

Regulations and Syllabus for the Pharm.D & Pharm.D (Post Baccalaureate)

(w.e.f. June 2018)



**ADICHUNCHANAGIRI UNIVERSITY
B.G.NAGARA**

REGULATIONS

1. Short Title and Commencement

These regulations shall be called as “The Revised Regulations for the Pharm D & Pharm D (Post Baccalaeureate) of the Pharmacy Council of India, New Delhi”. They shall come into effect from the Academic Year 2016-17. The regulations framed are subject to modifications from time to time by Pharmacy Council of India.

2. Minimum Qualification for Admission to -

2.1. Pharm.D Course First Year :

2.1.1. A pass in any of the following examinations –

- i. Students who have passed two years P.U.C. examination of Karnataka P.U.C. Board or an equivalent examination of any other approved Board or University established by law in India with English as one of the subject and Physics, Chemistry as compulsory subjects along with one of the following subject – Mathematics or Biology or P.C.M.B.
- ii. Students who have passed D. Pharm course from institutions approved by Pharmacy Council of India U/S 12 of Pharmacy Act, 1948
- iii. Any other qualification approved by the Pharmacy Council of India as equivalent to any of the above examinations.

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

2.2. Pharm.D Post Baccalaureate course –

2.2.1. A pass in B Pharm degree examination of ADICHUNCHANAGIRI UNIVERSITY or B Pharm examination of any other Indian University established by law in India or any other degree courses in Pharmacy recognized as equivalent to B Pharm by ACU and Pharmacy Council of India.

3. Duration of the course –

3.1. 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 speciality 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.

3.2. Pharm.D. (Post Baccalaureate):

Eligible B Pharm graduates shall be admitted to the first year of the Pharm.D (Post Baccalaureate) course. 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 speciality 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. Medium of Instruction

The medium of instruction and examination of Pharm.D & Pharm.D (Post Baccalaureate) courses shall be English.

Number of admissions in the above said courses shall be as prescribed by the Pharmacy Council of India from time to time and presently be restricted as below –

4.1 Pharm.D. Course – 30 students.

4.2 Pharm.D. (Post Baccalaureate) Course – 10 students.

- 5 Institutions running B.Pharm course approved under section 12 of the Pharmacy Act, will only be permitted to run Pharm.D course. Pharm.D. (Post Baccalaureate) course will be permitted only in those institutions which are permitted to run Pharm.D. course.

6 Course of study –

6.1 **Pharm.D Course** - 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 (3), (4) and (5) below.

6.2 The total number of years taken by the candidate to complete the Pharm.D course shall not exceed 10 years from the date of admission.

TABLES

First Year:

S. No	Name of Subject	No. of hours of Theory	No. of hours of Practical	No. of hours of Tutorial
(1)	(2)	(3)	(4)	(5)
1.1	Human Anatomy and Physiology	3	3	1
1.2	Pharmaceutics	2	3	1
1.3	Medicinal Biochemistry	3	3	1
1.4	Pharmaceutical Organic Chemistry	3	3	1
1.5	Pharmaceutical Inorganic Chemistry	2	3	1
1.6	Remedial Mathematics/ Biology**	3	3*	1
	Total Hours	16	18	6 = (40)

* For Biology

** For candidates who have studied PCMB in 10+2 course are exempted.

** Colleges to conduct the Examination.

Second Year:

S. No	Name of Subject	No. of hours of Theory	No. of hours of Practical	No. of hours of Tutorial
(1)	(2)	(3)	(4)	(5)
2.1	Pathophysiology	3	-	1
2.2	Pharmaceutical Microbiology	3	3	1
2.3	Pharmacognosy & Phytopharmaceuticals	3	3	1
2.4	Pharmacology-I	3	-	1
2.5	Community Pharmacy	2	-	1
2.6	Pharmacotherapeutics-I	3	3	1
	Total Hours	17	09	6 = (32)

Third Year:

S. No	Name of Subject	No. of hours of Theory	No. of hours of Practical	No. of hours of Tutorial
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(1)	(2)	(3)	(4)	(5)
3.1	Pharmacology-II	3	3	1
3.2	Pharmaceutical Analysis	3	3	1
3.3	Pharmacotherapeutics-II	3	3	1
3.4	Pharmaceutical Jurisprudence	2	-	-
3.5	Medicinal Chemistry	3	3	1
3.6	Pharmaceutical Formulations	2	3	1
	Total Hours	16	15	6 = (36)

Fourth Year:

S. No	Name of Subject	No. of hours of Theory	No. of hours of Practical	No. of hours of Tutorial
(1)	(2)	(3)	(4)	(5)
4.1	Pharmacotherapeutics-III	3	3	1
4.2	Hospital Pharmacy	2	3	1
4.3	Clinical Pharmacy	3	3	1
4.4	Biostatistics & Research Methodology	2	-	1
4.5	Biopharmaceutics & Pharmacokinetics	3	3	1
4.6	Clinical Toxicology	2	-	1
	Total Hours	15	12	6 = (33)

Fifth Year:

S. No	Name of Subject	No. of hours of Theory	No. of hours of Practical	No. of hours of Tutorial
(1)	(2)	(3)	(4)	(5)
5.1	Clinical Research	3	-	1
5.2	Pharmacoepidemiology and Pharmacoeconomics	3	-	1
5.3	Clinical Pharmacokinetics & Pharmacotherapeutic Drug Monitoring	2	-	1
5.4	Clerkship *	-	-	1

5.5	Project Work (Six Months)	-	20	-
	Total Hours	8	20	4 = (32)

* Attending ward rounds on daily basis.

Sixth Year:

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

6.2.1 Six months in General Medicine department, and

6.2.2 Two months each in three other speciality departments

6.3 Pharm.D (Post Baccalaureate) Course - The course of study for Pharm.D (Post Baccalaureate) 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 (3), (4) and (5) below.

6.4 The total number of years taken by the candidate to complete the Pharm.D (Post Baccalaureate) course shall not exceed 6 years from the date of admission.

T A B L E S

First Year (Post Baccalaureate):

S. No	Name of Subject	No. of hours of Theory	No. of hours of Practical	No. of hours of Tutorial
(1)	(2)	(3)	(4)	(5)
1.1	Pharmacotherapeutics- I & II	3	3	1
1.2	Hospital & Community Pharmacy	3	3	1
1.3	Clinical Pharmacy	3	3	1
1.4	Biostatistics & Research Methodology	2	-	1
1.5	Clinical Toxicology	2	-	1
1.6	Pharmacotherapeutics-III	3	3	1
1.7	Biopharmaceutics & Pharmacokinetics	3	3	1
	Total Hours	19	15	7 = (41)

Second Year (Post Baccalaureate):

S. No	Name of Subject	No. of	No. of	No. of
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		hours of Theory	hours of Practical	hours of Tutorial
(1)	(2)	(3)	(4)	(5)
2.1	Clinical Research	3	-	1
2.2	Pharmacoepidemiology and Pharmacoeconomics	3	-	1
2.3	Clinical Pharmacokinetics & Pharmacotherapeutic Drug Monitoring	2	-	1
2.4	Clerkship *	-	-	1
2.5	Project Work (Six Months)	-	20	-
	Total Hours	8	20	4 = (32)

* Attending ward rounds on daily basis.

Third Year:

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

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

- 7 Syllabus. – The syllabus for each subject of study in the said Tables shall be as specified in Appendix -A to these regulations.
- 8 Approval of institution conducting the course of study –
 - 8.1 The regular course for Pharm.D. I, II, III, IV, V and VI year shall be conducted by an institution approved by Pharmacy Council of India and affiliated to ADICHUNCHANAGIRI UNIVERSITY, Karnataka.
 - 8.2 Institution conducting Pharm D regular courses only shall be permitted to start Pharm D (Post Baccalaureate) course.
 - 8.3 The approval and affiliation will be granted only if adequate arrangements for teaching-infrastructural facilities, hospital, accommodation, equipments, chemicals, glassware, library, teaching and non- teaching staff are provided as prescribed by Pharmacy Council of India and as required under the norms of Rajiv Gandhi University of Health Science, Karnataka (as prescribed in Appendix B).
- 9 Examination –
 - 9.1 Every year there shall be an examination to examine the students.
 - 9.2 Each examination may be held twice every year. The first examination in a year shall be the annual examination and the second examination shall be supplementary examination.
 - 9.3 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:

T A B L E S

First Year Pharm.D examination:

S.No.	Name of Subject	Maximum marks for Theory			Maximum marks for Practicals		
		Examination	Sessional	Total	Examination	Sessional	Total
1.1	Human Anatomy and Physiology	70	30	100	70	30	100
1.2	Pharmaceutics	70	30	100	70	30	100
1.3	Medicinal Biochemistry	70	30	100	70	30	100
1.4	Pharmaceutical Organic Chemistry	70	30	100	70	30	100
1.5	Pharmaceutical Inorganic Chemistry	70	30	100	70	30	100
1.6	Remedial Mathematics/ Biology**	70	30	100	70*	30*	100*
				600			600 = 1200

*For Biology.

Second Year Pharm.D examination:

S.No.	Name of Subject	Maximum marks for Theory			Maximum marks for Practicals		
		Examination	Sessional	Total	Examination	Sessional	Total
2.1	Pathophysiology	70	30	100	-	-	-
2.2	Pharmaceutical Microbiology	70	30	100	70	30	100
2.3	Pharmacognosy & Phytopharmaceuticals	70	30	100	70	30	100
2.4	Pharmacology-I	70	30	100			
2.5	Community Pharmacy	70	30	100	-	-	-
2.6	Pharmacotherapeutics-I	70	30	100	70	30	100
				600			300 = 900

Third Year Pharm.D examination:

S.No.	Name of Subject	Maximum marks for Theory			Maximum marks for Practicals		
		Examination	Sessional	Total	Examination	Sessional	Total
3.1	Pharmacology-II	70	30	100	70	30	100
3.2	Pharmaceutical Analysis	70	30	100	70	30	100
3.3	Pharmacotherapeutics-II	70	30	100	70	30	100
3.4	Pharmaceutical Jurisprudence	70	30	100	-	-	-

3.5	Medicinal Chemistry	70	30	100	70	30	100
3.6	Pharmaceutical Formulations	70	30	100	70	30	100
				600			500 = 1100

Fourth Year Pharm.D examination:

S.No.	Name of Subject	Maximum marks for Theory			Maximum marks for Practicals		
		Examination	Sessional	Total	Examination	Sessional	Total
4.1	Pharmacotherapeutics-III	70	30	100	70	30	100
4.2	Hospital Pharmacy	70	30	100	70	30	100
4.3	Clinical Pharmacy	70	30	100	70	30	100
4.4	Biostatistics & Research Methodology	70	30	100	-	-	-
4.5	Biopharmaceutics & Pharmacokinetics	70	30	100	70	30	100
4.6	Clinical Toxicology	70	30	100	-	-	-
				600			400 = 1000

Fifth Year Pharm.D examination:

S.No.	Name of Subject	Maximum marks for Theory			Maximum marks for Practicals		
		Examination	Sessional	Total	Examination	Sessional	Total
5.1	Clinical Research	70	30	100	-	-	-
5.2	Pharmacoepidemiology and Pharmacoeconomics	70	30	100	-	-	-
5.3	Clinical Pharmacokinetics & Pharmacotherapeutic Drug Monitoring	70	30	100	-	-	-
5.4	Clerkship *	-	-	-	70	30	100
5.5	Project Work (Six Months)	-	-	-	100**	-	100
				300			200 = 500

*Attending ward rounds on daily basis.

**30 marks – viva-voce (oral); 70 marks – Thesis work

First Year Pharm.D (Post Baccalaureate) examination:

S.No.	Name of Subject	Maximum marks for Theory			Maximum marks for Practicals		
		Examination	Sessional	Total	Examination	Sessional	Total
1.1	Pharmacotherapeutics-I & II	70	30	100	70	30	100
1.2	Hospital & Community Pharmacy	70	30	100	70	30	100
1.3	Clinical Pharmacy	70	30	100	70	30	100
1.4	Biostatistics & Research Methodology	70	30	100	-	-	-
1.5	Clinical Toxicology	70	30	100	-	-	-
1.6	Pharmacotherapeutics-III	70	30	100	70	30	100

1.7	Biopharmaceutics & Pharmacokinetics	70	30	100	70	30	100
				700			500 = 1200

Second Year Pharm.D (Post Baccalaureate) examination:

S.No.	Name of Subject	Maximum marks for Theory			Maximum marks for Practicals		
		Examination	Sessional	Total	Examination	Sessional	Total
2.1	Clinical Research	70	30	100	-	-	-
2.2	Pharmacoepidemiology and Pharmacoeconomics	70	30	100	-	-	-
2.3	Clinical Pharmacokinetics & Pharmacotherapeutic Drug Monitoring	70	30	100	-	-	-
2.4	Clerkship *	-	-	-	70	30	100
2.5	Project Work (Six Months)	-	-	-	100**	-	100
				300			200 = 500

*Attending ward rounds on daily basis.

**30 marks – viva-voce (oral); 70 marks – Thesis work

10 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.

11 Mode of Examinations –

11.1 Theory examination shall be of three hours

11.2 Practical examination shall be as below:

11.2.1. Four hours duration for Pharm.D – I, II and III year course

11.2.2. Six hours duration for Pharm.D – IV and V year, and for Pharm.D (Post Baccalaureate) – I and II year.

11.3 Practical examination shall also consist of a viva –voce (Oral) examination.

11.4 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' capabilities in delivering clinical pharmacy services, pharmaceutical care planning and knowledge of therapeutics shall be assessed.

12 Award of sessional marks and maintenance of records –

12.1 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.

12.2 There shall be at least two periodic sessional examinations during each academic year and the highest aggregate of any two performances shall form the basis of calculating sessional marks.

12.3 The sessional marks in practicals shall be allotted on the following basis-

12.3.1 Actual performance in the sessional examination (20 marks);

12.3.2 Day to day assessment in the practical class work, promptness, viva-voce record maintenance, etc. (10 marks).

13 Minimum marks for passing examination —

13.1 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.

13.2 A Student who fails in theory or practical examination of a subject shall re-appear both in theory and practical of the same subject.

13.3 The students securing 60% marks or above in aggregate in all subjects in a single attempt at the Pharm.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.

14 Eligibility for promotion to next year –

14.1 Failure in more than three subjects shall debar him or her from promotion to the next year classes.

14.2 The candidates who become eligible after clearing subject/subjects in the supplementary examination shall only be eligible to get admission to the next higher class along with the students of annual pattern of examination.

14.3 Students of I Pharm.D who have appeared for all the subjects are permitted to carryover not more than three failed subjects to II Pharm.D and appear for II Pharm.D examination concurrently along with failed subjects of I Pharm.D However, these Students have to pass all the failed subjects of I Pharm.D to become eligible to proceed to III Pharm.D.

14.4 Students of II Pharm.D who have appeared for all the subjects are permitted to carryover not more than three failed subjects to III Pharm.D and appear for III Pharm.D examination concurrently along with failed subjects of II Pharm.D However, these Students have to pass all the failed subjects of II Pharm.D to become eligible to proceed to IV Pharm.D.

- 14.5 Students of III Pharm.D who have appeared for all the subjects are permitted to carryover not more than three failed subjects to IV Pharm.D and appear for IV Pharm.D examination concurrently along with failed subjects of III Pharm.D

However, these Students have to pass all the failed subjects of III Pharm.D to become eligible to proceed to V Pharm.D.

- 14.6 Students of IV Pharm.D who have appeared for all the subjects are permitted to carryover not more than three failed subjects to V Pharm.D and appear for V Pharm.D examination concurrently along with failed subjects of IV Pharm.D
- 14.7 Students of I Pharm.D (Post Baccalaureate) who have appeared for all the subjects are permitted to carryover not more than three failed subjects to II Pharm.D (Post Baccalaureate) and appear for II Pharm.D (Post Baccalaureate) examination concurrently along with failed subjects of I Pharm.D (Post Baccalaureate).
- 14.8 Students shall have to pass all the examinations of Pharm.D, or Pharm.D (Post Baccalaureate) as may be applicable, to be eligible to undergo Internship.

15. Internship –

- 15.1. 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 may become capable of functioning independently.
- 15.2. Every student has to undergo one-year internship as per Appendix-C to these regulations.

16. Award of Degree –

- 16.1. Every student who has passed the examinations for the Pharm.D or Pharm.D. (Post Baccalaureate) as the case may be, and has successfully completed one year internship, shall be eligible for the award of degree.

17. Practical training

- 17.1. Hospital posting — Every student shall be posted in constituent hospital for a period of not less than seventy five 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 daily basis as a part of clerkship. Theory teaching may be scheduled in the afternoon.
- 17.2. Project work—
- 17.2.1. 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.
- 17.2.2. Project work shall comprise of objectives of the work, methodology, results,

discussions and conclusions.

17.3. Objectives of project work— The main objectives 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.

17.4. Methodology— To complete the project work following methodology shall be adopted, namely:—

- (i) students shall work in groups of not more than four under an authorised teacher;
- (ii) The Head of the Department or Head of the Institution shall approve project topic;
- (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) The institutional ethics committee shall approve project work;
- (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.

17.5. Reporting—

17.5.1. 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 teacher, Head of the Department as well as by the Head of the Institution

17.5.2. Project report shall be computer typed in double space using Times Roman font on A4 paper. The title shall be in bold with font size 18, sub-titles in bold with font size 14 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 authorised teacher with font size 14.

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

17.6. 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 thirty 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 done on the following items:

	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.

APPENDIX-A

PHARM.D. SYLLABUS

First Year

1.1 HUMAN ANATOMY & PHYSIOLOGY (THEORY)

Theory: 3 Hrs. /Week

- 1. Scope and Objectives:** This course is designed to impart a fundamental knowledge on the structure and functions of the human body. It also helps in understanding both homeostasis mechanisms and homeostatic imbalances of various body systems. Since a medicament, which is produced by pharmacist, is used to correct the deviations in human body, it enhances the understanding of how the drugs act on the various body systems in correcting the disease state of the organs.
- 2. Upon completion of the course the student shall be able to:**
 - a. describe the structure (gross and histology) and functions of various organs of the human body;
 - b. describe the various homeostatic mechanisms and their imbalances of various systems;
 - c. identify the various tissues and organs of the different systems of the human body;
 - d. perform the hematological tests and also record blood pressure, heart rate, pulse and Respiratory volumes;
 - e. appreciate coordinated working pattern of different organs of each system; and
 - f. appreciate the interlinked mechanisms in the maintenance of normal functioning (homeostasis) of human body
- 3. Course materials:**

Text books

 - a. Tortora Gerard J. and Nicholas, P. Principles of anatomy and physiology Publisher Harpercollins college New York.
 - b. Wilson, K.J.W. Ross and Wilson's foundations of anatomy and physiology. Publisher: Churchill Livingstone, Edinburg.

Reference books

 - a. Guyton arthur, C. Physiology of human body. Publisher: Holtsaunders.
 - b. Chatterjee,C.C. Human physiology. Volume 1&11. Publisher: medical allied agency, Calcutta.
 - c. Peter L. Williams, Roger Warwick, Mary Dyson and Lawrence, H.
 - d. Gray's anatomy. Publisher:Churchill Livingstone, London

4. Lecture wise program: Topics

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Scope of anatomy and physiology, basic terminologies used in this subject (Description of the body as such planes and terminologies)	1 Hour
2	Structure of cell – its components and their functions.	1 Hour
3	Elementary tissues of the human body: epithelial, connective, Muscular and nervous tissues-their sub-types and characteristics	2 Hours
4	Osseous System a) Structure, composition of Bones and functions of the skeleton. b) Classification of joints, Types of movements of joints and disorders of joints (Definitions only)	2 Hours
5	Haemopoietic system a) Composition and functions of blood b) Haemopoiesis and disorders of blood components (definition of disorder) c) Blood groups d) Clotting factors and mechanism e) Platelets and disorders of coagulation	6 Hours
6	Lymph a) Lymph and lymphatic system, composition, formation and circulation. b) Spleen: structure and functions, Disorders c) Disorders of lymphatic system (definition only)	3 Hours
7	Cardiovascular system a) Anatomy and functions of heart b) Blood vessels and circulation (Pulmonary, coronary and systemic circulation) c) Electrocardiogram (ECG) d) Cardiac cycle and heart sounds e) Blood pressure – its maintenance and regulation f) Definition of the following disorders: Hypertension, Hypotension, Arteriosclerosis, Atherosclerosis, Angina, Myocardial infarction, Congestive heart failure, Cardiac arrhythmias	8 Hours
8	Respiratory system a) Anatomy of respiratory organs and functions b) Mechanism / physiology of respiration and regulation of respiration c) Transport of respiratory gases d) Respiratory volumes and capacities, and Definition of: Hypoxia, Asphyxia, Dybarism, Oxygen therapy and resuscitation.	5 Hours
9	Digestive system a) Anatomy and physiology of GIT b) Anatomy and functions of accessory glands of GIT c) Digestion and absorption d) Disorders of GIT (definitions only)	7 Hours

10	Nervous system a) Definition and classification of nervous system b) Anatomy, physiology and functional areas of cerebrum c) Anatomy and physiology of cerebellum d) Anatomy and physiology of mid brain e) Thalamus, hypothalamus and Basal Ganglia f) Spinal cord: Structure & reflexes – mono-poly-planter g) Cranial nerves – names and functions h) ANS – Anatomy & functions of sympathetic & parasympathetic N.S.	12 Hours
11	Urinary system a) Anatomy and physiology of urinary system b) Formation of urine c) Renin Angiotensin system – Juxtaglomerular apparatus - acid base Balance d) Clearance tests and micturition	6 Hours
12	Endocrine system a) Pituitary gland b) Adrenal gland c) Thyroid and Parathyroid glands d) Pancreas and gonads e) Penal gland f) Thymus	5 Hours
13	Reproductive system a) Male and female reproductive system b) Their hormones – Physiology of menstruation c) Spermatogenesis & Oogenesis d) Sex determination (genetic basis) e) Pregnancy and maintenance and parturition f) Contraceptive devices	7 Hours
14	Sense organs a) Eye b) Ear c) Skin d) Tongue & Nose	6 Hours
15	Skeletal muscles a) Histology b) Physiology of Muscle contraction c) Physiological properties of skeletal muscle and their disorders (definitions)	2 Hours
16	Sports physiology a) Muscles in exercise, Effect of athletic training on muscles and muscle performance, b) Respiration in exercise, CVS in exercise, Body heat in exercise, Body fluids and salts in exercise, c) Drugs and athletics	2 Hours

1.1 HUMAN ANATOMY & PHYSIOLOGY (PRACTICAL)

Practical : 3 Hrs./Week

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

Course materials:

Text books

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

Ranade VG, Text book of practical physiology, Latest edition, Publisher: PVG, Pune
Anderson Experimental Physiology, Latest edition, Publisher: NA

List of Experiments:

1. Study of tissues of human body
 - (a) Epithelial tissue.
 - (b) Muscular tissue.
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 & its clinical and diagnostic importance
5. Determination of R.B.C. count & its clinical and diagnostic importance
6. Determination of differential count & its clinical and diagnostic importance.
7. Determination of following with their clinical and diagnostic importance
 - (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 gastrocnemius sciatic nerve preparation.
- 14. To record simple summation curve using gastrocnemius sciatic nerve preparation.
- 15. To record simple effect of temperature using gastrocnemius sciatic nerve preparation.
- 16. To record simple effect of load & after load using gastrocnemius sciatic nerve preparation.
- 17. To record simple fatigue curve using gastrocnemius sciatic nerve preparation.

