FACULTY OF PHARMACY KARPAGAM ACADEMY OF HIGHER EDUCATION

Deemed to be University (*Established Under Section 3 of UGC Act 1956*) Eachanari Post, Pollachi Main Road, Coimbatore – 641021.

PHARM.D DEGREE COURSE (2024-2025) (DOCTOR OF PHARMACY)

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REGULATIONS 2024

COURSE OF STUDY AND SCHEME OF EXAMINATION

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SYLLABUS

CHAPTER-I

1. Short title and commencement

- (1) These regulations may be called the Pharm.D Regulations 2024.
- (2) They shall come into force from the date of their publication in the official Gazette.
- 2. Pharm.D shall consist of a certificate, having passed the course of study and examination as prescribed in these regulations, for the purpose of registration as a pharmacist to practice the profession under the Pharmacy Act, 1948.

CHAPTER-II

3. Duration of the course :

 a) Pharm.D : The duration of the course shall be six academic years (five years of study and one year of internship or residency) full time with each academic year spread over a period of not less than two hundred working days. The period of six years duration is divided into two phases –

Phase I – consisting of First, Second, Third, Fourth and Fifth academic year. Phase II – consisting of internship or residency training during sixth year involving posting in specialty units. It is a phase of training wherein a student is exposed to actual pharmacy practice or clinical pharmacy services and acquires skill under supervision so that he or she may become capable of functioning independently.

b) Pharm.D (Post Baccalaureate) : The duration of the course shall be for three academic years (two years of study and one year internship or residency) full time with each academic year spread over a period of not less than two hundred working days. The period of three years duration is divided into two phases:

Phase I — consisting of First and Second academic year.

Phase II – consisting of Internship or residency training during third year involving posting in specialty units. It is a phase of training wherein a student is exposed to actual pharmacy practice or clinical pharmacy services, and acquires skill under supervision so that he or she may become capable of functioning independently.

4. Minimum qualification for admission to:

- a) Pharm.D Part-I Course A pass in any of the following examinations :-
- 10+2 examination with Physics and Chemistry as compulsory subjects along withone of the following subjects: Mathematics or Biology.
- (2) A pass in D. Pharm course from an institution approved by the Pharmacy Councilof India under section 12 of the Pharmacy Act.
- (3) Any other qualification approved by the Pharmacy Council of India as equivalent to any of the above examinations.

Provided that a student should complete the age of 17 years on or before 31st December of the year of admission to the course.

Provided that there shall be reservation of seats for the students belonging to the Scheduled Castes, Scheduled Tribes and other Backward Classes in accordance with the instructions issued by the Central Government/State Government/Union Territory Administration as the case may be from time to time.

b) Pharm.D. (Post Baccalaureate) Course -

A pass in B.Pharm from an institution approved by the Pharmacy Council of India under section 12 of the Pharmacy Act:

Provided that there shall be reservation of seats for the students belonging to the Scheduled Castes, Scheduled Tribes and other Backward Classes in accordance with the instructions issued by the Central Government/State Government/Union Territory Administration as the case may be from time to time.

- **5.** Number of admissions in the above said programmes shall be as prescribed by the Pharmacy Council of India from time to time and presently be restricted as below :
 - i) Pharm.D. Programme 30 students
 - ii) Pharm.D. (Post Baccalaureate) Programme 10 students
- 6. Institutions running B.Pharm programme approved under section 12 of the Pharmacy Act, will only be permitted to run Pharm.D programme. Pharm.D (Post Baccalaureate) programme will be permitted only in those institutions which are permitted to run Pharm.D. Programme.

7. Attendance and progress:

A candidate is required to put in at least 80% attendance in individual courses considering Theory and Practical separately. The candidate shall complete the prescribed course satisfactorily to be eligible to appear for the respective examinations.

8. Program / Course credit structure:

As per the philosophy of Credit Based Semester System, certain quantum of academic work viz. Theory classes, tutorial hours, Practical classes, etc., are measured in terms of credits. On satisfactory completion of the courses, a candidate earns credits. The amount of credit associated with a course is dependent upon the number of hours of instruction per week in that course. Similarly, the credit associated with any of the other academic, co/extra-curricular activities is dependent upon the quantum of work expected to be put in for each of these activities per week.

Credit assignment:

Theory and Laboratory courses:

Courses are broadly classified as Theory and Practical. Theory courses consist of lecture (L) and/or tutorial (T) hours, and Practical (P) courses consist of hours spent in the laboratory. Credits (C) for a course is dependent on the number of hours of instruction per week in that course, and is obtained by using a multiplier of one (1) for lecture and tutorial hours, and a multiplier of half (1/2) for Practical (laboratory) hours. Thus, for example, a Theory course having three lectures and one tutorial per week throughout the semester carries a credit of 4. Similarly, a Practical having four laboratory hours per week throughout semester carries a credit of 2.

Minimum credit requirements:

The minimum credit points required for award of a Pharm.D degree is 160. These credits are divided into Theory courses, Tutorials, Practical, Clerkship and Project over the duration of six years. The credits are distributed year-wise as shown in Table VI. Courses generally progress in sequences, building competencies and their positioning indicates certain academic maturity on the part of the learners. Learners are expected to follow the year-wise schedule of courses given in the syllabus.

9. Academic work

A regular record of attendance both in Theory and Practical shall be maintained by the teaching staff of respective courses.

10. Course of study:

The course of study for Pharm.D shall include the subjects as given in the Tables below.

The number of hours in a week, devoted to each subject for its teaching in Theory,

Practical and tutorial shall not be less than that noted against it in columns below.

TABLES TABLE I : FIRST YEAR

Course Code	Name of Subject	No. of hours	Tutorial	Creditpoints
24PD101T	Human Anatomy and Physiology- Theory	3	1	4
24PD102T	Pharmaceutics - Theory	2	1	3
24PD103T	Medicinal Biochemistry - Theory	3	1	4
24PD104T	Pharmaceutical Organic Chemistry - Theory	3	1	4
24PD105T	Pharmaceutical Inorganic Chemistry - Theory	2	1	3
24PD106RMT /	Remedial Mathematics*/	2	1	Λ
24PD106RBT	RemedialBiology -Theory*	5	1	4
24PD107P	Human Anatomy and Physiology -Practical	3	-	2
24PD108P	Pharmaceutics -Practical	3	-	2
24PD109P	Medicinal Biochemistry -Practical	3	-	2
24PD110P	Pharmaceutical Organic Chemistry -Practical	3	-	2
24PD111P	Pharmaceutical Inorganic Chemistry- Practical	3	-	2
24PD112RBP	Remedial Biology -Practical**	3	-	2
24PD113ET /	Communication Skills -Theory***/	2	-	1
24PD114ET	Yoga for Youth Empowerment -Theory***			
	Total Hours	30/33*/ 36**	6	29/33*/ 35**

* For Remedial Mathematics / Remedial Biology (Non University Exam)

** For Remedial Biology Practical

*** Two Electives offered, students can choose any one.

TABLE II : SECOND YEAR

Course Code	Name of Subject	No. of hours	Tutorial	Creditpoints
24PD201T	Pathophysiology - Theory	3	1	4
24PD202T	Pharmaceutical Microbiology -Theory	3	1	4
24PD203T	Pharmacognosy and Phytopharmaceuticals-Theory	3	1	4
24PD204T	Pharmacology I-Theory	3	1	4
24PD205T	Community Pharmacy - Theory	2	1	3
24PD206T	Pharmacotherapeutics I- Theory	3	1	4
24PD207P	Pharmaceutical Microbiology - Practical	3	-	2
24PD208P	Pharmacognosy and Phytopharmaceuticals- Practical	3	-	2
24PD209P	Pharmacotherapeutics I- Practical	3	-	2
24PD210ET/	Health and Lifestyle -Theory*/	2	-	1
24PD211ET	NPTEL-1*			
	Total Hours	28	6	30

* Two Electives offered, students can choose any one.

TABLE III : THIRD YEAR

Course Code	Name of Subject	No. of hours	Tutorial	Credit points
24PD301T	Pharmacology II - Theory	3	1	4
24PD302T	Pharmaceutical analysis- Theory	3	1	4
24PD303T	Pharmacotherapeutics II - Theory	3	1	4
24PD304T	Pharmaceutical jurisprudence- Theory	2	-	2
24PD305T	Medicinal chemistry -Theory	3	1	4
24PD306T	Pharmaceutical formulations - Theory	2	1	3
24PD307P	Pharmacology II - Practical	3	-	2
24PD308P	Pharmaceutical analysis -Practical	3	-	2
24PD309P	Pharmacotherapeutics II- Practical	3	-	2
24PD310P	Medicinal chemistry- Practical	3	-	2
24PD311P	Pharmaceutical formulations -Practical	3	-	2
24PD312ET/	Indian Indigenous Medicine - Theory */	2	-	2
24PD313ET	NPTEL-2*			
	Total hours	33	5	33

*Two Electives offered, students can choose any one.

TABLE IV : FOURTH YEAR

Course Code	Name of Subject	No. of hours	Tutorial	Credit points
24PD401T	Pharmacotherapeutics III- Theory	3	1	4
24PD402T	Hospital Pharmacy- Theory	2	1	3
24PD403T	Clinical Pharmacy- Theory	3	1	4
24PD404T	Biostatistics and Research Methodology-Theory	2	1	3
24PD405T	Biopharmaceutics and Pharmacokinetics - Theory	3	1	4
24PD406T	Clinical Toxicology -Theory	2	1	3
24PD407P	Pharmacotherapeutics III- Practical	3	-	2
24PD408P	Hospital Pharmacy- Practical	3	-	2
24PD409P	Clinical Pharmacy -Practical	3	-	2
24PD410P	Biopharmaceutics and Pharmacokinetics- Practical	3	-	2
24PD411EP/	Statistical Software -Practical*/	3/1	-	2
24PD412ET	Ethical Leadership- Theory*			
		30/28*	6	31

*Two Electives offered, students can choose any one.

TABLE V : FIFTH YEAR

Course	Name of Subject	No. of	Seminar	Credit
Code		hours		points
24PD501T	Clinical Research - Theory	3	1	4
24PD502T	Pharmacoepidemiology and	3	1	4
	Pharmacoeconomics - Theory			
24PD503T	Clinical Pharmacokinetics &	2	1	3
	Pharmacotherapeutics Drug			
	Monitoring- Theory			
24PD504S	Clerkship *	-	1	1
24PD505P	Project work** (Six Months)	20	-	10
24PD506ET /	Medical Coding - Theory ***/	1/3	-	2
24PD507EP	Pharmaceutical Calculation - Practical***			
		29/31	4	24

* Attending ward rounds on daily basis ** **Two Electives offered, students can choose any one.**

SIXTH YEAR

Internship or residency training including postings in speciality units. Student

should independently provide the clinical pharmacy services to the allotted wards.

- (i) Six months in General Medicine department, and
- (ii) Two months each in three other speciality departments

TABLE VI: YEAR WISE CREDITS DISTRIBUTION

Year	Credit Points
Ι	29*/33**/35***
Π	30
III	33
IV	31
V	24
VI	15
Total credit points for the program	162*/166**/168***

11. Syllabus:

The syllabus for each subject of study in the said tables shall be as specified in **Chapter-IV** to these regulations.

12. Approval of the authority conducting the course of study:

- No person, institution, society or university shall start and conduct Pharm.D or Pharm.D
 (Post Baccalaureate) programme without the prior approval of the Pharmacy Council of India.
 - a. Any person or pharmacy college for the purpose of obtaining permission under subsection (1) of section 12 of the Pharmacy Act, shall submit a scheme as prescribed by the Pharmacy Council of India.
 - b. The scheme referred to in sub-regulation
- (2) above, shall be in such form and contain such particulars and be preferred in such manner and be accompanied with such fee as may be prescribed:

Provided that the Pharmacy Council of India shall not approve any institution under these regulations unless it provides adequate arrangements for teaching in regard to building, accommodation, labs, equipments, teaching staff, non-teaching staff, etc., as specified in Appendix-B to these regulations.

13. Examination:

- (1) Every year there shall be an examination to examine the students.
- (2) Each examination may be held twice every year. The first examination in a year shall be the annual examination and the second examination shall besupplementary examination.
- (3) The examinations shall be of written and Practical (including oral nature) carrying maximum marks for each part of a subject as indicated in Tables below:

TABLES

Course Coole	Norma de Carles d	Internal Assessment		Exami	Total	
Course Code	Name of Subject	Marks	Duration	Marks	Duration	Marks
24PD101T	Human Anatomy and Physiology -Theory	30	1Hr	70	3 Hrs	100
24PD102T	Pharmaceutics - Theory	30	1Hr	70	3 Hrs	100
24PD103T	Medicinal Biochemistry -Theory	30	1Hr	70	3 Hrs	100
24PD104T	Pharmaceutical Organic Chemistry -Theory	30	1Hr	70	3 Hrs	100
24PD105T	Pharmaceutical Inorganic Chemistry- Theory	30	1Hr	70	3 Hrs	100
24PD106RMT / 24PD106RBT	Remedial Mathematics - Theory*/ Remedial Biology - Theory*	30	1Hr	70	3 Hrs	100
24PD107P	Human Anatomy and Physiology- Practical	30	3Hrs	70	3 Hrs	100
24PD108P	Pharmaceutics- Practical	30	3Hrs	70	3 Hrs	100
24PD109P	Medicinal Biochemistry -Practical	30	3Hrs	70	3 Hrs	100
24PD110P	Pharmaceutical Organic Chemistry -Practical	30	3Hrs	70	3 Hrs	100
24PD111P	Pharmaceutical Inorganic Chemistry -Practical	30	3Hrs	70	3 Hrs	100
24PD112RBP	Remedial Biology -Practical**	30*	3Hrs	70*	3 Hrs	100*
24PD113ET / 24PD114ET	Communication Skills -Theory** */	30	1Hrs	70	3 Hrs	100
	Yoga for Youth Empowerment - Theory***					
	Total	360/390 390	25 Hrs	770/84 0/910	39 Hrs	1100/1200 /1300

TABLE VII: FIRST YEAR EXAMINATION

* Remedial Mathematics Theory/ Remedial Biology Theory

** For Remedial Biology Practical

*** Two Electives offered, students can choose any one.

Course Code	Name of Subject	Internal Assessment		Examination		Total
		Marks	Duration	Marks	Duration	Marks
24PD201T	Pathophysiology- Theory	30	1 Hr	70	3 Hrs	100
24PD202T	Pharmaceutical Microbiology -Theory	30	1 Hr	70	3 Hrs	100
24PD203T	Pharmacognosy and Phytopharmaceuticals -Theory	30	1 Hr	70	3 Hrs	100
24PD204T	Pharmacology I- Theory	30	1 Hr	70	3 Hrs	100
24PD205T	Community Pharmacy - Theory	30	1 Hr	70	3 Hrs	100
24PD206T	Pharmacotherapeutics I - Theory	30	1 Hr	70	3 Hrs	100
24PD207P	Pharmaceutical Microbiology - Practical	30	3 Hrs	70	3 Hrs	100
24PD208P	Pharmacognosy and Phytopharmaceutical -Practical	30	3 Hrs	70	3 Hrs	100
24PD209P	Pharmacotherapeutics I-Practical	30	3 Hrs	70	3 Hrs	100
24PD210ET/ 24PD211ET	Health and Lifestyle -Theory*/ NPTEL-1*	30	1 Hr	70	3 Hrs	100
	Total	300	16 Hrs	700	30 Hrs	1000

TABLE VIII: SECOND YEAR EXAMINATION

* Two Electives offered, students can choose any one.

TABLE IX: THIRD YEAR EXAMINATION

Course Code	Name of Subject	Internal Assessment		Examiı	Total	
Course Coue	Nume of Subject	Marks	Duration	Marks	Duration	Marks
24PD301T	Pharmacology II- Theory	30	1 Hr	70	3 Hrs	100
24PD302T	Pharmaceutical Analysis- Theory	30	1 Hr	70	3 Hrs	100
24PD303T	Pharmacotherapeutics II - Theory	30	1 Hr	70	3 Hrs	100
24PD304T	Pharmaceutical Jurisprudence - Theory	30	1 Hr	70	3 Hrs	100
24PD305T	Medicinal Chemistry- Theory	30	1 Hr	70	3 Hrs	100
24PD306T	Pharmaceutical Formulations -Theory	30	1 Hr	70	3 Hrs	100
24PD307P	Pharmacology II- Practical	30	3 Hrs	70	3 Hrs	100
24PD308P	Pharmaceutical Analysis- Practical	30	3 Hrs	70	3 Hrs	100
24PD309P	Pharmacotherapeutics II- Practical	30	3 Hrs	70	3 Hrs	100
24PD310P	Medicinal Chemistry- Practical	30	3 Hrs	70	3 Hrs	100
24PD311P	Pharmaceutical Formulations- Practical	30	3 Hrs	70	3 Hrs	100
24PD312ET/ 24PD313ET	Indian Indigenous Medicine -Theory*/ NPTEL-2*	30	1 Hr	70	3 Hrs	100

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** Two Electives offered, students can choose any one.

TABLE X: FOURTH YEAR EXAMINATION

Course Code	Name of Subject	Internal Assessment		Examination		Total Marks
		Marks	Duration	Marks	Duration	11111115
24PD401T	Pharmacotherapeutics III-	30	1 Hr	70	3 Hrs	100
	Theory					
24PD402T	Hospital Pharmacy- Theory	30	1 Hr	70	3 Hrs	100
24PD403T	Clinical Pharmacy- Theory	30	1 Hr	70	3 Hrs	100
24PD404T	Biostatistics and Research	30	1 Hr	70	3 Hrs	100
	Methodology- Theory					
24PD405T	Biopharmaceutics and	30	1 Hr	70	3 Hrs	100
	Pharmacokinetics- Theory					
24PD406T	Clinical Toxicology- Theory	30	1 Hr	70	3 Hrs	100
24PD407P	Pharmacotherapeutics III-	30	3 Hrs	70	3 Hrs	100
	Practical					
24PD408P	Hospital Pharmacy -Practical	30	3 Hrs	70	3 Hrs	100
24PD409P	Clinical Pharmacy- Practical	30	3 Hrs	70	3 Hrs	100
24PD410P	Biopharmaceutics and	30	3 Hrs	70	3 Hrs	100
	Pharmacokinetics-Practical					
24PD411EP/	Statistical Software- Practical**/		3 Hrs			
24PD412ET	Ethical Leadership- Theory**	30	1 Hr	70	3 Hrs	100
	Total	330	21/19 Hrs	770	33 Hrs	1100

** Two Electives offered, students can choose any one.

TABLE XI: FIFTH YEAR EXAMINATION

Course Code	Name of Subject	Internal Assessment		Examination		Total Marks
		Marks	Duration	Marks	Duration	
24PD501T	Clinical Research- Theory	30	1 Hr	70	3 Hrs	100
24PD502T	Pharmacoepidemiology and Pharmacoeconomics -Theory	30	1 Hr	70	3 Hrs	100
24PD503T	Clinical Pharmacokinetics and Pharmacotherapeutics Drug Monitoring -Theory	30	1 Hr	70	3 Hrs	100
24PD504S	Clerkship *	30	1 Hr	70	3 Hrs	100
24PD505P	Project work** (Six Months)	-	-	100	-	100
24PD506ET /	Medical Coding -Theory ***/		1 Hr			
24PD507EP	Pharmaceutical Calculation -Practical***	30	3 Hrs	70	3 Hrs	100
	Total	150	5 /7Hrs	450	15 Hrs	600

*Attending ward rounds on daily basis.

**30 marks – viva-voce (oral)

70 marks – Thesis work

*** Two Electives offered, students can choose any one.

CIA	1470/1500#/1530*	ESE	5000/5100#/5200*	Grand Total	6470/6600#/6730*				
# for Remedial Mathematics / Biology *for Remedial Biology Practicals									

14. Eligibility for appearing Examination:

Only such students who produce certificate from the Head of the Institution in which he or she has undergone the Pharm.D or as the case may be, the Pharm.D (Post Baccalaureate) course, in proof of his or her having regularly and satisfactorily undergone the course of study by attending not less than 80% of the classes held both in Theory and in Practical separately in each subject shall be eligible for appearing at examination.

15. Mode of examinations:

- (1) Theory examination shall be of three hours and Practical examination shall be of three hours duration.
- (2) A Student who fails in Theory or Practical examination of a subject shall re-appear both in Theory and Practical of the same subject.
- (3) Practical examination shall also consist of a viva -voce (Oral) examination.
- (4)Clerkship examination Oral examination shall be conducted after the completion of clerkship of students. An external and an internal examiner will evaluate the student. Students may be asked to present the allotted medical cases followed by discussion. Students' capabilities in delivering clinical pharmacy services, pharmaceutical care planning and knowledge of the rapeutics shall be assessed.

16. Award of sessional marks and maintenance of records:

A regular record of both Theory and Practical class work and examinations conducted in an institution imparting training for Pharm.D. or as the case may be, Pharm.D. (Post Baccalaureate) course, shall be maintained for each student in the institution and 30 marks for each Theory and 30 marks for each Practical subject shall be allotted as sessional. There shall be at least two periodic sessional examinations during each academic year and the average marks of at least twosessional exams shall be computed for internal assessment. The sessional marksin Practicals shall be allotted on the following basis:-

- i. Actual performance in the sessional examination (20 marks)
- Day to day assessment in the Practical class work, promptness, vivavocerecord maintenance, etc. (10 marks)

Minimum marks for passing examination:

A student shall not be declared to have passed examination unless he or she secures at least 50% marks in each of the subjects separately in the Theory examinations, including sessional marks and at least 50% marks in each of the Practical examinations including sessional marks. The students securing 60% marks or above in aggregate in all subjects in a single attempt at thePharm.D or as the case may be, Pharm. D. (Post Baccalaureate) course examination shall be declared to have passed in first class. Students securing 75% marks or above in any subject or subjects shall be declared to have passed with distinction in the subject or those subjects provided he or she passes in all the subjects in a single attempt.

Question paper pattern for Theory Sessional examinations

For 30 marks paper

I. II.	Short Answers (Answer 1 out of 2)	=	$1 \times 15 = 15$ $3 \times 5 = 15$	
	Total =			rks

Question paper pattern for Practical Sessional examinations

For 30 marks paper

I. II. III.	Synopsis Experiments Viva voce		= = =	5 13 (10+3) 2
		Total		20 marks
		iotai	_	av man No

A total sessional mark is 30 (20 for Practical sessional plus 10 marks for

regularity,

promptness, viva-voce and record maintenance).

Question paper pattern for Theory Examinations

For 70 marks paper

	Total	=	70 marks
111.	Short Allswers (Allswer 5 out of 7)	=	$3 \times 2 = 10$
ш	Short Answers (Answer 5 out of 7)		5 = 2 = 10
II.	Short Essay (Answer 6 out of 8)	=	$6 \ge 5 = 30$
I.	Long Essay (Answer 2 out of 3)	=	$2 \times 15 = 30$

Question paper pattern for Practical Examinations

For 70 marks paper

		Total	=	70 marks
III.	Viva voce		=	15
II.	Experiments		=	40 (25+15)
I.	Synopsis		=	15

17. Grading of performances

17.1. Letter grades and grade points allocations:

Based on the performances, each student shall be awarded a final letter grade at the end of the semester for each course. The letter grades and their corresponding grade points are given in Table – XII.

Table XII: LETTER GRADES AND GRADE POINTS EQUIVALENT TO PERCENTAGE OF MARKS AND PERFORMANCES

Letter Grade	Marks Range	Grade Point	Description
0	91 - 100	10	OUTSTANDING
A+	81 - 90	9	EXCELLENT
А	71-80	8	VERY GOOD
B+	66-70	7	GOOD
В	61-65	6	ABOVE AVERAGE
С	55-60	5	AVERAGE
D	50-54	4	PASS
RA	<50	0	REAPPEARANCE
AB		0	ABSENT

A learner who remains absent for any end semester examination shall be assigned a letter Grade of AB and a corresponding grade point of zero. He/she should reappear for the said evaluation/examination in due course.

18. Grade Sheet

After results are declared, Grade sheet will be issued to each student which will contain the following details:

- i. The list of courses enrolled during the year and the grade scored,
- ii. The Grade Point Average(GPA) for the year and

- iii. The Cumulative Grade Point Average (CGPA) of all courses enrolled from first years onwards.
- iv. The Cumulative Grade point Average (GPA) of all course enrolled from first year onwards.

GPA is ratio of the sum of the products of the number of credits © of courses enrolled and the Grade Points(GP) corresponding to the grades scored in those courses, taken for all the courses to the sum of the number of credits of all the courses in the year.

GPA= Sum of [C x GP]

Sum of C

CGPA will be calculated in a similar manner, considering all the courses enrolled from First year. RA grade and value added course will be excluded for calculating GPA and CGPA.

19. Revaluation

Revaluation and Re-totaling is allowed on representation. A candidate can apply for revaluation of his/her semester Examination answer paper in a Theory course, within 2 weeks from the declaration of results, on payment of a prescribed fee through proper application to the Controller of Examinations through the Head of the Department and Dean. A candidate can apply for revaluation of answer scripts for not exceeding 4 subjects at a time. The Controller of Examinations will arrange for the revaluation and the results will be intimated to the candidate through the Head of the Department and Dean. Revaluation is not permitted for Supplementary Examinations, Practical Examinations, Technical Seminars, In-plant Training and Project Work.

20. Declaration of class

The class shall be awarded on the basis of CGPA as follows:

First Class with Distinction	=	8 and above
First Class	=	6.50 to 7.99
Second Class	=	5.00 to 6.49

21. Eligibility for promotion to next year:

All students who have appeared for all the subjects and passed the first year annual examination are eligible for promotion to the second year and, so on. However, failure in more than two subjects shall debar him or her from promotion to the next year classes as per 2008.

Amendment: Ref. No. 14-194/2011-PCI dated 18/04/2012.

- A) There is a provision for supplementary examination under regulation 10 of the Pharm.D. Regulations, 2008.
- B) Regulation 15 and 10 of the Pharm.D. Regulations, 2008 are interwoven and are to be read together.
- C) Hence the failed students of annual examination are eligible to appear for supplementary examination under regulation 10 of the Pharm.D. Regulations, 2008 and if the failed students of annual examination pass in the supplementary examination they are eligible for promotion to next higher class without losing additional six months. However, failure in more than 2 subjects in the supplementary examination shall debar the students from promotion to the next year classes.
- C1]. Candidates of I Pharm.D are permitted to carry not more than any two subjects (Two Theory/ Two Practicals/ One Theory & one Practical of same or different subjects) to II Pharm.D and appear for II Pharm.D examination concurrently along with failed subjects of I Pharm.D. However, these candidates have to pass all the failed subjects of I Pharm.D to become eligible to III Pharm.D.
- C2]. Similarly, candidates of II Pharm.D who have completely passed all the subjects of I Pharm.D but have failed in II Pharm.D are permitted to carry not more than any two subjects (Two Theory/ Two Practicals/ One Theory & one Practical of same or different subjects) of II Pharm.D to III Pharm.D and appear for III Pharm.D concurrently along with failed subjects of II Pharm.D. However, these candidates have to pass all the failed subjects of II Pharm.D to become eligible to proceed to IV Pharm.D.
- C3]. Candidates of III Pharm.D who have completely passed all the subjects of II Pharm.D but have failed in III Pharm.D are permitted to carry not more than any two subjects (Two Theory/ Two Practicals/ One Theory & one Practical of same or different subjects) of III Pharm.D to IV Pharm.D and appear for IV Pharm.D examination concurrently along with failed subjects of III Pharm.D. However, these candidates have to pass all the failed subjects of III Pharm.D to become eligible to proceed to V Pharm.D.

C4]. Candidates of IV Pharm.D who have completely passed all the subjects of III Pharm.D but have failed in IV Pharm.D are permitted to carry not more than any two subjects (Two Theory/ Two Practicals/ One Theory & one Practical of same or different subjects) of IV Pharm.D to V Pharm.D and appear for V Pharm.D examination concurrently along with failed subject of IV Pharm.D. However, these candidates have to pass all the failed subjects of IV and V Pharm. D to become eligible to proceed to VI Pharm. D., to undergo internship.

22. Internship:

- (1) Internship is a phase of training wherein a student is expected to conduct actual practice of pharmacy and health care and acquires skills under the supervision so that he or she maybecome capable of functioning independently.
- (2) Every student has to undergo one year internship as per Appendix-C to these regulations.

23. Approval of examinations:

Examinations mentioned in regulations 10 to12 and 14 shall be held by the examining authority hereinafter referred to as the university, which shall be approved by the Pharmacy Council of India under sub-section (2) of section 12 of the Pharmacy Act, 1948. Such approval shall be granted only if the examining authority concerned fulfills the conditions as specified in Appendix–D to these regulations.

24. Certificate of passing examination:

Every student who has passed the examinations for the Pharm.D (Doctor of Pharmacy) orPharm.D (Post Baccalaureate) (Doctor of Pharmacy) as the case may be, shall be granted a certificate by the examining authority.

CHAPTER - III PRACTICAL TRAINING

25. Hospital posting:

Every student shall be posted in constituent hospital for a period of not less than fifty hours to be covered in not less than 200 working days in each of second, third & fourth year course. Each student shall submit report duly certified by the preceptor and duly attested by the Head of the Department or Institution as prescribed. In the fifth year, every student shall spend half a day in the morning hours attending ward rounds on dailybasis as a part of clerkship. Theory teaching may be scheduled in the afternoon.

26. Project work:

(1) To allow the student to develop data collection and reporting skills in the area of community, hospital and clinical pharmacy, a project work shall be carried out under the supervision of a teacher. The project topic must be approved by the Head of the Department or Head of the Institution. The same shall be announced to students within one month of commencement of the fifth year classes. Project work shall be presented in a written report and as a seminar at the end of the year. External and the internal examiners shall do the assessment of the project work.

(2) Project work shall comprise of objectives of the work, methodology, results, discussions and conclusions.

27. Objectives of project work:

The main objective of the project work is to --

- (i) show the evidence of having made accurate description of published work of others and of having recorded the findings in an impartial manner; and
- (ii) develop the students in data collection, analysis and reporting and interpretation skills.

28. Methodology:

- To complete the project work following methodology shall be adopted, namely,
- (i) students shall work in groups of not less than *two* and not more than *four* under an authorised lectures;

- (ii) project topic shall be approved by the Head of the Department or Head of the Institution;
- (iii) project work chosen shall be related to the pharmacy practice in community, hospital and clinical setup. It shall be patient and treatment (Medicine) oriented, like drug utilisation reviews, pharmacoepidemiology, Pharmacovigilance or Pharmacoeconomics;
- (iv) project work shall be approved by the institutional ethics committee;
- (v) student shall present at least three seminars, one in the beginning, one at middle and one at the end of the project work; and
- (vi) two-page write-up of the project indicating title, objectives,Methodology anticipated benefits and references shall be submitted to the Head of the Department or Head of the Institution.

29. Reporting:

(1) Student working on the project shall submit jointly to the Head of the Department or Head of the Institution a project report of about 40-50 pages. Project report should include a certificate issued by the authorised lecturer, Head of the Department as well as by the Head of the Institution.

(2) Project report shall be computer typed in double space using Times New Roman font onA4 paper. The title shall be in bold with font size 18, sub-tiles in bold with font size14 and the text with font size 12. The cover page of the project report shall contain details about the name of the student and the name of the authorized teacher with font size 14.

(3) Submission of the project report shall be done at least one month prior to the commencement of annual or supplementary examination.

30. Evaluation:

The following methodology shall be adopted for evaluating the project work.

- (i) Project work shall be evaluated by internal and external examiners.
- (ii) Students shall be evaluated in groups for four hours (i.e., about half an hour for a group of four students).
- (iii) Three seminars presented by students shall be evaluated for twenty marks each and the average of best two shall be forwarded to the university with marks of other subjects.

(iv) Evaluation shall be done on the following items:	Marks
a) Write up of the seminar	(7.5)
b) Presentation of work	(7.5)
c) Communication skills	(7.5)
d) Question and answer skills	(7.5)

	Total	(30 marks)
(v) Final evaluation of project work shall be do	one on the following it	ems: Marks
a) Write up of the seminar		(17.5)
b) Presentation of work		(17.5)
c) Communication skills		(17.5)
d) Question and answer skills		(17.5)
	Total	(70 marks)

Explanation:

For the purposes of differentiation in the evaluation in case of topic being the same for the group of students, the same shall be done based on item numbers b, c and d mentioned above.

31. a. Elective / Online Courses

Students shall study the value added elective courses / online courses are designed to enhance the capability of students beyond the general academic curriculum, which may help to improve the employability and equip the students with essential skills to succeed. The program offers five categories of elective courses such as Human value, Social Responsibility, Indian Knowledge system, Skill based and online course from NPTEL courses. All elective /online courses carry 2 credit points. Students can select any one elective courses offered in each year of study from the list. Examination shall be conducted at the end of the respective year of study. The student shall produce a pass certificate from the respective NPTEL organizations. The credit(s) earned by the students will be considered as additional credit(s) over and above the required credits earned from programme concerned.

31. b. Online Course Co-ordinator

To help students in planning their online courses and for general advice on online courses, the HOD shall nominate a co-ordinator for the online courses. The Online course co- ordinator shall identify the courses which students can select for their programme from the available online courses offered by the different agencies periodically and inform the same to the students. Further, the co-ordinator shall advice the students regarding the online courses and monitors their course.

32. Award of Ranks

Ranks and Medals shall be awarded on the basis of final CGPA. However, candidates who fail in one or more courses during the Pharm.D program shall not be eligible for award of ranks. Moreover, and the candidates should have completed the Pharm.D program in minimum prescribed number of years, (six years) for the award of Ranks.

33. Award of degree

Candidates who fulfill the requirements mentioned above shall be eligible for award of degree during the ensuing convocation.

34. Duration for completion of the program of study

The duration for the completion of the program shall be fixed as double the actual duration of the program and the students have to pass within the said period, otherwise they have to get fresh registration.

35. Re-admission after break of study

Candidate who seeks re-admission to the program after break of study has to get the approval from the university by paying a condonation fee.

