics and its socio				
Outcome 5	5 Learn op	rtunities in the hospital v	vaste managemen	t	K5
Suggested F		ł	8		
00	0	987). Management of Hosp	oital (4 Vols), (ed	.,).	
2. M.K.	Satheesh (1992).	Bioethics and Biosafety (ed	,)		
3. Jame	s A. William, Mc	illan (1991). Hospital Man	agement. (ed.,)		
•	Amin Tabish (20 ice (ed.,)). Hospital and Health Serv	ices Administratio	n – Princij	oals and
5. Nelso	on Thrones (2006)	Management in Health Care	e, 2nd (ed.,)		
		018).Hospital Management	and Administration	on. (ed.,)	
in/Biological Principles o	afety: Principles +Safety%3A+Pri f Hospital Admir	nd Practices, 5th Edition- iples+and+Practices%2C+5 tration And Management	oth+Edition-p-978	1683673132	
		ok/principles-of-hospital-a		-	<u>t</u>
K1-Remem	ber, K2-Understa	d, K3- Apply K4- Analyz	e, K5-Evaluate, H	K6- Create	

	Course Outcome VS Programme Outcomes										
СО	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10	
CO-1	M(2)	S(3)	S(3)	M(2)	S(3)	M(2)	M(2)	M(2)	M(2)	S(3)	
CO-2	L(1)	L(1)	M(2)	M(2)	S(3)	M(2)	L(1)	L(1)	M(2)	M(2)	
CO-3	M(2)	S(3)	S(3)	L(1)	S(3)	L(1)	M(2)	M(2)	L(1)	L(1)	
CO-4	M(2)	L(1)	M(2)	M(2)	S(3)	L(1)	L(1)	L(1)	M(2)	S(3)	
CO-5	M(2)	L(1)	L(1)	L(1)	S(3)	S(3)	L(1)	M(2)	L(1)	M(2)	
W.AV	1.8	1.8	2.2	1.6	3	1.8	1.4	1.6	1.6	2.2	

1. Low, 2. Medium, 3. Strong	1.	Low	, 2.	Medium,	3.	Strong
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Course Outcome VS Programme Specific Outcomes									
СО	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5				
CO-1	M(2)	S(3)	S(3)	M(2)	S(3)				
CO-2	L(1)	L(1)	M(2)	M(2)	S(3)				
CO-3	M(2)	S(3)	S(3)	L(1)	S(3)				
CO-4	M(2)	L(1)	M(2)	M(2)	S(3)				
CO-5	M(2)	L(1)	L(1)	L(1)	S(3)				
W.AV	1.8	1.8	2.2	1.6	3				
	2 1	Low 2 Me	dium 3 St	rong					