Scheme of Practical Examination:

	Sessionals	Annual
Identification	04	10
Synopsis	04	10
Major Experiment	07	20
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	04hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

1.2 PHARMACEUTICS (THEORY)

Theory: 2 Hrs. /Week

1. Scope and objectives: This course is designed to impart a fundamental knowledge on the art and science of formulating different dosage forms. It prepares the students for most basics of the applied field of pharmacy.

2. Upon the completion of the course the student should be able to:

- know the formulation aspects of different dosage forms;
- do different pharmaceutical calculation involved in formulation;
- formulate different types of dosage forms; and
- appreciate the importance of good formulation for effectiveness.

3. Course materials:

Text books

- Cooper and Gunns Dispensing for pharmacy students.
- A text book Professional Pharmacy by N.K.Jain and S.N.Sharma.

Reference books

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

4. Lecture wise programme:

Topics

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	a. Introduction to dosage forms - classification and definitions b. Prescription: definition, parts and handling c. Types of prescriptions and pricing of prescriptions d. Posology: Definition, Factors affecting dose selection. Calculation of children and infant doses.	8 +1 Hours
2	Historical back ground and development of profession of pharmacy and pharmaceutical industry in brief.	2 Hours
3	Development of Indian Pharmacopoeia and introduction to other Pharmacopoeias such as BP, USP, European Pharmacopoeia, Extra pharmacopoeia and Indian national formulary.	2 Hours

4	Weights and measures, Calculations involving percentage solutions, allegation, proof spirit, isotonic solutions etc.	2 Hours
5	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.	4 Hours
6	Monophasic Dosage forms: Theoretical aspects of formulation including adjuvant like stabilizers, colorants, flavours with examples. Study of Monophasic liquids like gargles, mouth washes, Throat paint, Ear drops, Nasal drops, Liniments and lotions, Enemas and collodions.	5 Hours
7	Biphasic dosage forms: Suspensions and emulsions, Definition, advantages and disadvantages, classification, test for the type of emulsion, formulation, stability and evaluation.	8 Hours
8	Suppositories and pessaries: Definition, advantages and disadvantages, types of base, method of preparation, Displacement value and evaluation.	5 Hours
9	Galenicals: Definition, equipment for different extraction processes like infusion, Decoction, Maceration and Percolation, methods of preparation of spirits, tinctures and extracts.	5 Hours
10	Pharmaceutical calculations.	2 Hours
11	Surgical aids: Surgical dressings, absorbable gelatin sponge, sutures, ligatures and medicated bandages.	4 Hours
12	Incompatibilities: Introduction, classification and methods to overcome the incompatibilities.	3 Hours

1.2 PHARMACEUTICS (PRACTICAL)

Practical: 3 Hrs./Week

List of Experiments:

1. Syrups

- a) Simple Syrup I.P
- b) Syrup of Ephedrine Hcl NF
- c) Syrup Vasaka IP
- d) Syrup of ferrous Phosphate IP
- e) Orange Syrup

2. Elixir

- a) Piperizine citrate elixir BP
- b) Cascara elixir BPC
- c) Paracetamol elixir BPC

3. Linctus

- a) Simple Linctus BPC
- b) Pediatric simple Linctus BPC

4. Solutions

- a) Solution of cresol with soap IP
- b) Strong solution of ferric chloride BPC
- c) Aqueous Iodine Solution IP
- d) Strong solution of Iodine IP
- e) Strong solution of ammonium acetate IP

5. Liniments

- a) Liniment of turpentine IP*
- b) Liniment of camphor IP

6. Suspensions*

- a) Calamine lotion
- b) Magnesium Hydroxide mixture BP

7. Emulsions*

- a) Cod liver oil emulsion
- b) Liquid paraffin emulsion

8. Powders

- a) Eutectic powder
- b) Explosive powder
- c) Dusting powder

d) Insufflations

9. Suppositories

- a) Boric acid suppositories
- b) Chloral suppositories

10. Incompatibilities

- a) Mixtures with Physical
- b) Chemical & Therapeutic incompatibilities

11. Salicylic acid collodion

* colourless bottles required for dispensing - Paper envelope (white), butter paper and white paper required for dispensing.

Scheme of Practical Examination:

	Sessionals	Annual
Synopsis	05	15
Major Experiment	10	25
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	04hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

1.3 MEDICINAL BIOCHEMISTRY (THEORY)

Theory : 3 Hrs. /Week

1. Scope of the Subject: Applied biochemistry deals with complete understanding of the molecular level of the chemical process associated with living cells. Clinical chemistry deals with the study of 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.

2. Objectives of the Subject (Know, do, appreciate):

The objective of the present course is providing biochemical facts and the principles to the students of pharmacy. Upon completion of the subject student shall be able to –

- understand the catalytic activity of enzymes and importance of isoenzymes in diagnosis of 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; and
- do the qualitative analysis and determination of biomolecules in the body fluids.

Text books (Theory)

- Harpers review of biochemistry - Martin
- Text book of biochemistry – D.Satyanarayana
- Text book of clinical chemistry- Alex kaplan &Laverve L.Szabo

Reference books (Theory)

- Principles of biochemistry -- Lehninger
- Text book of biochemistry -- Ramarao
- Practical Biochemistry-David T.Plummer.
- Practical Biochemistry-Pattabhiraman.

3.Lecture wise programme :Topics

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Introduction to biochemistry: Cell and its biochemical organization, transport process across the cell membranes. Energy rich compounds; ATP, Cyclic AMP and their biological significance.	3 Hours

2	Enzymes: Definition; Nomenclature; IUB classification; Factor affecting enzyme activity; Enzyme action; properties of enzymes, enzyme inhibition. Isoenzymes and their therapeutic and diagnostic applications; Coenzymes and their biochemical role and deficiency diseases.	12 (+1) hours
3	Carbohydrate metabolism: introduction to carbohydrates 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.	13 Hours (+1 hr)
4	Lipid metabolism: introduction to lipids, 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 (Atherosclerosis, fatty liver, hypercholesterolemia).	10 Hours (+1 hr)
5	Biological oxidation: Coenzyme system involved in Biological oxidation. Electron transport chain (its mechanism in energy capture; regulation and inhibition); Uncouplers of ETC; Oxidative phosphorylation;	5 Hours
6	Protein and amino acid metabolism: introduction to protein and amino acid, 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.	12 Hours (+1 hr)
7	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.	12 Hours
8	Introduction to clinical chemistry: Cell; composition; malfunction; Roll of the clinical chemistry laboratory.	2 Hours
9	The kidney function tests: Role of kidney; Laboratory tests for normal function includes a) Urine analysis (macroscopic and physical examination, quantitative and semiquantitative tests.) b) Test for NPN constituents. (Creatinine /urea clearance, determination of blood and urine creatinine, urea and uric acid) c) Urine concentration test d) Urinary tract calculi. (stones)	2 Hours

10	Liver function tests: Physiological role of liver, metabolic, storage, excretory, protective, circulatory functions and function in blood coagulation. a) Test for hepatic dysfunction-Bile pigments metabolism. b) Test for hepatic function test- Serum bilirubin, urine bilirubin, and urine urobilinogen. c) Dye tests of excretory function. d) Tests based upon abnormalities of serum proteins. Selected enzyme tests	2 Hours
11	Lipid profile tests: Lipoproteins, composition, functions. Determination of serum lipids, total cholesterol, HDL cholesterol, LDL cholesterol and triglycerides.	2 Hours
12	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)	2 Hours
13	Electrolytes: Body water, compartments, water balance, and electrolyte distribution. Determination of sodium, calcium potassium, chlorides, bicarbonates in the body fluids.	2 Hours

1.3 MEDICINAL BIOCHEMISTRY (PRACTICAL)

Practical : 3 Hrs./Week

Title of the Experiment:

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.

Scheme of Practical Examination:

	Sessionals	Annual
Synopsis	05	15
Major Experiment	10	25
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	04hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

1.4 PHARMACEUTICAL ORGANIC CHEMISTRY (THEORY)

Theory : 3 Hrs. /Week

- 1. Scope and objectives:** This course is designed to impart a very good knowledge about
 - a. IUPAC/Common system of nomenclature of simple organic compounds belonging to different classes of organic compounds;
 - b. Some important physical properties of organic compounds;
 - c. Free radical/ nucleophilic [alkyl/ acyl/ aryl] / electrophilic substitution, free radical/ nucleophilic / electrophilic addition, elimination, oxidation and reduction reactions with mechanism, orientation of the reaction, order of reactivity, stability of compounds;
 - d. Some named organic reactions with mechanisms; and
 - e. Methods of preparation, test for purity, principle involved in the assay, important medicinal uses of some important organic compounds.
- 2. Upon completion of the course the student shall be able to:**
 - a. Gain knowledge about the physical properties of organic compounds
 - b. Gain knowledge about the chemical properties of organic compounds
 - c. Gain knowledge about the synthesis of organic compounds
 - d. Gain knowledge about the nomenclature of organic compounds
 - e. Gain knowledge about the medicinal uses of pharmaceutically important organic compounds
- 3. Course materials:**

Text books

 - a. T.R.Morrison and R. Boyd - Organic chemistry,
 - b. Bentley and Driver-Text book of Pharmaceutical chemistry
 - c. I.L.Finer- Organic chemistry, the fundamentals of chemistry

Reference books

 - a. Organic chemistry – J.M.Cram and D.J.Cram
 - b. Organic chemistry- Brown
 - c. Advanced organic chemistry- Jerry March, Wiley
 - d. Organic chemistry- Cram and Hammered, Pine Hendrickson

4. Lecture wise programme :**Topics**

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Structures and Physical properties: a) 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, b) Acids and bases, Lowry bronsted and Lewis theories c) Isomerism	3 Hours
2	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.	8 Hours
3	Synthesis and Free radicals chain reactions of alkane : Mechanism, relative reactivity and stability	3 +1 Hours
4	Alicyclic compounds : Preparations of cyclo alkanes, Bayer strain theory and orbital picture of angle strain.	3 Hours
5	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, SN2 versus SN1 solvolyses, nucleophilic assistance by the solvents.	8 Hours
6	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, orientation and reactivity, E2 versus E1, elimination versus substitution, dehydration of alcohol, ease of dehydration, acid catalysis, reversibility, orientation.	7 Hours
7	Electrophilic and free radicals addition: Reactions at carbon -carbon, double bond, electrophile, hydrogenation, heat of hydrogenation and stability of alkenes, markownikoff rule, addition of hydrogen halides, addition of hydrogen bromides, peroxide effect, electrophilic addition, mechanism, rearrangement, absence of hydrogen exchange, orientation and reactivity, addition of halogen, mechanism, halohydrin formation, mechanism of free radicals addition, mechanism of peroxide initiated addition of hydrogen bromide, orientation of free addition, additions of carbene to alkene, cyclo addition reactions.	5 Hours

8	Carbon-carbon double bond as substituents: synthesis of alkenes Free radical halogenations of alkenes, comparison of free radical substitution with free radical addition, free radical substitution in alkenes, orientation and reactivity, allylic rearrangements.	2 +1 Hours
9	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, nucleophilic substitution in allylic substrate, SN1 reactivity, allylic rearrangement, resonance stabilisation of allyl cation, hyper conjugation, nucleophilic substitution in allylic substrate, SN2 nucleophilic substitution 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.	6 Hours
10	Electrophilic 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 craft acylation, reactivity and orientation, activating and deactivating O,P,M directing groups, electron release via resonance, effect of halogen on electrophilic aromatic substitution in alkyl benzene, side chain halogenation of alkyl benzene, resonance stabilization of benzyl radical.	8 Hours
11	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 carboxyl group, comparison of alkyl nucleophilic substitution with acyl nucleophilic substitution.	4 Hours
12	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.	4 Hours
13	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.	5 hours
14	Nucleophilic aromatic substitution: Bimolecular displacement mechanisms, orientation, comparison of aliphatic nucleophilic substitution with that of aromatic.	2 Hours
15	Oxidation reduction reaction and redox reaction	2 +1 Hours

16	Study of the following official compounds- preparation, test for purity, assay and medicinal uses of Chlorbutol, Dimercaprol, Glyceryl trinitrate, Urea, Ethylene diamine dihydrate, Vanillin, Paraldehyde, Ethylene chloride, Lactic acid, Tartaric acid, citric acid, salicylic acid, aspirin, methyl salicylate, ethyl benzoate, benzyl benzoate, dimethyl phthalate, sodium lauryl sulphate, saccharin sodium, mephensin.	5 Hours
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1.4 PHARMACEUTICAL ORGANIC CHEMISTRY (PRACTICAL)

Practical : 3 Hrs./Week

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

1. Acetanilide / 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-naphthol (Diazotisation and coupling)
6. Benzoic acid / salicylic acid (Hydrolysis of ester)
7. M-dinitro benzene (Nitration)
8. 9, 10 – Anthraquinone (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-chlorotoluene
14. Preparation of cyclohexanone from cyclohexanol

II. 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.

III. Introduction to the use of stereo models:

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

Scheme of Practical Examination:

	Sessionals	Annual
Synopsis	05	15
Major Experiment	10	25
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	04hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

1.5 PHARMACEUTICAL INORGANIC CHEMISTRY (THEORY)

Theory : 2 Hrs./Week

1. **Scope and objectives:** This course mainly deals with fundamentals of Analytical chemistry and also the study of inorganic pharmaceuticals regarding their monographs and also the course deals with basic knowledge of analysis of various pharmaceuticals.
2. **Upon completion of the course student shall be able to:**
 - a. understand the principles and procedures of analysis of drugs and also regarding the application of inorganic pharmaceuticals;
 - b. know the analysis of the inorganic pharmaceuticals their applications; and
 - c. appreciate the importance of inorganic pharmaceuticals in preventing and curing the disease.
 - d. basic fundamentals principles or theory of different titrimetric analysis
 - e. explain the mechanism of action of selected inorganic therapeutic and diagnostic agents.
 - f. understanding the biological functions of essential trace elements
 - g. describing the methods of preparation and properties of different inorganic compounds.

3. Course materials:

Text books

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

Reference books

- a. Inorganic Pharmaceutical Chemistry by Anand & Chetwal
- b. Pharmaceutical Inorganic chemistry by Dr.B.G.Nagavi
- c. Analytical chemistry principles by John H. Kennedy
- d. I.P.1985 and 1996, Govt. of India, Ministry of health

4. Lecture wise programme:

Topics

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Errors: Errors in quantitative analysis, classification of errors, concept of accuracy and precision, treatment of analytical, , methods to minimize the error	3+1 Hours

2	Volumetric analysis: Principle of volumetric analysis, different methods of analysis, different methods of expressing concentrations of solutions, Primary and secondary standards, applications of volumetric analysis in pharmaceutical industry	4 + 2 Hours
3	Acid-base titrations: Acid-base concepts, relative strength of acids and bases, law of mass action, common ion effect, ionic product of water, Henderson-Hasselbalch equation, buffer solutions, theory of indicators, neutralization curves, choice of indicators, mixed and universal indicators.	3+1 Hours
4	Redox titrations: Concepts of oxidation and reduction reactions. Redox reactions, theory of redox titrations, redox indicators, iodometry, iodimetry, titrations involving ceric sulphate, potassium iodate, potassium bromate, titanous chloride, Potassium permanganate.	3 Hours
5	Non aqueous titrations: Theoretical basis, types of solvents, preparations and standardization of titrant solutions, titration of weak acid, weak bases and indicators. Standardization of perchloric acid, lithium and sodium methoxide, tetrabutyl ammonium hydroxide.	2 +3 Hours
6	Precipitation titrations: Introduction, types of precipitation titrations, end point detection.	2 Hours
7	Complexometric titrations: Introduction, principle, types of titrations, end-point detection.	2 Hours
8	Theory of Indicators	1 Hour
9	Gravimetry: Basic concepts, precipitation techniques, co-precipitations, post-precipitation, various steps involved in gravimetric analysis, pharmaceutical applications.	2+2 Hours
10	Limit tests: Definition, importance, general procedure for limit test for chlorides, sulphates, iron, arsenic, lead and heavy metals, impurities in pharmaceutical substances and different sources of impurities	4 +2 Hours
11	Medicinal gases: Preparation and uses of the following oxygen, carbon dioxide, Helium, Nitrogen and Nitrous oxide. Method of preparation, assay, storage conditions and uses of Inorganic compounds listed in IP belonging to the following categories	2 Hours
12	Acidifiers: Dilute HCl, Sodium Phosphate, Ammonium chloride.	1+1 Hour
13	Antacids: Classification, qualities of an ideal antacid, side effects, advantages, combination therapy, acid neutralizing capacity, sodium bicarbonate, Potassium citrate, Aluminium hydroxide gel, Dried aluminium hydroxide gel, magnesium hydroxide, Light and Heavy magnesium trisilicate, Light and Heavy magnesium carbonate, Calcium carbonate, Magaldrate and Bismuth carbonate.	3 +2 Hours

14	Cathartics: Magnesium hydroxide, Magnesium sulphate, Magnesium carbonate, Sodium phosphate.	2 +1Hours
15	Electrolyte replenishers: Electrolytes used for replacement therapy: sodium chloride, potassium chloride, calcium chloride, calcium gluconate. Electrolytes used in acid-base therapy: Sodium acetate, potassium acetate, sodium bicarbonate, potassium bicarbonate, sodium citrate, sodium lactate, ammonium chloride, Electrolyte combination therapy, compound sodium chloride solution, sodium chloride injection and oral rehydration salt.	3 +2Hours
16	Essential Trace elements: Definition, physiological role of Iron, Copper, Zinc, Chromium, Manganese, Molybdenum, Selenium, Sulphur and Iodine.	2+2 Hours
17	Antimicrobials: Hydrogen peroxide, Potassium permanganate, Chlorinated Lime, Iodine, Boric acid, Silver nitrate, Selenium Sulphide.	2 Hours
18	Pharmaceutical Aids: Sodium bisulphate, sodium metabisulphite, bentonite, magnesium stearate, zinc stearate, aluminium sulphate, sodium carboxy methylcellulose, purified water, water for injection and sterile water for injection.	2 +2 Hours
19	Dental Products: Anti-caries agents: Role of Fluorides as anti-caries agents, Sodium Fluoride. Dentifrices: Calcium carbonate, dibasic calcium phosphate, Zinc chloride.	2+2 Hours
20	Miscellaneous compounds Sclerosing agents: Hypertonic saline, Sodium tetradecyl sulphate. Expectorants: Potassium citrate and Potassium iodide. Sedative: Potassium bromide. Antidotes: Sodium nitrite, Sodium thiosulphate and Charcoal Respiratory stimulant: Ammonium carbonate.	3+2 Hours
21	Radio Pharmaceuticals: Introduction, measurement of radioactivity, clinical applications and dosage, hazards and precautions.	2 +1Hours

1.5 PHARMACEUTICAL INORGANIC CHEMISTRY (PRACTICAL)

Practical : 3 Hrs./Week

1. 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

2. Assays (10 exercises)

- a. Ammonium chloride- Acid-base titration
- b. Ferrous sulphate- Cerimetry
- c. Copper sulphate- Iodometry
- d. Calcilugluconate-Complexometry
- e. Hydrogen peroxide – Permanganometry
- f. Sodium benzoate – Nonaqueous titration
- g. Sodium chloride – Modified volhard's method
- h. Assay of KI – KIO₃ titration
- i. Gravimetric estimation of barium as barium sulphate
- j. Sodium antimony gluconate or antimony potassium tartarate

3. Estimation of mixture (Any two exercises)

- a. Sodium hydroxide and sodium carbonate
- b. Boric acid and Borax
- c. Oxalic acid and sodium oxalate

4. Test for identity (Any three exercises)

- a. Sodium bicarbonate
- b. Barium sulphate
- c. Ferrous sulphate
- d. Potassium chloride

5. 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

6. Preparations (Any two exercises)

- a. Boric acids
- b. Potash alum
- c. Calcium lactate
- d. Magnesium sulphate

Scheme of Practical Examination:

	Sessionals	Annual
Synopsis	05	15
Major Experiment	10	25
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	04hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

1.6 REMEDIAL MATHEMATICS/BIOLOGY (THEORY)

Theory: 3 Hrs. /Week

REMEDIAL MATHEMATICS:

1. **Scope and objectives:** This is an introductory course in mathematics. This subjects deals with the introduction to matrices, determinants, trigonometry, analytical geometry, differential calculus, integral calculus, differential equations, laplace transform.
2. **Upon completion of the course the student shall be able to : –**
 - a. Know Trigonometry, Analytical geometry, Matrices, Determinant, Integration, Differential equation, Laplace transform and their applications;
 - b. solve the problems of different types by applying theory; and
 - c. appreciate the important applications of mathematics in pharmacy.

3. Course materials:

Text books

- a. Differential calculus By Shantinakaran
- b. Text book of Mathematics for second year pre-university by Prof.B.M.Sreenivas

Reference books

- a. Integral calculus By Shanthinarayan
- b. Engineering mathematics By B.S.Grewal
- c. Trigonometry Part-I By S.L.Loney

4. Lecture wise programme

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Algebra: Determinants, Matrices	
2	Trigonometry: Sides and angles of a triangle, solution of triangles	
3	Analytical Geometry: Points, Straight line, circle, parabola	

4	Differential calculus: Limit of a function, Differential calculus, Differentiation of a sum, Product, Quotient Composite, Parametric, Exponential, trigonometric and Logarithmic function. Successive differentiation, Leibnitz's theorem, Partial differentiation, Euler's theorem on homogeneous functions of two variables	
5	Integral Calculus: Definite integrals, integration by substitution and by parts, Properties of definite integrals.	
6	Differential equations: Definition, order, degree, variable separable, homogeneous, Linear, heterogeneous, linear, differential equation with constant coefficient, simultaneous linear equation of second order.	
7	Laplace transform: Definition, Laplace transform of elementary functions, Properties of linearity and shifting.	