No condonation is allowed for the candidate who has more than 2 years of breakup period and he/she has to rejoin the program by paying the required fees.

36. <u>Credit Transfer For Online Courses Through NPTEL, MOOCs Platform/</u> <u>International Institutions</u>

Students are encouraged to enroll in courses offered by MOOC platforms and international institutions of higher learning, either virtually or in person. The equivalent credits for these courses will be determined by a committee named Subject Equivalency Committee comprising the Dean, Head of Department (HoD), and one faculty member nominated by the Vice Chancellor. The committee's decision will be submitted for ratification/approval by the Board of Studies (BoS) and the Academic Council. Additionally, the equivalent grade points for marks/grades/grade points awarded by various MOOC platforms and international institutions of higher learning will be determined by a committee named Grade Equivalency Committee duly constituted by the Vice-Chancellor. The decisions of this committee will also be submitted for ratification/approval by the Academic Council. This shall be approved to be implemented from the even semester of the academic year 2024-25.

FACULTY OF PHARMACY PHARM.D PROGRAMME (2024–2025 Batch and onwards)

Course code	Name of the course		ectives 1 out mes	Instruction hours / week			cdit(s)	Max	Page no		
		EOs	POs	L	Т	Р	Cre	CIA	ESE	Total	
		A						30	70	100	
YEAR – I	1			I				T	1	1	
24PD101T	Human Anatomy and Physiology- Theory	2	a, k	3	1	-	4	30	70	100	1-4
24PD102T	Pharmaceutics- Theory	3	a	2	1	-	3	30	70	100	7-8
24PD103T	Medicinal Biochemistry -Theory	2	a,k	3	1	-	4	30	70	100	11-13
24PD104T	Pharmaceutical Organic Chemistry- Theory	3	a	3	1	-	4	30	70	100	16-19
24PD105T	Pharmaceutical Inorganic Chemistry- Theory	2	a	2	1	-	3	30	70	100	23-23
24PD106RM T/ 24PD106PPT	Remedial Mathematics**/ Remedial Biology**-	6	c/a	3	1	-	4	30	70	100	26-27 28-29
24FD100KD1	Human Anatomy and	2	ha			3	2	30	70	100	5.6
24PD107P	Physiology- Practical	2	0,0	-	-	3	2	30	70	100	5-0
24PD108P	Pharmaceutics- Practical	3	a,b	-	-	3	2	30	70	100	9-10
24PD109P	Medicinal Biochemistry -Practical	2	b,k	-	-	3	2	30	70	100	14-15
24PD110P	Pharmaceutical Organic Chemistry- Practical	2	b	-	-	3	2	30	70	100	20-21
24PD111P	Pharmaceutical Inorganic Chemistry- Practical	2	b	1	-	3	2	30	70	100	24-25
24PD112RBP	Remedial Biology -Practical**	2	b	-	-	3*	2*	30	70	100	30-31
24PD113ET	Communication Skills	4	f,h	2	-	-	1	30	70	100	
/ 24PD114ET	- Theory*** / Yoga for Youth Empowerment -Theory***	6	f, i, k								
	Total			18	6	15/ 18*	36/ 38*	390	910	1300	

*Applicable for Remedial Biology Practical

** For Remedial Mathematics / Remedial Biology (Non University Exam)

*** Two Electives offered, students can choose any one.

YEAR – II											
24PD201T	Pathophysiology- Theory	6	b,f,i	3	1	-	4	30	70	100	32-35
24PD202T	Pharmaceutical Microbiology - Theory	3	k	3	1	-	4	30	70	100	36-38
24PD203T	Pharmacognosy and Phytopharmaceuticals- Theory	2	a	3	1	-	4	30	70	100	41-42
24PD204T	Pharmacology I- Theory	3	a.d.k	3	1	-	4	30	70	100	45-48
24PD205T	Community Pharmacy - Theory	1,4	a,f,i	2	1	-	3	30	70	100	49-51
24PD206T	Pharmacotherapeutics I - Theory	3	a,f,k	3	1	-	4	30	70	100	52-54
24PD207P	Pharmaceutical Microbiology- Practical	3	a,b	-	-	3	2	30	70	100	39-40

Course code	Name of the course	Objecti out c	bjectives and out comes		Instruction hours / week		Instruction hours / week		Instruction hours / week		Instruction 10urs / week		(s)	Maximum Marks			Page no
		PEOs	POs	L	Т	Р	Credit	CIA	ESE	Total							
								30	70	100	1						
24PD208P	Pharmacognosy and Phytopharmaceuticals - P r a c t i c a l	2	a.b	-		3	2	30	70	100	43-44						
24PD209P	Pharmacotherapeutics I - Practical	3	b,c,g	-	-	3	2	30	70	100	55-56						
24PD210ET /	Health and Lifestyle -Theory*	4	g, j,k	2			2	20	70	100							
24PD211ET	NPTEL-1*	6	i,	2	-	-	2	50	70	100							
	Total			19	6	9	31	330	770	1000							
* Two Electives	s offered, students can choose any o	one.	•			•	•	•	•		·						

YEAR – III											
24PD301T	Pharmacology II - Theory	3	a,d,k	3	1	-	4	30	70	100	57-60
24PD302T	Pharmaceutical Analysis - Theory	2	с	3	1	-	4	30	70	100	63-66
24PD303T	Pharmacotherapeutics II - Theory	3	a,f,k	3	1	-	4	30	70	100	69-70
24PD304T	Pharmaceutical	5	a,e,g	2	-	-	2	30	70	100	73-75
	Jurisprudence-Theory										
24PD305T	Medicinal Chemistry- Theory	2	a.k	3	1	-	4	30	70	100	76-78
24PD306T	Pharmaceutical	4	a,c,k	2	1	-	3	30	70	100	81-82
	Formulations-Theory										
24PD307P	Pharmacology II- Practical	3	a,d,k			3	2	30	70	100	61-62
24PD308P	Pharmaceutical Analysis - Practical	2	с	ſ	-	3	2	30	70	100	67-68
24PD309P	Pharmacotherapeutics II- Practical	3	a,f,k	-	-	3	2	30	70	100	71-72
24PD310P	Medicinal Chemistry -Practical	2	a,b	ı	-	3	2	30	70	100	79-80
24PD311P	Pharmaceutical Formulations- Practical	4	a,c	-	1	3	2	30	70	100	83-84
24PD312ET/	Indian Indigenous Medicine	6	i, j				2	30	70	100	
24PD313ET	-Theory*/ NPTEL-2*	6	i								
	Total			16	5	15	33	360	840	1200	
VEAR – IV											
24PD401T											
	Pharmacotherapeutics III- Theory	3	a.f.k	3	1	-	4	30	70	100	85-86
24PD401T 24PD402T	Pharmacotherapeutics III- Theory Hospital Pharmacy- Theory	3 1,6	a,f,k a,f,g ,i,k	3	1	-	4 3	30 30	70 70	100 100	85-86 89-91
24PD401T 24PD402T 24PD403T	Pharmacotherapeutics III- Theory Hospital Pharmacy- Theory Clinical Pharmacy- Theory	3 1,6 1,6	a,f,k a,f,g ,i,k a,f,g,i, k	3 2 3	1 1 1	-	4 3 4	30 30 30	70 70 70	100 100 100	85-86 89-91 94-96
24PD402T 24PD402T 24PD403T 24PD404T	Pharmacotherapeutics III- TheoryHospital Pharmacy- TheoryClinical Pharmacy- TheoryBiostatistics and Research Methodology- Theory	3 1,6 1,6 2	a,f,k a,f,g ,i,k a,f,g,i, k b,c,d,k	3 2 3 2	1 1 1	-	4 3 4 3	30 30 30 30	70 70 70 70	100 100 100 100	85-86 89-91 94-96 99-101
24PD401T 24PD402T 24PD403T 24PD404T 24PD405T	Pharmacotherapeutics III- TheoryHospital Pharmacy- TheoryClinical Pharmacy- TheoryBiostatistics and Research Methodology- TheoryBiopharmaceutics and Pharmacokinetics -Theory	3 1,6 1,6 2 5	a,f,k a,f,g ,i,k a,f,g,i, k b,c,d,k a,c,k	3 2 3 2 3	1 1 1 1	-	4 3 4 3 4	30 30 30 30 30 30	70 70 70 70 70 70	100 100 100 100 100	85-86 89-91 94-96 99-101 102-104
24PD401T 24PD402T 24PD403T 24PD404T 24PD405T 24PD406T	Pharmacotherapeutics III- TheoryHospital Pharmacy- TheoryClinical Pharmacy- TheoryBiostatistics and Research Methodology- TheoryBiopharmaceutics and Pharmacokinetics - TheoryClinical Toxicology- Theory	3 1,6 1,6 2 5 3	a,f,k a,f,g ,i,k a,f,g,i, k b,c,d,k a,c,k a,g,k,i	3 2 3 2 3 2 3 2	1 1 1 1 1 1 1	-	4 3 4 3 4 3	30 30 30 30 30 30 30	70 70 70 70 70 70 70	100 100 100 100 100 100	85-86 89-91 94-96 99-101 102-104 107-109
24PD401T 24PD402T 24PD403T 24PD404T 24PD405T 24PD406T 24PD406T 24PD407P	Pharmacotherapeutics III- TheoryHospital Pharmacy- TheoryClinical Pharmacy- TheoryBiostatistics and Research Methodology- TheoryBiopharmaceutics and Pharmacokinetics -TheoryClinical Toxicology- TheoryPharmacotherapeutics III- Practical	3 1,6 1,6 2 5 3 3	a,f,k a,f,g ,i,k a,f,g,i, k b,c,d,k a,c,k a,c,k a,g,k,i a,f,k	3 2 3 2 3 2 -	1 1 1 1 1 -	3	4 3 4 3 4 3 2	30 30 30 30 30 30 30 30 30 30 30 30 30 30 30 30 30 30 30	70 70	100 100 100 100 100 100 100 100 100 100	85-86 89-91 94-96 99-101 102-104 107-109 87-88
24PD401T 24PD402T 24PD403T 24PD404T 24PD405T 24PD406T 24PD406T 24PD407P 24PD408P	Pharmacotherapeutics III- TheoryHospital Pharmacy- TheoryClinical Pharmacy- TheoryBiostatistics and Research Methodology- TheoryBiopharmaceutics and Pharmacokinetics -TheoryClinical Toxicology- TheoryPharmacotherapeutics III- PracticalHospital Pharmacy- Practical	3 1,6 1,6 2 5 3 3 1,6	a,f,k a,f,g i,k a,f,g,i, k b,c,d,k a,c,k a,c,k a,g,k,i a,f,k a,f,g,i, k	3 2 3 2 3 2 - -	1 1 1 1 1 1 -	- - - - 3 3	4 3 4 3 4 3 2 2	30 30	70 70	100 100 100 100 100 100 100 100	85-86 89-91 94-96 99-101 102-104 107-109 87-88 92-93

24PD410P	Biopharmaceutics and Pharmacokinetics- Practical	3,5	a,c,k	-	-	3	2	30	70	100	105-106
Course code	Name of the course	Obje andou	ectives It comes	Ir ho	nstruct ours / v	ion veek		Maxin	Maximum Marks		
		PEOs	POs	L	Т	Р	Credit(s)	CIA	ESE	Total	
								30	70	100	
24PD411EP/ 24PD412ET	Statistical Software -Practical*/ Ethical Leadership- Theory *	4	c,d,k		-	3	2	30	70	100	
	p	2	e,g,k	2		-	2				
	Total			17	5	15	33	330	770	1100	

* Two Electives offered, students can choose any one.

YEAR V											
24PD501T	Clinical Research - Theory	1,4,6	a,f,i,g, k	3	1	-	4	30	70	100	110-112
24PD502T	Pharmacoepidemiology and Pharmacoeconomics - Theory	3,6	a,d,j	3	1	-	4	30	70	100	113-115
24PD503T	Clinical Pharmacokinetics &Pharmacotherapeutic Drug Monitoring -Theory	3,5	a,c,k	2	1	-	3	30	70	100	116-118
24PD504S	Clerkship *	1,3,5,6	c,e,f,g ,h,i k	-	-	1	1	30	70	100	119
24PD505P	Project work (Six Months)	1,3,5, 6	a,b,c,d ,e,f,g, h,i,j,k	-	-	20	10	-	100**	100	120
24PD506ET/ 24PD507EP	Medical Coding - Theory***/ Pharmaceutical Calculation	4	f	1	-	-	2	30	70	100	
	-Practical***	6	c,d	-	-	3	2				
	Total			10	4	23	24	150	450	600	

*Attending ward rounds on daily basis

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** 30 marks – Viva- Voice (oral)
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70 marks – Thesis work
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YEAR VI

Internship or residency training including postings in speciality units. Student should independently provide the clinical pharmacy services to the allotted wards.

- (i) Six months in General Medicine department, and
- (ii) Two months each in three other speciality departments.

PROGRAM OUTCOMES (PO's)

- a. Pharmacy Knowledge: Possess knowledge and comprehension of the core and basic knowledge associated with the profession of pharmacy, including biomedical sciences; pharmaceutical sciences; behavioral, social, and administrative pharmacy sciences; and practice of pharmacy.
- b. **Planning Abilities:** Demonstrate effective planning abilities including timemanagement, resource management, delegation skills and organizational skills. Develop and implement plans and organize work to meet deadlines.
- c. Problem analysis: Utilize the principles of scientific enquiry, thinking analytically, clearly and critically, while solving problems and making decisions during daily practice.
 Find, analyze, evaluate and apply information systematically and shall make defensible decisions.
- d. **Modern tool usage:** Learn, select, and apply appropriate methods and procedures, resources, and modern pharmacy-related computing tools with an understanding of the limitations and adherence to regulatory standards.
- e. Leadership skills: Understand and consider the human reaction to change, motivation issues, leadership and team-building when planning changes required for fulfillment of practice, professional and societal responsibilities. Assume participatory roles as responsible citizens or leadership roles when appropriate to facilitate improvement in health and wellbeing.
- f. **Professional Identity:** Understand, analyze and communicate the value of their professional roles in society (e.g. health care professionals, promoters of health, educators, managers, employers, employees).
- g. **Pharmaceutical Ethics:** Honour personal values and apply ethical principles in professional and social contexts. Demonstrate behavior that recognizes cultural and personal variability in values, communication and lifestyles. Use ethical frameworks; apply ethical principles while making decisions and take responsibility for the outcomes associated with the decisions.
- h. **Communication:** Communicate effectively with patients, healthcare professionals, and the community, providing clear and accurate information on medications, treatments, and health promotion and able to comprehend and write effective reports, make effective documentation.
- i. **The Pharmacist and society:** Apply reasoning informed by the contextual knowledge to assess societal, health, safety and legal issues and the consequent responsibilities relevant to the professional pharmacy practice.

- j. **Environment and sustainability:** Understand the impact of the professional pharmacy solutions in societal and environmental contexts, and demonstrate the knowledge of, and need for sustainable development.
- k. Life-long learning: Recognize the need for, and have the preparation and ability to engage in independent and life-long learning in the broadest context of technological change. Selfassess and use feedback effectively from others to identify learning needs and to satisfy these needs on an ongoing basis.

PROGRAM SPECIFIC OUTCOMES (PSOs)

PSO I: Understand different classes of drugs, their mechanism of action, dynamics, kinetics, structure activity relationships, pathophysiology and pharmacotherapeutics of various diseases, ability to synthesize, develop and/or evaluate various pharmaceuticals and their formulations and cosmeceuticals products.

PSO m: Develop skills in qualitative and quantitative analysis of various pharmaceuticals. Acquire technical knowledge and hands on training on equipments, instruments and software used in the field of pharmaceutical sciences.

PSO n: To inculcate the practice of pharmacy and train pharmacists to play an important role in patient care, health and wellness and population-based care as members of the health care team

PSO o: To exhibit behaviors and values that are consistent with the trust given to the profession, professionalism in interactions with patients, professionalism in interactions with other healthcare providers, professionalism in interactions with society

PSO p: To strengthen the professional and ethical attitude, effective communication skills, teamwork skills, multidisciplinary approach, and an ability to relate pharmaceutical sciences issues to broader social context.

PSO q: To stream a lifelong career of personal and practicing professional growth with ethical codes and self-esteem.

PROGRAM EDUCATIONAL OBJECTIVES (PEOs)

PEO 1: Synthesize population-based drug information to address patient medication adherence, prescribing patterns, and treatment protocol adherence to document issues, alert prescribers, design interventions, and assess intervention effectiveness.

PEO 2: To provide students with a strong and well defined concepts in the various fields of pharmaceutical sciences viz., pharmaceutics, pharmaceutical chemistry, pharmacology and pharmacognosy according to the requirement of pharmaceutical industries, community and hospital pharmacy and also to develop a sense of teamwork and awareness amongst students towards the importance of interdisciplinary approach for developing competence in solving complex problems in the area of Pharmaceutical Sciences.

PEO 3: Identify physicochemical properties of drug substances that affect solubility, pharmacodynamic and pharmacokinetic properties, pharmacologic actions, and stability when designing patient-specific care plans.

PEO 4: Formulate and implement a care plan in cooperation with patients and other healthcare providers based on established, evidence-based standards of practice; provide medication therapy management services for patients with acute & chronic health problems. **PEO 5:** Integrate knowledge of chemical, physical, and biopharmaceutical principles to prepare safe and effective prescriptions (sterile and non-sterile) in conformity with all applicable federal and state laws and regulations.

PEO 6: Provide health care information regarding nutrition, lifestyle, and other non-drug measures that promote health or prevent the progression of a disease or medical condition. Demonstrate a comprehensive approach to practice and care, includes problem solving, educator, patient advocacy, inter-professional collaboration, cultural sensitivity, communication.

РО	a	b	c	d	e	f	g	h	i	j	k	PSO 1	PSO m	PSO n	PSO o	PSO p	PSO q
PEO 1	Х	Х		Х	Х	Х	Х	Х	Х		Х	Х		Х	Х	Х	Х
PEO 2	Х		Х		Х	Х					Х	Х	Х			Х	Х
PEO 3	Х	Х	Х			Х			Х		Х	Х	Х				
PEO 4	Х	Х	Х			Х	Х	Х		Х	Х			Х	Х	Х	Х
PEO 5	Х						Х	Х	Х	Х	Х		Х		Х		
PEO 6	Х		Х			Х	Χ	Х	Х	Х	Х			Х	Х	Х	Х

MAPPING

SYLLABUS

24PD101T

FIRST YEAR

HUMAN ANATOMY AND PHYSIOLOGY - THEORY 4H 4C

Instruction hours/ week: L: 3 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 4 Hrs / Week

Course Objectives: Students can able to

- Impart a fundamental knowledge on the structure and functions of the human body.
- Understanding both homeostasis mechanisms and homeostaticimbalances of various body systems.
- Know the metabolic process of biomolecules in health and illness (metabolic disorders);
- Understanding of how the drugs act on the various body systems incorrecting the disease state of the organs.
- Appreciate coordinated working pattern of different organs of each system;
- Know the biochemical principles of organ function tests of kidney, liver and endocrine gland.

Cos	Course Outcomes	Blooms Level
CO1	Describe the structure (gross and histology) and functions of various organs of thehuman body	Knowledge
CO2	Describe the various homeostatic mechanisms and their imbalances of varioussystems	Analyze
CO3	Discuss the various tissues and organs of the different systems in the human body	Understand
CO4	Perform the hematological tests and also record blood pressure, heart rate, pulse and respiratory volumes	Knowledge
CO5	Understand the coordinated working pattern of different organs of each system	Understand
CO6	Explain the interpret the various organ function tests.	Apply

Mapping with Programme Outcomes

Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12
CO1	S											
CO2	S	М										
CO3	S			Μ								
CO4			S	Μ								
CO5				S	М							
CO6										S		

S-Strong; M-Medium; L-Low

Course Content: UNIT I

- Scope of anatomy and physiology, basic terminologies used in this subject (Description of the body as such planes and terminologies)
- Structure of cell its components and their functions.
- Elementary tissues of the human body: epithelial, connective, Muscular and nervous tissues-their sub-types and characteristics

UNIT II

• Osseous system - structure, composition and functions of the skeleton Classification of joints, Types of movements of joints and disorders of joints(Definitions only)

UNIT III

Haemopoetic System

- Composition and functions of blood
- Haemopoesis and disorders of blood components (definition of disorder)
- Blood groups
- Clotting factors and mechanism
- Platelets and disorders of coagulation

<u>Lymph</u>

- Lymph and lymphatic system, composition, formation and circulation.
- Spleen: structure and functions, Disorders
- Disorders of lymphatic system (definition only)

UNIT IV

Cardiovascular system

- Anatomy and functions of heart
- Blood vessels and circulation (Pulmonary, coronary and systemic circulation)
- Electrocardiogram (ECG)
- Cardiac cycle and heart sounds
- Blood pressure its maintenance and regulation
- Definition of the following disorders
- Hypertension, Hypotension, Arteriosclerosis, Atherosclerosis, Angina, Myocardial infarction, Congestive heart failure, Cardiac arrhythmias

Respiratory system

- Anatomy of respiratory organs and functions
- Mechanism / physiology of respiration and regulation of respiration
- Transport of respiratory gases
- Respiratory volumes and capacities, and Definition of: Hypoxia, Asphyxia, Dybarism, Oxygen therapy and resuscitation.

UNIT V

Digestive system

- Anatomy and physiology of GIT
- Anatomy and functions of accessory glands of GIT
- Digestion and absorption
- Disorders of GIT (definitions only)

UNIT VI

Nervous system

- Definition and classification of nervous system
- Anatomy, physiology and functional areas of cerebrum
- Anatomy and physiology of cerebellum
- Anatomy and physiology of mid brain
- Thalamus, hypothalamus and Basal Ganglia
- Spinal cord: Structure & reflexes mono-poly-planter
- Cranial nerves names and functions
- ANS Anatomy & functions of sympathetic & parasympathetic N.S.

UNIT VII

Urinary system

- Anatomy and physiology of urinary system
- Formation of urine
- Renin Angiotensin system Juxtaglomerular apparatus acid base Balance
- Clearance tests and micturition

Endocrine system

- Pituitary gland
- Adrenal gland

Pharm.D

- Thyroid and Parathyroid glands
- Pancreas and gonads

UNIT VIII

Reproductive system

- Male and female reproductive system
- Their hormones Physiology of menstruation
- Spermatogenesis & Oogenesis
- Sex determination (genetic basis)
- Pregnancy and maintenance and parturition
- Contraceptive devices

UNIT IX

Sense organs

- Eye
- Ear
- Skin
- Tongue & Nose

Skeletal muscles

- Histology
- Physiology of Muscle contraction
- Physiological properties of skeletal muscle and their disorders (definitions)

UNIT X

Sports physiology

- Muscles in exercise, Effect of athletic training on muscles and muscleperformance,
- Respiration in exercise, CVS in exercise, Body heat in exercise, Body fluidsand salts in exercise,
- Drugs and athletics

Suggested Readings:

- 1. Tortora Gerard J. and Nicholas P. Principles of Anatomy and Physiology Publisher Harpercollins College New York.
- 2. Wilson, K.J.W. Ross and Wilson's foundations of Anatomy and Physiology Publisher: Churchill Livingstone, Edinburg.

Reference Books (Latest Editions):

- 1. Guyton Arthur, C. Physiology of human body. Publisher: Holtsaunders.
- 2. Chatterjee C C. Human physiology. Volume 1&11. Publisher: Medical Allied
- 3. Agency, Calcutta.
- 4. Peter L. Williams, Roger Warwick, Mary Dyson and Lawrence, H.
- 5. Gray's Anatomy. Publisher: Churchill Livingstone, London.

24PD102T

PHARMACEUTICS - THEORY

FIRST YEAR **3H 3C**

Instruction hours/ week: L: 2 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 3 Hrs /Week

Course Objectives: Students can able to

- Impart a fundamental knowledge on the preparatory pharmacywith arts of preparing the different conventional dosage forms.
- Understand the history of profession of pharmacy
- Formulate the basics of different dosage forms.
- Appreciate the importance of good formulation for effectiveness.
- Do different pharmaceutical calculation involved in formulation
- Develop a clear idea about Pharmaceutical incompatibility and different pharmaceutical calculations in pharmacy.

Course Outcomes (CO's): On successful completion of the course the student will

COs	Course Outcomes	Blooms Level
CO1	Understand and describe the formulation principles and	Understand
	characteristics of various pharmaceutical dosage forms	
CO2	Understand the professional way of handling the prescription	Apply
CO3	Develop and prepare different types of pharmaceutical dosage forms using appropriate techniques and standards	Create
CO4	Accurately perform and apply pharmaceutical calculations essential for the formulation process	Analyze
CO5	Comprehend the regulatory guidelines and quality assurance protocols governing pharmaceutical formulation practices to ensure compliance and product safety	Apply
CO6	Predict the instability & incompatibility problems in pharmaceutical	Apply

Mapping with Programme Outcomes

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1		S					Μ				
CO2		S					Μ		Μ	L	Μ
CO3	S		S	S	Μ	Μ	Μ	S	Μ		Μ
CO4		Μ	Μ	Μ	Μ	Μ	Μ		Μ	Μ	S
CO5			Μ	S			S		Μ		Μ
CO6			Μ	Μ	Μ						

S-Strong; M-Medium; L-Low

Course Content:

UNIT I

- Introduction to dosage forms classification and definitions
- Prescription: definition, parts and handling

UNIT II

Posology: Definition, Factors affecting dose selection. Calculation of children and infant doses.

UNIT III

Historical back ground and development of profession of pharmacy and pharmaceutical industry in brief.

UNIT IV

Development of Indian Pharmacopoeia and introduction to other Pharmacopoeias such asBP, USP, European Pharmacopoeia, Extra pharmacopoeia and Indian national formulary.

Weights and measures, Calculations involving percentage solutions, allegation, proof spirit, isotonic solutions etc.

UNIT V

Powders and Granules: Classification advantages and disadvantages, Preparation of simple, compound powders, Insufflations, Dusting powders, Eutectic and Explosive powders, Tooth powder and effervescent powders and granules.

UNIT VI

Monophasic Dosage forms: Theoretical aspects of formulation including adjuvant like stabilizers, colorants, flavors with examples. Study of Monophonic liquids like gargles, mouth washes, Throat paint, Ear drops, Nasal drops, Liniments and lotions, Enemas and collodions.

UNIT VII

Biphasic dosage forms: Suspensions and emulsions, Definition, advantages and disadvantages, classification, test for the type of emulsion, formulation, stability and evaluation.

UNIT VIII

Suppositories and pessaries: Definition, advantages and disadvantages, types of base, method of preparation, Displacement value and evaluation.

Galenicals: Definition, equipment for different extraction processes like infusion, Decoction, Maceration and Percolation, methods of preparation of spirits, tinctures and extracts.

UNIT IX

Pharmaceutical calculations.

Surgical aids: Surgical dressings, absorbable gelatin sponge, sutures, ligatures and
Pharm.D

medicated bandages.

UNIT X

Incompatibilities: Introduction, classification and methods to overcome the incompatibilities.

Suggested Readings:

- 1. Cooper and Gunn's Dispensing for pharmacy students.
- 2. A text book Professional Pharmacy by N.K.Jain and S.N.Sharma.

- 1. Introduction to Pharmaceutical dosage forms by Howard C. Ansel.
- 2. Remington's Pharmaceutical Sciences.Register of General Pharmacy by Cooper and Gunn.
- 3. General Pharmacy by M.L.Schroff.

Karpagam Academy of Higher Education

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MEDICINAL BIOCHEMISTRY - THEORY

Instruction hours/ week: L: 3 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Course Objectives: Students can able to

Theory: 4 Hrs. /Week

- Understand of the molecular level of the chemical process associated with living cells. •
- Know the chemical aspects of human life in health and illness and the • application of chemical laboratory methods to diagnosis, control of treatment, and prevention of diseases.
- Understand the catalytic activity of enzymes and importance of isoenzymes in diagnosisof diseases;
- Know the metabolic process of biomolecules in health and illness (metabolic disorders);
- Understand the genetic organization of mammalian genome; protein synthesis; replication; mutation and repair mechanism;
- Know the biochemical principles of organ function tests of kidney, liver and • endocrine gland

Course Outcomes At the end of this course, students will be able to

Cos	Course Outcomes	Blooms Level
CO1	Understand the catalytic activity of enzymes and importance of	Understand
	isoenzymes in diagnosis of diseases;	
CO2	Know the metabolic process of biomolecules in health and illness (metabolic disorders);	Apply
CO3	Understand the genetic organization of mammalian genome; protein synthesis; replication; mutation and repair mechanism;	Understand
CO4	Know the biochemical principles of organ function tests of kidney, liver and endocrine gland.	Apply
CO5	Perform the qualitative analysis and quantitative analysis of body fluids.	Apply
CO6	Understand the basic concepts of biological oxidation.	Understand

Mapping with Programme Outcomes

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12
CO1	Μ										Μ	
CO2	Μ										Μ	
CO3	S										S	
CO4	Μ										Μ	
CO5	S										S	
CO6	S										Μ	

S-Strong; M-Medium; L-Low

24PD103T

FIRST YEAR

2024-25

4C **4H**

Course content: UNIT I

• Introduction to biochemistry: Cell and its biochemical organization, transport process across the cell membranes. Energy rich compounds; ATP, Cyclic AMP and their biological significance.

• **Enzymes**: Definition; Nomenclature; IUB classification; Factor affecting enzyme activity;Enzyme action; enzyme inhibition. Isoenzymes and their therapeutic and diagnostic applications; Coenzymes and their biochemical role and deficiency diseases.

UNIT II

• **Carbohydrate metabolism**: Glycolysis, Citric acid cycle (TCA cycle), HMP shunt, Glycogenolysis, gluconeogenesis, glycogenesis. Metabolic disorders of carbohydrate metabolism (diabetes mellitus and glycogen storage diseases); Glucose, Galactose tolerance test and their significance; hormonal regulation of carbohydrate metabolism.

• Lipid metabolism: Oxidation of saturated (β -oxidation); Ketogenesis and ketolysis; biosynthesis of fatty acids, lipids; metabolism of cholesterol; Hormonal regulation of lipid metabolism. Defective metabolism of lipids (Atheroslerosis, fatty liver, hypercholesterolmiea).

UNIT III

Biological oxidation: Coenzyme system involved in Biological oxidation. Electron transport chain (its mechanism in energy capture; regulation and inhibition); Uncouplers of ETC; Oxidative phosphorylation;

UNIT IV

Protein and amino acid metabolism: protein turn over; nitrogen balance;

Catabolism of Amino acids (Transamination, deamination & decarboxylation). Urea cycle and its metabolic disorders; production of bile pigments; hyperbilirubinemia, porphoria, jaundice. Metabolic disorder of Amino acids.

UNIT V

Nucleic acid metabolism: Metabolism of purine and pyrimidine nucleotides; Protein synthesis; Genetic code; inhibition of protein synthesis; mutation and repair mechanism; DNA replication (semiconservative /onion peel models) and DNA repair mechanism.

UNIT VI

Introduction to clinical chemistry: Cell; composition; malfunction; Roll of the clinical chemistry laboratory.

UNIT VII

The kidney function tests: Role of kidney; Laboratory tests for normal function includes-

- Urine analysis (macroscopic and physical examination, quantitative and semiquantitative tests.)
- Test for NPN constituents. (Creatinine /urea clearance, determination of blood and urine creatinine, urea and uric acid)
- Urine concentration test
- Urinary tract calculi. (stones)

UNIT VIII

Liver function tests: Physiological role of liver, metabolic, storage, excretory,

protective, circulatory functions and function in blood coagulation.

- Test for hepatic dysfunction-Bile pigments metabolism.
- Test for hepatic function test- Serum bilirubin, urine bilirubin, and urineurobilinogen.
- Dye tests of excretory function.
- Tests based upon abnormalities of serum proteins.

UNIT IX

• **Lipid profile tests:** Lipoproteins, composition, functions. Determination of serum lipids, total cholesterol, HDL cholesterol, LDL cholesterol and triglycerides.

• **Immunochemical techniques** for determination of hormone levels and protein levels in serum for endocrine diseases and infectious diseases. Radio immuno assay (RIA) and Enzyme Linked Immuno Sorbent Assay (ELISA)

UNIT X

Electrolytes: Body water, compartments, water balance, and electrolyte distribution.

Determination of sodium, calcium potassium, chlorides, bicarbonates in the bodyfluids.

Suggested Readings:

- 1. Harpers Review of Biochemistry Martin
- 2. Text book of Biochemistry D. Satyanarayana
- 3. Text book of Clinical Chemistry- Alex Kaplan & Laverve L.Szabo

- 1. Principles of Biochemistry -- Lehninger
- 2. Text book of Biochemistry -- Ramarao
- 3. Practical Biochemistry-David T.Plummer

Pharm.D

24PD104 T

PHARMACEUTICAL ORGANIC CHEMISTRY - THEORY 4H 4C

Instruction hours/ week: L: 3 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 4 Hrs. /Week

Course Objectives: Students can able to

- Master organic chemistry fundamentals, covering polarity, solubility, acids and bases, and isomerism.
- Learn organic compound nomenclature and classification, including alkanes, alkenes, alcohols, and functional groups.
- Understand reaction mechanisms like nucleophilic substitution, elimination, and electrophilic addition.
- Explore resonance, allylic rearrangements, and electrophilic aromatic substitution.
- Gain proficiency in oxidation-reduction and nucleophilic aromatic substitution reactions.
- Study official compounds, focusing on preparation, purity tests, assay, and medicinal uses.

COs	Course Outcomes	Blooms Level
CO1	Develop a strong foundation in organic chemistry principles, enabling	Understand
	accurate analysis of chemical structures and reactions.	
CO2	Acquire proficiency in naming and categorizing organic compounds,	Analyse
	emaileng communication and comprehension within the field.	
CO3	Demonstrate understanding of reaction mechanisms, facilitating design	Understand
	of synthetic pathways and prediction of chemical transformations.	
CO4	Gain insight into advanced concepts like resonance and substitution	Apply
	reactions, enriching problem-solving skills.	
CO5	Apply knowledge to analyze and predict outcomes of chemical	Understand
	reactions, contributing to synthesis of novel compounds.	
CO6	Understand Practical applications in pharmaceutical science, improving	Understand
	ability to synthesize and analyze medicinal compounds.	

