University of Mumbai



No. UG/67 of 2019-20

CIRCULAR:-

Attention of the Principals of the Affiliated Colleges, Directors of the recognized Institutions in Science & Technology Faculty is invited to this office Circular No. UG/193 of 2016-17, dated 3rd December, 2016 relating to the Credit System Manual revised syllabus for B. Pharm. & M. Pharm. (Sem. I to IV).

They are hereby informed that the recommendations made by the Ad-hoc Board of Studies in Pharmacy at its meeting held on 14th June, 2019 have been accepted by the Academic Council at its meeting held on 26th July, 2019 vide item No. 4.39 and that in accordance therewith, the Manual and Revised Scheme and Syllabus as per the (CBCS) for the Bachelor of Pharmacy (Sem. I to VIII) has been brought into force with effect from the academic year 2019-20, accordingly. (The same is available on the University's website www.mu.ac.in). Linas

MUMBAI - 400 032 14 Maugust, 2019

(Dr. Ajay Deshmukh) REGISTRAR

To

The Principals of the affiliated Colleges, and Directors of the recognized Institutions in Science & Technology Faculty. (Circular No. UG/334 of 2017-18 dated 9th January, 2018.)

A.C/4.39/26/07/2019

No. UG/67 -A of 2019-20

MUMBAI-400 032

14 August, 2019

Copy forwarded with Compliments for information to:-

- 1) The I/c Dean, Faculty of Science & Technology,
- 2) The Chairman, Board of Studies in Pharmacy,
- 3) The Director, Board of Examinations and Evaluation,
- 4) The Director, Board of Students Development,
- 5) The Co-ordinator, University Computerization Centre,

(Dr. Ajay Deshmukh) REGISTRAR

AC 26 07/2019 Item No. 4-39

UNIVERSITY OF MUMBAI



Syllabus for Approval

Sr. No.	Heading	Particulars
1	Title of the Course	Buchelor of Pharmacy (B. Pharm.)
2	Eligibility for Admission	H.S.C /CET
3	Passing Marks	500%
4	Ordinances / Regulations (if any)	
5	No. of Years / Semesters	4 (Four) Years / 8 (class) Semisters
6	Level	P.G. / U.G./ Diploma / Certificate (Strike out which is not applicable)
7	Pattern	Yearly / Semester (Strike out which is not applicable)
8	Status	New / Revised (Strike out which is not applicable)
9	To be implemented from Academic Year	From Academic Year 2019-20

Date: 9/7/19

Signature:

Name of BOS Chairperson / Dean : 97 L 16 Lie

KRISHNA MER PLD.

UNIVERSITY OF MUMBAI



Manual on CHOICE BASED CREDIT SYSTEM

for

Undergraduate Programme (Bachelor of Pharmacy, B. Pharm.)

in

PHARMACY

Revised Course (Revised 2019) from the academic year 2019-2020

INTRODUCTION

RECOMMENDATIONS OF NATIONAL REGULATORY AUTHORITIES

The University Grants Commission (UGC), the National Assessment and Accreditation Council (NAAC), the Distance Education Council (DEC) and the National Knowledge Commission (NKC) have time and again come out with recommendations for improving the quality and effectiveness of Higher education provisions in the country. The ministry of Human Resource Development at the Central level and the Ministry of Higher & Technical Education, Govt. of Maharashtra have also repeatedly stressed on the need for universities to pay prompt attention to improve the quality of education. The National Knowledge Commission (NKC), in its report to the Prime Minister on 29th November 2006) has also reiterated the importance of higher education and the contribution it has made to economic development, social progress and political democracy in independent India.

An important concern voiced more strongly in recent times, is the need to develop a Choice-Based Credit System (CBCS) in tune with global trends and the adoption of a sound grading system for reflecting learner performance. This is in line with the recommendation of the UGC in its Action Plan for Academic and Administrative Reforms (Ref. UGC letters January 2008; March 2009) "...... Curricular flexibility and learners' mobility is an issue that warrants our urgent attention. These can be addressed by introducing credit based courses and credit accumulation. In order to provide with some degree of flexibility to learners, we need to provide flexibility in course selection and also a minimum as well as a maximum permissible span of time in which a course can be completed by a learner... The Choice-Based Credit System (CBCS) imminently fits into the emerging socioeconomic milieu, and could effectively respond to the educational and occupational aspirations of the upcoming generations. In view of this, institutions of higher education in India would do well to invest thought and resources into introducing CBCS. Aided by modern communication and information technology, CBCS has a high probability to be operationalized efficiently and effectively — elevating learners, institutions and higher education system in the country to newer heights...".

RATIONALE FOR INTRODUCTION OF CREDIT AND GRADING SYSTEM

The UGC while outlining the several unique features of the Choice-Based Credit System (CBCS) has, in fact, given in a nutshell, the rationale for its introduction. Among the features highlighted by the UGC are: Enhanced learning opportunities, ability to match learners' scholastic needs and aspirations, inter-institution transferability of learners (following the completion of a semester), part-completion of an academic programme in the institution of enrolment and part-completion in a specialized (and recognized) institution, improvement in educational quality and excellence, flexibility for working learners to complete the programme over an extended period of time, standardization and comparability of educational programmes across the country, etc.

This Choice Based Credit System enables a much-required shift in focus from teacher-centric to learner-centric education since the workload estimated is based on the investment of time in learning, not in teaching. It also focuses on continuous evaluation which will enhance the quality of education. It can be concluded from the above discussion that it is very much essential to implement the Choice Based Credit System in higher education in India. Course credit structure, examination/assessment and grading are mainly focused aspects of this manual and discussed in subsequent chapters.

DIRECTIVES OF PHARMACY COUNCIL OF INDIA

The Pharmacy Council of India (PCI) in exercise of the powers conferred to it under the sections 10 and 18 of the Pharmacy Act 1948 (8 of 1948), with the approval of the Central Government, had made the Bachelor of Pharmacy (B. Pharm.) Course Regulations, 2014 and Master of Pharmacy (M. Pharm.) Course regulations vide Gazette dated December 10, 2014. Further as per regulations 6 and 8 of the above course regulations the PCI has also prescribed the Rules and Syllabus for B. Pharm. course and Scheme and Syllabus for M. Pharm., its letter Ref 14-136/2016-PCI and Ref 14-154/2015 PCI dated December 21, 2016, with the subject heading "Statutory Scheme/Rules and syllabus for B. Pharm and M. Pharm. courses". It is thus mandatory to implement the directives of PCI with regard to the Rules/Regulations/Syllabus for recognition and extension of approval of B. Pharm. and M. Pharm. programs of institutes/Universities by the PCI

1. ADMISSION CRITERIA

Admission to the B. Pharm. program of University of Mumbai is governed by the rules and regulations of University of Mumbai and is as per norms of the Govt. of Maharashtra through the State CET-CELL (Maharashtra State), the All India Council for Technical Education (AICTE, New Delhi), and Pharmacy Council of India (PCI, New Delhi). Minimum qualification for admission into Bachelor of Pharmacy program would be according to the rules and regulations of AICTE, PCI, Government of Maharashtra and University of Mumbai in force at the time of admission.

Admission criteria for First Year B. Pharm is as follows:

In general, a learner who has passed HSC or its equivalent examination with Physics and Chemistry as compulsory subjects along with one of the Mathematics or Biotechnology or Biology and obtained at least 50% marks (at least 45 marks in case of candidates of backward category and persons with disability belonging to Maharashtra state only.) in the above subjects taken together and obtained score in CET/ NEET / any other equivalent exam is eligible for admission to Semester I of First Year B. Pharm. However, the rules/regulations and qualifications for admission will be those in effect at the day and time of admission.

➤ Admission criteria for admission into Semester III of Bachelor of Pharmacy (Lateral Entry to Second Year B. Pharm.) is as follows:

In general, a learner who has passed Diploma course in Pharmacy with an aggregate of 45% (at least 40 marks in case of candidates of backward category and persons with disability belonging to Maharashtra state only.) from an All India Council for Technical Education or Pharmacy Council of India or Central or State Government approved institutions or its equivalent. However, the rules/regulations and qualifications for admission will be those in effect at the day and time of admission.

2. COURSE STRUCTURE

2.1. Duration of the program

The course of study for B. Pharm. shall extend over a period of eight semesters (four academic years) and six semesters (three academic years) for lateral entry students. The medium of instruction shall be English.

2.2. Working days in each semester

Each semester shall consist of not less than 100 working days. The odd semesters shall be conducted from the month of July to November/December and the even semesters shall be conducted from December/January to May/June in every calendar year.

As the requirements for a particular degree (undergraduate or postgraduate), a certain quantum of academic work measured in terms of credits is laid down in general. Learner earns credits every semester by satisfactorily clearing courses/other academic activities. The amount of credit associated with a course is dependent upon the number of hours of instruction per week in that course. Similarly the credit associated with any of the other activities is dependent upon the quantum of work expected to be put in for each of the other activity per week.

2.3. Attendance and progress

A candidate is required to put in at least 75% attendance in individual courses considering theory and practical separately. The candidate shall complete the prescribed course satisfactorily to be eligible to appear for the respective examinations.

2.4. Credit Assignment

2.4.1. Theory and Laboratory Courses:

Courses are broadly classified as *Theory courses* and *Laboratory Courses*. Theory courses consist of lecture (**L**) and /or tutorial (**T**) hours. Laboratory courses consist of practical hours, but may have attached tutorial hours in special cases. Credit (**C**) for a course is dependent on the number of hours of instruction per week in that course, and is obtained by using a multiplier of one (**1**) for lecture and tutorial hours, and a multiplier of half (**1**/**2**) for laboratory hours. Thus, for example, a theory course having **four** lectures and **one** tutorial per week throughout the semester carries a credit of **5**. Similarly, a laboratory course having **two** laboratory hours per week throughout semester carries a credit of **1**.

For example -

Theory course				
L	C			
3	1		4	

Laboratory course					
P T =					
4	0		2		

2.4.2. Projects/Dissertations

Project is a requirement for the B. Pharm. degree, wherein under the guidance of a faculty member, a group of not more than five learners in the eighth semester, is required to do some innovative work with the application of knowledge gained while learning various courses in the earlier years. The learner/s is/are expected to do a survey of literature in the subject, work out a Project plan and carry it out through survey, experimentation and/or modeling / computation. Through the Project work the learner has to exhibit skills for both analysis and critical thinking. The complete details of the project have to submitted as a report of not less than 25 pages (A4, 1 inch margins, single line space, font Times Roman, font size 12, excluding count of reference pages) to the College before the prescribed date. The credits assigned for Project is 6 credits.

2.4.3. Practice School

In the VII semester, every candidate shall undergo practice school for a period of 150 hours evenly distributed throughout the semester. The student shall opt any one of the domains for practice school declared by the program committee from time to time.

At the end of the practice school, every student shall submit a printed report (in triplicate) on the practice school he/she attended (not more than 25 pages). Along with the exams of semester VII, the report submitted by the student, knowledge and skills acquired by the student through practice school shall be evaluated by the subject experts at college level and grade point shall be awarded. The credits assigned for Practice School is 6 credits.

2.4.4. Industrial training (Desirable)

Every candidate shall be required to work for at least 150 hours spread over four weeks in a Pharmaceutical Industry/Hospital. It includes Production unit, Quality Control department, Quality Assurance department, Analytical laboratory, Chemical manufacturing unit, Pharmaceutical R&D, Hospital (Clinical Pharmacy), Clinical Research Organization, Community Pharmacy, etc. After the Semester – VI and before the commencement of Semester – VII, and shall submit satisfactory report of such work and certificate duly signed by the authority of training organization to the head of the institute.

2.5. Minimum Credit Requirements

The minimum credit points required for award of a B. Pharm. degree is 208 plus credit of 1 for extra-curricular and co-curricular activities. These credits are divided into Theory courses, Tutorials, Practical, Practice School and Project over the duration of eight semesters. The credits are distributed semester wise as shown in the structure and syllabus manual. Courses generally progress in sequences, building competencies and their positioning indicates certain academic maturity on the part of the learners. Learners are expected to follow the semester wise schedule of courses given in the syllabus.

The lateral entry students shall get **52** credit points transferred from their D. Pharm program. Such students shall take up additional remedial courses of 'Communication Skills' (Theory and Practical) and 'Computer Applications in Pharmacy' (Theory and Practical) equivalent to 3 and 4 credit points respectively, a total of 7 credit points to attain 59 credit points, the maximum of I and II semesters.

2.6. Course/Subject codes:

In the syllabus manual of each programme of a particular discipline, subject code is assigned for each course as follows:

- First two characters are alphabets and indicate the program of a particular discipline (BP indicates B. Pharm.).
- Third digit indicates semester number (1 indicates first Semester).
- The next two digits indicate chronological order of the course in the list of the subjects of the respective semester (01 indicates first course in the list of all courses of the respective semester).
- Alphabets onward sixth indicates nature of the course i.e. T indicates Theory, P indicates Practical and ET indicates Elective Theory, R indicates Remedial, PS indicates Practice School, PW indicates Project Work.

For example -

- ✓ BP101T indicates a B. Pharm. course of semester one and first theory course
- ✓ BP107P indicates a B. Pharm. course of semester one and seventh practical course
- ✓ BP807ET indicates a B. Pharm. course of semester eight and seventh elective theory course

3. EXAMINATION / ASSESSMENT AND GRADING

Semester wise performance assessment of every registered learner is to be carried out through various modes of examinations, in both theory and laboratory classes. These include Internal Assessment and End Semester Examination.

3.1. End semester examinations

The End Semester Examinations in Semesters I, II, III, IV, V and VI of the B. Pharm. Degree course will be conducted by the respective institutions/colleges where the learner has been admitted following rules and regulations. The examinations in Semesters VII and VIII will be conducted by the university. All Non University Examination Subjects (NUES) marked with asterisk symbol (*) syllabus structure, will have examinations conducted by the subject experts at college level and the marks/grades shall be submitted to the university.

A common time-table and common question papers for all the theory examinations of different semesters will be prepared/set by the university as per the procedure.

The question papers for the Theory courses in Semesters I, II, III, IV, V, VI, VII, and VIII will be set by examiners and paper-setters appointed by the University.

The assessment and moderation of the answer booklets for the examinations in Theory courses in Semesters I to VI will be carried out by respective institutions/colleges by the examiners and moderators appointed by the principals of the institutions/colleges for each paper from the panel approved by the Ad-hoc Board of Studies in Pharmacy.

Principals of the respective institutions/colleges are authorized to appoint examiners in the Practical examinations at Semesters I to VI on behalf of the university, only from the panel of suitable persons for appointment as examiners prepared by the Adhoc Board of Studies in Pharmacy.

The assessment and moderation of the answer booklets of the Theory courses in Semesters VII and VIII will be conducted by the University through Central

Assessment Programme (CAP) or On Screen Marking (OSM) or as directed by the University of Mumbai.

The End Semester Examination for Laboratory classes for Semesters I to VI would be done at the institutional level by a pair of examiners appointed by the institution. For Semesters VII and VIII, the University would appoint two examiners for each Laboratory prescribed in Semesters VII and VIII. Evaluation would be done by the examiners appointed by the University at the place and time announced by the University.

3.2. Internal assessment: Continuous mode

The marks allocated for Continuous mode of Internal Assessment shall be awarded as per the scheme given below.

Table-1: Scheme for awarding internal assessment: Continuous mode

Theory		
Criteria	Maximu	m Marks
	100 M Course	50 M Course
Attendance (Refer Table – 2)	4	2
Academic activities (Average of any 3 activities e.g. quiz, assignment, open book test, field work, group discussion and seminar)	3	1.5
Student – Teacher interaction	3	1.5
Total	10	5
Practical		
Attendance (Refer Table – 2)	2	2
Based on Practical Records, Regular viva voce, etc.		3
Total		5

Table- 2: Guidelines for the allotment of marks for attendance

Percentage of		
Attendance	Theory	Practical
90 – 100	4	2
95 – 99	3	1.5
80 – 84	2	1
75 – 79	1	0.5

Less than 75	0	0

3.3. Sessional Exams

Two Sessional exams shall be conducted for each theory / practical course as per the schedule fixed by the college(s). The scheme of question paper for theory and practical Sessional examinations is given below. The average marks of two Sessional exams shall be computed for internal assessment as per the requirements given in tables of Schemes for internal assessments and end semester examinations.

Sessional exam shall be conducted for 30 marks for theory and shall be computed for 15 marks. Similarly Sessional exam for practical shall be conducted for 40 marks and shall be computed for 10 marks.

Question paper pattern for theory Sessional examinations

For subjects having University examination			
I. Multiple Choice Questions (MCQs)			
(Answer all the questions)	=	$10 \times 1 = 10$	
I. Long Answers (Answer 1 out of 2)	=	$1 \times 10 = 10$	
II. Short Answers (Answer 2 out of 3)	=	$2 \times 5 = 10$	
Total	=	30 marks	
➤ For subjects having Non University Examination			
I. Long Answers (Answer 1 out of 2)	=	$1 \times 10 = 10$	
II. Short Answers (Answer 4 out of 6)	=	$= 4 \times 5 = 20$	
Total	=	30 marks	
> Question paper pattern for practical sessional examinations	5		
I. Synopsis	=	10	
II. Experiments	=	25	
III. Viva voce	=	05	
Total	=	40 marks	

4. PROMOTION AND AWARD OF GRADES

A student shall be declared **PASS** and eligible for getting grade in a course of B. Pharm. programme if he/she secures at least **50%** marks in that particular course including internal assessment. For example, to be declared as PASS and to get grade, the student has to secure a minimum of 50 marks for the total of 100 including continuous mode of assessment and end semester theory examination and has to

secure a minimum of 25 marks for the total 50 including internal assessment and end semester practical examination.

5. CARRY FORWARD OF MARKS

In case a student fails to secure the minimum **50%** in any Theory or Practical course, then he/she shall reappear for the end semester examination of that course. However his/her marks of the Internal Assessment shall be carried over and he/she shall be entitled for grade obtained by him/her on passing.

6. IMPROVEMENT OF INTERNAL ASSESSMENT

A student shall have the opportunity to improve his/her performance only once in the Sessional exam component of the internal assessment. The re-conduct of the Sessional exam shall be completed before the commencement of next end semester theory examinations

7. RE-EXAMINATION OF END SEMESTER EXAMINATIONS

Reexamination of end semester examination shall be conducted as per the schedule given in table 3. The exact dates of examinations shall be notified from time to time.

Table-3: Tentative schedule of end semester examinations

Semester	For Regular Candidates	For Failed Candidates	
I, III, V and VII	November / December	May / June	
II, IV, VI and VIII	May / June	November / December	

Question paper pattern for end semester theory examinations

For 75 marks paper

I. Multiple Choice Questions(MCQs)

(Answer all the questions) = $20 \times 1 = 20$

I. Long Answers (Answer 2 out of 3) $= 2 \times 10 = 20$

II. Short Answers (Answer 7 out of 9) $= 7 \times 5 = 35$

Total = 75 marks

.....

For 50 marks paper

I. Long Answers (Answer 2 out of 3) $= 2 \times 10 = 20$

II. Short Answers (Answer 6 out of 8)
$$= 6 \times 5 = 30$$

.____

Total = 50 marks

For 35 marks paper

I. Long Answers (Answer 1 out of 2)
$$= 1 \times 10 = 10$$

II. Short Answers (Answer 5 out of 7)
$$= 5 \times 5 = 25$$

Total = 35 marks

Question paper pattern for end semester practical examinations

Total = 35 marks

8. ACADEMIC PROGRESSION:

Academic progression rules are applicable as follows:

A student shall be eligible to carry forward all the courses of I, II, and III semesters till the IV semester examinations. However, he/she shall not be eligible to attend the courses of V semester until all the courses of I and II semesters are successfully completed.

A student shall be eligible to carry forward all the courses of III, IV and V semesters till the VI semester examinations. However, he/she shall not be eligible to attend the courses of VII semester until all the courses of I, II, III and IV semesters are successfully completed.

A student shall be eligible to carry forward all the courses of V, VI and VII semesters till the VIII semester examinations. However, he/she shall not be eligible to get the course completion certificate until all the courses of I, II, III, IV, V and VI semesters

are successfully completed.

A student shall be eligible to get his/her CGPA upon successful completion of the courses of I to VIII semesters within the stipulated time period as per the norms specified in 10.

A lateral entry student shall be eligible to carry forward all the courses of III, IV and V semesters till the VI semester examinations. However, he/she shall not be eligible to attend the courses of VII semester until all the courses of III and IV semesters are successfully completed.

A lateral entry student shall be eligible to carry forward all the courses of V, VI and VII semesters till the VIII semester examinations. However, he/she shall not be eligible to get the course completion certificate until all the courses of III, IV, V and VI semesters are successfully completed.

A lateral entry student shall be eligible to get his/her CGPA upon successful completion of the courses of III to VIII semesters within the stipulated time period as per the norms specified in 10.

Any student who has given more than 4 chances for successful completion of I / III semester courses and more than 3 chances for successful completion of II / IV semester courses shall be permitted to attend V / VII semester classes ONLY during the subsequent academic year as the case may be. In simpler terms there shall NOT be any ODD BATCH for any semester.

Note: Grade AB should be considered as failed and treated as one head for deciding academic progression. Such rules are also applicable for those students who fail to register for examination(s) of any course in any semester.

9. GRADING OF PERFORMANCES

9.1. Letter grades and grade points allocations:

Based on the performances, each student shall be awarded a final letter grade at the end of the semester for each course. The letter grades and their corresponding grade

points are given in Table -4.

Table – 4: Letter grades and grade points equivalent to Percentage of marks and performances

Percentage of	Letter Grade	Grade Point	Performance
Marks Obtained			
90.00 – 100	0	10	Outstanding
80.00 - 89.99	A	9	Excellent
70.00 – 79.99	В	8	Good
60.00 - 69.99	C	7	Fair
50.00 - 59.99	D	6	Average
Less than 50	F	0	Fail
Absent	AB	0	Fail

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

9.2. The Semester grade point average (SGPA)

The performance of a student in a semester is indicated by a number called 'Semester Grade Point Average' (SGPA). The SGPA is the weighted average of the grade points obtained in all the courses by the student during the semester. For example, if a student takes five courses(Theory/Practical) in a semester with credits C1, C2, C3, C4 and C5 and the student's grade points in these courses are G1, G2, G3, G4 and G5, respectively, and then students' SGPA is equal to:

$$C_{1}G_{1} + C_{2}G_{2} + C_{3}G_{3} + C_{4}G_{4} + C_{5}G_{5}$$

$$SGPA = C_{1} + C_{2} + C_{3} + C_{4} + C_{5}$$

The SGPA is calculated to two decimal points. It should be noted that, the SGPA for any semester shall take into consideration the F and AB grade awarded in that semester. For example if a learner has an F or AB grade in course 4, the SGPA shall then be computed as:

$$C_{1}G_{1} + C_{2}G_{2} + C_{3}G_{3} + ZERO + C_{5}G_{5}$$

$$SGPA = C_{1} + C_{2} + C_{3} + C_{4} + C_{5}$$

9.3. Cumulative Grade Point Average (CGPA)

The CGPA is calculated with the SGPA of all the VIII semesters to two decimal points and is indicated in final grade report card/final transcript showing the grades of all VIII semesters and their courses. The CGPA shall reflect the failed status in case of F grade(s), till the course(s) is/are passed. When the course(s) is/are passed by obtaining a pass grade on subsequent examination(s) the CGPA shall only reflect the new grade and not the fail grades earned earlier. The CGPA is calculated as:

$$C_{1}S_{1} + C_{2}S_{2} + C_{3}S_{3} + C_{4}S_{4} + C_{5}S_{5} + C_{6}S_{6} + C_{7}S_{7} + C_{8}S_{8}$$

$$CGPA = C_{1} + C_{2} + C_{3} + C_{4} + C_{5} + C_{6} + C_{7} + C_{8}$$

where C_1 , C_2 , C_3 ,... is the total number of credits for semester I,II,III,... and S_1 , S_2 , S_3 ,... is the SGPA of semester I,II,III,....

9.4. Declaration of class

Although the GPA system is a stand-alone system of grading not amenable to facile conversion to percent marks, in general the conversion of CGPA to percent marks is: $CGPA \times 9.5 = Percent marks$.

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

First Class with Distinction = CGPA of. 7.37 and above

First Class = CGPA of 6.32 to 7.36

Second Class = CGPA of 6.00 to 6.31

9.5. Project work

The internal and external examiner appointed by the college but approved by the Board of Studies for Pharmacy shall evaluate the project at the time of the Practical examinations of other semester(s). Students shall be evaluated in groups for four hours (i.e., about half an hour for a group of five students). The projects shall be evaluated as per the criteria given below.

Evaluation of Dissertation Book:

Objective(s) of the work done	15 Marks
Methodology adopted	20 Marks
Results and Discussions	20 Marks
Conclusions and Outcomes	20 Marks
TOTAL	75 Marks

Evaluation of Presentation:

Presentation of work	25 Marks
Communication skills	20 Marks
Question and answer skills	30 Marks
TOTAL	75 Marks

Explanation: The 75 marks assigned to the dissertation book shall be same for all the students in a group. However, the 75 marks assigned for presentation shall be awarded based on the performance of individual students in the given criteria

9.6. Industrial training (Desirable)

Every candidate shall be required to work for at least 150 hours spread over four weeks in a Pharmaceutical Industry/Hospital. It includes Production unit, Quality Control department, Quality Assurance department, Analytical laboratory, Chemical manufacturing unit, Pharmaceutical R&D, Hospital (Clinical Pharmacy), Clinical Research Organization, Community Pharmacy, etc. After the Semester – VI and before the commencement of Semester – VII, and shall submit satisfactory report of such work and certificate duly signed by the authority of training organization to the head of the institute.

9.7. Practice School

In the VII semester, every candidate shall undergo practice school for a period of 150 hours evenly distributed throughout the semester. The student shall opt any one of the domains for practice school declared by the program committee from time to time.

At the end of the practice school, every student shall submit a printed report (in triplicate) on the practice school he/she attended (not more than 25 pages). Along with the exams of semester VII, the report submitted by the student, knowledge and skills acquired by the student through practice school shall be evaluated by the subject experts at college level and grade point shall be awarded. The credits assigned for Practice School is 6 credits.

9.8. Award of degree

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

10. Duration for completion of the program of study

The duration for the completion of the program shall be fixed as per the norms of the University of Mumbai

11. Re-admission after break of study

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

12. Program Committee

- 1. The B. Pharm. program shall have a Program Committee constituted by the Head of the institution in consultation with all the Heads of the departments.
- 2. The composition of the Program Committee shall be as follows: A senior teacher shall be the Chairperson; One Teacher from each department handling B.Pharm courses; and four student representatives of the program (one from each academic year), nominated by the Head of the institution.
- 3. Duties of the Program Committee:
- i. Periodically reviewing the progress of the classes.
- ii. Discussing the problems concerning curriculum, syllabus and the conduct of classes.
- iii. Discussing with the course teachers on the nature and scope of assessment for the course and the same shall be announced to the students at the beginning of respective semesters.
- iv. Communicating its recommendation to the Head of the institution on academic matters.
- v. The Program Committee shall meet at least thrice in a semester preferably at the end of each Sessional exam (Internal Assessment) and before the end semester exam.

SCHEME AND SYLLABUS for

CHOICE BASED CREDIT SYSTEM

for

Undergraduate Programme (Bachelor of Pharmacy, B. Pharm.)

in

PHARMACY

Revised Course (Revised 2019) from the academic year 2019-2020

COURSE OFSTUDY

The course of study for B. Pharm shall include Semester Wise Theory & Practical as given in Table - I to VIII. The number of hours to be devoted to each theory, tutorial and practical course in any semester shall not be less than that shown in Table - I to VIII.

Table-I: Course of study for semester I

Table-1. Course of study for semester 1					
Course code	Name of the course	No.of hours	Tuto rial	Credit points	
BP101T	Human Anatomy and Physiology I— Theory	3	1	4	
BP102T	Pharmaceutical Analysis I – Theory	3	1	4	
BP103T	Pharmaceutics I – Theory	3	1	4	
BP104T	Pharmaceutical Inorganic Chemistry – Theory	3	1	4	
BP105T	Communication skills – Theory *	2	-	2	
BP106RBT BP106RMT	Remedial Biology/ Remedial Mathematics – Theory*	2	-	2	
BP107P	Human Anatomy and Physiology – Practical	4	-	2	
BP108P	Pharmaceutical Analysis I – Practical	4	-	2	
BP109P	Pharmaceutics I – Practical	4	-	2	
BP110P	Pharmaceutical Inorganic Chemistry – Practical	4	-	2	
BP111P	Communication skills – Practical*	2	-	1	
BP112RBP	Remedial Biology – Practical*	2	-	1	
	Total	32/34\$/36#	4	27/29 ^{\$} /30 [#]	

[#]Applicable ONLY for the students who have studied Mathematics / Physics / Chemistry at HSC and appearing for Remedial Biology (RB)course.

^{\$}Applicable ONLY for the students who have studied Physics / Chemistry / Botany / Zoology at HSC and appearing for Remedial Mathematics (RM)course.

^{*} Non University Examination (NUE)

Table-II: Course of study for semester II

Course Code	Name of the course	No.of hours	Tutorial	Credit points
BP201T	Human Anatomy and Physiology II – Theory	3	1	4
BP202T	Pharmaceutical Organic Chemistry I – Theory	3	1	4
BP203T	Biochemistry – Theory	3	1	4
BP204T	Pathophysiology – Theory	3	1	4
BP205T	Computer Applications in Pharmacy – Theory *	3	-	3
BP206T	Environmental sciences – Theory *	3	-	3
BP207P	Human Anatomy and Physiology II –Practical	4	-	2
BP208P	Pharmaceutical Organic Chemistry I– Practical	4	-	2
BP209P	Biochemistry – Practical		-	2
BP210P	Computer Applications in Pharmacy – Practical*	2	-	1
	Total	32	4	29

^{*}Non University Examination (NUE)

Table-III: Course of study for semester III

Course	Name of the course		Tutorial	Credit
code		hours		points
BP301T	Pharmaceutical Organic Chemistry II – Theory	3	1	4
BP302T	Physical Pharmaceutics I – Theory	3	1	4
BP303T	Pharmaceutical Microbiology – Theory	3	1	4
BP304T	Pharmaceutical Engineering – Theory	3	1	4
BP305P	Pharmaceutical Organic Chemistry II – Practical	4	-	2
BP306P	Physical Pharmaceutics I – Practical	4	-	2
BP307P	Pharmaceutical Microbiology – Practical	4	-	2
BP 308P	Pharmaceutical Engineering –Practical	4	-	2
	Total	28	4	24

Table-IV: Course of study for semester IV

Course code	Name of the course	No.of hours	Tutorial	Credit points
BP401T	Pharmaceutical Organic Chemistry III– Theory	3	1	4
BP402T	Medicinal Chemistry I – Theory	3	1	4
BP403T	Physical Pharmaceutics II – Theory	3	1	4
BP404T	Pharmacology I – Theory	3	1	4
BP405T	Pharmacognosy and Phytochemistry I- Theory	3	1	4
BP406P	Medicinal Chemistry I – Practical	4	-	2
BP407P	Physical Pharmaceutics II – Practical	4		2
BP408P	Pharmacology I – Practical		-	2
BP409P	Pharmacognosy and Phytochemistry I – Practical	4	-	2
	Total	31	5	28

Table-V: Course of study for semester \boldsymbol{V}

Course code	Name of the course	No.of hours	Tutorial	Credit points
BP501T	Medicinal Chemistry II – Theory	3	1	4
BP502T	Industrial PharmacyI– Theory	3	1	4
BP503T	Pharmacology II – Theory	3	1	4
BP504T	Pharmacognosy and Phytochemistry II- Theory	3	1	4
BP505T	Pharmaceutical Jurisprudence – Theory	3	1	4
BP506P	Industrial PharmacyI – Practical	4	-	2
BP507P	Pharmacology II – Practical	4	-	2
BP508P	Pharmacognosy and Phytochemistry II –		-	2
	Practical			
	Total	27	5	26

Table-VI: Course of study for semester VI

Course code	Name of the course	No.of hours	Tutorial	Credit points
BP601T	Medicinal Chemistry III – Theory	3	1	4
BP602T	Pharmacology III – Theory	3	1	4
BP603T	Herbal Drug Technology – Theory	3	1	4
BP604T	Biopharmaceutics and Pharmacokinetics – Theory	3	1	4
BP605T	Pharmaceutical Biotechnology – Theory	3	1	4
BP606T	Quality Assurance –Theory	3	1	4
BP607P	Medicinal chemistry III – Practical	4	-	2
BP608P	Pharmacology III – Practical		-	2
BP609P	Herbal Drug Technology – Practical	4	-	2
	Total	30	6	30

Table-VII: Course of study for semester VII

Course code	Name of the course	No.of hours	Tutorial	Credit points
	T			
BP701T	Instrumental Methods of Analysis – Theory	3	1	4
BP702T	Industrial PharmacyII – Theory	3	1	4
BP703T	Pharmacy Practice – Theory	3	1	4
BP704T	Novel Drug Delivery System – Theory	3	1	4
BP705P	Instrumental Methods of Analysis –Practical	4	-	2
BP706PS	Practice School*	12	-	6
	Total	28	5	24

^{*} Non University Examination (NUE)

Table-VIII: Course of study for semester VIII

Course code	Name of the course	No.of hours	Tutorial	Credit points
BP801T	Biostatistics and Research Methodology	3	1	4
BP802T	Social and Preventive Pharmacy	3	1	4
BP803ET	Pharma Marketing Management			
BP804ET	Pharmaceutical Regulatory Science			
BP805ET	Pharmacovigilance			
BP806ET	Quality Control and Standardization of		1 + 1 = 2	
DF0U0E1	Herbals	3 + 3 =		4 + 4 =
BP807ET	Computer Aided Drug Design	6		8
BP808ET	Cell and Molecular Biology			
BP809ET	Cosmetic Science			
BP810ET	Experimental Pharmacology			
BP811ET	Advanced Instrumentation Techniques			
BP812ET	Dietary Supplements And			
DIGIZEI	Nutraceuticals - Theory			
BP813ET	Pharmaceutical Product Development -			
DIOISEI	Theory			
BP814PW	Project Work	12	-	6
	Total	24	4	22

Table-IX: Semester wise credits distribution

Semester	Credit Points
I	27/29 ^{\$} /30 [#]
II	29
III	26
IV	28
V	26
VI	26
VII	24
VIII	22
Extracurricular/ Co curricular activities	01*
Total credit points for the program	209/211\$/212#

^{*} The credit points assigned for extracurricular and or co-curricular activities shall be given by the Principals of the colleges and the same shall be submitted to the University. The criteria to acquire this credit point shall be defined by the colleges from time to time.

^{\$}Applicable ONLY for the students studied Physics / Chemistry / Botany / Zoology at HSC and appearing for Remedial Mathematics course.

*Applicable ONLY for the students studied Mathematics / Physics / Chemistry at HSC and appearing for
Remedial Biology course.
7

Schemes for internal assessments and end semester examinations semester wise

Semester I

Course			End Semes	End Semester Exams				
code	Name of the course	Continuous Sessional Exams			Total	Marks	Duration	Total Marks
		Mode	Marks	Duration	10141	17141115	Durunon	
BP101T	Human Anatomy and Physiology I– Theory	10	15	1 Hr	25	75	3 Hrs	100
BP102T	Pharmaceutical Analysis I – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP103T	Pharmaceutics I – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP104T	Pharmaceutical Inorganic Chemistry – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP105T	Communication skills – Theory *	5	10	1 Hr	15	35	1.5 Hrs	50
BP106RBT BP106RMT	Remedial Biology/ Mathematics – Theory*	5	10	1 Hr	15	35	1.5 Hrs	50
BP107P	Human Anatomy and Physiology – Practical	5	10	4 Hrs	15	35	4 Hrs	50
BP108P	Pharmaceutical Analysis I – Practical	5	10	4 Hrs	15	35	4 Hrs	50
BP109P	Pharmaceutics I – Practical	5	10	4 Hrs	15	35	4 Hrs	50
BP110P	Pharmaceutical Inorganic Chemistry – Practical	5	10	4 Hrs	15	35	4 Hrs	50
BP111P	Communication skills – Practical*	5	5	2 Hrs	10	15	2 Hrs	25
BP112RBP	Remedial Biology – Practical*	5	5	2 Hrs	10	15	2 Hrs	25
Total		70/75\$/80#	115/125\$/130#	23/24 ^{\$} /26 [#] H	185/200\$/210#	490/525 ^{\$} / 540 [#]	31.5/33 ^{\$} / 35 [#] Hrs	675/725 ^{\$} / 750 [#]

^{*}Applicable ONLY for the students studied Mathematics / Physics / Chemistry at HSC and appearing for Remedial Biology (RB)course.
\$Applicable ONLY for the students studied Physics / Chemistry / Botany / Zoology at HSC and appearing for Remedial Mathematics (RM) course.