BIOLOGY:

1. **Scope and objectives:** This is an introductory course in Biology, which gives detailed study of natural sources such as plant and animal origin. This subject has been introduced to the pharmacy course in order to make the student aware of various naturally occurring drugs and its history, sources, classification, distribution and the characters of the plants and animals. This subject gives basic foundation to Pharmacognosy.
2. Upon completion of the course the student shall be able to:
 - a. Gains the knowledge on Plant tissues, Plant kingdom and its classification, morphology of plants and Plant physiology.
 - b. Identify the Animal cell, animal tissues, poisonous animals
 - c. Knows the General organization of mammals.

3. Course materials:**Text books**

- a. Text book of Biology by S.B.Gokhale
- b. A Text book of Biology by Dr.Thulajappa and Dr. Seetaram.

Reference books

- a. A Text book of Biology by B.V.Sreenivasa Naidu
- b. A Text book of Biology by Naidu and Murthy
- c. Botany for Degree students By A.C.Dutta.
- d. Outlines of Zoology by M.Ekambaranatha ayyer and T.N.Ananthakrishnan.
- e. A manual for pharmaceutical biology practical by S.B.Gokhale and C.K.Kokate.

4. Lecture wise programme:**Topic**

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
PART-A		
1	Introduction	
2	General organization of plants and its inclusions	
3	Plant tissues	
4	Plant kingdom and its classification	
5	Morphology of plants	
6	Root, Stem, Leaf and Its modifications	
7	Inflorescence and Pollination of flowers	
8	Morphology of fruits and seeds	
9	Plant physiology	

10	Taxonomy of Leguminosae, umbelliferae, Solanaceae, Lilliacae, Zinziberaceae, Rubiaceae	
11	Study of Fungi, Yeast, Penicillin and Bacteria	
PART-B		
1	Study of Animal cell	
2	Study animal tissues	
3	Detailed study of frog	
4	Study of Pisces, Reptiles, Aves	
5	General organization of mammals	
6	Study of poisonous animals	

1.6 BIOLOGY (PRACTICAL)

Practical : 3 Hrs./Week

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

Scheme of Practical Examination:

	Sessionals	Annual
Identification	04	10
Synopsis	04	10
Major Experiment	07	20
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	04hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance)

Second Year:

2.1 PATHOPHYSIOLOGY (THEORY)

Theory : 3 Hrs. /Week

- 1. Scope of the Subject:** This course is designed to impart a thorough knowledge of the relevant aspects of pathology of various conditions with reference to its pharmacological applications, and understanding of basic Pathophysiological mechanisms. Hence it will not only help to study the syllabus of pathology, but also to get baseline knowledge of its application in other subject of pharmacy.
- 2. Objectives of the Subject:** Upon completion of the subject student shall be able to –
 - a. describe the etiology and pathogenesis of the selected disease states;
 - b. name the signs and symptoms of the diseases; and
 - c. mention the complications of the diseases.

Text books (Theory)

- a. Pathologic basis of disease by- Cotran, Kumar, Robbins
- b. Text book of Pathology- Harsh Mohan
- c. Text book of Pathology- Y.M. Bhide

Reference books (Theory)

- a. Clinical Pharmacy and Therapeutics; Second edition; Roger Walker; Churchill Livingstone publication

3. Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Basic principles of cell injury and Adaptation a) Causes, Pathogenesis and morphology of cell injury b) Abnormalities in lipoproteinaemia, glycogen infiltration and glycogen infiltration and glycogen storage diseases	6 Hours
2	Inflammation a) Pathogenesis of acute inflammation, Chemical mediators in inflammation, Types of chronic inflammation b) Repairs of wounds in the skin, factors influencing healing of wounds	7 Hours
3	Diseases of Immunity a) Introduction to T and B cells b) MHC proteins or transplantation antigens	15 Hours

	c) 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) - Amyloidosis	
4	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.	10 Hours
5	Types of shock, mechanisms, stages and management	2 Hours
6	Biological effects of radiation	1 Hour
7	Environmental and nutritional diseases a) Air pollution and smoking- SO₂, NO, NO₂, and CO b) Protein calorie malnutrition, vitamins, obesity, pathogenesis of starvation.	5 Hours
8	Pathophysiology of common diseases a) Parkinsonism b) Schizophrenia c) Depression and mania d) Hypertension, e) Stroke (ischaemic and hemorrhage) f) Angina, CCF, Atherosclerosis, Myocardial infarction g) Diabetes Mellitus h) Peptic ulcer and inflammatory bowel diseases i) Cirrhosis and Alcoholic liver diseases j) Acute and chronic renal failure k) Asthma and chronic obstructive airway diseases	19 Hours
9	Infectious diseases: Sexually transmitted diseases (HIV, Syphilis, Gonorrhea), Urinary tract infections, Pneumonia, Typhoid, Tuberculosis, Leprosy, Malaria Dysentery (bacterial and amoebic), Hepatitis- infective hepatitis.	10 Hours

4. 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.

2.2 PHARMACEUTICAL MICROBIOLOGY (THEORY)

Theory : 3 Hrs. /Week

1. **Scope of the Subject:** Microbiology has always been an essential component of pharmacy curriculum. This is because of the relevance of microbiology to pharmaceutical sciences and more specifically to pharmaceutical industry. Pharmaceutical biotechnology is the logical extension of pharmaceutical microbiology, which is expected to change the complete drug product scenario in the future.

This **subject deals about the history of microbiology** along with the various aspects of microorganisms, its classification, morphology, laboratory cultivation identification and maintenance. It also deals with sterilization of pharmaceutical products, equipment, media, various methods of disinfection, etc. **In addition to this the subject also enumerate** the immunological preparations, diseases its transmission, diagnosis, control and diagnostic tests.

2. Objectives of the Subject:

Upon completion of the subject student shall be able to –

- a. Know the history of microbiology and contribution of various scientists in the field of microbiology;
- b. Understand the anatomy, identification, isolation, growth factors and sterilization of microorganisms;
- c. Familiar with the cultivation and identification of the microorganisms in the laboratory;
- d. Acquaint with various disinfectants and their mode of action;
- e. Understand the different type of immunity, antigens, antibodies, antigen-antibody reactions and vaccines;
- f. Know the causative microorganism, mode of transmission, sign and symptoms, diagnostic tests and treatment aspects very common infectious diseases.

Text books (Theory)

- a. Vanitha Kale and Kishor Bhusari – Applied Microbiology – Himalaya Publishing House, Mumbai.
- b. Mary Louis Turgeon – Immunology and Serology in Laboratory Medicines - 2nd edition, 1996 Mosby- Year book Inc, St. Louis Missouri 63146.
- c. Harsh Mohan – Text book of Pathology – 3rd edition, 1998, B-3 Ansari road, Daryaganj, New Delhi.

Reference books (Theory)

- a. Prescott L.M., Jarley G.P, Klein D.A – Microbiology – 2nd edition, Mc Graw Hill Company Inc
- b. Rawlins E.A.. Bentley's Text Book of Pharmaceutics, Bailliere Tindals, 24 -28 London 1988
- c. Forbisher — Fundamentals of Microbiology_ Philadelphia W.B. Saunders.
- d. War Roitt, Jonathan Brostoff, David male, — Immunology_3rd edition 1996, Mosby- year book Europe Ltd, London.
- e. Pharmacopoeia of India, Govt of India, 1996.

3 Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Introduction to the science of microbiology. Major divisions of microbial world and Relationship among them.	3 Hours
2	Different methods of classification of microbes and study of Bacteria, Fungi, virus, Rickettsiae, Spirochetes.	5 Hours
3	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.	6 Hours
4	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.	4 Hours
5	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.	12 Hours
6	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.	10 Hours
7	Immunology- Immunity, Definition, Classification, General principles of natural immunity, Phagocytosis, acquired immunity (active and passive). 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.	13 Hours
8	Diagnostic tests : Schick's Test, Elisa test, Western Blot test, Southern Blot PCR Widal, QBC, Mantoux Peripheral smear. Study of malarial parasite	8 Hours
9	Microbial culture sensitivity Testing: Interpretation of results Principles and methods of different microbiological assays, microbiological assay of Penicillin, Streptomycin and vitamin B2 and B12. Standardisation of vaccines and sera.	6 Hours
10	Study of infectious diseases: Typhoid, Tuberculosis, Malaria, Cholera, Hepatitis, Meningitis, Syphilis & Gonorrhea and HIV.	8 Hours

2.2 PHARMACEUTICAL MICROBIOLOGY (PRACTICAL)

Practical: 3 Hrs./Week

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 sterilization 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.
 - b. 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.

Scheme of Practical Examination:

	Sessionals	Annual
Synopsis	05	15
Major Experiment	10	25
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	04hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

2.3 PHARMACOGNOSY & PHYTOPHARMACEUTICALS (THEORY)

Theory: 3 Hrs. /Week

1. **Scope and 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, distribution, method of cultivation, active constituents, medicinal uses, identification tests, preservation methods, substitutes and adulterants.
2. **Upon completion of the course student shall be able to:**
 - a. Know the various methods of classification of crude drugs;
 - b. Understand the various factors affecting the cultivation, collection , processing and storage of crude drugs;
 - c. Appreciate the applications of primary and secondary metabolites of the plant.

3. Course materials:

Text books

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

Reference books

- a. Pharmacognosy by Brady &Tyler.E.
- b. Pharmacognosy by T.E.Wallis.
- c. Pharmacognosy by C.S. Shah & Qadery.
- d. Pharmacognosy by M.A. Iyengar.

4. Lecture wise programme:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Introduction	2 Hours
2	Definition, history and scope of Pharmacognosy.	3 Hours
3	Classification of crude drugs.	5 Hours
4	Cultivation, collection, processing and storage of crude drugs.	4 Hours
5	Detailed method of cultivation of crude drugs.	13 Hours
6	Study of cell wall constituents and cell inclusions.	2 Hours
7	Microscopy and powder Microscopical study of crude drugs.	11 Hours
8	Study of natural pesticides.	3 Hours
9	Detailed study of various cell constituents.	2 Hours

10	Carbohydrates and related products.	2 Hours
11	Detailed study carbohydrates containing drugs.(11 drugs)	14 Hours
12	Definition sources, method extraction, chemistry and method of analysis of lipids.	2 Hours
13	Detailed study of oils.	3 Hours
14	Definition, classification, chemistry and method of analysis of protein.	2 Hours
15	Study of plants fibers used in surgical dressings and related products.	4 Hours
16	Different methods of adulteration of crude drugs.	3 Hours

2.3 PHARMACOGNOSY & PHYTOPHARMACEUTICALS (PRACTICAL)

Practical: 3 Hrs./Week

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.
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.
Macro, powder and microscopic study of Ginger.
16. Macro, powder and microscopic study of Podophyllum.
17. Determination of Iodine value.
18. Determination of Saponification value and unsaponifiable matter.
19. Determination of ester value.
20. Determination of Acid value.
21. Chemical tests for Acacia.
22. Chemical tests for Tragacanth.
23. Chemical tests for Agar.
24. Chemical tests for Starch.
25. Chemical tests for Lipids.(castor oil,sesame oil, shark liver oil,bees wax)
26. Chemical tests for Gelatin.

Scheme of Practical Examination:

	Sessionals	Annual
Identification	04	10
Synopsis	04	10
Major Experiment	07	20
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	04hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

2.4 PHARMACOLOGY – I (THEORY)

Theory: 3 Hrs. /Week

1. **Scope of the Subject:** 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, apart from general pharmacology, drugs acting on autonomic nervous system, cardiovascular system, central nervous system, blood and blood forming agents and renal system will be taught. In addition to theoretical knowledge, the basic practical knowledge relevant to therapeutics will be imparted.
2. **Objectives of the Subject :** Upon completion of the subject student shall be able to (Know, do, appreciate) -
 - a. understand the pharmacological aspects of drugs falling under the above mentioned chapters;
 - b. handle and carry out the animal experiments;
 - c. appreciate the importance of pharmacology subject as a basis of therapeutics; and
 - d. correlate and apply the knowledge therapeutically.

Text books (Theory) (Author, Title, Edition, Publication Place, Publisher, Year of Publication)

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

Reference books (Theory) (Author, Title, Edition, Publication Place, Publisher, Publication Year)

- a. 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 McGraw Hill, Pergamon press.
- b. Craig, C.R.&Stitzel, R.E. Modern Pharmacology. Latest edition. Publisher: Little Brown.Co
- c. Katzung, B.G. Basic and clinical pharmacology. Latest edition. Publisher: Prentice Hall, Int.
- d. Shargel and Leon. Applied Biopharmaceutics and pharmacokinetics. Latest edition. Publisher: Prentice Hall, London.

Text books (Practical):

Kulkarni, S. K. and Dandia, P. C. Hand book of experimental pharmacology. Latest edition, Publisher: Vallabh Prakashan, Delhi.

Reference books (Practical)

- a. Macleod, L.J. Pharmacological experiments on intact preparations. Latest edition, Publisher: Churchill livingstone.
- b. Macleod, L.J. Pharmacological experiments on isolated preparations. Latest edition, Publisher: Churchill livingstone.
- c. Ghosh, M.N. Fundamentals of experimental pharmacology. Latest edition, Publisher: Scientific book agency, Kolkata.
- d. Ian Kitchen. Textbook of in vitro practical pharmacology. Latest edition, Publisher: Black well Scientific.

3. Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	General Pharmacology a) Introduction, definitions and scope of pharmacology b) Routes of administration of drugs c) Pharmacokinetics (absorption, distribution, metabolism and excretion) d) Pharmacodynamics e) Factors modifying drug effects f) Drug toxicity - Acute, sub- acute and chronic toxicity as per OECD guidelines g) Pre-clinical evaluations h) 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.	18 Hours
2	Pharmacology of drugs acting on ANS a) Adrenergic and antiadrenergic drugs b) Cholinergic and anticholinergic drugs c) Neuromuscular blockers d) Mydriatics and miotics e) Drugs used in myasthenia gravis f) Drugs used in Parkinsonism	14 Hours
3	Pharmacology of drugs acting on cardiovascular system a) Antihypertensives b) Anti-anginal drugs c) Anti-arrhythmic drugs d) Drugs used for therapy of Congestive Heart Failure e) Drugs used for hyperlipidaemias	10 Hours

4	Pharmacology of drugs acting on Central Nervous System a) General anesthetics b) Sedatives and hypnotics c) Anticonvulsants d) Analgesic and anti-inflammatory agents e) Psychotropic drugs f) Alcohol and methyl alcohol g) CNS stimulants and cognition enhancers h) Pharmacology of local anaesthetics	20 Hours
5	Pharmacology of Drugs acting on Respiratory tract a) Bronchodilators b) Mucolytics c) Expectorants d) Antitussives e) Nasal Decongestants	3 Hours
6	Pharmacology of Hormones and Hormone antagonists a) Thyroid and Antithyroid drugs b) Insulin, Insulin analogues and oral hypoglycemic agents c) Sex hormones and oral contraceptives d) Oxytocin and other stimulants and relaxants	7 Hours
7	Pharmacology of autocoids and their antagonists a) Histamines and Antihistaminics b) 5-Hydroxytryptamine and its antagonists c) Lipid derived autocoids and platelet activating factor	

2.5 COMMUNITY PHARMACY (THEORY)

Theory: 2 Hrs. /Week

1. **Scope:** 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, responding to minor ailments by providing suitable safe medication, patient counselling, health screening services for improved patient care in the community set up.
2. **Objectives: Upon completion of the course, the student shall be able to –**
 - a. know pharmaceutical care services;
 - b. know the business and professional practice management skills in community pharmacies;
 - c. do patient counselling & provide health screening services to public in community pharmacy;
 - d. respond to minor ailments and provide appropriate medication;
 - e. show empathy and sympathy to patients; and
 - f. appreciate the concept of Rational drug therapy.

Text Books:

- a. Health Education and Community Pharmacy by N.S.Parmar.
- b. WHO consultative group report.
- c. Drug store & Business management by Mohammed Ali & Jyoti.

Reference books:

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

3.Special requirements:

1. 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.
3. Scheme of evaluation (80 Marks)
 1. Synopsis 10
 2. Major Experiment 30
(Counselling of patients with specific diseases – emphasis should be given on Counselling introduction, content, process and conclusion)
 3. Minor Experiment (Ability to measure B.P/ CBG / Lung function) 15
 4. Prescription Analysis (Analyzing the prescriptions for probable drug interaction and ability to tell the management) 15
 5. Viva – Voce 10

4. Lecture wise programme :

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Definition, scope, of community pharmacy Roles and responsibilities of Community pharmacist	2 Hours
2	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	4 Hours
3	Prescriptions – parts of prescription, legality & identification of medication related problems like drug interactions.	2 Hours
4	Inventory control in community pharmacy Definition, various methods of Inventory Control ABC, VED, EOQ, Lead time, safety stock	3 Hours
5	Pharmaceutical care Definition and Principles of Pharmaceutical care.	2 Hours
6	Patient counselling Definition, outcomes, various stages, barriers, Strategies to overcome barriers Patient information leaflets- content, design, & layouts, advisory labels	4 Hours
7	Patient medication adherence Definition, Factors affecting medication adherence, role of pharmacist in improving the adherence. Medication adherence scales	2 Hours + 1 hr
8	Health screening services Definition, importance, methods for screening - Blood pressure/ blood sugar/ lung function and Cholesterol testing	3 Hours
9	OTC Medication- Definition, OTC medication list & Counselling	2 Hours
10	Health Education WHO Definition of health, and health promotion, care for children, pregnant & breast feeding women, and geriatric patients. Commonly occurring Communicable Diseases, causative agents, Clinical presentations and prevention of communicable diseases – Tuberculosis, Hepatitis, Typhoid, Amoebiasis, Malaria, Leprosy, Syphilis, Gonorrhea and AIDS Balance diet, and treatment & prevention of deficiency disorders Family planning – role of pharmacist	14 Hours
11	Responding to symptoms of minor ailments	9 Hours

	Relevant pathophysiology, common drug therapy to – Pain, GI disturbances (Nausea, Vomiting, Dyspepsia, diarrhea, constipation), Pyrexia, Ophthalmic symptoms, worms infestations.	
12	Essential Drugs concept and Rational Drug Therapy Role of community pharmacist	2 Hours
13	Code of ethics for community pharmacists	1 Hours

2.6 PHARMACOTHERAPEUTICS I (THEORY)

Theory : 3 Hrs. /Week

1. **Scope of the Subject:** 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.
2. **Objectives:** At completion of this subject it is expected that students will be able to understand –
 - a. the pathophysiology of selected disease states and the rationale for drug therapy;
 - b. the therapeutic approach to management of these diseases;
 - c. the controversies in drug therapy;
 - d. the importance of preparation of individualised therapeutic plans based on diagnosis
 - e. 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);
 - f. describe the pathophysiology of selected disease states and explain the rationale for drug therapy;
 - g. summarise the therapeutic approach to management of these diseases including reference to the latest available evidence;
 - h. discuss the controversies in drug therapy;
 - i. discuss the preparation of individualised therapeutic plans based on diagnosis; and
 - j. 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).

Text Books:

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

Reference Books

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

3. Detailed syllabus and lecture wise schedule:

Etiopathogenesis and pharmacotherapy of diseases associated with systems/ diseases following

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Cardiovascular system: Hypertension, Congestive cardiac failure, Angina Pectoris, Myocardial infarction, Hyperlipidaemias, Electrophysiology of heart and Arrhythmias	13 Hours
2	Respiratory system: Introduction to Pulmonary function test, Asthma, Chronic obstructive airways disease, Drug induced pulmonary diseases	6 Hours
3	Endocrine system: Diabetes, Thyroid diseases, Oral contraceptives, Hormone replacement therapy, Osteoporosis	8 Hours
4	General prescribing guidelines for a. Paediatric patients b. Geriatric patients c. Pregnancy and breast feeding	4 Hours
5	Ophthalmology: Glaucoma, Conjunctivitis- viral & bacterial	3 Hours
6	Introduction to rational drug use Definition, Role of pharmacist Essential drug concept Rational drug formulations	2 Hours

2.6 PHARMACOTHERAPEUTICS - I (PRACTICAL)

Practical: 3 Hrs./Week

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.

Scheme of Practical Examination:

	Sessionals	Annual
Synopsis	05	15
Major Experiment	10	25
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	04hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

Third Year

3.1 PHARMACOLOGY – II (THEORY)

Theory: 3 Hrs. /Week

- 1. Scope of the Subject:** 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 autocoids 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.
- 2. Objectives of the Subject:** Upon completion of the subject student shall be able to:
 - a. understand the pharmacological aspects of drugs falling under the above mentioned chapters,
 - b. carry out the animal experiments confidently,
 - c. appreciate the importance of pharmacology subject as a basis of therapeutics, and
 - d. correlate and apply the knowledge therapeutically.
 - e. understand different aspects of genes and their regulatory functions.

Text books (Theory)

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

Reference books (Theory)

- a. 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: McGrawHill, Pergamon press.
- b. Craig, C.R. and Stitzel, R.E. Modern Pharmacology. Latest edition. Publisher: Little Brown and company.
- c. Katzung, B.G. Basic and clinical pharmacology. Latest edition. Publisher: Prentice Hall, International.

- d. 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.

Text books (Practical)

Kulkarni, S. K. and Dandia, P. C. Hand book of experimental pharmacology. Latest edition, Publisher: Vallabh Prakasan, Delhi.

Reference books (Practical):

- a. Macleod, L.J. Pharmacological experiments on intact preparations. Latest edition, Publisher: Churchill livingstone.
- b. Macleod, L.J. Pharmacological experiments on isolated preparations. Latest edition, Publisher: Churchill livingstone.
- c. Ghosh, M.N. Fundamentals of experimental pharmacology. Latest edition, Publisher: Scientific book agency, Kolkata.
- d. Ian Kitchen. Textbook of in vitro practical pharmacology. Latest edition, Publisher: Black well Scientific.

3. Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Pharmacology of Drugs acting on Blood and blood forming agents a) Anticoagulants b) Thrombolytics and antiplatelet agents c) Haemopoietics and plasma expanders	6 Hours
2	Pharmacology of drugs acting on Renal System a) Diuretics b) Antidiuretics	3 Hours
3	Chemotherapy a) Introduction b) Sulfonamides and co-trimoxazole c) Penicillins and Cephalosporins d) Tetracyclins and Chloramphenicol e) Macrolides, Aminoglycosides, Polyene & Polypeptide antibiotics f) Quinolines and Fluroquinolines g) Antifungal antibiotics h) Antiviral agents i) Chemotherapy of tuberculosis and leprosy j) Chemotherapy of Malaria k) Chemotherapy of protozoal infections (amoebiasis, Giardiasis)	31 Hours

	l) Pharmacology of Anthelmintic drugs m) Chemotherapy of cancer (Neoplasms)	
4	Immunopharmacology Pharmacology of immunosuppressants and stimulants	2 Hours
5	Principles of Animal toxicology Acute, sub-acute and chronic toxicity	2 Hours
6	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.)	14 Hours
7	The Gene: Genome structure and function: a) Gene structure: Organization and elucidation of genetic code. b) Gene expression: Expression systems (pro and eukaryotic), genetic elements that control gene expression (nucleosomes, histones, acetylation, HDACS, DNA binding protein families. c) Transcription and Transcription factors: Basic principles of transcription in pro and eukaryotes. Transcription factors that regulate transcription in pro and eukaryotes. 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	17 Hours

Books:

1. Molecular Biology of the Cell by Alberts B., Bray, D., Lewis, J., Raff M., Roberts K and Watson, JD, 3rd edition.
2. Molecular Cell Biology By Lodish, H., Baltimore, D., Berk, A et al., 5th edition.
3. Molecular Biology by Turner, PC., McLennan, AG., Bates, AD and White MRH. 2nd edition.
4. Genes VIII by Lewin, B., (2004)
5. Pharmaceutical Biotechnology, by Crommelin, DJA and Sindelar RD (1997)
6. Recombinant DNA by Watson, JD., Gilman, M., et al., (1996)
7. Biopharmaceutical: Biochemistry and Biotechnology by Walsh, G., (1998)

3.1 PHARMACOLOGY – II (PRACTICAL)

Practical : 3 Hrs./Week

List of Experiments:

1. Study of laboratory appliances used in experimental pharmacology.
2. Study of laboratory animals and their handling (a. Frogs, b. Mice, c. Rats, d. Guinea pigs, e. Rabbits)
3. To study the routes of administration of drugs in animals (Rats, Mice, Rabbits).
4. Study of use of anesthetics in laboratory animals.
5. Study of physiological salt solutions used in experimental pharmacology.
6. 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 rotorod.
 - f) Cardiotonic activity of drugs using isolated frog heart and mammalian heart preparations.
7. To record the dose response curve of Ach using isolated ileum/rectus abdominis muscle preparation.
8. To carry out bioassay of Ach using isolated ileum/rectus abdominis muscle by interpolation method.
9. To carry out bioassay of Ach using isolated ileum/rectus abdominis muscle preparation by three point method.
10. To record the dose response curve of Histamine using isolated guinea -pig ileum preparation.
11. To carry out bioassay of Histamine using isolated guinea -pig ileum preparation by interpolation method.
12. To carry out bioassay of Histamine using guinea -pig ileum preparation by three point method.
13. Study of agonistic and antagonistic effects of drugs using isolated guinea -pig ileum preparation.

Scheme of Practical Examination:

	Sessionals	Annual
Identification	02	10
Synopsis	04	10
Major Experiment (Bioassay)	08	30
Minor Experiment (Interpretation of given Graph or simulated experiment)	04	10
Viva	02	10
Max Marks	20	70
Duration	03hrs	04hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

3.2 PHARMACEUTICAL ANALYSIS (THEORY)

Theory: 3 Hrs. /Week

1. **Scope of the subject:** This course is designed to impart a fundamental knowledge on the art and science of testing drugs by various instrumental methods of analysis. This focuses on various modern instruments that are used for testing the purity of drugs in various dosage forms. This course also gives idea about modern instruments that are used for drug testing like NMR, IR, Mass, HPLC, HPTLC forms. etc,. It prepares the students for most basics of the applied field of pharmacy.
2. **Objectives of the subject:**
 - a. To understand the construction and working of various analytical instruments.
 - b. To know principle and mechanism of instrumentation.
 - c. To understand the different modern techniques of drug analysis.
 - d. To appreciate the advantages of instrumental methods of drug analysis.

Text Books:

Reference Books:

1. Text Book of Pharm. Analysis by Higuchi. T and Hasen. E. B., New York Inter Science Publishers.
2. Quantitative Pharma. Analysis by Jenkins, The Blakiston division, New York.
3. Quantitative Drug Analysis, by Garrot. D, Chapman & Hall Ltd., London.
4. Undergraduate Instrumental Analysis by James. E., CBS Publishers.
5. Instrumental Analysis by Willard and Merritt, EWP, East West Press Ltd., Delhi/Madras.
6. Pharm Analysis by Skoog and West, Sounders Manipl College Publishing.
7. Text Book of Chemical Analysis, by A.I.Vogel, ELBS with Macmillan press, Hampshire.
8. Textbook of Pharm. Analysis by K.A.Connors, John Wiley & Sons, New York, Brisbane, Singapore.
9. Textbook of Pharm. Analysis (Practical) by Beckett & Stenlake, CBS Publishers, Delhi.
10. Textbook of Drug Analysis by P.D. Sethi., CBS Publishers, Delhi.
11. Spectroscopy by Silverstein, John & Wiley & Sons. Inc., Canada & Singapore.
12. How to practise GMP-A Plan for total quality control by P.P. Sharma, Vandana Publications, Agra.
13. The Science & Practice of Pharmacy by Remington Vol -I & II, Mack Publishing Co. Pennsylvania.
14. TLC by Stahl, Spring Verlay.
15. Text Book of Pharm. Chemistry by Chatten, CBS Publications.
16. Spectroscopy by William Kemp, ELBS with Macmillan Press, Hampshire.
17. I.P.-1996, The Controller of Publications, New Delhi.
18. BPC- Dept. of Health, U.K. for HMSO.
19. USP - Mack Publishing Co., Easton, PA
20. The Extra Pharmacopoeia – The Pharm. Press, London.

4 Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Quality Assurance: - Introduction, sources of quality variation, control of quality variation. - Concept of statistical quality control. - Validation methods- quality of equipment, validation of equipment and validation of analytical instruments and calibration. - GLP, ISO 9000. - Total quality management, quality review and documentation. - ICH- international conference for harmonization-guidelines. - Regulatory control.	4 Hours
2	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. - Column Chromatography: Adsorption column chromatography, Operational technique, frontal analysis and elution analysis. Factors affecting column efficiency, applications and partition chromatography. - TLC: Introduction, principle, techniques, Rf value and applications. - PC: Introduction, principle, types of paper chromatography, preparation techniques, development techniques, applications. - Ion-exchange chromatography: Introduction, principles, types of ion exchange synthetic resins, physical properties, factors affecting ion exchange, methodology and applications. - HPLC: Introduction, theory, instrumentation, and applications. - HPTLC: Introduction, theory, instrumentation, and applications. - 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, programmed temperature gas chromatography, applications. - Electrophoresis: Principles of separation, equipment for paper and gel electrophoresis, and application. - Gel filtration and affinity chromatography: Introduction, technique, applications.	20 Hours

3	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.	14 Hours
4	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. Instrumentation – Photometer, U.V.-Visible spectrophotometer-sources of U.V.-Visible radiations, collimating systems, monochromators, samples cells and following detectors-Photocell, Barrier layer cell, Phototube, Diode array, applications of U.V.-Visible spectroscopy in pharmacy and spectrophotometric titrations. Infrared Spectroscopy: Vibrational transitions, frequency – structure correlations, Infrared absorption bands, Instrumentation–IR spectrometer – 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. Fluorimetric Analysis: Theory, luminescence, factors affecting fluorescence, quenching. Instrumentation, Applications, fluorescent indicators, study of pharmaceutically important compounds estimated by fluorimetry.	35 Hours

	b. Flame Photometry: Theory, nebulisation, flame and flame temperature, interferences, flame spectrometric techniques and instrumentation and pharmaceutical applications. c. Atomic Absorption Spectrometry: Introduction, Theory, types of electrodes, instrumentation and applications. d. Atomic Emission Spectroscopy: Spectroscopic sources, atomic emission spectrometers, photographic and photoelectric detection. e. NMR & ESR (introduction only): Introduction, theoretical aspects and applications. f. Mass Spectroscopy: (Introduction only) – Fragmentation, types of ions produced mass spectrum and applications. g. Polarimetry: (Introduction only) – Introduction to optical rotatory dispersion, circular dichroism, polarimeter. h. X-RAY Diffraction: (Introduction only) – Theory, reciprocal lattice concept, diffraction patterns and applications. i. Thermal Analysis: Introduction, instrumentation, applications, and DSC and DTA.	
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3.2 PHARMACEUTICAL ANALYSIS (PRACTICAL)

Practical : 3 Hrs./Week

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 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.
13. 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.

Scheme of Practical Examination:

	Sessionals	Annual
Synopsis	05	15
Major Experiment	10	25
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	04hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

3.3 PHARMACOTHERAPEUTICS – II (THEORY)

Theory : 3 Hrs. /Week

1. **Scope of the Subject:** 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.
2. **Objectives of the Subject: Upon completion of the subject student shall be able to –**
 - a. know the pathophysiology of selected disease states and the rationale for drug therapy
 - b. know the therapeutic approach to management of these diseases;
 - c. know the controversies in drug therapy;
 - d. know the importance of preparation of individualised therapeutic plans based on diagnosis; and
 - e. 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).

Text books (Theory)

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

Reference books (Theory)

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

3. Detailed syllabus and lecture wise schedule:

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

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Infectious disease: Guidelines for the rational use of antibiotics and surgical Prophylaxis, Tuberculosis, Meningitis, Respiratory tract infections, Gastroenteritis, Endocarditis, Septicemia, Urinary tract infections, Protozoal infection - Malaria, HIV & Opportunistic infections, Fungal infections, Viral infections, Gonorrhoea and Syphilis	38 Hours
2	Musculoskeletal disorders: Rheumatoid arthritis, Osteoarthritis, Gout, Spondylitis, Systemic lupus erythematosus.	10 Hours
3	Renal system: Acute Renal Failure, Chronic Renal Failure, Renal Dialysis, Drug induced renal disorders	11 Hours
4	Oncology: Basic principles of Cancer therapy, General introduction to cancer chemotherapeutic agents, Chemotherapy of breast cancer, leukemia, lung cancer, colorectal cancer, prostate cancer. Management of chemotherapy nausea and emesis	12 Hours (+3hrs)
5	Dermatology: Psoriasis, Scabies, Eczema, Impetigo	4 Hours

3.3 PHARMACOTHERAPEUTICS – II (PRACTICAL)

Practical: 3Hrs./Week

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.

Scheme of Practical Examination:

	Sessionals	Annual
Synopsis	05	15
Major Experiment	10	25
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	04hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

3.4 PHARMACEUTICAL JURISPRUDENCE (THEORY)

Theory: 2 Hrs. /Week

1. **Scope of the Subject:** 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 are 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.
2. **Objectives of the Subject:** Upon completion of the subject student shall be able to (Know, do, and appreciate) –
 - a. practice the Professional ethics;
 - b. understand the various concepts of the pharmaceutical legislation in India;
 - c. know the various parameters in the Drug and Cosmetic Act and rules;
 - d. know the Drug policy, DPCO, Patent and design act;
 - e. understand the labeling requirements and packaging guidelines for drugs and cosmetics;
 - f. be able to understand the concepts of Dangerous Drugs Act, Pharmacy Act and Excise duties Act; and
 - g. other laws as prescribed by the Pharmacy Council of India from time to time including International Laws.

Text books (Theory)

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

Reference books (Theory)

- a. Singh, KK, editor. Beotra's the Laws of Drugs, Medicines & cosmetics. Allahabad: Law Book House; 1984.
- b. Jain, NK. A Textbook of forensic pharmacy. Delhi: Vallabh prakashan ; 1995.
- c. Reports of the Pharmaceutical enquiry Committee
- d. I.D.M.A., Mumbai. DPCO 1995
- e. Various reports of Amendments.
- f. Deshapande, S.W. The drugs and magic remedies act 1954 and rules 1955. Mumbai: Susmit Publications; 1998.
- g. Eastern Book Company .The narcotic and psychotropic substances act 1985, Lucknow: Eastern; 1987.

3. Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Pharmaceutical Legislations – A brief review.	1 Hour
2	Principle and Significance of professional ethics. Critical study of the code of pharmaceutical ethics drafted by PCI.	2 Hours
3	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.	23 Hours
4	Pharmacy Act –1948. Objectives Legal Definitions, General Study, Constitution and Functions of State & Central Council, Registration & Procedure, ER.	3 Hours
5	Medicinal and Toilet Preparation Act –1955. Objectives, Legal Definitions, Licensing, Bonded and Non Bonded Laboratory, Ware Housing, Manufacture of Ayurvedic, Homeopathic, Patent & Proprietary Preparations.	3 Hours
6	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.	4 Hours
7	Study of Salient Features of Drugs and magic remedies Act and its rules.	2 Hours
8	Study of essential Commodities Act Relevant to drugs price control Order.	2 Hours
9	Drug Price control Order & National Drug Policy (Current).	2 Hours
10	Prevention Of Cruelty to animals Act-1960.	3 Hours
11	Patents & design Act-1970.	3 Hours
12	Brief study of prescription and Non-prescription Products.	2 Hours
13	Minimum wages Act	

4. Assignments:

Format of the assignment

1. Minimum & Maximum number of pages
2. It shall be a 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

Case studies relating to

1. 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.
2. Various prescription and non-prescription products.
3. Medical and surgical accessories.
4. Diagnostic aids and appliances available in the market.

3.5 MEDICINAL CHEMISTRY (THEORY)

Theory: 3 Hrs. /Week

1. **Scope of the subject:** This course is designed to impart a fundamental knowledge on the structure and functions of the different drugs. The course gives details of Chemistry, Mechanism of action, metabolism, adverse effects, Structure Activity Relationships (SAR) and uses of Chemotherapeutic Agents, Cardiovascular Drugs and synthesis of some important drugs. The course also covers modern techniques of drug design, which include Prodrug concept and combinatorial chemistry.
2. **Objectives of the subject:**
 - a. To understand the chemistry of drugs with respect to their biological activity.
 - b. To know the metabolism, adverse effect and therapeutic activity of drugs.
 - c. To understand the different modern techniques of drug design.
 - d. To appreciate the SAR of some important drug classes.

Text Books:

Reference Books:

1. Wilson and Gisvold's Text book of Organic, Medicinal and Pharmaceutical Chemistry, Lippincott-Raven Publishers-New York, Philadelphia.
2. William.O.Foye, Principles of Medicinal Chemistry, B.I. Waverly Pvt. Ltd., New Delhi.
3. Burgers, Medicinal Chemistry, M.E., Welly Med.Chemistry M.E. Walffed Johnwiley and Sons, Wiley-interscience Publication, New York, Toronto.
4. A Text Book of Medicinal Chemistry Vol. I and II by Surendra N. Pandeya, S.G. Publisher, 6, Dildayal Nagar, Varanasi -10.
5. Indian Pharmacopoeia 1985 and 1996. The Controller of Publications, Civil Lines, Delhi - 54.
6. Current Index of Medical Specialities (CIMS) and MIMS India, MIMS, A.E. Morgan Publications (I) Pvt. Ltd, New Delhi-19.
7. Organic Drug Synthesis-Ledniser Mitzsher Vol. I and II.
8. Pharmaceutical Chemistry drug Synthesis Vol. I and II by H. J. Roth and A. Kleemann.
9. The Science and Practice of Pharmacy Vol. 1 and 2, Remington, MACK Publishing Company, Easton, Pennsylvania.

3. Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Modern concept of rational drug design: A brief introduction to Quantitative Structure Activity Relationship (QSAR), prodrug, combinatorial chemistry and computer aided drug design (CADD) and concept of antisense molecules. 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.	6 Hours
2	Anti-infective agents a) Local anti-infective agents b) Preservatives c) Antifungal agents d) Urinary tract anti-infectives e) Antitubercular agents f) Antiviral agents and Anti AIDS agents g) Antiprotozoal agents h) Anthelmintics i) Antiscabies and Antipedicular agents	18 Hours
3	Sulphonamides and sulphones	4 Hours
4	Antimalarials	4 Hours
5	Antibiotics	12 Hours
6	Antineoplastic agents	5 Hours
7	Cardiovascular agents a) Antihypertensive agents b) Antianginal agents and vasodilators c) Antiarrhythmic agents d) Antihyperlipidemic agents e) Coagulants and Anticoagulants f) Endocrine	16 Hours
8	Hypoglycemic agents	2 Hours
9	Thyroid and Antithyroid agents	2 Hours
10	Diuretics	3 Hours
11	Diagnostic agents	1 Hour
12	Steroidal Hormones and Adrenocorticoids	2 Hours

3.5 MEDICINAL CHEMISTRY (PRACTICAL)

Practical: 3 Hrs./Week

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.

Scheme of Practical Examination:

	Sessionals	Annual
Synopsis	05	15
Major Experiment	10	25
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	04hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

3.6 PHARMACEUTICAL FORMULATIONS (THEORY)

Theory: 2 Hrs. /Week

1. **Scope of the Subject:** Subject deals with the formulation and evaluation of various pharmaceutical dosage forms.
2. **Objectives of the Subject:** Upon completion of the subject student shall be able to (Know, do, appreciate) –
 - a. understand the principle involved in formulation of various pharmaceutical dosage forms;
 - b. prepare various pharmaceutical formulation;
 - c. perform evaluation of pharmaceutical dosage forms; and
 - d. understand and appreciate the concept of bioavailability and bioequivalence, their role in clinical situations.

Text books (Theory)

- a. Pharmaceutical dosage forms, Vol, I,II and III by lachman
- b. Rowlings Text book of Pharmaceutics
- c. Tutorial Pharmacy – Cooper & Gun

Reference books (Theory)

- a. Remington's Pharmaceutical Sciences
- b. USP/BP/IP

3. Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Pharmaceutical dosage form- concept and classification	2 Hours
2	Tablets: Formulation of different types of tablets, tablet excipients, granulation techniques quality control and evaluation of tablets. Tablet coating, Type of coating, quality control tests for coated tablet.	10 Hours
3	Capsules; Production and filling of hard gelatin capsules, Raw material for shell, finishing, quality control tests for capsules. Production and filling of soft gelatin capsules, quality control tests for soft gelatin capsules.	8 Hours
4	Liquid orals: Formulation and evaluation of suspensions, emulsions and solutions. Stability of these preparations	4 Hours
5	Parenterals: Introduction Containers used for Parenterals (including official tests) Formulation of large and small volume Parenterals Sterilization	10 Hours

6	Semi- Solids: Introduction and classification Factors affecting absorption and anatomy of skin Packaging storage and labeling, Ointments. Types of Ointment Base Preparation of ointment, Jellies Types of jellies Formulation of jellies Suppositories, Method of preparation, Types Packaging	8 Hours
7	Definition and concept of Controlled and novel Drug delivery systems with available examples, viz. parenteral, trans dermal, buccal, rectal, nasal, implants, ocular	8 Hours

3.6 PHARMACEUTICAL FORMULATIONS (PRACTICAL)

Practical: 3 Hrs./Week

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. Salicylic 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)**

Scheme of Practical Examination:

	Sessionals	Annual
Synopsis	05	15
Major Experiment	20	25
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	04hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance)

Fourth Year

4.1 PHARMACOTHERAPEUTICS – III (THEORY)

Theory: 3 Hrs. /Week

1. **Scope:** 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.
2. **Objectives:** At completion of this subject it is expected that students will be able to understand –
 - a. the pathophysiology of selected disease states and the rationale for drug therapy;
 - b. the therapeutic approach to management of these diseases;
 - c. the controversies in drug therapy;
 - d. the importance of preparation of individualised therapeutic plans based on diagnosis;
 - e. 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);
 - f. describe the pathophysiology of selected disease states and explain the rationale for drug therapy;
 - g. to summarize the therapeutic approach to management of these diseases including reference to the latest available evidence;
 - h. to discuss the controversies in drug therapy;
 - i. to discuss the preparation of individualised therapeutic plans based on diagnosis; and
 - j. 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).

Text Books

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

Reference Books

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

3. Detailed syllabus and lecture wise schedule:

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

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Gastrointestinal system: Peptic ulcer disease, Gastro Esophageal Reflux Disease, Inflammatory bowel disease, Liver disorders - Alcoholic liver disease, Viral hepatitis including jaundice, and Drug induced liver disorders.	20 Hours
2	Haematological system: Anaemias, Venous thromboembolism, Drug induced blood disorders.	12 Hours
3	Nervous system: Epilepsy, Parkinsonism, Stroke, Alzheimer's disease	16 Hours
4	Psychiatry disorders: Schizophrenia, Affective disorders, Anxiety disorders, Sleep disorders, Obsessive Compulsive disorders	14 Hours
5	Pain management including Pain pathways, neuralgias, headaches.	8 Hours
6	Evidence Based Medicine	5 Hours

4.1 PHARMACOTHERAPEUTICS – III (PRACTICAL)

Practical: 3 Hrs./Week

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:

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

Scheme of Practical Examination:

	Sessionals	Annual
Synopsis	05	15
Major Experiment	10	25
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	06hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

4.2 HOSPITAL PHARMACY (THEORY)

Theory: 2 Hrs. /Week

1. **Scope:** 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, drug dispensing, manufacturing of parenteral preparations, drug information, patient counselling, and therapeutic drug monitoring for improved patient care.
2. **Objectives:** Upon completion of the course, the student shall be able to –
 - a. know various drug distribution methods;
 - b. know the professional practice management skills in hospital pharmacies;
 - c. provide unbiased drug information to the doctors;
 - d. know the manufacturing practices of various formulations in hospital set up;
 - e. appreciate the practice based research methods; and
 - f. appreciate the stores management and inventory control.