Course Outcomes: At the end of this course, students will be able to

Mapping with Programme Outcomes

Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12
CO1	Μ										Μ	
CO2	Μ										Μ	
CO3	Μ										Μ	
CO4	Μ										Μ	
CO5	Μ										Μ	
CO6	Μ										Μ	

S-Strong; M-Medium; L-Low

2024-25

FIRST YEAR

Course Content: UNIT I Structures and Physical properties:

- Polarity of bonds, polarity of molecules, M.P. Inter molecular forces, B.P.
- Solubility, non-ionic solutes and ionic solutes, protic and aprotic Solvents, ion pairs,
- Acids and bases, Lowry bronsted and Lewis theories
- Isomerism.

Nomenclature of organic compound belonging to the following classes Alkanes, Alkenes, Dienes, Alkynes, Alcohols, Aldehydes, Ketones, Amides, Amines, Phenols, Alkyl Halides, Carboxylic Acid, Esters, Acid Chlorides And cycloalkanes. Free radicals chain reactions of alkane: Mechanism, relative reactivity and stability

UNIT II

- Alicyclic compounds: Preparations of cyclo alkanes, Bayer strain Theory and orbitalpicture of angle strain.
- Nucleophilic aliphatic substitution mechanism: Nucleophiles and leaving groups, kinetics of second and first order reaction, mechanism and kinetics of SN₂ reactions. Stereochemistry and steric hindrance, role of solvents, phase transfer catalysis, mechanism and kinetics of SN1 reactions, stereochemistry, carbocation and their stability, rearrangement of carbocation, role of solvents in SN1 reaction, Ion dipole bonds, SN2versus SN1 solvolyses, nucleophilic assistance by the solvents.

UNIT III

Dehydro halogenation of alkyl halides: 1,2 elimination, kinetics, E2 and E1 mechanism, elimination via carbocation, evidence for E2 mechanism, absence of rearrangement isotope effect, absence hydrogen exchange, the element effect, orientationand reactivity, E2 versus E1, elimination versus substitution, dehydration of alcohol, ease of dehydration, acid catalysis, reversibility, orientation.

UNIT IV

Electrophillic and free radicals addition: Reactions at carbon-carbon, double bond, electrophile, hydrogenation, Heat of hydrogenation and stability of alkenes, Mark ownik off rule, addition of hydrogen halides, addition of hydrogen bromides, peroxide effect, electrophillic addition, mechanism, rearrangement, absence of hydrogen exchange, orientation and reactivity, addition of halogen, mechanism,

Halohydin formation, mechanism of free radicals additon, Mechanism of peroxide initiated addition of hydrogen bromide, orientation of free addition, Additions of carbene to alkene, cyclo addition reactions.

UNIT V

• **Carbon-carbon double bond as substituents:** Free radical halogenations of alkenes, comparision of free radical substitution with free radical addition, free radical substitution in alkenes, orientation and reactivity, allylic rearrangements.

• **Theory of resonance:** Allyl radical as a resonance hybrid, stability, orbital picture, resonance stabilisation of allyl radicals, hyper conjugation, allyl cation as a resonance hybrid, nucleophyllic substitution in allylic substrate, SN1 reactivity, allylic rearrangement, resonance stabilisation of allyl cation, hyper conjugation, nucleophilic substitution in allylic substrate, SN2 nucleophilic substituion in vinylic substrate, vinylic cation, stability of conjugated dienes, resonance in alkenes, hyper conjugation, ease of formation of conjugated dienes, orientation of elimination, electrophilic addition to conjugated dienes, 1,4- addition, 1,2-versus 1,4-addition, rate versus equilibrium, orientation and reactivity of free radical addition to conjugated dienes.

UNIT VI

Elecrophilic aromatic substitution: Effect of substituent groups, determination of orientation, determination of relative reactivity, classification of substituent group, mechanism of nitration, sulphonation, halogenation, friedel craft alkylation, friedel craftacylation, reactivity and orientation, activating and deactivating O,P,M directing groups, electron release via resonance, effect of halogen on electrophilic aromatic substitution inalkyl benzene, side chain halogination of alkyl benzene, resonance stabilization of benzylradical.

UNIT VII

Nucleophilic addition reaction: Mechanism, ionisation of carboxylic acids, acidity constants, acidity of acids, structure of carboxylate ions, effect of substituent on acidity, nucleophilic acyl substitution reaction, conversion of acid to acid chloride, esters, amide and anhydride. Role of caboxyl group, comparison of alkyl nucleophilic substitution with acyl nucleophilic substitution. Mechanism of aldol condensation, claisen condensation, cannizzaro reaction, crossed aldol condensation, crossed cannizzaro reaction, benzoin condensation, perkin condensation. Knoevenagel, Reformatsky reaction, Wittig reaction, Michael addition.

UNIT VIII

• **Hoffman rearrangement:** Migration to electron deficient nitrogen, Sandmeyer's reaction, basicity of amines, diazotisation and coupling, acidity of phenols, Williamson synthesis, Fries rearrangement, Kolbe reaction, Reimer tieman's reactions.

UNIT IX

- Nucleophilic aromatic substitution: Bimolecular displacement mechanisms, orientation, comparison of aliphatic nucleophilic substitution with that of aromatic.
- Oxidation reduction reaction.

UNIT X

 Study of the following official compounds- preparation, test for purity, assay and medicinal uses of Chlorbutol, Dimercaprol, Glyceryl trinitrate, Urea, Ethylene diamine dihyrate, Vanillin, Paraldehyde, Ethylene chloride, Lactic acid, Tartaric acid, citric acid, salicylic acid, aspirin, methyl salicylate, ethyl benzoate, benzyl benzoate, dimethyl pthalate, sodium lauryl sulphate, saccharin sodium, mephensin.

Suggested Readings:

- 1. T.R.Morrison and R. Boyd Organic chemistry
- 2. Bentley and Driver-Text book of Pharmaceutical chemistry
- 3. I.L.Finar- Organic chemistry, the fundamentals of chemistry

- 1. Organic chemistry J.M.Cram and D.J.Cram
- 2. Organic chemistry- Brown
- 3. Advanced organic chemistry- Jerry March, Wiley
- 4. Organic chemistry- Cram and Hammered, Pine Hendrickson

FIRST YEAR

PHARMACEUTICAL INORGANIC CHEMISTRY - THEORY 3H 3C

Instruction hours/ week: L: 2 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 3 Hrs. /Week

Course Objectives: Students can able to

- Understand the fundamentals of analytical chemistry and monographs of inorganic compounds.
- Understand the fundamentals of analytical chemistry and principles of electrochemical analysis of drugs
- Understand the basic concepts and Pharmacopoeial standards of pharmaceutical analysis
- Understand the principles of volumetric analysis and explore advanced analytical techniques for the detection and quantification of drug compounds
- Understand the real-world applications and challenges of utilizing inorganic pharmaceuticals in disease prevention and treatment strategies.

COs	Course Outcomes	Blooms Level
CO1	Understand the principles and procedures of analysis of	Understand
	drugs and also regarding the application of inorganic	
	pharmaceuticals	
CO2	Know the analysis of the inorganic pharmaceuticals their	Apply
	applications	
CO3	Appreciate the importance of inorganic pharmaceuticals	Apply
	in preventing and curing the disease.	
CO4	Identify the errors in analysis	Apply
CO5	Understand the principles of volumetric analysis and	Understand
	electrochemical analysis	
CO6	Applications of volumetric analysis.	Apply

Course Outcome (CO's): At the end of this course, students will be able to

Mapping with Programme Outcomes

Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1	Μ										
CO2	Μ										
CO3	Μ										
CO4	Μ										
CO5	Μ										
CO6	Μ										

S-Strong; M-Medium; L-Low

Course Content:

UNIT I

• Errors

UNIT II

• Volumetric analysis

UNIT III

- Acid-base titrations
- Redox titrations

UNIT IV

- Non aqueous titrations
- Precipitation titrations
- Complexometric titrations

UNIT V

• Theory of indicators

UNIT VI

• Gravimetry

UNIT VII

• Limit tests

UNIT VIII

• Medicinal gases

UNIT IX

- Acidifiers
- Antacids
- Cathartics
- Electrolyte replenishers
- Essential Trace elements
- Antimicrobials
- Pharmaceutical aids
- Dental Products
- Miscellaneous compounds

UNIT X

Radio Pharmaceuticals

Suggested Readings:

- 1. A text book Inorganic medicinal chemistry by Surendra N. Pandeya
- 2. A. H. Beckett and J. B. Stanlake's Practical Pharmaceutical chemistry Vol -I & Vol-II
- 3. Inorganic Pharmaceutical Chemistry III-Edition P. Gundu Rao

- 1. Inorganic Pharmaceutical Chemistry by Anand & Chetwal
- 2. Pharmaceutical Inorganic chemistry by Dr.B.G.Nagavi
- 3. Analytical Chemistry Principles by John H. Kennedy
- 4. I.P.1985 and 1996, Govt. of India, Ministry of Health

Pharm.D

24PD106RMT

REMEDIAL MATHEMATICS - THEORY

Instruction hours/ week: L: 3 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 4 Hrs. /Week

Course Objectives: Students can able to

- Understand the Trigonometry, Analytical geometry, Matrices, Determinant, Integration, Differential equation, Laplace transform and their applications.
- Solve the problems of different types by applying Theory.
- Understand important applications of mathematics in pharmacy.
- Proficiently use diverse mathematical tools for solving complex pharmaceutical problems.
- Analyze the mathematical theories to analyze pharmaceutical data, fostering critical thinking.
- Recognize mathematics' role in pharmaceutical applications, aiding understanding and decision-making in pharmacy

Course Outcome (CO's): On successful completion of the course the student will

COs	Course Outcomes	Blooms Level
CO1	Understand and perform the partial fraction, logarithms, function and limits, matrices and determinants differential and integral	Understand
	calculus	
CO2	Apply Theory to solve diverse problems.	Knowledge
CO3	Appreciate and analyze the math's importance in pharmacy.	Apply
CO4	Use math proficiently for complex pharmaceutical problem- solving.	Apply
CO5	Apply math theories for critical analysis of pharmaceutical data.	Apply
CO6	Appreciate the important application of mathematics in Pharmacy.	Apply

Mapping with Programme Outcomes

Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12
CO1	Μ											
CO2			Μ									
CO3			Μ									
CO4				Μ								
CO5			Μ									
CO6								Μ				

S- Strong; M-Medium; L-Low

2024-25

4H

FIRST YEAR

4C

Course Content:

UNIT I

Algebra: Determinants, Matrices

UNIT II

Trigonometry: Sides and angles of a triangle, solution of triangles

UNIT III

Analytical Geometry: Points, Straight line, circle, parabola

UNIT IV

Differential calculus: Limit of a function, Differential calculus, Differentiation of a sum, Product, Quotient Composite, Parametric, exponential, trigonometric and Logarithmic function.

UNIT V

Differential calculus: Successive differentiation, Leibnitz's theorem, Partial differentiation, Euler's theorem on homogeneous functions of two variables

UNIT VI

Integral Calculus: Definite integrals, integration by substitution and by parts,

Properties of definite integrals.

UNIT VII

Differential equations: Definition, order, degree, variable separable,homogeneous, Linear, heterogeneous, linear.

UNIT VIII

Differential equations: Differential equation with constant coefficient, simultaneous linear equation of second order.

UNIT IX

Laplace transform: Definition, Laplace transform of elementary functions

UNIT X

Laplace transform: Properties of linearity and shifting.

Suggested Readings:

- 1. Differential calculus By Shantinarayan
- 2. Text book of Mathematics for second year pre-university by Prof.B.M.Sreenivas

- 1. Integral calculus By Shanthinarayan
- 2. Engineering mathematics By B.S.Grewal
- 3. Trigonometry Part-I By S.L.Loney

Pharm.D

24PD106RBT

Course Objectives: Students can able to

- Understand the components of living world, structure and functional system of plant and animal kingdom.
- Know the classification and salient features of five kingdoms of life and morphology of flowering plants.

REMEDIAL BIOLOGY - THEORY

- Identify and differentiate various natural drugs to build a foundational understanding of Pharmacognosy.
- Know the significance of natural sources in pharmaceuticals, considering their diversity and potential therapeutic applications.
- Understand the photosynthesis and plant and mineral nutrition.
- Develop awareness of the ecological, ethical, and cultural aspects associated with the utilization of natural sources in pharmacy

Cos	Course Outcomes	Blooms Level
CO1	Explain the classification and salient feature s of five kingdoms of	Understand
	life and morphology of flowering plants.	
CO2	Develop awareness of the ecological, ethical, and cultural aspects associated with the utilization of natural sources in pharmacy.	Understand
CO3	Develop the ability to identify and distinguish various natural drugs, forming a foundational knowledge base in Pharmacognosy.	Knowledge
CO4	Analyze the significance of natural sources in pharmaceuticals, considering their diverse range and potential therapeutic benefits.	Understand
CO5	Evaluate the roles of plants and animals in providing in traditional and modern medicine.	Knowledge
CO6	Foster awareness of the ecological, ethical, and cultural dimensions associated with utilizing natural sources in pharmacy practice.	Knowledge

Course Outcomes (CO's): On successful completion of the course the student will

Instruction hours/ week: L: 3 T:1 P:0

Theory: 4 Hrs. /Week

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

4H

FIRST YEAR

4C

Mapping with Programme Outcomes

Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	S										
CO2	S										
CO3	S										
CO4	S										
CO5	S										
CO6	S										

S-Strong; M-Medium; L-Low

Course Content

UNIT I

- Introduction
- General organization of plants and its inclusions

UNIT II

- Plant tissues
- Plant kingdom and its classification
- Morphology of plants

UNIT III

- Root, Stem, Leaf and its modifications
- Inflorescence and Pollination of flowers
- Morphology of fruits and seeds

UNIT IV

• Plant physiology and Taxonomy of Leguminosae, Umbelliferae, Solanaceae, Liliaceae, Zinziberaceae, Rubiaceae

UNIT V

• Study of Fungi, Yeast, Penicillin and Bacteria

UNIT VI

- Study of Animal cell
- Study animal tissues

UNIT VII

• Detailed study of frog

UNIT VIII

• Study of Pisces, Raptiles, Aves

UNIT IX

• Genearal organization of mammals

UNIT X

• Study of poisonous animal

Suggested Readings:

- 1. Text book of Biology by S.B.Gokhale
- 2. A Text book of Biology by Dr. Thulajappa and Dr. Seetaram.

- 1. A Text book of Biology by B.V.Sreenivasa aidu
- 2. A Text book of Biology by Naidu and Murthy
- 3. Botany for Degree students By A.C.Dutta.
- 4. Outlines of Zoology by M.Ekambaranatha ayyer and T.N.Ananthakrishnan.
- 5. A manual for pharmaceutical biology Practical by S.B.Gokhale and C.K.Kokate.

24PD107P

HUMAN ANATOMY AND PHYSIOLOGY- PRACTICAL 3H 2C

Instruction hours/ week: L: 0 T:0 P:3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Practical : 3 Hrs/Week

Course Objectives: Students can able to

- Practical physiology is complimentary to the theoretical discussions in physiology.
- Practicals allow the verification of physiological processes discussed in Theory through experiments on living tissue, intact animals or normal human beings.
- To Identify epithelial, connective tissue, muscular, nervous tissues Microscopically and the axial, appendicular bones.
- Know about the bleeding time and clotting time.
- Know to Record the heart rate, pulse rate, blood pressure.
- Understand the WBC count and RBC count

Course Outcomes (CO's): On successful completion of the course the student will

	Course Outcomes	Blooms Level
COs		
CO1	Identify different types of tissues under microscope	Knowledge
CO2	Identify different types of bones	Knowledge
CO3	Identify different types of organs	Knowledge
CO4	Demonstration of blood analysis.	Apply
CO5	Demonstration of BP/heart rate/pulse rate	Apply
CO6	Explain different family planning techniques.	Understand

Mapping with Programme Outcomes

<u> </u>											
POs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
COs											
CO1	S		S	S							
CO2	S		S								
CO3	S		S								
CO4	S		S	S							
CO5	S										
CO6	S										

S-Strong; M-Medium; L-Low

General Requirements: Dissection box, Laboratory Napkin, muslin cloth, record, Observation book (100pages), Stationary items, Blood lancet.

List of Experiments:

- 1. Study of tissues of human body (a) Epithelial tissue
 - (b) Muscular tissue

FIRST YEAR

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- 2. Study of tissues of human body(a) Connective tissue(b) Nervous tissue
- 3. Study of appliances used in hematological experiments
- 4. Determination of W.B.C. count of blood
- 5. Determination of R.B.C. count of blood
- 6. Determination of differential count of blood
- 7. Determination of(a) Erythrocyte Sedimentation Rate(b) Hemoglobin content of Blood
 - (c) Bleeding time & Clotting time
- 8. Determination of
 - (a) Blood Pressure
 - (b) Blood group
- 9. Study of various systems with the help of charts, models & specimens
 - (a) Skeleton system part I-axial skeleton
 - (b) Skeleton system part II- appendicular skeleton
 - (c) Cardiovascular system
 - (d) Respiratory system
 - (e) Digestive system
 - (f) Urinary system
 - (g) Nervous system
 - (h) Special senses
 - (i) Reproductive system
- 10. Study of different family planning appliances.
- 11. To perform pregnancy diagnosis test.
- 12. Study of appliances used in experimental physiology.
- 13. To record simple muscle curve using gastroenemius sciatic nerve preparation.
- 14. To record simple summation curve using gastroenemius sciatic nerve preparation.
- 15. To record simple effect of temperature using gastroenemius sciatic nerve preparation.
- 16. To record simple effect of load & after load using gastroenemius sciatic nerve preparation.

17. To record simple fatigue curve using gastroenemius sciatic nerve preparation. **Suggested Readings:**

1. Goyal, R. K, Natvar M.P, and Shah S.A, Practical Anatomy, Physiology and Biochemistry, Latest Edition, Publisher: B.S Shah Prakashan, Ahmedabad.

Reference books

- 2. Ranade VG, Text Book of Practical Physiology, Latest Edition, Publisher: PVG,Pune.
- 3. Anderson Experimental Physiology, Latest Edition, Publisher: NA

24PD108P

PHARMACEUTICS- PRACTICAL

Instruction hours/ week: L: 0 T:0 P:3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Practical : 3 Hrs./Week

Course Objectives: Students can able to

- Know various internal liquid dosage forms preparations.
- Know various external liquid dosage forms.
- Prepare various solid dosage forms.
- Perform quality control tests for various dosage forms
- Know various formulations for body cavities.
- Pharmaceutical calculations in pharmacy.

Course Outcomes (CO's): At the end of this course, students will be able to

COs	Course Outcomes	Blooms
		Level
CO1	Prepare various internal liquid dosage forms	Analyze
CO2	Prepare various solid dosage forms	Analyze
CO3	Perform quality control tests for various dosage forms	Analyze
CO4	Prepare various formulations for body cavities	Analyze
CO5	Develop a clear idea about Pharmaceutical incompatibility and	Apply
	different pharmaceuticalcalculations in pharmacy.	

Mapping with Programme Outcomes

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1	S	Μ	Μ								
CO2	S	Μ	Μ	Μ					L		
CO3			S								
CO4	S	Μ		Μ					Μ		
CO5	S	Μ	S	Μ							

S-Strong; M-Medium; L-Low

List of Experiments:

Syrups

- Simple Syrup I.P
- Syrup of Ephedrine Hcl NF
- Syrup Vasaka IP
- Syrup of ferrous Phosphate IP
- Orange Syrup

Elixir Linctus

Solutions

- Piperizine citrate elixir BP
- Cascara elixir BPC
- Paracetamol elixir BPC

FIRST YEAR

2C

3H

- Simple Linctus BPC
- Pediatric simple Linctus BPC
- Solution of cresol with soap IP
- Strong solution of ferric chloride BPC
- Aqueous Iodine Solution IP
- Strong solution of Iodine IP
- Strong solution of ammonium acetate IP

Liniments

- Liniment of turpentine IP*
- Liniment of camphor IP

Suspensions*

- Calamine lotion
- Magnesium Hydroxide mixture BP

Emulsions*

- Cod liver oil emulsion
- Liquid paraffin emulsion

$\mathbf{Powders}^{\#}$

- Eutectic powder
- Explosive powder
- Dusting powder
- Insufflations

Suppositories[#]

- Boric acid suppositories
- Chloral suppositories

Incompatibilities

- Mixtures with Physical
- Chemical & Therapeutic incompatibilities
- * Colorless bottles required for dispensing
- [#] Paper envelope (white), butter paper and white paper required for dispensing.

Suggested Readings:

- 1. Cooper and Gunn's Dispensing for pharmacy students.
- 2. A text book Professional Pharmacy by N.K.Jain and S.N.Sharma.

- 1. Introduction to Pharmaceutical dosage forms by Howard C. Ansel.
- 2. Remington's Pharmaceutical Sciences.Register of General Pharmacy by Cooper and Gunn.
- 3. General Pharmacy by M.L.Schroff.

FIRST YEAR

MEDICINAL BIOCHEMISTRY -PRACTICAL 3H 2C

Instruction hours/ week: L: 0 T:0 P:3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Practical : 3 Hrs./Week

Course Objectives: Students can able to

- Know the Qualitative analysis of the biomolecules.
- Quantitatively analyze biochemical parameters and their importance indiagnosis of disease.
- Understand how to analyses the urine for abnormal constituents.
- Understand how to identify the biomolecules using chemical tests.
- Determine the enzymatic activity.
- Study the effect of physical parameters on the enzymatic activity.

Course Outcomes: At the end of this course, students will be able to

COs	Course Outcomes	Blooms Level
CO1	To perform the qualitatively analysis of biomolecules	Apply
CO2	To perform the Quantitatively analysis of biomolecules in	Apply
	blood, urine and serum	
CO3	Quantitative analysis of biomolecules by colorimetric method.	Apply
CO4	Quantitative analysis of biomolecules by titrimetric method.	Apply
CO5	Determine the enzymatic activity.	Apply
CO6	Study the effect of physical parameters on the enzymatic activity.	Apply

Mapping with Programme Outcomes

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12
CO1		S									Μ	
CO2		S									Μ	
CO3		S									Μ	
CO4		S									Μ	
CO5		S									Μ	
CO6		S									Μ	

S-Strong; M-Medium; L-Low

List of Experiments:

- 1 Qualitative analysis of normal constituents of urine.*
- 2 Qualitative analysis of abnormal constituents of urine.*
- 3 Quantitative estimation of urine sugar by Benedict's reagent method.**
- 4 Quantitative estimation of urine chlorides by Volhard's method.**
- 5 Quantitative estimation of urine creatinine by Jaffe's method.**
- 6 Quantitative estimation of urine calcium by precipitation method.**
- 7 Quantitative estimation of serum cholesterol by Libermann Burchard's method.**

- 8 Preparation of Folin Wu filtrate from blood.*
- 9 Quantitative estimation of blood creatinine.**
- 10 Quantitative estimation of blood sugar Folin-Wu tube method.**
- 11 Estimation of SGOT in serum.**
- 12 Estimation of SGPT in serum.**
- 13 Estimation of Urea in Serum.**
- 14 Estimation of Proteins in Serum.**
- 15 Determination of serum bilirubin**
- 16 Determination of Glucose by means of Glucoseoxidase.**
- 17 Enzymatic hydrolysis of Glycogen/Starch by Amylases.**
- 18 Study of factors affecting Enzyme activity. (pH & Temp.)**
- 19 Preparation of standard buffer solutions and its pH measurements (any two)*
- 20 Experiment on lipid profile tests**
- 21 Determination of sodium, calcium and potassium in serum.**
- ** Indicate major experiments & * Indicate minor experiments

Assignments: Format of the Assignment

- 1. Minimum & Maximum number of pages.
- 2. It shall be computer draft copy.
- 3. Reference(s) shall be included at the end.
- 4. Name and signature of the student.
- 5. Assignment can be a combined presentation at the end of the academic year.
- 6. Time allocated for presentation may be 8+2 Min.

Suggested Readings:

- 1. Harpers Review of Biochemistry Martin
- 2. Text book of Biochemistry D. Satyanarayana
- 3. Text book of Clinical Chemistry- Alex Kaplan & Laverve L.Szabo

- 1. Practical Biochemistry-David T.Plummer
- 2. Practical Biochemistry-Pattabhiraman

FIRST YEAR

PHARMACEUTICAL ORGANIC CHEMISTRY - PRACTICAL 3H 2C

Instruction hours/ week: L: 0 T:0 P: 3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Practical : 3 Hrs./Week

Course Objectives: Students can able to

- Understand the qualitative analysis of unknown organic compounds.
- Understand the special elements in an organic sample.
- Understand and confirm unknown compounds by m.p./b.p.
- Prepare derivatives of organic compounds.
- To know how to prepare the solid derivatives from organic compounds.
- Construct molecular models.

Course Outcomes (CO's): On successful completion of the course the student will

COs	Course Outcomes	Blooms Level
CO1	Synthesis of compounds using acetylation, Benzoylation and	Apply
	coupling reactions.	
CO2	Synthesis of compounds using Bromination, Nitration and	Apply
	condensation reactions.	
CO3	Synthesis of compounds using Oxidation, reduction and	Apply
	Hydrolysis reactions.	
CO4	Systematically perform qualitative analysis of unknown organic	Apply
	compounds.	
CO5	Prepare derivatives of organic compounds.	Apply
CO6	Construct molecular models.	Apply

Mapping with Programme Outcomes

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12
CO1		S										
CO2		S										
CO3		Μ										
CO4		Μ										
CO5		S										
CO6		S										

S- Strong; M-Medium; L-Low

Introduction to the various laboratory techniques through demonstration involving synthesis of the following compounds (at least 8 compounds to be synthesised):

- 1. Acetanilde / aspirin (Acetylation)
- 2. Benzanilide / Phenyl benzoate (Benzoylation)
- 3. P-bromo acetanilide / 2,4,6 tribromo aniline (Bromination)

- 4. Dibenzylidene acetone (Condensation)
- 5. 1-Phenylazo-2-napthol (Diazotisation and coupling)
- 6. Benzoic acid / salicylic acid (Hydrolysis of ester)
- 7. M-dinitro benzene (Nitration)
- 8. 9, 10 Antharaquinone (Oxidation of anthracene) / preparation of benzoic acid from toluene or benzaldehyde
- 9. M-phenylene diamine (Reduction of M-dinitrobenzene) / Aniline from nitrobenzene
- 10. Benzophenone oxime
- 11. Nitration of salicylic acid
- 12. Preparation of picric acid
- 13. Preparation of O-chlorobenzoic acid from O-chlorotolune
- 14. Preparation of cyclohexanone from cyclohexanol

Identification of organic compounds belonging to the following classes by :

Systematic qualitative organic analysis including preparation of derivatives Phenols, amides, carbohydrates, amines, carboxylic acids, aldehyde and ketones, Alcohols, esters, hydrocarbons, anilides, nitrocompounds.

Introduction to the use of stereo models:

Methane, Ethane, Ethylene, Acetylene, Cis alkene, Trans alkene, inversion of configuration.

Suggested Readings:

- 1. T.R.Morrison and R. Boyd Organic chemistry
- 2. Bentley and Driver-Text book of Pharmaceutical chemistry
- 3. I.L.Finar- Organic chemistry, the fundamentals of chemistry

- 1. Organic chemistry J.M.Cram and D.J.Cram
- 2. Organic chemistry- Brown
- 3. Advanced organic chemistry- Jerry March, Wiley
- 4. Organic chemistry- Cram and Hammered, Pine Hendrickson

24PD111P

FIRST YEAR

PHARMACEUTICAL INORGANIC CHEMISTRY -PRACTICAL 3H 2C

Instruction hours/ week: L: 0 T:0 P:3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Practical : 3 Hrs./Week

Course Objectives: students can able to

- Understand the limit test for samples.
- Understand how to prepare the solutions for volumetric and electro-analytical methods.
- Standardize the solutions by volumetric and electro-analytical methods.
- Know how to Perform the assay for chemical substances.
- Standardize the titrant used for the assay.
- Determine the strength of the solutions by electro-analytical methods.

Course Outcomes: At the end of this course, students will be able to

COs	Course Outcomes	Blooms Level
CO1	Analyze the limit test for samples.	Understand
CO2	Prepare the solutions for volumetric and electro-analytical methods.	Apply
CO3	Standardize the solutions by volumetric and electro-analytical methods.	Apply
CO4	Perform the assay for chemical substances.	Apply
CO5	Standardize the titrant used for the assay.	Apply
CO6	Determine the strength of the solutions by electro-analytical methods.	Understand

Mapping with Programme Outcomes

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12
CO1		Μ										
CO2		Μ										
CO3		Μ										
CO4		L										
CO5		L										
CO6		L										

S-Strong; M-Medium; L-Low

List of expriments: Limit test (6 exercises)

- a. Limit test for chlorides
- b. Limit test for sulphates
- c. Limit test for iron
- d. Limit test for heavy metals

- e. Limit test for arsenic
- f. Modified limit tests for chlorides and sulphates

Assays (10 exercises)

- g. Ammonium chloride- Acid-base titration
- h. Ferrous sulphate- Cerimetry
- i. Copper sulpahte- Iodometry
- j. Calcilugluconate- Complexometry
- k. Hydrogen peroxide Permanganometry
- 1. Sodium benzoate Nonaqueous titration
- m. Sodium chloride Modified volhard's method
- n. Assay of KI KIO3 titration
- o. Gravimetric estimation of barium as barium sulphate
- p. Sodium antimony gluconate or antimony potassium tartarate

Estimation of mixture (Any two exercises)

- q. Sodium hydroxide and sodium carbonate
- r. Boric acid and Borax
- s. Oxalic acid and sodium oxalate

Test for identity (Any three exercises)

- t. Sodium bicorbonate
- u. Barium sulphate
- v. Ferrous sulphate
- w. Potassium chloride

Test for purity (Any two exercises)

- a. Swelling power in Bentonite
- b. Acid neutralising capacity in aluminium hydroxide gel
- c. Ammonium salts in potash alum
- d. Adsorption power heavy Kaolin
- e. Presence of Iodates in KI

Preparations (Any two exercises)

- f. Boric acids
- g. Potash alum
- h. Calcium lactate
- i. Magnesium suphate

Suggested Readings:

- 1. A text book Inorganic medicinal chemistry by Surendra N. Pandeya
- 2. Inorganic Pharmaceutical Chemistry III-Edition P. Gundu Rao

- 1. Pharmaceutical Inorganic chemistry by Dr.B.G.Nagavi
- 2. Analytical Chemistry Principles by John H. Kennedy
- 3. I.P.1985 and 1996, Govt. of India, Ministry of Health

24PD112RBP

2024-25

FIRST YEAR

REMEDIAL BIOLOGY - PRACTICAL 3H 2C

Instruction hours/ week: L: 0 T:0 P:3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Practical: 3 Hrs./Week

Course Objectives:

- Students will be able to understand the microscope, cutting sections, mount, stain and slide preparation.
- To know about cell and its organelles
- To understand the parts of plant and their modifications
- Able to know the system using software
- Able to Identify types of bones.
- To determine blood group, blood pressure and tidal volume.

Course Outcomes (CO's): On successful completion of the course the student will

Cos	Course Outcomes	Blooms Level
CO1	Understand the microscope, cutting sections, mount, stain and	Understand
	slide preparation.	
CO2	Study cell and its organelles	Knowledge
CO3	Study the parts of plant and their modifications	Understand
CO4	Study the system in from using software	Knowledge
CO5	Identify types of bones.	Knowledge
CO6	Determine blood group, blood pressure and tidal volume.	Analyze

Mapping with Programme Outcomes

Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	S										
CO2	S										
CO3	S										
CO4	S										
CO5	S										
CO6				Μ							

S-Strong; M-Medium; L-Low

Title:

- 1. Introduction of biology experiments
- 2. Study of cell wall constituents and cell inclusions
- 3. Study of Stem modifications
- 4. Study of Root modifications

- 5. Study of Leaf modifications
- 6. Identification of Fruits and seeds
- 7. Preparation of Permanent slides
- 8. T.S. of Senna, Cassia, Ephedra, Podophyllum
- 9. Simple plant physiological experiments
- 10. Identification of animals
- 11. Detailed study of Frog
- 12. Computer based tutorials

Suggested Readings:

- 1. Text book of Biology by S.B.Gokhale
- 2. A Text book of Biology by Dr. Thulajappa and Dr. Seetaram.

- 1. A Text book of Biology by B.V.Sreenivasa aidu
- 2. A Text book of Biology by Naidu and Murthy
- 3. Botany for Degree students By A.C.Dutta.
- 4. Outlines of Zoology by M.Ekambaranatha ayyer and T.N.Ananthakrishnan.
- 5. A manual for pharmaceutical biology Practical by S.B.Gokhale and C.K.Kokate.

24PD201T

PATHOPHYSIOLOGY - THEORY

SECOND YEAR 4H 4C

Instruction hours/ week: L: 3 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 3 Hrs. /Week

Course Objectives:

- Pathophysiology is the study of causes of diseases and reactions of the body to such disease producing causes.
- This course is designed to impart a thorough knowledge of the relevant aspects of pathology of various conditions with reference to its pharmacological applications
- Understanding of basic pathophysiological mechanisms
- Hence it will not only help to study the syllabus of pathology,
- To get baseline knowledge required to practice medicine safely, confidently, rationally and effectively.
- To understand the etiology and pathogenesis of diseases.