			Seme	ster-III							
Cou	se code:	Molec		ced Diagnostics	Т	Credits 2	Hours 3				
			U	nit-I			I				
ve 1	Understand	the types	of diseases a	nd diagnosis							
	· ·	•		0							
						·	• • • •				
							clinical				
			of diseases of	lue to metabolic dis	orders	s and	K2				
Unit-II											
e 2	Know vario	us method	s of cytogen	etic analysis							
etics 1	Karyotype a	nalysis, blo	od , bone r	narrow, amniotic fl	uid, cl	norionic vil	lus samples,				
of co	nception Flu	uorescent in	n situ hybrid	lization, Cytogenetic	studi	es using mi	croarrays or				
beads.	Molecular I	DNA isolati	ion and quar	ntification, Probe and	1 prim	er designing	g, PCR -				
			·		.	U .	-				
e 2	Acquire kno	owledge on	the cytogen	etic studies			K4				
Unit III											
e 3	Acquire the	knowledge	e on health i	informatics							
ction	to pharmacy	informatio	cs, Medical	Transcription, Role	of in	formatics to	enhance the				
		utomation,	Informatics	applications in phar	macy,	survey and	evaluation of				
		• •		8.010			TZ A				
2.3	Recognise ti	ne informa					K4				
	-				0.1						
						<u> </u>					
			-			-	-				
		-			•						
		•	• ·	•		·					
-	-			-			pFLP, STR,				
			Multiplex PCR- Determination of Paternity- Human identification and sex determination. Southern								
-	blotting, isotopic and nonisotopic methods, Western blotting, DNA Sequencing, including massively										
parallel sequencing and microarrays.											
-		croarrays.	hods, Wester	n blotting, DNA Sec	quencii	ng, includin	g massively				
equen e 4		croarrays.	hods, Wester	n blotting, DNA Sec	quencii	ng, includin					
e 4	Learn the	croarrays. e various n	hods, Wester nethods ider Ur	n blotting, DNA Sec ntification of microl nit V	quencii	ng, includin	g massively				
e 4	Learn the	e various n	hods, Wester nethods ider Ur hniques in d	n blotting, DNA Sec ntification of micro nit V lisease diagnostics	quencii bial pa	ng, includin Ithogens	g massively K2				
e 4 e 5 w of	Learn the i	e various n mmunotec tem , Anti	hods, Wester nethods ider Ur hniques in o gens and ar	n blotting, DNA Sec ntification of micro nit V lisease diagnostics ntibodies , Antigen-	quencin bial pa	ng, includin thogens	g massively K2 ions, Major				
e 4 e 5 w of	Learn the i Learn the i immune sys bility Comp	e various n mmunotec tem , Anti lex (MHC	hods, Wester nethods ider Ur hniques in o gens and ar), HLA typ	n blotting, DNA Sec ntification of microl nit V lisease diagnostics ntibodies , Antigen- ping , Immunothera	quencin bial pa -antibo apy ar	ng, includin thogens ody interact nd immuno	g massively K2 ions, Major diagnostics.				
e 4 e 5 w of npatib	Learn the i Learn the i immune sys bility Comp ostics - Intro	e various n mmunotec tem , Anti lex (MHC oduction, a	hods, Wester nethods ider Ur hniques in o gens and ar), HLA typ ntigen- antib	n blotting, DNA Sec ntification of microl nit V lisease diagnostics ntibodies , Antigen- ning , Immunothera body binding interact	quencin bial pa -antibo apy an ctions	ng, includin thogens ody interact od immuno and assays;	g massively K2 ions, Major diagnostics. antibodies-				
e 4 e 5 w of npatib diagn al an	Learn the i immune sys bility Comp ostics - Intro d monoclon	e various n mmunotec tem , Anti lex (MHC oduction, a aal antibodi	hods, Wester nethods ider Ur hniques in d gens and ar), HLA typ ntigen- antib es, Immunoa	n blotting, DNA Sec ntification of micro nit V lisease diagnostics ntibodies , Antigen- bing , Immunothera body binding interaction ussays — types [RI	duencin bial pa -antibo apy ar ctions A, EL	ng, includin thogens ody interact ad immuno and assays; ISA, Chem	g massively K2 ions, Major diagnostics. antibodies- iluminescent				
e 4 e 5 w of npatib diagn al an] and	Learn the i immune sys bility Composition - Intro d monoclon specific ap	mmunotec immunotec tem , Anti lex (MHC oduction, a al antibodi plications;	hods, Wester nethods ider Ur hniques in o gens and ar), HLA typ ntigen- antib es, Immunoa Immunohis	n blotting, DNA Sec ntification of micro nit V lisease diagnostics ntibodies , Antigen- bing , Immunothera body binding interact assays — types [RI tochemistry — prin	duencin bial pa -antibo apy ar ctions A, EL	ng, includin thogens ody interact ad immuno and assays; ISA, Chem	g massively K2 ions, Major diagnostics. antibodies- iluminescent				