Text books: (latest editions)

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

References:

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

3. Lecture wise programme:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Hospital - its Organisation and functions	1 Hour
2	Hospital pharmacy-Organisation and management a) Organizational structure-Staff, Infrastructure & work load statistics b) Management of materials and finance c) Roles & responsibilities of hospital pharmacist	5 Hours
3	The Budget – Preparation and implementation	1 Hour

4	Hospital drug policy a) Pharmacy and Therapeutic committee (PTC) b) Hospital formulary c) Hospital committees - Infection committee - Research and ethical committee d) developing therapeutic guidelines	12 Hours
5	Hospital pharmacy services a) Procurement & warehousing of drugs and Pharmaceuticals b) Inventory control - Definition, various methods of Inventory Control - ABC, VED, EOQ, Lead time, safety stock c) Drug distribution in the hospital i) Individual prescription method ii) Floor stock method iii) Unit dose drug distribution method d) Distribution of Narcotic and other controlled substances e) Central sterile supply services – Role of pharmacist	12 Hours
6	Manufacture of Pharmaceutical preparations a) Sterile formulations – large and small volume parenterals b) Manufacture of Ointments, Liquids, and creams c) Manufacturing of Tablets, granules, capsules, and powders d) Total parenteral nutrition	12 Hours
7	Continuing professional development programs Education and training	3 Hours
8	Radio Pharmaceuticals – Handling and packaging	2 Hours
9	Professional Relations and practices of hospital pharmacist	3 Hours
10	Social & Preventive Pharmacy	4 Hours

4.2 HOSPITAL PHARMACY (PRACTICAL)

Practical: 3 Hrs./Week

List of Experiments

1. Assessment of drug interactions in the given prescriptions
2. Manufacture of parenteral formulations, powders.
3. Drug information queries.
4. 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.

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.

Scheme of Practical Examination:

	Sessionals	Annual
Synopsis	05	15
Major Experiment	10	25
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	06hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance.

4.3 CLINICAL PHARMACY (THEORY)

Theory: 3 Hrs. /Week

1. **Scope:** This course is designed to impart the basic knowledge and skills that required for practice of pharmacy including provision of various clinical pharmacy services to patients and healthcare professionals in clinical settings
2. **Objectives of the Subject :** Upon completion of the subject student shall be able to (Know, do, appreciate) –
 - a. monitor drug therapy of patient through medication chart review and clinical review;
 - b. obtain medication history interview and counsel the patients;
 - c. identify and resolve drug related problems;
 - d. detect, assess and monitor adverse drug reaction;
 - e. interpret selected laboratory results (as monitoring parameters in therapeutics) of specific disease states; and
 - f. retrieve, analyse, interpret and formulate drug or medicine information.

Text books (Theory)

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

References

- a. Australian drug information -Procedure manual. The Society of Hospital Pharmacists of Australia.
- b. Clinical Pharmacokinetics - Rowland and Tozer, Williams and Wilkins Publication.
- c. Pharmaceutical statistics. Practical and clinical applications. Sanford Bolton, Marcel Dekker, Inc.

2. Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Definitions, development and scope of clinical pharmacy	3 Hours
2	Healthcare ethics a. What is ethics? Ethics & law b. Beneficence; Non-beneficence; Respect for autonomy c. Ethical decision making Ethical dilemmas in healthcare of Pharmacy practice	2 Hours

3	Introduction to daily activities of a clinical pharmacist - Drug therapy monitoring (medication chart review, clinical review, pharmacist interventions) - Ward round participation - Adverse drug reaction management - Drug information and poisons information - Medication history - Patient counseling - Drug utilisation evaluation (DUE) and review (DUR) - Quality assurance of clinical pharmacy services	13 Hours
4	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.	3 Hours
5	Clinical laboratory tests used in the evaluation of disease states, and interpretation of test results - Haematological, Liver function, Renal function, thyroid function tests - Tests associated with cardiac disorders - Fluid and electrolyte balance - Microbiological culture sensitivity tests - Pulmonary Function Tests	15 Hours
5	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	8 Hours
6	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 d. Role of pharmacist in management of ADR.	10 Hours
7	Communication skills , including patient counselling techniques, medication history interview, presentation of cases.	10 Hours
8	Pharmaceutical care concepts	4 Hours
9	Critical evaluation of biomedical literature	6 Hours
10	Medication errors	3 Hours

4.3 CLINICAL PHARMACY (PRACTICAL)

Practical: 3 Hrs./Week

Students are expected to perform 15 practicals in the following areas covering the topics dealt in theory class.

- a. Answering drug information questions (4 Nos)
- b. Patient medication counselling (4 Nos)
- c. Case studies related to laboratory investigations (4 Nos)
- d. 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.

Scheme of Practical Examination:

	Sessionals	Annual
Synopsis	05	15
Major Experiment	10	25
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	06hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

4.4 BIOSTATISTICS AND RESEARCH METHODOLOGY (THEORY)

Theory: 2 Hrs. /Week

1. Scope of the subject:

This is an introductory course in statistics, research methodology and Computer application in hospital and community Pharmacy. This subject deals with Research methodology, Biostatistics, epidemiology and Computer application and clinical studies. Research methodology deal about types of clinical study, designing, sample size determination and power of study Statistics deals about frequency distribution, graphics, averages, measures of dispersion, Correlation, regression, Parametric and non-parametric tests. Incidence and prevalence, relative risk, attributable risk. Computer Application deals with application of Computer System in Hospital Pharmacy and Community Pharmacy.

2. Objectives of subject

1. Know the various statistical methods to solve different types of problems
2. Operate various statistical software packages
3. appreciate the importance of Computer in hospital and Community Pharmacy
4. appreciate the statistical technique in solving the pharmaceutical problems

Text Books:

Reference Books:

- a. Pharmaceutical statistics- practical and clinical applications, Sanford Bolton, 3rd edition, publisher Marcel Dekker Inc. NewYork.
- b. Drug Information- A Guide for Pharmacists, Patrick M Malone, Karen L Kier, John E Stanovich , 3rd edition, McGraw Hill Publications 2006

3. Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Research Methodology a) Types of clinical study designs: Case studies, observational studies, interventional studies, b) Designing the methodology c) 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 specified width, power of a study d) Report writing and presentation of data	10 Hours

2.1	Biostatistics a) Introduction b) Types of data distribution c) Measures describing the central tendency distributions- average, median, mode d) Measurement of the spread of data -range, variation of mean, standard deviation, variance, coefficient of variation, standard error of mean.	10 Hours
2.2	Data graphics Construction and labeling of graphs, histogram, pie-charts, scatter plots, semi-logarithmic plots	2 Hours
2.3	Basics of testing hypothesis a) Null hypothesis, level of significance, power of test, P value, statistical estimation of confidence intervals. b) Level of significance (Parametric data)- students t test (paired and unpaired), chi Square test, Analysis of Variance (one-way and two-way) a) Level of significance (Non-parametric data)- Sign test, Wilcoxon's signed rank test, Wilcoxon rank sum test, Mann Whitney U test, Kruskal -Wall is test (one way ANOVA) c) Linear regression and correlation- Introduction, Pearson's and Spearman's correlation and correlation co-efficient. d) Introduction to statistical software: SPSS, Epi Info, SAS.	15 Hours
2.4	Statistical methods in epidemiology Incidence and prevalence, relative risk, attributable risk	5 Hours
3	Computer applications in pharmacy a) <u>Computer System in Hospital Pharmacy</u>: 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. b) <u>Computer In Community Pharmacy</u>: Computerizing the Prescription Dispensing process; Use of Computers for Pharmaceutical Care in community pharmacy; Accounting and General ledger system c) <u>Drug Information Retrieval & Storage</u>: Introduction – Advantages of Computerized Literature Retrieval; Use of Computerized Retrieval	8 Hours

4.5 BIOPHARMACEUTICS AND PHARMACOKINETICS (THEORY)

Theory: 3 Hrs. /Week

1. Scope of the subject:

This course is designed to impart knowledge and skills necessary for dose calculations, dose adjustments and to apply biopharmaceutics theories in practical problem solving. Basic theoretical discussions of the principles of biopharmaceutics and pharmacokinetics are provided to help the students' to clarify the concepts

2.Objectives of the subject:

1. Define the basic concepts in biopharmaceutics and pharmacokinetics.
2. Use raw data and derive the pharmacokinetic models and parameters the best describe the process of drug absorption, distribution, metabolism and elimination.
3. Critically evaluate biopharmaceutic studies involving drug product equivalency
4. Design and evaluate dosage regimens of the drugs using pharmacokinetic and biopharmaceutic parameters.
5. Detect potential clinical pharmacokinetic problems and apply basic pharmacokinetic principles to solve them

Text Books:

Reference Books:

- a. Biopharmaceutics and Clinical Pharmacokinetics by, Milo Gibaldi
- b. Remington's Pharmaceutical Sciences, By Mack Publishing Company, Pennsylvania.
- c. Hand Book of Clinical Pharmacokinetics, By Milo Gibaldi and Laurie Prescott by ADIS Health Science Press.
- d. Biopharmaceutics and Pharmacokinetics; By Robert F Notari
- e. Biopharmaceutics; By Swarbrick
- f. Bio pharmaceutics and Pharmacokinetics-A Treatise, By D. M. Brahmkar and Sunil B.Jaiswal, Vallabh Prakashan Pitampura, Delhi
- g. Clinical Pharmacokinetics, Concepts and Applications: By Malcolm Rowland and Thomas, N. Tozen, Lea and Febrger, Philadelphia, 1995.
- h. Dissolution, Bioavailability and Bioequivalence, By Abdou H.M, Mack Publishing Company, Pennsylvania 1989.
- i. Biopharmaceutics and Clinical Pharmacokinetics-An introduction 4th edition Revised and expanded by Robert F Notari Marcel Dekker Inc, New York and Basel, 1987.
- j. Encyclopedia of Pharmaceutical Technology, Vol 13, James Swarbrick, James, C. Roylan, Marcel Dekker Inc, New York 1996.

3.Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Biopharmaceutics - Introduction to Biopharmaceutics - Absorption of drugs from gastrointestinal tract. - Drug Distribution. - Biotransformation - Drug Elimination. - Method to determine the pharmacokinetic parameters	16 +14Hours
Pharmacokinetics		
2	Introduction to Pharmacokinetics. - Mathematical model - Drug levels in blood. - Pharmacokinetic model - Compartment models e. Pharmacokinetic study.	10 Hours
3	One compartment open model. - Intravenous Injection (Bolus) - Intravenous infusion. - Extravascular administration - Urinary excretion data - Calculations	8 Hours
4	Multicompartment models. - Two compartment open model. - IV bolus, IV infusion and oral administration - Calculations	8- 2=6hrs
5	Multiple – Dosage Regimens. - Repetitive Intravenous injections – One Compartment Open Model - Repetitive Extravascular dosing – One Compartment Open model - Multiple Dose Regimen – Two Compartment Open Model - Calculations	10-4=6 hrs
6	Nonlinear Pharmacokinetics. - Introduction - Factors causing Non-linearity. - Michaelis-menton method of estimating parameters.	7-3=4 hrs

7	Noncompartmental Pharmacokinetics. - Statistical Moment Theory. - MRT for various compartment models. - Physiological Pharmacokinetic model.	8-2=6 hrs
8	Bioavailability and Bioequivalence. - Introduction. - Bioavailability study protocol. - Methods of Assessment of Bioavailability - Methods to improve bioavailability - Calculations	8+1 hrs

4.5 BIOPHARMACEUTICS AND PHARMACOKINETICS (PRACTICAL)

Practical: 3 Hrs./Week

1. Calibration curves (4 hrs)
2. Improvement of dissolution characteristics of slightly soluble drugs by some methods.
 - a. Surfactant addition
 - b. Co-solvent addition
 - c. pH change
3. Comparison of dissolution studies of two different marketed products of same drug.
4. Influence of polymorphism on solubility and dissolution.
5. Protein binding studies of a highly protein bound drug and poorly protein bound drug.
6. 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.
7. Bioavailability studies of some commonly used drugs on animal/human model.
8. Calculation of K_a , K_e , $t_{1/2}$, C_{max} , AUC, AUMC, MRT etc. from blood profile data.
9. Calculation of bioavailability from urinary excretion data for two drugs.
10. Calculation of AUC and bioequivalence from the given data for two drugs.
11. In vitro absorption studies.
12. Bioequivalency studies on the different drugs marketed. (eg) Tetracycline, Sulphamethoxazole, Trimethoprim, Aspirin etc., on animals and human volunteers.
13. Absorption studies in animal inverted intestine using various drugs.
14. Effect on contact time on the plasma protein binding of drugs.
15. Studying metabolic pathways for different drugs based on elimination kinetics data.
16. Calculation of elimination half-life for different drugs by using urinary elimination data and blood level data.
17. Determination of renal clearance.

4.6 CLINICAL TOXICOLOGY (THEORY)

Theory: 2 Hrs. /Week

1. Scope of the subject:

This course is designed to impart a thorough knowledge in the management of various poisoning cases thereby enabling the students to assist healthcare professionals / toxicologists in handling and managing the emergency case

2. Objectives of the subject:

1. Understand and deal with general principles involved in the management of poisoning
2. Recognize the clinical symptoms and manage poisoning cases
3. Educate public and healthcare professionals in the management of emergency cases
4. Minimize/ prevent the poisoning cases in local population

Text Books:

Reference Books:

- a. Matthew J Ellenhorn. ELLENHORNS MEDICAL TOXICOLOGY – DIAGNOSIS AND TREATMENT OF POISONING. Second edition. Williams and Willkins publication, London
- b. V V Pillay. HANDBOOK OF FORENSIC MEDICINE AND TOXICOLOGY. Thirteenth edition 2003 Paras Publication, Hyderabad

3. Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	General principles involved in the management of poisoning	3 Hours
2	Antidotes and the clinical applications.	2 Hours
3	Supportive care in clinical Toxicology.	2 Hours
4	Gut Decontamination.	2 Hours
5	Elimination Enhancement	2 Hours
6	Toxicokinetics	2 Hours

7	Clinical symptoms and management of acute poisoning with the following agents – a) Pesticide poisoning: organophosphorous compounds, carbamates, organochlorines, pyrethroids. b) Opiates overdose. c) Antidepressants d) Barbiturates and benzodiazepines. e) Alcohol: ethanol, methanol. f) Paracetamol and salicylates. g) Non-steroidal anti-inflammatory drugs. h) Hydrocarbons: Petroleum products and PEG. i) Caustics: inorganic acids and alkali. j) Radiation poisoning	16 Hours
8	Clinical symptoms and management of chronic poisoning with the following agents – Heavy metals: Arsenic, lead, mercury, iron, copper	6 Hours
9	Venomous snake bites: Families of venomous snakes, clinical effects of venoms, general management as first aid, early manifestations, complications and snake bite injuries.	4 Hours
10	Plants poisoning. Mushrooms, Mycotoxins.	2 Hours
11	Food Poisonings	2 Hours
12	Envenomations – Arthropod bites and stings.	2 Hours
13	Substance abuse: Signs and symptoms of substance abuse and treatment of dependence a) CNS stimulants :amphetamine b) Opioids c) CNS depressants d) Hallucinogens: LSD e) Cannabis group f) Tobacco	3 Hours

Fifth year

5.1 CLINICAL RESEARCH (THEORY)

Theory: 3 Hrs. /Week

1. Scope of the subject:

This course is designed to make the students to appreciate the principles and achieve ample knowledge regarding the various approaches to drug discovery including clinical phase of development. Also enables the students to recognize and execute all regulatory and ethical requirements in the process of drug development.

2. Objectives of the subject:

Upon completion of the course the student must be able to;

- a. Understand drug development process
- b. Understand and appreciate ethical guidelines in clinical research
- c. Exhibit good clinical practice and its guidelines
- d. Understand roles and responsibilities of clinical trial personnel
- e. Design clinical study documents and understand informed consent process
- f. Appreciate safety monitoring in clinical trial

Text Books:

Reference Books:

- a. Central Drugs Standard Control Organization. Good Clinical Practices Guidelines for Clinical Trials on Pharmaceutical Products in India. New Delhi: Ministry of Health; 2001.
- b. 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.
- c. Ethical Guidelines for Biomedical Research on Human Subjects 2000. Indian Council of Medical Research, New Delhi.
- d. Textbook of Clinical Trials edited by David Machin, Simon Day and Sylvan Green, March 2005, John Wiley and Sons.
- e. Principles of Clinical Research edited by Giovanna di Ignazio, Di Giovanna and Haynes.
- f. Clinical Data Management edited by R K Rondels, S A Varley, C F Webbs. Second Edition, Jan 2000, Wiley Publications.
- g. Goodman & Gilman: JG Hardman, LE Limbard, 10th Edn. McGraw Hill Publications, 2001.

3. Detailed syllabus and lecture wise schedule:

SL. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Drug development process: - Introduction - Various Approaches to drug discovery 1. Pharmacological 2. Toxicological 3. IND Application 4. Drug characterization 5. Dosage form	6 Hours
2	Clinical development of drug: 1. Introduction to Clinical trials 2. Various phases of clinical trial. 3. Methods of post marketing surveillance 4. Abbreviated New Drug Application submission. 5. Clinical trial approval process 6. Good Clinical Practice – ICH, GCP, Central drug standard control organization (CDSCO) guidelines 7. Challenges in the implementation of guidelines 8. Ethical guidelines in Clinical Research 9. Composition, responsibilities, procedures of IR B / IEC 10. Overview of regulatory environment in USA, Europe and India. 11. Role and responsibilities of clinical trial personnel as per ICH GCP a. Sponsor b. Investigators c. Clinical research associate d. Auditors e. Contract research coordinators f. Regulatory authority 12. Designing of clinical study documents (protocol, CRF, ICF, PIC with assignment) 13. Informed consent Process 14. Data management and its components 14. Safety monitoring in clinical trials.	44 Hours (+3hrs)

5.2 PHARMACOEPIDEMIOLOGY AND PHARMACOECONOMICS (THEORY)

Theory: 3 Hrs. /Week

1. Scope of the subject:

This course is designed to impart information regarding various methods and applications of pharmacoepidemiology and pharmacoeconomics in drug safety monitoring, drug approval and regulations

2. Objectives of the subject:

Upon completion of this course, it is expected that students will be able to -

1. Understand drugs use pattern and their outcome measures
2. carry out pharmacoepidemiological studies
3. implement the tools effectively in evaluating risk and benefit of therapy
4. perform pharmacoeconomic studies and evaluate the cost-benefit ratio

3. Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Pharmacoepidemiology :	
	Definition and scope: Origin and evaluation of pharmacoepidemiology need for pharmacoepidemiology, aims and applications.	6 Hours
	Measurement of outcomes in pharmacoepidemiology Outcome measure and drug use measures, Diagnosis and therapy surveys, Prevalence, incidence and incidence rate. Monetary units, number of prescriptions, units of drugs dispensed, defined daily doses and prescribed daily doses, medication adherence measurement	6 Hours
	Concept of risk in pharmacoepidemiology Measurement of risk, attributable risk and relative risk, time-risk relationship and odds ratio	6 Hours

	Pharmacoepidemiological methods Includes theoretical aspects of various methods and practical study of various methods with the help of case studies for individual methods Drug utilization review, case reports, case series, surveys of drug use, crosssectional studies, cohort studies, case control studies, case –cohort studies, calculation of odd’s ratio, meta-analysis studies, spontaneous reporting, prescription event monitoring and record linkage system.	22 Hours
	Sources of data for pharmacoepidemiological studies Ad Hoc data sources and automated data systems.	4 Hours
	Selected special applications of pharmacoepidemiology Studies of vaccine safety, hospital pharmacoepidemiology, pharmacoepidemiology and risk management, drug induced birth defects.	8 Hours
2	Pharmacoeconomics Definition, history, needs of pharmacoeconomic evaluations Role in formulary management decisions, Need of pharmacoeconomic studies in Indian healthcare system, cost categorisation and resources for cost estimation: Direct costs, Indirect costs, Intangible costs Pharmacoeconomic evaluation Outcome assessment and types of evaluation, Quality adjusted life years, Clinical outcomes, Economic outcomes, Humanistic outcomes, Disability adjusted life years incremental cost effective ratio, Average cost effective ratio, Person time, Willingness to pay Includes theoretical aspects of various methods and practical study of various methods with the help of case studies for individual methods: Cost – minimization, cost- benefit, cost – effectiveness, cost utility	20 Hours
3	Applications of Pharmacoeconomics Software and case studies Definition, steps involved, applications, advantages, disadvantages of the following: health related quality of life(HRQOL):Definition, need, and it’s measures Definitions, steps involved , applications of the following: Decision analysis and decision tree, Sensitivity analysis, Markov’s modelling	3 Hours

5.3 CLINICAL PHARMACOKINETICS AND PHARMACOTHERAPEUTIC DRUG MONITORING (THEORY)

Theory: 2 Hrs. /Week

1. Scope of the subject:

This course is designed to make the students to appreciate and apply pharmacokinetic principles in designing / individualizing dosage regimen. Also, permit the students to interpret the plasma drug range, and hepatic / renal function in optimizing the drug therapy.

2. Objectives of the subject:

Upon completion of the course the student must be able to;

- a. Design dosage regimens
- b. Identify and correlate pharmacokinetics of drug interactions
- c. Indicate and individualise TDM
- d. Perform dose adjustments in renal, hepatic, elderly & paediatric patients
- e. Analyse pharmacokinetic data Understand pharmacogenetics

3. Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Introduction to Clinical pharmacokinetics.	1 Hour
2	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.	7 Hours
3	Pharmacokinetics of Drug Interaction: a. Pharmacokinetic drug interactions b. Inhibition and Induction of Drug metabolism c. Inhibition of Biliary Excretion.	3 Hours

4	Therapeutic Drug monitoring: - Introduction - Individualization of drug dosage regimen (Variability – Genetic, Age and Weight, disease, Interacting drugs). - Indications for TDM. Protocol for TDM. - Pharmacokinetic/Pharmacodynamic Correlation in drug therapy. - TDM of drugs used in the following disease conditions: cardiovascular disease, Seizure disorders, Psychiatric conditions, and Organ transplantations.	20 Hours
5	Dosage adjustment in Renal and hepatic Disease. - Renal impairment - Pharmacokinetic considerations - General approach for dosage adjustment in Renal disease. - Measurement of Glomerular Filtration rate and creatinine clearance. - Dosage adjustment for uremic patients. - Extracorporeal removal of drugs. - Effect of Hepatic disease on pharmacokinetics. - Drug dosing in elderly - Drug dosing in paediatrics	10Hours (+2 hrs)
6	Population Pharmacokinetics. - Introduction to Bayesian Theory. - Adaptive method or Dosing with feedback. - Analysis of Population pharmacokinetic Data.	5 Hours
7	Pharmacogenetics - Genetic polymorphism in Drug metabolism: Cytochrome P-450 Isoenzymes. - Genetic Polymorphism in Drug Transport and Drug Targets. - Pharmacogenetics and Pharmacokinetics/Pharmacodynamic considerations	4 Hours

PHARM.D. (Post Baccalaureate) SYLLABUS

First Year

1.1 PHARMACOTHERAPEUTICS – I & II (THEORY)

Theory: 3 Hrs. /Week

1. **Scope:** 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.
2. **Objectives:** At completion of this subject it is expected that students will be able to understand –
 - a. the pathophysiology of selected disease states and the rationale for drug therapy;
 - b. the therapeutic approach to management of these diseases;
 - c. the controversies in drug therapy;
 - d. the importance of preparation of individualised therapeutic plans based on diagnosis;
 - e. 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);
 - f. describe the pathophysiology of selected disease states and explain the rationale for drug therapy;
 - g. summarise the therapeutic approach to management of these diseases including reference to the latest available evidence;
 - h. discuss the controversies in drug therapy;
 - i. discuss the preparation of individualised therapeutic plans based on diagnosis; and
 - j. 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).