Course Outcomes (CO's):

On successful completion of the course the student will

COs	Course Outcomes	Blooms Level
CO1	Describe the etiology and pathogenesis of the selected disease	Knowledge
	states;	
CO2	Name the signs and symptoms of the diseases; and	Knowledge
CO3	Mention the complications of the diseases.	Knowledge
CO4	Describe the mechanism of the diseases.	Knowledge
CO5	Understand the etiology and pathogenesis of diseases.	Understand
CO6	Discuss about the Sexually transmitted diseases	Understand

Mapping with Programme Outcomes

POs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
COs											
CO1	S										
CO2	S										
CO3	S										
CO4	S										
CO5	S										
CO6	S										

S-Strong; M-Medium; L-Low

UNIT I

Basic principles of cell injury and Adaptation

- Causes, Pathogenesis and morphology of cell injury
- Abnormalities in lipoproteinaemia, glycogen infiltration and glycogen

infiltration and glycogen infiltration and glycogen storage diseases

UNIT II

Inflammation

- Pathogenesis of acute inflammation, Chemical mediators ininflammation, Types of chronic inflammation
- Repairs of wounds in the skin, factors influencing healing of wounds

UNIT III

Diseases of Immunity

- Introduction to T and B cells
- MHC proteins or transplantation antigens

Immune tolerance

• Hypersensitivity

Hypersensitivity type I, II, III, IV, Biological significance, Allergy due to food, chemicals and drugs

• Autoimmunity

Criteria for autoimmunity, Classifications of autoimmune diseases in man, mechanism of autoimmunity, Transplantation and immunologic tolerance, allograft rejections, transplantation antigens, mechanism of rejection of allograft.

- Acquired immune deficiency syndrome (AIDS)
- Amylodosis

UNIT IV

Cancer: Differences between benign and malignant tumors, Histological diagnosis of malignancy, invasions and metastasis, patterns of spread,

- Disturbances of growth of cells, classification of tumors, general biology of tumors, spread of malignant tumors, etiology and pathogenesis of cancer.
- Types of shock, mechanisms, stages and management
- Biological effects of radiation

UNIT V

Environmental and nutritional diseases

- Air pollution and smoking- SO2,NO, NO2, and CO
- Protein calorie malnutrition, vitamins, obesity, pathogenesis of starvation.

UNIT VI

Pathophysiology of common diseases

Parkinsonism

Pharm.D

- Schizophrenia
- Depression and mania

UNIT VII

Pathophysiology of common diseases

- Hypertension,
- Stroke (ischaemic and hemorrhage)
- Angina, CCF, Atherosclerosis, Myocardial infarction

UNIT VIII

Pathophysiology of common diseases

- Diabetes Mellitus
- Peptic ulcer and inflammatory bowel diseases

UNIT IX

Pathophysiology of common diseases

- Cirrhosis and Alcoholic liver diseases
- Acute and chronic renal failure
- Asthma and chronic obstructive airway diseases

UNIT X

Infectious diseases :

Sexually transmitted diseases (HIV, Syphilis, Gonorrhea), Urinary tract infections, Pneumonia, Typhoid, Tuberculosis, Leprosy, Malaria Dysentery (bacterial and amoebic), Hepatitis- infective hepatitis.

Assignments :

Title of the Experiment

- 1 Chemical Mediators of inflammation
- 2 Drug Hypersensitivity
- 3 Cigarette smoking & its ill effects
- 4 Biological Effects of Radiation
- 5 Etiology and hazards of obesity
- 6 Complications of diabetes
- 7 Diagnosis of cancer
- 8 Disorders of vitamins
- 9 Methods in Pathology-Laboratory values of clinical significance
- 10 Pathophysiology of Dengue Hemorrhagic Fever (DHF)

Format of the assignment

- 1 Minimum & Maximum number of pages.
- 2. Reference(s) shall be included at the end.
- 3. Assignment can be a combined presentation at the end of the academic year
- 4. It shall be computer draft copy.
- 5. Name and signature of the student
- 6. Time allocated for presentation may be 8+2 Min.

Suggested Readings:

- 1. Pathologic basis of disease by- Cotran, Kumar, Robbins
- 2. Text book of Pathology- Harsh Mohan
- 3. Text book of Pathology- Y.M. Bhinde

Reference Books (Latest Editions):

1. Clinical Pharmacy and Therapeutics; Second edition; Roger Walker; Churchill Livingstone publication 24PD202T

2024-25

SECOND YEAR

PHARMACEUTICAL MICROBIOLOGY - THEORY4H4C

Instruction hours/ week: L: 3 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 4 Hrs. /WeekCourse

Objectives:

- Know the anatomy, identification, growth factors and sterilization of microorganisms;
- Know the mode of transmission of disease causing microorganism, symptoms of disease, and treatment aspect;
- Do estimation of RNA and DNA and there by identifying the source;
- Do cultivation and identification of the microorganisms in the laboratory;
- Do identification of diseases by performing the diagnostic tests; and
- Appreciate the behavior of motility and behavioral characteristics of microorganisms

Course Outcomes (CO's):

Upon completion of the subject student shall be able to

COs	Course Outcomes	Blooms	
		Level	
CO1	Know the anatomy, identification, growth factors and sterilization of	Understand	
	microorganisms.		
CO2	Know the mode of transmission of disease causing microorganism,	Understand	
	symptoms of disease, and treatment aspect.		
CO3	Do estimation of RNA and DNA and there by identifying the source.	Understand	
CO4	Do cultivation and identification of the microorganisms in the laboratory.	Understand	
CO5	Do identification of diseases by performing the diagnostic tests.	Apply	
CO6	Appreciate the behavior of motility and behavioral characteristics of	Knowledge	
	microorganisms.		

Mapping with Programme Outcomes

CO's	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1	S										
CO2	S										
CO3	S										
CO4	S										
CO5	S										
CO6	Μ										

S-Strong; M-Medium; L-Low
Course Content:

UNIT I

• Introduction to the science of microbiology. Major divisions of microbial world and Relationship among them.

UNIT II

• Different methods of classification of microbes and study of Bacteria, Fungi, virus, Rickettsiae, Spirochetes.

UNIT III

- Nutritional requirements, growth and cultivation of bacteria and virus.
- Study of different important media required for the growth of aerobic and anaerobic bacteria & fungi. Differential media, enriched media and selective media, maintenance of lab cultures.
- Different methods used in isolation and identification of bacteria with emphasis to different staining techniques and biochemical reactions. Counting of bacteria Total and Viable counting techniques.

UNIT IV

- Detailed study of different methods of sterilization including their merits and demerits. Sterilization methods for all pharmaceutical products. Detailed study of sterility testing of different pharmaceutical preparations.
- Brief information on Validation.

UNIT V

• **Disinfectants**- Study of disinfectants, antiseptics, fungicidal and virucidal agents factors affecting their activation and mechanism of action. Evaluation of bactericidal, bacteriostatic, virucidal activities, evaluation of preservatives in pharmaceutical preparations.

UNIT VI

• Immunology-Immunity, Definition, Classification, General principles of natural immunity, Phagocytosis, acquired immunity (active and passive)

UNIT VII

• Antigens, chemical nature of antigens structure and formation of Antibodies, Antigen-Antibody reactions. Bacterial exotoxins and endotoxins. Significance of toxoids in active immunity, Immunization programme, and importance of booster dose.

UNIT VIII

• Diagnostic tests : Schick's Test, Elisa test, Western Blot test, Southern Blot PCR Widal, QBC, Mantaux Peripheral smear. Study of malarial parasite.

UNIT IX

- Microbial culture sensitivity Testing: Interpretation of results Principles and methods of different microbiological assays, microbiological assay of
 - Penicillin, Streptomycin and vitamin B₂ and B₁₂.
 - Standardization of vaccines and sera.

UNIT X

• Study of infectious diseases: Typhoid, Tuberculosis, Malaria, Cholera, Hepatitis, Meningitis, Syphilis & Gonorrhea and HIV.

Reference Books (Latest Editions):

- Vanitha Kale and Kishor Bhusari Applied Microbiology || Himalaya Publishing houseMumbai.
- Mary Louis Turgeon Immunology and Serology in Laboratory Medicines 2nd edition, 1996 Mosby- Year book inc St. Louis Missouri 63146.
- Harsh Mohan, Text book of Pathology || 3rd edition, 1998, B-3 Ansari road Daryaganj N. Delhi.

Reference books (Theory)

- Prescot L.M., Jarley G.P Klein D.A Microbiology ¹2nd- edition Mc Graw Hill Company Inc
- Rawlins E.A. Bentley's Text Book of Pharmaceutics B ailliere Tindals 24-28 London 1988
- 3. Forbisher Fundamentals of Microbiology Philidelphia W.B. Saunders.
- Prescott L.M. Jarley G.P., Klein.D.A. Microbiology. ¹2nd edition WMC BrownPublishers, Oxford. 1993
- War Roitt, Jonathan Brostoff, David male, Immunology^{II}3rd edition 1996, Mosby-year book Europe Ltd, London.
- 6. Pharmacopoeia of India, Govt of India, 1996.

24PD203T

2024-25

SECOND YEAR

PHARMACOGNOSY AND PHYTOPHARMACEUTICALS -THEORY 4H 4C

Instruction hours/ week: L: 3 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 4 Hrs. /Week

Course Objectives:

- This subject has been introduced for the pharmacy course in order to make the student aware of medicinal uses of various naturally occurring drugs its history, sources, and distribution
- To know its method of cultivation,
- To know active constituents, medicinal uses
- To understand its preservation methods, identification tests, substitutes and adulterants.
- To understand the traditional system of medicine.
- To explain the Plant Products Primary metabolites Proteins, Enzymes, Lipids, Marine drugs.

Course Outcomes (CO's):

COs	Course Outcomes	Blooms
		Level
CO1	Explain the concepts and importance of Pharmacognosy.	Analyze
CO2	Illustrate the classification, Quality control of natural drugs & cultivation of	Apply
	crude drugs.	
CO3	Explain the applications of primary and secondary metabolites of the plant.	Analyze
CO4	Demonstrate the basic concepts of plant tissue culture.	Apply
CO5	Summarize the Traditional system of medicine.	Evaluate
CO6	Explain the importance of plant product sprimary metabolites Proteins,	Analyze
	Enzymes, Lipids, Marine drugs.	

Mapping with Programme Outcomes

CO's	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1	Μ										
CO2			Μ								
CO3						Μ					
CO4				Μ							
CO5						Μ					
CO6				Μ							

S-Strong; M-Medium; L-Low

UNIT I

- Introduction.
- Definition, history and scope of Pharmacognosy.

UNIT II

- Classification of crude drugs.
- Cultivation, collection, processing and storage of crude drugs.
- Detailed method of cultivation of crude drugs.

UNIT III

- Study of cell wall constituents and cell inclusions.
- Microscopical and powder Microscopical study of crude drugs.

UNIT IV

• Study of natural pesticides.

UNIT V

• Detailed study of various cell constituents.

UNIT VI

- Carbohydrates and related products.
- Detailed study carbohydrates containing drugs.(11 drugs)

UNIT VII

- Definition sources, method extraction, chemistry and method of analysis of lipids.
- Detailed study of oils.

UNIT VIII

• Definition, classification, chemistry and method of analysis of protein.

UNIT IX

• Study of plants fibers used in surgical dressings and related products.

UNIT X

• Different methods of adulteration of crude drugs.

Suggested Readings:

- 1. Pharmacognosy by G.E. Trease & W.C.Evans.
- 2. Pharmacognosy by C.K.Kokate,Gokhale & A.C.Purohit.

- 1. Pharmacognosy by Brady & Tyler.E.
- 2. Pharmacognosy by T.E.Wallis.
- 3. Pharmacognosy by C.S. Shah & Qadery.
- 4. Pharmacognosy by M.A. Iyengar.

24PD204T

SECOND YEAR

PHARMACOLOGY I-THEORY

Instruction hours/ week: L: 3 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

4H

Theory: 4 Hrs. /Week

Course Objectives:

- This subject will provide an opportunity for the student to learn about the drug with regard to classification, pharmacodynamic and pharmacokinetic aspects
- To know adverse effects, uses, dose
- To understand the route of administration, precautions, contraindications and interaction with other drugs.
- In this subject, apart from general pharmacology, drugs acting on autonomic nervous system, cardiovascular system, central nervous system
- To know the blood and blood forming agents and renal system will be taught.
- In addition to theoretical knowledge, the basic Practical knowledge relevant to therapeuticswill be imparted.

Course Outcome (CO's):

On successful completion of the course the student will

COs	Course Outcomes	Blooms Level
CO1	Understand the pharmacological aspects of	Understand
	drugs.	
CO2	Handle and carry out the animal experiments.	Apply
CO3	Appreciate the importance of pharmacology subject as a basis of	Knowledge
	therapeutics.	
CO4	Correlate and apply the knowledge therapeutically.	Analyze
CO5	Demonstrate the Pharmacology of drugs acting on various	Apply
	cardiovascular diseases.	
CO6	Illustrate the Pharmacology of Autocoids, Non-	Apply
	steroidalanti-inflammatory agents, Anti-goutdrugs	
	and Antirheumatic drugs	

Mapping with Programme Outcomes

POs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
COs											
CO1	S										
CO2	S	S	S				S				
CO3	S						S				
CO4	S		S								
CO5	S		S								
CO6	S										

S-Strong; M-Medium; L-Low

4C

UNIT I

General Pharmacology

- Introduction, definitions and scope of pharmacology
- Routes of administration of drugs
- Pharmacokinetics (absorption, distribution, metabolism and excretion)

UNIT II

- Pharmacodynamics
- Factors modifying drug effects
- Drug toxicity Acute, sub- acute and chronic toxicity.
- Pre-clinical evaluations
- Drug interactions

Note: The term Pharmacology used here refers to the classification, mechanism of action, pharmacokinetics, pharmacodynamics, adverse effects, contraindications, Therapeutic uses, interactions and dose and route of administration.

UNIT III Pharmacology of drugs acting on ANS

- Adrenergic and antiadrenergic drugs
- Cholinergic and anticholinergic drugs
- Neuromuscular blockers
- Mydriactics and miotics

UNIT IV

- Drugs used in myasthenia gravis
- Drugs used in Parkinsonism

UNIT V

Pharmacology of drugs acting on cardiovascular system

- Antihypertensives
- Anti-anginal drugs
- Anti-arrhythmic drugs
- Drugs used for therapy of Congestive Heart Failure
- Drugs used for hyperlipidaemias

UNIT VI

Pharmacology of drugs acting on Central Nervous System

- General anesthetics
- Sedatives and hypnotics

Anticonvulsants

• Analgesic and anti-inflammatory agents

UNIT VII

- Psychotropic drugs
- Alcohol and methyl alcohol
- CNS stimulants and cognition enhancers
- Pharmacology of local anaesthetics

UNIT VIII

Pharmacology of Drugs acting on Respiratory tract

- Bronchodilators
- Mucolytics
- Expectorants
- Antitussives
- Nasal Decongestants

UNIT IX

Pharmacology of Hormones and Hormone antagonists

- Thyroid and Antithyroid drugs
- Insulin, Insulin analogues and oral hypoglycemic agents
- Sex hormones and oral contraceptives
- Oxytocin and other stimulants and relaxants

UNIT X

Pharmacology of autocoids and their antagonists

- Histamines and Antihistaminics
- 5-Hydroxytryptamine and its antagonists
- Lipid derived autocoids and platelet activating factor
- The following details shall be submitted for any 25 selected drugs in the form of assignment/PPT.

**Name of the Drug, Brand name, Category, Therapeutic uses, Dose, Pharmacological action, Route of administration, Side effects, Contraindications, Drug Interactions with drug/food, Pharmacokinetics

Suggested Readings(Theory):

(Author, Title, Edition, Publication Place, Publisher, Year of Publication)

- Tripathi, K. D. Essentials of medical pharmacology. 4th Ed, 1999. Publisher: Jaypee, Delhi.
- 2. Satoskar, R.S. and Bhadarkar, S.D. Pharmacology and pharmacotherapeutics. 16th edition (single volume), 1999. Publisher: Popular, Dubai.
- Rang, H.P. & Dale, M.M. Pharmacology. 4th edition, 1999. Publisher: Churchill Living stone.

Reference Books (Theory-Latest Editions) :

(Author, Title, Edition, Publication Place, Publisher, Publication Year)

- Goodman Gilman, A., Rall, T.W., Nies, A.I.S. and Taylor, P. Goodman and Gilman's The pharmacological Basis of therapeutics. 9th Ed, 1996. Publisher Mc Graw Hill, Pergamon press.
- Craig, C.R.&Stitzel, R.E. Modern Pharmacology. Latest edition. Publisher: Little Brown.Co
- 3. Katzung, B.G. Basic and clinical pharmacology. Latest edition. Publisher: Prentice Hall, Int.
- 4. Shargel and Leon. Applied Biopharmaceutics and pharmacokinetics. Latest edition. Publisher: Prentice Hall, London.

24PD205T

COMMUNITY PHARMACY - THEORY

Instruction hours/ week: L:2 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 3 Hrs.

/WeekCourse

Objectives:

- In the changing scenario of pharmacy practice in India, Community Pharmacists are expected to offer various pharmaceutical care services.
- In order to meet this demand, students will be learning various skills such as dispensing of drugs
- To know responding to minor ailments by providing suitable safe medication
- To understand patient counseling
- To know health screening services for improved patient care in the community set up.
- To study the concept of Rational drug therapy

Course Outcome (CO's):

Upon completion of the course, the student shall be able to -

Cos	Course Outcomes	Blooms Level
CO1	Describe the rules and responsibilities of community pharmacy	Understanding
CO2	Explain community pharmacy management	Applying
CO3	Give an Outline of the concept of pharmaceutical care	Understanding
CO4	What is the importance of patient counselling and health screening	Remembering&
	services?	Applying
CO5	Discuss health education.	Creating
CO6	Explain the management of minor ailments	Applying

Mapping with Programme Outcomes

Cos	PO1	PO2	PO3	PO4	P05	PO6	PO7	PO8	PO9	PO10	PO11
CO1	S				S	S	S	L	S	L	
CO2	S	S	Μ	Μ			Μ	L	Μ	S	S
CO3	Μ				Μ	L		L		L	
CO4	Μ	Μ	Μ	S		S		L	Μ	Μ	S
CO5	М			М		S		S		Μ	S
CO6	М	S	Μ	S	Μ	Μ	Μ	Μ	Μ	S	S

S-Strong; M-Medium; L-Low

Special requirements:

 Either the college is having model community pharmacy (meeting the schedule N Requirement) or sign MoU with at least 4-5 community pharmacies nearby to the

college for training the students on dispensing and counselling activities.

2. Special equipments like B.P apparatus, Glucometer, Peak flow meter, and apparatus for cholesterol estimation.

2024-25

3C

SECOND YEAR

3H

UNIT I

Definition, scope, of community pharmacy

Roles and responsibilities of Community pharmacist

UNIT II

Community Pharmacy Management

- a) Selection of site, Space layout, and design
- b) Staff, Materials- coding, stocking
- c) Legal requirements
- d) Maintenance of various registers
- e) Use of Computers: Business and health care soft wares

UNIT III

Prescriptions – parts of prescription, legality & identification of medication related problems like drug interactions.

Inventory control in community pharmacy Definition,

various methods of Inventory Control

ABC, VED, EOQ, Lead time, safety stock

UNIT IV

Pharmaceutical care

Definition and Principles of Pharmaceutical care.

Patient counselling

Definition, outcomes, various stages, barriers, Strategies to overcome barriers

Patient information leaflets- content, design, & layouts, advisory labels

UNIT V

Patient medication adherence

Definition, Factors affecting medication adherence, role of Pharmacist in improving the adherence.

Health screening services

Definition, importance, methods for screening Blood pressure/ blood sugar/ lung function and Cholesterol testing

UNIT VI

OTC Medication- Definition, OTC medication list & Counselling

Health Education

WHO Definition of health, and health promotion, care for children, pregnant & breast feeding women, and geriatric patients.

UNIT VII

Introduction to Social and Preventive Pharmacy:

Commonly occurring Communicable Diseases, causative agents, Clinical presentations and prevention of Communicable diseases – Tuberculosis, Hepatitis, Typhoid, Amoebiasis, Malaria, Leprosy, Syphilis, Gonorrhea and AIDS, HIV & AIDS control programme, National leprosy control programme.

UNIT VIII

Balance diet, and treatment & prevention of deficiency disorders

Family planning – role of pharmacist

UNIT IX

Responding to symptoms of minor ailments Relevant pathophysiology, common drug therapy

to, Pain, GI disturbances (Nausea, Vomiting, Dyspepsia, diarrhea, constipation), Pyrexia,

Opthalmic symptoms, worms infestations.

UNIT X

Essential Drugs concept and Rational Drug Therapy Role of community pharmacist

Code of ethics for community pharmacists

Suggested Readings:

- 1. Health Education and Community Pharmacy by N.S.Parmar.
- 2. WHO consultative group report.
- 3. Drug store & Business management by Mohammed Ali & Jyoti.

- 1. Handbook of pharmacy health care.Edt. Robin J Harman. The Pharmaceutical press.
- Comprehensive Pharmacy Review Edt. Leon Shargel. Lippincott Williams & Wilkins.

Pharm.D		2024-25	_
24PD206T		SECOND YEAR	-
	PHARMACOTHERAPEUTICS I -THEORY	4H 4C	

Instruction hours/ week: L: 3 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 4 Hrs.

/WeekCourse

Objectives:

- This course is designed to impart knowledge and skills necessary for contribution to quality use of medicines.
- Chapters dealt cover briefly pathophysiology and mostly therapeutics of various diseases.
- This will enable the student to understand the pathophysiology of common diseases and their management.
- To know the importance of preparation of individualized therapeutic plans based on diagnosis;
- To understand patient-specific parameters relevant in initiating drug therapy
- To know the pathophysiology of selected disease states and explain the rationale for drug therapy

Course Outcome (CO's):

A successful completion of this subject it is expected that students will be able to

COs	Course Outcomes	Blooms Level
CO1	Understand the etiologic factors, signs and symptoms of	Understand
	selected diseases	
CO2	Explain pathophysiology of selected diseases.	Knowledge
CO3	Explain drug - drug interactions for selected drugs	Knowledge
CO4	Explain drug- food interactions for selected drugs	Knowledge
CO5	Illustrate the therapeutic approach of management of selected disease.	Apply
CO6	Understand the basic concepts of different guidelines for prescribing.	Apply

Mapping with Programme Outcomes

Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12
CO1	S											
CO2	S											
CO3					М			М				
CO4					М			М				
CO5						S			S			
CO6									М		S	

Pharm.D

Strong; M-Medium; L-Low

Identify the patient-specific parameters relevant in initiating drug therapy, and monitoring therapy (including alternatives, time-course of clinical and laboratory indices of therapeutic response and adverse effects).

Detailed syllabus and lecture wise schedule:

Etiopathogenesis and pharmacotherapy of diseases associated with following systems/

diseases

UNIT I

Cardiovascular system: Hypertension, Congestive cardiac failure, Angina Pectoris

UNIT II

Myocardial infarction, Hyperlipidaemias

UNIT III

Electrophysiology of heart and Arrhythmias

UNIT IV

Respiratory system : Introduction to Pulmonary function test, Asthma,

UNIT V

Chronic obstructive airways disease, Drug induced pulmonary diseases

UNIT VI

Endocrine system : Diabetes, Thyroid diseases, Oral contraceptives

UNIT VII

Hormone replacement therapy, Osteoporosis

UNIT VIII

General prescribing guidelines for

- a. Paediatric patients
- b. Geriatric patients
- c. Pregnancy and breast feeding

UNIT IX

Ophthalmology: Glaucoma, Conjunctivitis- viral & bacterial

UNIT X

Introduction to rational drug use

Definition, Role of pharmacist Essential drug concept Rational drug formulations

Suggested Readings:

- 1. Clinical Pharmacy and Therapeutics Roger and Walker, Churchill Livingstone publication.
- Pharmacotherapy: A Pathophysiologic approach Joseph T. Dipiro et al. Appleton & Lange.

- 1. Pathologic basis of disease Robins SL, W.B.Saunders publication.
- 2. Pathology and therapeutics for Pharmacists: A Basis for Clinical Pharmacy Practice - Green and Harris, Chapman and Hall publication.
- 3. Clinical Pharmacy and Therapeutics Eric T. Herfindal, Williams and Wilkins Publication.
- Applied Therapeutics: The clinical Use of Drugs. Lloyd Young and Koda-Kimble MA
- 5. Avery's Drug Treatment, 4th Edn, 1997, Adis International Limited.
- 6. Relevant review articles from recent medical and pharmaceutical literature

24PD207P

SECOND YEAR

PHARMACEUTICAL MICROBIOLOGY - PRACTICAL 3H 2C

Instruction hours/ week: L: 0 T:0 P:3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Practical : 3 Hrs./Week

Course Objectives:

- To discuss about the instruments used in experimental microbiology
- To understand the sterilization methods followed in laboratory.
- To know the staining techniques used in microbiology.
- To carry out assay of different antibiotics
- To understand the mechanism of action opf antibiotics.
- To perform different sterility tests and bacteriological analysis of water

Course Outcomes (CO's):

At the end of this course, students will be able to

COs	Course Outcomes	Blooms
		Level
CO1	Discuss about the instruments used in experimental microbiology	Understand
CO2	Understand the sterilization methods followed in laboratory.	Understand
CO3	Discover the staining techniques used in microbiology.	Understand
CO4	Carry out assay of different antibiotics	Apply
CO5	Understand the mechanism of action of antibiotics.	Understand
CO6	Execute different terility tests and bacteriological analysis of water	Knowledge

Mapping with Programme Outcomes

CO's	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1	S										
CO2	S										
CO3	S										
CO4	S										
CO5	S										
CO6	Μ										

S-Strong; M-Medium; L-Low

Title of the Experiment:

- 1 Study of apparatus used in experimental microbiology*.
- 2 Sterilisation of glass ware's. Preparation of media and sterilisation.*
- 3 Staining techniques Simple staining ; Gram's staining ; Negative staining**
- 4 Study of motility characters*.
- 5 Enumeration of micro-organisms (Total and Viable)*
- 6 Study of the methods of isolation of pure culture.*
- 7 Bio chemical testing for the identification of micro*-organisms

- 8 Cultural sensitivity testing for some micro-organisms.*
- 9 Sterility testing for powders and liquids.*
- 10 Determination of minimum inhibitory concentration.*
- 11 Microbiological assay of antibiotics by cup plate method.*
- 12 Microbiological assay of vitamins by Turbidometric method**
- 13 Determination of RWC.**
- 14 Diagnostic tests for some common diseases, Widal, malarial parasite.**

* Indicate minor experiment & ** indicate major experiment

Assignments:

- 1 Visit to some pathological laboratories & study the activities and equipment/instruments used and reporting the same.
- 2. Visit to milk dairies (Pasturization) and microbial laboratories(other sterization methods) & study the activities and equipment/instruments used and reporting the same.
- 3. Library assignments
 - a. Report of recent microbial techniques developed in diagnosing some common diseases.
 - Latest advancement developed in identifying, cultivating & handling of microorganisms.

Format of the assignment:

- 1. Minimum & Maximum number of pages.
- 2. It shall be computer draft copy.
- 3. Reference(s) shall be included at the end.
- 4. Name and signature of the student.
- 5. Assignment can be a combined presentation at the end of the academic year.
- 6. Time allocated for presentation may be 8+2 Min.

- Vanitha Kale and Kishor Bhusari Applied Microbiology || Himalaya Publishing house Mumbai.
- Harsh Mohan, Text book of Pathology ^I 3rd edition, 1998, B-3 Ansari road Darya ganj N. Delhi.

Reference books

- Prescott L.M. Jarley G.P., Klein.D.A. Microbiology. ¹2nd edition WMC Brown Publishers, Oxford. 1993.
- War Roitt, Jonathan Brostoff, David male, Immunology∥3rd edition 1996, Mosby-year book Europe Ltd, London.
- 3. Pharmacopoeia of India, Govt of India, 1996.

24PD208P

PHARMACOGNOSY AND PHYTOPHARMACEUTICALS -PRACTICAL 3H 2C

Instruction hours/ week: L: 0 T:0 P:3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Practical : 3 Hrs./Week

Course Objectives:

- Analyze the crude drugs by chemical tests.
- Determine the stomatal number and index and vein islet number, vein islet termination and palisade ratio.
- Determine the starch grains, calcium oxalate crystals by eye piece micrometer.
- Perform the Fiber length and width starch grains by Lycopodium spore method.
- Analyze the purity of crude drugs by ash value and extractive value.
- Determine the moisture content, swelling index and foaming index.

Course Outcomes (CO's): On successful completion of the course the student will

COs	Course Outcomes	Blooms Level
CO1	Explain the importance basic fundamentals of Pharmacognosy	Analyze
CO2	Analyze the macroscopy of various crude drugs	Analyze
CO3	Prepare the microscopy& powder characteristics of various crude drugs	Create
CO4	Judge the Iodinevalue & saponification value and unsaponifiable matter	Evaluate
CO5	Judge the ester & acid values	Evaluate
CO6	Analyze the phytoconsituents by chemical test	Analyze

Mapping with Programme Outcomes

POs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
COs											
CO1	Μ										
CO2				Μ							
CO3				Μ							
CO4				Μ							
CO5				Μ							
CO6				Μ							

S-Strong; M-Medium; L-Low

General Requirements: Laboratory Napkin, Observation Book 150 pages Zero brush, Needle, Blade, Match box.

List of experiments:

- 1 Introduction of Pharmacognosy laboratory and experiments.
- 2 Study of cell wall constituents and cell inclusions.

SECOND YEAR

Pharm.D

- 3 Macro, powder and microscopic study of Datura.
- 4 Macro, powder and microscopic study of Senna.
- 5 Macro, powder and microscopic study of Cassia, Cinnamon.
- 6 Macro, powder and microscopic study of Cinchona.
- 7 Macro, powder and microscopic study of Ephedra.
- 8 Macro, powder and microscopic study of Quassia.
- 9 Macro, powder and microscopic study of Clove
- 10 Macro, powder and microscopic study of Fennel.
- 11 Macro, powder and microscopic study of Coriander.
- 12 Macro, powder and microscopic study of Isapgol.
- 13 Macro, powder and microscopic study of Nux vomica.
- 14 Macro, powder and microscopic study of Rauwolfia.
- 15 Macro, powder and microscopic study of Liquorice.
- 16 Macro, powder and microscopic study of Ginger.
- 17 Macro, powder and microscopic study of Podophyllum.
- 18 Determination of Iodine value.
- 19 Determination of Saponification value and unsaponifiable matter.
- 20 Determination of ester value.
- 21 Determination of Acid value.
- 22 Chemical tests for Acacia.
- 23 Chemical tests for Tragacanth.
- 24 Chemical tests for Agar.
- 25 Chemical tests for Starch.
- 26 Chemical tests for Lipids.(castor oil, sesame oil, shark liver oil, bees wax)
- 27 Chemical tests for Gelatin.

Suggested Readings:

- 1. Pharmacognosy by G.E. Trease & W.C.Evans.
- 2. Pharmacognosy by C.K.Kokate,Gokhale & A.C.Purohit.

- 1. Pharmacognosy by Brady & Tyler.E.
- 2. Pharmacognosy by T.E.Wallis.
- 3. Pharmacognosy by C.S. Shah & Qadery.
- 4. Pharmacognosy by M.A. Iyengar.

SECOND YEAR

PHARMACOTHERAPEUTICS I -PRACTICAL3H2C

Instruction hours/ week: L: 0 T:0 P:3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Practical : 3 Hrs./Week Course Objectives:

- This course is designed to impart knowledge and skills necessary for contribution to quality use of medicines.
- Chapters dealt cover briefly pathophysiology and mostly therapeutics of various diseases.
- This will enable the student to understand the pathophysiology of common diseases and their management.
- To know the importance of preparation of individualized therapeutic plans based on diagnosis;
- To understand patient-specific parameters relevant in initiating drug therapy
- To know the pathophysiology of selected disease states and explain the rationale for drug therapy

Course Outcome (CO's):

A successful completion of this subject it is expected that students will be able to

Cos	Course Outcomes	Blooms Level
CO1	Understand the etiologic factors, signs and symptoms of selected diseases	Understand
CO2	Explain pathophysiology of selected diseases.	Knowledge
CO3	Explain drug - drug interactions for selected drugs	Knowledge
CO4	Explain drug- food interactions for selected drugs	Knowledge
CO5	Illustrate the therapeutic approach of management of selected disease.	Apply
CO6	Understand the basic concepts of different guidelines for prescribing.	Apply

Mapping with Programme Outcomes

Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12
CO1	S											
CO2	S											
CO3					М			М				
CO4					М			М				
CO5						S			S			
CO6									М		S	

S-Strong; M-Medium; L-Low

Practicals:

• Hospital postings in various departments designed to complement the lectures by providing Practical clinical discussion; attending ward rounds; follow up the progress and changes made in drug therapy in allotted patients; case presentation upon discharge. Students are required to maintain a record of cases presented and the same should be submitted at the end of the course for evaluation.

• A minimum of 20 cases should be presented and recorded covering most common diseases.

Assignments:

Students are required to submit written assignments on the topics given to them. Topics allotted should cover recent developments in drug therapy of various diseases. A minimum of THREE assignments [1500 – 2000 words] should be submitted for evaluation

Format of the assignment:

- 1. Minimum & Maximum number of pages.
- 2. Reference(s) shall be included at the end.
- 3. Assignment can be a combined presentation at the end of the academic year.
- 4. It shall be computer draft copy.
- 5. Name and signature of the student.
- 6. Time allocated for presentation may be 8+2 Min.

Suggested Readings:

- 1. Clinical Pharmacy and Therapeutics Roger and Walker, Churchill Livingstone publication.
- Pharmacotherapy: A Pathophysiologic approach Joseph T. Dipiro et al. Appleton & Lange.