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Molecular DNA isolation and quara and various modifications, Real time PCR e 2 Acquire knowledge on the cytogene Un e 3 Acquire the knowledge on health is provided by pharmaceutical care gives ment, Medical Coding Systems Telemed cy systems and automation, Informatics resources. e 3 Recognise the informatics applicated and the informatics of human bacterial patial DNA Restriction analysis (ARDRA)-C the of human bacterial patial DNA Restriction analysis of fungal pathonologens through PCR. RAPD for animal and the informatical and pathonologens through PCR. 	 mode of transmission in infections, factors predisposing t ous diseases- bacterial, viral, fungal, protozoans and other para collection, transport and processing of samples, interpretation Inderstand the causes of diseases due to metabolic dis infectious pathogens Unit-II Example 1 Understand the causes of diseases due to metabolic dis infectious pathogens Unit-II Example 2 Know various methods of cytogenetic analysis etics Karyotype analysis, blood , bone marrow, amniotic fl of conception Fluorescent in situ hybridization, Cytogenetic beads. Molecular DNA isolation and quantification, Probe and and various modifications, Real time PCR, MLPA analysis, S 2 Acquire knowledge on the cytogenetic studies Unit III e 3 Acquire the knowledge on health informatics ction to pharmacy informatics, Medical Transcription, Role provided by pharmaceutical care givers. Health Information and automation, Informatics applications in phar resources. a Recognise the informatics application in pharmacy Unit IV e 4 Describe various molecular methods of identification analysis (ARDRA)-Culture independent TLP. Molecular diagnosis of fungal pathogens based on 18St mogens through PCR. RAPD for animal and plants. PCR in formation of the process of the state of the state	 infectious, physiological and metabolic errors, genetic basis of di mode of transmission in infections, factors predisposing to microus diseases- bacterial, viral, fungal, protozoans and other parasites. In collection, transport and processing of samples, interpretation of restination of the processing of samples, interpretation of restinations pathogens Understand the causes of diseases due to metabolic disorders infectious pathogens Unit-II e 2 Know various methods of cytogenetic analysis etics Karyotype analysis, blood , bone marrow, amniotic fluid, cloof conception Fluorescent in situ hybridization, Cytogenetic studies and various modifications, Real time PCR, MLPA analysis, SNP, SS e 2 Acquire knowledge on the cytogenetic studies Unit III e 3 Acquire the knowledge on health informatics ction to pharmacy informatics, Medical Transcription, Role of in provided by pharmaceutical care givers. Health Information Ament, Medical Coding Systems Telemedicine and Telehealth, Ethicy systems and automation, Informatics applications in pharmacy, resources. a Recognise the informatics application in pharmacy unit IV e 4 Describe various molecular methods of identification of human bacterial pathogens- PCR, 16S rRNA al DNA Restriction analysis (ARDRA)-Culture independent analy LP. Molecular diagnosis of fungal pathogens based on 18SrRNA inogens through PCR. RAPD for animal and plants. PCR in forensics 	 - infectious, physiological and metabolic errors, genetic basis of diseases, inhe - mode of transmission in infections, factors predisposing to microbial patho ous diseases- bacterial, viral, fungal, protozoans and other parasites. Methods of a collection, transport and processing of samples, interpretation of results. ne 1 Understand the causes of diseases due to metabolic disorders and infectious pathogens Unit-II e 2 Know various methods of cytogenetic analysis etics Karyotype analysis, blood , bone marrow, amniotic fluid, chorionic vill of conception Fluorescent in situ hybridization, Cytogenetic studies using mi beads. Molecular DNA isolation and quantification, Probe and primer designing and various modifications, Real time PCR, MLPA analysis, SNP, SSCP, e 2 Acquire knowledge on the cytogenetic studies Unit III e 3 Acquire the knowledge on health informatics ction to pharmace utical care givers. Health Information Architecture, ment, Medical Coding Systems Telemedicine and Telehealth, Ethics in medic cy systems and automation, Informatics applications in pharmacy 				