Text Books

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

Reference Books

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

3. Detailed syllabus and lecture wise schedule:

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

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Cardiovascular system Hypertension, Congestive cardiac failure, Angina Pectoris, Myocardial infarction, Hyperlipidemia, Electrophysiology of heart and Arrhythmias.	13 Hours
2	Respiratory system Introduction to Pulmonary function test, Asthma, Chronic obstructive airways disease, Drug induced pulmonary diseases.	06 Hours
3	Endocrine system Diabetes, Thyroid diseases, Oral contraceptives, Hormone replacement therapy, Osteoporosis	08 Hours
4	General prescribing guidelines for 4.1 Paediatric patients 4.2 Geriatric patients 4.3 Pregnancy and breast feeding	04 Hours
5	Ophthalmology Glaucoma, Conjunctivitis- viral & bacterial.	03 Hours
6	Introduction to rational drug use Definition, Role of pharmacist Essential drug concept Rational drug formulations.	02 Hours
7	Infectious disease Guidelines for the rational use of antibiotics and surgical prophylaxis, Tuberculosis, Meningitis, Respiratory tract infections, Gastroenteritis,	18 Hours

	Endocarditis, Septicemia, Urinary tract infections, Protozoal infection-Malaria, HIV & Opportunistic infections, Fungal infections, Viral infections, Gonorrhea and Syphilis.	
8	Musculoskeletal disorders Rheumatoid arthritis, Osteoarthritis, Gout, Spondylitis, Systemic lupus erythematosus.	06 Hours
9	Renal system Acute Renal Failure, Chronic Renal Failure, Renal Dialysis, Drug induced renal disorders.	05 Hours
10	Oncology Basic principles of Cancer therapy, General introduction to cancer chemotherapeutic agents, Chemotherapy of breast cancer, leukemia, lung cancer, colorectal cancer, prostate cancer . Management of chemotherapy nausea and emesis.	06+3 Hours
11	Dermatology Psoriasis, Scabies, Eczema, Impetigo	04 Hours

1.1 PHARMACOTHERAPEUTICS – I & II (PRACTICAL)

Practical: 3 Hrs./Week

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.

Scheme of Practical Examination:

	Sessionals	Annual
Synopsis	05	15
Major Experiment	10	25
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	06hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

1.2 HOSPITAL & COMMUNITY PHARMACY (THEORY)

Theory: 3 Hrs. /Week

1. **Scope:** 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, drug dispensing, manufacturing of parenteral preparations, drug information, patient counselling, and therapeutic drug monitoring for improved patient care.
2. **Objectives:** Upon completion of the course, the student shall be able to –
 - a. know various drug distribution methods;
 - b. know the professional practice management skills in hospital pharmacies;
 - c. provide unbiased drug information to the doctors;
 - d. know the manufacturing practices of various formulations in hospital set up;
 - e. appreciate the practice based research methods; and
 - f. appreciate the stores management and inventory control.

Text books: (latest editions)

- a. Hospital pharmacy by William .E. Hassan
- b. A text book of Hospital Pharmacy by S.H.Merchant & Dr. J.S. Qadry. Revised by R.K.Goyal & R.K. Parikh
- c. Health Education and Community Pharmacy by N.S.Parmar.
- d. WHO consultative group report.
- e. Drug store & Business management by Mohammed Ali & Jyoti.

References:

- a. WHO consultative group report.
- b. R.P.S. Vol.2. Part –B; Pharmacy Practice section.
- c. Handbook of pharmacy – health care. Edt. Robin J Harman. The Pharmaceutical press.
- d. Comprehensive Pharmacy Review – Edt. Leon Shargel. Lippincott Williams & Wilkins.

3. Lecture wise programme:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
A. HOSPITAL PHARMACY		
1.	Hospital - its Organization and functions	1 Hour

2.	Hospital Pharmacy - Organization and management a) Organizational Structure-Staff, Infrastructure & work load statistic b) Management of materials and finance c) Roles & responsibilities of hospital pharmacist	6 Hours
3.	Hospital drug policy a) Pharmacy and Therapeutic committee (PTC) b) Hospital formulary c) Hospital committees - Infection committee - Research and ethical committee d) Developing therapeutic guidelines e) Hospital pharmacy communication - Newsletter	15 Hours
4.	Hospital pharmacy services a) Procurement & warehousing of drugs and Pharmaceuticals b) Inventory control Definition, various methods of Inventory Control ABC, VED, EOQ, Lead time, safety stock c) Drug distribution in the hospital • Individual prescription method • Floor stock method • Unit dose drug distribution method d) Distribution of Narcotic and other controlled substances e) Central sterile supply services – Role of pharmacist	15 Hours
5.	Enteral and parenteral nutrition preparations Total Parenteral Nutrition and I.V. admixtures	4 Hours
6.	Continuing professional development programs Education and training	3 Hours
7.	Radio Pharmaceuticals – Handling and packaging	2 Hours
8.	Professional Relations and practices of hospital pharmacist	3 Hours
9.	Social & Preventive Pharmacy	4Hours

B. COMMUNITY PHARMACY		
1.	Definition, scope, of community pharmacy Roles and responsibilities of Community pharmacist	2 Hours
2.	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 softwares	5 Hours
3.	Responding to symptoms of minor ailments Relevant pathophysiology, common drug therapy to Pain, GI disturbances (Nausea, Vomiting, Dyspepsia, diarrhea, constipation), Pyrexia, Ophthalmic symptoms, worm's infestations.	12 Hours
4.	Code of ethics for community pharmacists	2 Hour
5.	Good Pharmacy Practice	3 Hours
6.	Pharmaceutical care	2 Hours
7.	Patient medication adherence	2 Hours

1.2 HOSPITAL & COMMUNITY PHARMACY (PRACTICAL)

Practical: 3 Hrs./Week

List of Experiments

1. Assessment of drug interactions in the given prescriptions
2. Manufacture of parenteral formulations, powders.
3. Inventory control
4. Prescription Analysis (Analyzing the prescriptions for probable drug interactions and ability to tell the management)

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.

Special requirements:

1. Each college should sign MoU with nearby local hospital having minimum 300 beds for providing necessary training to the students' on hospital pharmacy activities.
2. Well equipped with various resources of drug information.

Scheme of Practical Examination:

	Sessionals	Annual
Synopsis	05	15
Major Experiment	10	25
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	06hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

1.3 CLINICAL PHARMACY (THEORY)

Theory: 3 Hrs./Week

1. Objectives of the Subject :

Upon completion of the subject student shall be able to (Know, do, appreciate) –

- a. To impart ethical values in clinical pharmacy;
- b. monitor drug therapy of patient through medication chart review and clinical review;
- c. obtain medication history interview and counsel the patients;
- d. identify and resolve drug related problems;
- e. detect, assess and monitor adverse drug reaction;
- f. interpret selected laboratory results (as monitoring parameters in therapeutics) of specific disease states; and
- g. retrieve, analyse, interpret and formulate drug or medicine information.

Text books (Theory)

- a. Practice Standards and Definitions - The Society of Hospital Pharmacists of Australia.
- b. Basic skills in interpreting laboratory data - Scott LT, American Society of Health System Pharmacists Inc.
- c. Biopharmaceutics and Applied Pharmacokinetics - Leon Shargel, Prentice Hall publication.
- d. A text book of Clinical Pharmacy Practice; Essential concepts and skills, Dr.G.Parthasarathi et al, Orient Orient Langram Pvt.Ltd. ISSN8125026

References

- a. Australian drug information -Procedure manual. The Society of Hospital Pharmacists of Australia.
- b. Clinical Pharmacokinetics - Rowland and Tozer, Williams and Wilkins Publication.
- c. Pharmaceutical statistics. Practical and clinical applications. Sanford Bolton, Marcel Dekker, Inc.

2. Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Definitions, development and scope of clinical pharmacy Healthcare ethics a. What is ethics? Ethics & law b. Beneficence; Non-beneficence; Respect for autonomy c. Ethical decision making d. Ethical dilemmas in healthcare of Pharmacy practice	3 + 2 Hours

2	Introduction to daily activities of a clinical pharmacist - Drug therapy monitoring (medication chart review, clinical review, pharmacist interventions) - Ward round participation - Adverse drug reaction management - Drug information and poisons information - Medication history - Patient counseling - Drug utilisation evaluation (DUE) and review (DUR) - Quality assurance of clinical pharmacy services	13 Hours
3	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.	3 Hours
4	Clinical laboratory tests used in the evaluation of disease states, and interpretation of test results - Haematological, Liver function, Renal function, thyroid function tests - Tests associated with cardiac disorders - Fluid and electrolyte balance - Microbiological culture sensitivity tests - Pulmonary Function Tests	15 Hours
5	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	8 Hours
6	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 d. Role of pharmacist in management of ADR.	10 Hours
7	Communication skills , including patient counselling techniques, medication history interview, presentation of cases.	10 Hours
8	Pharmaceutical care concepts	4 Hours
9	Critical evaluation of biomedical literature	6 Hours
10	Medication errors	3 Hours

1.3 CLINICAL PHARMACY (PRACTICAL)

Practical: 3 Hrs./Week

Students are expected to perform 15 practicals in the following areas covering the topics dealt in theory class.

- a. Answering drug information questions (4 Nos)
- b. Patient medication counselling (4 Nos)
- c. Case studies related to laboratory investigations (4 Nos)
- d. 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:

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

Scheme of Practical Examination:

	Sessionals	Annual
Synopsis	05	15
Major Experiment	10	25
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	06hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

1.4 BIOSTATISTICS AND RESEARCH METHODOLOGY (THEORY)

Theory: 2 Hrs. /Week

Reference Books:

1. Pharmaceutical statistics- practical and clinical applications, Sanford Bolton, 3rd edition, publisher Marcel Dekker Inc. NewYork.
2. Drug Information- A Guide for Pharmacists, Patrick M Malone, Karen L Kier, John E Stanovich , 3rd edition, McGraw Hill Publications 2006

Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Research Methodology e) Types of clinical study designs: Case studies, observational studies, interventional studies, f) Designing the methodology g) 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 specified width, power of a study h) Report writing and presentation of data	10 Hours
2.1	Biostatistics e) Introduction f) Types of data distribution g) Measures describing the central tendency distributions- average, median, mode h) Measurement of the spread of data -range, variation of mean, standard deviation, variance, coefficient of variation, standard error of mean.	10 Hours
2.2	Data graphics Construction and labeling of graphs, histogram, piecharts, scatter plots, semilogarithmic plots	2 Hours
2.3	Basics of testing hypothesis e) Null hypothesis, level of significance, power of test, P value, statistical estimation of confidence intervals. f) Level of significance (Parametric data)- students t test (paired and unpaired), chi Square test, Analysis of Variance (one-way and two-way)	15 Hours

	b) Level of significance (Non-parametric data)- Sign test, Wilcoxon's signed rank test, Wilcoxon rank sum test, Mann Whitney U test, Kruskal -Wall is test (one way ANOVA) g) Linear regression and correlation- Introduction, Pearson's and Spearmann's correlation and correlation co-efficient. h) Introduction to statistical software: SPSS, Epi Info, SAS.	
2.4	Statistical methods in epidemiology Incidence and prevalence, relative risk, attributable risk	5 Hours
3	Computer applications in pharmacy d) <u>Computer System in Hospital Pharmacy</u>: 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. e) <u>Computer In Community Pharmacy</u>: Computerizing the Prescription Dispensing process; Use of Computers for Pharmaceutical Care in community pharmacy; Accounting and General ledger system f) <u>Drug Information Retrieval & Storage</u>: Introduction – Advantages of Computerized Literature Retrieval; Use of Computerized Retrieval	8 Hours

1.5 CLINICAL TOXICOLOGY (THEORY)

Theory: 2 Hrs. /Week

1. Scope of the subject:

The study of the adverse effects of exogenous agents by assimilating knowledge and techniques from most branches of biochemistry, biology, chemistry, genetics, mathematics, medicine, pharmacology, physiology, and physics and applies safety evaluation and risk assessment to the discipline. In all branches of toxicology, scientists explore the mechanisms by which chemicals produce adverse effects in biological systems. Activities in these broad subjects complement toxicologic research, thereby contributing to the application of this knowledge to the science and art of toxicology.

2. Upon completion of the course the student shall be able to:

- a. Deal with emergencies such as overdoses, poisonings, attempted suicides.
- b. Identify and quantify poisonous compound.
- c. Manage the sign and symptom of over dose
- d. Emergency care-home poisoning.
- e. Knows and appreciates Poison control

Text Books:

Reference Books:

1. Matthew J Ellenhorn. ELLENHORNS MEDICAL TOXICOLOGY – DIAGNOSIS AND TREATMENT OF POISONING. Second edition. Williams and Willkins publication, London
2. V V Pillay. HANDBOOK OF FORENSIC MEDICINE AND TOXICOLOGY. Thirteenth edition 2003 Paras Publication, Hyderabad

3. Detailed syllabus and lecture wise schedule:

I. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	General principles involved in the management of poisoning	3 Hours
2	Antidotes and the clinical applications.	2 Hours
3	Supportive care in clinical Toxicology.	2 Hours
4	Gut Decontamination.	2 Hours
5	Elimination Enhancement	2 Hours
6	Toxicokinetics	2 Hours

7	Clinical symptoms and management of acute poisoning with the following agents – a) Pesticide poisoning: organophosphorous compounds, carbamates, organochlorines, pyrethroids. b) Opiates overdose. c) Antidepressants d) Barbiturates and benzodiazepines. e) Alcohol: ethanol, methanol. f) Paracetamol and salicylates. g) Non-steroidal anti-inflammatory drugs. h) Hydrocarbons: Petroleum products and PEG. i) Caustics: inorganic acids and alkali. j) Radiation poisoning	16 Hours
8	Clinical symptoms and management of chronic poisoning with the following agents – Heavy metals: Arsenic, lead, mercury, iron, copper	6 Hours
9	Venomous snake bites: Families of venomous snakes, clinical effects of venoms, general management as first aid, early manifestations, complications and snake bite injuries.	4 Hours
10	Plants poisoning. Mushrooms, Mycotoxins.	2 Hours
11	Food Poisonings	2 Hours
12	Envenomations – Arthropod bites and stings.	2 Hours
13	Substance abuse: Signs and symptoms of substance abuse and treatment of dependence g) CNS stimulants :amphetamine h) Opioids i) CNS depressants j) Hallucinogens: LSD k) Cannabis group l) Tobacco	3 Hours

1.6 PHARMACOTHERAPEUTICS – III (THEORY)

Theory: 3 Hrs. /Week

1. **Scope:** 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.
2. **Objectives:** At completion of this subject it is expected that students will be able to understand –
 - a. the pathophysiology of selected disease states and the rationale for drug therapy;
 - b. the therapeutic approach to management of these diseases;
 - c. the controversies in drug therapy;
 - d. the importance of preparation of individualised therapeutic plans based on diagnosis;
 - e. 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);
 - f. describe the pathophysiology of selected disease states and explain the rationale for drug therapy;
 - g. to summarize the therapeutic approach to management of these diseases including reference to the latest available evidence;
 - h. to discuss the controversies in drug therapy;
 - i. to discuss the preparation of individualised therapeutic plans based on diagnosis; and
 - j. 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).

Text Books

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

Reference Books

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

3. Detailed syllabus and lecture wise schedule:

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

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Gastrointestinal system: Peptic ulcer disease, Gastro Esophageal Reflux Disease, Inflammatory bowel disease, Liver disorders - Alcoholic liver disease, Viral hepatitis including jaundice, and Drug induced liver disorders.	20 Hours
2	Haematological system: Anaemias, Venous thromboembolism, Drug induced blood disorders.	12 Hours
3	Nervous system: Epilepsy, Parkinsonism, Stroke, Alzheimer's disease	16 Hours
4	Psychiatry disorders: Schizophrenia, Affective disorders, Anxiety disorders, Sleep disorders, Obsessive Compulsive disorders	14 Hours
5	Pain management including Pain pathways, neuralgias, headaches.	8 Hours
6	Evidence Based Medicine	5 Hours

1.6 PHARMACOTHERAPEUTICS – III (PRACTICAL)

Practical: 3 Hrs./Week

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.

Scheme of Practical Examination:

	Sessionals	Annual
Synopsis	05	15
Major Experiment	10	25
Minor Experiment	03	15
Viva	02	15
Max Marks	20	70
Duration	03hrs	04hrs

Note: Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity, promptness, viva-voce and record maintenance).

1.7 BIOPHARMACEUTICS AND PHARMACOKINETICS (THEORY)

Theory: 3 Hrs. /Week

1. Scope of the subject:

This course is designed to impart knowledge and skills necessary for dose calculations, dose adjustments and to apply biopharmaceutics theories in practical problem solving. Basic theoretical discussions of the principles of biopharmaceutics and pharmacokinetics are provided to help the students' to clarify the concepts

2.Objectives of the subject:

- a. Define the basic concepts in biopharmaceutics and pharmacokinetics.
- b. Use raw data and derive the pharmacokinetic models and parameters the best describe the process of drug absorption, distribution, metabolism and elimination.
- c. Critically evaluate biopharmaceutic studies involving drug product equivalency
- d. Design and evaluate dosage regimens of the drugs using pharmacokinetic and biopharmaceutic parameters.
- e. Detect potential clinical pharmacokinetic problems and apply basic pharmacokinetic principles to solve them

Text Books:

Reference Books:

- a. Biopharmaceutics and Clinical Pharmacokinetics by, Milo Gibaldi
- b. Remington's Pharmaceutical Sciences, By Mack Publishing Company, Pennsylvania.
- c. Pharmacokinetics: By Milo Gibaldi Donald, R. Mercel Dekker Inc.
- d. Hand Book of Clinical Pharmacokinetics, By Milo Gibaldi and Laurie Prescott by ADIS Health Science Press.
- e. Biopharmaceutics and Pharmacokinetics; By Robert F Notari
- f. Biopharmaceutics; By Swarbrick
- g. Bio pharmaceutics and Pharmacokinetics-A Treatise, By D. M. Brahmanekar and Sunil B.Jaiswal, Vallabh Prakashan Pitampura, Delhi
- h. Clinical Pharmacokinetics, Concepts and Applications: By Malcolm Rowland and Thomas, N. Tozen, Lea and Febiger, Philadelphia, 1995.
- i. Dissolution, Bioavailability and Bioequivalence, By Abdou H.M, Mack Publishing Company, Pennsylvania 1989.
- j. Biopharmaceutics and Clinical Pharmacokinetics-An introduction 4th edition Revised and expanded by Robert F Notari Marcel Dekker Inc, New York and Basel, 1987.
- k. Encyclopedia of Pharmaceutical Technology, Vol 13, James Swarbrick, James, C. Roylan, Marcel Dekker Inc, New York 1996.

3. Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Biopharmaceutics - Introduction to Biopharmaceutics - Absorption of drugs from gastrointestinal tract. - Drug Distribution. - Biotransformation - Drug Elimination. - Method to determine the pharmacokinetic parameters	16 +14Hours
Pharmacokinetics		
2	Introduction to Pharmacokinetics. - Mathematical model - Drug levels in blood. - Pharmacokinetic model - Compartment models a. e. Pharmacokinetic study.	10 Hours
3	One compartment open model. - Intravenous Injection (Bolus) - Intravenous infusion. - Extravascular administration - Urinary excretion data - Calculations	8 Hours
4	Multicompartment models. - Two compartment open model. - IV bolus, IV infusion and oral administration - Calculations	8- 2=6hrs
5	Multiple – Dosage Regimens. - Repetitive Intravenous injections – One Compartment Open Model - Repetitive Extravascular dosing – One Compartment Open model - Multiple Dose Regimen – Two Compartment Open Model - Calculations	10-4=6 hrs
6	Nonlinear Pharmacokinetics. - Introduction - Factors causing Non-linearity. - Michaelis-menton method of estimating parameters.	7-3=4 hrs

7	Noncompartmental Pharmacokinetics. - Statistical Moment Theory. - MRT for various compartment models. - Physiological Pharmacokinetic model.	8-2=6 hrs
8	Bioavailability and Bioequivalence. - Introduction. - Bioavailability study protocol. - Methods of Assessment of Bioavailability - Methods to improve bioavailability - Calculations	8+1 hrs

1.7 BIOPHARMACEUTICS AND PHARMACOKINETICS (PRACTICAL)

Practical: 3+1=4 Hrs./Week

1. Calibration curves
2. Improvement of dissolution characteristics of slightly soluble drugs by some methods.
 - a. Surfactant addition
 - b. Co-solvent addition
 - c. pH change
3. Comparison of dissolution studies of two different marketed products of same drug.
4. Influence of polymorphism on solubility and dissolution.
5. Protein binding studies of a highly protein bound drug and poorly protein bound drug.
6. 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.
7. Bioavailability studies of some commonly used drugs on animal/human model.
8. Calculation of K_a , K_e , $t_{1/2}$, C_{max} , AUC, AUMC, MRT etc. from blood profile data.
9. Calculation of bioavailability from urinary excretion data for two drugs. Calculation of AUC and bioequivalence from the given data for two drugs.
10. In vitro absorption studies. Absorption studies in animal inverted intestine using various drugs.
11. Bioequivalency studies on the different drugs marketed. (eg) Tetracycline, Sulphamethoxazole, Trimethoprim, Aspirin etc., on animals and human volunteers.
12. Effect on contact time on the plasma protein binding of drugs.
13. Studying metabolic pathways for different drugs based on elimination kinetics data.
14. Calculation of elimination half-life for different drugs by using urinary elimination data and blood level data.
15. Determination of renal clearance.