- 1. Pathologic basis of disease Robins SL, W.B.Saunders publication.
- 2. Relevant review articles from recent medical and pharmaceutical literature

24PD301T

PHARMACOLOGY II-THEORY

Instruction hours/ week: L: 3 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 4 Hrs. /Week

Course Objectives:

- This subject will provide an opportunity for the student to learn about the drug with regard to classification, pharmacodynamic and pharmacokinetic aspects, adverse effects, uses, dose, route of administration, precautions, contraindications and interaction with other drugs.
- In this subject, drugs acting on autacoids, respiratory system, GIT, immune system and hormones, and pharmacology of autacoids and hormones will be concentrated.
- In addition, pharmacology of chemotherapeutic agents, vitamins, essential minerals and principles of toxicology are also taught.
- In addition to theoretical knowledge, the basic Practical knowledge relevant to therapeutics will be imparted.
- To understand drugs acting on Blood and blood forming agents
- To study structures and functions of the components of the cell

Course Outcomes (CO's):

On successful completion of the course the student will

	Course Outcomes	Blooms Level		
COs				
CO1	Understand the pharmacological aspects of drugs	Understand		
	falling under the above mentioned chapters,			
CO2	Carryout the animal experiments confidently,	Apply		
CO3	Appreciate the importance of pharmacology subject as a basis of	Knowledge		
	therapeutics, and			
CO4	Correlate and apply the knowledge therapeutically.	Analyze		
CO5	Drugs acting on Blood and blood forming agents	Understand		
C06	Structures and functions of the components of the	Understand		
	cell			

Mapping with Programme Outcomes

POs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
COs											
CO1	S										
CO2	S	S	S	S							
CO3	S										
CO4	S		S								
CO5	S										
CO6	S										

S-Strong; M-Medium; L-Low

2024-25

UNIT I

Pharmacology of Drugs acting on Blood and blood forming agents

- Anticoagulants
- Thrombolytics and antiplatelet agents
- Haemopoietics and plasma expanders

UNIT II

Pharmacology of drugs acting on Renal System

- Diuretics
- Antidiuretics

UNIT III

Chemotherapy

- Introduction
- Sulfonamides and co-trimoxazole
- Penicillins and Cephalosporins
- Tetracyclines and Chloramphenicol
- Macrolides, Aminoglycosides, Polyene & Polypeptide antibiotics
- Quinolines and Fluroquinolines

UNIT IV

Chemotherapy

- Antifungal antibiotics
- Antiviral agents
- Chemotherapy of tuberculosis and leprosy
- Chemotherapy of Malaria
- Chemotherapy of protozoal infections (amoebiasis, Giardiasis)
- Pharmacology of Anthelmintic drugs

UNIT V

• Chemotherapy : Chemotherapy of cancer (Neoplasms)

UNIT VI

- Immuno pharmacology
- Pharmacology of immune suppressants and stimulants

UNIT VII

• Principles of Animal toxicology Acute, sub acute and chronic toxicity

UNIT VIII

• The dynamic cell: The structures and functions of the components of the cell

- a) Cell and macromolecules: Cellular classification, subcellular organelles, macromolecules, large macromolecular assemblies
- b) Chromosome structure: Pro and eukaryotic chromosome structures, chromatin structure, genome complexity, the flow of genetic information.
- c) DNA replication: General, bacterial and eukaryotic DNA replication.
- d) The cell cycle: Restriction point, cell cycle regulators and modifiers.
- e) Cell signaling: Communication between cells and their environment, ion- channels, signal transduction pathways (MAP kinase, P38 kinase, JNK, Ras and PI3- kinase pathways, biosensors.

UNIT IX

• The Gene: Genome structure and function:

Gene structure: Organization and elucidation of genetic code.

• Gene expression: Expression systems (pro and eukaryotic), genetic elements that control gene expression (nucleosomes, histones, acetylation, HDACS, DNA binding protein families. Transcription and Transcription factors: Basic principles of transcription in pro andeukaryotes. Transcription factors that regulate transcription in pro and eukaryotes.

UNIT X

- **RNA processing:** rRNA, tRNA and mRNA processing.
- **Protein synthesis:** Mechanisms of protein synthesis, initiation in eukaryotes, translation control and post-translation events Altered gene functions: Mutations, deletions, amplifications, LOH, traslocations, trinucleotide repeats and other genetic abnormalities.
- Oncogenes and tumor suppressor genes.
- The gene sequencing, mapping and cloning of human disease genes.Introduction to gene therapy and targeting.
- **Recombinant DNA technology**: principles. Processes (gene transfer technology) and applications

The following details shall be submitted for any 25 selected drugs in the form of assignment/PPT.

**Name of the Drug, Brand name, Category, Therapeutic uses, Dose, Pharmacological action, Route of administration, Side effects, Contraindications, Drug Interactions with drug/food, Pharmacokinetics

Suggested Readings (Theory):

- Tripathi, K. D. Essentials of medical pharmacology. 4th edition, 1999. Publisher: Jaypee, Delhi.
- ^{2.} Satoskar, R.S. and Bhadarkar, S.D. Pharmacology and pharmacotherapeutics. 16th edition (single volume), 1999. Publisher: Popular, Dubai.
- Rang, H.P. and Dale, M.M. Pharmacology. 4th edition, 1999. Publisher: Churchill Living stone.

- Goodman Gilman, A., Rall, T.W., Nies, A.I.S. and Taylor, P. Goodman and Gilman's The pharmacological Basis of therapeutics. 9th edition, 1996. Publisher: Mc Graw Hill, Pergamon press.
- 2. Craig, C.R. and Stitzel, R.E. Modern Pharmacology. Latest edition. Publisher: Little Brown and company.
- 3. Katzung, B.G. Basic and clinical pharmacology. Latest edition. Publisher: Prentice Hall, International.
- 4. Gupta, P.K. and Salunkhe, D.K. Modern Toxicology. Volume I, II and III. Latest edition. Publisher: B.V. Gupta, Metropolitan Book Co. (p) Ltd, New Delhi.

24PD302T

2024-25

THIRD YEAR

PHARMACEUTICAL ANALYSIS - THEORY4H4C

Instruction hours/ week: L: 3 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory : 4 Hrs. /Week

Course Objectives:

- This subject deals with the application of instrumental methods in qualitative and quantitative analysis of drugs.
- This subject is designed to impart a fundamental knowledge on the principles and instrumentation of spectroscopic and chromatographic technique.
- This also emphasizes on theoretical and Practical knowledge on modern analytical instruments that are used for drug testing.
- To discuss the applications of analytical techniques.
- To perform quantitative analysis of drugs using various analytical instruments.
- To perform qualitative analysis of drugs using various analytical instruments

Course Outcomes (CO's):

At the end of this course, students will be able to

COs	Course Outcomes	Blooms Level
CO1	Understand the interaction of matter with electromagnetic	Understand
	radiations and its applications in drug analysis.	
CO2	Describe the instrumentation of spectroscopy techniques.	Apply
CO3	Understand the chromatographic separation and analysis of drugs.	Understand
CO4	Discuss the applications of analytical techniques.	Apply
CO5	Perform quantitative analysis of drugs using various analytical	Apply
	instruments.	
CO6	Perform qualitative analysis of drugs using various analytical	Apply
	instruments.	

Mapping with Programme Outcomes

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12
CO1			Μ		L							
CO2			S		S							
CO3			Μ		Μ							
CO4			S		S							
CO5			S		S							
CO6			S		Μ							

S-Strong; M-Medium; L-Low

Course Content:

UNIT I

Quality Assurance:

- a. Introduction, sources of quality variation, control of quality variation.
- b. Concept of statistical quality control.
- c. Validation methods- quality of equipment, validation of equipment and validation of analytical instruments and calibration.
- d. GLP, ISO 9000.
- e. Total quality management, quality review and documentation.
- f. ICH- international conference for harmonization-guidelines.
- g. Regulatory control.

UNIT II

Chromatography:

Introduction, history, classification, separation techniques, choice of methods. The following techniques be discussed with relevant examples of pharmaceutical products involving principles and techniques of separation of drugs from excipients.

- h. **Column Chromatography**: Adsorption column chromatography, Operational technique, frontal analysis and elution analysis. Factors affecting column efficiency, applications and partition chromatography.
- i. TLC: Introduction, principle, techniques, R_f value and applications.
- j. **PC:** Introduction, principle, types of paper chromatography, preparation techniques, development techniques, applications.
- k. **Ion-exchange chromatography**: Introduction, principles, types of ion exchange synthetic resins, physical properties, factors affecting ion exchange, methodology and applications.
- 1. **HPLC**: Introduction, Theory, instrumentation, and applications.
- m. **HPTLC**: Introduction, Theory, instrumentation, and applications.
- n. **Gas Chromatography**: Introduction, Theory, instrumentation-carrier gases,types of columns, stationary phases in GLC & GSC. Detectors- Flame ionization detectors, electron capture detector, thermal conductivity detector. Typical gas chromatogram, derivatisation techniques, programmedtemperature gas chromatography, applications.
 - h. **Electrophoresis**: Principles of separation, equipment for paper and gelelectrophoresis, and application.
 - i. Gel filtration and affinity chromatography: Introduction, technique, applications.

UNIT III

Electrometric Methods:

Theoretical aspects, instrumentation, interpretation of data/spectra and analytical applications be discussed on the following topics.

- **Potentiometry**: Electrical potential, electrochemical cell, reference electrodes, indicator electrodes, measurement of potential and pH, construction and working of electrodes, Potentiometric titrations, methods of detecting end point, Karl Fischer titration.
- **Conductometry**: Introduction, conductivity cell, conductometric titrations and applications.
- **Polarography**: Instrumentation, DME, residual current, diffusion current and limiting current, polarographic wave, Ilkovic's equation, Effect of oxygen on polarographic wave, Polarographic maxima and suppressors and applications.
- Amperometric Titrations: Introduction, types of electrodes used, reference and indicator electrode, instrumentation, titration procedure, advantages and disadvantages of Amperometry over potentiometry. Pharma applications.

UNIT IV

Spectroscopy:

Theoretical aspects, instrumentation, elements of interpretation of data/spectra and application of analytical techniques be discussed on.

Absorption Spectroscopy:

Theory of electronic, atomic and molecular spectra. Fundamental laws of photometry, Beer-Lambert's Law, application and its deviation, limitation of Beer law, application of the law to single and multiple component analysis, measurement of equilibrium constant and rate constant by spectroscopy. Spectra of isolated chromophores, auxochromes, batho-chromic shift, hypsochromic shift, hyperchromic and hypochromic effect, effect of solvent on absorption spectra,molecular structure and infrared spectra.

UNIT V

Instrumentation – Photometer, UV Visible spectrometer – sources of UV – Visible radiations, collimating systems, monochromators, sample cells and following detectors – photocell Barrier layer cell, Phototube, Diode array, applications of U.V.-Visible spectroscopy in pharmacy and spectrophotometric titrations.

UNIT VI

Infrared Spectroscopy: Vibrational transitions, frequency – structure correlations, Infrared absorption bands, Instrumentation–IR spectro-meter – sources of IR, Collimating systems, monochromators, sample cells, sample handling in IR spectroscopy and detectors– Thermocouple, Golay Cells, Thermistor, Bolometer, Pyroelectric detector, Applications of IR in pharmacy.

UNIT VII

Fluorimetric Analysis: Theory, luminescence, factors affecting fluorescence, quenching. Instrumentation, Applications, fluorescent indicators, study of pharmaceutically important compounds estimated by fluorimetry.

Flame Photometry: Theory, nebulisation, flame and flame temperature, interferences, flame spectrometric techniques and instrumentation and pharmaceutical applications.

Atomic Absorption Spectrometry: Introduction, Theory, types of electrodes, instrumentation and applications.

UNIT VIII

Atomic Emission Spectroscopy: Spectroscopic sources, atomic emission spectrometers, photographic and photoelectric detection.

NMR & ESR (introduction only): Introduction, theoretical aspects and applications.

UNIT IX

Mass Spectroscopy: (**Introduction only**) – Fragmentation, types of ions produced mass spectrum and applications.

Polarimetry: (Introduction only) – Introduction to optical rotatory dispersion, circular dichroism, polarimeter.

UNIT X

X-RAY Diffraction: (Introduction only) – Theory, reciprocal lattice concept, diffraction patterns and applications.

Thermal Analysis: Introduction, instrumentation, applications, and DSC and DTA.

Suggested Readings:

- A.H.Beckett& J.B. Stenlake's, Practical Pharmaceutical Chemistry Vol I&II, StahlonePress of University of London.
- 2. A.I.Vogel, TextBook of Quantitative Inorganic analysis.
- 3. P.Gundu Rao, Inorganic Pharmaceutical Chemistry.
- 4. Bentley and Driver's Textbook of Pharmaceutical Chemistry.

Reference Books

- 1. John H. Kennedy, Analytical chemistry principles.
- 2. Indian Pharmacopoeia.

24PD303T

THIRD YEAR

PHARMACOTHERAPEUTICS II - THEORY 4H 4C

Instruction hours/ week: L: 3 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 4 Hrs. /Week

Course Objectives:

- This course is designed to impart knowledge and skills necessary for contribution to quality use of medicines.
- Chapters dealt cover briefly pathophysiology and mostly therapeutics of various diseases. This will enable the student to understand the pathophysiology of common diseases and their management.
- To know the importance of preparation of individualized therapeutic plans based on diagnosis
- To appreciate the needs to identify the patient-specific parameters relevant in initiating drug therapy, and monitoring therapy (including alternatives, time-course of clinical and laboratory indices of therapeutic response and adverse effects).
- To develop patient case based assessment skills

Course Outcomes (CO's):

At the end of this course, students will be able to

COs	Course Outcomes	Blooms Level
CO1	Know the pathophysiology ofselected disease states and the rationale for drug therapy	Knowledge
CO2	Know the therapeutic approach to management of these diseases;	Analyse
CO3	Know the controversies in drug therapy;	Evaluate
CO4	Know the importance of preparation of individualised therapeutic plans based on diagnosis; and	Apply
CO5	Appreciate the needs to identify the patient-specific parameters relevant in initiating drug therapy, and monitoring therapy (including alternatives, time-course of clinical and laboratory indices of therapeutic response and adverse effects).	Evaluate
CO6	Students will be developing patient case based assessment skills	Create

Mapping with Programme Outcome:

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1	S	S	S		Μ	S	S		Μ		S
CO2		S			S	М					L
CO3	S	S	М		L						S
CO4											
					S		S			Μ	L
CO5	Μ				S	S		L		L	Μ
CO6					S			S		L	М

S-Strong; M-Medium; L-Low

Course Outcomes (CO's):

- 1. Know the pathophysiology of selected disease states and the rationale for drug therapy
- 2. Know the therapeutic approach to management of these diseases;
- 3. Know the controversies in drug therapy;
- 4. Know the importance of preparation of individualised therapeutic plans based on diagnosis; and
- 5. Appreciate the needs to identify the patient-specific parameters relevant in initiating drug therapy, and monitoring therapy (including alternatives, time-course of clinical and laboratory indices of therapeutic response and adverse effects).
- 6. Students will be developing patient case based assessment skills

Detailed syllabus and lecture wise schedule : Etiopathogenesis and pharmacotherapy of diseases associated with following systems / diseases – UNIT I

Infectious disease: Guidelines for the rational use of antibiotics and surgical Prophylaxis,

Tuberculosis, Meningitis, Respiratory tract infections

UNIT II

Infectious disease: Gastroenteritis, Endocarditis, Septicemia

UNIT III

Infectious disease: Urinary tract infections, Protozoal infection- Malaria, HIV &

Opportunistic infections

UNIT IV

Infectious disease: Fungal infections, Viral infections, Gonarrhoea and Syphillis

UNIT V

Musculoskeletal disorders

UNIT VI

Renal system

Acute Renal Failure, Chronic Renal Failure, Renal Dialysis, Drug induced renal disorders

UNIT VII

Oncology: Basic principles of Cancer therapy, General introduction to cancer chemotherapeutic agents

UNIT VIII

Chemotherapy of breast cancer, leukemia.

UNIT IX

Infectious disease: Management of chemotherapy nausea and emesis

UNIT X

Dermatology: Psoriasis, Scabies, Eczema, Impetigo

Suggested Readings:

1. Clinical Pharmacy and Therapeutics - Roger and Walker, Churchill Livingstone publication

- Pharmacotherapy: A Pathophysiologic approach Joseph T. Dipiro et al. Appleton & Lange
- Clinical Pharmacy and Therapeutics Eric T. Herfindal, Williams and Wilkins Publication
- Applied Therapeutics: The clinical Use of Drugs. Lloyd Young and Koda-Kimble MA]

24PD304T

THIRD YEAR

PHARMACEUTICAL JURISPRUDENCE - THEORY2H2C

Instruction hours/ week: L: 2 T:0 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory : 2 Hrs.

/WeekCourse

Objectives:

- This course exposes the student to several important legislations related to the profession of pharmacy in India.
- The Drugs and Cosmetics Act, along with its amendments is the core of this course.
- Other acts, which are covered, include the Pharmacy Act, dangerous drugs, medicinal and toilet preparation Act etc.
- Besides this the new drug policy, professional ethics, DPCO, patent and design Act will be discussed.
- To acquire clear idea on drug price control order and its implication in India.
- To acquire information regarding the licenses to be achieved for a new drug discovery process.

Course Outcomes (CO's):

At the end of this course, students will be able to

COs	Course Outcomes	Blooms
		Level
CO1	Practice the Professional ethics;	Apply
CO2	Understand the various concepts of the pharmaceutical legislation in	Understand
	India;	
CO3	Know the various parameters in the Drug and Cosmetic Act and rules;	Knowledge
CO4	Know the Drug policy, DPCO, Patent and design act;	Knowledge
CO5	Understand the labeling requirements and packaging guidelines for	Understand
	drugs and cosmetics;	
CO6	Be able to understand the concepts of Dangerous Drugs Act, Pharmacy	Understand
	Act and Exciseduties Act.	

Mapping with Programme Outcomes

CO's	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9
CO1							S		
CO2									Μ
CO3									Μ
CO4									Μ
CO5									Μ
CO6									Μ

S-Strong; M-Medium; L-Low

Detailed syllabus and lecture wise schedule:

UNIT I

- **Pharmaceutical Legislations** A brief review.
- Principle and Significance of professional ethics. Critical study of the code of pharmaceutical ethics drafted by PCI.

UNIT II

Drugs and Cosmetics Act, 1940, and its rules 1945.

Objectives, Legal definition, Study of Schedule's with reference to Schedule B,C&C1, D, E1,

F&F1, F2, F3, FF, G, H, J, K, M, N, P, R, V, W, X, Y.

Sales, Import, labeling and packaging of Drugs And Cosmetics Provisions Relating to Indigenous Systems.

Constitution and Functions of DTAB, DCC, CDL.

Qualification and duties -Govt. analyst and Drugs Inspector.

UNIT III

Pharmacy Act -1948.

Objectives Legal Definitions, General Study, Constitution and Functions of State & Central Council, Registration & Procedure, ER.

UNIT IV

Medicinal and Toilet Preparation Act -1955.

Objectives, Legal Definitions, Licensing, Bonded and Non Bonded Laboratory,

Ware Housing, Manufacture of Ayurvedic, Homeopathic, Patent & Proprietory Preparations.

UNIT V

Narcotic Drugs and Psychotropic substances Act-1985 and Rules. Objectives, Legal Definitions, General Study, Constitution and Functions of narcotic & Psychotropic Consultative Committee, National Fund for Controlling the Drug Abuse, Prohibition, Control and regulations, Schedules to the Act.

UNIT VI

Study of Salient Features of Drugs and magic remedies Act and its rules.

UNIT VII

Study of essential Commodities Act Relevant to drugs price control Order.

UNIT VIII

Drug Price control Order & National Drug Policy (Current).

UNIT IX

Prevention of Cruelty to animals Act-1960.

Patents & design Act-1970.

UNIT X

Brief study of prescription and Non-prescription Products.

Assignments:

Format of the assignment

- Minimum & Maximum number of pages
- It shall be a computer draft copy
- Reference(s) shall be included at the end.
- Name and signature of the student
- Assignment can be a combined presentation at the end of the academic year.
- Time allocated for presentation may be 8+2 Min

Case studies relating to

- Drugs and Cosmetics Act and rules along with its amendments, Dangerous Drugs Act, Medicinal and Toilet preparation Act, New Drug Policy, Professional Ethics, Drugs (Price control) Order, Patent and Design Act.
- Various prescription and non-prescription products.
- Medical and surgical accessories.
- Diagnostic aids and appliances available in the market.

Suggested Readings:

1. Mithal, B M. Textbook of Forensic Pharmacy. Calcutta :National; 1988.

Reference Books (Latest Editions):

- Singh, KK, editor. Beotra's the Laws of Drugs, Medicines & cosmetics. Allahabad: Law Book House; 1984.
- 2. Jain, NK. A Textbook of forensic pharmacy. Delhi: Vallabh prakashan ; 1995.
- 3. Reports of the Pharmaceutical enquiry Committee
- 4. I.D.M.A., Mumbai. DPCO 1995
- 5. Various reports of Amendments.
- 6. Deshapande, S.W. The drugs and magic remedies act 1954 and rules 1955. Mumbai: Susmit

Publications; 1998.

 Eastern Book Company .The narcotic and psychotropic substances act 1985, Lucknow: Eastern; 1987.
2024-25

24PD305T

MEDICINAL CHEMISTRY - THEORY

THIRD YEAR4H4C

Instruction hours/ week: L: 3 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 4 Hrs. /Week

Course Objectives:

- This subject is designed to impart fundamental knowledge on the structure, chemistry and therapeutic value of drugs.
- The subject emphasis on modern techniques of rational drug design like quantitative structure activity relationship (QSAR)
- To understand the prodrug concept, combinatorial chemistry and Computer aided drug design (CADD).
- The subject also emphasizes on the chemistry, mechanism of action, metabolism,
- To know adverse effects, Structure Activity Relationships (SAR) and therapeutic uses
- To synthesis of important drugs.

Course Outcomes (CO's):

At the end of this course, students will be able to

COs	Course Outcomes	Blooms Level
CO1	Illustrate the classification of drugs.	Understand
CO2	Explain the mechanism of action of drugs.	Apply
CO3	Understand the chemistry of drugs with respect to their biological activity.	Understand
CO4	Know the metabolism, adverse effects and therapeutic value of drugs.	Understand
CO5	Discuss the importance of SAR of drugs.	Understand
CO6	Understand the importance of drug design and different techniques of drug design.	Understand

Mapping with Programme Outcomes

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12
CO1	Μ										Μ	
CO2	Μ										Μ	
CO3	Μ										Μ	
CO4	Μ										Μ	
CO5	Μ										Μ	
CO6	Μ										Μ	

S-Strong; M-Medium; L-Low

UNIT I

Modern concept of rational drug design: A brief introduction to Quantitative Structure Activity Relationaship (QSAR), prodrug, combinatorial chemistry and computer aided drug design (CADD) and concept of antisense molecules.

UNIT II

A study of the development of the following classes of drugs including SAR, mechanism of action, synthesis of important compounds, chemical nomenclature, brand names of important marketed products and their side effects.

UNIT III

Anti-infective agents

- Local anti-infective agents
- Preservatives
- Antifungal agents
- Urinary tract anti-infectives

UNIT IV

Anti-infective agents

- Antitubercular agents
- Antiviral agents and Anti AIDS agents
- Antiprotozoal agents
- Anthelmentics
- Antiscabies and Antipedicular agents

UNIT V

- Sulphonamides and sulphones
- Antimalarials

UNIT VI

- Antibiotics
- Antineoplastic agents

UNIT VII

Cardiovascular agents

- Antihypertensive agents
- Antianginal agents and vasodilators

UNIT VIII

Cardiovascular agents

- Antiarrhythmic agents
- Antihyperlipidemic agents
- Coagulants and Anticoagulants

• Endocrine

UNIT IX

- Hypoglycemic agents
- Thyroid and Antithyroid agents

UNIT X

- Diureties
- Diagnostic agents
- Steroidal Hormones and Adrenocorticoids

Suggested Readings:

- 1. Wilson and Gisvold's Text book of Organic, Medicinal and Pharmaceutical
- 2. Chemistry, Lippincott-Raven Publishers-New York, Philadelphia.
- William.O.Foye, Principles of Medicinal Chemistry, B.I. Waverly Pvt. Ltd., New Delhi.
- 4. Burgers, Medicinal Chemistry, M.E., Welly Med. Chemistry M.E. Walffed Johnwilley and Sons, Wiley-interscience Publication, New York, Toranto.
- A Text Book of Medicinal Chemistry Vol. I and II by Surendra N. Pandeya, S.G. Publisher, 6, Dildayal Nagar, Varanasi -10.
- Indian Pharmacopoeia 1985 and 1996. The Controller of Publications, Civil Lines, Delhi - 54.

- Current Index of Medical Specialities (CIMS) and MIMS India, MIMS, A.E. Morgan Publications (I) Pvt. Ltd, New Delhi-19.
- 2. Organic Drug Synthesis-Ledniser Mitzsher Vol. I and II.
- Pharmaceutical Chemistry drug Synthesis Vol. I and II by H. J. Roth and A. Kleemann.
- The Science and Practice of Pharmacy Vol. 1 and 2, Remington, MACK Publishing Company, Easton, Pennsylvania.

THIRD YEAR

PHARMACEUTICAL FORMULATIONS - THEORY 3H 3C

Instruction hours/ week: L: 2 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 3 Hrs. /Week

Course Objectives:

- Course enables the student to understand and appreciate the influence of pharmaceutical additives and various pharmaceutical dosage forms on the performance of the drug product.
- To discover various formulation considerations in development of pharmaceutical dosage forms like tablets, capsules, etc.
- To Understand the quality control tests for the dosage forms.
- To know parenterals and novel drug delivery system.
- To understand clearly about packaging and cosmetic preparations.
- To interpret the various pharmaceutical additives to be included in all dosage forms.

Course Outcomes (CO's):

At the end of this course, students will be able to

COs	Course Outcomes	Blooms	
		Level	
CO1	Aquire knowledge about the variouspharmaceutical dosage	Knowledge	
	forms and their manufacturing techniques.		
CO2	Discover various formulation considerations in development of	Create	
	pharmaceutical dosageforms like tablets, capsules, etc.		
CO3	Understand the quality control tests for the dosage forms.	Understand	
CO4	Detail on parenterals and Novel Drug Delivery System.	Apply	
CO5	Understand clearly about packaging and cosmetic preparations.	Apply	
CO6	Interpret the various pharmaceutical additives to be included in all dosage	Analyze	
	forms		

Mapping with Programme Outcomes

CO's	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9
CO1	L								
CO2	L								
CO3	L								
CO4	L								
CO5	L								
CO6	L								

S-Strong; M-Medium; L-Low

Detailed syllabus and lecture wise schedule:

UNIT I

Pharmaceutical dosage form- concept and classification

UNIT II

Tablets: Formulation of different types of tablets, tablet excipients, granulation techniques

quality control and evaluation of tablets.

UNIT III

Tablet coating, Type of coating, quality control tests for coated tablet.

UNIT IV

Capsules; Production and filling of hard gelatin capsules, Raw material for shell, finishing, quality control tests for capsules.

UNIT V

Production and filling of soft gelatin capsules, quality control tests for soft gelatin capsules.

UNIT VI

Liquid orals: Formulation and evaluation of suspensions, emulsions and solutions. Stability of these preparations

UNIT VII

Parenterals Introduction Containers used for Parenterals (including official tests) Formulation of large and small volume Parenterals Sterilization

UNIT VIII

Ophthalmic preparations (Semi – Solids): Introduction and classification Factors affecting absorption and anatomy of eye Packaging storage and labeling

UNIT IX

Ointments Types of Ointment Base Preparation of ointment, Jellies Types of jellies Formulation of jellies Suppositories, Method of preparation, Types Packaging

UNIT X

Definition and concept of Controlled and novel Drug delivery systems with available examples, viz. parentral, trans dermal, buccal, rectal, nasal, implants, ocular

Suggested Readings:

- 1. Pharmaceutical dosage forms, Vol, I,II and III by lachman
- 2. Rowlings Text book of Pharmaceutics
- 3. Tutorial Pharmacy Cooper & Gunn

- 1. Remington's Pharmaceutical Sciences
- 2. USP/BP/IP

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THIRD YEAR

PHARMACOLOGY II - PRACTICAL 3H 2C

Instruction hours/ week: L: 0 T:0 P:3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Practical : 3 Hrs./Week

Course Objectives:

- To Calculate the dose in pharmacological experiments.
- To Perform various pharmacological screening studies.
- To Demonstrate the toxicity studies in animal models.
- To know the student's t test, ANOVA, Chi square test, Wilcoxon Signed Rank test.
- To determine the pharmacokinetic parameters by using the data.
- To evaluate the acute skin irritation, acute eye irritation and corrosion of a test substance.

Course Outcomes (CO's):

On successful completion of the course the student will

COs	Course Outcomes	Blooms Level			
CO1	Understand the pharmacological aspects of drugs	Understand			
	falling under the above mentioned chapters,				
CO2	Carryout the animal experiments confidently,	Apply			
CO3	Appreciate the importance of pharmacology subject as a basis of	Knowledge			
	therapeutics, and				
CO4	Correlate and apply the knowledge therapeutically.	Analyze			
CO5	Drugs acting on Blood and blood forming agents	Understand			
CO6	Structures and functions of the components of the cell	Understand			

Mapping with Programme Outcomes

POs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
COs											
CO1			S								
CO2			S	S							
CO3	S										
CO4	S			S							
CO5	S		S								
CO6			S								

S-Strong; M-Medium; L-Low

List of Experiments:

- 1. Study of laboratory animals and their handling (a. Frogs, b. Mice, c. Rats, d. Guinea pigs, e. Rabbits).
- 2. Study of physiological salt solutions used in experimental pharmacology.
- 3. Study of laboratory appliances used in experimental pharmacology.
- 4. Study of use of anesthetics in laboratory animals.
- 5. To record the dose response curve of Ach using isolated ileum/rectus abdominis muscle preparation.
- 6. To carry out bioassay of Ach using isolated ileum/rectus abdominis muscle preparation by interpolation method.
- To carry out bioassay of Ach using isolated ileum/rectus abdominis muscle preparation by three point method.
- 8. To record the dose response curve of Histamine using isolated guinea -pig ileum preparation.
- 9. Study of agonistic and antagonistic effects of drugs using isolated guinea-pig ileum preparation.
- 10. To carry out bioassay of Histamine using isolated guinea -pig ileum preparation by interpolation method.
- 11. To carry out bioassay of Histamine using guinea-pig ileum preparation by three point method.
- 12. To study the routes of administration of drugs in animals (Rats, Mice, Rabbits).
- 13. Study of Theory, principle, procedure involved and interpretation of given results for the following experiments:
 - a) Analgesic property of drug using analgesiometer.
 - b) Antiinflammatory effect of drugs using rat-paw edema method.
 - c) Anticonvulsant activity of drugs using maximal electroshock and pentylene tetrazole methods.
 - d) Antidepressant activity of drugs using pole climbing apparatus and pentobarbitone induced sleeping time methods.
 - e) Locomotor activity evaluation of drugs using actophotometer and rotarod.
 - f) Cardiotonic activity of drugs using isolated frog heart and mammalian heart preparations.

Suggested Readings

- Tripathi, K. D. Essentials of medical pharmacology. 4th edition, 1999. Publisher: Jaypee, Delhi.
- Satoskar, R.S. and Bhadarkar, S.D. Pharmacology and pharmacotherapeutics. 16th edition (single volume), 1999. Publisher: Popular, Dubai.
- Rang, H.P. and Dale, M.M. Pharmacology. 4th edition, 1999. Publisher: Churchill Living stone.

Reference Books (Latest Editions):

1. Gupta, P.K. and Salunkhe, D.K. Modern Toxicology. Volume I, II and III. Latest edition. Publisher: B.V. Gupta, Metropolitan Book Co. (p) Ltd, New Delhi.

THIRD YEAR

PHARMACEUTICAL ANALYSIS -PRACTICAL 3H 2C

Instruction hours/ week: L: 0 T:0 P:3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Practical: 3 Hrs./Week

Course Objectives:

- To estimate the samples using analytical instruments.
- To perform assay of drug samples using analytical intruments
- To determine the effect of solvents on absorption maxima.
- To separate the mixtures of sample using chromatographic techniques.
- To demonstrate HPLC.
- To demonstrate gas chromatography.

Course Outcomes

At the end of this course, students will be able to

COs	Course Outcomes	Blooms Level
CO1	Estimate the samples using analytical instruments	Apply
CO2	Perform assay of drug samples using analytical instruments	Apply
CO3	Determine the effect of solvents on absorption maxima.	Apply
CO4	Separate the mixtures of sample using chromatographic techniques.	Apply
CO5	Demonstrate HPLC.	Apply
CO6	Demonstrate gas chromatography.	Apply

Mapping with Programme Outcomes

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12
CO1			S	S								
CO2			S	S								
CO3			S	S								
CO4			S	Μ								
CO5			S	Μ								
CO6			S	Μ								

S-Strong; M-Medium; L-Low

Course Content:

List of Experiments:

- 1. Separation and identification of Amino Acids by Paper Chromatography.
- 2. Separation and identification of Sulpha drugs by TLC technique.

- 3. Effect of pH and solvent on the UV spectrum of given compound.
- 4. Comparison of the UV spectrum of a compound with that of its derivatives.
- 5. Determination of dissociation constant of indicators using UV-Visible spectroscopy.
- 6. Conductometric titration of mixture of acids with a strong base.
- 7. Potentiometric titration of a acid with a strong base.
- 8. Estimation of drugs by Fluorimetric technique.
- 9. Study of quenching effect in fluorimetry.
- 10. Colourimetric estimation of Supha drugs using BMR reagent.
- 11. Simultaneous estimation of two drugs present in given formulation.
- 12. Assay of Salicylic Acid by colourimetry.
- Determination of Chlorides and Sulphates in Calcium gluconate by Nepheloturbidimetric Method.
- 14. Determination of Na/K by Flame Photometry.
- 15. Determination of pKa using pH meter.
- 16. Determination of specific rotation.
- 17. Comparison of the IR spectrum of a compound with that of its derivatives.
- 18. Demonstration of HPLC.
- 19. Demonstration of HPTLC.
- 20. Demonstration of GC-MS.
- 21. Demonstration of DSC.
- 22. Interpretation of NMR spectra of any one compound.