^{*} Non-University Examination (NUE)

Semester II

Course		Internal Assessment				End Seme	Total	
code	Name of the course	Continuous Sessional Exams			Total	Marks	Duration	Marks
couc		Mode	Marks	Duration	Total	With KS	Duration	1,141,119
BP201T	Human Anatomy and Physiology II – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP202T	Pharmaceutical Organic Chemistry I – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP203T	Biochemistry – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP204T	Pathophysiology – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP205T	Computer Applications in Pharmacy – Theory	10	15	1 Hr	25	50	2 Hrs	75
BP206T	Environmental sciences – Theory	10	15	1 Hr	25	50	2 Hrs	75
BP207P	Human Anatomy and Physiology II –Practical	5	10	4 Hrs	15	35	4 Hrs	50
BP208P	Pharmaceutical Organic Chemistry I— Practical	5	10	4 Hrs	15	35	4 Hrs	50
BP209P	Biochemistry – Practical	5	10	4 Hrs	15	35	4 Hrs	50
BP210P	Computer Applications in Pharmacy – Practical	5	5	2 Hrs	10	15	2 Hrs	25
	Total	80	125	20 Hrs	205	520	30 Hrs	725

Semester III

Course			Internal Assessment				End Semester Exams		
code	Name of the course	Continuous	Continuous Sessional Exams			Marks	Duration	Total Marks	
code		Mode	Marks	Duration	Total	Watks	Duration	TVICEI INS	
BP301T	Pharmaceutical Organic	10	15	1 Hr	25	75	3 Hrs	100	
DI 3011	Chemistry II – Theory	10	13	1 111	23	13	31118	100	
BP302T	PhysicalPharmaceuticsI –Theory	10	15	1 Hr	25	75	3 Hrs	100	
BP303T	Pharmaceutical Microbiology –	10	15	1 Hr	25	75	3 Hrs	100	
DI 3031	Theory	10	15	1 Hr	25	/3	3 Hrs	100	
BP304T	Pharmaceutical Engineering –	10	15	1 Hr	25	75	3 Hrs	100	
DI 3041	Theory	10	13	1 П	23	13	3 1118	100	
BP305P	Pharmaceutical Organic	5	10	4 Hr	15	35	4 Hrs	50	
DF 303F	Chemistry II – Practical	3	10	4 nr	13	33	4 HIS	30	
BP306P	Physical Pharmaceutics I –	5	10	4 Hr	15	35	4 Hrs	50	
DI 3001	Practical	3	10	4111	13	33	71115	30	
BP307P	Pharmaceutical Microbiology –	5	10	4 Hr	15	35	4 Hrs	50	
D 1 3071	Practical	3	10	7111	13	33	71115	30	
BP308P	Pharmaceutical Engineering –	5	10	4 Hr	15	35	4 Hrs	50	
וטטכ ום	Practical	3	10	4 111	13	33	4 1115	30	
	Total	60	100	20	160	440	28Hrs	600	

Semester IV

Course			Internal As	End Seme	Total			
code	Name of the course	Continuous Sessional Exams		Total	Marks	Duration	Marks	
Couc		Mode	Marks	Duration	Total	IVIAI KS	Duration	Marks
BP401T	Pharmaceutical Organic Chemistry III– Theory	10	15	1 Hr	25	75	3 Hrs	100
BP402T	Medicinal Chemistry I – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP403T	Physical Pharmaceutics II – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP404T	Pharmacology I – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP405T	Pharmacognosy I – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP406P	Medicinal Chemistry I – Practical	5	10	4 Hr	15	35	4 Hrs	50
BP407P	Physical Pharmaceutics II – Practical	5	10	4 Hrs	15	35	4 Hrs	50
BP408P	Pharmacology I – Practical	5	10	4 Hrs	15	35	4 Hrs	50
BP409P	Pharmacognosy I – Practical	5	10	4 Hrs	15	35	4 Hrs	50
	Total	70	115	21 Hrs	185	515	31 Hrs	700

Semester V

Course			Internal As	sessment		End Seme	Total	
code	Name of the course	Continuous	Sessiona	al Exams	Total	Marks	Duration	Marks
Couc		Mode	Marks	Duration	Total	Marks	Duration	
BP501T	Medicinal Chemistry II – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP502T	Industrial PharmacyI— Theory	10	15	1 Hr	25	75	3 Hrs	100
BP503T	Pharmacology II – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP504T	Pharmacognosy and	10	15	1 Hr	25	75	3 Hrs	100
	Phytochemistry II – Theory							
BP505T	Pharmaceutical Jurisprudence—	10	15	1 Hr	Hr 25 75	75	3 Hrs	100
DF 303 1	Theory		13	13 1111		13		100
BP506P	Industrial PharmacyI– Practical	5	10	4 Hr	15	35	4 Hrs	50
BP507P	Pharmacology II – Practical	5	10	4 Hr	15	35	4 Hrs	50
BP508P	Pharmacognosy II – Practical	5	10	4 Hr	15	35	4 Hrs	50
	Total	65	105	17 Hr	170	480	27 Hrs	650

Semester VI

Course			Internal As	sessment		End Semester Exams		Total Marks
code	Name of the course	Continuous	ontinuous Sessional Exams		Total	Marks	Duration	
		Mode	Marks	Duration	1 Otal	Wiai Ks	Duration	14141143
BP601T	Medicinal Chemistry III – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP602T	Pharmacology III – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP603T	Herbal Drug Technology – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP604T	Biopharmaceutics and Pharmacokinetics – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP605T	Pharmaceutical Biotechnology— Theory	10	15	1 Hr	25	75	3 Hrs	100
BP606T	Pharmaceutical Quality Assurance—Theory	10	15	1 Hr	25	75	3 Hrs	100
BP607P	Medicinal chemistry III – Practical	5	10	4 Hrs	15	35	4 Hrs	50
BP608P	Pharmacology III – Practical	5	10	4 Hrs	15	35	4 Hrs	50
BP609P	Herbal Drug Technology – Practical	5	10	4 Hrs	15	35	4 Hrs	50
	Total	75	120	18 Hrs	195	555	30 Hrs	750

Semester VII

Course	Name of the course]	Internal As	End Semester Exams		Total		
code	Name of the course	Continuous	Session	al Exams	Total	Marks	Duration	Marks
		Mode	Marks	Duration	Total	Wiai KS	Duration	
BP701T	Instrumental Methods of Analysis – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP702T	Industrial Pharmacy – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP703T	Pharmacy Practice – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP704T	Novel Drug Delivery System – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP705 P	Instrumental Methods of Analysis – Practical	5	10	4 Hrs	15	35	4 Hrs	50
BP706 PS	Practice School*	25	-	-	25	125	5 Hrs	150
	Total	70	70	8Hrs	140	460	21 Hrs	600

^{*} The subject experts at college level shall conductexaminations

Semester VIII

Course			Interr	nal Assessme	nt	End Semester Exams		Total
code	Name of the course	Continuou	Session	nal Exams	Total	Marks	Duration	Marks
code		s Mode	Marks	Duration	Total	Maiks	Duration	
BP801T	Biostatistics and Research Methodology – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP802T	Social and Preventive Pharmacy – Theory	10	15	1 Hr	25	75	3 Hrs	100
BP803ET	Pharmaceutical Marketing – Theory							
BP804ET	Pharmaceutical Regulatory Science – Theory							
BP805ET	Pharmacovigilance – Theory							
BP806ET	Quality Control and Standardization of Herbals – Theory	10 + 10	15 + 15	1 + 1 =	25 + 25	75 +	3 + 3 = 6	100 +
BP807ET	Computer Aided Drug Design – Theory	= 20	= 30	2 Hrs	= 50	75 = 150	Hrs	100 = 200
BP808ET	Cell and Molecular Biology – Theory							
BP809ET	Cosmetic Science – Theory							
BP810ET	Experimental Pharmacology – Theory							
BP811ET	Advanced Instrumentation Techniques – Theory							
BP812ET	Dietary Supplements And Nutraceuticals - Theory							
BP813ET	Pharmaceutical Product Development - Theory							
BP814PW	Project Work	-	-	-	-	150	4 Hrs	15 0

Total	40	60	4 Hrs	100	450	16 Hrs	550

SEMESTER I

BP101T HUMAN ANATOMY AND PHYSIOLOGY-I (Theory)3 L + 1T / Week

Scope: This subject is designed to impart fundamental knowledge on the structure andfunctions of the various systems of the human body. It also helps in understanding both homeostatic mechanisms. The subject provides the basic knowledge required to understand the various disciplines of pharmacy.

Objectives: Upon completion of this course the student should be able to

- 1. Explain the gross morphology, structure and functions of various organs of the human body.
- 2. Describe the various homeostatic mechanisms and their imbalances.
- 3. Identify the various tissues and organs of different systems of human body.
- 4. Perform the various experiments related to special senses and nervous system.
- 5. Appreciate coordinated working pattern of different organs of each system

Sr.	Content	Hours
No.		
1	UNIT I	10
1.1	Introduction to human body	2
	Definition and scope of anatomy and physiology, levels of structural	
	organization and body systems, basic life processes, homeostasis, basic	
	anatomical terminology	
1.2	Cellular level of organization	4
	Structure and functions of cell, transport across cell membrane, cell	
	division, cell junctions. General principles of cell communication,	
	intracellular signaling pathway activation by extracellular signal molecule,	
	Forms of intracellular signaling: a) Contact-dependent b) Paracrine c)	
	Synaptic d) Endocrine	
1.3	Tissue level of organization	4
	Classification of tissues, structure, location and functions of epithelial,	
	muscular and nervous and connective tissues	
2	UNIT II	10
2.1	Integumentary system Structure and functions of skin	1
2.2	Skeletal system	6
	Divisions of skeletal system, types of bone, salient features and functions	
	of bones of axial and appendicular skeletal system.	
	Organization of skeletal muscle, physiology of muscle contraction,	
	neuromuscular junction	
2.3	Joints	3
	Structural and functional classification, types of joints movements and its	

	articulation	
3	UNIT III	10
3.1	Body fluids and blood	6
	Body fluids, composition and functions of blood, hemopoeisis, formation	
	of hemoglobin, anemia, mechanisms of coagulation, blood grouping, Rh	
	factors, transfusion, its significance and disorders of blood, Reticulo	
	endothelial system.	
3.2	Lymphatic system	4
	Lymphatic organs and tissues, lymphatic vessels, lymph circulation and	
	functions of lymphatic system	
4	UNIT IV	08
4.1	Peripheral nervous system:	3
	Classification of peripheral nervous system: Structure and functions of	
	sympathetic and parasympathetic nervous system. Origin and functions of	
	spinal and cranial nerves.	
4.2	Special senses	5
	Structure and functions of eye, ear, nose and tongue and their disorders.	
5	UNIT V	07
	Cardiovascular system	4
	Heart – anatomy of heart, blood circulation, blood vessels, structure and	
	functions of artery, vein and capillaries, elements of conduction system of	
	heart and hear beat, its regulation by autonomic nervous system, cardiac	
	output, cardiac cycle.	
	Regulation of blood pressure, pulse, electrocardiogram and disorders of	3
	heart.	
	Total	45

Recommended Books (Latest Editions)

- 1. Essentials of Medical Physiology by K. Sembulingam and P. Sembulingam. Jaypee brothers medical publishers, New Delhi.
- 2. Anatomy and Physiology in Health and Illness by Kathleen J.W. Wilson, Churchill Livingstone, New York
- 3. Physiological basis of Medical Practice-Best and Tailor. Williams & Wilkins Co,Riverview,MI USA
- 4. Text book of Medical Physiology- Arthur C, Guyton and John. E. Hall. Miamisburg, OH, U.S.A.
- 5. Principles of Anatomy and Physiology by Tortora Grabowski. Palmetto, GA, U.S.A.

- 6. Textbook of Human Histology by Inderbir Singh, Jaypee brother's medical publishers, New Delhi.
- 7. Textbook of Practical Physiology by C.L. Ghai, Jaypee brother's medical publishers, New Delhi.
- 8. Practical workbook of Human Physiology by K. Srinageswari and Rajeev Sharma, Jaypee brother's medical publishers, New Delhi.

Reference Books (Latest Editions)

- 1. Physiological basis of Medical Practice-Best and Tailor. Williams & Wilkins Co, Riverview, MI USA
- 2. Text book of Medical Physiology- Arthur C, Guyton and John. E. Hall. Miamisburg, OH, U.S.A.
- 3. Human Physiology (vol 1 and 2) by Dr. C.C. Chatterrje ,Academic Publishers Kolkata

BP102T PHARMACEUTICAL ANALYSIS (Theory)3 L + 1T / Week

Scope: This course deals with the fundamentals of analytical chemistry and principles of electrochemical analysis of drugs

Objectives: Upon completion of the course student shall be able to

- 1. understand the principles of volumetric and electro chemical analysis
- 2. carryout various volumetric and electrochemical titrations
- 3. develop analytical skills

Sr. No.	Content	Hours
1	UNIT I	10
	Pharmaceutical analysis- Definition and scope	6
	i) Different techniques of analysis	
1.1	ii) Methods of expressing concentration	
	iii) Primary and secondary standards.	
	iv) Preparation and standardization of various molar and normal	
	solutions-Oxalic acid, sodium hydroxide, hydrochloric acid, sodium	
	thiosulphate, sulphuric acid, potassium permanganate and ceric	
	ammonium sulphate	
1.2	Errors: Sources of errors, types of errors, methods of minimizing	2
	errors,accuracy, precision and significant figures	
1.3	Pharmacopoeia, Sources of impurities in medicinal agents, limit tests.	2
2	UNIT II	10
2.1	Acid base titration: Theories of acid base indicators, classification	
	ofacid base titrations and theory involved in titrations of strong, weak,	5
	and very weak acids and bases, neutralization curve	
2.2	Non aqueous titration: Solvents, acidimetry and alkalimetry titration	5
	andestimation of Sodium benzoate and Ephedrine HCl	
3	UNIT III	10
3.1	Precipitation titrations: Mohr's method, Volhard's, Modified	2
3.2	Complexometric titration: Classification, metal ion indicators,	4
	maskingand demasking reagents, estimation of Magnesium sulphate,	
	and calcium gluconate.	
3.3	Gravimetry: Principle and steps involved in gravimetric analysis.	2
	Purity of the precipitate: co-precipitation and post precipitation,	
	Estimation of barium sulphate	
3.4	Basic Principles,methods and application of diazotisation titration	2

4	UNIT IV	08
4.1	Redox titrations	
	Concepts of oxidation and reduction	
4.2	Types of redox titrations (Principles and applications)	
	Cerimetry, Iodimetry, Iodometry, Bromatometry, Dichrometry,	
	Titration with potassium iodate	
5	UNIT V - Electrochemical methods of analysis	07
5.1	Conductometry- Introduction, Conductivity cell,	2
	Conductometric titrations, applications.	
5.2	Potentiometry - Electrochemical cell, construction and workingof	3
	reference (Standard hydrogen, silver chloride electrode and calomel	
	electrode) and indicator electrodes (metal electrodes and glass	
	electrode), methods to determine end point of potentiometric titration	
	and applications.	
5.3	Polarography - Principle, Ilkovic equation, construction and working of	2
	dropping mercury electrode and rotating platinum electrode,	
	applications	
	Total	45

Recommended Books: (Latest Editions)

- 1. A.H. Beckett & J.B. Stenlake's, Practical Pharmaceutical Chemistry Vol I & II, Stahlone Press of University of London
- 2. A.I. Vogel, Text Book of Quantitative Inorganic analysis
- 3. P. Gundu Rao, Inorganic Pharmaceutical Chemistry
- 4. Bentley and Driver's Textbook of Pharmaceutical Chemistry
- 5. John H. Kennedy, Analytical chemistry principles
- 6. Indian Pharmacopoeia.

BP103T PHARMACEUTICS- I (Theory)3 L + 1T / Week

Scope: This course is designed to impart a fundamental knowledge on the preparatorypharmacy with arts and science of preparing the different conventional dosage forms.

Objectives: Upon completion of this course the student should be able to:

- 1. Know the history of profession of pharmacy
- 2. Understand the basics of different dosage forms, pharmaceutical incompatibilities and pharmaceutical calculations
- 3. Understand the professional way of handling the prescription
- 4. Preparation of various conventional dosage forms

Sr.	Content	Hours
No.		
1	UNIT I	10
1.1	Historical background and development of profession of pharmacy:	3
	Historyof profession of Pharmacy in India in relation to pharmacy	
	education, industry and organization, Pharmacy as a career,	
	Pharmacopoeias: Introduction to IP, BP, USP and Extra Pharmacopoeia.	
1.2	Dosage forms: Introduction to dosage forms, classification and definitions	3
1.3	Prescription: Definition, Parts of prescription, handling of	2
	Prescription and Errors in prescription.	
1.4	Posology: Definition, Factors affecting posology. Pediatric dose	2
	calculationsbased on age, body weight and body surface area.	
2	UNIT II	10
2.1	Pharmaceutical calculations: Weights and measures–Imperial &	4
	Metricsystem, Calculations involving percentage solutions, alligation, proof	
	spirit and isotonic solutions based on freezing point and molecular weight.	
2.2	Powders: Definition, classification, advantages and disadvantages, Simple	3
	&compound powders – official preparations, dusting powders, effervescent,	
	efflorescent and hygroscopic powders, eutectic mixtures. Geometric	
	dilutions.	
2.3	Liquid dosage forms: Advantages and disadvantages of liquid dosage	3
	forms. Excipients used in formulation of liquid dosage forms. Solubility	
	enhancement techniques	
	UNIT III	10
3.1	Monophasic liquids: Definitions and preparations of Gargles,	3
	Mouthwashes, Throat Paint, Eardrops, Nasal drops, Enemas, Syrups, Elixirs,	

	Liniments and Lotions.	
3.2	Biphasic liquids:	7
	o Suspensions: Definition, advantages and disadvantages,	
	classifications,Preparation of suspensions; Flocculated and	
	Deflocculated suspension & stability problems and methods to	
	overcome.	
	o Emulsions: Definition, classification, emulsifying agent, test for the	
	identification of type of Emulsion, Methods of preparation & stability	
	problems and methods to overcome.	
4	UNIT IV	08
4.1	Suppositories: Definition, types, advantages and disadvantages, types of	5
	bases,methods of preparations. Displacement value & its calculations,	
	evaluation of suppositories.	
4.2	Pharmaceutical incompatibilities: Definition, classification, physical,	3
	chemicaland therapeutic incompatibilities with examples.	
5	UNIT V	07
	Semisolid dosage forms: Definitions, classification, mechanisms and	
	factorsinfluencing dermal penetration of drugs. Preparation of ointments,	
	pastes, creams and gels. Excipients used in semi solid dosage forms.	
	Evaluation of semi solid dosages forms	
	Total	45

Recommended Books (Latest edition)

- 1. H. C. Ansel et al., Pharmaceutical Dosage Form and Drug Delivery System, Lippincott Williams and Walkins, New Delhi.
- 2. Carter S.J., Cooper and Gunn's-Dispensing for Pharmaceutical Students, CBS publishers, New Delhi.
- 3. M.E. Aulton, Pharmaceutics, The Science& Dosage Form Design, Churchill Livingstone, Edinburgh.
- 4. Indian pharmacopoeia.
- 5. British pharmacopoeia.
- 6. Lachmann. Theory and Practice of Industrial Pharmacy, Lea& Febiger Publisher, The University of Michigan.
- 7. Alfonso R. Gennaro Remington. The Science and Practice of Pharmacy, Lippincott Williams, New Delhi.
- 8. Carter S.J., Cooper and Gunn's. Tutorial Pharmacy, CBS Publications, New Delhi.
- 9. E.A. Rawlins, Bentley's Text Book of Pharmaceutics, English Language Book Society, Elsevier Health Sciences, USA.
- 10. Isaac Ghebre Sellassie: Pharmaceutical Pelletization Technology, Marcel Dekker, INC, New York.

- 11. Dilip M. Parikh: Handbook of Pharmaceutical Granulation Technology, Marcel Dekker, INC, New York.
- 12. Francoise Nieloud and Gilberte Marti-Mestres: Pharmaceutical Emulsions and Suspensions, Marcel Dekker, INC, New York.

BP104T PHARMACEUTICAL INORGANIC CHEMISTRY (Theory)3 L + 1T / Week

Scope: This subject deals with the monographs of inorganic drugs and pharmaceuticals.

Objectives: Upon completion of course student shall be able to

- 1. know the sources of impurities and methods to determine the impurities in inorganic drugs and pharmaceuticals
- 2. understand the medicinal and pharmaceutical importance of inorganic compounds

Sr. No.	Content	Hours
1	UNIT I	10
1.1	Impurities in pharmaceutical substances: History of Pharmacopoeia, Sources and types of impurities, principle involved in	4
	the limit test for Chloride, Sulphate, Iron, Arsenic, Lead and Heavy metals, modified limit test for Chloride and Sulphate	
1.2	General methods of preparation, assay for the compounds	6
1.2	superscripted with asterisk (*), properties and medicinal uses of	
	inorganic compounds belonging to the following classes	
2	UNIT II	10
2.1	Acids, Bases and Buffers: Buffer equations and buffer capacity in	4
	general, buffers in pharmaceutical systems, preparation, stability,	
	buffered isotonic solutions, measurements of tonicity, calculations and	
	methods of adjusting isotonicity.	
2.2	Major extra and intracellular electrolytes: Functions of	4
	major □ physiological ions, Electrolytes used in the replacement	
	therapy: Sodium chloride*, Potassium chloride, Calcium gluconate*	
	and Oral Rehydration Salt (ORS), Physiological acid base balance.	
2.3	Dental products : Dentifrices, role of fluoride in the treatment of	2
	dentalcaries, Desensitizing agents, Calcium carbonate, Sodium	
	fluoride, and Zinc eugenol cement.	
3	UNIT III - Gastrointestinal agents	10
3.1	Acidifiers: Ammonium chloride* and Dil. HCl	1
3.2	Antacid: Ideal properties of antacids, combinations of antacids,	3
	Sodium Bicarbonate*, Aluminum hydroxide gel, Magnesium	
	hydroxide mixture2	
3.3	Cathartics: Magnesium sulphate, Sodium orthophosphate, Kaolin and Bentonite	2

3.4	Antimicrobials: Mechanism, classification, Potassium permanganate,	4
	Boricacid, Hydrogen peroxide*, Chlorinated lime*, Iodine and its	
	preparations	
4	UNIT IV - Miscellaneous compounds	08
4.1	Expectorants: Potassium iodide, Ammonium chloride*.	1
4.2	Emetics: Copper sulphate*, Sodium potassium tartarate	2
4.3	Haematinics: Ferrous sulphate*, Ferrous gluconate	2
4.4	Poison and Antidote: Sodium thiosulphate*, Activated charcoal,	2
	Sodiumnitrite	
4.5	Astringents: Zinc Sulphate, Potash Alum	1
5	UNIT V	07
	Radiopharmaceuticals: Radio activity, Measurement of	
	radioactivity, Properties of α , β , γ radiations, Half life, radio isotopes	
	and study of radio isotopes - Sodium iodide I ¹³¹ , Storage conditions,	
	precautions & pharmaceutical application of radioactive substances.	
	Total	45

Recommended Books: (Latest Editions)

- 1. A.H. Beckett & J.B. Stenlake's, Practical Pharmaceutical Chemistry Vol I & II, Stahlone Press of University of London, 4th edition.
- 2. A.I. Vogel, Text Book of Quantitative Inorganic analysis
- 3. P. Gundu Rao, Inorganic Pharmaceutical Chemistry, 3rd Edition
- 4. M.L Schroff, Inorganic Pharmaceutical Chemistry
- 5. Bentley and Driver's Textbook of Pharmaceutical Chemistry
- 6. Anand & Chatwal, Inorganic Pharmaceutical Chemistry
- 7. Indian Pharmacopoeia

BP105T COMMUNICATION SKILLS (Theory) 2hours/week

Scope: This course will prepare the young pharmacy student to interact effectively withdoctors, nurses, dentists, physiotherapists and other health workers. At the end of this course the student will get the soft skills set to work cohesively with the team as a team player and will add value to the pharmaceutical business.

Objectives: Upon completion of the course the student shall be able to

- 1. Understand the behavioural needs for a Pharmacist to function effectively in the areas of pharmaceutical operation
- 2. Communicate effectively (Verbal and Non-Verbal)
- 3. Effectively manage the team as a team player
- 4. Develop interview skills
- 5. Develop Leadership qualities and essentials

Sr. No.	Content	Hours
1	UNIT I	7
1.1	Communication Skills: Introduction, Definition, The Importance of	3
	Communication, The Communication Process - Source, Message,	
	Encoding, Channel, Decoding, Receiver, Feedback, Context	
1.2	Barriers to communication: Physiological Barriers, Physical Barriers,	2
	CulturalBarriers, Language Barriers, Gender Barriers, Interpersonal	
	Barriers, Psychological Barriers, Emotional barriers	
1.3	Perspectives in Communication: Introduction, Visual Perception,	2
	Language, Otherfactors affecting our perspective - Past Experiences,	
	Prejudices, Feelings, Environment	
2	UNIT II	7
2.1	Elements of Communication: Introduction, Face to Face	3
	Communication - Tone of Voice, Body Language (Non-verbal	
	communication), Verbal Communication, Physical Communication	
2.2	Communication Styles: Introduction, The Communication Styles	4
	Matrix with example for each -Direct Communication Style, Spirited	
	Communication Style, Systematic Communication Style, Considerate	
	Communication Style	
3	UNIT III	7
3.1	Basic Listening Skills: Introduction, Self-Awareness, Active Listening,	2
	Becoming anActive Listener, Listening in Difficult Situations	
3.2	Effective Written Communication: Introduction, When and When Not	3

	to Use WrittenCommunication - Complexity of the Topic, Amount of	
	Discussion' Required, Shades of Meaning, Formal Communication	
3.3	Writing Effectively: Subject Lines, Put the Main Point First, Know	2
	Your Audience, Organization of the Message	
4	UNIT IV	5
4.1	Interview Skills: Purpose of an interview, Do's and Dont's of an	2
	interview	
4.2	Giving Presentations: Dealing with Fears, Planning your Presentation,	3
	Structuring YourPresentation, Delivering Your Presentation,	
	Techniques of Delivery	
5	UNIT V	4
	Group Discussion: Introduction, Communication skills in group	
	discussion, Do's andDont's of group discussion	
	Total	30

Recommended Books: (Latest Edition)

- 1. Basic communication skills for Technology, Andreja. J. Ruther Ford, 2nd Edition, Pearson Education, 2011
- 2. Communication skills, Sanjay Kumar, Pushpalata, 1stEdition, Oxford Press, 2011
- 3. Organizational Behaviour, Stephen .P. Robbins, 1st Edition, Pearson, 2013
- 4. Brilliant- Communication skills, Gill Hasson, 1st Edition, Pearson Life, 2011
- 5. The Ace of Soft Skills: Attitude, Communication and Etiquette for success, Gopala Swamy Ramesh, 5thEdition, Pearson, 2013
- 6. Developing your influencing skills, Deborah Dalley, Lois Burton, Margaret, Green hall, 1st Edition Universe of Learning LTD, 2010
- 7. Communication skills for professionals, Konar nira, 2ndEdition, New arrivals PHI, 2011
- 8. Personality development and soft skills, Barun K Mitra, 1stEdition, Oxford Press, 2011
- 9. Soft skill for everyone, Butter Field, 1st Edition, Cengage Learning india pvt.ltd, 2011
- 10. Soft skills and professional communication, Francis Peters SJ, 1stEdition, Mc Graw Hill Education, 2011
- 11. Effective communication, John Adair, 4thEdition, Pan Mac Millan, 2009
- 12. Bringing out the best in people, Aubrey Daniels, 2ndEdition, Mc Graw Hill, 1999

BP106RBT REMEDIAL BIOLOGY (Theory) 2hours/week

Scope: To learn and understand the components of living world, structure and functional system of plant and animal kingdom.

Objectives: Upon completion of the course, the student shall be able to

- 1. know the classification and salient features of five kingdoms of life
- 2. understand the basic components of anatomy & physiology of plant
- 3. know understand the basic components of anatomy & physiology animal with special reference to human

Sr. No.	Content	Hours
1	UNIT I	7
1.1	Living world:	4
	Definition and characters of living organisms	
	Diversity in the living world	
	Binomial nomenclature	
	• Five kingdoms of life and basis of classification. Salient features of	
	Monera, Potista, Fungi, Animalia and Plantae, Virus,	
1.2	Morphology of Flowering plants	3
	• Morphology of different parts of flowering plants – Root, stem,	
	inflorescence, flower, leaf, fruit, seed.	
	General Anatomy of Root, stem, leaf of monocotyledons &	
	Dicotylidones.	
2	UNIT II	7
2.1	Body fluids and circulation	4
	Composition of blood, blood groups, coagulation of blood	
	Composition and functions of lymph	
	Human circulatory system	
	Structure of human heart and blood vessels	
	Cardiac cycle, cardiac output and ECG	
	Human alimentary canal and digestive glands	
	Role of digestive enzymes	
	Digestion, absorption and assimilation of digested food	
2.2	Breathing and respiration	3
	Human respiratory system	
	Mechanism of breathing and its regulation	

	Exchange of gases, transport of gases and regulation of respiration	
	Respiratory volumes	
3	UNIT III	7
3.1	Excretory products and their elimination	2
	Modes of excretion	
	Human excretory system- structure and function	
	Urine formation	
	Rennin angiotensin system	
3.2	Neural control and coordination	2
	Definition and classification of nervous system	
	Structure of a neuron	
	Generation and conduction of nerve impulse	
	Structure of brain and spinal cord	
	Functions of cerebrum, cerebellum, hypothalamus and medulla	
	oblongata	
3.3	Chemical coordination and regulation	2
	Endocrine glands and their secretions	
	Functions of hormones secreted by endocrine glands	
3.4	Human reproduction	1
	Parts of female reproductive system	
	Parts of male reproductive system	
	Spermatogenesis and Oogenesis	
	Menstrual cycle	
4	UNIT IV	5
4.1	Plants and mineral nutrition:	3
	Essential mineral, macro and micronutrients	
	Nitrogen metabolism, Nitrogen cycle, biological nitrogen fixation	
4.2	Photosynthesis	2
	Autotrophic nutrition, photosynthesis, Photosynthetic pigments,	
	Factors affecting photosynthesis.	
5	UNIT V	4
5.1	Plant respiration: Respiration, glycolysis, fermentation (anaerobic).	1
5.2	Plant growth and development	1
	Phases and rate of plant growth, Condition of growth, Introduction to	
	plant growth regulators	
5.3	Cell - The unit of life	1
	Structure and functions of cell and cell organelles. Cell division	

5.4	Tissues	1
	Definition, types of tissues, location and functions.	
	Total	30

Text Books

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

Reference Books

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

BP106RMT REMEDIAL MATHEMATICS (Theory) 2hours/week

Scope: This is an introductory course in mathematics. This subject deals with theintroduction to Partial fraction, Logarithm, matrices and Determinant, Analytical geometry, Calculus, differential equation and Laplace transform.

Objectives:Upon completion of the course the student shall be able to:

- 1. Know the theory and their application in Pharmacy
- 2. Solve the different types of problems by applying theory
- 3. Appreciate the important application of mathematics in Pharmacy

Sr.	Content	Hours
No.		
1	UNIT I	6
1.1	Partial fraction	
	Introduction, Polynomial, Rational fractions, Proper and Improper	
	fractions, Partial fraction, Resolving into Partial fraction, Application of	
	Partial Fraction in Chemical Kinetics and Pharmacokinetics	
1.2	Logarithms	
	Introduction, Definition, Theorems/Properties of logarithms, Common	
	logarithms, Characteristic and Mantissa, worked examples, application of	
	logarithm to solve pharmaceutical problems.	
1.3	Function:	
	Real Valued function, Classification of real valued functions	
1.4	Limits and continuity :	
	Introduction, Limit of a function, Definition of limit of a function (∈ - δ	
	definition), $\lim_{x\to a} \frac{x^n - a^n}{x - a} = na^{n-1}$, $\lim_{\theta \to 0} \frac{\sin \theta}{\theta} = 1$,	
	definition), $\lim_{x \to a} \frac{1}{x - a} = na^{n-1}$, $\lim_{a \to 0} \frac{1}{a} = 1$,	
	x - u	
2	UNIT II	6
_	Matrices and Determinant:	
	Introduction matrices, Types of matrices, Operation on matrices,	
	Transpose of a matrix, Matrix Multiplication, Determinants, Properties of	
	determinants, Product of determinants, Minors and co-Factors, Adjoint or	
	adjugate of a square matrix, Singular and non-singular matrices,	
	Inverse of a matrix, Solution of system of linear of equations using matrix	
	method, Cramer's rule, Characteristic equation and roots of a square	
	matrix, Cayley–Hamilton theorem, Application of Matrices in solving	
L		

Pharmacokinetic equations	
3 UNIT III	6
Calculus	
Differentiation: Introductions, Derivative of a function, Derivative o	f a
constant, Derivative of a product of a constant and a function, Derivat	ive
of the sum or difference of two functions, Derivative of the product of tw	
functions (product formula), Derivative of the quotient of two functions	
(Quotient formula) – Without Proof, Derivative of xn w.r.tx, where n is a	-
rational number, Derivative of ex,, Derivative of loge x, Derivat	
of ax,Derivative of trigonometric functions from first principles (with	
Proof), Successive Differentiation, Conditions for a function to be	a
maximum or a minimum at a point. Application	
4 UNIT IV - Analytical Geometry	6
4.1 Introduction: Signs of the Coordinates, Distance formula,	
4.2 Straight Line : Slope or gradient of a straight line, Condition	ons
forparallelism and perpendicularity of two lines, Slope of a line joining t	wo
points, Slope – intercept form of a straight line	
4.3 Integration: Introduction, Definition, Standard formulae, Rules of	
integration, Method of substitution, Method of Partial fractions,	
Integration by parts, definite integrals, application	
5 UNIT V	6
5.1 Differential Equations : Some basic definitions, Order a	and
degree, Equations in separable form, Homogeneous equations, Lin	ear
Differential equations, Exact equations, Application	in
solvingPharmacokinetic equations	
5.2 Laplace Transform : Introduction, Definition, Properties	of
Laplacetransform, Laplace Transforms of elementary functions, Inve	
Laplace transforms, Laplace transform of derivatives, Application to so	
Linear differential equations, Application in solving Chemicalkinet	ics
and Pharmacokinetics equations	
Total	30

Recommended Books (Latest Edition)

- 1. Differential Calculus by Shanthinarayan
- 2. Pharmaceutical Mathematics with application to Pharmacy by Panchaksharappa Gowda D.H.
- 3. Integral Calculus by Shanthinarayan
- 4. Higher Engineering Mathematics by Dr.B.S.Grewal

BP107P HUMAN ANATOMY AND PHYSIOLOGY (Practical) 4 Hours/week

- 1. Study of compound microscope.
- 2. Microscopic study of epithelial and connective tissue
- 3. Microscopic study of muscular and nervous tissue
- 4. Identification of axial bones
- 5. Identification of appendicular bones
- 6. Introduction to hemocytometry.
- 7. Enumeration of white blood cell (WBC) count
- 8. Enumeration of total red blood corpuscles (RBC) count
- 9. Determination of bleeding time
- 10. Determination of clotting time
- 11. Estimation of hemoglobin content
- 12. Determination of blood group.
- 13. Determination of erythrocyte sedimentation rate (ESR).
- 14. Determination of heart rate and pulse rate.
- 15. Recording of blood pressure.

Recommended Books (Latest Editions)

- 1. Essentials of Medical Physiology by K. Sembulingam and P. Sembulingam. Jaypee brothers medical publishers, New Delhi.
- 2. Anatomy and Physiology in Health and Illness by Kathleen J.W. Wilson, Churchill Livingstone, New York
- 3. Physiological basis of Medical Practice-Best and Tailor. Williams & Wilkins Co,Riverview,MI USA
- 4. Text book of Medical Physiology- Arthur C,Guyton andJohn.E. Hall. Miamisburg, OH, U.S.A.
- 5. Principles of Anatomy and Physiology by Tortora Grabowski. Palmetto, GA, U.S.A
- 6. Textbook of Human Histology by Inderbir Singh, Jaypee brother's medical publishers, New Delhi.
- 7. Textbook of Practical Physiology by C.L. Ghai, Jaypee brother's medical publishers, New Delhi.
- 8. Practical workbook of Human Physiology by K. Srinageswari and Rajeev Sharma, Jaypee brother's medical publishers, New Delhi.

Reference Books (Latest Editions)

- Physiological basis of Medical Practice-Best and Tailor. Williams & Wilkins Co, Riverview, MI USA
- 2. Text book of Medical Physiology- Arthur C, Guyton and John. E. Hall. Miamisburg, OH, U.S.A.
- 3. Human Physiology (vol 1 and 2) by Dr. C.C. Chatterrje, Academic Publishers Kolkata.

BP108P PHARMACEUTICAL ANALYSIS (Practical)

4 Hours / Week

I Limit Test of the following

- (1) Chloride
- (2) Sulphate
- (3) Iron
- (4) Arsenic

II Preparation and standardization of

- (1) Sodium hydroxide
- (2) Sulphuric acid
- (3) Sodium thiosulfate
- (4) Potassium permanganate
- (5) Ceric ammonium sulphate

III Assay of the following compounds along with Standardization of Titrant

- (1) Ammonium chloride by acid base titration
- (2) Ferrous sulphate by Cerimetry
- (3) Copper sulphate by Iodometry
- (4) Calcium gluconate by complexometry
- (5) Hydrogen peroxide by Permanganometry
- (6) Sodium benzoate by non-aqueous titration
- (7) Sodium Chloride by precipitation titration

IV Determination of Normality by electro-analytical methods

- (1) Conductometric titration of strong acid against strong base
- (2) Conductometric titration of strong acid and weak acid against strong base
- (3) Potentiometric titration of strong acid against strong base

Recommended Books: (Latest Editions)

- 1. A.H. Beckett & J.B. Stenlake's, Practical Pharmaceutical Chemistry Vol I & II, Stahlone Press of University of London
- 2. A.I. Vogel, Text Book of Quantitative Inorganic analysis
- 3. P. Gundu Rao, Inorganic Pharmaceutical Chemistry
- 4. Bentley and Driver's Textbook of Pharmaceutical Chemistry
- 5. John H. Kennedy, Analytical chemistry principles
- 6. Indian Pharmacopoeia.