Suggested Readings:

Bailey & Scott's Diagnostic Microbiology (2002), Betty A. Forbes, Daniel F. Sahm, Alice S.Weissfeld, Ernest A. Trevino, Published by C.V. Mosby

Jawetz, Melnick, & Adelberg's Medical Microbiology (2004), Geo F. Brooks, Stephen Morse, Janet S. Butel.

- Fundamentals of Molecular Diagnostics (2007). David E. Bruns, Edward R. Ashwood, Carl A.Burtis. Saunders Group.
- Molecular Diagnostics: Fundamentals, Methods & Clinical applications (2007). Lele Buckingham and Maribeth L. Flaws.
- Fundamentals of Molecular Diagnostics (2007). David E. Bruns, Edward R. Ashwood, Carl A.Burtis. Saunders Group.

Expert Review of Proteomics and Molecular Diagnostics (Journals) Basic Concepts of Molecular Pathology Series: Molecular Pathology Library, Vol. 2; Cagle, Philip T. Allen, Timothy C. (Eds.);Springer 2009

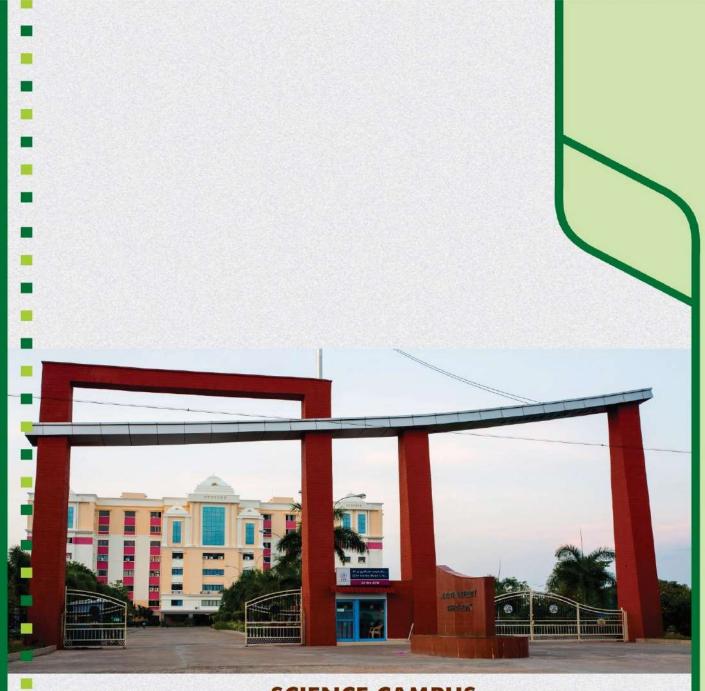
Online resources

Molecular Diagnostics- https://link.springer.com/book/10.1385/1592599281 Principles and Applications of Molecular Diagnostics- https://shop.elsevier.com/books/principlesand-applications-of-molecular-diagnostics/rifai/978-0-12-816061-9 K1-Remember, K2-Understand, K3- Apply K4- Analyze, K5-Evaluate, K6- Create

	Course Outcome VS Programme Outcomes											
CO	PO-1	PO-2	PO-3	PO-4	PO-5	PO-6	PO-7	PO-8	PO-9	PO-10		
CO-1	S(3)	S(3)	S(3)	M (2)	M(2)	M(2)	M(2)	S(3)	M(2)	S(3)		
CO-2	M(2)	S(3)	S(3)	M(2)	M(2)	M(2)	L(1)	S(3)	M(2)	M(2)		
CO-3	L(1)	M(2)	M(2)	L(1)	S(3)	L(1)	M(2)	S(3)	L(1)	L(1)		
CO-4	M(2)	S(3)	S(3)	M(2)	M(2)	L(1)	L(1)	S(3)	M(2)	S(3)		
CO-5	L(1)	M(2)	S(3)	L(1)	L(1)	<mark>S(</mark> 3)	L(1)	S (3)	L(1)	M(2)		
W.AV	1.8	2.6	2.8	1.6	2	1.8	1.4	3	1.6	2.2		

1. Low, 2. Medium, 3. Strong

Course Outcome VS Programme Specific Outcomes										
CO	PSO-1	PSO-2	PSO-3	PSO-4	PSO-5					
CO-1	S(3)	S(3)	S(3)	S(3)	M(2)					
CO-2	M(2)	S(3)	S(3)	S(3)	M(2)					
CO-3	L(1)	M(2)	M(2)	S(3)	S(3)					
CO-4	M(2)	S(3)	S(3)	S(3)	M(2)					
CO-5	L(1)	M(2)	S(3)	S(3)	L(1)					
W.AV	1.8	2.6	2.8	3	2					



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