Suggested Readings:

1. A.H.Beckett& J.B. Stenlake's, Practical Pharmaceutical Chemistry Vol I&II, Stahlone

Press of University of London.

- 2. A.I.Vogel, TextBook of Quantitative Inorganic analysis.
- 3. P.Gundu Rao, Inorganic Pharmaceutical Chemistry.
- 4. Bentley and Driver's Textbook of Pharmaceutical Chemistry.

- 1. John H. Kennedy, Analytical chemistry principles.
- 2. Indian Pharmacopoeia.

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THIRD YEAR

PHARMACOTHERAPEUTICS II - PRACTICAL 3H 2C

Instruction hours/ week: L: 0 T:0 P:3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Practical : 3 Hrs./Week

Course Objectives:

- This course is designed to impart knowledge and skills necessary for contribution to quality use of medicines.
- Chapters dealt cover briefly pathophysiology and mostly therapeutics of various diseases. This will enable the student to understand the pathophysiology of common diseases and their management.
- To know the importance of preparation of individualised therapeutic plans based on diagnosis
- To appreciate the needs to identify the patient-specific parameters relevant in initiating drug therapy, and monitoring therapy (including alternatives, time-course of clinical and laboratory indices of therapeutic response and adverse effects).
- To develop patient case based assessment skills

Course Outcomes (CO's):

At the end of this course, students will be able to

COs	Course Outcomes	Blooms Level
CO1	Know the pathophysiology of selected disease states and the	Knowledge
	rationale for drug therapy	
CO2	Know the therapeutic approach to management of these	Analyse
	diseases;	
CO3	Know the controversies in drug therapy	Evaluate
CO4	Know the importance of preparation of individualised	Apply
	therapeutic plans based on diagnosis; and	
CO5	Appreciate the needs to identify the patient-specific parameters	Evaluate
	relevant in initiating drug therapy, and monitoring therapy	
	(including alternatives, time-course of clinical and laboratory	
	indices of therapeutic response and adverse effects).	
CO6	Students will be developing patient case based assessment skills	Create

	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1	S	S	S		Μ	S	S		М		S
CO2		S		Μ	S	Μ			М		S
CO3	S	S	М		L						
CO4					S	S			М	М	
CO5	М			S	S		S	L			L
CO6				S	S			S		М	L

Mapping with Programme Outcome:

S-Strong; M-Medium; L-Low

Practicals :

• Hospital postings in various departments designed to complement the lectures by providing Practical clinical discussion; attending ward rounds; follow up the progress and changes made in drug therapy in allotted patients; case presentation upon discharge. Students are required to maintain a record of cases presented and the same should be submitted at the end of the course for evaluation.

• The student shall be trained to understand the principle and practice involved in selection of drug therapy including clinical discussion.

• A minimum of 20 cases should be presented and recorded covering most common diseases.

Assignments :

Students are required to submit written assignments on the topics given to them. Topics allotted should cover recent developments in drug therapy of various diseases. A minimum of THREE assignments [1500 – 2000 words] should be submitted for evaluation.

Format of the assignment :

- 1. Minimum & Maximum number of pages.
- 2. Reference(s) shall be included at the end.
- 3. Assignment can be a combined presentation at the end of the academic year.
- 4. It shall be computer draft copy.
- 5. Name and signature of the student.
- 6. Time allocated for presentation may be 8+2 Min.

Suggested Readings:

1. Clinical Pharmacy and Therapeutics - Roger and Walker, Churchill Livingstone publication

Reference Books (Latest Editions):

1. Pharmacotherapy: A Pathophysiologic approach - Joseph T. Dipiro et al. Appleton &

Lange

Practical : 3 Hrs./Week

Course Objectives:

• To prepare drugs and medicinally important compounds by traditional and microwave method.

MEDICINAL CHEMISTRY - PRACTICAL

- To prepare drug intermediates by traditional and microwave method.
- To perform assay of drug substances.
- To draw structures of chemicals using soft ware's.
- To determine physicochemical properties for drugs using software.
- To screen drug likeliness.

Course Outcomes

At the end of this course, students will be able to

COs	Course Outcomes	Blooms Level
CO1	Prepare drugs and medicinally important compounds by traditional and microwave method.	Apply
CO2	Prepare drug intermediates by traditional and micro wave method.	Apply
CO3	Perform assay of drug substances.	Apply
CO4	Draw structures of chemicals using softwares.	Apply
CO5	Determine physicochemical properties for drugs using software.	Understand
CO6	Screen drug like liness.	Understand

Mapping with Programme Outcomes

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12
CO1	S	Μ										
CO2	S	Μ										
CO3	S	Μ										
CO4	L	L										
CO5	Μ	L										
CO6	L	L										

S-Strong; M-Medium; L-Low

24PD310P

THIRD YEAR

3H 2C

External Semester Exam: 3 Hours

Marks: Internal: 30 External: 70 Total: 100

2024-25

CONTENT

- 1. Assays of important drugs from the course content.
- 2. Preparation of medicinally important compounds or intermediates required for synthesis of drugs.
- 3. Monograph analysis of important drugs.
- 4. Determination of partition coefficients, dissociation constants and molar refractivity of compounds for QSAR analysis.

Suggested Readings:

- 1. Wilson and Gisvold's Text book of Organic, Medicinal and Pharmaceutical
- 2. Chemistry, Lippincott-Raven Publishers-New York, Philadelphia.
- Indian Pharmacopoeia 1985 and 1996. The Controller of Publications, Civil Lines, Delhi - 54.

- Current Index of Medical Specialities (CIMS) and MIMS India, MIMS, A.E. Morgan Publications (I) Pvt. Ltd, New Delhi-19.
- 2. Organic Drug Synthesis-Ledniser Mitzsher Vol. I and II.
- 3. Pharmaceutical Chemistry drug Synthesis Vol. I and II by H. J. Roth and A. Kleemann.
- 4. The Science and Practice of Pharmacy Vol. 1 and 2, Remington, MACK Publishing Company, Easton, Pennsylvania.

THIRD YEAR

PHARMACEUTICAL FORMULATIONS - PRACTICAL 3H 2C

Instruction hours/ week: L: 0 T:0 P:3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Practical: 3 Hrs./Week

Course Objectives:

- To manufacture tablets.
- To understand the strict formulation considerations in parenteral and ophthalmic manufacturing.
- To demonstrate the evaluations of different packaging materials in pharmaceutical industry.
- To achieve skills in making a pharmaceutical product.
- To demonstrate the manufacturing of capsules.
- To exploit the formulation of various cosmetics.

Course Outcomes (CO's):

At the end of this course, students will be able to

COs	Course Outcomes	Blooms Level
CO1	Manufacture tablets.	Apply
CO2	Understand the strict formulation considerations in parenteral and ophthalmic manufacturing.	Understand
CO3	Demonstrate the evaluations of different packaging materials in pharmaceuticalindustry.	Evaluation
CO4	Achieve skills in making a pharmaceutical product.	Apply
CO5	Demonstrate the manufacturing of capsules.	Apply
CO6	Exploit the formulation of various cosmetics.	Analyze

Mapping with Programme Outcomes

CO's	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9
CO1	L								
CO2	L								
CO3	L								
CO4	L								
CO5	L								
CO6	L								

S-Strong; M-Medium; L-Low

List of Experiments :

1. Manufacture of Tablets

- a. Ordinary compressed tablet-wet granulation
- **b.** Tablets prepared by direct compression.
- **c.** Soluble tablet.
- d. Chewable tablet.

2. Formulation and filling of hard gelatin capsules

3. Manufacture of parenterals

- **a.** Ascorbic acid injection
- **b.** Calcium gluconate injection
- **c.** Sodium chloride infusion.
- d. Dextrose and Sodium chloride injection/ infusion.

4. Evaluation of Pharmaceutical formulations (QC tests)

- **a.** Tablets
- **b.** Capsules
- **c.** Injections

5. Formulation of two liquid oral preparations and evaluation by assay

- a. Solution: Paracetamol Syrup
- b. Antacid suspensions- Aluminum hydroxide gel

6. Formulation of semisolids and evaluation by assay

- a. Salicyclic acid and benzoic acid ointment
- **b.** Gel formulation Diclofenac gel

7. Cosmetic preparations

- a. Lipsticks
- **b.** Cold cream and vanishing cream
- c. Clear liquid shampoo
- **d.** Tooth paste and tooth powders.

8. Tablet coating (demonstration)

Suggested Readings:

- 1. Pharmaceutical dosage forms, Vol, I,II and III by lachman
- 2. Rowlings Text book of Pharmaceutics
- 3. Tutorial Pharmacy Cooper & Gunn

- 1. Remington's Pharmaceutical Sciences
- 2. USP/BP/IP

24PD401T

FOURTH YEAR

PHARMACOTHERAPEUTICS III - THEORY 4H 4C

Instruction hours/ week: L: 3 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory : 4 Hrs. /Week

Course Objectives:

- This course is designed to impart knowledge and skills necessary for contribution to quality use of medicines.
- Chapters dealt cover briefly pathophysiology and mostly therapeutics of various diseases.
- This will enable the student to understand the pathophysiology of common diseases and their management.
- To summarize the therapeutic approach to management of these diseases including reference to the latest available evidence;
- To diagnose a disease using biochemical & pathological Parameters.
- To discuss the preparation of individualized therapeutic plans based on diagnosis

Course Outcome (CO's):

On successful completion of the course the student will

COs	Course Outcomes	Blooms Level
CO1	Explain Pathophysiology of selected disease states and the rationale	Evaluating
	for drug therapy;	
CO2	Understand the therapeutic approach to manage of these diseases;	Understanding
CO3	Diagnose a dynamic based on biochemical &Pathological Parameters;	Analyzing
CO4	Evaluate the importance of individualized therapeutic plans based on	Evaluating
	diagnosis;	
CO5	Identify the patient-specific parameters relevant in initiating drug	Applying
	therapy, and monitoring drug therapy (including alternatives, time-	
	course of clinical andlaboratory indices of therapeutic response and	
	adverse effects);	
CO6	Demonstrate patient counselling points for various diseases;	Understanding

Mapping with Programme Outcomes

COs	PO1	PO2	PO3	PO4	P05	PO6	PO7	PO8	PO9	PO10	PO11
CO1	М										
CO2			S								
CO3			S								
CO4						Μ					
CO5			Μ								
CO6								М			

S-Strong; M-Medium; L-Low

Etiopathogenesis and pharmacotherapy of diseases associated with following systems/ diseases:

UNIT I

• Gastrointestinal system: Peptic ulcer disease, Gastro Esophageal Reflux Disease

UNIT II

• Inflammatory bowel disease

UNIT III

• Liver disorders - Alcoholic liver disease, Viral hepatitis including jaundice

UNIT IV

• Drug induced liver disorders.

UNIT V

• Haematological system: Anaemias, Venous thromboembolism

UNIT VI

• Drugs induced blood disorders.

UNIT VII

• Nervous system: Epilepsy, Parkinsonism, Stroke, Alzheimer's disease,

UNIT VIII

• **Psychiatry disorders:** Schizophrenia, Affective disorders, Anxiety disorders, Sleepdisorders, Obsessive Compulsive disorders

UNIT IX

• Pain management including Pain pathways, neuralgias, headaches.

UNIT X

• Evidence Based Medicine

The following details shall be submitted for any 25 selected drugs in the form of assignment/PPT.

**Name of the Drug, Brand name, Category, Therapeutic uses, Dose, Pharmacological action, Route of administration, Side effects, Contraindications, Drug Interactions with drug/food, Pharmacokinetics

Suggested Readings:

- 1. Clinical Pharmacy and Therapeutics Roger and Walker, Churchill Livingstone publication
- Pharmacotherapy: A Pathophysiologic approach Joseph T. Dipiro et al. Appleton & Lange

- 1. Pathologic basis of disease Robins SL, W.B.Saunders publication
- 2. Pathology and therapeutics for Pharmacists: A Basis for Clinical Pharmacy Practice -Green and Harris, Chapman and Hall publication
- Clinical Pharmacy and Therapeutics Eric T. Herfindal, Williams and Wilkins Publication
- Applied Therapeutics: The clinical Use of Drugs. Lloyd Young and Koda -Kimble MA
- 5. Avery's Drug Treatment, 4th Edn, 1997, Adis International Limited.

24PD402T

HOSPITAL PHARMACY - THEORY

Instruction hours/ week: L: 2 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory : 3 Hrs. /Week

Course Objectives:

- In the changing scenario of pharmacy practice in India, for successful practice of Hospital Pharmacy,
- the students are required to learn various skills like drug distribution,
- to know drug dispensing, manufacturing of parenteral preparations,
- to understand drug information, patient counselling,
- to learn therapeutic drug monitoring for improved patient care.
- To know The Stores Management and Inventory Control.

Course Outcome (CO's):

Upon completion of the course, the student will -

COs	Course Outcomes	Blooms Level
CO1	Outline on hospital pharmacy Organize & understand the	Knowledge
	managementof materials & finance;	
CO2	Understand budget prepare and implementation in Hospital	Knowledge
	Pharmacy;	
CO3	Illustrate the Stores Management and Inventory Control.	Knowledge
CO4	Discuss various drug distribution methods in hospital & for	Evaluate
	scheduleddrugs;	
CO5	Describe the manufacturing practices of various formulations	Knowledge
	inhospitalset up & radio Pharmaceuticals;	
CO6	Understand the professional practice management skills in	Analyse
	hospitalpharmacies;.	

Mapping with Programme Outcome

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1	S	S					S		Μ		S
CO2		S	S	S		М	S		Μ		
CO3						S	S		S		S
CO4		S		S			S	М			
CO5	М	М					S	L			
CO6	М	М			S		S	L			

S-Strong; M-Medium; L-Low

3Н 3С

FOURTH YEAR

UNIT I

Hospital - its Organization and functions

UNIT II

Hospital pharmacy-Organization and management

- a) Organizational structure-Staff, Infrastructure & work load statistics
- b) Management of materials and finance
- c) Roles & responsibilities of hospital pharmacist

UNIT III

The Budget - Preparation and implementation

Hospital drug policy

- a) Pharmacy and Therapeutic committee (PTC)
- b) Hospital formulary

UNIT IV

Hospital committees

- Infection committee
- Research and ethical committee
- c) developing therapeutic guidelines
- d) Hospital pharmacy communication Newsletter

UNIT V

Hospital pharmacy services

- Procurement & warehousing of drugs and Pharmaceuticals
- Inventory control
- Definition, various methods of Inventory Control ABC, VED, EOQ, Lead time, safety stock

UNIT VI

Drug distribution in the hospital

- Individual prescription method
- Floor stock method
- Unit dose drug distribution method

UNIT VII

Distribution of Narcotic and other controlled substances

Central sterile supply services - Role of pharmacist

UNIT VIII

Manufacture of Pharmaceutical preparations

Sterile formulations - large and small volume parenteralsmanufacture

Of Ointments, Liquids, and cream, Pharmaceutical compounding

Sterile preparations: Sterile compounding, Good Compounding Practices, Responsibility of Compounding personnel.

UNIT IX

Manufacturing of Tablets, granules, capsules, and powders

Total parenteral nutrition

Continuing professional development programs

Education and training

Radio Pharmaceuticals – Handling and packaging

UNIT X

Professional Relations and practices of hospital pharmacist

Suggested Readings:

- 1. Hospital pharmacy by William E. Hassan
- A text book of Hospital Pharmacy by S.H.Merchant & Dr. J.S. Qadry. Revised by R.K.Goyal & R.K. Parikh

- 1. WHO consultative group report.
- 2. R.P.S. Vol.2. Part –B; Pharmacy Practice section.
- 3. Handbook of pharmacy health care. Edt. Robin J Harman. The Pharmaceutical press.

Pharm.D	2024-25
24PD403T	FOURTH YEAR

CLINICAL PHARMACY - THEORY 4H 4C

Instruction hours/ week: L: 3 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory : 4 Hrs.

/WeekCourse

objective:

- To Monitor drug therapy of patient through medication chart review and clinical review;
- To Obtain medication history interview and counsel the patients;
- To Identify and resolve drug related problems;
- To detect, assess and monitor adverse drug reaction;
- To Interpret selected laboratory results (as monitoring parameters in therapeutics) of specific disease states; and
- To Retrieve, analyse, interpret and formulate drug or medicine information.

Course Outcomes (CO's):

On successful completion of the course the student will

Cos	Course Outcomes	Blooms Level
CO1	Describe the daily activities of clinical Pharmacist in	Understand
	improving the patient case.	
CO2	Understand the basis for evaluating the biomedical	Evaluate
	literatures;	
CO3	Identify and resolve drug related problems;	Apply
CO4	Detect, assess and monitor adverse drug reaction;	Apply
CO5	Interpret selected laboratory results (as monitoring parameters	Evaluate
	in therapeutics) of specific disease states;	
CO6	Retrieve, analyse, interpret and formulate drug or medicine	Create
	information.	

Mapping with Programme Outcomes

POs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	S									S	S
CO2	S										
CO3	S			S			S				
CO4	S		S							S	Μ
CO5	S		S				S			Μ	
CO6	S	S	S		S	Μ	S			Μ	

S-Strong; M-Medium; L-Low

Detailed syllabus and lecture wise schedule:

UNIT I

Definitions, development and scope of clinical pharmacy

Introduction to daily activities of a clinical pharmacist

- a. Drug therapy monitoring (medication chart review, clinical review, pharmacist interventions)
- b. Ward round participation
- c. Adverse drug reaction management
- d. Drug information and poisons information

UNIT II

- Medication history
- Patient counseling

UNIT III

• Drug utilisation evaluation (DUE) and review (DUR)

UNIT IV

• Quality assurance of clinical pharmacy services

UNIT V

• Patient data analysis

The patient's case history, its structure and use in evaluation of drug therapy & Understanding common medical abbreviations and terminologies used in clinical practices.

• Clinical laboratory tests used in the evaluation of disease states, and interpretation oftest results

- a. Haematological, Liver function, Renal function, thyroid function tests
- b. Tests associated with cardiac disorders
- c. Fluid and electrolyte balance
- d. Microbiological culture sensitivity tests
- e. Pulmonary Function Tests

UNIT VI

- Drug & Poison information
 - Introduction to drug information resources available
 - Systematic approach in answering DI queries

- Critical evaluation of drug information and literature
- Preparation of written and verbal reports
- Establishing a Drug Information Centre
- Poisons information- organization & information resources

UNIT VII

Pharmacovigilance

- Scope, definition and aims of pharmacovigilance
- Adverse drug reactions Classification, mechanism, predisposing factors, causality assessment [different scales used]
- Reporting, evaluation, monitoring, preventing & management of ADRs
- Role of pharmacist in management of ADR.

UNIT VIII

• Communication skills, including patient counselling techniques, medication historyinterview, presentation of cases.

UNIT IX

- Pharmaceutical care concepts
- Critical evaluation of biomedical literature

UNIT X

Medication errors

Suggested Readings:

- 1. Practice Standards and Definitions The Society of Hospital Pharmacists of Australia.
- 2. Basic skills in interpreting laboratory data Scott LT, American Society of Health System Pharmacists Inc.
- 3. Biopharmaceutics and Applied Pharmacokinetics Leon Shargel, Prentice Hall publication.
- A text book of Clinical Pharmacy Practice; Essential concepts and skills, Dr.G.Parthasarathi etal, Orient Orient Langram Pvt. Ltd. ISSBN8125026

- 1 Australian drug information -Procedure manual. The Society of Hospital Pharmacists of Australia.
- 2 Clinical Pharmacokinetics Rowland and Tozer, Williams and Wilkins Publication.
- Pharmaceutical statistics. Practical and clinical applications. Sanford Bolton, MarcelDekker, Inc.

24PD404T

FOURTH YEAR

BIOSTATISTICS AND RESEARCH METHODOLOGY - THEORY 3H 3C

Instruction hours/ week: L: 2 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory : 3 Hrs. /WeekCourse Objectives:

- To understand some basic concepts of research and its methodologies
- To identify appropriate research topics
- To select and define appropriate research problem and parameters
- To prepare a project proposal
- To organize and conduct research in a more appropriate manner
- To write a research report and thesis

Course Outcomes (CO's):

A successful completion of this subject it is expected that students will be able to

COs	Course Outcomes	Blooms Level
CO1	Understand some basic concepts of research and its	Understand
	methodologies.	
CO2	Execute computer applications in Hospital Pharmacy.	Knowledge
CO3	Select and define appropriate research problem and parameters.	Knowledge
CO4	Construct various data graphics in research.	Knowledge
CO5	Organize and conduct research in a more appropriate manner.	Apply
CO6	Implement statistical parameter in research.	Apply

Mapping with Programme Outcomes:

Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12
CO1	S											
CO2			S									
CO3					М							
CO4								М				
CO5											S	
CO6			S									

S-Strong; M-Medium; L-Low

Course Content:

UNIT I

Research Methodology

- Types of clinical study designs:
- Case studies, observational studies, interventional studies,
- Designing the methodology
- Sample size determination and Power of a study
- Determination of sample size for simple comparative experiments, determination of sample size to obtain a confidence interval of specifiedwidth, power of a study
- Report writing and presentation of data

UNIT II

Biostatistics

- Introduction
- Types of data distribution
- Measures describing the central tendency distributions- average, median, mode
- Measurement of the spread of data-range, variation of mean, standard deviation, variance, coefficient of variation, standard error of mean.

UNIT III

Data graphics

• Construction and labeling of graphs, histogram, piecharts, scatter plots, semilogarthimic plots

UNIT IV

- Basics of testing hypothesis
 - Null hypothesis, level of significance, power of test, P value, statisticalestimation of confidence intervals.
 - Level of significance (Parametric data)- students t test (paired and unpaired), chi Square test, Analysis of Variance (one-way and two-way)
 - Level of significance (Non-parametric data)- Sign test, Wilcoxan's signed rank test, Wilcoxan rank sum test, Mann Whitney U test, Kruskal-Wall is test (one way ANOVA)
 - Linear regression and correlation- Introduction, Pearsonn's and Spearmann's correlation and correlation co-efficient.
 - Introduction to statistical software: SPSS, Epi Info, SAS.

UNIT V

• Statistical methods in epidemiology

Incidence and prevalence, relative risk, attributable risk

UNIT VI

• Computer applications in pharmacy

UNIT VII

• Computer System in Hospital Pharmacy:

Patterns of Computer use in Hospital Pharmacy –Patient record database management, Medication order entry – Drug labels and list – Intravenous solution and admixture, patient medication profiles, Inventory control, Management report & Statistics.

UNIT VIII

• Computer In Community Pharmacy

Computerizing the Prescription Dispensing process

Use of Computers for Pharmaceutical Care in community pharmacy

UNIT IX

• Accounting and General ledger system

UNIT X

• Drug Information Retrieval & Storage : Introduction – Advantages of Computerized Literature Retrieval

Use of Computerized Retrieval

Suggested Readings:

1. Pharmaceutical statistics- Practical and clinical applications, Sanford

Bolton 3rd edition, publisher Marcel Dekker Inc. NewYork.

Reference Books (Latest Editions):

1. Drug Information- A Guide for Pharmacists, Patrick M Malone, Karen L Kier, John E

Stanovich , 3rd edition, McGraw Hill Publications 2006

FOURTH YEAR

BIOPHARMACEUTICS AND PHARMACOKINETICS - THEORY 4H 4C

Instruction hours/ week: L: 3 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 4 Hrs. /Week

Course Objectives:

- This subject is designed to impart knowledge and skills of Biopharmaceutics and pharmacokinetics and their applications in pharmaceutical development, design of dose and dosage regimen and in solving the problems raised therein.
- To explain the use of plasma drug concentration-time data to calculate the pharmacokinetic parameters.
- To understand the concepts of bioavailability and bioequivalence of drug products and their significance.
- To understand various pharmacokinetic parameters, their significance & applications.
- To demonstrate a clear information on compartmental models and methods to assess the models.
- To describe the kinetics of drug absorption, distribution, metabolism, excretion, elimination.

COs	Course Outcomes	Blooms Level
CO1	Understand the basic concepts in biopharmaceutics and	Understand
	pharmacokinetics and their significance.	
CO2	Explain the use of plasma drug concentration-time data	Understand
	to calculate the pharmacokinetic parameters.	
CO3	Understand the concepts of bioavailability and	Understand
	bioequivalence of drug products and their significance.	
CO4	Understand various pharmacokinetic parameters, their	Analyze
	significance & applications.	
CO5	Demonstrate a clear information on compartmental	Apply
	models and methods to assess the models.	
CO6	Describe the kinetics of drug absorption, distribution,	Understand
	metabolism, excretion, elimination.	

Course Outcomes (CO's): At the end of this course, students will be able to

Mapping with Programme Outcomes

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1		S					Μ				
CO2		S					Μ		Μ	L	Μ
CO3	S		S	S	Μ	Μ	Μ	S	Μ		Μ
CO4		Μ	Μ	Μ	Μ	Μ	Μ		Μ	Μ	S
CO5			Μ	S			S		Μ		Μ
CO6			Μ	Μ	Μ						

S-Strong; M-Medium; L-Low

UNIT I

Biopharmaceutics

Introduction to Biopharmaceutics

- Absorption of drugs from gastrointestinal tract.
- Drug Distribution.
- Drug Elimination.

UNIT II

Introduction to Pharmacokinetics.

- Mathematical model
- Drug levels in blood.
- Pharmacokinetic model
- Compartment models
- Pharmacokinetic study.

UNIT III

One compartment open model.

- Intravenous Injection (Bolus)
- Intravenous infusion.

UNIT IV

Multicompartment models.

- Two compartment open model.
- IV bolus, IV infusion and oral administration

UNIT V

Multiple – Dosage Regimens.

- Repititive Intravenous injections One Compartment Open Model
- Repititive Extravascular dosing One Compartment Open model
- Multiple Dose Regimen Two Compartment Open Model

Nonlinear Pharmacokinetics.

- Introduction
- Factors causing Non-linearity.
- Michaelis-menton method of estimating parameters.

UNIT VII

Non-compartmental Pharmacokinetics.

- Statistical Moment Theory.
- MRT for various compartment models.

UNIT VIII

• Physiological Pharmacokinetic model.

UNIT IX

Bioavailability and Bioequivalence.

- Introduction.
- Bioavailability study protocol.

UNIT X

• Methods of Assessment of Bioavailability

Suggested Readings:

- 1 Biopharmaceutics and Clinical Pharmacokinetics by, Milo Gibaldi
- 2 Remington's Pharmaceutical Sciences, By Mack Publishing Company, Pennsylvnia.
- 3 Pharmacokinetics: By Milo Glbaldi Donald, R. Mercel Dekker Inc.
- 4 Hand Book of Clinical Pharmacokinetics, By Milo Gibaldi and Laurie Prescott by ADIS Health Science Press.
- 5 Biopharmaceutics and Pharmacokinetics; By Robert F Notari

- 1. Biopharmaceutics; By Swarbrick
- Bio pharmaceutics and Pharmacokinetics-A Treatise, By D. M. Brahmankar and Sunil B.Jaiswal, Vallabh Prakashan Pitampura, Delhi
- 3. Cilincal Pharmacokinetics, Concepts and Applications: By Malcolm Rowland and Thomas, N. Tozen, Lea and Febrger, Philadelphia, 1995.

24PD406T

CLINICAL TOXICOLOGY - THEORY 3H

Instruction hours/ week: L: 2 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory : 3 Hrs. /Week Course Objectives:

- To understand the general working knowledge of the principles and practice of clinical toxicology
- To know the health implications of toxic exposures and commonly involved chemicals for toxicity
- To understand principles of general toxicology and clinical management practice.
- To demonstrate and apply history, assessment and therapy considerations associated with the management of a toxic exposure.
- To understand treatment guidelines for specific toxic substances.
- To get knowledge about preventive approaches to reduce unintentional poisoning.

Course Outcomes (CO's):

On successful completion of the course the student will

	Course Outcomes	Blooms Level	
COs			
CO1	Understandthegeneralworkingprinciplesandpracticeofclin icaltoxicology	Understand	
CO2	The health implications of toxicexposures and commonly involved chemicals for toxicity	Apply	
CO3	Understand principles of general toxicology and clinical management practice.	Understand	
CO4	Demonstrate the assessment and therapy considerations associated with the management of a toxic exposure.	Apply	
CO5	Understand treatment guidelines for specific toxic substances.	Understand	
CO6	Knowledge about preventive approaches to reduce unintentional poisoning	Knowledge	

Mapping with Programme Outcomes

POs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
COs											
CO1	S										
CO2	S										
CO3	S										
CO4	S										
CO5	S										
CO6	S										

S-Strong; M-Medium; L-Low

2024-25

3C

FOURTH YEAR

UNIT I

- General principles involved in the management poisoning
- Antidotes and the clinical applications.
- Supportive care in clinical Toxicology.

UNIT II

- Gut Decontamination.
- Elimination Enhancement.
- Toxicokinetics.

UNIT III

Clinical symptoms and management of acute poisoning with the following agents -

- Pesticide poisoning: organophosphorous compounds, carbamates, organochlorines, pyrethroids.
- Opiates overdose.
- Antidepressants
- Barbiturates and benzodiazepines.

UNIT IV

Clinical symptoms and management of acute poisoning with the following agents – Alcohol: ethanol, methanol.

- Paracetamol and salicylates.
- Non-steroidal anti-inflammatory drugs.

UNIT V

- Clinical symptoms and management of acute poisoning with the following agents –Hydrocarbons: Petroleum products and PEG.
- Caustics: inorganic acids and alkali.
- Radiation poisoning

UNIT VI

• Clinical symptoms and management of chronic poisoning with the following agents - Heavymetals: Arsenic, lead, mercury, iron, copper

UNIT VII

 Clinical symptoms and management of acute poisoning with the following agents – Venomous snake bites: Families of venomous snakes, clinical effects of venoms, generalmanagement as first aid, early manifestations, complications and snake bite injuries.

UNIT VIII

- Plants poisoning. Mushrooms, Mycotoxins.Food poisonings
- Envenomations Arthropod bites and stings.

UNIT IX

Substance abuse:

Signs and symptoms of substance abuse and treatment of dependence

- CNS stimulants :amphetamine
- Opioids
- CNS depressants

UNIT X

Hallucinogens: LSD

- Cannabis group
- Tobacco

The following details shall be submitted for any 25 selected drugs in the form of assignment/PPT.

**Name of the Drug, Brand name, Category, Therapeutic uses, Dose, Pharmacological action, Route of administration, Side effects, Contraindications, Drug Interactions with drug/food, Pharmacokinetics.

Suggested Readings:

- 1. Biopharmaceutics and Clinical Pharmacokinetics by, Milo Gibaldi
- Matthew J Ellenhorn. Ellenhorns Medical Toxicology Diagnosis and Treatment of Poisoning. Second edition. Williams and Willkins publication, London

Reference Books (Latest Editions):

 V V Pillay. Handbook of Forensic Medicine and Toxicology. Thirteenth edition 2003 Paras Publication, Hyderabad

Pharm.D

24PD407P

Instruction hours/ week: L: 0 T:0 P:3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Practical: 3 Hrs./Week

Course Objectives:

- This course is designed to impart knowledge and skills necessary for contribution to quality use of medicines.
- Chapters dealt cover briefly pathophysiology and mostly therapeutics of various diseases.
- This will enable the student to understand the pathophysiology of common diseases and their management.
- To summarize the therapeutic approach to management of these diseases including reference to the latest available evidence;
- To discuss the controversies in drug therapy;
- To discuss the preparation of individualized therapeutic plans based on diagnosis

Course Outcome (CO's): On successful completion of the course the student will

COs	Course Outcomes	Blooms Level
CO1	Analyze individual cases for various diseases;	Analyzing
CO2	Present case reports for various diseases;	Analyzing
CO3	Demo Patient counselling points;	Understanding
CO4	Identify advance drug reactions;	Applying
CO5	Analyze various drug interactions;	Analyzing
CO6	Manage selection of drug therapy during ward rounds participations;	Evaluating

Mapping with Programme Outcomes

Cos	PO1	PO2	PO3	PO4	P05	PO6	PO7	PO8	PO9	PO10	PO11
CO1			S								
CO2			Μ					Μ			
CO3								Μ			
CO4			Μ								
CO5			Μ								
CO6			Μ								

S Strong; M-Medium; L-Low

Practicals:

Hospital postings for a period of at least 50 hours is required to understand the principles and practice involved in ward round participation and clinical discussion on selection of drug therapy. Students are required to maintain a record of 15 cases observed in the ward

and the same should be submitted at the end of the course for evaluation. Each student should present at least two medical cases they have observed and followed in the wards.

Assignments:

Students are required to submit written assignments on the topics given to them. Topics allotted should cover recent developments in drug therapy of various diseases. A minimum of THREE assignments [1500 - 2000 words] should be submitted for evaluation.

Format of the assignment:

- 1. Minimum & Maximum number of pages
- 2. Reference(s) shall be included at the end.
- 3. Assignment can be a combined presentation at the end of the academic year
- 4. It shall be computer draft copy
- 5. Name and signature of the student
- 6. Time allocated for presentation may be 8+2 Min.

Suggested Readings:

1. Clinical Pharmacy and Therapeutics - Roger and Walker, Churchill Livingstone publication

- 1. Pathologic basis of disease Robins SL, W.B.Saunders publication
- 2. Pathology and therapeutics for Pharmacists: A Basis for Clinical Pharmacy Practice -Green and Harris, Chapman and Hall publication
24PD408P

2024-25

FOURTH YEAR

HOSPITAL PHARMACY -PRACTICAL 3H 2C

Instruction hours/ week: L: 0 T:0 P:3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Practical : 3 Hrs./Week Course Objectives:

- In the changing scenario of pharmacy practice in India, for successful practice of Hospital Pharmacy,
- the students are required to learn various skills like drug distribution,
- to know drug dispensing, manufacturing of parenteral preparations,
- to understand drug information, patient counselling,
- to learn therapeutic drug monitoring for improved patient care.
- To know The Stores Management And Inventory Control.