BP109P PHARMACEUTICSI (Practical) 4 Hours / week

1.Syrups

- a) Syrup IP'66
- b) Compound syrup of Ferrous Phosphate BPC'68

2. Elixirs

- a) Piperazine citrate elixir
- b) Paracetamol pediatric elixir

3.Linctus

- a) Terpin Hydrate Linctus IP'66
- b) Iodine Throat Paint (Mandles Paint)

4. Solutions

- a) Strong solution of ammonium acetate
- b) Cresol with soap solution
- c) Lugol's solution

5. Suspensions

- a) Calamine lotion
- b) Magnesium Hydroxide mixture
- c) Aluminimum Hydroxide gel

6. Emulsions

- a) Turpentine Liniment
- b) Liquid paraffin emulsion

7. Powders and Granules

- a) ORS powder (WHO)
- b) Effervescent granules
- c) Dusting powder
- d) Divded powders

8. Suppositories

- a) Glycero gelatin suppository
- b) Coca butter suppository
- c) Zinc Oxide suppository

8. Semisolids

- a) Sulphur ointment
- b) Non staining-iodine ointment with methyl salicylate
- c) Carbopal gel

9. Gargles and Mouthwashes

- a) Iodine gargle
- b) Chlorhexidine mouthwash

Recommended Books: (Latest Editions)

- 1. H. C. Ansel et al., Pharmaceutical Dosage Form and Drug Delivery System, Lippincott Williams and Walkins, New Delhi.
- 2. Carter S.J., Cooper and Gunn's-Dispensing for Pharmaceutical Students, CBS publishers, New Delhi
- 3. M.E. Aulton, Pharmaceutics, The Science& Dosage Form Design, Churchill Livingstone, Edinburgh.
- 4. Indian pharmacopoeia.
- 5. British pharmacopoeia.
- 6. Lachmann. Theory and Practice of Industrial Pharmacy, Lea& Febiger Publisher, The University of Michigan.
- 7. Alfonso R. Gennaro Remington. The Science and Practice of Pharmacy, Lippincott Williams, New Delhi.
- 8. Carter S.J., Cooper and Gunn's. Tutorial Pharmacy, CBS Publications, New Delhi.
- 9. E.A. Rawlins, Bentley's Text Book of Pharmaceutics, English Language Book Society, Elsevier Health Sciences, USA.
- 10. Isaac Ghebre Sellassie: Pharmaceutical Pelletization Technology, Marcel Dekker, INC, New York.
- 11. Dilip M. Parikh: Handbook of Pharmaceutical Granulation Technology, Marcel Dekker, INC, New York.
- 12. Françoise Nieloud and Gilberte Marti-Mestres: Pharmaceutical Emulsions and Suspensions, Marcel Dekker, INC, New York.

BP110P PHARMACEUTICAL INORGANIC CHEMISTRY (Practical)

4 Hours / Week

I Limit tests for following ions

- a) Limit test for Chlorides and Sulphates
- b) Modified limit test for Chlorides and Sulphates Limit test for Iron
- c) Limit test for Heavy metals Limit test for Lead
- d) Limit test for Arsenic

II Identification test

a) Magnesium hydroxide Ferrous sulphate Sodium Bicarbonate Calcium gluconate Copper sulphate

III Test for purity

- a) Swelling power of Bentonite
- b) Neutralizing capacity of aluminum hydroxide gel
- c) Determination of potassium iodate and iodine in potassium Iodide

VI Preparation of inorganic pharmaceuticals

- a) Boric acid
- b) Potash alum
- c) Ferrous sulphate

Recommended Books (Latest Editions)

- 1. A.H. Beckett & J.B. Stenlake's, Practical Pharmaceutical Chemistry Vol I & II, Stahlone Press of University of London, 4th edition.
- 2. A.I. Vogel, Text Book of Quantitative Inorganic analysis
- 3. P. Gundu Rao, Inorganic Pharmaceutical Chemistry, 3rd Edition
- 4. M.L Schroff, Inorganic Pharmaceutical Chemistry
- 5. Bentley and Driver's Textbook of Pharmaceutical Chemistry
- 6. Anand & Chatwal, Inorganic Pharmaceutical Chemistry
- 7. IndianPharmacopoeia

BP111P COMMUNICATION SKILLS (Practical) 2 Hours / week

The following learning modules are to be conducted using wordsworth English language lab software

1. Basic communication covering the following topics

- a. Meeting People
- b. Asking Questions
- c. Making Friends
- d. What did you do?
- e. Do's and Dont's

2. Pronunciations covering the following topics

- a. Pronunciation (Consonant Sounds)
- b. Pronunciation and Nouns
- c. Pronunciation (Vowel Sounds)

3. Advanced Learning

- a. Listening Comprehension / Direct and Indirect Speech
- b. Figures of Speech
- c. Effective Communication
- d. Writing Skills
- e. Effective Writing
- f. Interview Handling Skills
- g. E-Mail etiquette
- h. Presentation Skills

Recommended Books: (Latest Edition)

- 1. Basic communication skills for Technology, Andreja. J. Ruther Ford, 2nd Edition, Pearson Education, 2011
- 2. Communication skills, Sanjay Kumar, Pushpalata, 1stEdition, Oxford Press, 2011
- 3. Organizational Behaviour, Stephen .P. Robbins, 1stEdition, Pearson, 2013
- 4. Brilliant- Communication skills, Gill Hasson, 1stEdition, Pearson Life, 2011
- 5. The Ace of Soft Skills: Attitude, Communication and Etiquette for success, Gopala Swamy Ramesh, 5thEdition, Pearson, 2013
- 6. Developing your influencing skills, Deborah Dalley, Lois Burton, Margaret, Green hall, 1st Edition Universe of Learning LTD, 2010
- 7. Communication skills for professionals, Konar nira, 2ndEdition, New arrivals PHI, 2011
- 8. Personality development and soft skills, Barun K Mitra, 1stEdition, Oxford Press, 2011
- 9. Soft skill for everyone, Butter Field, 1st Edition, Cengage Learning India pvt.ltd, 2011

- 10. Soft skills and professional communication, Francis Peters SJ, 1stEdition, Mc Graw Hill Education, 2011
- 11. Effective communication, John Adair, 4thEdition, Pan Mac Millan, 2009
- 12. Bringing out the best in people, Aubrey Daniels, 2ndEdition, Mc Graw Hill, 1999

BP112RBP REMEDIAL BIOLOGY (Practical)

2 Hours / week

- 1. Introduction to experiments in biology
 - a) Study of Microscope
 - b) Section cutting techniques
 - c) Mounting and staining
 - d) Permanent slide preparation
- 2. Study of cell and its inclusions
- 3. Study of Stem, Root, Leaf, seed, fruit, flower and their modifications
- 4. Detailed study of frog by using computer models
- 5. Microscopic study and identification of tissues pertinent to Stem, Root Leaf, seed, fruit and flower
- 6. Identification of bones
- 7. Determination of blood group
- 8. Determination of blood pressure
- 9. Determination of tidal volume

Reference Books

- 1. Practical human anatomy and physiology. by S.R.Kale and R.R.Kale.
- 2. A Manual of pharmaceutical biology practical by S.B.Gokhale, C.K.Kokate and S.P.Shriwastava.
- 3. Biology practical manual according to National core curriculum .Biology forum of Karnataka. Prof .M.J.H.Shafi

SEMESTER II

BP201T HUMAN ANATOMY AND PHYSIOLOGY-II (Theory)3 L + 1T / Week

Scope: This subject is designed to impart fundamental knowledge on the structure andfunctions of the various systems of the human body. It also helps in understanding both homeostatic mechanisms. The subject provides the basic knowledge required to understand the various disciplines of pharmacy.

Objectives: Upon completion of this course the student should be able to:

- 1. Explain the gross morphology, structure and functions of various organs of the human body.
- 2. Describe the various homeostatic mechanisms and their imbalances.
- 3. Identify the various tissues and organs of different systems of human body.
- 4. Perform the hematological tests like blood cell counts, haemoglobin estimation, bleeding/clotting time etc and also record blood pressure, heart rate, pulse and respiratory volume.
- 5. Appreciate coordinated working pattern of different organs of each system
- 6. Appreciate the interlinked mechanisms in the maintenance of normal functioning (homeostasis) of human body.

Sr.	Content	Hours
No.		
1	UNIT I - Nervous system	10
1.1	Organization of nervous system, neuron, neuroglia, classification	4
	and properties of nerve fibre, electrophysiology, action potential,	
	nerve impulse, receptors, synapse, neurotransmitters.	
1.2	Central nervous system: Meninges, ventricles of brain and cerebrospinal	6
	fluid. Structure and functions of brain (cerebrum, brain stem, and	
	cerebellum), spinal cord (gross structure, functions of afferent and efferent	
	nerve tracts,reflex activity).	
2	UNIT II	6
2.1	Digestive system	4
	Anatomy of GI Tract with special reference to anatomy and functions of	
	stomach, (Acid production in the stomach, regulation of acid production	
	through parasympathetic nervous system, pepsin role in protein digestion)	
	small intestine and large intestine, anatomy and functions of salivary	
	glands, pancreas and liver, movements of GIT, digestion and absorption of	
	nutrients and disorders of GIT.	

2.2	Energetics	2
	Formation and role of ATP, Creatinine Phosphate and BMR.	
3	UNIT III	10
3.1	Respiratory system	5
	Anatomy of respiratory system with special reference to anatomy of lungs,	
	mechanism of respiration, regulation of respiration	
	Lung Volumes and capacities transport of respiratory gases, artificial	
	respiration, and resuscitation methods.	
3.2	Urinary system	5
	Anatomy of urinary tract with special reference to anatomy of kidney and	
	nephrons, functions of kidney and urinary tract, physiology of urine	
	formation, micturition reflex and role of kidneys in acid base balance, role	
	of RAS in kidney and disorders of kidney.	
4	UNIT IV	10
	Endocrine system	
	Classification of hormones, mechanism of hormone action, structure and	
	functions of pituitary gland, thyroid gland, parathyroid gland, adrenal	
	gland, pancreas, pineal gland, thymus and their disorders	
5	UNIT V	9
5.1	Reproductive system	6
	Anatomy of male and female reproductive system, Functions of male and	
	female reproductive system, sex hormones, physiology of menstruation,	
	fertilization, spermatogenesis, oogenesis, pregnancy and parturition	
5.2	Introduction to genetics	4
	Chromosomes, genes and DNA, protein synthesis, genetic pattern of	
	inheritance	
	Total	45

Recommended Books (Latest Editions)

- 1. Essentials of Medical Physiology by K. Sembulingam and P. Sembulingam. Jaypee brothers medical publishers, New Delhi.
- 2. Anatomy and Physiology in Health and Illness by Kathleen J.W. Wilson, Churchill Livingstone, New York
- 3. Physiological basis of Medical Practice-Best and Tailor. Williams & Wilkins Co,Riverview,MI USA
- 4. Text book of Medical Physiology- Arthur C,Guyton andJohn.E. Hall. Miamisburg, OH, U.S.A.

- 5. Principles of Anatomy and Physiology by Tortora Grabowski. Palmetto, GA, U.S.A.
- 6. Textbook of Human Histology by Inderbir Singh, Jaypee brothers medical publishers, New Delhi.
- 7. Textbook of Practical Physiology by C.L. Ghai, Jaypee brothers medical publishers, New Delhi.
- 8. Practical workbook of Human Physiology by K. Srinageswari and Rajeev Sharma, Jaypee brother's medical publishers, New Delhi.

Reference Books:

- 1. Physiological basis of Medical Practice-Best and Tailor. Williams & Wilkins Co, Riverview, MI USA
- 2. Text book of Medical Physiology- Arthur C, Guyton and John. E. Hall. Miamisburg, OH, U.S.A.
- 3. Human Physiology (vol 1 and 2) by Dr. C.C. Chatterrje ,Academic Publishers Kolkata

BP202T PHARMACEUTICAL ORGANIC CHEMISTRY –I(Theory)

3L + 1T / Week

Scope: This subject deals with classification and nomenclature of simple organiccompounds, structural isomerism, intermediates forming in reactions, important physical properties, reactions and methods of preparation of these compounds. The syllabus also emphasizes on mechanisms and orientation of reactions.

Objectives: Upon completion of the course the student shall be able to:

- 1. write the structure, name and the type of isomerism of the organic compound
- 2. write the reaction, name the reaction and orientation of reactions
- 3. account for reactivity/stability of compounds,
- 4. identify/confirm the identification of organic compound

Sr. No.	Content	Hours
	General methods of preparation and reactions of compounds superscripted	
	with asterisk (*) to be explained	
	To emphasize on definition, types, classification,	
	principles/mechanisms,applications, examples and differences	
1	UNIT I	7
	Classification, nomenclature and isomerism Classification of Organic	
	Compounds, Common and IUPAC systems of nomenclature of organic	
	compounds (up to 10 Carbons open chain and carbocyclic compounds)	
	Structural isomerisms in organic compounds.	
2	UNIT II - Alkanes*, Alkenes* and Conjugated dienes*	10
2.1	SP ³ hybridization in alkanes, Halogenation of alkanes, uses of paraffins.	3
	Stabilities of alkenes, SP ² hybridization in alkenes	
2.2	E ₁ and E ₂ reactions – kinetics, order of reactivity of alkyl halides,	7
	rearrangement of carbocations, Saytzeffs orientation and evidences. E ₁	
	verses E ₂ reactions, Factors affecting E ₁ and E ₂ reactions. Ozonolysis,	
	electrophilic addition reactions of alkenes, Markownikoff's orientation,	
	free radical addition reactions of alkenes, Anti Markownikoff's	
	orientation.	
	Stability of conjugated dienes, Diel-Alder, electrophilic addition, free	
	radical addition reactions of conjugated dienes, allylic rearrangement	
3	UNIT III	10
3.1	Alkylhalides*	5
	SN ₁ and SN ₂ reactions - kinetics, order of reactivity of alkyl halides,	
	stereochemistry and rearrangement of carbocations.	
	SN ₁ versus SN ₂ reactions, Factors affecting SN ₁ and SN ₂ reactions	

	Structure and uses of ethylchloride, Chloroform, trichloroethylene,	
	tetrachloroethylene, dichloromethane, tetrachloromethane and iodoform.	
3.2	Alcohols*- Qualitative tests, Structure and uses of Ethyl alcohol, Methyl	5
	alcohol, chlorobutanol, Cetosteryl alcohol, Benzyl alcohol, Glycerol,	
	Propylene glycol	
4	UNIT IV	10
	Carbonyl compounds* (Aldehydes and ketones)	
	Nucleophilic addition, Electromeric effect, aldol condensation, Crossed	
	Aldol condensation, Cannizzaro reaction, Crossed Cannizzaro reaction,	
	Benzoin condensation, Perkin condensation, qualitative tests, Structure	
	and uses of Formaldehyde, Paraldehyde, Acetone, Chloral hydrate,	
	Hexamine, Benzaldehyde, Vanilin, Cinnamaldehyde	
5	UNIT V	8
5.1	Carboxylic acids*	5
	Acidity of carboxylic acids, effect of substituents on acidity, inductive	
	effect and qualitative tests for carboxylic acids ,amide and ester	
	Structure and Uses of Acetic acid, Lactic acid, Tartaric acid, Citric acid,	
	Succinic acid. Oxalic acid, Salicylic acid, Benzoic acid, Benzyl benzoate,	
	Dimethyl phthalate, Methyl salicylate and Acetyl salicylic acid	
5.2	Aliphatic amines* - Basicity, effect of substituent on Basicity. Qualitative	3
	test, Structure anduses of Ethanolamine, Ethylenediamine, Amphetamine	
	Total	45

Recommended Books (Latest Editions)

- 1. Organic Chemistry by Morrison and Boyd
- 2. Organic Chemistry by I.L. Finar, Volume-I
- 3. Textbook of Organic Chemistry by B.S. Bahl & Arun Bahl.
- 4. Organic Chemistry by P.L.Soni
- 5. Practical Organic Chemistry by Mann and Saunders.
- 6. Vogel's text book of Practical Organic Chemistry
- 7. Advanced Practical organic chemistry by N.K.Vishnoi.
- 8. Introduction to Organic Laboratory techniques by Pavia, Lampman and Kriz.
- 9. Reaction and reaction mechanism by Ahluwaliah/Chatwal.

BP203T BIOCHEMISTRY (Theory)3 L + 1T / Week

Scope: Biochemistry deals with complete understanding of the molecular levels of thechemical process associated with living cells. The scope of the subject is providing biochemical facts and the principles to understand metabolism of nutrient molecules in physiological and pathological conditions. It is also emphasizing on genetic organization of mammalian genome and hetero & autocatalytic functions of DNA.

Objectives: Upon completion of course student shell able to:

- 1. Understand the catalytic role of enzymes, importance of enzyme inhibitors in design of new drugs, therapeutic and diagnostic applications of enzymes.
- 2. Understand the metabolism of nutrient molecules in physiological and pathological conditions.
- 3. Understand the genetic organization of mammalian genome and functions of DNA in the synthesis of RNAs and proteins.

Sr.	Content	Hours
No.		
1	UNIT I	8
1.1	Biomolecules	
	Introduction, classification, chemical nature and biological role of	
	carbohydrate, lipids, nucleic acids, amino acids and proteins	
1.2	Bioenergetics	
	Concept of free energy, endergonic and exergonic reaction, Relationship	
	between free energy, enthalpy and entropy; Redox potential.	
1.3	Energy rich compounds; classification; biological significances of ATP and	
	cyclic AMP.	
2	UNIT II	10
2.1	Carbohydrate metabolism	
	Glycolysis - Pathway, energetics and significance Citric acid cycle-	
	Pathway, energetics and significance	
	HMP shunt and its significance; Glucose-6-Phosphate dehydrogenase	
	(G6PD) deficiency	
	Glycogen metabolism Pathways and glycogen storage diseases (GSD)	
	Gluconeogenesis- Pathway and its significance	
	Hormonal regulation of blood glucose level and Diabetes mellitus	
2.2	Biological oxidation	
	Electron transport chain (ETC) and its mechanism	
	Oxidative phosphorylation & its mechanism and substrate level	

	phosphorylation	
	Inhibitors ETC and oxidative phosphorylation/Uncouplers	
3	UNIT III	10
3.1	Lipid metabolism	
	β-Oxidation of saturated fatty acid (Palmitic acid)	
	Formation and utilization of ketone bodies; ketoacidosis De novo synthesis	
	of fatty acids (Palmitic acid)	
	Biological significance of cholesterol and conversion of cholesterol into bile	
	acids, steroid hormone and vitamin D	
	Disorders of lipid metabolism: Hypercholesterolemia, atherosclerosis, fatty	
	liver and obesity.	
3.2	Amino acid metabolism	
	General reactions of amino acid metabolism: Transamination, deamination	
	& decarboxylation, urea cycle and its disorders	
	Catabolism of phenylalanine and tyrosine and their metabolic disorders	
	(Phenyketonuria, Albinism, alkeptonuria, tyrosinemia)	
	Synthesis and significance of biological substances; 5-HT, melatonin,	
	dopamine, noradrenaline, adrenaline	
	Catabolism of heme; hyperbilirubinemia and jaundice	
4	UNIT IV	10
	Nucleic acid metabolism and genetic information transfer Biosynthesis	
	of purine and pyrimidine nucleotides	
	Catabolism of purine nucleotides and Hyperuricemia and Gout disease	
	Organization of mammalian genome	
	Structure of DNA and RNA and their functions DNA replication (semi	
	conservative model) Transcription or RNA synthesis	
	Genetic code, Translation or Protein synthesis and inhibitors	
5	UNIT V	07
	Enzymes	
	Introduction, properties, nomenclature and IUB classification of enzymes	
	Enzyme kinetics (Michaelis plot, Line Weaver Burke plot)	
	Enzyme inhibitors with examples	
	Regulation of enzymes: enzyme induction and repression, allosteric	
	enzymes regulation	
	Therapeutic and diagnostic applications of enzymes and isoenzymes	
	Coenzymes –Structure and biochemical functions	
	Total	45

Recommended Books: (Latest Editions)

1. Principles of Biochemistry by Lehninger.

- 2. Harper's Biochemistry by Robert K. Murry, Daryl K. Granner and Victor W. Rodwell.
- 3. Biochemistry by Stryer.
- 4. Biochemistry by D. Satyanarayan and U.Chakrapani
- 5. Textbook of Biochemistry by Rama Rao.
- 6. Textbook of Biochemistry by Deb.
- 7. Outlines of Biochemistry by Conn and Stumpf
- 8. Practical Biochemistry by R.C. Gupta and S. Bhargavan.
- 9. Introduction of Practical Biochemistry by David T. Plummer. (3rd Edition)
- 10. Practical Biochemistry for Medical students by Rajagopal and Ramakrishna.
- 11. Practical Biochemistry by Harold Varley.

BP204T PATHOPHYSIOLOGY (Theory) 4 hours/week

Scope: Pathophysiology is the study of causes of diseases and reactions of the body tosuch disease producing causes. This course is designed to impart a thorough knowledge of the relevant aspects of pathology of various conditions with reference to its pharmacological applications, and understanding of basic pathophysiological mechanisms. Hence it will not only help to study the syllabus of pathology, but also to get baseline knowledge required to practice medicine safely, confidently, rationally and effectively.

Objectives: Upon completion of the subject student shall be able to:

- 1. Describe the etiology and pathogenesis of the selected disease states;
- 2. Name the signs and symptoms of the diseases; and
- 3. Mention the complications of the diseases.

Sr. No.	Content	Hours
1	UNIT I	10
1.1	Basic principles of Cell injury and Adaptation:	5
	Introduction, definitions, Homeostasis, Components and Types of	
	Feedback systems, Causes of cellular injury, Pathogenesis (Cell	
	membrane damage, Mitochondrial damage, Ribosome damage,	
	Nuclear damage), Morphology of cell injury - Adaptive changes	
	(Atrophy, Hypertrophy, hyperplasia, Metaplasia, Dysplasia), Cell	
	swelling, Intra cellular accumulation, Calcification, Enzyme leakage	
	and Cell Death Acidosis & Alkalosis, Electrolyte imbalance	
1.2	Basicmechanism involved in the process of inflammation and	5
	repair:	
	Introduction, Clinical signs of inflammation, Different types of	
	Inflammation, Mechanism of Inflammation – Alteration in vascular	
	permeability and blood flow, migration of WBC's, Mediators of	
	inflammation,Basic principles of wound healing in the	
	skin,Pathophysiology of Atherosclerosis	
2	UNIT II	10
2.1	Cardiovascular System:	4
	Hypertension, congestive heart failure, ischemic heart disease	
	(angina,myocardial infarction, atherosclerosis and arteriosclerosis)	
2.2	Respiratory system: Asthma, Chronic obstructive airways diseases.	3
2.3	Renal system: Acute and chronic renal failure	3
3	UNIT III	10

3.1	Haematological Diseases:	3
	Iron deficiency, megaloblastic anemia (Vit B12 and folic acid),	
	sickle cell anemia, thalasemia, hereditary acquired anemia,	
	hemophilia	
3.2	Endocrine system: Diabetes, thyroid diseases, disorders of sex hormones	3
3.3	Nervous system: Epilepsy, Parkinson's disease, stroke,	3
	psychiatric disorders:depression, schizophrenia and	
	Alzheimer's disease.	
3.4	Gastrointestinal system: Peptic Ulcer	1
4	UNIT IV	08
4.1	Inflammatory bowel diseases, jaundice, hepatitis (A,B,C,D,E,F)	2
	alcoholic liver disease.	
4.2	Disease of bones and joints: Rheumatoid arthritis, osteoporosis and	2
	gout	
4.3	Principles of cancer: classification, etiology and pathogenesis of	4
	cancer	
5	UNIT V	07
5.1	Infectious diseases: Meningitis, Typhoid, Leprosy,	4
	TuberculosisUrinary tract infections	
5.2	Sexually transmitted diseases: AIDS, Syphilis, Gonorrhea	3
	Total	45

Recommended Books (Latest Editions)

- 1. Vinay Kumar, Abul K. Abas, Jon C. Aster; Robbins & Cotran Pathologic Basis of Disease; South Asia edition; India; Elsevier; 2014.
- 2. Harsh Mohan; Text book of Pathology; 6th edition; India; Jaypee Publications; 2010.
- 3. Laurence B, Bruce C, Bjorn K.; Goodman Gilman's The Pharmacological Basis of Therapeutics; 12th edition; New York; McGraw-Hill; 2011.
- 4. Best, Charles Herbert 1899-1978; Taylor, Norman Burke 1885-1972; West, John B (John Burnard); Best and Taylor's Physiological basis of medical practice; 12th ed; united states;
- 5. William and Wilkins, Baltimore;1991 [1990 printing].
- 6. Nicki R. Colledge, Brian R. Walker, Stuart H. Ralston; Davidson's Principles and Practice of Medicine; 21st edition; London; ELBS/Churchill Livingstone; 2010.
- 7. Guyton A, John .E Hall; Textbook of Medical Physiology; 12th edition; WB Saunders Company; 2010.

- 8. Joseph DiPiro, Robert L. Talbert, Gary Yee, Barbara Wells, L. Michael Posey;
- 9. Pharmacotherapy: A Pathophysiological Approach; 9th edition; London; McGraw-Hill Medical; 2014.
- 10. V. Kumar, R. S. Cotran and S. L. Robbins; Basic Pathology; 6th edition; Philadelphia; WB Saunders Company; 1997.
- 11. Roger Walker, Clive Edwards; Clinical Pharmacy and Therapeutics; 3rd edition; London; Churchill Livingstone publication; 2003.

• Recommended Journals

- 1. The Journal of Pathology. ISSN: 1096-9896 (Online)
- 2. The American Journal of Pathology. ISSN: 0002-9440
- 3. Pathology. 1465-3931 (Online)
- 4. International Journal of Physiology, Pathophysiology and Pharmacology. ISSN: 1944-8171 (Online)
- 5. Indian Journal of Pathology and Microbiology. ISSN-0377-4929.

BP205T COMPUTER APPLICATIONS IN PHARMACY (Theory)3hours/week

Scope: This subject deals with the introduction Database, Database Management system, and computer application in clinical studies and use of databases.

Objectives: Upon completion of the course the student shall be able to:

- 1. know the various types of application of computers in pharmacy
- 2. know the various types of databases
- 3. know the various applications of databases in pharmacy

Sr.	Content	Hours
No.		
1	UNIT I	6
1.1	Number system: Binary number system, Decimal number system,	
	Octalnumber system, Hexadecimal number systems, conversion decimal	
	to binary, binary to decimal, octal to binary etc, binary addition, binary	
	subtraction - One's complement ,Two's complement method, binary	
	multiplication, binary division	
1.2	Concept of Information Systems and Software: Information gathering,	
	requirement and feasibility analysis, data flow diagrams, process	
	specifications, input/output design, process life cycle, planning and	
	managing the project.	
2	UNIT II	6
2.1	Web technologies: Introduction to HTML, XML,CSS and Programming	
	languages, introduction to web servers and Server Products	
2.2	Introduction to databases, MYSQL, MS ACCESS, Pharmacy Drug	
	database	
3	UNIT III	
	Application of computers in Pharmacy -Drug information storage	6
	andretrieval, Pharmacokinetics, Mathematical model in Drug design,	Hours
	Hospital and Clinical Pharmacy, Electronic Prescribing and discharge	
	(EP) systems, barcode medicine identification and automated dispensing	
	of drugs, mobile technology and adherence monitoring	
	Diagnostic System, Lab-diagnostic System, Patient Monitoring System,	
	Pharma Information System	
4	UNIT IV	6
	Bioinformatics: Introduction, Objective of Bioinformatics,	
	BioinformaticsDatabases, Concept of Bioinformatics, Impact of	
	Bioinformatics in Vaccine Discovery	

5	UNIT V	
	Computers as data analysis in Preclinical	6
	development: Chromatographic dada analysis (CDS), Laboratory	Hours
	Information management System (LIMS) and Text Information	
	Management System(TIMS)	
	Total	30

Recommended Books: (Latest Editions)

- 1. Computer Application in Pharmacy William E.Fassett –Lea and Febiger, 600 South Washington Square, USA, (215) 922-1330.
- 2. Computer Application in Pharmaceutical Research and Development –Sean Ekins Wiley-Interscience, A John Willey and Sons, INC., Publication, USA
- 3. Bioinformatics (Concept, Skills and Applications) S.C.Rastogi-CBS Publishers and Distributors, 4596/1- A, 11 Darya Gani, New Delhi 110 002(INDIA)
- 4. Microsoft office Access 2003, Application Development Using VBA, SQL Server, DAP and Infopath Cary N.Prague Wiley Dreamtech India (P) Ltd., 4435/7, Ansari Road, Daryagani, New Delhi 110002

BP206T ENVIRONMENTAL SCIENCES (Theory) 3 hours/week

Scope: Environmental Sciences is the scientific study of the environmental system and the status of its inherent or induced changes on organisms. It includes not only the study of physical and biological characters of the environment but also the social and cultural factors and the impact of man on environment.

Objectives: Upon completion of the course the student shall be able to:

- 1. Create the awareness about environmental problems among learners.
- 2. Impart basic knowledge about the environment and its allied problems.
- 3. Develop an attitude of concern for the environment.
- 4. Motivate learner to participate in environment protection and environment improvement.
- 5. Acquire skills to help the concerned individuals in identifying and solving environmental problems.
- 6. Strive to attain harmony with Nature
- 7. Acquire skills to help the concerned individuals in identifying and solving environmental problems.

Sr.	Content	Hours
No.		
1	UNIT I	10
	The Multidisciplinary nature of environmental studies	
	Natural Resources	
	Renewable and non-renewable resources:	
	Natural resources and associated problems	
	a) Forest resources; b) Water resources; c) Mineral resources; d)	
	Food resources; e) Energy resources; f) Land resources: Role of	
	an individual in conservation of natural resources.	
2	UNIT II	10
	Ecosystems	
	Concept of an ecosystem.	
	Structure and function of an ecosystem.	
	Introduction, types, characteristic features, structure and function of the	
	ecosystems: Forest ecosystem; Grassland ecosystem; Desert ecosystem;	
	Aquatic ecosystems (ponds, streams, lakes, rivers, oceans, estuaries)	
3	UNIT III	10
	Environmental Pollution: Air pollution; Water pollution; Soil pollution	
	Total	30

Recommended Books (Latest edition):

- 1. Y.K. Sing, Environmental Science, New Age International Pvt, Publishers, Bangalore
- 2. Agarwal, K.C. 2001 Environmental Biology, Nidi Publ. Ltd. Bikaner.
- 3. Bharucha Erach, The Biodiversity of India, Mapin Pu blishing Pvt. Ltd., Ahmedabad 380 013, India,
- 4. Brunner R.C., 1989, Hazardous Waste Incineration, McGraw Hill Inc. 480p
- 5. Clark R.S., Marine Pollution, Clanderson Press Oxford
- 6. Cunningham, W.P. Cooper, T.H. Gorhani, E & Hepworth, M.T. 2001, Environmental Encyclopedia, Jaico Publ. House, Mumbai, 1196p
- 7. De A.K., Environmental Chemistry, Wiley Eastern Ltd.
- 8. Down of Earth, Centre for Science and Environment

BP207P HUMAN ANATOMY AND PHYSIOLOGY (Practical) 4 Hours/week

Practical physiology is complimentary to the theoretical discussions in physiology. Practicals allow the verification of physiological processes discussed in theory classes through experiments on living tissue, intact animals or normal human beings. This is helpful for developing an insight on the subject.

- 1. To study the integumentary and special senses using specimen, models, etc.,
- 2. To study the nervous system using specimen, models, etc.,
- 3. To study the endocrine system using specimen, models, etc
- 4. To demonstrate the general neurological examination
- 5. To demonstrate the function of olfactory nerve
- 6. To examine the different types of taste.
- 7. To demonstrate the visual acuity
- 8. To demonstrate the reflex activity
- 9. Recording of body temperature
- 10. To demonstrate positive and negative feedback mechanism.
- 11. Determination of tidal volume and vital capacity.
- 12. Study of digestive, respiratory, cardiovascular systems, urinary and reproductive systems with the help of models, charts and specimens.
- 13. Recording of basal mass index .
- 14. Study of family planning devices and pregnancy diagnosis test.
- 15. Demonstration of total blood count by cell analyser
- 16. Permanent slides of vital organs and gonads.

Recommended Books (Latest Editions)

- 1. Essentials of Medical Physiology by K. Sembulingam and P. Sembulingam. Jaypee brothers medical publishers, New Delhi.
- 2. Anatomy and Physiology in Health and Illness by Kathleen J.W. Wilson, Churchill Livingstone, New York
- 3. Physiological basis of Medical Practice-Best and Tailor. Williams & Wilkins Co,Riverview,MI USA
- 4. Text book of Medical Physiology- Arthur C,Guyton andJohn.E. Hall. Miamisburg, OH, U.S.A.
- 5. Principles of Anatomy and Physiology by Tortora Grabowski. Palmetto, GA, U.S.A.
- 6. Textbook of Human Histology by Inderbir Singh, Jaypee brothers medical publishers, New Delhi.
- 7. Textbook of Practical Physiology by C.L. Ghai, Jaypee brothers medical publishers, New Delhi.
- 8. Practical workbook of Human Physiology by K. Srinageswari and Rajeev Sharma, Jaypee brother's medical publishers, New Delhi.

Reference Books:

- 1. Physiological basis of Medical Practice-Best and Tailor. Williams & Wilkins Co, Riverview, MI USA
- 2. Text book of Medical Physiology- Arthur C, Guyton and John. E. Hall. Miamisburg, OH, U.S.A.
- 3. Human Physiology (vol 1 and 2) by Dr. C.C. Chatterrje ,Academic Publishers Kolkata

BP208P PHARMACEUTICAL ORGANIC CHEMISTRY -I (Practical) 4 Hours / week

- 1. Systematic qualitative analysis of unknown organic compounds like
 - a) Preliminary test: Color, odour, aliphatic/aromatic compounds, saturation and unsaturation, etc.
 - b) Detection of elements like Nitrogen, Sulphur and Halogen by Lassaigne's test
 - c) Solubility test
 - d) Functional group test like Phenols, Amides/ Urea, Carbohydrates, Amines, Carboxylic acids, Aldehydes and Ketones, Alcohols, Esters, Aromatic and Halogenated Hydrocarbons, Nitro compounds and Anilides.
 - e) Melting point/Boiling point of organic compounds
 - f) Identification of the unknown compound from the literature using melting point/boiling point.
 - g) Preparation of the derivatives and confirmation of the unknown compound by melting point/ boiling point.
 - h) Minimum 5 unknown organic compounds to be analysed systematically.
- 2. Preparation of suitable solid derivatives from organic compounds
- 3. Construction of molecular models

Recommended Books (Latest Editions)

- 1. Organic Chemistry by Morrison and Boyd
- 2. Organic Chemistry by I.L. Finar, Volume-I
- 3. Textbook of Organic Chemistry by B.S. Bahl & Arun Bahl.
- 4. Organic Chemistry by P.L.Soni
- 5. Practical Organic Chemistry by Mann and Saunders.
- 6. Vogel's text book of Practical Organic Chemistry
- 7. Advanced Practical organic chemistry by N.K.Vishnoi.
- 8. Introduction to Organic Laboratory techniques by Pavia, Lampman and Kriz.
- 9. Reaction and reaction mechanism by Ahluwaliah/Chatwal.

BP209P BIOCHEMISTRY (Practical) 4 Hours / Week

- 1. Qualitative analysis of carbohydrates (Glucose, Fructose, Lactose, Maltose, Sucrose and starch)
- 2. Identification tests for Proteins (albumin and Casein)
- 3. Quantitative analysis of reducing sugars (DNSA method) and Proteins (Biuret method)
- 4. Qualitative analysis of urine for abnormal constituents
- 5. Determination of blood creatinine
- 6. Determination of blood sugar
- 7. Determination of serum total cholesterol
- 8. Preparation of buffer solution and measurement of pH
- 9. Study of enzymatic hydrolysis of starch
- 10. Determination of Salivary amylase activity
- 11. Study the effect of Temperature on Salivary amylase activity.
- 12. Study the effect of substrate concentration on salivary amylase activity.

Recommended Books (Latest Editions)

- 1. Principles of Biochemistry by Lehninger.
- 2. Harper's Biochemistry by Robert K. Murry, Daryl K. Granner and Victor W. Rodwell.
- 3. Biochemistry by Stryer.
- 4. Biochemistry by D. Satyanarayan and U.Chakrapani
- 5. Textbook of Biochemistry by Rama Rao.
- 6. Textbook of Biochemistry by Deb.
- 7. Outlines of Biochemistry by Conn and Stumpf
- 8. Practical Biochemistry by R.C. Gupta and S. Bhargavan.
- 9. Introduction of Practical Biochemistry by David T. Plummer. (3rd Edition)
- 10. Practical Biochemistry for Medical students by Rajagopal and Ramakrishna
- 11. Practical Biochemistry by Harold Varley.