Second year

2.1 CLINICAL RESEARCH (THEORY)

Theory: 3 Hrs. /Week

1. Scope of the subject: This course is designed to make the students to appreciate the principles and achieve ample knowledge regarding the various approaches to drug discovery including clinical phase of development. Also enables the students to recognize and execute all regulatory and ethical requirements in the process of drug development.

2. Objectives of the subject:

Upon completion of the course the student must be able to;

- a. Understand drug development process
- b. Understand and appreciate ethical guidelines in clinical research
- c. Exhibit good clinical practice and its guidelines
- d. Understand roles and responsibilities of clinical trial personnel
- e. Design clinical study documents and understand informed consent process
- f. Appreciate safety monitoring in clinical trial

Text Books:

Reference Books:

1. 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.
5. Principles of Clinical Research edited by Giovanna di Ignazio, Di Giovanna and Haynes.
6. Clinical Data Management edited by R K Rondels, S A Varley, C F Webbs. Second Edition, Jan 2000, Wiley Publications.
7. Goodman & Gilman: JG Hardman, LE Limbard, 10th Edn. McGraw Hill Publications, 2001.

3. Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Drug development process: - Introduction Various Approaches to drug discovery 1. Pharmacological 2. Toxicological 3. IND Application 4. Drug characterization 5. Dosage form	6 Hours
2	Clinical development of drug: 1. Introduction to Clinical trials 2. Various phases of clinical trial. 3. Methods of post marketing surveillance 4. Abbreviated New Drug Application submission. 5. Clinical trial approval process 6. Good Clinical Practice – ICH, GCP, Central drug standard control organization (CDSCO) guidelines 7. Challenges in the implementation of guidelines 8. Ethical guidelines in Clinical Research 9. Composition, responsibilities, procedures of IR B / IEC 10. Overview of regulatory environment in USA, Europe and India. 11. Role and responsibilities of clinical trial personnel as per ICH GCP a. Sponsor b. Investigators c. Clinical research associate d. Auditors e. Contract research coordinators f. Regulatory authority 12. Designing of clinical study documents (protocol, CRF, ICF, PIC with assignment) 13. Informed consent Process 14. Data management and its components 15. Safety monitoring in clinical trials.	44+3 Hours

2.2 PHARMACOEPIDEMIOLOGY AND PHARMACOECONOMICS (THEORY)

Theory: 3 Hrs./Week

1. Scope of the subject:

This course is designed to impart information regarding various methods and applications of pharmacoepidemiology and pharmacoeconomics in drug safety monitoring, drug approval and regulations

2. Objectives of the subject:

Upon completion of this course, it is expected that students will be able to -

1. Understand drugs use pattern and their outcome measures
2. carry out pharmacoepidemiological studies
3. implement the tools effectively in evaluating risk and benefit of therapy
4. perform pharmacoeconomic studies and evaluate the cost-benefit ratio

3. Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Pharmacoepidemiology :	
	Definition and scope: Origin and evaluation of pharmacoepidemiology need for pharmacoepidemiology, aims and applications.	6 Hours
	Measurement of outcomes in pharmacoepidemiology Outcome measure and drug use measures, Diagnosis and therapy surveys, Prevalence, incidence and incidence rate. Monetary units, number of prescriptions, units of drugs dispensed, defined daily doses and prescribed daily doses, medication adherence measurement	6 Hours
	Concept of risk in pharmacoepidemiology Measurement of risk, attributable risk and relative risk, time-risk relationship and odds ratio	6 Hours
	Pharmacoepidemiological methods Includes theoretical aspects of various methods and practical study of various methods with the help of case studies for individual methods Drug utilization review, case reports, case series, surveys of drug use, cross-sectional studies, cohort studies, case control studies, case –cohort studies, calculation of odd's ratio, meta-analysis studies, spontaneous reporting, prescription event monitoring and record linkage system.	22 Hours

	Sources of data for pharmacoepidemiological studies Ad Hoc data sources and automated data systems.	4 Hours
	Selected special applications of pharmacoepidemiology Studies of vaccine safety, hospital pharmacoepidemiology, pharmacoepidemiology and risk management, drug induced birth defects.	8 Hours
2	Pharmacoeconomics Definition, history, needs of pharmacoeconomic evaluations Role in formulary management decisions, Need of pharmacoeconomic studies in Indian healthcare system, cost categorisation and resources for cost estimation: Direct costs, Indirect costs, Intangible costs Pharmacoeconomic evaluation Outcome assessment and types of evaluation Quality adjusted life years, Clinical outcomes, Economic outcomes, Humanistic outcomes, Disability adjusted life years incremental cost effective ratio, Average cost effective ratio, Person time, Willingness to pay Includes theoretical aspects of various methods and practical study of various methods with the help of case studies for individual methods: Cost – minimization, cost- benefit, cost – effectiveness, cost utility	20 Hours
3	Applications of Pharmacoeconomics Software and case studies Definition, steps involved, applications, advantages, disadvantages of the following: health related quality of life(HRQOL): Definition, need, and it's measures Definitions, steps involved , applications of the following: Decision analysis and decision tree, Sensitivity analysis, Markov's modelling	3 Hours

2.3 CLINICAL PHARMACOKINETICS AND PHARMACOTHERAPEUTIC DRUG MONITORING (THEORY)

Theory: 2 Hrs. /Week

- 1. Scope of the subject:** This course is designed to make the students to appreciate and apply pharmacokinetic principles in designing / individualizing dosage regimen. Also, permit the students to interpret the plasma drug range, and hepatic / renal function in optimizing the drug therapy.
- 2. Objectives of the subject:**
Upon completion of the course the student must be able to;
 - a. Design dosage regimens
 - b. Identify and correlate pharmacokinetics of drug interactions
 - c. Indicate and individualise TDM
 - d. Perform dose adjustments in renal, hepatic, elderly & paediatric patients
 - e. Analyse pharmacokinetic data Understand pharmacogenetics

3.Detailed syllabus and lecture wise schedule:

Sl. NO	TITLE OF THE TOPIC	TOTAL HOURS
1	Introduction to Clinical pharmacokinetics.	1 Hour
2	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.	7 Hours
3	Pharmacokinetics of Drug Interaction: a. Pharmacokinetic drug interactions b. Inhibition and Induction of Drug metabolism c. Inhibition of Biliary Excretion.	3 Hours
4	Therapeutic Drug monitoring: a. Introduction b. Individualization of drug dosage regimen (Variability – Genetic, Age and Weight, disease, Interacting drugs). c. Indications for TDM. Protocol for TDM. d. Pharmacokinetic/Pharmacodynamic Correlation in drug therapy. e. TDM of drugs used in the following disease conditions: cardiovascular disease, Seizure disorders, Psychiatric conditions, and Organ transplantations.	20 Hours

5	Dosage adjustment in Renal and hepatic Disease. - Renal impairment - Pharmacokinetic considerations - General approach for dosage adjustment in Renal disease. - Measurement of Glomerular Filtration rate and creatinine clearance. - Dosage adjustment for uremic patients. - Extracorporeal removal of drugs. - Effect of Hepatic disease on pharmacokinetics. - Drug dosing in elderly - Drug dosing in paediatrics	10 +2 Hours
6	Population Pharmacokinetics. - Introduction to Bayesian Theory. - Adaptive method or Dosing with feedback. - Analysis of Population Pharmacokinetic Data.	5 Hours
7	Pharmacogenetics - Genetic polymorphism in Drug metabolism: Cytochrome P-450 Isoenzymes. - Genetic Polymorphism in Drug Transport and Drug Targets. - Pharmacogenetics and Pharmacokinetics/Pharmacodynamic considerations	4 Hours

APPENDIX-B
CONDITIONS TO BE FULFILLED BY THE
ACADEMIC TRAINING INSTITUTION

1. Any authority or institution in India applying to the Pharmacy Council of India for approval of courses of study for Pharm.D. and Pharm.D. (Post Baccalaureate) under subsection (1) of section 12 of the Pharmacy Act, 1948 shall comply with the infrastructural facilities as prescribed by the Pharmacy Council of India from time to time.
2. Pharm.D. and Pharm.D. (Post Baccalaureate) programmes shall be conducted only in those institutions which -
 - a) are approved by the Pharmacy Council of India for Pharm. D course as provided under section 12 of the Pharmacy Act, 1948;
 - b) have 300 bedded hospital attached to it.

(i) Hospital Details

1. Institution with their own hospital of minimum 300 beds.
2. Teaching hospital recognised by the Medical Council of India or University, or a Government hospital not below the level of district headquarter hospital with 300 beds with clearly defined Memorandum of Understanding including housing pharmacy practice department with minimum carpet area of 30 square feet per student along with consent to provide the professional manpower to support the programme.
3. Corporate type hospital with minimum 300 beds with clearly defined Memorandum of Understanding including housing pharmacy practice department with minimum carpet area of 30 square feet per student along with consent to provide the professional manpower to support the programme.
4. The hospital may not enter into similar agreement(s) with any other Pharmacy institution/s or department/s offering or intend to offer Pharm.D programs.

(ii) Specialty

- a) Tertiary care hospitals are desirable
- b) Medicine [compulsory], and any three specializations like -
 1. Surgery
 2. Pediatrics
 3. Gynecology and obstetrics
 4. Psychiatry
 5. Skin and VD
 6. Orthopedics

(iii) Location of the Hospital

Within the same limits of Corporation or Municipality or Campus with Medical Faculty involvement as adjunct faculty.

3. TEACHING STAFF REQUIREMENT

- (i) Staff Pattern: All faculty shall be full time. However, part time preceptors in hospital shall be allowed.
- (ii) Subject wise specialization of the Teaching Staff:

S.No.	Subject	Specialization required
1.	Pharmacy Practice	M.Pharm in Pharmacy Practice or Pharmacology or Pharmaceutics.
2.	Human Anatomy & Physiology	M.Pharm in Pharmacology or Pharmacy practice
3.	Pharmaceutics (Dispensing & General Pharmacy)	M.Pharm in Pharmaceutics
4.	Pharmacognosy-I	M.Pharm in Pharmacognosy
5.	Pharmaceutical Organic Chemistry-I	M.Pharm in Pharmaceutical chemistry or Pharmaceutical Analysis or Quality assurance or Bulk Drug
6.	Pharmaceutical Inorganic Chemistry	M.Pharm in Pharmaceutical chemistry or Pharmaceutical Analysis or Quality assurance or Bulk Drug
7.	Pharmaceutical microbiology	M.Pharm in Pharmaceutics or Pharmaceutical Biotechnology
8.	Pathophysiology	M.Pharm Pharmacy practice or Pharmacology
9.	Applied Biochemistry & Clinical Chemistry	M.Pharm in Pharmacology or Pharmacy practice or Pharmaceutical chemistry
10.	Pharmacology-I	M.Pharm in Pharmacology or Pharmacy practice
11.	Pharmaceutical Jurisprudence	M.Pharm in Pharmaceutics
12.	Pharmacology-II	M.Pharm in Pharmacology or Pharmacy practice
13.	Pharmaceutical Dosage Forms	M.Pharm in Pharmaceutics or Industrial Pharmacy
14.	Pharmacotherapeutics–I, II and III	M.Pharm Pharmacy practice or Pharmacology
15.	Community Pharmacy	M.Pharm in Pharmacy practice or Pharmacology or Pharmaceutics

16.	Hospital Pharmacy	M.Pharm in Pharmacy practice or Pharmacology or Pharmaceutics
17.	Clinical Pharmacy	M.Pharm in Pharmacy practice
18.	Computer Science or Computer Application in pharmacy	MCA
19.	Mathematics	M.Sc. (Maths)

iii) Teaching Staff :

Department/Division	Name of the post	No.
Department of Pharmaceutics	Professor	1
	Asst. Professor	1
	Lecturer	2
Department of Pharmaceutical Chemistry (Including Pharmaceutical Analysis)	Professor	1
	Asst. Professor	1
	Lecturer	3
Department of Pharmacology	Professor	1
	Asst. Professor	1
	Lecturer	2
Department of Pharmacognosy	Professor	1
	Asst. Professor	1
	Lecturer	1
Department of Pharmacy Practice	Professor	1
	Asst. Professor	2
	Lecturer	3

iv) Prescribed qualifications and experience for Professor, Assistant Professor, Lecturer and others :

Sl. No.	CADRE	QUALIFICATIONS	EXPERIENCE
1.	Lecturer/Assistant Professor	First Class B.Pharm with Master's degree in Pharmacy (M.Pharm) in appropriate branch of specialization in Pharmacy (Qualification must be PCI recognized).	A lecturer will be re-designated as Assistant Professor after 2 years of teaching experience in PCI approved/recognized Pharmacy College.

		A PCI recognized Pharm.D degree holder shall also be eligible for the posts of Lecturer/Assistant Professor in the subjects of pathophysiology, pharmacology and pharmacy practice.	
2.	Associate Professor	First Class B.Pharm with Master's degree in Pharmacy (M.Pharm) in appropriate branch of specialization in Pharmacy (Qualification must be PCI recognized). A PCI recognized Pharm.D degree holder shall also be eligible for the posts of Associate Professor in the subjects of pathophysiology, pharmacology sand pharmacy practice. Associate Professor shall acquire PCI recognized Ph.D in any of Pharmacy subjects within 7 years to become eligible for the post of Professor.	3 years' experience in teaching or research at the level of Assistant Professor or equivalent in PCI approved/recognized Pharmacy College.
3.	Professor	First Class B.Pharm With Master's degree in Pharmacy (M.Pharm) in appropriate branch of specialization in Pharmacy or Pharm.D (Qualifications must be PCI recognized). With Ph.D degree in any of Pharmacy subjects (Ph.D.	Essential 10 years' experience in teaching in PCI approved/recognized Pharmacy College or research experience out of which 5 years must be as Associate Professor in PCI approved/recognized Pharmacy College.

		Qualifications must be PCI recognized).	
4.	Director or Principal or Head of institute	First Class B.Pharm with Master's degree in Pharmacy (M.Pharm) in appropriate branch of specialization in Pharmacy or Pharm.D (Qualifications must be PCI recognized). With Ph.D degree in any of Pharmacy subjects (Ph.D. Qualifications must be PCI recognized).	Essential 15 years' experience in teaching or research out of which 5 years must be as Professor/HOD in a PCI approved/recognized pharmacy college. Desirable Administrative experience in a responsible position.

Note : If a class or division is not awarded at Master's level, a minimum of 60% marks in aggregate or equivalent cumulative grade point average shall be considered equivalent to first class or division, as the case may be.

v) Workload of Faculty :

Professor – 8 hrs. per week

Assistant Professor – 12 hrs. per week

Lecturers – 16 hrs. per week

vi) Training of Pharmacy Practice Faculty:

a) Teaching staff will be trained as per the module prescribed by the Central Council.

b) Duration of training – Minimum 3 months.

c) Training sites – Institutions running pharmacy practice Programmes for at least five years.

d) Trainer – Professor or Assistant Professor with minimum of five years of clinical pharmacy teaching and practice experience.

4) NON-TEACHING STAFF:

Sl.No.	Designation	Required (Minimum)	Required Qualification
1	Laboratory Technician	1 for each Dept	D. Pharm

2	Laboratory Assistants or Laboratory Attenders	1 for each Lab (minimum)	SSLC
3	Office Superintendent	1	Degree
4	Accountant	1	Degree
5	Store keeper	1	D.Pharm or a Bachelor degree recognized by a University or institution.
6	Computer Data Operator	1	BCA or Graduate with Computer Course
7	Office Staff I	1	Degree
8	Office Staff II	2	Degree
9	Peon	2	SSLC
10	Cleaning personnel	Adequate	---
11	Gardener	Adequate	---

5) ACCOMMODATION

Suitable and sufficient accommodation with adequate ventilation, lighting and other hygienic conditions should be provided to the rooms for Principal or the Head of the department, office, class rooms, library, staff, staff common room, students common room, museum, laboratories, stores, etc.

At least two lecture halls along with eight laboratories as specified below should be provided for:

1.	Pharmaceutics and Pharmacokinetics Lab	2
2.	Life Science (Pharmacology, Physiology, Pathophysiology)	2
3.	Phytochemistry / Pharmaceutical Chemistry	2
4.	Pharmacy Practice	2
	TOTAL	8

In addition to the laboratories, balance room, aseptic room or cabinet, animal house and a machine room shall also be provided.

Floor area of the laboratory should not be less than 30 square feet per student required to work in the laboratory at any given time subject to a minimum of 750 square feet.

Laboratories should be fitted and constructed in a manner that these can be kept reasonably clean. Gas and water fittings, shelves, fuming cupboards be provided wherever necessary.

6. EQUIPMENT AND APPARATUS:

Department wise list of minimum equipments

A. DEPARTMENT OF PHARMACOLOGY :

I. Equipment:

S.No.	Name	Minimum required Nos.
1	Microscopes	15
2	Haemocytometer with Micropipettes	20
3	Sahli's haemocytometer	20
4	Hutchinson's spirometer	01
5	Sphygmomanometer	05
6	Stethoscope	05
7	Permanent Slides for various tissues	One pair of each tissue Organs and endocrine glands One slide of each organ system
8	Models for various organs	One model of each organ system
9	Specimen for various organs and systems	One model for each organ system
10	Skeleton and bones	One set of skeleton and one spare bone
11	Different Contraceptive Devices and Models	One set of each device
12	Muscle electrodes	01
13	Lucas moist chamber	01
14	Myographic lever	01
15	Stimulator	01
16	Centrifuge	01
17	Digital Balance	01
18	Physical /Chemical Balance	01
19	Sherrington's Kymograph Machine or Polyrite	10

20	Sherrington Drum	10
21	Perspex bath assembly (single unit)	10
22	Aerators	10
23	Computer with LCD	01
24	Software packages for experiment	01
25	Standard graphs of various drugs	Adequate number
26	Actophotometer	01
27	Rotarod	01
28	Pole climbing apparatus	01
29	Analgesiometer (Eddy's hot plate and radiant heat methods)	01
30	Convulsiometer	01
31	Plethysmograph	01
32	Digital pH meter	01

II. Apparatus:

S.No	Name	Minimum required Nos.
1	Folin-Wu tubes	60
2	Dissection Tray and Boards	10
3	Haemostatic artery forceps	10
4	Hypodermic syringes and needles of size 15,24,26G	10
5	Levers, cannulae	20

NOTE: Adequate number of glassware commonly used in the laboratory should be provided in each laboratory and department.

B. DEPARTMENT OF PHARMACOGNOSY :

I. Equipment:

S.No.	Name	Minimum required Nos.
1	Microscope with stage micrometer	15
2	Digital Balance	02
3	Autoclave	02
4	Hot air oven	02
5	B.O.D.incubator	01
6	Refrigerator	01

7	Laminar air flow	01
8	Colony counter	02
9	Zone reader	01
10	Digital pH meter	01
11	Sterility testing unit	01
12	Camera Lucida	15
13	Eye piece micrometer	15
14	Incinerator	01
15	Moisture balance	01
16	Heating mantle	15
17	Flourimeter	01
18	Vacuum pump	02
19	Micropipettes (Single and multi channeled)	02
20	Micro Centrifuge	01
21	Projection Microscope	01

II. Apparatus:

S.No.	Name	Minimum required Nos.
1	Reflux flask with condenser	20
2	Water bath	20
3	Clavengers apparatus	10
4	Soxhlet apparatus	10
6	TLC chamber and sprayer	10
7	Distillation unit	01

NOTE: Adequate number of glassware commonly used in the laboratory should be provided in each laboratory and department.

C. DEPARTMENT OF PHARMACEUTICAL CHEMISTRY :

I. Equipment:

S.No.	Name	Minimum required Nos.
1	Hot plates	05
2	Oven	03
3	Refrigerator	01
4	Analytical Balances for demonstration	05
5	Digital balance 10mg sensitivity	10

6	Digital Balance (1mg sensitivity)	01
7	Suction pumps	06
8	Muffle Furnace	01
9	Mechanical Stirrers	10
10	Magnetic Stirrers with Thermostat	10
11	Vacuum Pump	01
12	Digital pH meter	01
13	Microwave Oven	02

II. Apparatus:

S.No.	Name	Minimum required Nos.
1	Distillation Unit	02
2	Reflux flask and condenser single necked	20
3	Reflux flask and condenser double/triple necked	20
4	Burettes	40
5	Arsenic Limit Test Apparatus	20
6	Nessler's Cylinders	40

NOTE: Adequate number of glassware commonly used in the laboratory should be provided in each laboratory and department.

D. DEPARTMENT OF PHARMACEUTICS:

I. Equipment:

S.No	Name	Minimum required Nos.
1	Mechanical stirrers	10
2	Homogenizer	05
3	Digital balance	05
4	Microscopes	05
5	Stage and eye piece micrometers	05
6	Brookfield's viscometer	01
7	Tray dryer	01
8	Ball mill	01
9	Sieve shaker with sieve set	01
10	Double cone blender	01
11	Propeller type mechanical agitator	05

12	Autoclave	01
13	Steam distillation still	01
14	Vacuum Pump	01
15	Standard sieves, sieve no. 8, 10, 12,22,24, 44, 66, 80	10 sets
16	Tablet punching machine	01
17	Capsule filling machine	01
18	Ampoule washing machine	01
19	Ampoule filling and sealing machine	01
20	Tablet disintegration test apparatus IP	01
21	Tablet dissolution test apparatus IP	01
22	Monsanto's hardness tester	01
23	Pfizer type hardness tester	01
24	Friability test apparatus	01
25	Clarity test apparatus	01
26	Ointment filling machine	01
27	Collapsible tube crimping machine	01
28	Tablet coating pan	01
29	Magnetic stirrer, 500ml and 1 liter capacity with speed control	05 EACH 10
30	Digital pH meter	01
31	All purpose equipment with all accessories	01
32	Aseptic Cabinet	01
33	BOD Incubator	02
34	Bottle washing Machine	01
35	Bottle Sealing Machine	01
36	Bulk Density Apparatus	02
37	Conical Percolator (glass/copper/ stainless steel)	10
38	Capsule Counter	02
39	Energy meter	02
40	Hot Plate	02
41	Humidity Control Oven	01
42	Liquid Filling Machine	01
43	Mechanical stirrer with speed regulator	02
44	Precision Melting point Apparatus	01
45	Distillation Unit	01

II. Apparatus:

S.No	Name	Minimum required Nos.
1	Ostwald's viscometer	15
2	Stalagmometer	15
3	Desiccator*	05
4	Suppository moulds	20
5	Buchner Funnels (Small, medium, large)	05 each
6	Filtration assembly	01
7	Permeability Cups	05
8	Andreason's Pipette	03
9	Lipstick moulds	10

NOTE: Adequate number of glassware commonly used in the laboratory should be provided in each laboratory and department.