Course Outcome (CO's):

Upon completion of the course, the student shall be able to -

COs	Course Outcomes	Blooms Level
CO1	Evaluate drug interactions in outpatient prescriptions;	Analyse
CO2	Evaluate drug interactions in ward prescription;	Analyse
CO3	Defending drug information queries;	Evaluate
CO4	Identify adverse drug reaction	Understand
CO5	Assessing drug inventory control in Hospital Pharmacy	Create
CO6	Assessing drug inventory control in Nursing Station.	Create

Mapping with Programme Outcome

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12
CO1	М	М	S		М	S	S	L			L	
CO2	М	М	S		М	S	S	L			L	
CO3	М	S			S	S	S	S			Μ	
CO4		S		S	S			S				
CO5	S	S	М		М	S		М	L			
CO6	S	S	М		М	S		М	L			

S-Strong; M-Medium; L-Low

COURSE CONTENT

- 1. Assessment of drug interactions in the given prescriptions
- 2. Drug information queries.
- 3. Inventory control

List of Assignments:

- 1. Design and Management of Hospital pharmacy department for a 300 bedded hospital.
- 2. Pharmacy and Therapeutics committee Organization, functions, and limitations.
- 3. Development of a hospital formulary for 300 bedded teaching hospital
- 4. Preparation of ABC analysis of drugs sold in one month from the pharmacy.
- 5. Different phases of clinical trials with elements to be evaluated.
- 6. Various sources of drug information and systematic approach to provide unbiased drug information.
- 7. Evaluation of prescriptions generated in hospital for drug interactions and find out the suitable management
- 8. Sterile compounding

Special requirements:

- 1. Each college should sign MoU with nearby local hospital having minimum 150 beds for providing necessary training to the students' on hospital pharmacy activities.
- 2. Well equipped with various resources of drug information.

Suggested Readings:

1. Hospital pharmacy by William E. Hassan

Reference Books (Latest Editions):

- 1. WHO consultative group report.
- 2. R.P.S. Vol.2. Part –B; Pharmacy Practice section.

Pharm.D

2024-25

FOURTH YEAR

CLINICAL PHARMACY - PRACTICAL 3H 2C

Instruction hours/ week: L: 0 T:0 P:3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Practical : 3 Hrs./Week

Course objective:

- To Monitor drug therapy of patient through medication chart review and clinicalreview
- To Obtain medication history interview and counsel the patients;
- To Identify and resolve drug related problems;
- To detect, assess and monitor adverse drug reaction;
- To Interpret selected laboratory results (as monitoring parameters in therapeutics) of specific disease states; and
- To Retrieve, analyse, interpret and formulate drug or medicine information.

Course Outcomes (CO's):

On successful completion of the course the student will

COs	Course Outcomes	Blooms Level
CO1	Demonstrate patient meditation history	Understand
CO2	Illustrate patient counseling interview;	Apply
CO3	Defending drug information queries;	Evaluate
CO4	Detect, assess and monitor adverse drug reaction;	Evaluate
CO5	Interpret selected laboratory results (as monitoring parameters in therapeutics) of specific disease states;	Apply
CO6	Analyze and interpret drug or medicine information.	Evaluate

Mapping with Programme Outcomes

Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
COs											
CO1	S	S	S	L		Μ	Μ			Μ	Μ
CO2	S	S	S		S	Μ		S	Μ	L	
CO3	S		S	Μ				Μ			Μ
CO4	S		S		L	L	L			S	Μ
CO5		S								Μ	
CO6	S	S	S		S	Μ	S			Μ	

S-Strong; M-Medium; L-Low

Students are expected to perform 15 Practicals in the following areas covering the topics dealt in Theory class.

- 1 Answering drug information questions (4 Nos)
- 2 Patient medication counseling (4 Nos)
- 3 Case studies related to laboratory investigations (4 Nos)
- 4 Patient medication history interview (3 Nos)

Assignment:

Students are expected to submit THREE written assignments (1500 - 2000 words) on the topics given to them covering the following areas dealt in Theory class.

Drug information, Patient medication history interview, Patient medication counselling, Critical appraisal of recently published articles in the biomedical literature which deals with a drug or therapeutic issue.

Format of the assignment:

- 1. Minimum & Maximum number of pages.
- 2. Reference(s) shall be included at the end.
- 3. Assignment can be a combined presentation at the end of the academic year.
- 4. It shall be computer draft copy.
- 5. Name and signature of the student.
- 6. Time allocated for presentation may be 8+2 Min.

Suggested Readings:

- 1. Practice Standards and Definitions The Society of Hospital Pharmacists of Australia.
- 2. A text book of Clinical Pharmacy Practice; Essential concepts and skills, Dr.G.Parthasarathi etal, Orient Orient Langram Pvt. Ltd. ISSBN8125026

Reference Books (Latest Editions):

- 1 Australian drug information -Procedure manual. The Society of Hospital Pharmacists of Australia.
- 2 Clinical Pharmacokinetics Rowland and Tozer, Williams and Wilkins Publication.

24PD410P

BIOPHARMACEUTICS AND PHARMACOKINETICS -PRACTICAL 3H

Instruction hours/ week: L: 0 T:0 P:3

Practical: 3 Hrs./Week

Course Objectives:

- This subject is designed to impart knowledge and skills of Biopharmaceutics and pharmacokinetics and their applications in pharmaceutical development, design of dose and dosage regimen and in solving the problems raised therein.
- To explain the use of plasma drug concentration-time data to calculate the pharmacokinetic parameters.
- To understand the concepts of bioavailability and bioequivalence of drug products and their significance.
- To understand various pharmacokinetic parameters, their significance & applications.
- To demonstrate a clear information on compartmental models and methods to assess the models.
- To describe the kinetics of drug absorption, distribution, metabolism, excretion, • elimination.

COs	Course Outcomes	Blooms Level
CO1	Understand the basic concepts in biopharmaceutics and	Analyze
	pharmacokinetics and their significance.	
CO2	Explain the use of plasma drug concentration-time data to calculate the pharmacokinetic parameters.	Analyze
CO3	Understand the concepts of bioavailability and bioequivalence of drug products and their significance.	Analyze
CO4	Understand various pharmacokinetic parameters, their significance & applications.	Analyze
CO5	Demonstrate a clear information on compartmental models and methods to assess the models.	Understand
CO6	Describe the kinetics of drug absorption, distribution, metabolism, excretion, elimination.	Understand

Mapping with Programme Outcomes

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1	S	Μ	Μ								
CO2	S	Μ	Μ	Μ					L		
CO3			S								
CO4	S	Μ		Μ					Μ		
CO5	S	Μ	S	Μ							
CO6	S	М	S	M							

S-Strong; M-Medium; L-Low



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FOURTH YEAR

Marks: Internal: 30 External: 70 Total: 100

External Semester Exam: 3 Hours

2C

- 1. Improvement of dissolution characteristics of slightly soluble drugs by some methods.
- 2. Comparison of dissolution studies of two different marketed products of same drug.
- 3. Influence of polymorphism on solubility and dissolution.
- 4. Protein binding studies of a highly protein bound drug and poorly protein bound drug.
- 5. Extent of plasma-protein binding studies on the same drug (i.e. highly and poorly protein bound drug) at different concentrations in respect of constant time.
- 6. Bioavailability studies of some commonly used drugs on animal/human model.
- 7. Calculation of Ka, Ke , t1/2, Cmax , AUC, AUMC, MRT etc. from blood profile data.
- 8. Calculation of bioavailability from urinary excretion data for two drugs.
- 9. Calculation of AUC and bioequivalence from the given data for two drugs.
- 10. In vitro absorption studies.
- 11. Bioequivalency studies on the different drugs marketed. (eg) Tetracycline, Sulphamethoxzole, Trimethoprim, Aspirin etc., on animals and human volunteers.
- 12. Absorption studies in animal inverted intestine using various drugs.
- 13. Effect on contact time on the plasma protein binding of drugs.
- 14. Studying metabolic pathways for different drugs based on elimination kinetics data.
- 15. Calculation of elimination half-life for different drugs by using urinary elimination data and blood level data.
- 16. Determination of renal clearance.

Suggested Readings:

- 1. Biopharmaceutics and Clinical Pharmacokinetics by, Milo Gibaldi
- 2. Biopharmaceutics and Pharmacokinetics; By Robert F Notari

Reference Books (Latest Editions):

- 1. Biopharmaceutics; By Swarbrick
- 2. Bio pharmaceutics and Pharmacokinetics-A Treatise, By D. M. Brahmankar and SunilB.Jaiswal, Vallabh Prakashan Pitampura, Delhi.

24PD501T

CLINICAL RESEARCH - THEORY

Instruction hours/ week: L: 3 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory : 4 Hrs. /Week Course Objectives:

- Understand various approaches to drug discovery
- Understand the regulatory and ethical process
- Describe the new drug development process
- Estimate and Discuss the clinical trials activities
- Estimate safety monitoring and reporting in clinical trials
- Identify preventive approaches to reduce unintentional poisoning.
- Related the roles & responsibilities of Clinical trial Personnels.

Course Outcomes (CO's):

On successful completion of the course the student will

COs	Course Outcomes	Blooms Level
CO1	Summarize the various approaches to drug discovery	Understanding
CO2	Explain the regulatory and ethical process	Understanding &
		Applying
CO3	Describe the new drug development process	Applying
CO4	Estimate and Discuss the clinical trials activities	Remembering&
		Applying
CO5	Estimate safety monitoring and reporting in clinical trials	Analyzing and
		Evaluating
CO6	Identify preventive approaches to reduce unintentional poisoning	Creating
CO7	Describe the roles & responsibilities of Clinical trial Personnel.	Understanding

Mapping with Programme Outcomes

Cos	PO1	PO2	PO3	PO4	P05	PO6	PO7	PO8	PO9	PO10	PO11
CO1	S	S		S	S				S	L	S
CO2	S		S		S	S	S		S		
CO3	S	S	S	S	Μ				S	М	S
CO4	S		М	S	Μ	S		S	М		L
CO5	S	М	S	М	Μ	М	S		S	S	
CO6	М	S	S			S		S	Μ		L
CO7	S		М			М	М	S	М	М	

S-Strong; M-Medium; L-Low

4C

FIFTH YEAR

4H

UNIT I

- Drug development process: IntroductionVarious Approaches to drug discovery
- Pharmacological
- Toxicological

UNIT II

- IND Application
- Drug characterization
- Dosage form

UNIT III

Clinical development of drug:

- Introduction to Clinical trials
- Various phases of clinical trial.
- Methods of post marketing surveillance

UNIT IV

Abbreviated New Drug Application submission.

- Good Clinical Practice ICH, GCP, Central drug standard control organisation (CDSCO) guidelines
- Challenges in the implementation of guidelines
- Ethical guidelines in Clinical Research

UNIT V

- Composition, responsibilities, procedures of IRB / IEC
- Overview of regulatory environment in USA, Europe and India.

UNIT VI

Role and responsibilities of clinical trial personnel as per ICH GCP

- Sponsor
- Investigators
- Clinical research associate
- Auditors
- Contract research coordinators
- Regulatory authority

UNIT VII

• Designing of clinical study documents (protocol, CRF, ICF, PIC with assignment)

UNIT VIII

• Informed consent Process

UNIT IX

• Data management and its components

UNIT X

• Safety monitoring in clinical trials.

Suggested Readings:

- Central Drugs Standard Control Organization. Good Clinical Practices-Guidelines for Clinical Trials on Pharmaceutical Products in India. New Delhi: Ministry of Health; 2001.
- 2 International Conference on Harmonisation of Technical requirements for registration of Pharmaceuticals for human use. ICH Harmonised Tripartite Guideline. Guideline for Good Clinical Practice.E6; May 1996.
- 3 Ethical Guidelines for Biomedical Research on Human Subjects 2000. Indian Council of Medical Research, New Delhi.
- 4 Textbook of Clinical Trials edited by David Machin, Simon Day and Sylvan Green, March 2005, John Wiley and Sons.

Reference Books (Latest Editions):

- 1. Principles of Clinical Research edited by Giovanna di Ignazio, Di Giovanna and Haynes.
- 2. Clinical Data Management edited by R K Rondels, S A Varley, C F Webbs. Second Edition, Jan 2000, Wiley Publications.

FIFTH YEAR

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

PHARMACOEPIDEMIOLOGY AND PHARMACOECONOMICS -THEORY 4H 4C

Instruction hours/ week: L: 3 T: 1 P: 0

Theory: 4 Hrs. /Week Course Objectives:

- Identify the application pharmacoepidemiology and pharmacoeconomics in clinicalsettings
- Discuss the various pharmacoepidemiology outcome measures.
- Understand the concept of risk in pharmacoepidemiology and different methods of measuring risk
- Examine various pharmacoepidemiological methods
- Interpret various types case studies
- Asses about current pharmacoenomic evaluation methods

Course Outcomes (CO's):

On successful completion of the course the student will

COs	Course Outcomes	Blooms Level
CO1	Identify the application pharmacoepidemiology and	Analyse
	pharmacoeconomics in clinical setting	
CO2	Discuss the various pharmacoepidemiology outcome	Evaluate
	measures.	
CO3	Understand the concept of risk in	Knowledge
	pharmacoepidemiology and different methods of	
	measuring risk	
CO4	Examine various pharmacoepidemiological methods	Analyse
CO5	Interpret various types case studies	Evaluate
CO6	Asses about current pharmacoenomic evaluation methods	Create

Mapping with Programme Outcome

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1					Μ	S		S		S	М
CO2	S	S		S						S	S
CO3	S	Μ			Μ						S
CO4	S	Μ				L	Μ				S
C05	M	S	S			S	S	S	S	Μ	Μ
CO6	Μ			S	S	S	S	Μ	S	Μ	М

S-Strong; M-Medium; L-Low

UNIT I

Pharmacoepidemiology:

- Definition and scope:
- Origin and evaluation of pharmacoepidemiology need for pharmacoepidemiology, aims and applications.

UNIT II

• Measurement of outcomes in pharmacoepidemiology Outcome measure and drug use measures Prevalence, incidence and incidence rate. Monetary units, number of prescriptions, units of drugs dispensed, defined daily doses and prescribed daily doses, medication adherence measurement

UNIT III

- Concept of risk in pharmacoepidemiology
- Measurement of risk, attributable risk and relative risk, time-risk relationship and odds ratio

UNIT IV

Pharmacoepidemiological methods

• Includes theoretical aspects of various methods and Practical study of variousmethods with the help of case studies for individual methods

UNIT V

• Drug utilization review, case reports, case series, surveys of drug use, cross – sectional studies, cohort studies, case control studies, case –cohort studies, meta –analysis studies, spontaneous reporting, prescription event monitoring and record linkagesystem.

UNIT VI

• Sources of data for pharmacoepidemiological studies Ad Hoc data sources and automated data systems

UNIT VII

• Selected special applications of pharmacoepidemiology

Studies of vaccine safety, hospital pharmacoepidemiology, pharmacoepidemiology and risk management, drug induced birth defects.

UNIT VIII

Pharmacoeconomics:

- Definition, history, needs of pharmacoeconomic evaluations
- Role in formulary management decisions

UNIT IX

Pharmacoeconomic evaluation

- Outcome assessment and types of evaluation
- Includes theoretical aspects of various methods and Practical study of variousmethods with the help of case studies for individual methods:
- Cost minimization, cost- benefit, cost effectiveness, cost utility

UNIT X

• Applications of Pharmacoeconomics Software and case studies

Suggested Readings:

- Rascati K L. Essentials of Pharmacoeconomics, 2nd ed. Philadelphia: Woulters Kluwer Lippincott Williams & Wilkins, 2013
- Thomas E Getzen. Health economics. Fundamentals and Flow of Funds. 1997, 2003. John Wiley & Sons, second edition
- Andrew Briggs, Karl Claxton, Mark Sculpher. Decision Modelling for Health Economic Evaluation, Published by the Oxford University Press2006

Reference Books

- MichaelDrummond, MarkSculpher,George Torrence, BernieO'BrienandGreg Stoddart. Methods for the Economic Evaluation of Health Care Programmes Oxford University Press (Third Edition) – 2005.
- 2. George E Mackinnon III. Understanding health outcomes and pharmacoeconomics, 2013.

24PD503T FIFTH YEAR CLINICAL PHARMACOKINETICS AND PHARMACOTHERAPEUTIC DRUG MONITORING -THEORY 3H 3C

Instruction hours/ week: L: 2 T:1 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 3 Hrs. /Week

Course Objectives:

- This subject is designed to impart knowledge and skills of Biopharmaceutics and pharmacokinetics and their applications in pharmaceutical development, design of dose and dosage regimen and in solving the problems arised therein.
- To explain the use of plasma drug concentration-time data to calculate the pharmacokinetic parameters.
- To understand the concepts of bioavailability and bioequivalence of drug products and their significance.
- To understand various pharmacokinetic parameters, their significance & applications.
- To demonstrate a clear information on compartmental models and methods to assess the models.
- To describe the kinetics of drug absorption, distribution, metabolism, excretion, elimination.

Course Outcomes (CO's):

On successful completion of the course the student will

COs	Course Outcomes	Blooms Level
CO1	Understand the basic concepts in biopharmaceutics and pharmacokinetics and their significance.	Understand
CO2	Demonstrate compartmental models its methods.	Understand
CO3	Describe Therapeutic drug monitoring of drugs in various disease conditions	Apply
CO4	Discuss various pharmacokinetic parameters, their significance & applications.	Apply
CO5	Evaluate dosage adjustment in renal and hepatic impairment	Apply
CO6	Discuss genetic polymorphisim in drug metabolism	Knowledge

Mapping with Programme Outcomes

Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12
CO1	S											
CO2	S			М								
CO3			М									
CO4								М			М	
CO5	S					М			М			
CO6									S			

S-Strong; M-Medium; L-Low

UNIT I

• Introduction to Clinical pharmacokinetics.

UNIT II

- Design of dosage regimens:
- Nomograms and Tabulations in designing dosage regimen, Conversion from intravenous to oral dosing, Determination of dose and dosing intervals, Drug dosing in the elderly and pediatrics and obese patients.

UNIT III

Pharmacokinetics of Drug Interaction:

- Pharmacokinetic drug interactions
- Inhibition and Induction of Drug metabolism
- Inhibition of Biliary Excretion.

UNIT IV

Therapeutic Drug monitoring:

- Introduction
- Individualization of drug dosage regimen (Variability Genetic, Age and Weight, disease, Interacting drugs).
- Indications for TDM. Protocol for TDM.

UNIT IV

Pharmacokinetic/Pharmacodynamic Correlation in drug therapy.

• TDM of drugs used in the following disease conditions: cardiovascular disease, Seizure disorders, Psychiatric conditions, and Organ transplantations.

UNIT V

Dosage adjustment in Renal and hepatic Disease.

• Renal impairment, Pharmacokinetic considerations

Pharm.D

UNIT VI

General approach for dosage adjustment in Renal disease.

- Measurement of Glomerular Filtration rate and creatinine clearance.
- Dosage adjustment for uremic patients.

UNIT VII

- Extracorporeal removal of drugs.
- Effect of Hepatic disease on pharmacokinetics.

UNIT VIII

Population Pharmacokinetics.

- Introduction to Bayesian Theory.
- Adaptive method or Dosing with feed back.
- Analysis of Population pharmacokinetic Data.

UNIT IX

Pharmacogenetics

- Genetic polymorphism in Drug metabolism: Cytochrome P-450 Isoenzymes.
- Genetic Polymorphism in Drug Transport and Drug Targets.

UNIT X

• Pharmacogenetics and Pharmacokinetics / Pharmacodynamic considerations

The following details shall be submitted for any 25 selected drugs in the form of assignment/PPT.

**Name of the Drug, Brand name, Category, Therapeutic uses, Dose, Pharmacological action, Route of administration, Side effects, Contraindications, Drug Interactions with drug/food, Pharmacokinetics

Suggested Readings:

- 1. Leon Shargel, Susanna Wu-Pong, Andrew Yu. Applied Biopharmaceutics & Pharmacokinetics. 6th edition. New York: Mc Graw Hill; 2012.
- 2. Peter L. Bonate. Pharmacokinetic Pharmacodynamic Modeling and Simulation. 2nd edition. USA: Springer; 2011.
- Michael E. Burton, Leslie M. Shaw, Jerome J. Schentag, William E.Evans. Applied Pharmacokinetics & Pharmacodynamics: Principles of Therapeutic Drug Monitoring.4th edition. US: Lippincott Williams & Wilkins; 2005.

Reference Books (Latest Editions):

- Steven How-Yan Wong, Irving Sunshine. Handbook of Analytical Therapeutic Drug Monitoring and Toxicology. 1st edition. USA: CRC Press; 1996.
- 2. Soraya Dhillon, Andrzej Kostrzewski. Clinical pharmacokinetics. 1st edition. London: Pharmaceutical Press;2006.

24PD504S

2024-25

FIFTH YEAR

CLERKSHIP

1H 1C

Instruction hours/ week: L:0 T:0 P:1

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Seminar: 1 Hrs. /Week

SEMINAR & CASE PRESENTATION ASSESSMENT RUBRIC

Ware	d:		Date:					
Title	of presentation:							
Nam	e:	ID No:				Group		
Scop	be of Evaluation			_				
				Po	oor		Excelle	ent
Obje	ectives			1	2	3	4	5
1.	Provide clear objectives and able to ans the end of the presentation.	wer them a	t	1	2	3	4	5
Case	e Summary							
2.	Case presented with adequate informati chief complaints and histories.	on on bio-d	ata,	1	2	3	4	5
3.	Case presented with adequate informati of systems, lab investigations and revie course.	on on revie w of hospit	w al	1	2	3	4	5
Eval	luation of drug therapy							
4.	Able to identify, describe and explain the of patient-specific PCIs.	ne significat	nce	1	2	3	4	5
5.	Able to assess through the hospital cour important occurrences, therapeutic inter appropriate subjective and objective da demonstrate the therapeutic efficacy or	rse, any vention, ta to toxicity		1	2	3	4	5
6.	Able to correlate pharmacokinetics (AD drugs which may affect the patient's the	OME) of the erapy.		1	2	3	4	5
7.	Able to recommend how the manageme improved. (Drug choice, monitoring parameters, th outcome, patient education)	ent may be		1	2	3	4	5
App	lication of literature or references to c	ase discuss	ion				•	
8.	Able to select updated and relevant liter correlate the finding to the discussion.	ature and		1	2	3	4	5
Deli	very of presentation					1		
9.	Able to present in a professional manne appropriate flow in the allocated time.	r with		1	2	3	4	5
10.	Able to handle Q&A session and rationa answer.	alize the		1	2	3	4	5
]	Fotal	=	/5	0 = 10	0%	/

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Sign and stamp of examiner

24PD505P

PROJECT WORK

FIFTH YEAR **20H 10C**

Instruction hours/ week: L:0 T:0 P:20

Practicals : 20 Hrs. /Week

Marks: External: 100 Total: 100 External Semester Exam: 3 Hours

	Excellent (4)	Good (3)	Average (2)	Poor (1)
Organization	 Writing shows high degree of attention to logic and reasoning of points. Unity clearly leads the reader to the conclusion and stirs thought regarding the topic 	 Writing is coherent and logically organized with transitions used between ideas and paragraphs to create coherence. Overall unity of ideas is present 	 Writing is coherent and logically organized. Some points remain misplaced and stray from the topic. Transitions evident but not used throughout essay. 	 Writing lacks logical organization. It shows some coherence but ideas lack unity. Serious errors.
Level of Content	• Content indicates synthesis of ideas, in- depth analysis and evidences original thought and support for the topic.	 Content indicates original thinking Develops ideas with sufficient and firm evidence 	• Content indicates thinking and reasoning applied with original thought on a few ideas.	• Shows some thinking and reasoning but most ideas are underdeveloped and unoriginal.
Conclusion	• Conclusion is a concise, well- written summary of the argument	• Conclusion is somewhat related to the thesis and argument.	• There is a conclusion but it is not obviously related to the thesis or argument	No conclusion
References and citation	• Uniform style of referencing and updated information provided	• Few different style of referencing and quite updated references	• Noticeable different style of referencing and only few updated references	Wrong or different style of referencing.Outdated references
Format	• Meets all formal and assignment requirements and evidences attention to detail; all margins, spacing and indentations are correct; essay is neat and correctly assembled with professional look.	 Meets format and assignment requirements; margins, spacing and indentations are correct Essay is neat and correctly assembled. 	 Meets format and assignment requirements; generally correct margins, spacing and indentations Essay is neat but may have some assembly errors 	 Fails to follow format and assignment requirements. incorrect margins, spacing and indentation. neatness of essay

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Sign and stamp of examiner

S No	Category	Course code	Course title	No of	Credit	Year in which
				hrs	points	course can be taken
1.	Skills	24PD113ET	Communication	4	2	Ι
			Skills Theory			
2.	Human Value	24PD114ET	Yoga for Youth	4	2	Ι
			Empowerment			
			Theory			
3.	Human Value	24PD210ET	Health and	4	2	II
			Lifestyle Theory			
4.	MOOC	24PD211ET	NPTEL-1	4	2	II
5.	Indian	24PD312ET	Indian Indigenous	4	2	III
	Knowledge		Medicine			
	system					
6.	MOOC	24PD312ET	NPTEL-2	4	2	III
7.	Skills	24PD411EP	Statistical	4	2	IV
			Software Practical			
8.	Social	24PD412ET	Ethical leadership	4	2	IV
	Responsibility		Theory			
9.	Skills	24PD506ET	Medical Coding	4	2	V
			Theory			
10.	Skills	24PD507EP	Pharmaceutical	4	2	V
			calculation-			
			Theory			

VALUE ADDED ELECTIVE COURSES FOR PHARM.D PROGRAMME

Value added elective courses are those designed to enhance the capability of students beyond the general academic curriculum, which may help to improve the employability and equip the students with essential skills to succeed in life of the student. The program offers five categories of Elective courses – Human value, Social Responsibility, Indian Knowledge system, Skill based and NPTEL courses. All Elective courses carry 2 credits each.

A student can select any 1 elective course from the 2 Human value courses offered and 1 other social responsibility courses out of the 2 courses in the respective category. Under the Indian Knowledge system Courses category 1 course, and a student can select any 1 of them. Skill-based courses are Practical-oriented ones to provide the necessary skills to increase the employability quotient of the student. The program offers 4 skill-based courses and 2 MOOC courses are offered. The student must complete 2 MOOC courses and in the year 4 offered skill-based course is compulsory remaining 3 skill-based courses can select any 1 of them. Above all selected electives courses are considered for the CGPA calculation and the grades of all electives completed will be included in the respective year grade sheet.

Pharm.D		20	24-25
24PD113ET		FIR	ST YEAR
	COMMUNICATION SKILLS	2H	1C

Instruction hours/ week: L: 2 T:0 P:0

Theory: 3 Hrs / Week

Course Objectives:

• Understand the communication models and their application in real-life communication scenarios.

Marks: Internal: 30 External: 70 Total: 100

External Semester Exam: 3 Hours

- Analyze different communication styles and strategies, and apply appropriate strategies for effective communication in diverse contexts.
- Explore verbal and nonverbal communication, including body language, tone of voice, facial expressions, and gestures, and understand how they impact the message conveyed
- Develop active listening skills to enhance understanding, rapport-building, and conflict resolution in interpersonal communication
- Examine the culture, gender, ethnicity, and other factors on interpersonal communication, and develop strategies for effective cross-cultural communication and relationship-building
- Apply communication theories to improve communication dynamics in various personal and professional contexts.

Course Outcomes (CO's):

On successful completion of the course the student will

COs	Course Outcomes	Blooms Level
CO1	Understand the communication theories and models, including their	Understand
	application and mediated communication contexts.	
CO2	Apply communication strategies and techniques to achieve desired	Apply
	outcomes in personal and professional interactions.	
CO3	Analyse communication dynamics, evaluate messages and sources,	Analyze
	and make decisions in communication situations	
CO4	Enhance interpersonal competence by improving ability to build	Analyze
	rapport, establish trust, manage relationships, and navigate cultural	
	differences in diverse personal and professional context.	
CO5	Develop presentation skills, including speech organization, audience	Apply
	analysis, visual design, and delivery techniques, to deliver engaging	
	and persuasive presentations in various settings.	
CO6	Understand the ethical communication practices, and promote	Understand
	constructive and collaboration in personal, professional and contexts.	

Mapping with Programme Outcomes

Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1								S			
CO2					М			S			
CO3					S				М		
CO4								М	S		М
CO5	S					М				М	
CO6								S			

S-Strong; M-Medium; L-Low

Course Content:

Unit 1 Introduction to Communication Theory

• Overview of communication as a field of study, introduction to foundational theories and models, including the Shannon-Weaver model, and basic concepts such as sender, receiver, message, channel, noise, and feedback.

Unit 2 Interpersonal Communication:

• Exploration of communication dynamics in one-on-one interactions, including verbal and nonverbal cues, listening skills, self-disclosure, empathy, and conflict resolution strategies

Unit 3 Group Communication:

• Examination of communication processes within small groups and teams, including roles, norms, cohesion, decision-making, leadership, and managing group dynamics.

Unit 4 Public Speaking and Presentation Skills:

• Development of effective public speaking and presentation skills, including speech organization, audience analysis, delivery techniques, visual aids, and overcoming public speaking anxiety.

Unit 5 Intercultural Communication:

• Understanding the impact of culture on communication patterns and norms, exploring cultural differences in verbal and nonverbal communication, and developing strategies for successful intercultural communication.

Unit 6 Mass Communication and Media Effects:

• Analysis of mass communication theories and media effects on society, including agenda-setting, cultivation Theory, media literacy, and the role of media in shaping perceptions and attitudes

Unit 7 Communication in Organizations:

• Examination of communication structures and processes within organizations, including formal and informal communication channels, organizational culture, leadership communication, and conflict management strategies.

Unit 8 Communication Ethics and Law:

• Exploration of ethical issues in communication, including privacy, accuracy, fairness, and responsible use of media, as well as an overview of legal frameworks governing communication, such as defamation, copyright, and freedom of speech.

Unit 9 Technology and Communication:

• Discussion of the impact of technology on communication practices, including social media, online communication tools, digital etiquette, and the challenges and opportunities of mediated communication.

Unit 10 Applied Communication Theories:

• Application of communication theories and concepts to real- world scenarios, case studies, and Practical exercises aimed at improving communication skills in personal, professional contexts.

Suggested Readings:

- 1. Basic communication skills for Technology, Andreja.J.RutherFord, 2ndEdition, Pearson Education, 2111
- 2. Communication skills, SanjayKumar, Pushpalata, 1st Edition, Oxford Press, 2011
- 3. Organizational Behaviour, Stephen.P.Robbins, 1st Edition, Pearson, 2013
- 4. Brilliant-Communication skills, GillHasson, 1st Edition, Pearson Life, 2011
- 5. The Ace of Soft Skills: Attitude, Communication and Etiquette for success, Gopala Swamy Ramesh, 5thEdition, Pearson, 2013.
- Developing your influencing skills, Deborah Dalley, Lois Burton, Margaret, Green hall,1st Edition Universe of Learning LTD,2010
- 7. Communication skills for professionals, Konarnira, 2nd Edition, Newarrivals-PHI, 2011
- 8. Personality development and soft skills, Barun KMitra, 1st Edition, Oxford Press, 2011
- 9. Soft skill for everyone, Butter Field, 1st Edition, Cengage Learning india pvt.ltd, 2011
- Soft skills and professional communication, rancis Peters SJ, 1st Edition, McGraw Hill Education, 2011
- 11. Effective communication, John Adair, 4th Edition, Pan Mac Millan, 2009.

24PD114ET

YOGA FOR YOUTH EMPOWERMENT - THEORY 2H 1C

Instruction hours/ week: L: 2 T:0 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 3 Hrs / Week Course Objectives: To make the students

- To create awareness about Yoga and Physical Health
- To providing Value Education to improve the students character understanding Greatness of Life force and Mind
- To know about five aspects of life and to develop good Qualities and eliminating bad ones
- Learning introspection practices like Analysis of Thoughts, Moralization of Desires, Neutralization of Anger and Eradication of Worries Diversity in Men (Why Men Differ).
- To understand about the yoga, life and practice Yogasanas
- Develop teaching skills, embody Yoga philosophy, and apply learnings through practicum and reflection.