BP210P COMPUTER APPLICATIONS IN PHARMACY (Practical) 2 hours/ week

- 1. Design a questionnaire using a word processing package to gather information about a particular disease.
- 2. Create a HTML web page to show personal information.
- Retrieve the information of a drug and its adverse effects using online tools
- 4 Creating mailing labels Using Label Wizard, generating label in MS WORD
- 5 Create a database in MS Access to store the patient information with the required fields Using access
- 6. Design a form in MS Access to view, add, delete and modify the patient record in the database
- 7. Generating report and printing the report from patient database
- 8. Creating invoice table using MS Access
- 9. Drug information storage and retrieval using MS Access
- 10. Creating and working with queries in MS Access
- 11. Exporting Tables, Queries, Forms and Reports to web pages
- 12. Exporting Tables, Queries, Forms and Reports to XML pages

Recommended books (Latest edition):

- 1. Computer Application in Pharmacy William E.Fassett –Lea and Febiger, 600 South Washington Square, USA, (215) 922-1330.
- 2. Computer Application in Pharmaceutical Research and Development –Sean Ekins Wiley-Interscience, A John Willey and Sons, INC., Publication, USA
- 3. Bioinformatics (Concept, Skills and Applications) S.C.Rastogi-CBS Publishers and Distributors, 4596/1- A, 11 Darya Gani, New Delhi 110 002(INDIA)
- 4. Microsoft office Access 2003, Application Development Using VBA, SQL Server, DAP and Infopath Cary N.Prague Wiley Dreamtech India (P) Ltd., 4435/7, Ansari Road, Daryagani, New Delhi 110002

SEMESTR III

BP301TPHARMACEUTICAL ORGANIC CHEMISTRY-II (Theory)

3L + 1T / Week

Scope: This subject deals with general methods of preparation and reactions of someorganic compounds. Reactivity of organic compounds are also studied here. The syllabus emphasizes on mechanisms and orientation of reactions. Chemistry of fats and oils are also included in the syllabus.

Objectives: Upon completion of the course the student shall be able to

- 1. write the structure, name and the type of isomerism of the organic compound
- 2. write the reaction, name the reaction and orientation of reactions
- 3. account for reactivity/stability of compounds,
- 4. prepare organic compounds

Sr.	Content	Hours
No.	General methods of preparation and reactions of compounds superscripted	
	with asterisk (*) to be explained	
	To emphasize on definition, types, classification, principles/mechanisms,	
	applications, examples and differences	
1	UNIT I- Benzene and its derivatives	10
1.1	Analytical, synthetic and other evidences in the derivation of structure of	3
	benzene, Orbital picture, resonance in benzene, aromatic characters,	
	Huckel's rule	
1.2	Reactions of benzene - nitration, sulphonation, halogenation-reactivity,	3
	Friedelcrafts alkylation- reactivity, limitations, Friedelcrafts acylation.	
1.3	Substituents, effect of substituents on reactivity and orientation of mono	3
	substituted benzene compounds towards electrophilic substitution reaction	
1.4	Structure and uses of DDT, Saccharin, BHC and Chloramine	1
2	UNIT II	10
2.1	Phenols* - Acidity of phenols, effect of substituents on acidity, qualitative	5
	tests, Structure and uses of phenol, cresols, resorcinol, naphthols	
2.2	Aromatic Amines* - Basicity of amines, effect of substituents on basicity,	3
	and synthetic uses of aryl diazonium salts	
2.3	Aromatic Acids* –Acidity, effect of substituents on acidity and important	2
	reactions of benzoic acid.	
3	UNIT III - Fats and Oils	10

3.1	Fatty acids – reactions.	4
3.2	Hydrolysis, Hydrogenation, Saponification and Rancidity of oils, Drying oils	3
3.3	Analytical constants – Acid value, Saponification value, Ester value, Iodine	3
	value, Acetyl value, Reichert Meissl (RM) value – significance and principle	
	involved in their determination.	
4	UNIT IV- Polynuclear hydrocarbons:	08
4.1	Synthesis, reactions	4
4.2	Structure and medicinal uses of Naphthalene, Phenanthrene, Anthracene,	4
	Diphenylmethane, Triphenylmethane and their derivatives	
5	UNIT V	07
	Cyclo alkanes	
	Stabilities - Baeyer's strain theory, limitation of Baeyer's strain theory,	
	Coulson and Moffitt's modification, Sachse Mohr's theory (Theory of	
	strainless rings), reactions of cyclopropane and cyclobutane only	
	Total	45

- 1. Organic Chemistry by R.T. Morrison and R.N.Boyd, 6th edition, Prentice Hall Publications
- 2. Organic Chemistry by Pine, Stanley H.; Hendrickson, James B.; Cram, Donald J.; Hammond, George S., 4th edition. The Macgraw hill publications
- 3. Organic Chemistry by I.L. Finar, Vol 1& 2, 6th edition, Pearson education
- 4. Advanced Organic Chemistry: Reactions, Mechanisms, Structures by Jerry March, John Wiley and sons
- 5. Organic Chemistry, Part A: Structures and Mechanism, Part B: Reactions and Synthesis, Francis and Carry, Richard J Sundberg. Springer publications
- 6. A Guidebook to Mechanism in Organic Chemistry, 6th edition, Peter Sykes, Pearson Education Peter Sykes, Essentials of Organic chemistry by Paul M Dewick, Wiley, Pine
- 7. Essentials of Organic chemistry by Paul M Dewick, Wiley
- 8. Eliel, Kalsi, Organic Chemistry by L.G.Wade, Jr., Maya Shankar Singh, Pearson Education, 6th Ed, Organic Chemistry, 2nd Ed., Thomas Sorrell, University Science Books
- 9. Stereochemistry: Conformation and Mechanism, b) Organic Reactions And Their Mechanisms. By P. S. Kalsi. New age International
- 10. Organic Chemistry through Solved Problems, Goutam Brahmachari. Edition, Morgan & Claypool
- 11. Organic Name Reactions: A Unified Approach. Goutam Brahmachari. Alpha Science publications

BP302T PHYSICAL PHARMACEUTICS-I (Theory) 3 L + 1T / Week

Scope:

The course deals with the various physical and physicochemical properties, and principles involved in dosage forms/formulations. Theory and practical components of the subject help the student to get a better insight into various areas of formulation research and development, and stability studies of pharmaceutical dosage forms.

Objectives: Upon the completion of the course student shall be able to:

- 1. Understand various physicochemical properties of drug molecules in the designing the dosage forms
- 2. Know the principles of chemical kinetics & to use them for stability testing and determination of expiry date of formulations
- 3. Demonstrate use of physicochemical properties in the formulation development and evaluation of dosage forms.

Sr.	Content	Hours
No.		
1	UNIT I- Solubility of drugs	10
1.1	Solubility expressions, mechanisms of solute solvent interactions, ideal	3
	solubility parameters, solvation & association, quantitative approach to the	
	factors influencing solubility of drugs,	
1.2	Dissolution & drug release, diffusion principles in biological systems.	3
	Solubility of gas in liquids, solubility of liquids in liquids, (Binary	
	solutions, ideal solutions)	
1.3	Raoult's law, real solutions, azeotropic mixtures, fractional distillation.	4
	Partially miscible liquids, Critical solution temperature and applications.	
	Distribution law, its limitations and applications	
2	UNIT II	10
2.1	States of Matter and properties of matter: State of matter, changes in the	5
	state of matter, latent heats, vapour pressure, sublimation critical point,	
	eutectic mixtures, gases, aerosols - inhalers, relative humidity, liquid	
	complexes, liquid crystals, glassy states, solid-crystalline, amorphous &	
	polymorphism.	
2.2	Physicochemical properties of drug molecules: Refractive index, optical	5
	rotation, dielectric constant, dipole moment, dissociation constant,	

	determinations and applications	
3	UNIT III	10
	Surface and interfacial phenomenon: Liquid interface, surface &	
	interfacial tensions,	
	surface free energy, measurement of surface & interfacial tensions,	
	spreading coefficient, adsorption at liquid interfaces, surface active agents,	
	HLB Scale, solubilisation, detergency, adsorption at solid interface.	
4	UNIT IV	08
	Complexation and protein binding: Introduction, Classification of	
	Complexation, Applications, methods of analysis, protein binding,	
	Complexation and drug action, crystalline structures of complexes and	
	thermodynamic treatment of stability constants.	
5	UNIT V	07
	pH, buffers and Isotonic solutions: Sorensen's pH scale, pH	
	determination (electrometric and calorimetric), applications of buffers,	
	buffer equation, buffer capacity, buffers in pharmaceutical and biological	
	systems, buffered isotonic solutions.	
	Total	45

- 1. Physical pharmacy by Alfred Martin
- 2. Experimental pharmaceutics by Eugene, Parott.
- 3. Tutorial pharmacy by Cooper and Gunn.
- 4. Stocklosam J. Pharmaceutical calculations, Lea &Febiger, Philadelphia.
- 5. Liberman H.A, Lachman C., Pharmaceutical Dosage forms, Tablets, Volume-1 to 3, MarcelDekkar Inc.
- 6. Liberman H.A, Lachman C, Pharmaceutical dosage forms. Disperse systems, volume 1, 2, 3. Marcel Dekkar Inc.
- 7. Physical pharmaceutics by Ramasamy C and Manavalan R.

$BP303T\,PHARMACEUTICAL\,MICROBIOLOGY\,(Theory) \hspace{0.5cm} 3\,L+1T\,/\,Week$

Scope:

Study of all categories of microorganisims especially for the production of alchol antibiotics, vaccines, vitamins enzymes etc..

Objectives: Upon completion of the subject student shall be able to:

- 1. Understand methods of identification, cultivation and preservation of various microorganisms
- 2. To understand the importance and implementation of sterlization in pharmaceutical processing and industry
- 3. Learn sterility testing of pharmaceutical products.
- 4. Carry out microbiological standardization of Pharmaceuticals.
- 5. Understand the cell culture technology and its applications in pharmaceutical industries.

Sr.	Content	Hour
No.		\mathbf{s}
1	UNIT I	10
1.1	Introduction, history of microbiology, its branches, scope and its importance	1
1.2	Introduction to Prokaryotes and Eukaryotes	1
1.3	Study of ultra-structure and morphological classification of bacteria, nutritional requirements, raw materials used for culture media and physical parameters for growth, growth curve, isolation and preservation methods for pure cultures, cultivation of anaerobes, quantitative measurement of bacterial growth (total & viable count).	6
1.4	Study of different types of phase constrast microscopy, dark field microscopy and electron microscopy	2
2	UNIT II	10
2.1	Identification of bacteria using staining techniques (simple, Gram's &Acid fast staining) and biochemical tests (IMViC).	3
2.2	Study of principle, procedure, merits, demerits and applications of Physical, chemical and mechanical method of sterilization, Evaluation of the efficiency of sterilization method, Equipments employed in large scale sterilization, Sterility indicators	7
3	UNIT III	10
3.1	Study of morphology, classification, reproduction/replication and cultivation	3

	of Fungi and Virus.	
3.2	Classification and mode of action of disinfectants. Factors influencing	4
	disinfection, antiseptics and their evaluation, for bacteriostatic and	
	bactericidal actions	
3.3	Sterility testing of products (solids, liquids, ophthalmic and other sterile	3
	products) according to IP, BP and USP	
4	UNIT IV	08
4.1	Designing of aseptic area, laminar flow equipments; study of different	3
	sources of contamination in an aseptic area and methods of prevention, clean	
	area classification.	
4.2	Principles and methods of different microbiological assay. Methods for	3
	standardization of antibiotics, vitamins and amino acids.	
4.3	Assessment of a new antibiotic and testing of antimicrobial activity of a new	2
	substance.	
5	UNIT V	07
5.1	Types of spoilage, factors affecting the microbial spoilage of pharmaceutical	2
	products, sources and types of microbial contaminants, assessment of	
	microbial contamination and spoilage.	
5.2	Preservation of pharmaceutical products using antimicrobial agents,	2
	evaluation of microbial stability of formulations.	
5.3	Growth of animal cells in culture, general procedure for cell culture,	2
	Primary, established and transformed cell cultures.	
5.4	Application of cell cultures in pharmaceutical industry and research.	1
	Total	45

- 1. W.B. Hugo and A.D. Russel: Pharmaceutical Microbiology, Blackwell Scientific publications, Oxford London.
- 2. Prescott and Dunn., Industrial Microbiology, 4th edition, CBS Publishers & Distributors, Delhi.
- 3. Pelczar, Chan Kreig, Microbiology, Tata McGraw Hill edn.
- 4. Malcolm Harris, Balliere Tindall and Cox: Pharmaceutical Microbiology.
- 5. Rose: Industrial Microbiology.
- 6. Probisher, Hinsdill et al: Fundamentals of Microbiology, 9th ed. Japan
- 7. Cooper and Gunn's: Tutorial Pharmacy, CBS Publisher and Distribution.
- 8. Peppler: Microbial Technology.
- 9. I.P., B.P., U.S.P.- latest editions.
- 10. Ananthnarayan: Text Book of Microbiology, Orient-Longman, Chennai
- 11. Edward: Fundamentals of Microbiology.
- 12. N.K.Jain: Pharmaceutical Microbiology, Vallabh Prakashan, Delhi

BP304T PHARMACEUTICAL ENGINEERING (Theory) 3 L + 1T / Week

Scope: This course is designed to impart a fundamental knowledge on the art and scienceof various unit operations used in pharmaceutical industry.

Objectives: Upon completion of the course student shall be able:

- 1. To know various unit operations used in Pharmaceutical industries.
- 2. To understand the material handling techniques.
- 3. To perform various processes involved in pharmaceutical manufacturing process.
- 4. To carry out various test to prevent environmental pollution.
- 5. To appreciate and comprehend significance of plant lay out design for optimum use of resources.
- 6. To appreciate the various preventive methods used for corrosion control in Pharmaceutical industries

Sr. No	Content	Hours
1	UNIT I	10
1.1	Flow of fluids: Types of manometers, Reynolds number and its significance,	3
	Bernoulli's theorem and its applications, Energy losses, Orifice meter,	
	Venturimeter, Pitot tube and Rotometer.	
1.2	Size Reduction: Objectives, Mechanisms & Laws governing size reduction,	4
	factors affecting size reduction, principles, construction, working, uses,	
	merits and demerits of Hammer mill, ball mill, fluid energy mill, Edge	
	runner mill & end runner mill.	
1.3	Size Separation: Objectives, applications & mechanism of size separation,	3
	official standards of powders, sieves, size separation Principles, construction,	
	working, uses, merits and demerits of Sieve shaker, cyclone separator, Air	
	separator, Bag filter & elutriation tank	
2	UNIT II	10
2.1	Evaporation: Objectives, applications and factors influencing evaporation,	4
	differences between evaporation and other heat process. principles,	
	construction, working, uses, merits and demerits of Steam jacketed kettle,	
	horizontal tube evaporator, climbing film evaporator, forced circulation	
	evaporator, multiple effect evaporator& Economy of multiple effect	
	evaporator.	
2.2	Heat Transfer: Objectives, applications & Heat transfer mechanisms.	3
	Fourier's law, Heat transfer by conduction, convection & radiation. Heat	
	interchangers & heat exchangers.	
2.3	Distillation : Basic Principles and methodology of simple distillation, flash	3
	distillation, fractional distillation, distillation under reduced pressure, steam	

	distillation & molecular distillation	
3	UNIT III	10
3.1	Drying: Objectives, applications & mechanism of drying process,	5
	measurements & applications of Equilibrium Moisture content, rate of	
	drying curve. principles, construction, working, uses, merits and demerits of	
	Tray dryer, drum dryer spray dryer, fluidized bed dryer, vacuum dryer,	
	freeze dryer	
3.2	Mixing: Objectives, applications & factors affecting mixing, Difference	5
	between solid and liquid mixing, mechanism of solid mixing, liquids mixing	
	and semisolids mixing. Principles, Construction, Working, uses, Merits and	
	Demerits of Double cone blender, twin shell blender, ribbon blender, Sigma	
	blade mixer, planetary mixers, Propellers, Turbines, Paddles & Silverson	
	Emulsifier	
4	UNIT IV	08
4.1	Filtration: Objectives, applications, Theories & Factors influencing	4
	filtration, filter aids, filter medias. Principle, Construction, Working, Uses,	
	Merits and demerits of plate & frame filter, filter leaf, rotary drum filter,	
	Meta filter & Cartridge filter, membrane filters and Seidtz filter.	
4.2	Centrifugation: Objectives, principle & applications of Centrifugation,	4
	principles, construction, working, uses, merits and demerits of Perforated	
	basket centrifuge, Non-perforated basket centrifuge, semi continuous	
	centrifuge & super centrifuge.	
5	UNIT V	07
	Materials of pharmaceutical plant construction, Corrosion and its	
	prevention: Factors affecting during materials selected for Pharmaceutical	
	plant construction, Theories of corrosion, types of corrosion and there	
	prevention. Ferrous and nonferrous metals, inorganic and organic non	
	metals, basic of material handling system	
	Total	45
	10tti	73

- 1. Introduction to chemical engineering Walter L Badger & Julius Banchero, Latest edition.
- 2. Solid phase extraction, Principles, techniques and applications by Nigel J.K. Simpson-Latest edition.
- 3. Unit operation of chemical engineering Mcabe Smith, Latest edition.
- 4. Pharmaceutical engineering principles and practices C.V.S Subrahmanyam et al., Latest edition.
- 5. Remington practice of pharmacy- Martin, Latest edition.

- 6. Theory and practice of industrial pharmacy by Lachmann., Latest edition.
- 7. Physical pharmaceutics- C.V.S Subrahmanyam et al., Latest edition.
- 8. Cooper and Gunn's Tutorial pharmacy, S.J. Carter, Latest edition.

BP305P PHARMACEUTICAL ORGANIC CHEMISTRY -II (Practical)

4 Hours / Week

- I Experiments involving laboratory techniques
 - Recrystallization
 - Steam distillation
- II Determination of following oil values (including standardization of reagents)
 - Acid value
 - Saponification value
 - Iodine value

III Preparation of compounds

- Benzanilide/Phenyl benzoate/Acetanilide from Aniline/Phenol/Aniline by acylation reaction.
- 2,4,6-Tribromo aniline/Para bromo acetanilide from Aniline/
- Acetanilide by halogenation (Bromination) reaction.
- 5-Nitro salicylic acid/Meta di nitro benzene from Salicylic acid / Nitro benzene by nitration reaction.
- Benzoic acid from Benzyl chloride by oxidation reaction.
- Benzoic acid/ Salicylic acid from alkyl benzoate/ alkyl salicylate by hydrolysis reaction.
- 1-Phenyl azo-2-napthol from Aniline by diazotization and coupling reactions.
- Benzil from Benzoin by oxidation reaction.
- Dibenzal acetone from Benzaldehyde by Claison Schmidt reaction
- Cinnammic acid from Benzaldehyde by Perkin reaction
- P-Iodo benzoic acid from P-amino benzoic acid

- 1. Organic Chemistry by Morrison and Boyd
- 2. Organic Chemistry by I.L. Finar, Volume-I
- 3. Textbook of Organic Chemistry by B.S. Bahl & Arun Bahl.
- 4. Organic Chemistry by P.L.Soni
- 5. Practical Organic Chemistry by Mann and Saunders.
- 6. Vogel's text book of Practical Organic Chemistry
- 7. Advanced Practical organic chemistry by N.K. Vishnoi.
- 8. Introduction to Organic Laboratory techniques by Pavia, Lampman and Kriz.

BP306P PHYSICAL PHARMACEUTICS – I (Practical) 4 Hours/week

- 1. Determination the solubility of drug at room temperature
- 2. Determination of pKa value by Half Neutralization/ Henderson Hasselbalch equation.
- 3. Determination of Partition co- efficient of benzoic acid in benzene and water
- 4. Determination of Partition co- efficient of Iodine in CCl4 and water
- 5. Determination of % composition of NaCl in a solution using phenol-water system by CST method
- 6. Determination of surface tension of given liquids by drop count and drop weight method
- 7. Determination of HLB number of a surfactant by saponification method
- 8. Determination of Freundlich and Langmuir constants using activated char coal
- 9. Determination of critical micellar concentration of surfactants
- 10. Determination of stability constant and donor acceptor ratio of PABA-Caffeine complex by solubility method
- 11. Determination of stability constant and donor acceptor ratio of Cupric-Glycine complex by pH titration method

Recommended Books: (Latest Editions)

- 1. Physical pharmacy by Alfred Martin
- 2. Experimental pharmaceutics by Eugene, Parott.
- 3. Tutorial pharmacy by Cooper and Gunn.
- 4. Stocklosam J. Pharmaceutical calculations, Lea & Febiger, Philadelphia.
- 5. Liberman H.A, Lachman C., Pharmaceutical Dosage forms, Tablets, Volume-1 to 3, MarcelDekkar Inc.
- 6. Liberman H.A, Lachman C, Pharmaceutical dosage forms. Disperse systems, volume 1, 2, 3. Marcel Dekkar Inc.
- 7. Physical pharmaceutics by Ramasamy C and ManavalanR.
- 8. Laboratory manual of physical pharmaceutics, C.V.S. Subramanyam, J. Thimma settee

BP307P PHARMACEUTICAL MICROBIOLOGY (Practical)4 Hours/week

- 1. Introduction and study of different equipments and processing, e.g., B.O.D. incubator, laminar flow, aseptic hood, autoclave, hot air sterilizer, deep freezer, refrigerator, microscopes used in experimental microbiology.
- 2. Sterilization of glassware, preparation and sterilization of media.
- 3. Sub culturing of bacteria and fungus. Nutrient stabs and slants preparations.
- 4. Staining methods- Simple, Grams staining and acid fast staining (Demonstration with practical).
- 5. Isolation of pure culture of micro-organisms by multiple streak plate technique and other techniques.
- 6. Microbiological assay of antibiotics by cup plate method and other methods
- 7. Motility determination by Hanging drop method.
- 8. Sterility testing of pharmaceuticals.
- 9. Bacteriological analysis of water
- 10. Biochemical test (IMViC reactions)
- 11. Revision Practical Class

Recommended Books (Latest edition)

- 1. W.B. Hugo and A.D. Russel: Pharmaceutical Microbiology, Blackwell Scientific publications, Oxford London.
- 2. Prescott and Dunn., Industrial Microbiology, 4th edition, CBS Publishers & Distributors, Delhi.
- 3. Pelczar, Chan Kreig, Microbiology, Tata McGraw Hill edn.
- 4. Malcolm Harris, Balliere Tindall and Cox: Pharmaceutical Microbiology.
- 5. Rose: Industrial Microbiology.
- 6. Probisher, Hinsdill et al: Fundamentals of Microbiology, 9th ed. Japan
- 7. Cooper and Gunn's: Tutorial Pharmacy, CBS Publisher and Distribution.
- 8. Peppler: Microbial Technology.
- 9. I.P., B.P., U.S.P.- latest editions.
- 10. Ananthnarayan: Text Book of Microbiology, Orient-Longman, Chennai
- 11. Edward: Fundamentals of Microbiology.
- 12. N.K.Jain: Pharmaceutical Microbiology, Vallabh Prakashan, Delhi
- 13. Bergeys manual of systematic bacteriology, Williams and Wilkins- A Waverly company

BP308P PHARMACEUTICAL ENGINEERING (Practical) 4 Hours/week

- 1. Determination of radiation constant of brass, iron, unpainted and painted glass.
- 2. Steam distillation To calculate the efficiency of steam distillation.
- 3. To determine the overall heat transfer coefficient by heat exchanger.
- 4. Construction of drying curves (for calcium carbonate and starch).
- 5. Determination of moisture content and loss on drying.
- 6. Determination of humidity of air -i) From wet and dry bulb temperatures –use of Dew point method.
- 7. Description of Construction working and application of Pharmaceutical Machinery such as rotary tablet machine, fluidized bed coater, fluid energy mill, de humidifier.
- 8. Size analysis by sieving To evaluate size distribution of tablet granulations Construction of various size frequency curves including arithmetic andlogarithmic probability plots.
- 9. Size reduction: To verify the laws of size reduction using ball mill and determining Kicks, Rittinger's, Bond's coefficients, power requirement and critical speed of Ball Mill.
- 10. Demonstration of colloid mill, planetary mixer, fluidized bed dryer, freeze dryer and such othermajor equipment.
- 11. Factors affecting Rate of Filtration and Evaporation (Surface area, Concentration and Thickness/ viscosity
- 12. To study the effect of time on the Rate of Crystallization.
- 13. To calculate the uniformity Index for given sample by using Double Cone Blender.

Recommended Books: (Latest Editions)

- 1. Introduction to chemical engineering Walter L Badger & Julius Banchero, Latest edition.
- 2. Solid phase extraction, Principles, techniques and applications by Nigel J.K. Simpson-Latest edition.
- 3. Unit operation of chemical engineering Mcabe Smith, Latest edition.
- 4. Pharmaceutical engineering principles and practices C.V.S Subrahmanyam et al., Latest edition.
- 5. Remington practice of pharmacy- Martin, Latest edition.
- 6. Theory and practice of industrial pharmacy by Lachmann., Latest edition.
- 7. Physical pharmaceutics- C.V.S Subrahmanyam et al., Latest edition.

SEMESTER IV

BP401T PHARMACEUTICAL ORGANIC CHEMISTRY –III (Theory)

3L + 1T / Week

Scope: This subject imparts knowledge on stereo-chemical aspects of organic compounds and organic reactions, important named reactions, chemistry of important hetero cyclic compounds. It also emphasizes on medicinal and other uses of organic compounds.

Objectives: At the end of the course, the student shall be able to:

- 1. understand the methods of preparation and properties of organic compounds
- 2. explain the stereo chemical aspects of organic compounds and stereo chemical reactions
- 3. know the medicinal uses and other applications of organic compounds

Sr.	Content	Hours
No.	Note: To emphasize on definition, types, mechanisms, examples,	
	uses/applications	
1	UNIT I	10
	Stereo isomerism	
	Optical isomerism –	
	i. Optical activity, enantiomerism, diastereoisomerism, meso compounds	
	ii. Elements of symmetry, chiral and achiral molecules	
	iii. DL system of nomenclature of optical isomers, sequence rules, RS	
	system of nomenclature of optical isomers	
	iv. Reactions of chiral molecules	
	v. Racemic modification and resolution of racemic mixture.	
	vi. Asymmetric synthesis: partial and absolute	
2	UNIT II	10
	Geometrical isomerism	
	i. Nomenclature of geometrical isomers (Cis Trans, EZ, Syn Anti systems)	
	ii. Methods of determination of configuration of geometrical isomers.	
	iii. Conformational isomerism in Ethane, n-Butane and Cyclohexane.	
	iv. Stereo isomerism in biphenyl compounds (Atropisomerism) and	
	conditions for optical activity.	
	v. Stereospecific and stereoselective reactions	
3	UNIT III - Heterocyclic compounds:	10
	Heterocyclic compounds:	
	Nomenclature and classification	
	Synthesis, reactions and medicinal uses of following compounds/derivatives	
	Pyrrole, Furan, and Thiophene	
	Relative aromaticity and reactivity of Pyrrole, Furan and Thiophene	

4	UNIT IV	08
	Synthesis, reactions and medicinal uses of following compounds/derivatives	
	Pyrazole, Imidazole, Oxazole and Thiazole.	
	Pyridine, Quinoline, Isoquinoline, Acridine and Indole. Basicity of pyridine	
	Synthesis and medicinal uses of Pyrimidine, Purine, azepines and their	
	derivatives	
5	UNIT V - Reactions of synthetic importance	07
5.1	Metal hydride reduction (NaBH ₄ and LiAlH ₄), Clemmensen reduction,	2
	Birch reduction, Wolff Kishner reduction	
5.2	Oppenauer-oxidation and Dakin reaction.	2
5.3	Beckmanns rearrangement and Schmidt rearrangement	2
5.4	Claisen-Schmidt condensation	1
	Total	45

- 1. Organic chemistry by I.L. Finar, Volume-I & II.
- 2. A text book of organic chemistry Arun Bahl, B.S. Bahl.
- 3. Heterocyclic Chemistry by Raj K. Bansal
- 4. Organic Chemistry by Morrison and Boyd
- 5. Heterocyclic Chemistry by T.L. Gilchrist

BP402T MEDICINAL CHEMISTRY – I (Theory) 3 L + 1T / Week

Scope: This subject is designed to impart fundamental knowledge on the structure, chemistry and therapeutic value of drugs. The subject emphasizes on structure activity relationships of drugs, importance of physicochemical properties and metabolism of drugs. The syllabus also emphasizes on chemical synthesis of important drugs under each class.

Objectives: Upon completion of the course the student shall be able to:

- 1. understand the chemistry of drugs with respect to their pharmacological activity
- 2. understand the drug metabolic pathways, adverse effect and therapeutic value of drugs
- 3. know the Structural Activity Relationship (SAR) of different class of drugs
- 4. write the chemical synthesis of some drugs

Sr.	Content	Hours
No.	Study of the development of the following classes of drugs, Classification,	
	mechanism of action, uses of drugs mentioned in the course, Structure	
	activity relationship of selective class of drugs as specified in the course and	
	synthesis of drugs superscripted*	
1	UNIT I - Introduction to Medicinal Chemistry	10
1.1	History and development of medicinal chemistry	1
1.2	Physicochemical properties in relation to biological action	4
	Ionization, Solubility, Partition Coefficient, Hydrogen bonding, Protein	
	binding, Chelation, Bioisosterism, Optical and Geometrical isomerism	
1.3	Drug metabolism	5
	• Drug metabolism principles- Phase I and Phase II.	
	Factors affecting drug metabolism including stereo chemical aspects	
2	UNIT II - Drugs acting on Autonomic Nervous System	10
2.1	Adrenergic Neurotransmitters:	2
	Biosynthesis and catabolism of catecholamine.	
	• Adrenergic receptors (Alpha & Beta) and their distribution.	
2.2	Sympathomimetic agents: SAR of Sympathomimetic agents	4
	• Direct acting: Nor-epinephrine, Epinephrine, Phenylephrine*,	
	Dopamine	
	Methyldopa, Clonidine, Dobutamine, Isoproterenol, Terbutaline,	
	Salbutamol*, Bitolterol, Naphazoline, Oxymetazoline and	

	Xylometazoline	
	Indirect acting agents: Hydroxyamphetamine, Pseudoephedrine,	
	Propylhexedrine.	
	Agents with mixed mechanism: Ephedrine, Metaraminol.	
2.3	Adrenergic Antagonists:	4
	Alpha adrenergic blockers: Tolazoline*, Phentolamine,	
	Phenoxybenzamine, Prazosin, Dihydroergotamine, Methysergide.	
	Beta adrenergic blockers: SAR of beta blockers, Propranolol*,	
	Metibranolol, Atenolol, Betazolol, Bisoprolol, Esmolol, Metoprolol,	
	Labetolol, Carvedilol.	
3	UNIT III - Cholinergic neurotranimitters	10
3.1	Biosynthesis and catabolism of acetylcholine.	2
	Cholinergic receptors (Muscarinic & Nicotinic) and their distribution	
3.2	Parasympathomimetic agents: SAR of Parasympathomimetic agents	4
	• Direct acting agents: Acetylcholine, Carbachol*, Bethanechol,	
	Methacholine, Pilocarpine.	
	Indirect acting/ Cholinesterase inhibitors (Reversible &	
	Irreversible): Physostigmine, Neostigmine*, Pyridostigmine,	
	Edrophonium chloride, Tacrine hydrochloride, Ambenonium chloride,	
	Isofluorphate, Echothiophate iodide, Parathione, Malathion.	
	Cholinesterase reactivator: Pralidoxime chloride.	
3.3	Cholinergic Blocking agents: SAR of cholinolytic agents	4
	Solanaceous alkaloids and analogues: Atropine sulphate,	
	Hyoscyamine sulphate, Scopolamine hydrobromide, Homatropine	
	hydrobromide, Ipratropium bromide*.	
	• Synthetic cholinergic blocking agents: Tropicamide, Cyclopentolate	
	hydrochloride, Clidinium bromide, Dicyclomine hydrochloride*,	
	Glycopyrrolate, Methantheline bromide, Propantheline bromide,	
	Benztropine mesylate, Orphenadrine citrate, Biperidine hydrochloride,	
	Procyclidine hydrochloride*, Tridihexethyl chloride, Isopropamide	
	iodide, Ethopropazine hydrochloride.	
4	LINUT IV Dans of cather on Control Norman Creatons	00
4	UNIT IV - Drugs acting on Central Nervous System	08
4.1	Sedatives and Hypnotics: Paradiagarinas SAP of Paradiagarinas Chlordiagaravida	3
	Benzodiazepines: SAR of Benzodiazepines, Chlordiazepoxide, Diazapam* Ovazapam Chlorazapata Largazapam Almazalam	
	Diazepam*, Oxazepam, Chlorazepate, Lorazepam, Alprazolam,	
	Zolpidem Powbituwtog: SAP of barbiturates, Porbital* Phonobarbital	
	Barbiturtes: SAR of barbiturates, Barbital*, Phenobarbital,	

	Mephobarbital, Amobarbital, Butabarbital, Pentobarbital, Secobarbital	
	3.61	
	Miscelleneous: Amides and imides:Glutethmide.	
	Aldebyde & their derivatives: Trielefes and ym Pereldebyde	
	Aldehyde & their derivatives: Triclofos sodium, Paraldehyde.	
4.2	Antipsychotics	3
	• Phenothiazeines: SAR of Phenothiazeines - Promazine hydrochloride,	
	Chlorpromazine hydrochloride*, Triflupromazine, Thioridazine	
	hydrochloride, Piperacetazine hydrochloride, Prochlorperazine maleate,	
	Trifluoperazine hydrochloride.	
	• Ring Analogues of Phenothiazeines: Chlorprothixene, Thiothixene,	
	Loxapine succinate, Clozapine.	
	• Fluro buterophenones: Haloperidol, Droperidol, Risperidone.	
	Beta amino ketones: Molindone hydrochloride.	
	Benzamides: Sulpieride.	
4.3	Anticonvulsants: SAR of Anticonvulsants, mechanism of anticonvulsant	2
	action	
	Barbiturates: Phenobarbitone, Methabarbital.	
	• Hydantoins: Phenytoin*, Mephenytoin, Ethotoin	
	Oxazolidine diones: Trimethadione, Paramethadione	
	Succinimides: Phensuximide, Methsuximide, Ethosuximide*	
	Urea and monoacylureas: Phenacemide, Carbamazepine*	
	Benzodiazepines: Clonazepam	
	Miscellaneous: Primidone, Valproic acid , Gabapentin, Felbamate	
5	UNIT V- Drugs acting on Central Nervous System	07
5.1	General anesthetics:	3
	• Inhalation anesthetics: Halothane*, Methoxyflurane, Enflurane,	
	Sevoflurane, Isoflurane, Desflurane.	
	• Ultra short acting barbitutrates: Methohexital sodium*, Thiamylal	
	sodium, Thiopental sodium.	
	Dissociative anesthetics: Ketamine hydrochloride.*	
5.2	Narcotic and non-narcotic analgesics	2
	Morphine and related drugs: SAR of Morphine analogues, Morphine	
	sulphate, Codeine, Meperidine hydrochloride, Anilerdine hydrochloride,	
	Diphenoxylate hydrochloride, Loperamide hydrochloride, Fentanyl	
	citrate*, Methadone hydrochloride*, Propoxyphene hydrochloride,	
	Pentazocine, Levorphanol tartarate.	

	Narcotic antagonists: Nalorphine hydrochloride, Levallorphan	
	tartarate, Naloxone hydrochloride	
5.3	Anti-inflammatory agents: Sodium salicylate, Aspirin, Mefenamic acid*,	2
	Meclofenamate, Indomethacin, Sulindac, Tolmetin, Zomepriac, Diclofenac,	
	Ketorolac, Ibuprofen*, Naproxen, Piroxicam, Phenacetin, Acetaminophen,	
	Antipyrine, Phenylbutazone	
	Total	45

- 1. Wilson and Giswold's Organic medicinal and Pharmaceutical Chemistry.
- 2. Foye's Principles of Medicinal Chemistry.
- 3. Burger's Medicinal Chemistry, Vol I to IV.
- 4. Introduction to principles of drug design- Smith and Williams.
- 5. Remington's Pharmaceutical Sciences.
- 6. Martindale's extra pharmacopoeia.
- 7. Organic Chemistry by I.L. Finar, Vol. II.
- 8. The Organic Chemistry of Drug Synthesis by Lednicer, Vol. 1-5.
- 9. Indian Pharmacopoeia.
- 10. Text book of practical organic chemistry- A.I.Vogel.

BP403T PHYSICAL PHARMACEUTICS-II (Theory) 3 L + 1T / Week

Scope: The course deals with the various physical and physicochemical properties, and principles involved in dosage forms/formulations. Theory and practical components of the subject help the student to get a better insight into various areas of formulation research and development, and stability studies of pharmaceutical dosage forms.