E. DEPARTMENT OF PHARMACEUTICAL BIOTECHNOLOGY :

S.No.	Name	Minimum required Nos.
1	Orbital shaker incubator	01
2	Lyophilizer (Desirable)	01
3	Gel Electrophoresis (Vertical and Horizontal)	01
4	Phase contrast/Trinocular Microscope	01
5	Refrigerated Centrifuge	01
6	Fermenters of different capacity (Desirable)	01
7	Tissue culture station	01
8	Laminar airflow unit	01
9	Diagnostic kits to identify infectious agents	01
10	Rheometer	01
11	Viscometer	01
12	Micropipettes (single and multi channeled)	01 each
13	Sonicator	01
14	Respinometer	01
15	BOD Incubator	01

16	Paper Electrophoresis Unit	01
17	Micro Centrifuge	01
18	Incubator water bath	01
19	Autoclave	01
20	Refrigerator	01
21	Filtration Assembly	01
22	Digital pH meter	01

NOTE: Adequate number of glassware commonly used in the laboratory should be provided in each laboratory and department.

F. DEPARTMENT OF PHARMACY PRACTICE:

Equipment:

S.No.	Name	Minimum required Nos.
1	Colorimeter	2
2	Microscope	Adequate
3	Permanent slides (skin, kidney, pancreas, smooth muscle, liver etc.,)	Adequate
4	Watch glass	Adequate
5	Centrifuge	1
6	Biochemical reagents for analysis of normal and pathological constituents in urine and blood facilities	Adequate
7	Filtration equipment	2
8	Filling Machine	1
9	Sealing Machine	1
10	Autoclave Sterilizer	1
11	Membrane Filter	1 Unit
12	Sintered glass funnel with complete filtering assemble	Adequate
13	Small disposable membrane filter for IV admixture filtration	Adequate
14	Laminar airflow bench	1
15	Vacuum pump	1
16	Oven	1
17	Surgical Dressing	Adequate
18	Incubator	1

19	PH Meter	1
20	Disintegration test apparatus	1
21	Hardness tester	1
22	Centrifuge	1
23	Magnetic stirrer	1
24	Thermostatic bath	1

NOTE:

1. **Computers and Internet connection (Broadband), six computers for students with internet and staff computers as required.**
2. **Adequate number of glassware commonly used in the laboratory should be provided in each laboratory and the department.**

G. CENTRAL INSTRUMENTATION ROOM :

S.No.	Name	Minimum required Nos.
1	Colorimeter	01
2	Digital pH meter	01
3	UV- Visible Spectrophotometer	01
4	Flourimeter	01
5	Digital Balance (1mg sensitivity)	01
6	Nephelo Turbidity meter	01
7	Flame Photometer	01
8	Potentiometer	01
9	Conductivity meter	01
10	Fourier Transform Infra Red Spectrometer (Desirable)	01
11	HPLC	01
12	HPTLC (Desirable)	01
13	Atomic Absorption and Emission spectrophotometer (Desirable)	01
14	Biochemistry Analyzer (Desirable)	01
15	Carbon, Hydrogen, Nitrogen Analyzer (Desirable)	01
16	Deep Freezer (Desirable)	01
17	Ion- Exchanger	01
18	Lyophilizer (Desirable)	01

APPENDIX-C

INTERNSHIP

1) SPECIFIC OBJECTIVES:

- i) to provide patient care in cooperation with patients, prescribers, and other members of an interprofessional health care team based upon sound therapeutic principles and evidence-based data, taking into account relevant legal, ethical, social cultural, economic, and professional issues, emerging technologies, and evolving biomedical, pharmaceutical, social or behavioral or administrative, and clinical sciences that may impact therapeutic outcomes.
- ii) to manage and use resources of the health care system, in cooperation with patients, prescribers, other health care providers, and administrative and supportive personnel, to promote health; to provide, assess, and coordinate safe, accurate, and time-sensitive medication distribution; and to improve therapeutic outcomes of medication use.
- iii) to promote health improvement, wellness, and disease prevention in co-operation with patients, communities, at-risk population, and other members of an interprofessional team of health care providers.
- iv) to demonstrate skills in monitoring of the National Health Programmes and schemes, oriented to provide preventive and promotive health care services to the community.
- v) to develop leadership qualities to function effectively as a member of the health care team organised to deliver the health and family welfare services in existing socio-economic, political and cultural environment.
- vi) to communicate effectively with patients and the community.

2) OTHER DETAILS :

- i) All parts of the internship shall be done, as far as possible, in institutions in India. Internship or residency training not more than 3 months may be allowed in some other hospital / super-specialty hospital in India or abroad.
- ii) Where an intern is posted to district hospital for training, there shall be a committee consisting of representatives of the college or university, and the district hospital

administration, who shall regulate the training of such trainee. For such trainee a certificate of satisfactory completion of training shall be obtained from the relevant administrative authorities which shall be countersigned by the Principal or Dean of College.

- iii) Every candidate shall be required, after passing the final Pharm.D. or Pharm.D. (Post Baccalaureate) examination as the case may be to undergo compulsory rotational internship to the satisfaction of the College authorities and University concerned for a period of twelve months so as to be eligible for the award of the degree of Pharm.D. or Pharm.D. (Post Baccalaureate) as the case may be.

3. ASSESSMENT OF INTERNSHIP :

- i) The intern shall maintain a record of work which is to be verified and certified by the preceptor (teacher practitioner) under whom he works. Apart from scrutiny of the record of work, assessment and evaluation of training shall be undertaken by an objective approach using situation tests in knowledge, skills and attitude during and at the end of the training. Based on the record of work and date of evaluation, the Dean or Principal shall issue certificate of satisfactory completion of training, following which the university shall award the degree or declare him eligible for it.
- ii) Satisfactory completion of internship shall be determined on the basis of the following:-
- (1) Proficiency of knowledge required for each case management SCORE 0-5
 - (2) The competency in skills expected for providing Clinical Pharmacy Services
SCORE 0-5
 - (3) Responsibility, punctuality, work up of case, involvement in patient care
SCORE 0-5
 - (4) Ability to work in a team (Behavior with other healthcare professionals including medical doctors, nursing staff and colleagues). SCORE 0-5
 - (5) Initiative, participation in discussions, research aptitude. SCORE 0-5

Poor	Fair	Below Average	Average	Above Average	Excellent
0	1	2	3	4	5

A Score of less than 3 in any of above items will represent unsatisfactory completion of internship.

APPENDIX-D

CLERKSHIP

Goals:

1. To understand clinical dynamics by attending ward round on daily basis
2. To develop a sufficient, relevant, and experiential knowledge base to utilize appropriate resources necessary to provide direct patient care regarding individualized Therapeutic planning, intervention, and evaluation.
3. To master the clinical skills necessary to assume accountability and responsibility for therapeutic outcomes in the process of providing pharmaceutical care.
4. To develop professionalism and interpersonal skills as a member of team of health care providers, necessary for the provision of optimal patient care and pharmacy services

Objectives:

After successful completion of the clerkship, the student will be able to confidently do

- A. Medication Therapy management
- B. Treatment Chart Review
- C. Drug Interaction Monitoring
- D. Therapeutic Intervention
- E. Adverse Drug Reaction Monitoring
- F. Patient Counseling and Education
- G. Drug Information
- H. Prescription Audit
- I. Health/Community Outreach
- J. Journal club

Reporting format

The sufficient number of clerkship activities as per the above-mentioned objectives must be documented in a bound format and assessed in internal and final examination as per the “Pharm D student’s clerkship activity report and assessment form” annexed in the syllabus book

Textbooks / References

1. DiPiro JT, Talbert RL, Yee GC, Matzke GR, Wells BG, Posey LM, eds. Pharmacotherapy, a Pathophysiologic Approach, Latest Ed. McGraw-Hill, New York, NY.
2. Ruth & Karen. Pharmacy Clerkship Manual McGraw-Hill, New York, NY, 2002
3. Boh L, ed. Pharmacy Practice Clinical Manual, 2ed. Lippincott, 2002.
4. Robert J. Cipolle, Peter C. Morley: Pharmaceutical Care Practice: The Clinician Guide

Scheme of Practical examination

	Sessional	Final Examination
Major Experiment	Perform live Clinical Pharmacy Activity as per defined objectives 10 marks	Perform live Clinical Pharmacy Activity as per defined objectives 30 marks
Minor Experiment	Reviewing clerkship activities carried out by the student as per ACU template by presentation and evaluating their contribution 5 marks	Reviewing clerkship activities carried out by the student as per ACU template and evaluating their contribution 25 marks
Viva	5 marks	15 marks
Maximum Marks	20 marks	70 marks
Duration	3 hours	4 hours
	Total sessional marks is 30 (20 for practical sessional plus 10 marks for regularity& promptness)	

APPENDIX-E
ADICHUNCHANAGIRI UNIVERSITY
Pharm D Student's Clerkship Activity Report & Assessment Form
 Clerkship Summary Sheet
 (To be filled by preceptor at the end of the clerkship)

Name of the Student:

Reg. No.:

Activity	Total Number	Performance (On a 0 to 5 Scale)
1. Patient History Review (Medical & Medication)		
2. Treatment Chart Review		
3. Drug Interaction Monitoring (Major)		
4. Therapeutic Intervention		
5. Adverse Drug Reaction & Hemovigilance Monitoring and Reporting		
6. Patient Counseling and Education		
7. Drug Information		
8. Case Presentation		
9. Seminars and assignments		
10. Journal Club participation and presentation		
11. TDM (optional)		
12. Prescription Audit		
13. Health Outreach Programs or camps/conducted/attended/participated		

(Above activities should be assessed in the Internals and Final examinations)

Signature of the Preceptor	Signature of Head of the Department	Signature of the Principal
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STUDENT'S INTERNSHIP ROTATION REPORT & ASSESSMENT FORM
(Six Reports should be submitted to the University along with Internship completion certificate for the issue of Provisional degree certificate)

Name of the Student:

Reg.No:

Activity	Total Number	Performance (On a 0 to 5 Scale)
1. Patient History Review (Medical & Medication)		
2. Treatment Chart Review		
3. Drug Interaction Monitoring (Major)		
4. Therapeutic Intervention		
5. Adverse Drug Reaction & Hemovigilance Monitoring and Reporting		
6. Patient Counseling & Education		
7. Drug Information		
8. Responsibility, punctuality, work up of case, involvement in patient care		
9. Proficiency of knowledge required for each case management		
10. Initiative, participation in discussions, research aptitude.		
11. Journal Club Attended and Presented		
12. Prescription Audit		
13. Contribution to the newsletter (In terms of articles, news items etc)		
14. Health Outreach Programs or camps/conducted/attended/participated		
15. Conferences /CMEs Attended/ volunteered		
16. Paper/s Presented in Conferences/and published in journals		
17. Ability to work in a team (Behavior with other healthcare professionals including medical doctors, nursing staff and colleagues).		
18. The competency in skills expected for providing Clinical Pharmacy Services		
19. TDM (optional)		

Signature of the Preceptor	Signature of Head of the Department	Signature of the Principal
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ADICHUNCHANAGIRI UNIVERSITY

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Department of Pharmacy Practice

DRUG INFORMATION QUERY REPORT FORM

DIQRF No (Office use):-		QUERY RESPONSE DATE & TIME:-	
QUERY CATEGORY:			
☐ ADMINISTRATION	☐ ADR	☐ AVAILABILITY	☐ COST
☐ DOSAGE	☐ EFFICACY	☐ INTERACTIONS	☐ INDICATION
☐ LACTATION	☐ PHARMACOKINETIC	☐ PREGNANCY	☐ POISONING
☐ THERAPY	☐ OTHERS (SPECIFY):		
RESPONSE PROVIDED:			
☐ IMMEDIATELY	☐ WITHIN 2-4 HRS	☐ WITHIN A DAY	☐ WITHIN 1-2 DAYS
RESPONSE FORMAT:			
☐ ORAL	☐ WRITTEN	☐ EMAIL	☐ OTHERS (SPECIFY)
QUERY RESPONSE IN BRIEF:			
SUPPORTING DOCUMENTS (IF ANY):			
REFERENCE SOURCES:			
ENQUIRER SIGNATURE:			
NAME & SIGNATURE OF THE STUDENT:		NAME & SIGNATURE OF THE STAFF INCHARGE:	

ADICHUNCHANAGIRI UNIVERSITY
 4th T Block, Jayanagar, Bengaluru, Karnataka 560041
Department of Pharmacy Practice
DRUG INFORMATION QUERY REQUEST FORM

DIQRF No (Office use):-	DATE:-	TIME:-
WARD:-	WING:-	SPECIALITY:-
QUERY: 		
QUERY TYPE: ☐ DISEASE MANAGEMENT ☐ CLINICAL STUDIES ☐ DRUG EFFICACY ☐ DRUG SAFETY ☐ POISON MANAGEMENT ☐ OTHERS (SPECIFY):		
PURPOSE OF THE QUERY: ☐ UPDATING KNOWLEDGE ☐ BETTER PATIENT CARE ☐ OTHERS (SPECIFY):		
PATIENT DETAILS (If Required): Attach Patient Profile From		
MODE OF REQUEST: ☐ DIRECT ACCESS ☐ QUERY BOX ☐ EMAIL ☐ TELEPHONE ☐ POST		
RESPONSE TIME: ☐ IMMEDIATELY ☐ WITHIN 2-4 HRS ☐ WITHIN A DAY ☐ WITHIN 1-2 DAYS		
RESPONSE REQUIRED: ☐ ORALLY ☐ WRITTEN ☐ DETAILED INFORMATION ☐ EMAIL ☐ OTHERS (SPECIFY):		
ENQUIRER PROFESSIONAL STATUS: ☐ CONSULTANT ☐ RESIDENT ☐ PHARMACIST ☐ NURSE ☐ OTHERS (SPECIFY):		
NAME OF THE ENQUIRER:		
EXTN NO.:	MOBILE NO.:	EMAIL:

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Department of Pharmacy Practice

DRUG INTERACTION DOCUMENTATION FORM

DICF No (Office use):-		DATE & TIME:-		IP/OP NO.:-		DOA:-	
AGE:-		SEX:-		UNIT:-		SPECIALITY:-	
PATIENT TREATMENT DETAILS : Attach Patient Profile From							
DRUG INTERACTION CHART:							
INTERACTING DRUGS				INTERACTING DRUGS			
DRUG INTERACTION CLASSIFICATION:							
PHARMACOKINETIC				PHARMACODYNAMIC			
☐ ABSORPTION		☐ DISTRIBUTION		☐ SYNERGISM		☐ ANTAGONISM	
☐ METABOLISM		☐ EXCRETION		☐ NEUTRALISATION			
DRUG INTERACTION EFFECT:							
PHARMACOLOGICAL				CLINICAL			
BRIEF DESCRIPTION OF DRUG INTERACTION:							
TYPE OF DRUG INTERACTION:							
☐ DRUG-DRUG ☐ DRUG-FOOD ☐ DRUG-HERBAL ☐ DRUG-DISEASE ☐ ANY OTHER							
DRUG INTERACTION SEVERITY:							
☐ MAJOR ☐ MODERATE ☐ MINOR							
DRUG INTERACTION ASSESSMENT:							
☐ DEFINITE ☐ PROBABLE ☐ SUSPECTED ☐ POSSIBLE ☐ UNLIKELY							
DRUG INTERACTION MANAGEMENT:							
☐ DRUG WITHDRAWN		☐ DOSE REDUCED		☐ TIME SPACING			
☐ CONTINUE WITH MONITOR		☐ OTHERS (SPECIFY):					
SUGGESTIONS ACCEPTED: ☐ YES ☐ NO							
IF NO, GIVE REASON:							
REFERENCES:							
NAME & SIGNATURE OF THE STUDENT:				NAME & SIGNATURE OF THE STAFF INCHARGE:			

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Department of Pharmacy Practice

PATIENT COUNSELING DOCUMENTATION FORM

PCDF No (Office use):-		DATE & TIME:-		IP /OPNO.:-		DOA:-	
AGE:-		SEX:-		WARD & WING:-		SPECIALITY:-	
PATIENT AND TREATMENT DETAILS : Attach Patient Profile From							
COUNSELING PROVIDED TO: ☐ PATIENT ☐ PATIENT REPRESENTATIVE							
IF PATIENT REPRESENTATIVE, GIVE REASON:							
☐		☐		☐			
☐		☐					
COUNSELED ABOUT:							
☐ DRUGS		☐ DISEASE		☐ LIFE STYLE			
COUNSELING AIDS USED:							
☐ PICTOGRAMS		☐ DUMMY INHALERS		☐ SPACER			
☐ NONE		☐ OTHERS (SPECIFY):					
COUNSELING MATERIAL PROVIDED:							
☐ PATIENT INFORMATION LEAFLETS		☐ PRODUCT INFORMATION LEAFLET					
☐ PAMPHLET		☐ NONE		☐ OTHERS (SPECIFY):			
MAJOR BARRIERS INVOLVED: ☐ YES ☐ NO							
IF YES, ☐ PATIENT BASED ☐ PROVIDER BASED ☐ SYSTEM BASED							
COUNSLEING STEPS				INFORMATION PROVIDED			
CASE SHEET REVIEWED ☐				INFORMATION ON SIGNS AND SYMPTOMS/DISEASE ☐			
SELF-INTRODUCTION ☐				NAME AND PURPOSE OF MEDICATION ☐			
PURPOSE OF COUNSELING EXPLAINED ☐				DOSAGE REGIMEN ADVICE ON MISSED DOSE ☐			
INITIAL DRUG RELATED INFORMATION OBTAINED ☐				POTENT SIDE EFFECTS ☐			
PATIENT WAS WARNED ABOUT TAKING OTHER MEDICATION INCLUDING OTC'S, HERBAL DRUGS ETC ☐				SIGNIFICANT INTERACTIONS (DRUG-DRUG, DRUG-FOOD, DRUG-DISEASE) ☐			
ACTUAL COUNSELING DONE ☐				PRECAUTIONS TO BE TAKEN ☐			
PATIENT UNDERSTANDINGS TOWARDS THERAPY WAS ASCERTAINED ☐				DRUG STORAGE RECOMMEN ☐ TIONS ☐			
COUNSELING POINTS SUMMARIZED ☐				BENEFITS OF COMPLETING THE MEDICATION COURSE ☐			
				LIFE STYLE MODIFICATION ☐			
REMARKS:-							
NAME & SIGNATURE OF THE STUDENT:				NAME & SIGNATURE OF THE STAFF INCHARGE:			

ADICHUNCHANAGIRI UNIVERSITY

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Department of Pharmacy Practice

THERAPEUTIC INTERVENTION FORM

TIF No (Office use):-		DATE & TIME:-	IP NO.:-	DOA:-
AGE:-	SEX:-	WARD & WING:-	SPECIALITY:-	
PATIENT TREATMENT DETAILS : Attach Patient Profile From				
BRIEF DESCRIPTION OF THE INTERVENTION:-				
INTERVENTION IDENTIFIED DURING:				
☐ PRE ROUNDS ☐ WARD ROUNDS ☐ SPECIAL VISITS				
☐ OTHERS (SPECIFY):				
SUGGESTIONS MADE:				
☐ DRUG WITHDRAWN ☐ DOSE TAPERED ☐ DOSE INCREASED				
☐ DRUG READMINISTRATION ☐ NEW DRUG ADDED ☐ OTHERS (SPECIFY):				
SUGGESTIONS ACCEPTED: ☐ YES ☐ NO				
IF NO, GIVE REASON:				
DRUG THERAPY CHANGED: ☐ YES ☐ NO				
IF NO, GIVE REASON:				
INTERVENTION CATEGORY:				
☐ ADR ☐ CONTRA INDICATION ☐ DRUG DUPLICATION ☐ DOSAGE FORM				
☐ DRUG USE WITHOUT INDICATION ☐ DOSE ADJUSTMENT IN RENAL IMPAIRMENT				
☐ DOSE ADJUSTMENT IN SPECIAL POPULATION ☐ DRUG SHORTAGE				
☐ DOSE ADJUSTMENT IN HEPATIC IMPAIRMENT ☐ DRUG INTERACTION				
☐ EXPIRED DRUG ☐ INCOMPATIBILITY ☐ INAPPROPRIATE DRUG SELECTION				
☐ MEDICATION ADHERENCE ☐ OVERDOSE ☐ SUB THERAPEUTIC DOSE				
☐ UNTREATED INDICATION ☐ UNACCEPTED ABBREVIATION ☐ OTHERS				
REFERENCES:				
NAME & SIGNATURE OF THE STUDENT:		NAME & SIGNATURE OF THE STAFF INCHARGE:		

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Department of Pharmacy Practice

PATIENT PROFILE FORM

IP No:-	DOA:-	WARD/BED No:-	SPECIALITY:-
AGE:-	SEX: -	HT IN CM:-	WT IN KG:-
	ALLERGIES:-		
PRESENTING COMPLAINTS:-		HISTORY OF PRESENTING COMPLAINTS:-	
FAMILY HISTORY:-		PAST HISTORY:-	
PERSONAL HISTORY:-		OBSTETRIC HISTORY:- (WHENEVER RELEVANT) DEVELOPMENT HISTORY:- (WHENEVER RELEVANT)	
CURRENT MEDICATIONS:-		GENERAL PHYSICAL EXAMINATION:-	
BUILT:- WELL/MODERATE/POOR	BP:-	PULSE:-	RR:-
NUTRITION:- WELL/MODERATE/POOR	GCS:-	CLUBBING:-	CYANOSIS:-
LYMPHADENOPATHY:-	PALLOR:-	EDEMA:-	ICTERUS:-
CLINICAL EXAMINATION:-			

	Plts:					Lipid Profile			
	PT/PTT					LFT			
	ESR:-					Others			
	Others								
Microbiology	Urine				Radiology	CXR			
	Faeces					Ultrasound			
	Cultures					ECG			
	Cavity fluids					Others			
Others					Others				

PHYSICIAN/NURSE NOTES

DAY 1	DAY 2	DAY 3
DAY 4	DAY 5	DAY 6

SIGNATURE OF STUDENT:-	SIGNATURE OF IN CHARGE:-
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SOAP ANALYSIS