Course Outcomes (CO's):

On successful completion of the course the student will

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Cos	Course Outcomes	Blooms Level
CO1	Understand the concepts of about Yoga and Physical Health	Understand
CO2	Study the concepts a Greatness of Life force and Mind	Understand
CO3	Learn the aspects of Personality Development - Sublimation	Understand
CO4	Practices Human Resource Development	Apply
CO5	Understand about the yoga, life and Law of Nature	Apply
CO6	Develop effective teaching skills, embody Yoga philosophy in	Apply
	daily life, and demonstrate competence through practicum and	
	reflective practice sessions	

2024-25

FIRST YEAR

Mapping with Programme Outcomes

Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1										S	М
CO2											S
CO3										S	
CO4										S	S
CO5											S
CO6								S			

S-Strong; M-Medium; L-Low

Course Content:

Unit 1: Introduction to Yoga

- History and Philosophy of Yoga
- Benefits of Yoga
- Ethical Principles in Yoga Practice
- Introduction to Breath Awareness

Unit 2: Foundation of Asana Practice

- Basic Yoga Poses (Asanas)
- Alignment Principles and Modifications
- Sun Salutations (Surya Namaskar)
- Introduction to Mindful Movement

Unit 3: Deepening the Asana Practice

- Standing Poses and Balance
- Forward Bends and Hip Openers
- Backbends and Heart Openers
- Twists and Core Strength

Unit 4: Exploring Pranayama and Breathwork

- Principles of Pranayama
- Ujjayi Breath and Breath Awareness
- Introduction to Kapalabhati and Nadi Shodhana
- Integrating Breath with Movement

Unit 5: Cultivating Meditation and Mindfulness

- Introduction to Meditation
- Mindfulness Practices
- Mantra Meditation

• Loving-Kindness Meditation

Unit 6: Understanding Anatomy and Physiology

- Functional Anatomy for Yogis
- Biomechanics of Yoga Poses
- Effects of Yoga on the Nervous System
- Physiology of Stress and Relaxation

Unit 7: Teaching Methodology

- Effective Verbal Communication and Cueing
- Demonstration and Hands-on Adjustments
- Creating Sequences and Class Plans
- Cultivating a Supportive Teaching Environment

Unit 8: Yoga Philosophy and Lifestyle

- The Eight Limbs of Yoga
- Yogic Diet and Nutrition
- Practicing Yoga Off the Mat: Ethics and Values
- Integrating Yoga Philosophy into Daily Life

Unit 9: Practicum and Teaching Experience

- Teaching Practicum Sessions
- Peer Feedback and Reflection
- Finalizing Sequences and Lesson Plans
- Teaching Demonstration and Evaluation

Unit 10: Integration and Reflection

- Integration of Course Learnings
- Reflective Practice and Self-Inquiry
- Goal Setting for Continued Practice and Growth
- Celebration and Closing Ceremony

Textbooks and Readings:

- 1. "The Heart of Yoga: Developing a Personal Practice" by T.K.V. Desikachar
- 2. "The Yoga Sutras of Patanjali" by Swami Satchidananda
- 3. "Light on Yoga" by B.K.S. Iyengar
- 4. "The Secret Power of Yoga" by Nischala Joy Devi
- 5. "The Bhagavad Gita" translated by Eknath Easwaran

Anatomy and Physiology:

- 1. "The Key Muscles of Yoga" by Ray Long
- 2. "The Key Poses of Yoga" by Ray Long
- 3. "Yoga Anatomy" by Leslie Kaminoff and Amy Matthews
- 4. "The Anatomy of Yoga: An Instructor's Inside Guide to Improving Your Poses" by Abby Ellsworth
- 5. "Functional Anatomy of Yoga: A Guide for Practitioners and Teachers" by David Keil

Teaching Methodology:

- 1. "Teaching Yoga: Essential Foundations and Techniques" by Mark Stephens
- 2. "Yoga Sequencing: Designing Transformative Yoga Classes" by Mark Stephens
- 3. "The Art of Teaching Yoga: Teacher Training Manual and Workbook" by Amy Ippoliti and Taro Smith
- 4. "The Breathing Book: Good Health and Vitality Through Essential Breath Work" by Donna Farhi
- 5. "Yoga Adjustments: Philosophy, Principles, and Techniques" by Mark Stephens

Meditation and Mindfulness:

- 1. "The Miracle of Mindfulness: An Introduction to the Practice of Meditation" by Thich Nhat Hanh
- 2. "Wherever You Go, There You Are: Mindfulness Meditation in Everyday Life" by Jon Kabat-Zinn
- 3. "The Mind Illuminated: A Complete Meditation Guide Integrating Buddhist Wisdom and Brain Science" by Culadasa (John Yates, Ph.D.), Matthew Immergut, and Jeremy Graves
- 4. "Meditation for Beginners" by Jack Kornfield
- 5. "Real Happiness: The Power of Meditation: A 28-Day Program" by Sharon Salzber

24PD210ET

HEALTH AND LIFESTYLE -THEORY

Instruction hours/ week: L: 2 T:0 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 3 Hrs / Week

Course Objectives:

- Understand the relationship between lifestyle factors and health outcomes.
- Identify modifiable risk factors for chronic diseases and medication-related complications.
- Explore evidence-based strategies for promoting healthy behaviors and lifestyle modifications.
- Develop communication skills to engage patients in discussions about health and lifestyle choices.
- Evaluate the role of pharmacists in interdisciplinary healthcare teams for lifestyle management and preventive care.
- To Foster critical thinking and decision-making skills regarding health-related information and trends

Course Outcomes (CO's):

On successful con	npletion of the cou	urse the student will
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Cos	Course Outcomes	Blooms Level
CO1	To articulate and evaluate different philosophical perspectives on the	Evaluate
	nature of health and well-being.	
CO2	To demonstrate an understanding of ethical considerations in health-	Understand
	related decisions and lifestyle choices	
CO3	To analyze the philosophical implications of the mind-body	Analyze
	relationship and its significance for overall well-being.	
CO4	To assess the ethical dimensions of dietary choices and their impact	Analyze
	on personal health and environmental sustainability.	
CO5	Students will evaluate the philosophical significance of physical	Evaluate
	activity and exercise for promoting health and well-being.	
CO6	To apply mindfulness practices and meditation techniques to enhance	Apply
	self-awareness and mental health.	

2024-25 SECOND YEAR

2C

Mapping with Programme Outcomes

Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1									S		М
CO2										S	
CO3										S	
CO4		М							М		S
CO5											S
CO6		S									

S-Strong; M-Medium; L-Low

Course Content:

Unit 1: Introduction to Health and Lifestyle Medicine

- Definition of lifestyle medicine
- Overview of lifestyle factors influencing health
- The role of pharmacists in promoting healthy lifestyles

Unit 2: Nutrition and Dietary Habits

- Principles of healthy eating
- Dietary guidelines and recommendations
- Nutrition-related diseases and interventions

Unit 3: Physical Activity and Exercise Prescription

- Benefits of physical activity for health and well-being
- Exercise recommendations for different age groups and populations
- Exercise prescription and counseling skills for pharmacists

Unit 4: Stress Management and Mental Health

- Understanding the impact of stress on health
- Stress management techniques and coping strategies
- Mental health resources and referral pathways

Unit 5: Sleep Hygiene and Circadian Rhythms

- Importance of quality sleep for overall health
- Strategies for improving sleep hygiene
- Circadian rhythms and their implications for health

Unit 6: Tobacco Cessation and Substance Use Disorders

- Health consequences of tobacco and substance use
- Evidence-based interventions for tobacco cessation
- Pharmacist's role in addressing substance use disorders

Unit 7: Weight Management and Obesity Prevention

- Obesity as a chronic disease
- Strategies for weight management and prevention
- Behavioral approaches to promoting healthy weight

Unit 8: Chronic Disease Prevention and Management

- Lifestyle interventions for preventing and managing chronic diseases
- Patient-centered approaches to chronic disease management
- Medication adherence and lifestyle modifications

Unit 9: Health Promotion and Community Outreach

- Designing health promotion initiatives for diverse populations
- Community resources and partnerships for promoting healthy lifestyles
- Evaluating the effectiveness of health promotion programs

Unit 10: Integrative Approaches to Lifestyle Medicine

- Integrating lifestyle medicine into pharmacy practice
- Interdisciplinary collaboration for holistic patient care
- Ethical considerations in promoting lifestyle modifications

Textbooks and Readings:

- "Prescribing Lifestyle Medicine" by Mark H. Houston
- "Lifestyle Medicine: A Manual for Clinical Practice" edited by James M. Rippe
- "Principles and Practice of Sleep Medicine" by Meir H. Kryger

Pharm.D	2024-25
24PD211ET	SECOND YEAR
NPTEL COURSE	2H 2C

24PD312ET

INDIAN INDIGENOUS MEDICINE - THEORY

2H

Instruction hours/ week: L: 2 T:0 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Course Objectives: Students can able to

- Understand the philosophical foundations, historical development, and cultural significance of Indian indigenous medicine systems.
- Familiarize students with the diagnostic methods, principles of treatment, and pharmacological practices of Ayurveda, Siddha, and Unani
- Explore the integration of Indian indigenous medicine with modern healthcare systems and its cultural significance in Indian society.
- Analyze global perspectives on traditional medicine and identify emerging trends and career pathways in Indian indigenous medicine.
- Develop critical thinking skills and the ability to apply indigenous medical principles to contemporary health challenges.
- Apply the ethical engagement with indigenous healing practices and foster a respectful understanding of cultural diversity in healthcare.

Course Outcomes (CO's):

On successful completion of the course the student will

Cos	Course Outcomes	Blooms Level
CO1	Explain the philosophical underpinnings of Ayurveda, Siddha,	Understand
	Unani, and folk medicine traditions.	
CO2	Identify and describe various diagnostic techniques used in	Knowledge
	Ayurveda, Siddha, and Unani medicine.	
CO3	Analyze the principles of treatment and pharmacological practices	Analyze
	employed in Ayurveda, Siddha, and Unani systems.	
CO4	Explore emerging trends, research opportunities, and career	Apply
	pathways in the field of Indian indigenous medicine.	
CO5	Analyze and synthesize information from various sources to	Analyze
	critically evaluate indigenous medical practices and their	
	applications.	
CO6	Apply indigenous medical principles to propose solutions for	Apply
	contemporary health challenges, such as non-communicable	
	diseases and mental health	

THIRD YEAR

2C

Mapping with Programme Outcomes

Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1	S									S	М
CO2	S								М		М
CO3		S								S	
CO4									S		М
CO5				М							S
CO6							S	М			

S-Strong; M-Medium; L-Low

Course Content: Unit 1: Introduction to Indian Indigenous Medicine

- Overview of Indian indigenous medicine systems: Ayurveda, Siddha, Unani, and folk medicine traditions
- Historical development and cultural context of indigenous medicine in India
- Philosophical underpinnings and basic principles shared across Indian indigenous medicine systems

Unit 2: Philosophical Foundations and Diagnostic Methods

- Principles of Ayurveda: Doshas, dhatus, and malas
- Siddha philosophy: Body-mind-spirit integration and vitalism
- Unani humoral Theory and temperament types
- Diagnostic methods in Ayurveda, Siddha, and Unani: Pulse diagnosis, tongue examination, and clinical examination

Unit 3: Principles of Treatment and Pharmacological Practices

- Ayurvedic treatment modalities: Herbal medicine, panchakarma, and dietary recommendations
- Siddha therapeutic interventions: Marma therapy, yoga, and siddha external therapies
- Unani therapeutic modalities: Regimental therapy, pharmacotherapy, and dietotherapy
- Pharmacology in Ayurveda, Siddha, and Unani: Herbal formulations, mineral preparations, and traditional alchemical processes

Unit 4: Integrative Medicine and Modern Healthcare Systems

- Challenges and opportunities in integrating indigenous medicine with modern healthcare systems
- Case studies and examples of successful integration models in India and abroad
- Regulatory frameworks and quality standards for traditional medicine products and practitioners
- Cultural significance of indigenous medicine in Indian society: Rituals, festivals, and healing traditions

Unit 5: Global Perspectives and Recognition of Traditional Medicine

- International recognition and acceptance of traditional medicine: WHO Traditional Medicine Strategy 2014-2023
- Role of traditional medicine in addressing global health challenges: Access to healthcare, non-communicable diseases, and mental health
- Emerging trends, research opportunities, and career pathways in Indian indigenous medicine

Unit 6: Ethics and Cultural Competency

- Ethical considerations in engaging with indigenous healing practices
- Respectful understanding of cultural diversity in healthcare
- Communication skills and cultural competency in engaging with diverse communities and healthcare traditions

Unit 7: Ayurvedic Pharmacology and Herbal Medicine

- Principles of Ayurvedic pharmacology: Rasa, guna, veerya, vipaka, and prabhava
- Classification of medicinal plants and herbal preparations
- Ayurvedic pharmacopoeia and quality standards for herbal products

Unit 8: Siddha Medicine: Alchemy and Vitalism

- Siddha alchemical processes: Preparation of mercury and minerals
- Siddha herbal formulations and external therapies

• Siddha pharmacopoeia and quality standards for medicinal substances

Unit 9: Unani Medicine: Greco-Arabic Influences

- Historical development and origins of Unani medicine
- Principles of Unani humoral Theory and treatment modalities
- Unani pharmacopoeia and quality standards for herbal medicines

Unit 10: Case Studies and Research Opportunities

- Analysis of case studies and clinical trials in Indian indigenous medicine
- Research opportunities and future directions in Ayurveda, Siddha, and Unani
- Career pathways and professional development in the field of Indian indigenous medicine

Reference Books:

- 1. "Encyclopedia of Indian Medicine" edited by C.R. Rao
- "History of Indian Medicine: Containing Notices, Biographical Sketches, and Bibliographic Lists of Medical Works in Sanskrit, Arabic, Persian, Pali, Prakrit, and Other Eastern Languages" by G. S. Mehta
- 3. "The Roots of Ayurveda" by Dominik Wujastyk
- 4. "Indian Medicinal Plants: An Illustrated Dictionary" by C.P. Khare
- 5. "Indian Alchemy: Soma in the Veda" by David Gordon White
- 6. "Indian Medicinal Plants: A Compendium of 500 Species" by Ashok K. Jain

Textbooks:

- 1. "Ayurveda: The Science of Self-Healing" by Dr. Vasant Lad
- 2. "The Yoga of Herbs: An Ayurvedic Guide to Herbal Medicine" by Dr. David Frawley and Dr. Vasant Lad
- 3. "Essentials of Siddha Medicine" by M.A. Venkatakrishna Rao
- 4. "Introduction to Unani Medicine" by Hakim Syed Zillur Rahman
- 5. "Principles and Practice of Unani Medicine" by S.M. Bokhari
- 6. "Textbook of Pharmacognosy and Phytochemistry" by Biren Shah and A.K. Seth
- 7. "Ayurveda and the Mind: The Healing of Consciousness" by Dr. David Frawley
- 8. "Siddha Medicine: The Secrets of Ancient Siddha Vaidyas" by Dr. David Frawley and Dr. Subhash Ranade
- 9. "Unani Medicine: Introduction and Clinical Application" by Mohammad Zaki Ansari
- 10. "The Legacy of Charaka" by Veerabhadran Ramanathan
- 11. "Traditional Medicine: A Global Perspective" edited by Steven B. Kayne

Pharm.D	2024-25
24PD312ET	THIRD YEAR
NPTEL COURSE	2H 2C

-
FOURTH YEAR

24PD411EP

STATISTICAL SOFTWARE -PRACTICAL

3H

Instruction hours/ week: L: 0 T:0 P:3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Course Objectives:

- Understand the importance of statistical software in pharmacy research and practice.
- Familiarize with the interface and functionalities of statistical software packages.
- Perform basic statistical analyses, including descriptive statistics and hypothesis testing.
- Explore advanced statistical techniques relevant to pharmacy research, such as regression analysis and survival analysis.
- Develop skills in data visualization and presentation using statistical software tools.

Course Outcomes (CO's):

On successful completion of the course the student will

Cos	Course Outcomes	Blooms Level
CO1	Understand the significance of statistical software in pharmacy	Understand
	research, including its role in data analysis, decision-making and	
	evidence-based practice.	
CO2	Demonstrate proficiency in navigating the interface and utilizing	Apply
	the key functionalities of statistical software packages commonly	
	used in pharmacy research and practice settings.	
CO3	Apply fundamental statistical techniques, including descriptive	Analyse
	statistics and hypothesis testing, to analyze pharmacy-related data	
	sets effectively, interpret findings and draw appropriate	
	conclusions.	
CO4	Applying advanced statistical methods relevant to pharmacy	Understand
	research to investigate complex relationships and outcomes in	
	pharmaceutical studies.	
CO5	Develop in data visualization and presentation techniques using	Understand
	statistical software tools, enabling the creation of clear and	
	informative visual representations of pharmacy-related data for	
	communication.	

2C

Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1				S							
CO2	S		М								
CO3	S			М							
CO4	М		М	S							
CO5	М			S							
CO6	S			М							

Mapping with Programme Outcomes

S-Strong; M-Medium; L-Low

Course Outline:

Unit 1: Introduction to Statistical Software

- Overview of statistical software packages (e.g., SPSS, R, SAS)
- Installing and setting up statistical software
- Introduction to the user interface and basic functionalities

Unit 2: Data Management and Importing

- Data types and formats
- Importing data from different sources (e.g., Excel, CSV files)
- Data cleaning and preparation techniques

Unit 3: Descriptive Statistics

- Measures of central tendency and variability
- Frequency distributions and graphical representations
- Interpreting descriptive statistics output

Unit 4: Inferential Statistics: Parametric Tests

- Introduction to hypothesis testing
- One-sample and independent samples t-tests
- Paired samples t-test and analysis of variance (ANOVA)

Unit 5: Inferential Statistics: Non-parametric Tests

- Wilcoxon signed-rank test and Mann-Whitney U test
- Kruskal-Wallis test and Friedman test
- Choosing between parametric and non-parametric tests

Unit 6: Regression Analysis

- Simple linear regression
- Multiple linear regression

• Interpreting regression output and assessing model fit

Unit 7: Survival Analysis

- Kaplan-Meier survival curves
- Log-rank test and Cox proportional hazards model
- Applications of survival analysis in pharmacy and healthcare research

Unit 8: Data Visualization

- Creating basic and advanced plots (e.g., histograms, scatterplots, box plots)
- Customizing plot aesthetics and labels
- Exporting plots for publication and presentations

Unit 9: Advanced Topics in Statistical Software

- Factor analysis and principal component analysis (PCA)
- Cluster analysis and discriminant analysis
- Time series analysis and forecasting techniques

Unit 10: Applied Research Projects

- Designing and executing a research project using statistical software
- Analyzing and interpreting research findings
- Presenting research results using appropriate statistical visualizations and techniques

Textbooks and Readings:

- "Discovering Statistics Using SPSS" by Andy Field
- "R for Data Science" by Hadley Wickham and Garrett Grolemund
- "Applied Survival Analysis: Regression Modeling of Time-to-Event Data" by David W. Hosmer Jr., Stanley Lemeshow, and Susanne May.

Suggested reading books

- 1. "An Introduction to Statistical Learning: with Applications in R" by Gareth James, Daniela Witten, Trevor Hastie, and Robert Tibshirani. 2nd Edition (2021)
- 2. "Discovering Statistics Using IBM SPSS Statistics" by Andy Field 5th Edition (2021)
- 3. "R for Data Science" by Hadley Wickham and Garrett Grolemund. 1st Edition (2017)
- 4. "Stata Survival Manual" by David Pevalin and Karen Robson. 5th Edition (2020)
- 5. "SPSS Survival Manual" by Julie Pallant. 7th Edition (2020)

6. "SAS Essentials: Mastering SAS for Data Analytics" by Alan Elliott and Wayne A. Woodward. 2nd Edition (2020)

7. "R Graphics Cookbook" by Winston Chang. 2nd Edition (2018)

8. "Data Analysis Using Regression and Multilevel/Hierarchical Models" by Andrew Gelman and Jennifer Hill. 2nd Edition (2020)

9. "Modern Applied Statistics with S" by William N. Venables and Brian D. Ripley 4th Edition (2002)

24PD412ET

ETHICAL LEADERSHIP - THEORY

Instruction hours/ week: L: 1 T:0 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Course Objective

- 1. Understand a comprehensive ethical leadership principles, theories, and models, including the importance of integrity, honesty, fairness, and accountability.
- 2. Identify, analyze, and evaluate ethical dilemmas in leadership contexts, applying ethical decision-making frameworks.
- 3. Apply ethical principles in ethical decision-making, recognize cognitive biases, and to resolve complex dilemmas.
- 4. Enhance the communication skills, learning to communicate ethically and effectively and ethical challenges with integrity and transparency.
- 5. Understand the importance of diversity, equity, and inclusion in ethical leadership, learning to promote a culture of respect, dignity, and fairness that values diverse perspectives and experiences.
- 6. Fostering a culture of ethical leadership within organizations, including promoting ethical behavior, values alignment, and ethical decision-making processes.

Course Outcome (CO's):

Upon completion of the course, the student shall be able to

Cos	Course Outcomes	Blooms Level
CO1	Understand of Ethical Leadership Concepts, key principles and theories of	Understand
	ethical leadership, and accountability in leadership roles.	
CO2	Apply ethical decision-making models and frameworks to analyze complex	Apply
	dilemmas, evaluate alternative courses of action, and make ethical decisions	
	in leadership contexts.	
CO3	Communicate ethically and transparently with stakeholders, address	Analyse
	conflicts with integrity, and foster open dialogue in organizational settings.	
CO4	Understand the ethical imperative of promoting diversity, equity and	Understand
	develop strategies for creating inclusive environments that value diverse	
	perspectives	
CO5	Cultivate a culture of ethical leadership within organizations, including	Apply
	ethical behavior, values and ethical decision-making processes among team	
	members.	
CO6	Engage in self-reflection and introspection, assessing their own values,	Apply
	beliefs, and ethical principles, and develop a personal ethical leadership	
	philosophy and action plan.	



FOURTH YEAR

1H 2C

Mapping with Programme Outcomes

Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1					S		S				
CO2					М		S				
CO3					S		М				
CO4		М					S			S	
CO5		S					М				
CO6		S			S						

S-Strong; M-Medium; L-Low

Course contents

Unit 1: Introduction to Ethical Leadership

- Definition and importance of ethical leadership
- Historical perspectives on ethical leadership
- Theoretical frameworks for understanding ethical leadership
- Ethical dilemmas in leadership roles

Unit 2: Moral Reasoning and Decision Making

- Ethical theories and principles (e.g., utilitarianism, deontology, virtue ethics)
- Moral development and ethical decision-making models
- Cognitive biases and ethical decision-making
- Case studies on ethical decision-making in leadership contexts

Unit 3: Values-Based Leadership

- Identifying personal and organizational values
- Aligning values with leadership actions and decisions
- Building a values-driven organizational culture
- Ethical leadership and corporate social responsibility

Unit 4: Ethical Communication

- Importance of transparency and honesty in communication
- Ethical considerations in persuasive communication
- Active listening and empathetic communication skills
- Addressing conflicts and resolving ethical communication challenges

Unit 5: Ethical Decision-Making Processes

- Ethical decision-making frameworks (e.g., Rest's Four Component Model, Kidder's Ethical checkpoints)
- Stakeholder analysis and ethical impact assessment
- Implementing ethical decision-making processes in organizational contexts
- Ethical leadership in crisis management and decision-making under pressure

Unit 6: Ethical Leadership in Diversity and Inclusion

- Promoting diversity and inclusion in leadership roles
- Ethical considerations in managing diverse teams
- Addressing bias and discrimination in leadership practices
- Creating inclusive organizational cultures through ethical leadership

Unit 7: Ethical Leadership and Organizational Justice

- Principles of organizational justice (distributive, procedural, interactional)
- Ethical implications of power dynamics in organizations
- Fairness in resource allocation and decision-making processes
- Ethical leadership and fostering a culture of trust and fairness

Unit 8: Ethical Leadership Development

- Assessing and developing ethical leadership competencies
- Ethical leadership training and development programs
- Coaching and mentoring for ethical leadership
- Personal reflection and growth as an ethical leader

Unit 9: Ethical Leadership in Global Contexts

- Cross-cultural perspectives on ethical leadership
- Ethical challenges in international business and diplomacy
- Cultural differences in ethical values and practices
- Strategies for promoting ethical leadership in global organizations

Unit 10: Ethical Leadership in Action: Case Studies and Applications

- Analyzing real-world ethical leadership challenges and dilemmas
- Ethical leadership in specific industries or sectors (e.g., healthcare, finance, technology)
- Developing action plans for ethical leadership initiatives
- Reflection on personal values and commitments as ethical leaders

Pharm.D

References

- 1. "Ethical Leadership and Decision Making in Education: Applying Theoretical Perspectives to Complex Dilemmas" by Joan Poliner Shapiro and Steven Jay Gross
- 2. "Ethical Leadership in Schools: Creating Community in an Environment of Accountability" by Kenneth A. Strike and Camille A. Farrington
- 3. "Ethical Leadership and Decision Making in Education: Applying Theoretical Perspectives to Complex Dilemmas" by Joan Poliner Shapiro and Steven Jay Gross
- 4. "Ethical Leadership: A Primer" by Gerald C. Kane and Bret Simmons: A concise introduction to ethical leadership, covering key concepts, theories, and Practical strategies for fostering ethical behavior in organizations.
- 5. **"The Ethics of Leadership" by Joanne B. Ciulla** This book provides an in-depth exploration of the ethical dimensions of leadership, examining historical perspectives, contemporary issues, and case studies from various fields.
- "Leading with Integrity: Character-Based Leadership" by John J. Sosik and William A. Gentry - Focusing on character-based leadership, this text explores the role of integrity, honesty, and authenticity in ethical leadership practices.
- "Ethical Leadership: Global Challenges and Perspectives" edited by Aycan Kara and Ayşegül Özbebek Tunç - Offering global perspectives on ethical leadership, this book addresses cross-cultural challenges, ethical dilemmas, and best practices in leadership ethics.
- 8. **"The Ethics of Authenticity" by Charles Taylor** This philosophical work explores the concept of authenticity in leadership, examining how leaders can align their actions with their values and maintain integrity in a complex world.
- 9. "The Responsible Leader: Developing a Culture of Responsibility in an Uncertain World" by Tim Richardson - Focusing on responsibility and accountability in leadership, this book offers Practical insights and tools for ethical decision-making and organizational governance.
- 10. "Ethical Leadership: Progress with a Moral Compass" by Joanne B. Ciulla, Terry L. Price, and Susan E. Murphy - This book provides a comprehensive overview of ethical leadership research, theories, and practices, offering guidance for leaders to navigate ethical challenges and build trust within organizations.

Pharm.D		2024-2	25
24PD506ET		FIFTH `	YEAR
	MEDICAL CODING -THEORY	1H	2C

Instruction hours/ week: L: 1 T:0 P:0

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Theory: 3 Hrs / Week **Course Objectives**

- Course Objectives
 - Understand the concept of the structure, functioning and regulations of the healthcare system.
 - Identify the influence of fundamental terminology used in healthcare documentation and coding.
 - Acquire proficiency in assigning accurate medical codes using various classification systems such as ICD-10-CM, CPT, and HCPCS Level II.
 - Explore strategies of coding guidelines, conventions, and rules to ensure code assignment and compliance with regulatory.
 - Develop critical thinking skills by analyzing complex medical scenarios and determining the appropriate codes based on documentation, clinical indicators, and coding guidelines.
 - Critically evaluate ethical coding practices, patient confidentiality and coding regulations:

Course Outcomes (CO's):

COs	Course Outcomes	Blooms Level
CO1	Understand the diagnosis and procedure codes using the	Understand
	appropriate code set, including ICD-10-CM, CPT, and HCPCS	
	Level II.	
CO2	Applying the coding guidelines, conventions, and principles by	Apply
	assign correct codes in various medical scenarios	
CO3	Analyze medical documentation, including physician notes,	Analyze
	operative reports, and diagnostic test results, to extract relevant	
	information for coding purposes	
CO4	Evaluate the accuracy and compliance of coded medical records	Understand
	with coding guidelines, regulatory requirements.	
CO5	Understand the ability to use coding reference materials, such as	Understand
	codebooks, encoding software, and online databases, to research	
	and resolve coding queries.	
CO6	Communicate effectively with healthcare professionals, including	Apply
	physicians, nurses, and administrators, to clarify documentation	
	and ensure accurate code assignment	

Mapping with Programme Outcomes

Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1	S	М									
CO2		S									М
CO3	М	S									
CO4		S		М							
CO5	М						S				
CO6	S	М									

S-Strong; M-Medium; L-Low

Course Content

Unit 1: Introduction to Medical Coding

- Overview of medical coding and its importance in healthcare
- Historical perspective and evolution of coding systems
- Role of medical coders in the healthcare industry

Unit 2: Medical Terminology and Anatomy

- Fundamentals of medical terminology, including prefixes, suffixes, and root words
- Anatomy and physiology basics relevant to medical coding
- Common diseases, conditions, and procedures encountered in medical coding practice

Unit 3: ICD-10-CM Coding

- Introduction to the International Classification of Diseases, 10th Revision, Clinical Modification (ICD-10-CM)
- Structure and organization of the ICD-10-CM code set
- Guidelines for accurate diagnosis code assignment and documentation requirements

Unit 4: CPT Coding

- Introduction to the International Classification of Diseases, 10th Revision, Clinical Modification (ICD-10-CM)
- Structure and organization of the ICD-10-CM code set
- Guidelines for accurate diagnosis code assignment and documentation requirements

Unit 5: HCPCS Level II Coding

- Introduction to the Healthcare Common Procedure Coding System (HCPCS) Level II
- Coverage of supplies, durable medical equipment (DME), and other healthcare services not included in CPT
- Application of HCPCS Level II codes in coding and billing processes

Unit 6: Coding Guidelines and Conventions

- Detailed exploration of coding guidelines, conventions, and official coding rules
- Importance of accuracy, specificity, and consistency in code assignment
- Coding scenarios and exercises to reinforce guideline application

Unit 7: Modifiers and Edits

- Understanding modifiers and their role in modifying or clarifying procedure codes
- Commonly used modifiers and their appropriate application
- Identifying and resolving coding edits, including National Correct Coding Initiative (NCCI) edits

Unit 8: Evaluation and Management (E/M) Coding

- Overview of E/M coding for outpatient and inpatient encounters
- Components of E/M services and key documentation requirements
- Guidelines for selecting the appropriate E/M code based on level of service provided

Unit 9: Regulatory Compliance and Ethics

- Importance of ethical coding practices and compliance with regulatory requirements
- Overview of healthcare regulations, including HIPAA, Stark Law, and False Claims Act
- Consequences of fraudulent coding practices and strategies for maintaining compliance

Unit 10: Coding Practice and Case Studies

- Application of coding knowledge and skills through Practical coding exercises
- Analysis of real-world coding scenarios and case studies
- Feedback and discussion to reinforce learning and address coding challenges.

Reference Books:

- ICD-10-CM Official Guidelines for Coding and Reporting: Published by the American Hospital Association, this book provides official coding guidelines for diagnosis coding using the International Classification of Diseases, 10th Revision, Clinical Modification (ICD-10-CM).
- CPT Professional Edition: Published by the American Medical Association (AMA), this book provides the most current procedural coding guidelines and codes using the Current Procedural Terminology (CPT) system.
- 3. HCPCS Level II Professional: Also published by the AMA, this book contains the Healthcare Common Procedure Coding System (HCPCS) Level II codes, which are used for procedures, services, and supplies not covered by CPT codes.
- 4. Principles of Healthcare Reimbursement: This book covers the basics of healthcare reimbursement, including insurance policies, billing processes, and regulatory requirements related to medical coding and billing
- 5. Step-by-Step Medical Coding: This textbook provides a comprehensive overview of medical

coding principles and practices, including step-by-step instructions for assigning diagnostic and procedural codes.

- 6. Understanding ICD-10-CM and ICD-10-PCS: A Worktext: This book focuses specifically on understanding and applying ICD-10-CM and ICD-10-PCS coding systems, with Practical exercises and case studies.
- 7. Medical Coding Certification Exam Preparation: This book is designed to help students prepare for medical coding certification exams, such as the Certified Professional Coder (CPC) exam offered by the American Academy of Professional Coders (AAPC).
- 8. Coding Basics: Understanding Medical Collections: This book covers the basics of medical coding and collections, including coding principles, reimbursement methodologies, and strategies for maximizing revenue

Pharm.D	2024-25
24PD507EP	FIFTH YEAR

PHARMACEUTICAL CALCULATION - PRACTICAL 3H 2C

Instruction hours/ week: L:0 T:0 P:3

Marks: Internal: 30 External: 70 Total: 100 External Semester Exam: 3 Hours

Course Objective:

- Develop accurate pharmaceutical calculation skills.
- Understand dosage determination principles.
- Apply pharmacokinetic concepts for dosage optimization.
- Practice pharmaceutical compounding techniques.
- Perform quality control tests for product potency and uniformity.
- Promote awareness of medication safety through precise calculations.

Course Outcomes (CO's):

On successful completion of the course the student will

Cos	Course Outcomes	Blooms Level
CO1	Achieve accuracy in pharmaceutical calculations	Apply
CO2	Optimize dosage regimens using pharmacokinetic concepts	Apply
CO3	Master pharmaceutical compounding techniques	Apply
CO4	Ensure product potency and uniformity through quality control	Apply
CO5	Grasp dosage determination principles for precise prescriptions.	Apply
CO6	Enhance medication safety awareness through meticulous calculations.	Apply

Mapping with Programme Outcomes

Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1	S	М	М								
CO2	S										
CO3	S	М									
CO4	S										
CO5	S		S								
CO6	S			М							

S-Strong; M-Medium; L-Low

Course Content

- 1. Drug Dosage Calculation
- 2. Pharmaceutical Solution Preparation
- 3. Percent Composition Calculation

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- 4. Drug Stability Testing
- 5. Pharmaceutical Compounding
- 6. Pharmacokinetic Parameter Calculation
- 7. Pharmaceutical Dosing Regimen Optimization
- 8. Pharmaceutical Excipient Compatibility Study
- 9. Pharmaceutical Quality Control
- 10. Pharmaceutical Formulation Cost Analysis
- 11. Pharmaceutical Dilution Calculation
- 12. Pediatric Dosage Calculation
- 13. Intravenous Infusion Rate Calculation
- 14. Drug Concentration Adjustment Calculation
- 15. Pharmaceutical Manufacturing Yield Calculation
- 16. Bioavailability and Bioequivalence Calculation
- 17. Pharmacokinetic Modeling and Simulation
- 18. Pharmaceutical Expiry Date Calculation
- 19. Drug Solubility Calculation
- 20. Pharmaceutical Process Validation Calculation

References

- Remington: The Science and Practice of Pharmacy" edited by David B. Troy and Joseph Price Remington
- 2. "Pharmaceutical Calculations" by Howard C. Ansel and Mitchell J. Stoklosa
- "Goodman & Gilman's: The Pharmacological Basis of Therapeutics" edited by Laurence Brunton, Bruce Chabner, and Björn Knollmann
- 4. "Pharmaceutical Dosage Forms and Drug Delivery Systems" by Howard C. Ansel and Loyd V. Allen Jr.
- "Principles of Physical Biochemistry" by Kensal E. van Holde, W. Curtis Johnson, and P. Shing Ho
- "Pharmacokinetics: Processes, Mathematics, and Applications" by Peter Welling, Thomas
 E. Saltzman, and William E. Hassan
- 7. "Pharmaceutical Compounding and Dispensing" by John F. Marriott and Keith A. Wilson
- 8. "Pharmaceutical Calculations" by Payal Agarwal and Sarvesh Paliwal
- 9. "Pharmaceutical Analysis: A Textbook for Pharmacy Students and Pharmaceutical Chemists" by David G. Watson
- "Pharmaceutical Manufacturing Handbook: Production and Processes" edited by Shayne Cox Gad