Objectives: Upon the completion of the course student shall be able to:

- 1. Understand various physicochemical properties of drug molecules in the designing the dosage forms
- 2. Know the principles of chemical kinetics & to use them for stability testing and determination of expiry date of formulations
- 3. Demonstrate use of physicochemical properties in the formulation development and evaluation of dosage forms

Sr.	Content	Hours
No.		
1	UNIT I	7
	Colloidal dispersions: Classification of dispersed systems & their general	
	characteristics, size & shapes of colloidal particles, classification of colloids	
	& comparative account of their general properties. Optical, kinetic &	
	electrical properties. Effect of electrolytes, coacervation, peptization&	
	protective action.	
2	UNIT II	10
2.1	Rheology: Newtonian systems, law of flow, kinematic viscosity, effect of	7
	temperature, non-Newtonian systems, pseudoplastic, dilatants, plastic,	
	thixotropy, thixotropy in formulation, determination of viscosity, capillary,	
	falling Sphere, rotational viscometers	
2.2	Deformation of solids: Plastic and elastic deformation, Heckel equation,	3
	Stress, Strain, Elastic Modulus	
3	UNIT III - Coarse dispersion	10
3.1	Suspension, interfacial properties of suspended particles, settling in	7
	suspensions, formulation of suspensions. Emulsions and theories of	
	emulsification, microemulsion and multiple emulsions;	
3.2	Physical stability of emulsions, preservation of emulsions, rheological	3
	properties of emulsions, phase equilibria and emulsion formulation.	
4	UNIT IV	8
	Micromeretics: Particle size and distribution, mean particle size, number	
	and weight distribution, particle number, methods for determining particle	
	size by different methods, counting and separation method, particle shape,	

	specific surface, methods for determining surface area, permeability,	
	adsorption, derived properties of powders, porosity, packing arrangement,	
	densities, bulkiness & flow properties.	
5	UNIT V	10
5.1	Drug stability: Reaction kinetics: zero, pseudo-zero, first & second order,	
	units of basic rate constants, determination of reaction order. Physical and	
	chemical factors influencing the chemical degradation of pharmaceutical	
	product: temperature, solvent, ionic strength, dielectric constant, specific &	
	general acid base catalysis, Simple numerical problems. Stabilization of	
	medicinal agents against common reactions like hydrolysis & oxidation.	
	Accelerated stability testing in expiration dating of pharmaceutical dosage	
	forms. Photolytic degradation and its prevention	
	Total	45

- 1. Physical Pharmacy by Alfred Martin, Sixth edition
- 2. Experimental pharmaceutics by Eugene, Parott.
- 3. Tutorial pharmacy by Cooper and Gunn.
- 4. Stocklosam J. Pharmaceutical calculations, Lea & Febiger, Philadelphia.
- 5. Liberman H.A, Lachman C., Pharmaceutical Dosage forms, Tablets, Volume-1 to 3, Marcel Dekkar Inc.
- 6. Liberman H.A, Lachman C, Pharmaceutical dosage forms. Disperse systems, volume 1, 2, 3. Marcel Dekkar Inc.
- 7. Physical Pharmaceutics by Ramasamy C, and Manavalan R.

BP404T PHARMACOLOGY-I (Theory) 3 L + 1T / Week

Scope: The main purpose of the subject is to understand what drugs do to the livingorganisms and how their effects can be applied to therapeutics. The subject covers the information about the drugs like, mechanism of action, physiological and biochemical effects (pharmacodynamics) as well as absorption, distribution, metabolism and excretion (pharmacokinetics) along with the adverse effects, clinical uses, interactions, doses, contraindications and routes of administration of different classes of drugs.

Objectives: Upon completion of this course the student should be able to

- 1. Understand the pharmacological actions of different categories of drugs
- 2. Explain the mechanism of drug action at organ system/sub cellular/ macromolecular levels.
- 3. Apply the basic pharmacological knowledge in the prevention and treatment of various diseases.
- 4. Observe the effect of drugs on animals by simulated experiments
- 5. Appreciate correlation of pharmacology with other bio medical sciences

Sr. No.	Content	Hours
1	UNIT I - General Pharmacology	8
1.1	Introduction to Pharmacology- Definition, historical landmarks and scope	4
	of pharmacology, nature and source of drugs, essential drugs concept and	
	routes of drug administration, Agonists, antagonists(competitive and non-	
	competitive), spare receptors, addiction, tolerance, dependence,	
	tachyphylaxis, idiosyncrasy, allergy	
1.2	Pharmacokinetics- Membrane transport, absorption, distribution, metabolism and excretion of drugs .Enzyme induction, enzyme inhibition,	4
	kinetics of elimination	
2	UNIT II - General Pharmacology	12
2.1	Pharmacodynamics- Principles and mechanisms of drug action. Receptor	6
	theories and classification of receptors, regulation of receptors. drug	
	receptors interactions signal transduction mechanisms, G-protein-coupled	
	receptors, ion channel receptor, transmembrane enzyme linked receptors,	
	transmembrane JAK-STAT binding receptor and receptors that regulate	
	transcription factors, dose response relationship, therapeutic index,	
	combined effects of drugs and factors modifying drug action.	
2.2	Adverse drug reactions.	2
2.3	Drug interactions (pharmacokinetic and pharmacodynamic)	2
2.4	Drug discovery and clinical evaluation of new drugs -Drug discovery phase,	2
	preclinical evaluation phase, clinical trial phase, phases of clinical trials and	

	pharmacovigilance	
3	UNIT III -Pharmacology of peripheral nervous system	10
3.1	Organization and function of ANS, Neurohumoral transmission,co-	1
	transmission and classification of neurotransmitters	
3.2	Parasympathomimetics, Parasympatholytics, Sympathomimetics,	3
	sympatholytics	
3.3	Neuromuscular blocking agents and skeletal muscle relaxants (peripheral).	2
3.4	Local anesthetic agents	3
3.5	Drugs used in myasthenia gravis and glaucoma	1
4	UNIT IV - Pharmacology of central nervous system	08
4.1	Neurohumoral transmission in the C.N.S.special emphasis on importance of	1
	various neurotransmitters like with GABA, Glutamate, Glycine, serotonin,	
	dopamine.	
4.2	General anesthetics and pre-anesthetics.	2
4.3	Sedatives, hypnotics and centrally acting muscle relaxants	2
4.4	Anti-epileptics	2
4.5	Alcohols and disulfiram	1
5	UNIT V - Pharmacology of central nervous system	07
5.1	Psychopharmacological agents: Antipsychotics, antidepressants, anti-	2
	anxiety agents, anti-manics and hallucinogens	
5.2	Drugs used in Parkinsons disease and Alzheimer's disease.	1
5.3	CNS stimulants and nootropics	1
5.4	Opioid analgesics and antagonists	2
5.5	Drug addiction, drug abuse, tolerance and dependence.	1
	Total	45

- 1. Rang H. P., Dale M. M., Ritter J. M., Flower R. J., Rang and Dale's Pharmacology, Churchil Livingstone Elsevier
- 2. Katzung B. G., Masters S. B., Trevor A. J., Basic and clinical pharmacology, Tata Mc Graw-Hill
- 3. Goodman and Gilman's, The Pharmacological Basis of Therapeutics
- 4. Marry Anne K. K., Lloyd Yee Y., Brian K. A., Robbin L.C., Joseph G. B., Wayne A. K., Bradley R.W., Applied Therapeutics, The Clinical use of Drugs, The Point Lippincott Williams & Wilkins
- 5. Mycek M.J, Gelnet S.B and Perper M.M. Lippincott's Illustrated Reviews- Pharmacology
- 6. K.D.Tripathi. Essentials of Medical Pharmacology, JAYPEE Brothers Medical Publishers (P) Ltd, New Delhi.
- 7. Sharma H. L., Sharma K. K., Principles of Pharmacology, Paras medical publisher

- 8. Modern Pharmacology with clinical Applications, by Charles R.Craig& Robert,
- 9. Ghosh MN. Fundamentals of Experimental Pharmacology. Hilton & Company, Kolkata.
- 10. Kulkarni SK. Handbook of experimental pharmacology. VallabhPrakashan

BP405T PHARMACOGNOSY AND PHYTOCHEMISTRY I (Theory)

3L + 1T / Week

Scope: The subject involves the fundamentals of Pharmacognosy like scope, classification of crude drugs, their identification and evaluation, phytochemicals present in them and their medicinal properties.

Objectives: Upon completion of the course, the student shall be able:

- 1. to know the techniques in the cultivation and production of crude drugs
- 2. to know the crude drugs, their uses and chemical nature
- 3. know the evaluation techniques for the herbal drugs
- 4. to carry out the microscopic and morphological evaluation of crude drugs

Sr.	Content	Hour
No.		S
1	UNIT I	10
1.1	Introduction to Pharmacognosy:	3
	(a) Definition, history, scope and development of Pharmacognosy	
	(b) Sources of Drugs – Plants, Animals, Marine & Tissue culture	
	(c) Organized drugs, unorganized drugs (dried latex, dried juices, dried	
	extracts, gums and mucilages, oleoresins and oleo- gum -resins).	
1.2	Classification of drugs:	2
	Alphabetical, morphological, taxonomical, chemical, pharmacological,	
	chemo and sero taxonomical classification of drugs	
1.3	Quality control of Drugs of Natural Origin:	5
	Adulteration of drugs of natural origin. Evaluation by organoleptic,	
	microscopic, physical, chemical and biological methods and properties.	
	Quantitative microscopy of crude drugs including lycopodium spore method,	
	leafconstants, camera lucida and diagrams of microscopic objects to scale	
	with camera lucida.	
2	UNIT II	12
2.1	Cultivation, Collection, Processing and storage of drugs of natural	10
	origin:	
	Cultivation and Collection of drugs of natural origin	
	Factors influencing cultivation of medicinal plants.	
	Plant hormones and their applications.	
	Polyploidy, mutation and hybridization with reference to medicinal plants	
2.2	Conservation of medicinal plants	2
3	UNIT III	7

	-	
	Plant tissue culture:	
	Historical development of plant tissue culture, types of cultures, Nutritional	
	requirements, growth and their maintenance.	
	Applications of plant tissue culture in pharmacognosy.	
	Edible vaccines	
4	UNIT IV	10
4.1	Pharmacognosy in various systems of medicine:	3
	Role of Pharmacognosy in allopathy and traditional systems of medicine	
	namely, Ayurveda, Unani, Siddha, Homeopathy and Chinese systems of	
	medicine	
4.2	Introduction to secondary metabolites:	7
	Definition, classification, properties and test for identification of Alkaloids,	
	Glycosides, Flavonoids, Tannins, Volatile oil and Resins	
5	UNIT V - Study of biological source, chemical nature and uses of drugs of	08
	natural origin containing following drugs	
	(a) Plant Products:	3
	Fibers - Cotton, Jute, Hemp	
	Hallucinogens, Teratogens, Natural allergens	
	(b) Primary metabolites:	3
	General introduction, detailed study with respect to chemistry, sources,	
	preparation, evaluation, preservation, storage, therapeutic used and	
	commercial utility as Pharmaceutical Aids and/or Medicines for the	
	following Primary metabolites:	
	(c) Carbohydrates: Acacia, Agar, Tragacanth, Honey	
	(d) Proteins and Enzymes: Gelatin, casein, proteolytic enzymes (Papain,	
	bromelain, serratiopeptidase, urokinase, streptokinase, pepsin).	
	(e) Lipids(Waxes, fats, fixed oils): Castor oil, Chaulmoogra oil, Wool Fat,	2
	Bees Wax	
	(f) Marine Drugs:	
	Novel medicinal agents from marine sources	
		4.5
	Total	45

- 1. W.C.Evans, Trease and Evans Pharmacognosy, 16th edition, W.B. Sounders & Co., London, 2009.
- 2. Tyler, V.E., Brady, L.R. and Robbers, J.E., Pharmacognosy, 9th Edn., Lea and Febiger, Philadelphia, 1988.

- 3. Text Book of Pharmacognosy by T.E. Wallis
- 4. Mohammad Ali. Pharmacognosy and Phytochemistry, CBS Publishers & Distribution, New Delhi.
- 5. Text book of Pharmacognosy by C.K. Kokate, Purohit, Gokhlae (2007), 37th Edition, Nirali Prakashan, New Delhi.
- 6. Herbal drug industry by R.D. Choudhary (1996), Ist Edn, Eastern Publisher, New Delhi.
- 7. Essentials of Pharmacognosy, Dr.SH.Ansari, IInd edition, Birla publications, New Delhi, 2007
- 8. Practical Pharmacognosy: C.K. Kokate, Purohit, Gokhlae
- 9. Anatomy of Crude Drugs by M.A. Iyengar

BP406P MEDICINAL CHEMISTRY – I (Practical) 3 L + 1T / Week

I Preparation of drugs/intermediates

- 1 1,3-pyrazole
- 2 1,3-oxazole
- 3 Benzimidazole
- 4 Benztriazole
- 5 2,3- diphenyl quinoxaline
- 6 Benzocaine
- 7 Phenytoin
- 8 Phenothiazine
- 9 Barbiturate

II Assay of drugs

- 1 Chlorpromazine
- 2 Phenobarbitone
- 3 Atropine
- 4 Ibuprofen
- 5 Aspirin
- 6 Furosemide

III Determination of Partition coefficient for any two drugs

Recommended Books (Latest Editions)

- 1. Wilson and Giswold's Organic medicinal and Pharmaceutical Chemistry.
- 2. Foye's Principles of Medicinal Chemistry.
- 3. Burger's Medicinal Chemistry, Vol I to IV.
- 4. Introduction to principles of drug design- Smith and Williams.
- 5. Remington's Pharmaceutical Sciences.
- 6. Martindale's extra pharmacopoeia.
- 7. Organic Chemistry by I.L. Finar, Vol. II.
- 8. The Organic Chemistry of Drug Synthesis by Lednicer, Vol. 1-5.
- 9. Indian Pharmacopoeia.
- 10. Text book of practical organic chemistry- A. I. Vogel.

- 1. Determination of particle size, particle size distribution using sieving method
- 2. Determination of particle size, particle size distribution using Microscopic method
- 3. Determination of bulk density, true density and porosity
- 4. Determine the angle of repose and influence of lubricant on angle of repose
- 5. Determination of viscosity of liquid using Ostwald's viscometer
- 6. Determination sedimentation volume with effect of different suspending agent
- 7. Determination sedimentation volume with effect of different concentration of single suspending agent
- 8. Determination of viscosity of semisolid by using Brookfield viscometer
- 9. Determination of reaction rate constant first order.
- 10. Determination of reaction rate constant second order
- 11. Accelerated stability studies

Recommended Books: (Latest Editions)

- 1. Physical Pharmacy by Alfred Martin, Sixth edition
- 2. Experimental pharmaceutics by Eugene, Parott.
- 3. Tutorial pharmacy by Cooper and Gunn.
- 4. Stocklosam J. Pharmaceutical calculations, Lea & Febiger, Philadelphia.
- 5. Liberman H.A, Lachman C., Pharmaceutical Dosage forms, Tablets, Volume-1 to 3, Marcel Dekkar Inc.
- 6. Liberman H.A, Lachman C, Pharmaceutical dosage forms. Disperse systems, volume 1.
- 7. 2. 3. Marcel Dekkar Inc.
- 8. Physical Pharmaceutics by Ramasamy C, and Manavalan R.

BP408P PHARMACOLOGY I (Practical) 4Hours/Week

- 1. Introduction to experimental pharmacology.
- 2. Commonly used instruments in experimental pharmacology.
- 3. Study of common laboratory animals.
- 4. Maintenance of laboratory animals as per CPCSEA guidelines.
- 5. Common laboratory techniques. Blood withdrawal, serum and plasma separation, anesthetics and euthanasia used for animal studies.
- 6. Study of different routes of drugs administration in mice/rats.

- 7. Study of effect of hepatic microsomal enzyme inducers on the phenobarbitone sleeping time in mice.
- 8. Effect of drugs on ciliary motility of frog oesophagus
- 9. Effect of drugs on rabbit eye.
- 10. Effects of skeletal muscle relaxants using rota-rod apparatus.
- 11. Effect of drugs on locomotor activity using actophotometer.
- 12. Anticonvulsant effect of drugs by MES and PTZ method.
- 13. Study of stereotype and anti-catatonic activity of drugs on rats/mice.
- 14. Study of anxiolytic activity of drugs using rats/mice.
- 15. Study of local anesthetics by different methods

Note: All laboratory techniques and animal experiments are demonstrated by simulated experiments by softwares and videos

Recommended Books (Latest Editions)

- 1. Rang H. P., Dale M. M., Ritter J. M., Flower R. J., Rang and Dale's Pharmacology, Churchil Livingstone Elsevier
- 2. Katzung B. G., Masters S. B., Trevor A. J., Basic and clinical pharmacology, Tata Mc Graw-Hill
- 3. Goodman and Gilman's, The Pharmacological Basis of Therapeutics
- 4. Marry Anne K. K., Lloyd Yee Y., Brian K. A., Robbin L.C., Joseph G. B., Wayne A. K., Bradley R.W., Applied Therapeutics, The Clinical use of Drugs, The Point Lippincott Williams & Wilkins
- 5. Mycek M.J, Gelnet S.B and Perper M.M. Lippincott's Illustrated Reviews- Pharmacology
- 6. K.D.Tripathi. Essentials of Medical Pharmacology, JAYPEE Brothers Medical Publishers (P) Ltd, New Delhi.
- 7. Sharma H. L., Sharma K. K., Principles of Pharmacology, Paras medical publisher
- 8. Modern Pharmacology with clinical Applications, by Charles R.Craig& Robert,
- 9. Ghosh MN. Fundamentals of Experimental Pharmacology. Hilton & Company, Kolkata.
- 10. Kulkarni SK. Handbook of experimental pharmacology. VallabhPrakashan,

BP409P PHARMACOGNOSY AND PHYTOCHEMISTRY I (Practical)

4 Hours/Week

- 1. Analysis of crude drugs by chemical tests: (i)Tragaccanth (ii) Acacia (iii)Agar (iv) Gelatin (v) starch (vi) Honey (vii) Castor oil
- 2. Determination of stomatal number and index
- 3. Determination of vein islet number, vein islet termination and paliside ratio.
- 4. Determination of size of starch grains, calcium oxalate crystals by eye piece micrometer
- 5. Determination of Fiber length and width
- 6. Determination of number of starch grains by Lycopodium spore method
- 7. Determination of Ash value
- 8. Determination of Extractive values of crude drugs
- 9. Determination of moisture content of crude drugs
- 10. Determination of swelling index and foaming

Recommended Books: (Latest Editions)

- 1. W.C.Evans, Trease and Evans Pharmacognosy, 16th edition, W.B. Sounders & Co., London, 2009.
- 2. Tyler, V.E., Brady, L.R. and Robbers, J.E., Pharmacognosy, 9th Edn., Lea and Febiger, Philadelphia, 1988.
- 3. Text Book of Pharmacognosy by T.E. Wallis
- 4. Mohammad Ali. Pharmacognosy and Phytochemistry, CBS Publishers & Distribution, New Delhi.
- 5. Text book of Pharmacognosy by C.K. Kokate, Purohit, Gokhlae (2007), 37th Edition, Nirali Prakashan, New Delhi.
- 6. Herbal drug industry by R.D. Choudhary (1996), Ist Edn, Eastern Publisher, New Delhi.
- 7. Essentials of Pharmacognosy, Dr.SH.Ansari, IInd edition, Birla publications, New Delhi, 2007
- 8. Practical Pharmacognosy: C.K. Kokate, Purohit, Gokhlae
- 9. Anatomy of Crude Drugs by M.A. Iyengar

SEMESTER V

BP501T MEDICINAL CHEMISTRY – II (Theory)

3L + 1T / Week

Scope: This subject is designed to impart fundamental knowledge on the structure, chemistry and therapeutic value of drugs. The subject emphasizes on structure activity relationships of drugs, importance of physicochemical properties and metabolism of drugs. The syllabus also emphasizes on chemical synthesis of important drugs under each class.

Objectives: Upon completion of the course the student shall be able to

- 1. Understand the chemistry of drugs with respect to their pharmacological activity
- 2. Understand the drug metabolic pathways, adverse effect and therapeutic value of drugs
- 3. Know the Structural Activity Relationship of different class of drugs
- 4. Study the chemical synthesis of selected drugs

Sr.	Content:	Hours
No.	Study of the development of the following classes of drugs, Classification,	
	mechanism of action, uses of drugs mentioned in the course, Structure	
	activity relationship of selective class of drugs as specified in the course and	
	synthesis of drugs superscripted (*)	
1	UNIT I	10
1.1	Antihistaminic agents: Histamine, receptors and their distribution in	4
	thehumanbody	
	H ₁ –antagonists: Diphenhydramine hydrochloride*,	
	Dimenhydrinate, Doxylamines cuccinate, Clemastine fumarate,	
	Diphenylphyraline hydrochloride, Tripelenamine hydrochloride,	
	Chlorcyclizine hydrochloride, Meclizine hydrochloride, Buclizine	
	hydrochloride, Chlorpheniramine maleate, Triprolidine hydrochloride*,	
	Phenidamine tartarate, Promethazine hydrochloride*, Trimeprazine tartrate,	
	Cyproheptadine hydrochloride, Azatidine maleate, Astemizole, Loratadine,	
	Cetirizine, Levocetrazine Cromolyn sodium	
	H₂-antagonists: Cimetidine*, Famotidine, Ranitidin.	
1.2	Gastric Proton pump inhibitors: Omeprazole, Lansoprazole,	1
	Rabeprazole, Pantoprazole	
1.3	Anti-neoplastic agents:	5
	Alkylatingagents: Meclorethamine* Cyclophosphamide, Melphalan,	
	Chlorambucil, Busulfan, Thiotepa	
	Antimetabolites: Mercaptopurine*, Thioguanine, Fluorouracil,	
	Floxuridine, Cytarabine, Methotrexate*, Azathioprine	
	Antibiotics: Dactinomycin, Daunorubicin, Doxorubicin, Bleomycin	

	Plant products: Etoposide, Vinblastin sulphate, Vincristin sulphate	
	Miscellaneous: Cisplatin, Mitotane.	
2	UNIT II	10
2.1	Anti-anginal:	7
	Vasodilators: Amyl nitrite, Nitroglycerin*, Pentaerythritol tetranitrate,	
	Isosorbidedinitrite*, Dipyridamole.	
	Calcium channel blockers: Verapamil, Bepridil hydrochloride,	
	Diltiazemhydrochloride, Nifedipine, Amlodipine, Felodipine, Nicardipine,	
	Nimodipine.	
	Diuretics:	
	Carbonic anhydrase inhibitors: Acetazolamide*, Methazolamide,	
	Dichlorphenamide. Thiazides: Chlorthiazide*, Hydrochlorothiazide, Hydroflumethiazide,	
	Cyclothiazide Cyclothiazide Cyclothiazide	
	Loop diuretics: Furosemide*, Bumetanide, Ethacrynic acid.	
	Potassium sparing Diuretics: Spironolactone, Triamterene, Amiloride.	
	Osmotic Diuretics: Mannitol	
2.2	Anti-hypertensive Agents: Timolol, Captopril, Lisinopril, Enalapril,	3
	Benazeprilhydrochloride, Quinapril hydrochloride, Methyldopate	_
	hydrochloride,* Clonidine hydrochloride, Guanethidine monosulphate,	
	Guanabenz acetate, Sodium nitroprusside, Diazoxide, Minoxidil, Reserpine,	
	Hydralazine hydrochloride.	
3	UNIT III	10
3.1	Anti-arrhythmic Drugs: Quinidine sulphate, Procainamide	4
	hydrochloride, Disopyramide phosphate*, Phenytoin sodium, Lidocaine	
	hydrochloride, Tocainide hydrochloride, Mexiletine hydrochloride,	
	Lorcainide hydrochloride, Amiodarone, Sotalol.	
3.2	Anti-hyperlipidemic agents: Clofibrate, Lovastatin, Cholesteramine	2
	andCholestipol	
3.3	Coagulant & Anticoagulants: Menadione, Acetomenadione,	2
	Warfarin*, Anisindione, clopidogrel	
3.4	Drugs used in Congestive Heart Failure: Digoxin	2
	Digitoxin, Nesiritide, Bosentan, Tezosentan.	
4	UNIT IV	08
4.1	Drugs acting on Endocrine system	2
4.5	Nomenclature, Stereochemistry and metabolism of steroids	
4.2	Sex hormones: Testosterone, Nandralone, Progestrones, Oestriol,	1
	Oestradiol,Oestrione, Diethyl stilbestrol.	

4.3	Drugs for erectile dysfunction: Sildenafil, Tadalafil	1
4.4	Oral contraceptives: Mifepristone, Norgestril, Levonorgestrol	1
4.5	Corticosteroids: Cortisone, Hydrocortisone, Prednisolone,	2
	Betamethasone, Dexamethasone	
4.6	Thyroid and antithyroid drugs: L-Thyroxine, L-Thyronine,	1
	Propylthiouracil, Methimazole.	
5	UNIT V	07
5.1	Antidiabetic agents:	2
	Insulin and its preparations	
	Sulfonyl ureas: Tolbutamide*, Chlorpropamide, Glipizide, Glimepiride.	
	Biguanides: Metformin.	
	Thiazolidinediones: Pioglitazone, Rosiglitazone.	
	Meglitinides: Repaglinide, Nateglinide.	
	Glucosidase inhibitors: Acrabose, Voglibose.	
5.2	Local Anesthetics: SAR of Local anesthetics	5
	Benzoic Acid derivatives; Cocaine, Hexylcaine, Meprylcaine,	
	Cyclomethycaine, Piperocaine.	
	Amino Benzoic acid derivatives: Benzocaine*, Butamben, Procaine*,	
	Butacaine, Propoxycaine, Tetracaine, Benoxinate.	
	Lidocaine/Anilide derivatives: Lignocaine, Mepivacaine, Prilocaine,	
	Etidocaine.	
	Miscellaneous: Phenacaine, Diperodon, Dibucaine.*	
	Total	45

- 1. Wilson and Giswold's Organic medicinal and Pharmaceutical Chemistry.
- 2. Foye's Principles of Medicinal Chemistry.
- 3. Burger's Medicinal Chemistry, Vol I to IV.
- 4. Introduction to principles of drug design- Smith and Williams.
- 5. Remington's Pharmaceutical Sciences.
- 6. Martindale's extra pharmacopoeia.
- 7. Organic Chemistry by I.L. Finar, Vol. II.
- 8. The Organic Chemistry of Drug Synthesis by Lednicer, Vol. 1to 5.
- 9. Indian Pharmacopoeia.
- 10. Text book of practical organic chemistry- A.I.Vogel.

BP502T INDUSTRIAL PHARMACY I (Theory) 3 L + 1T / Week

Scope: Course enables the student to understand and appreciate the influence ofpharmaceutical additives and various pharmaceutical dosage forms on the performance of the drug product.

Objectives: Upon completion of the course the student shall be able to

- 1. Know the various pharmaceutical dosage forms and their manufacturing techniques.
- 2. Know various considerations in development of pharmaceutical dosage forms
- 3. Formulate solid, liquid and semisolid dosage forms and evaluate them for their quality

Sr.	Content	Hours
No.		_
1	UNIT I - Preformulation Studies	7
1.1	Introduction to preformulation, goals and objectives, study ofphysicochemical characteristics of drug substances	1
1.2	Physical properties: Physical form (crystal & amorphous), particle size, shape, flowproperties, solubility profile (pKa, pH, partition coefficient), polymorphism	2
1.3	Chemical Properties: Hydrolysis, oxidation, reduction, racemisation, polymerizationBCS classification of drugs	2
1.4	Application of preformulation considerations in the development of solid, liquid oral and parenteral dosage forms and its impact on stability of dosage forms.	2
2	UNIT II	10
2.1	Tablets	8
	 a.Introduction, ideal characteristics of tablets, classification of tablets. Excipients, Formulation of tablets, granulation methods, compression and processing problems. Equipments and tablet tooling. b.Tablet coating: Types of coating, coating materials, formulation of coating composition, methods of coating, equipment employed and defects in coating. c.Quality control tests: In process and finished product tests 	
2.2	Liquid orals: Formulation and manufacturing consideration of solutions,	2
	suspensions and emulsions; Filling and packaging; evaluation of liquid orals official in pharmacopoeia	
3	UNIT III	8
3.1	Hard gelatin capsules: Introduction, Extraction of gelatin and production of	3

	hardgelatin capsule shells. size of capsules, Filling, finishing and special techniques of formulation of hard gelatin capsules. In process and final	
	product quality control tests for capsules.	
3.2	Soft gelatin capsules: Nature of shell and capsule content, size	3
	ofcapsules,importance of base adsorption and minimum/gram factors,	
	production, in process and final product quality control tests. Packing,	
	storage and stability testing of soft gelatin capsules	
3.3	Pellets: Introduction, formulation requirements, pelletization process,	2
	equipments formanufacture of pellets	
4	UNIT IV	10
4.1	Definition, types, advantages and limitations. Preformulation factors and	2
	essential requirements, vehicles, additives, importance of isotonicity	
4.2	Production procedure, production facilities and controls.	1
4.3	Formulation of injections, sterile powders, emulsions, suspensions, large	3
	volume parenterals and lyophilized products, Sterilization.	
4.4	Containers and closures selection, filling and sealing of ampoules, vials and	1
	infusion fluids. Quality control tests	
4.5	Ophthalmic Preparations: Introduction, formulation considerations;	3
	formulation of eyedrops, eye ointments and eye lotions; methods of	
	preparation; labeling, containers; evaluation of ophthalmic preparations	
5	UNIT V	10
5.1	Cosmetics: Formulation and preparation of the following cosmetic	3
	preparations:lipsticks, shampoos, cold cream and vanishing cream, tooth	
	pastes, hair dyes and sunscreens.	
5.2	Pharmaceutical Aerosols: Definition, propellants, containers, valves, types	3
	of aerosolsystems; formulation and manufacture of aerosols; Evaluation of	
	aerosols; Quality control and stability studies.	
5.3	Packaging Materials Science: Materials used for packaging of	4
	pharmaceutical products, factors influencing choice of containers, legal and	
	official requirements for containers, stability aspects of packaging materials,	
	quality control tests	
	Total	45

- 1. Pharmaceutical dosage forms Tablets, volume 1 -3 by H.A. Liberman, Leon Lachman &J.B.Schwartz
- 2. Pharmaceutical dosage form Parenteral medication vol- 1&2 by Liberman & Lachman
- 3. Pharmaceutical dosage form disperse system VOL-1 by Liberman & Lachman
- 4. Modern Pharmaceutics by Gilbert S. Banker & C.T. Rhodes, 3rd Edition

- 5. Remington: The Science and Practice of Pharmacy, 20th edition Pharmaceutical Science (RPS)
- 6. Theory and Practice of Industrial Pharmacy by Liberman & Lachman
- 7. Pharmaceutics- The science of dosage form design by M.E.Aulton, Churchill livingstone, **Latest edition**
- 8. Introduction to Pharmaceutical Dosage Forms by H. C.Ansel, Lea &Febiger, Philadelphia, 5thedition, 2005
- 9. Drug stability Principles and practice by Cartensen & C.J. Rhodes, 3rd Edition, Marcel Dekker Series, Vol 107.

BP503T PHARMACOLOGY-II (Theory)

3L + 1T / Week

Scope: This subject is intended to impart the fundamental knowledge on various aspects(classification, mechanism of action, therapeutic effects, clinical uses, side effects and contraindications) of drugs acting on different systems of body and in addition, emphasis on the basic concepts of bioassay.

Objectives: Upon completion of this course the student should be able to:

- 1. Understand the mechanism of drug action and its relevance in the treatment of different diseases
- 2. Demonstrate isolation of different organs/tissues from the laboratory animals by simulated experiments
- 3. Demonstrate the various receptor actions using isolated tissue preparation
- 4. Appreciate correlation of pharmacology with related medical sciences

Sr.	Content	Hours
No.		
1	UNIT I	10
	Pharmacology of drugs acting on cardio vascular system	
	a. Introduction to hemodynamic and electrophysiology of heart.	
	b. Drugs used in congestive heart failure	
	c. Anti-hypertensive drugs.	
	d. Anti-anginal drugs.	
	e. Anti-arrhythmic drugs.	
	f. Anti-hyperlipidemic drugs.	
2	UNIT II	10
2.1	Pharmacology of drugs acting on cardio vascular system	6
	a Drug used in the therapy of shock.	
	b Hematinics, coagulants and anticoagulants.	
	c Fibrinolytics and anti-platelet drugs	
	d Plasma volume expanders	
2.2	Pharmacology of drugs acting on urinary system	4
	a Diuretics	
	b Anti-diuretics	
3	UNIT III	10
	Autocoids and related drugs	
	a Introduction to autacoids and classification	
	b Histamine, 5-HT and their antagonists.	
	c Prostaglandins, Thromboxanes and Leukotrienes.	
	d Angiotensin, Bradykinin and Substance P.	

	e Non-steroidal anti-inflammatory agents	
	f Anti-gout drugs	
	g Antirheumatic drugs	
	h Histamine, 5-HT and their antagonists.	
	8 ,	
	j Angiotensin, Bradykinin and Substance P.	
	k Non-steroidal anti-inflammatory agents	
	l Anti-gout drugs	
	m Antirheumatic drugs	0.0
4	UNIT IV	08
	Pharmacology of drugs acting on endocrine system	
	a Basic concepts in endocrine pharmacology.	
	b Anterior Pituitary hormones- analogues and their inhibitors.	
	c Thyroid hormones- analogues and their inhibitors.	
	d Hormones regulating plasma calcium level-Parathormone, calcitonin	
	and Vitamin-D.	
	e Insulin, Oral Hypoglycemic agents and glucagon.	
	f ACTH and corticosteroids.	
5	UNIT V	07
5.1	Pharmacology of drugs acting on endocrine system	4
	a. Androgens and Anabolic steroids.	
	b. Estrogens, progesterone and oral contraceptives.	
	c. Drugs acting on the uterus.	
5.2	Bioassays	3
	a. Principles and applications of bioassay.	
	b. Types of bioassay	
	c. Bioassay of insulin, oxytocin, vasopressin, ACTH, d-	
	tubocurarine, digitalis, histamine and 5-HT	
	Total	45

- 1. Rang H. P., Dale M. M., Ritter J. M., Flower R. J., Rang and Dale's Pharmacology, Churchil Livingstone Elsevier
- 2. Katzung B. G., Masters S. B., Trevor A. J., Basic and clinical pharmacology, Tata Mc Graw-Hill.
- 3. Goodman and Gilman's, The Pharmacological Basis of Therapeutics
- 4. Marry Anne K. K., Lloyd Yee Y., Brian K. A., Robbin L.C., Joseph G. B., Wayne A. K., Bradley R.W., Applied Therapeutics, The Clinical use of Drugs, The Point Lippincott Williams & Wilkins.

- 5. Mycek M.J, Gelnet S.B and Perper M.M. Lippincott's Illustrated Reviews-Pharmacology.
- 6. K.D.Tripathi. Essentials of Medical Pharmacology, , JAYPEE Brothers Medical Publishers (P) Ltd, New Delhi.
- 7. Sharma H. L., Sharma K. K., Principles of Pharmacology, Paras medical publisher
- 8. Modern Pharmacology with clinical Applications, by Charles R.Craig& Robert.
- 9. Ghosh MN. Fundamentals of Experimental Pharmacology. Hilton & Company, Kolkata.
- 10. Kulkarni SK. Handbook of experimental pharmacology. Vallabh Prakashan

BP504T PHARMACOGNOSY AND PHYTOCHEMISTRY II (Theory)

3L + 1T / Week

Scope: The main purpose of subject is to impart the students the knowledge of how thesecondary metabolites are produced in the crude drugs, how to isolate and identify and produce them industrially. Also this subject involves the study of producing the plants and phytochemicals through plant tissue culture, drug interactions and basic principles of traditional system of medicine

Objectives: Upon completion of the course, the student shall be able:

- 1. to know the modern extraction techniques, characterization and identification of the herbal drugs and phytoconstituents
- 2. to understand the preparation and development of herbal formulation.
- 3. to understand the herbal drug interactions
- 4. to carryout isolation and identification of phytoconstituents

Sr.	Content	Hour
No.		S
1	UNIT I - Metabolic pathways in higher plants and their determination	7
1.1	Brief study of basic metabolic pathways and formation of different	4
	secondary metabolites through these pathways- Shikimic acid pathway,	
	Acetate pathways and Amino acid pathway.	
1.2	Study of utilization of radioactive isotopes in the investigation of Biogenetic	3
	studies.	
2	UNIT II – General introduction, composition, chemistry & chemical	14
	classes, general methods of extraction & analysis, biosources, therapeutic	
	uses and commercial applications of following secondary metabolites:	
2.1	Alkaloids: Vinca, Rauwolfia, Belladonna, Opium,	2
2.2	Phenylpropanoids and Flavonoids: Lignans, Tea, Ruta	2
2.3	Steroids, Cardiac Glycosides & Triterpenoids: Liquorice, Dioscorea,	2
	Digitalis	
2.4	Volatile oils: Mentha, Clove, Cinnamon, Fennel, Coriander,	2
2.5	Tannins: Catechu, Pterocarpus	1
2.6	Resins: Benzoin, Guggul, Ginger, Asafoetida, Myrrh, Colophony	2
2.7	Glycosides: Senna, Aloes, Bitter Almond	1
2.8	Iridoids, Other terpenoids & Naphthaquinones: Gentian, Artemisia,	2
	taxus, carotenoids	
3	UNIT III	6

	Isolation ,identifaction and analysis of phytoconstituents	
	1. Terpenoids: Menthol, Citral, Artemisin	
	2. Glycosides: Glycyrhetinic Acid and Rutin	
	3. Alkaloids: Atropine, Quinene, reserpine, Caffeine	
	4. Resins: Phodophyllatoxin, curcumin	
4	UNIT IV	6
	Industrial production, estimation and utilization of the following	
	phytoconstituents: Forskolin, Sennoside, Artemisinin, Diosgenin, Digoxin,	
	Atropine, Podophyllotoxin, Caffeine, Taxol, Vincristine and Vinblastine	
5	UNIT V	10
	Basics of Phytochemistry	
	Modern methods of extraction, application of latest techniques like	
	Spectroscopy, chromatography and electrophoresis in the isolation,	
	purification and identification of crude drugs	
	Total	45

- 1. W.C.Evans, Trease and Evans Pharmacognosy, 16th edition, W.B. Sounders & Co., London, 2009.
- 2. Mohammad Ali. Pharmacognosy and Phytochemistry, CBS Publishers & Distribution, New Delhi.
- 3. Text book of Pharmacognosy by C.K. Kokate, Purohit, Gokhlae (2007), 37th Edition, Nirali Prakashan, New Delhi.
- 4. Herbal drug industry by R.D. Choudhary (1996), Ist Edn, Eastern Publisher, New Delhi.
- 5. Essentials of Pharmacognosy, Dr.SH.Ansari, IInd edition, Birla publications, New Delhi, 2007
- 6. Herbal Cosmetics by H.Pande, Asia Pacific Business press, Inc, New Delhi.
- 7. A.N. Kalia, Textbook of Industrial Pharmacognosy, CBS Publishers, New Delhi, 2005.
- 8. R Endress, Plant cell Biotechnology, Springer-Verlag, Berlin, 1994.
- 9. Pharmacognosy & Pharmacobiotechnology. James Bobbers, Marilyn KS, VE Tylor.
- 10. The formulation and preparation of cosmetic, fragrances and flavours.
- 11. Remington's Pharmaceutical sciences.
- 12. Text Book of Biotechnology by Vyas and Dixit.
- 13. Text Book of Biotechnology by R.C. Dubey.

BP505T PHARMACEUTICAL JURISPRUDENCE (Theory) 3 L + 1T / Week

Scope: This course is designed to impart basic knowledge on importantlegislations related to the profession of pharmacy in India.

Objectives: Upon completion of the course, the student shall be able to understand:

- 1. The Pharmaceutical legislations and their implications in the development and marketing of pharmaceuticals.
- 2. Various Indian pharmaceutical Acts and Laws
- 3. The regulatory authorities and agencies governing the manufacture and sale of pharmaceuticals
- 4. The code of ethics during the pharmaceutical practice

Sr. No.	Content	Hours
1	UNIT I - Drugs and Cosmetics Act, 1940 and its rules 1945	10
1.1	Objectives, Definitions, Legal definitions of schedules to the act and rules	3
1.2	Import of drugs – Classes of drugs and cosmetics prohibited from import, Import under license or permit. Offences and penalties.	2
1.3	Manufacture of drugs – Prohibition of manufacture and sale of certain drugs	2
1.4	Conditions for grant of license and conditions of license for manufacture of	3
	drugs, Manufacture of drugs for test, examination and analysis, manufacture	
	of new drug, loan license and repacking license	
2	UNIT II - Drugs and Cosmetics Act, 1940 and its rules 1945.	10
2.1	Detailed study of Schedule G, H, M, N, P,T,U, V, X, Y, Part XII B, Sch F & DMR (OA)	4
2.2	Sale of Drugs – Wholesale, Retail sale and Restricted license. Offences and penalties	1
2.3	Labeling & Packing of drugs- General labeling requirements and specimen labels for drugs and cosmetics, List of permitted colors. Offences and penalties.	2
2.4	Administration of the act and rules – Drugs Technical Advisory Board, Central drugs Laboratory, Drugs Consultative Committee, Government drug analysts, Licensing authorities, controlling authorities, Drugs Inspectors	3
3	UNIT III	10
3.1	Pharmacy Act –1948 : Objectives, Definitions, Pharmacy Council of India; its constitution and functions, Education Regulations, State and Joint state pharmacy councils; constitution and functions, Registration of Pharmacists, Offences and Penalties	3
3.2	Medicinal and Toilet Preparation Act –1955: Objectives, Definitions, Licensing, Manufacture In bond and Outside bond, Export of alcoholic	3

	preparations, Manufacture of Ayurvedic, Homeopathic, Patent & Proprietary	
	Preparations. Offences and Penalties	
3.3	Narcotic Drugs and Psychotropic substances Act-1985 and Rules:	4
	Objectives, Definitions, Authorities and Officers, Constitution and Functions	
	of narcotic & Psychotropic Consultative Committee, National Fund for	
	Controlling the Drug Abuse, Prohibition, Control and Regulation, opium	
	poppy cultivation and production of poppy straw, manufacture, sale and	
	export of opium, Offences and Penalties	
4	UNIT IV	08
4.1	Study of Salient Features of Drugs and magic remedies Act and its	2
	rules: Objectives, Definitions, Prohibition of certain advertisements, Classes	
	of Exempted advertisements, Offences and Penalties	
4.2	Prevention of Cruelty to animals Act-1960: Objectives, Definitions,	3
	InstitutionalAnimal Ethics Committee, Breeding and Stocking of Animals,	
	Performance of Experiments, Transfer and acquisition of animals for	
	experiment, Records, Power to suspend or revoke registration, Offences and	
	Penalties	
4.3	National Pharmaceutical Pricing Authority: Drugs Price Control Order	3
	(DPCO)-2013. Objectives, Definitions, Sale prices of bulk drugs, Retail	
	price of formulations, Retail price and ceiling price of scheduled	
	formulations, National List of Essential Medicines (NLEM)	
5	UNIT V	07
5.1	Pharmaceutical Legislations – A brief review, Introduction, Study of drugs	1
	enquirycommittee, Health survey and development committee, Hathi	
	committee and Mudaliar committee	
5.2	Code of Pharmaceutical ethics D efinition, Pharmacist in relation to his	1
	job, trade,medical profession and his profession, Pharmacist's oath	
5.3	Medical Termination of pregnancy act	1
5.4	Right to information Act	1
5.5	Introduction to Intellectual Property Rights (IPR)	3
	Total	45
L		

- 1. Forensic Pharmacy by B. Suresh
- 2. Text book of Forensic Pharmacy by B.M. Mithal
- 3. Hand book of drug law-by M.L. Mehra
- 4. A text book of Forensic Pharmacy by N.K. Jain
- 5. Drugs and Cosmetics Act/Rules by Govt. of India publications.
- 6. Medicinal and Toilet preparations act 1955 by Govt. of India publications.
- 7. Narcotic drugs and psychotropic substances act by Govt. of India publications

- 8. Drugs and Magic Remedies act by Govt. of India publication
- 9. Bare Acts of the said laws published by Government. Reference books (Theory)

BP506P INDUSTRIAL PHARMACY(Practical)4 Hours/week

- 1. Preformulation studies on paracetamol/aspirin/or any other drug
- 2. Preparation and evaluation of Paracetamol tablets
- 3. Preparation and evaluation of Aspirin tablets
- 4. Coating of tablets- film coating of tables/granules
- 5. Preparation and evaluation of Tetracycline capsules
- 6. Preparation of Calcium Gluconate injection
- 7. Preparation of Ascorbic Acid injection
- 8. Quality control test of (as per IP) marketed tablets and capsules
- 9. Preparation of Eye drops/ and Eye ointments
- 10. Preparation of Creams (cold / vanishing cream)
- 11. Evaluation of Glass containers (as per IP)

Recommended Books: (Latest Editions)

- 1. Pharmaceutical dosage forms Tablets, volume 1 -3 by H.A. Liberman, Leon Lachman &J.B.Schwartz
- 2. Pharmaceutical dosage form Parenteral medication vol- 1&2 by Liberman & Lachman
- 3. Pharmaceutical dosage form disperse system VOL-1 by Liberman & Lachman
- 4. Modern Pharmaceutics by Gilbert S. Banker & C.T. Rhodes, 3rd Edition
- 5. Remington: The Science and Practice of Pharmacy, 20th edition Pharmaceutical Science (RPS)
- 6. Theory and Practice of Industrial Pharmacy by Liberman & Lachman
- 7. Pharmaceutics- The science of dosage form design by M.E.Aulton, Churchill livingstone, Latest edition
- 8. Introduction to Pharmaceutical Dosage Forms by H. C.Ansel, Lea &Febiger, Philadelphia, 5thedition, 2005
- 9. Drug stability Principles and practice by Cartensen & C.J. Rhodes, 3rd Edition, Marcel Dekker Series, Vol 107.

- 1. Introduction to *in-vitro* pharmacology and physiological salt solutions.
- 2. Effect of drugs on isolated frog heart.
- 3. Effect of drugs on blood pressure and heart rate of dog.
- 4. Study of diuretic activity of drugs using rats/mice.
- 5. DRC of acetylcholine using frog rectus abdominis muscle.
- 6. Effect of physostigmine and atropine on DRC of acetylcholine using frog rectus abdominis muscle and rat ileum respectively.
- 7. Bioassay of histamine using guinea pig ileum by matching method.
- 8. Bioassay of oxytocin using rat uterine horn by interpolation method.
- 9. Bioassay of serotonin using rat fundus strip by three point bioassay.
- 10. Bioassay of acetylcholine using rat ileum/colon by four point bioassay.
- 11. Determination of PA₂ value of prazosin using rat anococcygeus muscle (by Schilds plot method).
- 12. Determination of PD₂ value using guinea pig ileum.
- 13. Effect of spasmogens and spasmolytics using rabbit jejunum.
- 14. Anti-inflammatory activity of drugs using carrageenan induced paw-edema model.
- 15. Analgesic activity of drug using central and peripheral methods

Note: All laboratory techniques and animal experiments are demonstrated by simulated experiments by softwares and videos

Recommended Books (Latest Editions)

- 1. Rang H. P., Dale M. M., Ritter J. M., Flower R. J., Rang and Dale's Pharmacology, Churchil Livingstone Elsevier
- 2. Katzung B. G., Masters S. B., Trevor A. J., Basic and clinical pharmacology, Tata Mc Graw-Hill.
- 3. Goodman and Gilman's, The Pharmacological Basis of Therapeutics
- 4. Marry Anne K. K., Lloyd Yee Y., Brian K. A., Robbin L.C., Joseph G. B., Wayne A. K., Bradley R.W., Applied Therapeutics, The Clinical use of Drugs, The Point Lippincott Williams & Wilkins.
- 5. Mycek M.J, Gelnet S.B and Perper M.M. Lippincott's Illustrated Reviews-Pharmacology.
- 6. K.D.Tripathi. Essentials of Medical Pharmacology, , JAYPEE Brothers Medical Publishers (P) Ltd, New Delhi.
- 7. Sharma H. L., Sharma K. K., Principles of Pharmacology, Paras medical publisher
- 8. Modern Pharmacology with clinical Applications, by Charles R.Craig& Robert.
- 9. Ghosh MN. Fundamentals of Experimental Pharmacology. Hilton & Company, Kolkata.
- 10. Kulkarni SK. Handbook of experimental pharmacology. Vallabh Prakashan.

BP508P PHARMACOGNOSY AND PHYTOCHEMISTRY II (Practical) 4 Hours/Week

- 1. Morphology, histology and powder characteristics & extraction & detection of: Cinchona, Cinnamon, Senna, Clove, Ephedra, Fennel and Coriander
- 2. Exercise involving isolation & detection of active principles
 - a. Caffeine from tea dust.
 - b. Diosgenin from Dioscorea
 - c. Atropine from Belladonna
 - d. Sennosides from Senna
- 3. Separation of sugars by Paper chromatography
- 4. TLC of herbal extract
- 5. Distillation of volatile oils and detection of phytoconstitutents by TLC
- 6. Analysis of crude drugs by chemical tests: (i) Asafoetida (ii) Benzoin (iii) Colophony (iv) Aloes (v) Myrrh

Recommended Books: (Latest Editions)

- 1. W.C.Evans, Trease and Evans Pharmacognosy, 16th edition, W.B. Sounders & Co., London, 2009.
- 2. Mohammad Ali. Pharmacognosy and Phytochemistry, CBS Publishers & Distribution, New Delhi.
- 3. Text book of Pharmacognosy by C.K. Kokate, Purohit, Gokhlae (2007), 37th Edition, Nirali Prakashan, New Delhi.
- 4. Herbal drug industry by R.D. Choudhary (1996), Ist Edn, Eastern Publisher, New Delhi.
- 5. Essentials of Pharmacognosy, Dr.SH.Ansari, IInd edition, Birla publications, New Delhi, 2007
- 6. Herbal Cosmetics by H.Pande, Asia Pacific Business press, Inc, New Delhi.
- 7. A.N. Kalia, Textbook of Industrial Pharmacognosy, CBS Publishers, New Delhi, 2005.
- 8. R Endress, Plant cell Biotechnology, Springer-Verlag, Berlin, 1994.
- 9. Pharmacognosy & Pharmacobiotechnology. James Bobbers, Marilyn KS, VE Tylor.
- 10. The formulation and preparation of cosmetic, fragrances and flavours.
- 11. Remington's Pharmaceutical sciences.
- 12. Text Book of Biotechnology by Vyas and Dixit.
- 13. Text Book of Biotechnology by R.C. Dubey.

SEMESTER VI

BP601T MEDICINAL CHEMISTRY – III (Theory)

3L + 1T / Week

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

Objectives: Upon completion of the course student shall be able to

- 1. Understand the importance of drug design and different techniques of drug design.
- 2. Understand the chemistry of drugs with respect to their biological activity.
- 3. Know the metabolism, adverse effects and therapeutic value of drugs.
- 4. Know the importance of SAR of drugs.

Sr.	Content	Hours
No.	Study of the development of the following classes of drugs, Classification,	
	mechanism of action, uses of drugs mentioned in the course, Structure	
	activity relationship of selective class of drugs as specified in the course and	
	synthesis of drugs superscripted by (*)	
1	UNIT I	10
	Antibiotics	
	Historical background, Nomenclature, Stereochemistry, Structure activity	
	relationship, Chemical degradation classification and important products of	
	the following classes.	
	(a) β-Lactam antibiotics: Penicillin, Cepholosporins, β Lactamase	
	inhibitors,Monobactams	
	(b) Aminoglycosides: Streptomycin, Neomycin, Kanamycin	
	(c) Tetracyclines: Tetracycline,Oxytetracycline,Chlortetracycline,Minocycl	
	ine, Doxycycline	
2	UNIT II	10
	Antibiotics	
	Historical background, Nomenclature, Stereochemistry, Structure activity	
	relationship, Chemical degradation classification and important products of	
	the following classes.	
	(a) Macrolide: Erythromycin Clarithromycin, Azithromycin.	
	(b) Miscellaneous: Chloramphenicol*, Clindamycin.	

 (c) Prodrugs: Basic concepts and application of prodrugs design. (d) Antimalarials: Etiology of malaria. (e) Quinolines: SAR, Quinine sulphate, Chloroquine* 	
(e) Quinolines: SAR, Quinine sulphate, Chloroquine*	
Amodiaquine, Primaquine phosphate, Pamaquine*, Quinacrine	
hydrochloride, Mefloquine.	
(f) Biguanides and dihydro triazines: Cycloguanil pamoate, Proguanil.	
(g) Miscellaneous: Pyrimethamine, Artesunete, Artemether, Atovoquone.	
3 UNIT III	10
3.1 Anti-tubercular Agents :	3
(a) Synthetic anti tubercular agents: Isoniozid*, Ethionamide	
Ethambutol, Pyrazinamide, Para amino salicylic acid.*	
(b) Anti-tubercular antibiotics: Rifampicin, Rifabutin	
CycloserineStreptomycine, Capreomycin sulphate.	
3.2 Urinary tract anti-infective agents :	3
(a) Quinolones: SAR of quinolones, Nalidixic Acid, Norfloxacin	
Enoxacin, Ciprofloxacin*, Ofloxacin, Lomefloxacin, Sparfloxacin	
Gatifloxacin, Moxifloxacin	
(b) Miscellaneous: Furazolidine, Nitrofurantoin*, Methanamine.	
3.3 Antiviral agents:	4
Amantadine hydrochloride, Rimantadine hydrochloride, Idoxuridine	
trifluoride, Acyclovir*, Gancyclovir, Zidovudine, Didanosine, Zalcitabine	
Lamivudine, Loviride, Delavirding, Ribavirin, Saquinavir, Indinavir Ritonavir.	
4 UNIT IV	08
4.1 Antifungal agents: (a) Antifungal antibiotics: Amphotericin-B, Nystatin, Natamycin	2
(a) Antifungal antibiotics: Amphotericin-B, Nystatin, Natamycin Griseofulvin.	
(b) Synthetic Antifungal agents: Clotrimazole, Econazole	
Butoconazole, Oxiconazole Tioconozole, Miconazole*, Ketoconazole	
Terconazole, Itraconazole, Fluconazole, Naftifine hydrochloride	
Tolnaftate*.	
4.2 Anti-protozoal Agents: Metronidazole*, Tinidazole, Ornidazole	1
Diloxanide, Iodoquinol, Pentamidine Isethionate, Atovaquone, Eflornithine.	
4.3 Anthelmintics: Diethylcarbamazine citrate*, Thiabendazole	1
Mebendazole*, Albendazole, Niclosamide, Oxamniquine, Praziquinal	
Ivermectin	
4.4 Sulphonamides and Sulfones:	4
Historical development, chemistry, classification and SAR of Sulfonamides.	
Sulphamethizole, Sulfisoxazole, Sulphamethizine, Sulfacetamide*	

	Sulphapyridine, Sulfamethoxaole*, Sulphadiazine, Mefenide acetate,	
	Sulfasalazine.	
	Folate reductase inhibitors: Trimethoprim*, Cotrimoxazole	
	Sulfones: Dapsone*.	
5	UNIT V	07
5.1	Introduction to Drug Design	5
	Various approaches used in drug design.	
	Physicochemical parameters used in quantitative structure activity	
	relationship (QSAR) such as partition coefficient, Hammet's electronic	
	parameter, Tafts steric parameter and Hansch analysis.	
	Pharmacophore modeling and docking techniques	
5.2	Combinatorial Chemistry: Concept and applications of	2
	combinatorialchemistry: solid phase and solution phase synthesis.	
	Total	45

- 1. Wilson and Giswold's Organic medicinal and Pharmaceutical Chemistry.
- 2. Foye's Principles of Medicinal Chemistry.
- 3. Burger's Medicinal Chemistry, Vol I to IV.
- 4. Introduction to principles of drug design- Smith and Williams.
- 5. Remington's Pharmaceutical Sciences.
- 6. Martindale's extra pharmacopoeia.
- 7. Organic Chemistry by I.L. Finar, Vol. II.
- 8. The Organic Chemistry of Drug Synthesis by Lednicer, Vol. 1-5.
- 9. Indian Pharmacopoeia.
- 10. Text book of practical organic chemistry- A.I.Vogel.

BP602T PHARMACOLOGY-III (Theory)

3L + 1T / Week

Scope: This subject is intended to impart the fundamental knowledge on various aspects(classification, mechanism of action, therapeutic effects, clinical uses, side effects and contraindications) of drugs acting on respiratory and gastrointestinal system, infectious diseases, immuno-pharmacology and in addition, emphasis on the principles of toxicology and chronopharmacology.

Objectives: Upon completion of this course the student should be able to:

- 1. understand the mechanism of drug action and its relevance in the treatment of different infectious diseases
- 2. comprehend the principles of toxicology and treatment of various poisonings and
- 3. appreciate correlation of pharmacology with related medical sciences

Sr.	Content	Hours
No.		
1	UNIT I	10
1.1	Pharmacology of drugs acting on Respiratory system	5
	a. Anti -asthmatic drugs	
	b. Drugs used in the management of COPD	
	c. Expectorants and antitussives	
	d. Nasal decongestants	
	e. Respiratory stimulants	
1.2	Pharmacology of drugs acting on the Gastrointestinal Tract	5
	a. Antiulcer agents.	
	b. Drugs for constipation and diarrhoea.	
	c. Appetite stimulants and suppressants.	
	d. Digestants and carminatives.	
	e. Emetics and anti-emetics.	
2	UNIT II	10
	Chemotherapy	
	a. General principles of chemotherapy.	
	b. Sulfonamides and cotrimoxazole.	
	c. Antibiotics- Penicillins, cephalosporins, chloramphenicol, macrolides,	
	quinolones and fluoroquinolins, tetracycline and aminoglycosides	
3	UNIT III	10
	Chemotherapy	
	a. Antitubercular agents	
	b. Antileprotic agents	

	c. Antifungal agents	
	d. Antiviral drugs	
	e.Anthelmintics	
	f. Antimalarial drugs	
	g. Antiamoebic agents	
4	UNIT IV	NO.
		08
4.1	Chemotherapy	3
	a. Urinary tract infections and sexually transmitted diseases.	
	b. Chemotherapy of malignancy	
4.2	Immunopharmacology	5
	a. Immunostimulants	
	b. Immunosuppressant	
	c. Protein drugs, monoclonal antibodies, target drugs to	
	d. antigen, biosimilars	
5	UNIT V	07
5.1	Principles of toxicology	6
	a. Definition and basic knowledge of acute, subacute and chronic toxicity.	
	b. Definition and basic knowledge of genotoxicity, carcinogenicity,	
	teratogenicity and mutagenicity	
	c. General principles of treatment of poisoning	
	d. Clinical symptoms and management of barbiturates, morphine,	
	organophosphorus compound and lead, mercury and arsenic	
	poisoning	
5.2	Chronopharmacology	1
	a. Definition of rhythm and cycles.	
	b. Biological clock and their significance leading to chronotherapy	
	Total	45

- 1. Rang H. P., Dale M. M., Ritter J. M., Flower R. J., Rang and Dale's Pharmacology, Churchil Livingstone Elsevier
- 2. Katzung B. G., Masters S. B., Trevor A. J., Basic and clinical pharmacology, Tata Mc Graw-Hill
- 3. Goodman and Gilman's, The Pharmacological Basis of Therapeutics
- 4. Marry Anne K. K., Lloyd Yee Y., Brian K. A., Robbin L.C., Joseph G. B., Wayne A. K., Bradley R.W., Applied Therapeutics, The Clinical use of Drugs. The Point Lippincott Williams & Wilkins
- 5. Mycek M.J, Gelnet S.B and Perper M.M. Lippincott's Illustrated Reviews-Pharmacology

- 6. K.D.Tripathi. Essentials of Medical Pharmacology, , JAYPEE Brothers Medical Publishers (P) Ltd, New Delhi.
- 7. Sharma H. L., Sharma K. K., Principles of Pharmacology, Paras medical publisher Modern Pharmacology with clinical Applications, by Charles R.Craig& Robert,
- 8. Ghosh MN. Fundamentals of Experimental Pharmacology. Hilton & Company, Kolkata,
- 9. Kulkarni SK. Handbook of experimental pharmacology. VallabhPrakashan,
- 10. N.Udupa and P.D. Gupta, Concepts in Chronopharmacology.

BP603T HERBAL DRUG TECHNOLOGY (Theory)

3L + 1T / Week

Scope: This subject gives the student the knowledge of basic understanding of herbal drugindustry, the quality of raw material, guidelines for quality of herbal drugs, herbal cosmetics, natural sweeteners, nutraceutical etc. The subject also emphasizes on Good Manufacturing Practices (GMP), patenting and regulatory issues of herbal drugs

Objectives: Upon completion of this course the student should be able to:

- 1. understand raw material as source of herbal drugs from cultivation to herbal drug product
- 2. know the WHO and ICH guidelines for evaluation of herbal drugs
- 3. know the herbal cosmetics, natural sweeteners, nutraceuticals
- 4. appreciate patenting of herbal drugs, GMP.

Sr. No.	Content	Hours
1	UNIT I	11
1.1	Herbs as raw materials	3
	Definition of herb, herbal medicine, herbal medicinal product, herbal drug	
	preparation Source of Herbs Selection, identification and authentication of	
	herbal materials Processing of herbal raw material	
1.2	Biodynamic Agriculture	3
	Good agricultural practices in cultivation of medicinal plants including	
	Organic farming.	
	Pest and Pest management in medicinal plants: Biopesticides/Bioinsecticides	
1.3	Indian Systems of Medicine	5
	Basic principles involved in Ayurveda, Siddha, Unani and Homeopathy	
	Preparation and standardization of Ayurvedic formulations viz Aristas and	
	Asawas, Gjutika, Churna, Lehya and Bhasma	
2	UNIT II	07
2.1	Neutraceuticals	2
	General aspects, Market, growth, scope and types of products available in	
	the market. Health benefits and role of Nutraceuticals in ailments like	
	Diabetes, CVS diseases, Cancer, Irritable bowel syndrome and various	
	Gastro intestinal diseases.	
2.2	Study of following herbs as health food: Alfaalfa, Chicory, Ginger,	2
	Fenugreek, Garlic, Honey, Amla, Ginseng, Ashwagandha, Spirulina	
2.3	Herbal-Drug and Herb-Food Interactions: General introduction to	3
	interaction and classification. Study of following drugs and their possible side	
	effects and interactions: Hypercium, kava-kava, Ginkobiloba, Ginseng,	
	Garlic, Pepper & Ephedra	

3	UNIT III	10
3.1	Herbal Cosmetics	4
	Sources and description of raw materials of herbal origin used via, fixed oils,	
	waxes, gums colours, perfumes, protective agents, bleaching agents,	
	antioxidants in products such as skin care, hair care and oral hygiene	
	products.	
3.2	Herbal excipients:	3
	Herbal Excipients – Significance of substances of natural origin as excipients	
	- colorants, sweeteners, binders, diluents, viscosity builders, disintegrants,	
	flavors & perfumes.	
3.3	Herbal formulations :	3
	Conventional herbal formulations like syrups, mixtures and tablets and	
	Novel dosage forms like phytosomes	
4	UNIT IV	10
4.1	Evaluation of Drugs WHO & ICH guidelines for the assessment of herbal	2
	drugsStability testing of herbal drugs.	
4.2	Patenting and Regulatory requirements of natural products:	5
	a. Definition of the terms: Patent, IPR, Farmers right, Breeder's right,	
	Bioprospecting and Biopiracy	
	b. Patenting aspects of Traditional Knowledge and Natural Products.	
	Case study of Curcuma & Neem.	
4.3	Regulatory Issues - Regulations in India (ASU DTAB, ASU DCC),	3
	Regulation ofmanufacture of ASU drugs - Schedule Z of Drugs & Cosmetics	
	Act for ASU drugs	
5	UNIT V	07
5.1	General Introduction to Herbal Industry	3
	Herbal drugs industry: Present scope and future prospects.	
	A brief account of plant based industries and institutions involved in work on	
	medicinal and aromatic plants in India.	
5.2	Schedule T-Good Manufacturing Practice of Indian systems of medicine	4
	Components of GMP (Schedule – T) and its objectives	
	Infrastructural requirements, working space, storage area, machinery and	
	equipments, standard operating procedures, health and hygiene,	
	documentation and records.	
	Total	45

- 1. Textbook of Pharmacognosy by Trease & Evans.
- 2. Textbook of Pharmacognosy by Tyler, Brady & Robber.

- 3. Pharmacognosy by Kokate, Purohit and Gokhale
- 4. Essential of Pharmacognosy by Dr.S.H.Ansari
- 5. Pharmacognosy & Phytochemistry by V.D.Rangari
- 6. Pharmacopoeal standards for Ayurvedic Formulation (Council of Research in Indian Medicine & Homeopathy)
- 7. Mukherjee, P.W. Quality Control of Herbal Drugs: An Approach to Evaluation of Botanicals. Business Horizons Publishers, New Delhi, India, 2002.

BP604T BIOPHARMACEUTICS AND PHARMACOKINETICS (Theory)

3L + 1T / Week

Scope: This subject is designed to impart knowledge and skills of Biopharmaceutics and pharmacokinetics and their applications in pharmaceutical development, design of dose and dosage regimen and in solving the problems arised therein.

Objectives: Upon completion of the course student shall be ableto:

- 1. Understand the basic concepts in biopharmaceutics and pharmacokinetics and their significance.
- 2. Use of plasma drug concentration-time data to calculate the pharmacokinetic parameters to describe the kinetics of drug absorption, distribution, metabolism, excretion, elimination.
- 3. To understand the concepts of bioavailability and bioequivalence of drug products and their significance.
- 4. Understand various pharmacokinetic parameters, their significance & applications

Sr.	Content	Hour
No.		S
1	UNIT I -	10
1.1	Introduction to Biopharmaceutics	1
1.2	Absorption; Mechanisms of drug absorption through GIT, factors	5
	influencing drugabsorption though GIT, absorption of drug from Non per	
	oral extra-vascular routes	
1.3	Distribution of Tissue permeability of drugs, binding of drugs, apparent,	4
	volume of drug distribution, protein binding of drugs, factors affecting	
	protein-drug binding. Kinetics of protein binding, Clinical significance of	
	protein binding of drugs	
2	UNIT II	10
2.1	Drug Elimination renal excretion of drugs, factors affecting renal excretion	3
	of drugs,renal clearance, Non-renal routes of drug excretion of drugs	
2.2	Bioavailability and Bioequivalence: Definition andObjectives of	7
	bioavailability studies, absolute andrelative bioavailability, measurement of	
	bioavailability, in-vitro drug dissolution models, in-vitro, in-vivo	
	correlations, bioequivalence studies, methods to enhance the bioavailability	
	of poorly soluble drugs.	
3	UNIT III	10
	Pharmakokinetics: Definition and introduction of pharmacokinetics,	

	compartment models, Non-compartment models, physiological models, One	
	compartment open model. a. Intravenous Injection (Bolus) b. Intravenous	
	infusion, extra vascular administrations, calculations of Ka, K _E , t1/2, Vd,	
	AUC Ka, Clt and CLr- definition methods of elimination, understanding of	
	their significance and application.	
4	UNIT IV	08
	Multicompartment models: Two compartment open model. IV bolus	
	kinetics of Multiple dosing, steady state drug level, calculation of loading	
	and maintenancedose and their significance in clinical setting	
5	UNIT V	07
	Nonlinear Pharmacokinetics: a. Introduction, b. Factors causing	
	Non-linearity.c. Michaelis-menton method of estimating parameters,	
	Biotransformation of drugs	
	Total	45

- 1. Biopharmaceutics and Clinical Pharmacokinetics by, Milo Gibaldi.
- 2. Biopharmaceutics and Pharmacokinetics; By Robert F Notari
- 3. Applied biopharmaceutics and pharmacokinetics, Leon Shargel and Andrew B.C.YU 4th edition, Prentice-Hall Inernational edition. USA
- 4. Bio pharmaceutics and Pharmacokinetics-A Treatise, By D. M. Brahmankar and Sunil B.Jaiswal, Vallabh Prakashan Pitampura, Delhi
- 5. Pharmacokinetics: By Milo Glbaldi Donald, R. Mercel Dekker Inc.
- 6. Hand Book of Clinical Pharmacokinetics, By Milo Gibaldi and Laurie Prescott by ADIS Health Science Press.
- 7. Biopharmaceutics; By Swarbrick
- 8. Clinical Pharmacokinetics, Concepts and Applications: By Malcolm Rowland and
- 9. Thomas, N. Tozen, Lea and Febrger, Philadelphia, 1995.
- 10. Dissolution, Bioavailability and Bioequivalence, By Abdou H.M, Mack, Publishing Company, Pennsylvania 1989.
- 11. Biopharmaceutics and Clinical Pharmacokinetics-An introduction 4th edition Revised and expanded by Rebort F Notari Marcel Dekker Inn, New York and Basel, 1987.
- 12. Remington's Pharmaceutical Sciences, By Mack Publishing Company, Pennsylvnia

BP605T PHARMACEUTICAL BIOTECHNOLOGY (Theory) 3 L + 1T / Week

Scope:

- Biotechnology has a long promise to revolutionize the biological sciences and technology.
- Scientific application of biotechnology in the field of genetic engineering, medicine and fermentation technology makes the subject interesting.
- Biotechnology is leading to new biological revolutions in diagnosis, prevention and cure of diseases, new and cheaper pharmaceutical drugs.
- Biotechnology has already produced transgenic crops and animals and the future promises lot more.
- It is basically a research-based subject.

Objectives: Upon completion of the subject student shall be able to;

- 1. Understanding the importance of Immobilized enzymes in Pharmaceutical Industries
- 2. Genetic engineering applications in relation to production of pharmaceuticals
- 3. Importance of Monoclonal antibodies in Industries
- 4. Appreciate the use of microorganisms in fermentation technology

Sr.	Content	Hours
No.		
1	UNIT I	10
1.1	Brief introduction to Biotechnology with reference to Pharmaceutical Sciences	1
1.2	Enzyme Biotechnology- Methods of enzyme immobilization and applications.	2
1.3	Biosensors- Working and applications of biosensors in Pharmaceutical Industries.	1
1.4	Brief introduction to Protein Engineering.	2
1.5	Use of microbes in industry. Production of Enzymes- General consideration - Amylase, Catalase, Peroxidase, Lipase, Protease, Penicillinase.	2
1.6	Basic principles of genetic engineering.	2
2	UNIT II	10
2.1	Study of cloning vectors, restriction endonucleases and DNA ligase.	2
2.2	Recombinant DNA technology. Application of genetic engineering in	2
	medicine.	
2.3	Application of r DNA technology and genetic engineering in the products:	2
2.4	Interferon b) Vaccines- hepatitis- B c) Hormones- Insulin.	2
2.5	Brief introduction to PCR	2

3	UNIT III	10
	Types of immunity- humoral immunity, cellular immunity	
	a. Structure of Immunoglobulins	
	b. Structure and Function of MHC	
	c. Hypersensitivity reactions, Immune stimulation and Immune suppressions	
	d. General method of the preparation of bacterial vaccines, toxoids, viral	
	vaccine, antitoxins, serum-immune blood derivatives and other products	
	relative to immunity	
	e. Storage conditions and stability of official vaccines	
	f. Hybridoma technology- Production, Purification and Applications	
	g. Blood products and Plasma Substitutes	
4	UNIT IV	08
4.1	Immuno blotting techniques- ELISA, Western blotting, Southern blotting.	2
4.2	Genetic organization of Eukaryotes and Prokaryotes	1
4.3	Microbial genetics including transformation, transduction, conjugation,	2
	plasmids and transposons	
4.4	Introduction to Microbial biotransformation and applications	2
4.5	Mutation.: Types of mutation/ mutants	1
5	UNIT V	07
5.1	Fermentation methods and general requirements, study of media, equipments,	2
	sterilization methods, aeration process, stirring.	
5.2	Large scale production fermenter design and its various controls.	1
5.3	Study of the production of - penicillins, citric acid, Vitamin B12, Glutamic	2
	acid, Griseofulvin	
5.4	Blood product collection, Processing and storage of whole volume blood,	2
	dries=d human plasma, plasma substituents	
	Total	45

Reference Books (Latest Editions to be adopted)

- 1. B.R. Glick and J.J. Pasternak: Molecular Biotechnology: Principles and Applicationsof RecombinantDNA: ASM Press Washington D.C.
- 2. RA Goldshy et. al., : Kuby Immunology.
- 3. J.W. Goding: Monoclonal Antibodies.
- 4. J.M. Walker and E.B. Gingold: Molecular Biology and Biotechnology by Royal Society of Chemistry.
- 5. Zaborsky: Immobilized Enzymes, CRC Press, Degraland, Ohio.
- 6. S.B. Primrose: Molecular Biotechnology (Second Edition) Blackwell Scientific Publication.
- 7. Stanbury F., P., Whitakar A., and Hall J., S., Principles of fermentation technology, 2nd edition, Aditya books Ltd., New Delhi

BP606T PHARMACEUTICAL QUALITY ASSURANCE (Theory)

3L + 1T / Week

Scope: This course deals with the various aspects of quality control and qualityassurance aspects of pharmaceutical industries. It deals with the important aspects like cGMP, QC tests, documentation, quality certifications and regulatory affairs.

Objectives: Upon completion of the course student shall be able to:

- 1. understand the cGMP aspects in a pharmaceutical industry
- 2. appreciate the importance of documentation
- 3. understand the scope of quality certifications applicable to pharmaceutical industries
- 4. understand the responsibilities of QA & QC departments

Sr.	Content	Hours
No.		
1	UNIT I	10
1.1	Quality Assurance and Quality Management concepts: Definition and	4
	concept of Quality control, Quality assurance and GMP	
1.2	Total Quality Management (TQM): Definition, elements, philosophies	2
1.3	ICH Guidelines: purpose, participants, process of harmonization, Brief	2
	overview of QSEM, with special emphasis on Q-series guidelines, ICH	
	stability testing guidelines	
1.4	QbD : Definition, overview, elements of QbD program, tools	1
	ISO 9000 & ISO14000: Overview, Benefits, Elements, steps for	
	registration	
1.5	NABL accreditation : Principles and procedure	1
2	UNIT II	10
2.1	Organization and personnel: Personnel responsibilities, training, hygiene	5
	and personal records. Premises: Design, construction and plant layout,	
	maintenance, sanitation, environmentalcontrol, utilities and maintenance of	
	sterile areas, control of contamination.	
2.2	Equipments and raw materials: Equipments selection, purchase	5
	specifications, maintenance, purchase specifications and maintenance of	
	stores for raw materials	
3	UNIT III	10
3.1	Quality Control: Quality control test for containers, rubber closures and	5
	secondary packing materials	
3.2	Good Laboratory Practices: General Provisions, Organization and	5
j	Personnel, Facilities, Equipment, Testing Facilities Operation, Test and	

	Control Articles, Protocol for Conduct of a Nonclinical Laboratory Study,	
	Records and Reports, Disqualification of Testing Facilities	
4	UNIT IV	08
4.1	Complaints: Complaints and evaluation of complaints, Handling of return	2
	good, recalling andwaste disposal.	
4.2	Document maintenance in pharmaceutical industry: Batch Formula	6
	Record, Master Formula	
	Record, SOP, Quality audit, Quality Review and Quality documentation,	
	Reports and documents, distribution records.	
5	UNIT V	07
5.1	Calibration and Validation: Introduction, definition and general principles	6
	of calibration, qualification and validation, importance and scope of	
	validation, types of validation, validation master plan. Calibration of pH	
	meter, Qualification of UV-Visible spectrophotometer, General principles	
	of Analytical method Validation.	
5.2	Warehousing: Good warehousing practice, materials management	1
	Total	45

Reference Books (Latest Editions to be adopted)

- 1. Quality Assurance Guide by organization of Pharmaceutical Products of India.
- 2. Good Laboratory Practice Regulations, 2nd Edition, Sandy Weinberg Vol. 69.
- 3. Quality Assurance of Pharmaceuticals- A compendium of Guide lines and Related materials Vol I WHO Publications.
- 4. A guide to Total Quality Management- Kushik Maitra and Sedhan K Ghosh
- 5. How to Practice GMP's P P Sharma.
- 6. ISO 9000 and Total Quality Management Sadhank G Ghosh
- 7. The International Pharmacopoeia Vol I, II, III, IV- General Methods of Analysis and Quality specification for Pharmaceutical Substances, Excipients and Dosage forms
- 8. Good laboratory Practices Marcel Deckker Series
- 9. ICH guidelines, ISO 9000 and 14000 guidelines

BP607P MEDICINAL CHEMISTRY- III (Practical) 4 Hours / week

I Preparation of drugs and intermediates

- 1 Sulphanilamide
- 2 7-Hydroxy, 4-methyl coumarin
- 3 Chlorobutanol
- 4 Triphenyl imidazole
- 5 Tolbutamide
- 6 Hexamine

II Assay of drugs

- 1 Isonicotinic acid hydrazide
- 2 Chloroquine
- 3 Metronidazole
- 4 Dapsone
- 5 Chlorpheniramine maleate
- 6 Benzyl penicillin
- III Preparation of medicinally important compounds or intermediates by Microwave irradiation technique
- IV Drawing structures and reactions using chem draw®
- V Determination of physicochemical properties such as logP, clogP, MR, Molecular weight, Hydrogen bond donors and acceptors for class of drugs course content using drug design software Drug likeliness screening (Lipinskies RO5)

Recommended Books (Latest Editions)

- 1. Wilson and Giswold's Organic medicinal and Pharmaceutical Chemistry.
- 2. Foye's Principles of Medicinal Chemistry.
- 3. Burger's Medicinal Chemistry, Vol I to IV.
- 4. Introduction to principles of drug design- Smith and Williams.
- 5. Remington's Pharmaceutical Sciences.
- 6. Martindale's extra pharmacopoeia.
- 7. Organic Chemistry by I.L. Finar, Vol. II.
- 8. The Organic Chemistry of Drug Synthesis by Lednicer, Vol. 1-5.
- 9. Indian Pharmacopoeia.
- 10. Text book of practical organic chemistry- A.I.Vogel.

BP608P PHARMACOLOGY-III (Practical) 4Hrs/Week

- 1. Dose calculation in pharmacological experiments
- 2. Antiallergic activity by mast cell stabilization assay
- 3. Study of anti-ulcer activity of a drug using pylorus ligand (SHAY) rat model and NSAIDS induced ulcer model.
- 4. Study of effect of drugs on gastrointestinal motility
- 5. Effect of agonist and antagonists on guinea pig ileum
- 6. Estimation of serum biochemical parameters by using semi autoanalyser
- 7. Effect of saline purgative on frog intestine
- 8. Insulin hypoglycemic effect in rabbit
- 9. Test for pyrogens (rabbit method)
- 10. Determination of acute oral toxicity (LD50) of a drug from a given data
- 11. Determination of acute skin irritation / corrosion of a test substance
- 12. Determination of acute eye irritation / corrosion of a test substance
- 13. Calculation of pharmacokinetic parameters from a given data
- 14. Biostatistics methods in experimental pharmacology(student's t test, ANOVA)
- 15. Biostatistics methods in experimental pharmacology (Chi square test, Wilcoxon Signed Rank test)
 - *Experiments are demonstrated by simulated experiments/videos

Recommended Books (Latest Editions)

- 1. Rang H. P., Dale M. M., Ritter J. M., Flower R. J., Rang and Dale's Pharmacology, Churchil Livingstone Elsevier
- 2. Katzung B. G., Masters S. B., Trevor A. J., Basic and clinical pharmacology, Tata Mc Graw-Hill
- 3. Goodman and Gilman's, The Pharmacological Basis of Therapeutics
- 4. Marry Anne K. K., Lloyd Yee Y., Brian K. A., Robbin L.C., Joseph G. B., Wayne A. K., Bradley R.W., Applied Therapeutics, The Clinical use of Drugs. The Point Lippincott Williams & Wilkins
- 5. Mycek M.J, Gelnet S.B and Perper M.M. Lippincott's Illustrated Reviews-Pharmacology
- 6. K.D.Tripathi. Essentials of Medical Pharmacology, , JAYPEE Brothers Medical Publishers (P) Ltd, New Delhi.
- 7. Sharma H. L., Sharma K. K., Principles of Pharmacology, Paras medical publisher Modern Pharmacology with clinical Applications, by Charles R.Craig& Robert,
- 8. Ghosh MN. Fundamentals of Experimental Pharmacology. Hilton & Company, Kolkata,

- 9. Kulkarni SK. Handbook of experimental pharmacology. VallabhPrakashan,
- 10. N.Udupa and P.D. Gupta, Concepts in Chronopharmacology.

BP609P HERBAL DRUG TECHNOLOGY (Practical) 4 hours/ week

- 1. To perform preliminary phytochemical screening of crude drugs.
- 2. Determination of the alcohol content of Asava and Arista
- 3. Evaluation of excipients of natural origin
- 4. Incorporation of prepared and standardized extract in cosmetic formulations like creams, lotions and shampoos and their evaluation.
- 5. Incorporation of prepared and standardized extract in formulations like syrups, mixtures and tablets and their evaluation as per Pharmacopoeial requirements.
- 6. Monograph analysis of herbal drugs from recent Pharmacopoeias
- 7. Determination of Aldehyde content
- 8. Determination of Phenol content
- 9. Determination of total alkaloids

Recommended Books: (Latest Editions)

- 1. Textbook of Pharmacognosy by Trease & Evans.
- 2. Textbook of Pharmacognosy by Tyler, Brady & Robber.
- 3. Pharmacognosy by Kokate, Purohit and Gokhale
- 4. Essential of Pharmacognosy by Dr.S.H.Ansari
- 5. Pharmacognosy & Phytochemistry by V.D.Rangari
- 6. Pharmacopoeal standards for Ayurvedic Formulation (Council of Research in Indian Medicine & Homeopathy)
- 7. Mukherjee, P.W. Quality Control of Herbal Drugs: An Approach to Evaluation of Botanicals. Business Horizons Publishers, New Delhi, India, 2002.

SEMESTER VII

BP701T INSTRUMENTAL METHODS OF ANALYSIS (Theory) 3L + 1T / week

Scope: This subject deals with the application of instrumental methods in qualitative andquantitative analysis of drugs. This subject is designed to impart a fundamental knowledge on the principles and instrumentation of spectroscopic and chromatographic technique. This also emphasizes on theoretical and practical knowledge on modern analytical instruments that are used for drug testing.

Objectives: Upon completion of the course the student shall be able to:

- 1. Understand the interaction of matter with electromagnetic radiations and its applications in drug analysis
- 2. Understand the chromatographic separation and analysis of drugs.
- 3. Perform quantitative & qualitative analysis of drugs using various analytical instruments.

Sr.	Content	Hours
No.		
1.	UNIT –I-UV Visible spectroscopy	10
1.1	Electronic transitions, chromophores, auxochromes, spectral shifts, solvent	3
	effect on absorption spectra, Beer and Lambert's law, Derivation and	
	deviations.	
1.2	Instrumentation - Sources of radiation, wavelength selectors, sample cells, detectors-Photo tube, Photomultiplier tube, Photo voltaic cell, Silicon Photodiode.	2
1.3	Applications - Spectrophotometric titrations, Single component and multi component analysis	2
1.4	Fluorimetry	
	Theory, Concepts of singlet, doublet and triplet electronic states, internal	3
	and external conversions, factors affecting fluorescence, quenching,	
	instrumentation and applications	
2	UNIT –II -IR spectroscopy	10
2.1	Introduction, fundamental modes of vibrations in poly atomic molecules,	2
	sample handling, factors affecting vibrations	
2.2	Instrumentation - Sources of radiation, wavelength selectors, detectors -	2
	Golay cell, Bolometer, Thermocouple, Thermister, Pyroelectric detector	
	and applications	
2.3	Flame Photometry-Principle, interferences, instrumentation and	2
	applications	

2.4	Atomic absorption spectroscopy- Principle, interferences,	2
	instrumentation and applications	
2.5	Nepheloturbidometry- Principle, instrumentation and applications	2
3	UNIT -III Introduction to chromatography	10
3.1	Adsorption and partition column chromatography-	4
	Methodology, advantages, disadvantages and applications.	
3.2	Thin layer chromatography- Introduction, Principle, Methodology,	2
	Rf values, advantages, disadvantages and applications.	
3.3	Paper chromatography-Introduction, methodology, development	2
	advantages, disadvantages and applications	
3.4	Electrophoresis—Introduction, factors affecting electrophoretic mobility,	2
	Techniques of paper, gel, capillary electrophoresis, applications	
4	UNIT –IV	8
4.1	Gas chromatography - Introduction, theory, instrumentation,	4
	derivatization,temperature programming, advantages, disadvantages and	
	applications	
4.2	High performance liquid chromatography (HPLC)-Introduction,	4
	theory,instrumentation, advantages and applications.	
5	UNIT -V	7
5.1	Ion exchange chromatography- Introduction, classification, ion	3
	exchange resins, properties, mechanism of ion exchange process,	
	factors affecting ion exchange, methodology and applications	
5.2	Gel chromatography- Introduction, theory, instrumentation and	2
	applications	
5.3	Affinity chromatography- Introduction, theory, instrumentation and	2
	applications	
		45

Reference Books:

- 1. Instrumental Methods of Chemical Analysis by B.K Sharma
- 2. Organic spectroscopy by Y.R Sharma
- 3. Text book of Pharmaceutical Analysis by Kenneth A. Connors
- 4. Vogel's Text book of Quantitative Chemical Analysis by A.I. Vogel
- 5. Practical Pharmaceutical Chemistry by A.H. Beckett and J.B. Stenlake
- 6. Organic Chemistry by I. L. Finar
- 7. Organic spectroscopy by William Kemp
- 8. Quantitative Analysis of Drugs by D. C. Garrett
- 9. Quantitative Analysis of Drugs in Pharmaceutical Formulations by P. D. Sethi
- 10. Spectrophotometric identification of Organic Compounds by Silverstein

BP702T INDUSTRIAL PHARMACY II (Theory) 3L + 1T / week

Scope: This course is designed to impart fundamental knowledge on pharmaceutical product development and translation from laboratory to market

Objectives: Upon completion of the course, the student shall be able to:

- 1. Know the process of pilot plant and scale up of pharmaceutical dosage forms
- 2. Understand the process of technology transfer from lab scale to commercial batch
- 3. Know different Laws and Acts that regulate pharmaceutical industry
- 4. Understand the approval process and regulatory requirements for drug products

Sr.	Content	Hours
No.		
1.	UNIT –I	10
	Pilot plant scale up techniques: General considerations - including	
	significance ofpersonnel requirements, space requirements, raw	
	materials, Pilot plant scale up considerations for solids, liquid orals,	
	semi solids and relevant documentation, SUPAC guidelines,	
	Introduction to Platform technology	
2	UNIT –II	10
	Technology development and transfer: WHO guidelines for	
	Technology Transfer: Terminologies, Technology transfer protocol,	
	Quality risk management, Transfer from R & D to production	
	(Process, packaging and cleaning), Granularity of TT Process (API,	
	excipients, finished products, packing materials) Documentation,	
	Premises and equipments, qualification and validation, quality control,	
	analytical method transfer, Approved regulatory bodies and agencies,	
	Commercialization - practical aspects and problems (case studies),	
	TOT agencies in India - APCTD, NRDC, TIFAC, BCIL, TBSE /	
	SIDBI; Technology of Transfer (TOT) related documentation -	
	confidentiality agreements, licensing, MoUs, legal issues	
3	UNIT –III	10
3.1	Regulatory affairs: Introduction, Historical overview of Regulatory	2
	Affairs, Regulatoryauthorities, Role of Regulatory affairs department,	
	Responsibility of Regulatory Affairs Professionals	
3.2	Regulatory requirements for drug approval: Drug Development	8
	Teams, Non-ClinicalDrug Development, Pharmacology, Drug	
	Metabolism and Toxicology, General considerations of Investigational	
	New Drug (IND) Application, Investigator's Brochure (IB) and New	

	Drug Application (NDA), Clinical research / BE studies, Clinical	
	Research Protocols, Biostatistics in Pharmaceutical Product	
	Development, Data Presentation for FDA Submissions, Management	
	of Clinical Studies.	
4	UNIT –IV	8
	Quality management systems: Quality management & Certifications:	
	Concept of Quality, Total Quality Management, Quality by design, Six	
	Sigma concept, Out of Specifications (OOS), Change control,	
	Introduction to ISO 9000 series of quality systems standards, ISO	
	14000, NABL, GLP	
5	UNIT -V	7
	Indian Regulatory Requirements: Central Drug Standard Control	
	Organization(CDSCO) and State Licensing Authority: Organization,	
	Responsibilities, Common Technical Document (CTD), Certificate of	
	Pharmaceutical Product (COPP), Regulatory requirements and	
	approval procedures for New Drugs.	
		45

Recommended Books: (Latest Editions)

- 1. Regulatory Affairs from Wikipedia, the free encyclopedia modified on 7th April available at http,//en.wikipedia.org/wiki/Regulatory_ Affairs.
- 2. International Regulatory Affairs Updates, 2005. available at http://www.iraup.com/about.php
- 3. Douglas J Pisano and David S. Mantus. Text book of FDA Regulatory Affairs A Guide for Prescription Drugs, Medical Devices, and Biologics' Second Edition.
- 4. Regulatory Affairs brought by learning plus, inc. available at http://www.cgmp.com/ra.htm.

BP703T PHARMACY PRACTICE (Theory) 3L + 1T / week

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 information, and therapeutic drug monitoring for improved patient care. In community pharmacy, students will be learning various skills such as dispensing of drugs, responding to minor ailments by providing suitable safe medication, patient counselling for improved patient care in the community set up.

Objectives: Upon completion of the course, the student shall be able to

- 1. know various drug distribution methods in a hospital
- 2. appreciate the pharmacy stores management and inventory control
- 3. monitor drug therapy of patient through medication chart review and clinical review
- 4. obtain medication history interview and counsel the patients
- 5. identify drug related problems
- 6. detect and assess adverse drug reactions
- 7. interpret selected laboratory results (as monitoring parameters in therapeutics) of specific disease states
- 8. know pharmaceutical care services
- 9. do patient counseling in community pharmacy;
- 10. appreciate the concept of Rational drug therapy.

Sr.	Content	Hours
No.		
1.	UNIT –I	10
1.1	Hospital and it's organization	2
	Definition, Classification of hospital- Primary, Secondary and Tertiary	
	hospitals, Classification based on clinical and non-clinical basis,	
	Organization Structure of a Hospital, and Medical staffs involved in the	
	hospital and their functions.	
1.2	Hospital pharmacy and its organization	2
	Definition, functions of hospital pharmacy, Organization structure,	
	Location, Layout and staff requirements, and Responsibilities and	
	functions of hospital pharmacists.	
1.3	Adverse drug reaction	3
	Classifications - Excessive pharmacological effects, secondary	
	pharmacological effects, idiosyncrasy, allergic drug reactions, genetically	
	determined toxicity, toxicity following sudden withdrawal of drugs, Drug	
	interaction- beneficial interactions, adverse interactions, and	
	pharmacokinetic drug interactions, Methods for detecting drug interactions,	
	spontaneous case reports and record linkage studies, and Adverse drug	

1.4	Community Pharmacy	3
	Organization and structure of retail and wholesale drug store, types and	
	design, Legal requirements for establishment and maintenance of a drug	
	store, Dispensing of proprietary products, maintenance of records of retail	
	and wholesale drug store	
2	UNIT –II	10
2.1	Drug distribution system in a hospital	2
	Dispensing of drugs to inpatients, types of drug distribution systems,	
	charging policy and labelling, Dispensing of drugs to ambulatory patients,	
	and Dispensing of controlled drugs.	
2.2	Hospital formulary	2
	Definition, contents of hospital formulary, Differentiation of hospital	
	formulary and Drug list, preparation and revision, and addition and	
	deletion of drug from hospital formulary.	
2.3	Therapeutic drug monitoring	2
	Need for Therapeutic Drug Monitoring, Factors to be considered during the	
	Therapeutic Drug Monitoring, and Indian scenario for Therapeutic Drug	
	Monitoring.	
2.4	Medication adherence	1
	Causes of medication non-adherence, pharmacist role in the medication	
	adherence, and monitoring of patient medication adherence.	
2.5	Patient medication history interview	1
	Need for the patient medication history interview, medication interview	
•	forms.	
2.6	Community pharmacy management	2
	Financial, materials, staff, and infrastructure requirements.	4.0
3	UNIT -III	10
3.1	Pharmacy and therapeutic committee	2
	Organization, functions, Policies of the pharmacy and therapeutic	
	committee in including drugs into formulary, inpatient and outpatient	
	prescription, automatic stop order, and emergency drug list preparation.	
3.2	Drug information services	1
	Drug and Poison information centre, Sources of drug information,	
2.2	Computerised services, and storage and retrieval of information	
3.3	Patient counseling	2
	Definition of patient counseling; steps involved in patient	
2.1	counseling, and Special cases that require the pharmacist	
3.4	Education and training program in the hospital	3

	Role of pharmacist in the education and training program, Internal and	
	external training program, Services to the nursing homes/clinics, Code	
	of ethics for community pharmacy, and Role of pharmacist in the	
	interdepartmental communication and community health education	
3.5	Prescribed medication order and communication skills	2
	Prescribed medication order- interpretation and legal requirements, and	
	Communication skills- communication with prescribers and patients.	
4	UNIT –IV	8
4.1	Budget preparation and implementation	2
	Budget preparation and implementation	
4.2	Clinical Pharmacy	5
	Introduction to Clinical Pharmacy, Concept of clinical pharmacy, functions	
	and responsibilities of clinical pharmacist, Drug therapy monitoring -	
	medication chart review, clinical review, pharmacist intervention, Ward	
	round participation, Medication history and Pharmaceutical care	
4.3	Over the counter (OTC) sales	1
	Introduction and sale of over the counter, and Rational use of common	
	over the counter medications.	
5	UNIT -V	7
5.1	Drug store management and inventory control	3
	Organisation of drug store, types of materials stocked and storage	
	conditions, Purchase and inventory control: principles, purchase procedure,	
	purchase order, procurement and stocking, Economic order quantity,	
	Reorder quantity level, and Methods used for the analysis of the drug	
	expenditure	
5.2	Investigational use of drugs	2
	Description, principles involved, classification, control, identification, role	
	of hospital pharmacist, advisory committee	
5.3	Interpretation of Clinical Laboratory Tests	2
	Blood chemistry, hematology, and urinalysis	
		45

Recommended Books (Latest Edition):

- 1. Merchant S.H. and Dr. J.S.Quadry. *A textbook of hospital pharmacy*, 4th ed. Ahmadabad: B.S. Shah Prakakshan; 2001.
- 2. Parthasarathi G, Karin Nyfort-Hansen, Milap C Nahata. *A textbook of ClinicalPharmacy Practice- essential concepts and skills*, 1sted. Chennai: OrientLongman Private Limited; 2004.
- 3. William E. Hassan. Hospital pharmacy, 5th ed. Philadelphia: Lea & Febiger; 1986.
- 4. Tipnis Bajaj. *Hospital Pharmacy*, 1st ed. Maharashtra: Career Publications; 2008.

- 5. Scott LT. *Basic skills in interpreting laboratory data*, 4thed. American Society of Health System Pharmacists Inc; 2009.
- 6. Parmar N.S. *Health Education and Community Pharmacy*, 18th ed. India: CBS Publishers & Distributers; 2008.

Journals:

- 1. Therapeutic drug monitoring. ISSN: 0163-4356
- 2. Journal of pharmacy practice. ISSN: 0974-8326
- 3. American journal of health system pharmacy. ISSN: 1535-2900 (online)
- 4. Pharmacy times (Monthly magazine)

BP704T NOVEL DRUG DELIVERY SYSTEMS (Theory) 3L + 1T / week

Scope: This subject is designed to impart basic knowledge on the area of novel drugdelivery systems.

Objectives: Upon completion of the course student shall be able

- 1. To understand various approaches for development of novel drug delivery systems.
- 2. To understand the criteria for selection of drugs and polymers for the development of Novel drug delivery systems, their formulation and evaluation

Sr.	Content	Hours
No.	UNIT I	10
1.1	Controlled drug delivery systems: Introduction, terminology/definitions	7
1.1	and rationale, advantages, disadvantages, selection of drug candidates.	,
	Approaches to design controlled release formulations based on diffusion,	
	dissolution and ion exchange principles. Physicochemical and biological	
	properties of drugs relevant to controlled release formulations	
1.2	Polymers: Introduction, classification, properties, advantages and	3
	application ofpolymers in formulation of controlled release drug delivery	
	systems	
2	UNIT –II	10
2.1	Microencapsulation: Definition, advantages and disadvantages,	3
	microspheres/microcapsules, microparticles, methods of	
	microencapsulation, applications	
2.2	Mucosal Drug Delivery system: Introduction, Principles of bioadhesion	4
	/mucoadhesion, concepts, advantages and disadvantages, transmucosal	
	permeability and formulation considerations of buccal delivery systems	2
2.3	Implantable Drug Delivery Systems:Introduction, advantages and	3
2	disadvantages,concept of implantsand osmotic pump	10
3	UNIT -III	10
3.1	Transdermal Drug Delivery Systems: Introduction, Permeation through	3
	skin, factorsaffecting permeation, permeation enhancers, basic components of TDDS, formulation approaches	
3.2	Gastroretentive drug delivery systems: Introduction, advantages,	3
3.4	disadvantages, approaches for GRDDS – Floating, high density systems,	3
	inflatable and gastroadhesive systems and their applications	
3.3	1 11	4
3.3	Nasopulmonary drug delivery system: Introduction to Nasal and	4

	Pulmonary routes ofdrug delivery, Formulation of Inhalers (dry powder and	
	metered dose), nasal sprays, nebulizers	
4	UNIT –IV	8
	Nanotechnology and its Concepts: Concepts and approaches for targeted drug deliverysystems, advantages and disadvantages, introduction to liposomes, niosomes, nanoparticles, monoclonal antibodies and their applications	
5	UNIT –V	7
5.1	Ocular Drug Delivery Systems: Introduction, intra ocular barriers and methods toovercome –Preliminary study, ocular formulations and ocuserts	5
5.2	Intrauterine Drug Delivery Systems: Introduction, advantages and disadvantages, development of intra uterine devices (IUDs) and	2
		45

Recommended Books: (Latest Editions)

- 1. Y W. Chien, Novel Drug Delivery Systems, 2nd edition, revised and expanded, Marcel Dekker, Inc., New York, 1992.
- 2. Robinson, J. R., Lee V. H. L, Controlled Drug Delivery Systems, Marcel Dekker, Inc., New York, 1992.
- 3. Encyclopedia of Controlled Delivery. Edith Mathiowitz, Published by Wiley Interscience Publication, John Wiley and Sons, Inc, New York. Chichester/Weinheim
- 4. N.K. Jain, Controlled and Novel Drug Delivery, CBS Publishers & Distributors, New Delhi, First edition 1997 (reprint in 2001).
- 5. S.P. Vyas and R.K. Khar, Controlled Drug Delivery -concepts and advances, Vallabh Prakashan, New Delhi, First edition 2002.

Journals

- 1. Indian Journal of Pharmaceutical Sciences (IPA)
- 2. Indian Drugs (IDMA)
- 3. Journal of Controlled Release (Elsevier Sciences)
- 4. Drug Development and Industrial Pharmacy (Marcel & Decker)
- 5. International Journal of Pharmaceutics (Elsevier Sciences)

BP705P INSTRUMENTAL METHODS OF ANALYSIS (Practical) 4 Hours/Week

- Determination of absorption maxima and effect of solvents on absorption maxima of organic compounds
- 2 Estimation of dextrose by colorimetry
- 3 Estimation of sulfanilamide by colorimetry
- 4 Simultaneous estimation of ibuprofen and paracetamol by UV spectroscopy
- 5 Assay of paracetamol by UV- Spectrophotometry
- 6 Estimation of quinine sulfate by fluorimetry
- 7 Study of quenching of fluorescence
- 8 Determination of sodium by flame photometry
- 9 Determination of potassium by flame photometry
- Determination of chlorides and sulphates by nephelo turbidometry
- 11 Separation of amino acids by paper chromatography
- 12 Separation of sugars by thin layer chromatography
- 13 Separation of plant pigments by column chromatography
- 14 Demonstration experiment on HPLC
- 15 Demonstration experiment on Gas Chromatography

Recommended Books (Latest Editions)

- 1. Instrumental Methods of Chemical Analysis by B.K Sharma
- 2. Organic spectroscopy by Y.R Sharma
- 3. Text book of Pharmaceutical Analysis by Kenneth A. Connors
- 4. Vogel's Text book of Quantitative Chemical Analysis by A.I. Vogel
- 5. Practical Pharmaceutical Chemistry by A.H. Beckett and J.B. Stenlake
- 6. Organic Chemistry by I. L. Finar
- 7. Organic spectroscopy by William Kemp
- 8. Quantitative Analysis of Drugs by D. C. Garrett
- 9. Quantitative Analysis of Drugs in Pharmaceutical Formulations by P. D. Sethi
- 10. Spectrophotometric identification of Organic Compounds by Silverstein

BP706PS PRACTICE SCHOOL 12 Hours/Week

Every candidate shall undergo practice school for a period of 150hours evenly distributed throughout the semester. The student shall opt any one of the domains for practice school declared by the program committee from time to time. At the end of the practice school, every student shall submit a printed report (in triplicate) on the practice school he/she attended (not more than 25 pages). Along with the exams of semester VII, the report submitted by the student, knowledge and skills acquired by the student through practice school shall be evaluated by the subject experts at collegeleveland grade point shall be awarded.

SEMESTER VIII

BP801T BIOSTATISITCS AND RESEARCH METHODOLOGY (Theory) 3L + 1T / week

Scope: To understand the applications of Biostatics in Pharmacy. This subject deals withdescriptive statistics, Graphics, Correlation, Regression, logistic regression Probability theory, Sampling technique, Parametric tests, Non-Parametric tests, ANOVA, Introduction to Design of Experiments, Phases of Clinical trials and Observational and Experimental studies, SPSS, R and MINITAB statistical software's, analysing the statistical data using Excel.

Objectives: Upon completion of the course the student shall be able to:

- 1. Know the operation of M.S. Excel, SPSS, R and MINITAB®, DoE (Design of Experiment)
- 2. Know the various statistical techniques to solve statistical problems
- 3. Appreciate statistical techniques in solving the problems

Sr.	Content	Hours
No.		
1	UNIT I	10
1.1	Introduction: Statistics, Biostatistics, Frequency distribution	2
1.2	Measures of central tendency: Mean, Median, Mode- Pharmaceutical	3
	examples	
1.3	Measures of dispersion: Dispersion, Range, standard deviation,	2
	Pharmaceuticalproblems	
1.4	Correlation: Definition, Karl Pearson's coefficient ofcorrelation, Multiple	3
	correlation -Pharmaceuticals examples	
2	UNIT –II	10
2.1	Regression: Curve fitting by the method of least squares, fitting the lines y=	3
	a + bx and $x = a + by$, Multiple regression, standard error of regression–	
	Pharmaceutical Examples	
2.2	Probability: Definition of probability, Binomial distribution, Normal	4
	distribution, Poisson's distribution, properties – problems	
	Sample, Population, large sample, small sample, Null hypothesis,	
	alternative hypothesis, sampling, essence of sampling, types of sampling,	
	Error-I type, Error-II type, Standard error of mean (SEM) - Pharmaceutical	
	examples	
2.3	Parametric test: t-test(Sample, Pooled or Unpaired and Paired), ANOVA,	3
	(One wayand Two way), Least Significance difference	
3	UNIT –III	10

3.1	Non Parametric tests: Wilcoxon Rank Sum Test, Mann-Whitney U test,	2
	Kruskal-Wallistest, Friedman Test	
3.2	Introduction to Research: Need for research, Need for design of	3
	Experiments, Experiential Design Technique, plagiarism	
3.3	Graphs: Histogram, Pie Chart, Cubic Graph, response surface plot, Counter	2
	Plot graph	
3.4	Designing the methodology: Sample size determination and Power of a	3
	study, Reportwriting and presentation of data, Protocol, Cohorts studies,	
	Observational studies, Experimental studies, Designing clinical trial, various	
	phases.	
4	UNIT –IV	8
4.1	Blocking and confounding system for Two-level factorials	2
4.2	Regression modeling: Hypothesis testing in Simple and Multiple	2
	regressionmodels	
4.3	Introduction to Practical components of Industrial and Clinical Trials	4
	problems:Statistical Analysis Using Excel, SPSS, MINITAB®, DESIGN	
	OF EXPERIMENTS, R - Online Statistical Software's to Industrial and	
	Clinical trial approach	
5	UNIT -V	7
5.1	Design and Analysis of experiment- Factorial Design: Definition, 2 ² ,	3
	2 ³ design. Advantage of factorial design	
5.2	Response Surface methodology: Central composite design, Historical	4
	design,Optimization Techniques	
		45

Recommended Books (Latest edition):

- 1. Pharmaceutical statistics- Practical and clinical applications, Sanford Bolton, publisher Marcel Dekker Inc. NewYork.
- 2. Fundamental of Statistics Himalaya Publishing House- S.C.Guptha
- 3. Design and Analysis of Experiments –PHI Learning Private Limited, R. Pannerselvam,
- 4. Design and Analysis of Experiments Wiley Students Edition, Douglas and C. Montgomery

BP802T SOCIAL AND PREVENTIVE PHARMACY 3L + 1T / week

Scope: The purpose of this course is to introduce to students a number of health issues and their challenges. This course also introduced a number of national health programmes. The roles of the pharmacist in these contexts are also discussed.

Objectives: After the successful completion of this course, the student shall be able to:

- 1. Acquire high consciousness/realization of current issuesrelated to health and pharmaceutical problems within the country and worldwide.
- 2. Have a critical way of thinking based on current healthcare development.
 - 3. Evaluate alternative ways of solving problems related tohealth and pharmaceutical issues

Sr.	Content	Hours
No.		
1	UNIT I	10
1.1	Concept of health and disease: Definition, concepts and evaluation of	
	public health.Understanding the concept of prevention and control of	
	disease, social causes of diseases and social problems of the sick.	
1.2	Social and health education: Food in relation to nutrition and health,	
	Balanced diet, Nutritional deficiencies, Vitamin deficiencies, Malnutrition	
	and its prevention.	
1.3	Sociology and health: Socio cultural factors related to health and disease,	
	Impact of urbanization on health and disease, Poverty and health	
1.4	Hygiene and health: personal hygiene and health care; avoidable habits	
2	UNIT –II	10
2.1	Preventive medicine: General principles of prevention and control of	
	diseases such ascholera, SARS, Ebola virus, influenza, acute respiratory	
	infections, malaria, chicken guinea, dengue, lymphatic filariasis,	
	pneumonia, hypertension, diabetes mellitus, cancer, drug addiction-drug	
	substance abuse	
3	UNIT –III	10
	National health programs, its objectives, functioning and outcome of	
	the following: HIV AND AIDS control programme, TB, Integrated disease	
	surveillance program (IDSP), National leprosy control programme, National	
	mental health program, National programme for prevention and control of	
	deafness, Universal immunization programme, National programme for	
	control of blindness, Pulse polio programme	
4	UNIT –IV	8
	National health intervention programme for mother and child, National	

		45
	promotion and education in school.	
	Improvement in rural sanitation, national urban health mission, Health	
	Community services in rural, urban and school health: Functions of PHC,	
5	UNIT –V	7
	elderly, Social health programme; role of WHO in Indian national program	
	Malaria Prevention Program, National programme for the health care for the	
	family welfare programme, National tobacco control programme, National	

Recommended Books (Latest edition):

- 1. Short Textbook of Preventive and Social Medicine, Prabhakara GN, 2nd Edition, 2010, ISBN: 9789380704104, JAYPEE Publications
- 2. Textbook of Preventive and Social Medicine (Mahajan and Gupta), Edited by Roy Rabindra Nath, Saha Indranil, 4th Edition, 2013, ISBN: 9789350901878, JAYPEE Publications
- 3. Review of Preventive and Social Medicine (Including Biostatistics), Jain Vivek, 6th Edition, 2014, ISBN: 9789351522331, JAYPEE Publications
- 4. Essentials of Community Medicine—A Practical Approach, Hiremath Lalita D, Hiremath Dhananjaya A, 2nd Edition, 2012, ISBN: 9789350250440, JAYPEE Publications
- 5. Park Textbook of Preventive and Social Medicine, K Park, 21st Edition, 2011, ISBN-14: 9788190128285, BANARSIDAS BHANOT PUBLISHERS.
- 6. Community Pharmacy Practice, Ramesh Adepu, BSP publishers, Hyderabad

Recommended Journals:

1. Research in Social and Administrative Pharmacy, Elsevier, Ireland

BP803ET PHARMACEUTICAL MARKETING MANAGEMENT(Theory) - ELECTIVE 3L + 1T Hours / week

Scope: The pharmaceutical industry not only needs highly qualified researchers, chemists and, technical people, but also requires skilled managers who can take the industry forward by managing and taking the complex decisions which are imperative for the growth of the industry. The Knowledge and Know-how of marketing management groom the people for taking a challenging role in Sales and Product management.

Objective: The course aims to provide an understanding of marketing conceptsand techniques and their applications in the pharmaceutical industry.

Sr.	Content	Hours
No.		
1	UNIT I	10
1.1	Marketing:	
	Definition, general concepts, and scope of marketing; Distinction between	
	marketing & selling; Marketing environment; Industry and competitive	
	analysis; Analyzing consumer buying behavior; industrial buying behavior	
1.2	Pharmaceutical market:	
	Quantitative and qualitative aspects; size and composition of the market;	
	demographic descriptions and socio-psychological characteristics of the	
	consumer; market segmentation& targeting. Consumer profile; Motivation	
	and prescribing habits of the physician; patients' choice of physician and	
	retail pharmacist. Analyzing the Market; Role of market research.	
2	UNIT –II	10
	Product decision:	
	Meaning, Classification, product line and product mix decisions, product	
	life cycle, product portfolio analysis; product positioning; New product	
	decisions; Product branding, packaging and labeling decisions, Product	
	management in pharmaceutical industry.	
3	UNIT –III	10
	Promotion:	
	Meaning and methods, determinants of promotional mix, promotional	
	budget; An overview of personal selling, advertising, direct mail, journals,	
	sampling, retailing, medical exhibition, public relations, online promotional	
	techniques for OTC Products.	
4	UNIT –IV	8
4.1	Pharmaceutical marketing channels:	

	Designing channel, channel members, selecting the appropriate channel,	
	conflict in channels, physical distribution management: Strategic	
	importance, tasks in physical distribution management.	
4.2	Professional sales representative (PSR):	
	Duties of PSR, purpose of detailing, selection and training, supervising,	
	norms for customer calls, motivating, evaluating, compensation and future	
	prospects of the PSR.	
5	UNIT –V	7
5.1	Pricing:	
	Meaning, importance, objectives, determinants of price; pricing methods	
	and strategies, issues in price management in pharmaceutical industry. An	
	overview of DPCO (Drug Price Control Order) and NPPA (National	
	Pharmaceutical Pricing Authority).	
5.2	Emerging concepts in marketing:	
	Vertical & Horizontal Marketing; Rural Marketing; Consumerism;	
	Industrial Marketing; Global Marketing.	
-		45

Recommended Books: (Latest Editions)

- 1. Philip Kotler and Kevin Lane Keller: Marketing Management, Prentice Hall of India, New Delhi
- 2. Walker, Boyd and Larreche: Marketing Strategy- Planning and Implementation, Tata MC GrawHill, New Delhi.
- 3. Dhruv Grewal and Michael Levy: Marketing, Tata MC Graw Hill
- 4. Arun Kumar and N Menakshi: Marketing Management, Vikas Publishing, India
- 5. Rajan Saxena: Marketing Management; Tata MC Graw-Hill (India Edition)
- 6. Ramaswamy, U.S & Nanakamari, S: Marketing Managemnt: Global Perspective, IndianContext, Macmilan India, New Delhi.
- 7. Shanker, Ravi: Service Marketing, Excell Books, New Delhi
- 8. Subba Rao Changanti, Pharmaceutical Marketing in India (GIFT Excel series) Excel Publications.

BP804ETPHARMACEUTICAL REGULATORY SCIENCE (Theory) - ELECTIVE 3L + 1T Hours / week

Scope: This course is designed to impart the fundamental knowledge on the regulatoryrequirements for approval of new drugs, and drug products in regulated markets of India & other countries like US, EU, Japan, Australia, UK etc. It prepares the students to learn in detail on the regulatory requirements, documentation requirements, and registration procedures for marketing the drug products.

Objectives: Upon completion of the subject student shall be able to;

- 1. Know about the process of drug discovery and development
- 2. Know the regulatory authorities and agencies governing the manufacture and sale of pharmaceuticals
- 3. Know the regulatory approval process and their registration in Indian and international markets

Sr.	Content	Hours
No.		
1	UNIT I	10
	New Drug Discovery and development	
	Stages of drug discovery, Drug development process, pre-clinical	
	studies, non-clinical activities, clinical studies, Innovator and generics,	
	Concept of generics, Generic drug product development.	
2	UNIT –II	10
2.1	Regulatory Approval Process	6
	Approval processes and timelines involved in Investigational New	
	Drug (IND), New Drug Application (NDA), Abbreviated New Drug	
	Application (ANDA) in US. Changes to an approved NDA / ANDA.	
2.2	Regulatory authorities and agencies	4
	Overview of regulatory authorities of United States, European Union,	
	Australia, Japan, Canada (Organization structure and types of	
	applications)	
3	UNIT –III	10
	Registration of Indian drug product in overseas market	
	Procedure for export of pharmaceutical products, Technical	
	documentation, Drug Master Files (DMF), Common Technical	
	Document (CTD), electronic Common Technical Document (eCTD),	
	ASEAN Common Technical Document (ACTD)research	
4	UNIT –IV	8
	Clinical trials	

	Developing clinical trial protocols, Institutional Review Board /	
	Independent Ethics committee - formation and working procedures,	
	Informed consent process and procedures, GCP obligations of	
	Investigators, sponsors & Monitors, Managing and Monitoring clinical	
	trials, Pharmacovigilance - safety monitoring in clinical trials	
5	UNIT –V	7
		-
	Regulatory Concepts	-
	Regulatory Concepts Basic terminologies, guidance, guidelines, regulations, laws and acts,	,
		7
	Basic terminologies, guidance, guidelines, regulations, laws and acts,	,

Recommended books (Latest edition):

- 1. Drug Regulatory Affairs by Sachin Itkar, Dr. N.S. Vyawahare, Nirali Prakashan.
- 2. The Pharmaceutical Regulatory Process, Second Edition Edited by Ira R. Berry and Robert P. Martin, Drugs and the Pharmaceutical Sciences, Vol. 185. Informa Health care Publishers.
- 3. New Drug Approval Process: Accelerating Global Registrations By Richard A Guarino, MD, 5th edition, Drugs and the Pharmaceutical Sciences, Vol. 190.
- 4. Guidebook for drug regulatory submissions / Sandy Weinberg. By John Wiley & Sons. Inc.
- 5. FDA Regulatory Affairs: a guide for prescription drugs, medical devices, and biologics /edited by Douglas J. Pisano, David Mantus.
- 6. Generic Drug Product Development, Solid Oral Dosage forms, Leon Shargel and Isader Kaufer, Marcel Dekker series, Vol.143
- 7. Clinical Trials and Human Research: A Practical Guide to Regulatory Compliance By Fay A. Rozovsky and Rodney K. Adams
- 8. Principles and Practices of Clinical Research, Second Edition Edited by John I. Gallin and Frederick P. Ognibene
- 9. Drugs: From Discovery to Approval, Second Edition By Rick Ng

BP805ET PHARMACOVIGILANCE (Theory) - ELECTIVE 3L + 1T Hours / week

Scope: This paper will provide an opportunity for the student to learn about development ofpharmacovigilance as a science, basic terminologies used in pharmacovigilance, global scenario of Pharmacovigilance, train students on establishing pharmacovigilance programme in an organization, various methods that can be used to generate safety data and signal detection. This paper also develops the skills of classifying drugs, diseases and adverse drug reactions.

Objectives:

At completion of this paper it is expected that students will be able to (know, do, and appreciate):

- 1. Why drug safety monitoring is important?
- 2. History and development of pharmacovigilance
- 3. National and international scenario of pharmacovigilance
- 4. Dictionaries, coding and terminologies used in pharmacovigilance
- 5. Detection of new adverse drug reactions and their assessment
- 6. International standards for classification of diseases and drugs
- 7. Adverse drug reaction reporting systems and communication in pharmacovigilance
- 8. Methods to generate safety data during preclinical, clinical and post approval phases of drugs' life cycle
- 9. Drug safety evaluation in paediatrics, geriatrics, pregnancy and lactation
- 10. Pharmacovigilance Program of India (PvPI) requirement for ADR reporting in India
- 11. ICH guidelines for ICSR, PSUR, expedited reporting, pharmacovigilance planning
- 12. CIOMS requirements for ADR reporting
- 13. Writing case narratives of adverse events and their quality.

Sr. No.	Content	Hours
1	UNIT I	10
1.1	Introduction to Pharmacovigilance	4
	History and development of Pharmacovigilance	
	Importance of safety monitoring of Medicine	
	WHO international drug monitoring programme	
	Pharmacovigilance Program of India(PvPI)	
1.2	Introduction to adverse drug reactions	4
	Definitions and classification of ADRs	
	Detection and reporting	
	Methods in Causality assessment	
	Severity and seriousness assessment	
	Predictability and preventability assessment	

	Management of adverse drug reactions	
1.3	Basic terminologies used in pharmacovigilance	2
	Terminologies of adverse medication related events	
	Regulatory terminologies	
2	UNIT –II	10
2.1	Drug and disease classification	3
	Anatomical, therapeutic and chemical classification of drugs	
	International classification of diseases	
	Daily defined doses	
	International Non proprietary Names for drugs	
2.2	Drug dictionaries and coding in pharmacovigilance	3
	WHO adverse reaction terminologies	
	MedDRA and Standardised MedDRA queries	
	WHO drug dictionary	
	Eudravigilance medicinal product dictionary	
2.3	Information resources in pharmacovigilance	2
	Basic drug information resources	
	Specialised resources for ADRs	
2.4	Establishing pharmacovigilance programme	2
	Establishing in a hospital	
	Establishment & operation of drug safety department in industry	
	Contract Research Organisations (CROs)	
	Establishing a national programme	
3	UNIT –III	10
3.1	Vaccine safety surveillance	3
	Vaccine Pharmacovigilance	
	Vaccination failure	
	Adverse events following immunization	
3.2	Pharmacovigilance methods	5
	Passive surveillance – Spontaneous reports and case series	
	Stimulated reporting	
	Active surveillance – Sentinel sites, drug event monitoring and	
	registries	
	• Comparative observational studies – Cross sectional study, case	
	control study and cohort study	
	Targeted clinical investigations	

 Effective communication in Pharmacovigilance Communication in Drug Safety Crisis management 	
Communication in Drug Safety Crisis management	•
Communicating with Regulatory Agencies, Business Partners,	
Healthcare facilities & Media	
4 UNIT –IV	8
4.1 Statistical methods for evaluating medication safety data Safety	y 3
data generation	
Preclinical phase	
Clinical phase	
Post approval phase	
4.2 ICH Guidelines for Pharmacovigilance	5
Organization and objectives of ICH	
Expedited reporting	
Individual case safety reports	
Periodic safety update reports	
Post approval expedited reporting	
Pharmacovigilance planning	
Good clinical practice in pharmacovigilance studies	
5 Unit V	7
5.1 Pharmacogenomics of adverse drug reactions	3
Genetics related ADR with example focusing PK parameters	
5.2 Drug safety evaluation in special population	2
Paediatrics	
Pregnancy and lactation	
Geriatrics	
5.3 CIOMS	1
CIOMS Working Groups	
CIOMS Form	
5.4 CDSCO (India) and Pharmacovigilance	1
D&C Act and Schedule Y	
Differences in Indian and global pharmacovicilance requirement	nts
Differences in Indian and global pharmacovigilance requirement	

Recommended Books (Latest edition):

- 1. Textbook of Pharmacovigilance: S K Gupta, Jaypee Brothers, Medical Publishers.
- 2. Practical Drug Safety from A to Z By Barton Cobert, Pierre Biron, Jones and Bartlett Publishers.

- 3. Mann's Pharmacovigilance: Elizabeth B. Andrews, Nicholas, Wiley Publishers.
- 4. Stephens' Detection of New Adverse Drug Reactions: John Talbot, Patrick Walle, Wiley Publishers.
- 5. An Introduction to Pharmacovigilance: Patrick Waller, Wiley Publishers.
- 6. Cobert's Manual of Drug Safety and Pharmacovigilance: Barton Cobert, Jones & Bartlett Publishers.
- 7. Textbook of Pharmacoepidemiolog edited by Brian L. Strom, Stephen E Kimmel, Sean Hennessy, Wiley Publishers.
- 8. A Textbook of Clinical Pharmacy Practice -Essential Concepts and Skills:G. Parthasarathi, Karin NyfortHansen,Milap C. Nahata
- 9. National Formulary of India
- 10. Text Book of Medicine by Yashpal Munjal
- 11. Text book of Pharmacovigilance: concept and practice by GP Mohanta and PK Manna
- 12. http://www.whoumc.org/DynPage.aspx?id=105825&mn1=7347&mn2=7259&mn3=7297
- 13. http://www.ich.org/
- 14. http://www.cioms.ch/
- 15. http://cdsco.nic.in/
- 16. http://www.who.int/vaccine_safety/en/
- 17. http://www.ipc.gov.in/PvPI/pv_home.html

BP806ETQUALITY CONTROL AND STANDARDIZATION OF HERBALS (Theory)-ELECTIVE 3L + 1T Hours / week

Scope: In this subject the student learns about the various methods and guidelines forevaluation and standardization of herbs and herbal drugs. The subject also provides anopportunity for the student to learn cGMP, GAP and GLP in traditional system of medicines.

Objectives: Upon completion of the subject student shall be able to;

- 1. know WHO guidelines for quality control of herbal drugs
- 2. know Quality assurance in herbal drug industry
- 3. know the regulatory approval process and their registration in Indian and international markets
- 4. appreciate EU and ICH guidelines for quality control of herbal drugs

Sr.	Content	Hours
No.		
1	UNIT I	10
	Basic tests for drugs – Pharmaceutical substances, Medicinal plants	
	materials and dosage forms	
	WHO guidelines for quality control of herbal drugs.	
	Evaluation of commercial crude drugs intended for use	
2	UNIT –II	10
2.1	Quality assurance in herbal drug industry of cGMP, GAP, GMP	6
	and GLP intraditional system of medicine	
2.2	WHO Guidelines on current good manufacturing Practices (cGMP) for	4
	Herbal Medicines WHO Guidelines on GACP for Medicinal Plants.	
3	UNIT –III	10
	EU and ICH guidelines for quality control of herbal drugs.	
	Research Guidelines for Evaluating the Safety and Efficacy of Herbal	
	Medicines	
4	UNIT –IV	8
	Stability testing of herbal medicines. Application of various	
	chromatographic techniques in standardization of herbal products.	
	Preparation of documents for new drug application and export	
	registration	
	GMP requirements and Drugs & Cosmetics Act provisions.	
5	UNIT –V	7
	Regulatory requirements for herbal medicines.	

products	45
Role of chemical and biological markers in standardization of herbal	
Comparison of various Herbal Pharmacopoeias.	
pharmacovigilance systems	
WHO guidelines on safety monitoring of herbal medicines in	

Recommended Books: (Latest Editions

- 1. Pharmacognosy by Trease and Evans
- 2. Pharmacognosy by Kokate, Purohit and Gokhale
- 3. Rangari, V.D., Text book of Pharmacognosy and Phytochemistry Vol. I, Carrier Pub., 2006.
- 4. Aggrawal, S.S., Herbal Drug Technology. Universities Press, 2002.
- 5. EMEA. Guidelines on Quality of Herbal Medicinal Products/Traditional Medicinal Products,
- 6. Mukherjee, P.W. Quality Control of Herbal Drugs: An Approach to Evaluation of Botanicals. Business Horizons Publishers, New Delhi, India, 2002.
- 7. Shinde M.V., Dhalwal K., Potdar K., Mahadik K. Application of quality control principles to herbal drugs. International Journal of Phytomedicine 1(2009); p. 4-8.
- 8. WHO. Quality Control Methods for Medicinal Plant Materials, World Health Organization, Geneva, 1998. WHO. Guidelines for the Appropriate Use of Herbal Medicines. WHO Regional Publications, Western Pacific Series No 3, WHO Regional office for the Western Pacific, Manila, 1998.
- 9. WHO. The International Pharmacopeia, Vol. 2: Quality Specifications, 3rd edn. World Health Organization, Geneva, 1981.
- 10. WHO. Quality Control Methods for Medicinal Plant Materials. World Health Organization, Geneva, 1999.
- 11. WHO. WHO Global Atlas of Traditional, Complementary and Alternative Medicine. 2 vol. set. Vol. 1 contains text and Vol. 2, maps. World Health Organization, Geneva, 2005.
- 12. WHO. Guidelines on Good Agricultural and Collection Practices (GACP) for Medicinal Plants. World Health Organization, Geneva, 2004

BP807ET COMPUTER AIDED DRUG DESIGN (Theory)- ELECTIVE

3L + 1T / week

Scope: This subject is designed to provide detailed knowledge of rational drug designprocess and various techniques used in rational drug design process.

Objectives: Upon completion of the course, the student shall be able to understand

- 1. Design and discovery of lead molecules
- 2. The role of drug design in drug discovery process
- 3. The concept of QSAR and docking
- 4. Various strategies to develop new drug like molecules.
- 5. The design of new drug molecules using molecular modelling software

Sr.	Content	Hours
No.		
1	UNIT I	10
1.1	Introduction to Drug Discovery and Development	2
	Stages of drug discovery and development	
1.2	Lead discovery and Analog Based Drug Design	4
	Rational approaches to lead discovery based on traditional medicine,	
	Random screening, Non-random screening, serendipitous drug	
	discovery, lead discovery based on drug metabolism, lead discovery	
	based on clinical observation.	
1.3	Analog Based Drug Design: Bioisosterism, Classification,	4
	Bioisostericreplacement. Any three case studies	
2	UNIT –II	10
	Quantitative Structure Activity Relationship (QSAR)	
	SAR versus QSAR, History and development of QSAR, Types of	
	physicochemical parameters, experimental and theoretical approaches	
	for the determination of physicochemical parameters such as Partition	
	coefficient, Hammet's substituent constant and Tafts steric constant.	
	Hansch analysis, Free Wilson analysis, 3D-QSAR approaches like	
	COMFA and COMSIA.	
3	UNIT -III - Molecular Modeling and virtual screening techniques	10
3.1	Virtual Screening techniques: Drug likeness screening, Concept	6
	ofpharmacophore mapping and pharmacophore based Screening,	
3.2	Molecular docking: Rigid docking, flexible docking, manual	4
	docking, Docking based screening. De novo drug design.	
4	UNIT –IV	8
	Informatics & Methods in drug design	

	Introduction to Bioinformatics, chemoinformatics. ADME databases,	
	chemical, biochemical and pharmaceutical databases.	
5	UNIT –V	7
	Molecular Modeling: Introduction to molecular mechanics and quantummechanics. Energy Minimization methods and Conformational Analysis, global conformational minima determination.	
		45

Recommended Books (Latest Editions)

- 1. Robert GCK, ed., "Drug Action at the Molecular Level" University Prak Press Baltimore.
- 2. Martin YC. "Quantitative Drug Design" Dekker, New York.
- 3. Delgado JN, Remers WA eds "Wilson & Gisvolds's Text Book of Organic Medicinal & Pharmaceutical Chemistry" Lippincott, New York.
- 4. Foye WO "Principles of Medicinal chemistry 'Lea & Febiger.
- 5. Koro lkovas A, Burckhalter JH. "Essentials of Medicinal Chemistry" Wiley Interscience.
- 6. Wolf ME, ed "The Basis of Medicinal Chemistry, Burger's Medicinal Chemistry" John Wiley & Sons, New York.
- 7. Patrick Graham, L., An Introduction to Medicinal Chemistry, Oxford University Press
- 8. Smith HJ, Williams H, eds, "Introduction to the principles of Drug Design" Wright Boston.
- 9. Silverman R.B. "The organic Chemistry of Drug Design and Drug Action" Academic Press New York.

BP808ET CELL AND MOLECULAR BIOLOGY (Theory)- ELECTIVE 3L + 1T / week

Scope:

Cell biology is a branch of biology that studies cells – their physiological properties, their
structure, the organelles they contain, interactions with their environment, their life cycle,
division, death and cell function.
This is done both on a microscopic and molecular level.
Cell biology research encompasses both the great diversity of single-celled organisms like
bacteria and protozoa, as well as the many specialized cells in multi-cellular organisms such
as humans, plants, and sponges.

Objectives: Upon completion of the subject student shall be able to;

- 1. Summarize cell and molecular biology history.
- 2. Summarize cellular functioning and composition.
- 3. Describe the chemical foundations of cell biology.
- 4. Summarize the DNA properties of cell biology.
- 5. Describe protein structure and function.
- 6. Describe cellular membrane structure and function.
- 7. Describe basic molecular genetic mechanisms.
- 8. Summarize the Cell Cycle

Sr.	Content	Hours
No.		
1	UNIT I	10
	a) Cell and Molecular Biology: Definitions theory and basics and	
	Applications.	
	b) Cell and Molecular Biology: History and Summation.	
	c) Theory of the Cell? Properties of cells and cell membrane.	
	d) Prokaryotic versus Eukaryotic	
	e) Cellular Reproduction	
	f) Chemical Foundations – an Introduction and Reactions (Types)	
2	UNIT –II	10
	a) DNA and the Flow of Molecular Structure	
	b) DNA Functioning	
	c) DNA and RNA	
	d) Types of RNA	
	e) Transcription and Translation	

3	UNIT –III	10
	a) Proteins: Defined and Amino Acids	
	b) Protein Structure	
	c) Regularities in Protein Pathways	
	d) Cellular Processes	
	e) Positive Control and significance of Protein Synthesis	
4	UNIT –IV	8
	a) Science of Genetics	
	b) Transgenics and Genomic Analysis	
	c) Cell Cycle analysis	
	d) Mitosis and Meiosis	
	e) Cellular Activities and Checkpoints	
5	UNIT –V	7
	a) Cell Signals: Introduction	
	b) Receptors for Cell Signals	
	c) Signaling Pathways: Overview	
	d) Misregulation of Signaling Pathways	
	e) Protein-Kinases: Functioning	
		45

Recommended Books (latest edition):

- 1. W.B. Hugo and A.D. Russel: Pharmaceutical Microbiology, Blackwell Scientific publications, Oxford London.
- 2. Prescott and Dunn., Industrial Microbiology, 4th edition, CBS Publishers & Distributors, Delhi.
- 3. Pelczar, Chan Kreig, Microbiology, Tata McGraw Hill edn.
- 4. Malcolm Harris, Balliere Tindall and Cox: Pharmaceutical Microbiology.
- 5. Rose: Industrial Microbiology.
- 6. Probisher, Hinsdill et al: Fundamentals of Microbiology, 9th ed. Japan
- 7. Cooper and Gunn's: Tutorial Pharmacy, CBS Publisher and Distribution.
- 8. Peppler: Microbial Technology.
- 9. Edward: Fundamentals of Microbiology.
- 10. N.K.Jain: Pharmaceutical Microbiology, Vallabh Prakashan, Delhi
- 11. Bergeys manual of systematic bacteriology, Williams and Wilkins- A Waverly company
- 12. B.R. Glickand J.J. Pasternak: Molecular Biotechnology: Principles and Applications of RecombinantDNA: ASM Press Washington D.C.
- 13. R. A Goldshy et. al.,: Kuby Immunology

Sr.	Content	Hours
No.		
1	UNIT I	10
1.1	Classification of cosmetic and cosmeceutical products	2
	Defination of cosmetics as per Indian and EU regulations, Evolution of	
	cosmoceuticals from cosmetics, cosmetics as quasi and OTC drugs	
1.2	Cosmetic excipients: Surfactants, rheology modifiers, humectants,	8
	emollients,preservatives. Classification and application	
	Skin: Basic structure and function of skin.	
	Hair: Basic structure of hair. Hair growth cycle	
	Oral Cavity: Common problem associated with teeth and gums.	
2	UNIT –II	10
2.1	Principles of formulation and building blocks of skin care	5
	products:	
	Face wash, Moisturizing cream, Cold Cream, Vanishing cream their	
	relative skin sensory, advantages and disadvantages. Application of	
	these products in formulation of cosmecuticals.	
	Principles of formulation and building blocks of Hair care	5
	products:	
	Conditioning shampoo, Hair conditioners, antidandruff shampoo.	
	Hair oils.	
	Chemistry and formulation of Para-phylene diamine based hair dye.	
	Principles of formulation and building blocks of oral care products:	
	Toothpaste for bleeding gums, sensitive teeth. Teeth whitening,	
	Mouthwash	
3	UNIT –III	10
3.1	Sun protection, Classification of Sunscreens and SPF	2
3.2	Role of herbs in cosmetics:	6
	Skin Care: Aloe and turmeric	
	Hair care: Henna and amla.	
	Oral care: Neem and clove	
3.3	Analytical cosmetics: BIS specification and analytical methods for	2
	shampoo, skin-cream and toothpaste	
4	UNIT –IV	8
	Principles of Cosmetic Evaluation: Principles of sebumeter,	
	corneometer. Measurement of TEWL, Skin Color, Hair tensile strength,	

		45
	Antiperspirants and Deodorants- Actives and mechanism of action	
	acne, prickly heat and body odor.	
	Cosmetic problems associated with skin: blemishes, wrinkles,	
	Hair fall causes	
	Cosmetic problems associated with Hair and scalp: Dandruff,	
	Basic understanding of the terms Comedogenic, dermatitis.	
	Oily and dry skin, causes leading to dry skin, skin moisturisation.	
5	UNIT –V	7
	beneits.	
	Hair combing properties Soaps, and syndet bars. Evolution and skin	

References

- 1. Harry's Cosmeticology, Wilkinson, Moore, Seventh Edition, George Godwin.
- 2. Cosmetics Formulations, Manufacturing and Quality Control, P.P. Sharma, 4th Edition, Vandana Publications Pvt. Ltd., Delhi.
- 3. Text book of cosmelicology by Sanju Nanda & Roop K. Khar, Tata Publishers.

BP810ET EXPERIMENTAL PHARMACOLOGY- ELECTIVE 3L + 1T/ week

Scope:This subject is designed to impart the basic knowledge of preclinical studies inexperimental animals including design, conduct and interpretations of results.

Objectives: Upon completion of the course the student shall be able to,

- 1. Appreciate the applications of various commonly used laboratory animals.
- 2. Appreciate and demonstrate the various screening methods used in preclinical research
- 3. Appreciate and demonstrate the importance of biostatistics and research methodology
- 4. Design and execute a research hypothesis independently

Sr.	Content	Hours
No.		
1	UNIT I	8
	Laboratory Animals:	
	Study of CPCSEA and OECD guidelines for maintenance,	
	breeding and conduct of experiments on laboratory animals,	
	Common lab animals: Description and applications of different	
	species and strains of animals. Popular transgenic and mutant	
	animals.	
	• Techniques for collection of blood and common routes of	
	drug administration in laboratory animals, Techniques of blood	
	collection and euthanasia	
2	UNIT –II	13
2.1	Preclinical screening models	6
	a. Introduction: Dose selection, calculation and conversions,	
	preparation of drug solution/suspensions, grouping of animals	
	and importance of sham negative and positive control groups.	
	Rationale for selection of animal species and sex for the study.	
	b. Study of screening animal models for	
	Diuretics, nootropics, anti-Parkinson's, antiasthmatics,	
2.2	Preclinical screening models: for CNS activity- analgesic, antipyretic,	7
	anti-inflammatory, general anaesthetics, sedative and hypnotics,	
	antipsychotic, antidepressant, antiepileptic, antiparkinsonism,	
	alzheimer's disease	
3	UNIT –III	12
	Preclinical screening models: for ANS activity, sympathomimetics,	
	sympatholytics, parasympathomimetics, parasympatholytics, skeletal	
	muscle relaxants, drugs acting on eye, local anaethetics	

4	UNIT –IV	12
4.1	Preclinical screening models: for CVS activity- antihypertensives,	6
	diuretics, antiarrhythmic, antidyslepidemic, anti aggregatory,	
	coagulants, and anticoagulants	
	Preclinical screening models for other important drugs like antiulcer,	
	antidiabetic, anticancer and antiasthmatics.	
4.2	Research methodology and Bio-statistics	6
	Selection of research topic, review of literature, research hypothesis	
	and study design	
	Pre-clinical data analysis and interpretation using Student's 't' test and	
	One-way ANOVA. Graphical representation of data	

Recommended Books (latest edition):

- 1. Fundamentals of experimental Pharmacology-by M.N.Ghosh
- 2. Hand book of Experimental Pharmacology-S.K.Kulakarni
- 3. CPCSEA guidelines for laboratory animal facility.
- 4. Drug discovery and Evaluation by Vogel H.G.
- 5. Drug Screening Methods by Suresh Kumar Gupta and S. K. Gupta
- 6. Introduction to biostatistics and research methods by PSS Sundar Rao and J Richard

BP8011ETADVANCED INSTRUMENTATION TECHNIQUES (Theory) – ELECTIVE3L + 1T / week

Scope: This subject deals with the application of instrumental methods in qualitative andquantitative analysis of drugs. This subject is designed to impart advanced knowledge on the principles and instrumentation of spectroscopic and chromatographic hyphenated techniques. This also emphasizes on theoretical and practical knowledge on modern analytical instruments that are used for drug testing.

Objectives: Upon completion of the course the student shall be able to

- 1. understand the advanced instruments used and its applications in drug analysis
- 2. understand the chromatographic separation and analysis of drugs.
- 3. understand the calibration of various analytical instruments
- 4. know analysis of drugs using various analytical instruments

Sr. No.	Content	Hours
1	UNIT I	10
1.1	Nuclear Magnetic Resonance spectroscopy	
	Principles of H-NMR and C-NMR, chemical shift, factors affecting	
	chemical shift, coupling constant, Spin - spin coupling, relaxation,	
	instrumentation and applications	
1.2	Mass Spectrometry- Principles, Fragmentation, Ionization techniques—	
	Electron impact, chemical ionization, MALDI, FAB, Analyzers-Time of	
	flight and Quadrupole, instrumentation, applications	
2	UNIT –II	10
2.1	Thermal Methods of Analysis: Principles, instrumentation and applications	
	of ThermogravimetricAnalysis (TGA), Differential Thermal Analysis	
	(DTA), Differential Scanning Calorimetry (DSC)	
2.2	X-Ray Diffraction Methods: Origin of X-rays, basic aspects of crystals, X-	
	ray Crystallography, rotating crystal technique, single crystal	
	diffraction, powder diffraction, structural elucidation and applications.	
3	UNIT –III	10
3.1	Calibration and validation-as per ICH and USFDA guidelines	
3.2	Calibration of following Instruments- Electronic balance,	
	UV-Visible spectrophotometer, IR spectrophotometer, Fluorimeter, Flame	
	Photometer, HPLC and GC	
4	UNIT –IV	8
4.1	Radio immune assay:Importance, various components, Principle, different	
	methods, Limitation and Applications of Radio immuno assay	
4.2	Extraction techniques:General principle and procedure involved in the	
	solid phase extraction and liquid-liquid extraction	
5	UNIT-V	7

Hyphenated techniques-LC-MS/MS, GC-MS/MS, HPTLC-MS.	
Total	45

Recommended Books (Latest Editions)

- 1. Instrumental Methods of Chemical Analysis by B.K Sharma
- 2. Organic spectroscopy by Y.R Sharma
- 3. 3. Text book of Pharmaceutical Analysis by Kenneth A. Connors
- 4. Vogel's Text book of Quantitative Chemical Analysis by A.I. Vogel
- 5. Practical Pharmaceutical Chemistry by A.H. Beckett and J.B. Stenlake
- 6. Organic Chemistry by I. L. Finar
- 7. Organic spectroscopy by William Kemp
- 8. Quantitative Analysis of Drugs by D. C. Garrett
- 9. Quantitative Analysis of Drugs in Pharmaceutical Formulations by P. D. Sethi
- 10. Spectrophotometric identification of Organic Compounds by Silverstein

BP8012ET DIETARY SUPPLEMENTS AND NUTRACEUTICALS (Theory) - ELECTIVE

3L + 1T/ Week

Scope: This subject covers foundational topic that are important for understanding the need and requirements of dietary supplements among different groups in the population.

Objective: This module aims to provide an understanding of the concepts behind the theoretical applications of dietary supplements. By the end of the course, students should be able to:

- 1. Understand the need of supplements by the different group of people to maintain healthy life.
- 2. Understand the outcome of deficiencies in dietary supplements.
- 3. Appreciate the components in dietary supplements and the application.
- 4. Appreciate the regulatory and commercial aspects of dietary supplements including health claims.

Sr. No.	Content	Hours
1	UNIT I	10
	a) Definitions of Functional foods, Nutraceuticals and Dietary supplements.	
	Classification of Nutraceuticals, Health problems and diseases that can be	
	prevented or cured by Nutraceuticals i.e. weight control, diabetes, cancer,	
	heart disease, stress, osteoarthritis,	
	hypertension etc.	
	b) Public health nutrition, maternal and child nutrition, nutrition and	
	ageing, nutrition education in community.	
	c) Source, Name of marker compounds and their chemical nature,	
	Medicinal uses and health benefitsof following used as	
	nutraceuticals/functional foods: Spirulina, Soyabean, Ginseng, Garlic,	
	Broccoli, Gingko, Flaxseeds.	
2	UNIT –II	10

	Phytochemicals as nutraceuticals: Occurrence and characteristic features(chemical nature medicinal benefits) of following: a) Carotenoids- α and β-Carotene, Lycopene, Xanthophylls, leutin b) Sulfides: Diallyl sulfides, Allyl trisulfide. c) Polyphenolics: Reservetrol d) Flavonoids- Rutin , Naringin, Quercitin, Anthocyanidins, catechins, Flavones e) Prebiotics / Probiotics.: Fructo oligosaccharides, Lacto bacillum f) Phyto estrogens: Isoflavones, daidzein, Geebustin, lignans g) Tocopherols	
	h) Proteins, vitamins, minerals, cereal, vegetables and beverages as functional foods: oats, wheat bran, rice bran, sea foods, coffee, tea and the like.	
3	UNIT –III	10
	 a) Introduction to free radicals: Free radicals, reactive oxygen species, production of free radicals in cells, damaging reactions of free radicals on lipids, proteins, Carbohydrates, nucleic acids. b) Dietary fibres and complex carbohydrates as functional food ingredients 	
4	UNIT –IV	8
	 a) Free radicals in Diabetes mellitus, Inflammation, Ischemic reperfusion injury, Cancer, Atherosclerosis, Free radicals in brain metabolism and pathology, kidney damage, muscle damage. Free radicals involvement in other disorders. Free radicals theory of ageing. b) Antioxidants: Endogenous antioxidants – enzymatic and nonenzymatic antioxidant defence, Superoxide dismutase, catalase, Glutathione peroxidase, Glutathione Vitamin C, Vitamin E, α- Lipoic acid, melatonin Synthetic antioxidants: Butylated hydroxy Toluene, Butylated hydroxy Anisole. c) Functional foods for chronic disease prevention 	
5	UNIT-V	7
	 a) Effect of processing, storage and interactions of various environmental factors on the potential of nutraceuticals. b) Regulatory Aspects; FSSAI, FDA, FPO, MPO, AGMARK. HACCP and GMPs on Food Safety. Adulteration of foods. c) Pharmacopoeial Specifications for dietary supplements and nutraceuticals. 	

References:

1. Dietetics by Sri Lakshmi

- 2. Role of dietary fibres and neutraceuticals in preventing diseases by K.T Agusti and
- 3. P.Faizal: BSPunblication.
- 4. Advanced Nutritional Therapies by Cooper. K.A., (1996).
- 5. The Food Pharmacy by Jean Carper, Simon & Schuster, UK Ltd., (1988).
- 6. Prescription for Nutritional Healing by James F.Balch and Phyllis A.Balch 2 nd Edn.,
- 7. Avery Publishing Group, NY (1997).
- 8. G. Gibson and C.williams Editors 2000 Functional foods Woodhead Publ.Co.London.
- 9. Goldberg, I. Functional Foods. 1994. Chapman and Hall, New York.
- 10. Labuza, T.P. 2000 Functional Foods and Dietary Supplements: Safety, Good
- 11. Manufacturing Practice (GMPs) and Shelf Life Testing in Essentials of Functional
- 12. Foods M.K. Sachmidl and T.P. Labuza eds. Aspen Press.
- 13. Handbook of Nutraceuticals and Functional Foods, Third Edition (Modern Nutrition)
- 14. Shils, ME, Olson, JA, Shike, M. 1994 Modern Nutrition in Health and Disease.
- 15. Eighth edition. Lea and Febiger

$\label{eq:BP803ETPHARMACEUTICAL PRODUCT DEVELOPMENT} BP803ETPHARMACEUTICAL PRODUCT DEVELOPMENT (Theory) ELECTIVE \\ 3L + 1T/WEEK$

Sr.	Topic	Hours
No.		
1	UNIT I	10
1.1	Introduction to pharmaceutical product development, objectives,	
	regulations related to preformulation, formulation development,	
	stability assessment, manufacturing and quality control testing of	
	different types of dosage forms	10
2	UNIT -II	10
2.1	An advanced study of Pharmaceutical Excipients in pharmaceutical	
	product development with a special reference to the following	
	i. Solvents and solubilizers	
	ii. Cyclodextrins and their applications	
	iii. Non - ionic surfactants and their applications	
	iv. Polyethylene glycols and sorbitols	
	v. Suspending and emulsifying agents	
	vi. Semi solid excipients	
3	UNIT –III	10
3.1	An advanced study of Pharmaceutical Excipients in pharmaceutical	
	product development with a special reference to the following	
	categories	
	i. Tablet and capsule excipients	
	ii. Directly compressible vehicles	
	iii. Coat materials	
	iv. Excipients in parenteral and aerosols products	
	v. Excipients for formulation of NDDS	
	Selection and application of excipients in pharmaceutical formulations	
_	with specific industrial applications	
4	UNIT –IV	8
	Optimization techniques in pharmaceutical product development.A	
	study of various optimization techniques for pharmaceutical product	
	development with specific examples. Optimization by factorial designs	
	and their applications. A study of QbD and its application in pharmaceutical product development.	
5	UNIT-V	7
3	Selection and quality control testing of packaging materials for	
	pharmaceutical product development- regulatory considerations.	
	Total	45
	1000	70

Recommended Books (Latest editions)

- 1. Pharmaceutical Statistics Practical and Clinical Applications by Stanford Bolton, CharlesBon; Marcel Dekker Inc.
- 2. Encyclopedia of Pharmaceutical Technology, edited by James swarbrick, Third Edition,Informa Healthcare publishers.
- 3. Pharmaceutical Dosage Forms, Tablets, Volume II, edited by Herbert A. Lieberman and Leon Lachman; Marcel Dekker, Inc.
- 4. The Theory and Practice of Industrial Pharmacy, Fourth Edition, edited by Roop kKhar, S P Vyas, Farhan J Ahmad, Gaurav K Jain; CBS Publishers and Distributors Pvt.Ltd. 2013.
- 5. Martin's Physical Pharmacy and Pharmaceutical Sciences, Fifth Edition, edited by Patrick J. Sinko, BI Publications Pvt. Ltd.
- 6. Targeted and Controlled Drug Delivery, Novel Carrier Systems by S. P. Vyas and R. K.Khar, CBS Publishers and Distributors Pvt. Ltd, First Edition 2012.
- 7. Pharmaceutical Dosage Forms and Drug Delivery Systems, Loyd V. Allen Jr., Nicholas B.Popovich, Howard C. Ansel, 9th Ed. 40
- 8. Aulton's Pharmaceutics The Design and Manufacture of Medicines, Michael E. Aulton,3rd Ed.
- 9. Remington The Science and Practice of Pharmacy, 20th Ed.
- 10. Pharmaceutical Dosage Forms Tablets Vol 1 to 3, A. Liberman, Leon Lachman and Joseph B. Schwartz
- 11. Pharmaceutical Dosage Forms Disperse Systems Vol 1 to 3, H.A. Liberman, Martin, M.R and Gilbert S. Banker.
- 12. Pharmaceutical Dosage Forms Parenteral Medication Vol 1 & 2, Kenneth E. Avis and H.A. Libermann.
- 13. Advanced Review Articles related to the topics.

BP814PW PROJECT WORK 12 Hours/Week

Project is a requirement for the B. Pharm. degree, wherein under the guidance of a faculty member, a group of not more than five learners in the eighth semester, is required to do some innovative work with the application of knowledge gained while learning various courses in the earlier years. The area of the project shall directly relate any one of the elective subject opted by the student in semester VIII. The learner/s is/are expected to do a survey of literature in the subject, work out a Project plan and carry it out through survey, experimentation and/or modeling / computation. Through the Project work the learner has to exhibit skills for both analysis and critical thinking. The complete details of the project have to submitted as a report of not less than 25 pages (A4, 1 inch margins, single line space, font Times Roman, font size 12, excluding count of reference pages) to the College before the prescribed date. The credits assigned for Project is 6 credits.