Himachal Pradesh Technical University, Hamirpur (H.P.)



CURRICULUM (PCI) BACHELOR OF PHARMCY (B.PHARMACY) 1st TO 8th SEMESTER





Teaching and Evaluation Scheme

SEMESTER- I (B. Pharmacy)

S.	Catego	Paper	Subject	L	T	P/D	Credits		Eva	luatio	n Sche	me
N.	ry	Code						Internal Assessment		ESE	Subject Total	
								CT		Total		
	Theory:				1		Į					
1	PC	BP101 T	Human Anatomy and Physiology I— Theory	3	1	-	4	15	10	25	75	100
2	PC	BP102 T	BP102T Pharmaceutical Analysis I – Theory	3	1	1	4	15	10	25	75	100
3	PC	BP103 T	BP103T Pharmaceutics I – Theory	3	1	-	4	15	10	25	75	100
4	PC	BP104 T	Pharmaceutical Inorganic Chemistry – Theory	3	1	-	4	15	10	25	75	100
5	MC	BP105 T	BP105T Communication skills – Theory *	2	-	1	2	10	5	15	35	50
6	MC	BP106 RBT BP106 RMT	Remedial Biology/ Remedial Mathematics - Theory*	2	-	1	2	10	5	15	35	50
		Labs:										
1	PC	BP107 P	Human Anatomy and Physiology – Practical	-	-	4	2	10	5	15	35	50
2	PC	BP108 P	BP108P Pharmaceutical Analysis I – Practical	-	-	4	2	10	5	15	35	50
3	PC	BP109 P	BP109P Pharmaceutics I – Practical	-	-	4	2	10	5	15	35	50
4	PC	BP110 P	Pharmaceutical Inorganic Chemistry – Practical	-	-	4	2	10	5	15	35	50
	MC	BP111 P	Communication skills – Practical*	-	-	2	1	5	5	10	15	25
	MC	BP112 RBP	Remedial Biology – Practical*	-	-	2	1	5	5	10	15	25
Total	Total				4	18/20	29/30					675/725 \$/ 750#
			Total work Load=27 H	Irs.			1	Total (Credi	t = 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)

Legend:

L	Lecture	ESE	End Semester Examination
T	Tutorial	PC	Program Core Courses
P	Practical	FC	Foundation Courses
CT	Class Test	HS	Humanities and social science
TA	Teacher's Assessment	MC	Mandatory Course



SEMESTER – II (B. Pharmacy)

S.	Categor	Paper	Subject	L	T	P/	Credits		Eva	aluation	Schen	ne	
N.	y	Code				D		Inter	rnal ssmer	ıt.	ESE	Subject Total	
								CT	TA	Total		Total	
	Theory:												
1	MC	BP201 T	Human Anatomy and Physiology II – Theory	3	1	-	4	15	10	25	75	100	
2	PC	BP202 T	Pharmaceutical Organic Chemistry I – Theory	3	1	-	4	15	10	25	75	100	
3	PC	BP203 T	Biochemistry – Theory	3	1	-	4	15	10	25	75	100	
4	PC	BP204 T	Pathophysiology – Theory	3	1	-	4	15	10	25	75	100	
5	PC	BP205 T	Computer Applications in Pharmacy – Theory *	3	-	-	3	15	10	25	75	100	
6	FC	BP206 T	Environmental sciences – Theory *	3	-	-	3	15	10	25	75	100	
1	PC	Labs: BP207 P	Human Anatomy and Physiology II –		-	4	2	10	5	15	35	50	
2	PC	BP208 P	Practical Pharmaceutical Organic Chemistry I— Practical			4	2	10	5	15	35	50	
3	PC	BP209 P	Biochemistry – Practical		-	4	2	10	5	15	35	50	
4	PC	BP210 P	Computer Applications in Pharmacy – Practical*		-	2	1	5	5	10	15	25	
Total	Cotal			18	0	14	29					725	
	Tota	Total Work Load 32 Hrs.			Total Credit 29								

^{*} The subject experts at college level shall conduct examinations

Legend:

L	Lecture	ESE	End Semester Examination
T	Tutorial	PC	Program Core Courses
P	Practical	FC	Foundation Courses
CT	Class Test	HS	Humanities and social science
TA	Teacher's Assessment	MC	Mandatory Course

SEMESTER- III (B. Pharmacy)

S.	Categor	Paper	Subject	L	T	P /	Cred		Eva	luatio	n Schei	ne
N.	У	Code				D	its	Inter Asse	rnal ssmen	ıt	ESE	Subject Total
								CT	TA	Tota l		
	Theo	ry:				•	•	•		•		
1	PC	BP301 T	Pharmaceutical Organic Chemistry II – Theory	3	1		4	15	10	25	75	100
2	PC	BP302 T	Physical Pharmaceutics I – Theory	3	1		4	15	10	25	75	100
3	PC	BP303 T	Pharmaceutical Microbiology – Theory	3	1		4	15	10	25	75	100
4	PC	BP304 T	Pharmaceutical Engineering – Theory	3	1		4	15	10	25	75	100
		Labs:										
1	PC	BP305 P	Pharmaceutical Organic Chemistry II – Practical			4	2	10	5	15	35	50
2	PC	BP306 P	Physical Pharmaceutics I — Practical			4	2	10	5	15	35	50
3	PC	BP307 P	Pharmaceutical Microbiology – Practical			4	2	10	5	15	35	50
4	PC	BP 308P	Pharmaceutical Engineering –Practical			4	2	10	5	15	35	50
Total				12	4	16	24					600
	Total Work Load=32 Hrs.					Total Credit 24						

Legend:

L	Lecture	ESE	End Semester Examination
T	Tutorial	PC	Program Core Courses
P	Practical	FC	Foundation Courses
CT	Class Test	HS	Humanities and social science
TA	Teacher's Assessment	MC	Mandatory Course

SEMESTER- IV (B. Pharmacy)

S. N.	Categor	Paper	Subject	L	T	P /	Credits		Eval	uation	Schem	ie
	y	Code				D		Inter	rnal ssmer	ıt	ESE	Subj ect
								CT	TA	Tota		Total
	Theo	rv:								1		
1	FC	BP401 T	Pharmaceutical Organic Chemistry III– Theory	3	1		4	15	10	25	75	100
2	PC	BP402 T	Medicinal Chemistry I – Theory	3	1		4	15	10	25	75	100
3	PC	BP403 T	Physical Pharmaceutics II – Theory	3	1		4	15	10	25	75	100
4	PC	BP404 T	Pharmacology I – Theory	3	1		4	15	10	25	75	100
5	PC	BP405 T	Pharmacognosy and Phytochemistry I– Theory	3	1		4	15	10	25	75	100
		Labs:										
1	CC	BP406 P	Medicinal Chemistry I – Practical			4	2	10	5	15	35	50
2	PC	BP407 P	Physical Pharmaceutics II– Practical			4	2	10	5	15	35	50
3	PC	BP408 P	Pharmacology I – Practical			4	2	10	5	15	35	50
4	PC	BP409 P	Pharmacognosy and Phytochemistry I – Practical			4	2	10	5	15	35	50
Total				15	5	16	28					700
		7	Fotal Working = 36 Hrs.	•	•	•	7	Total (Credit	=28		

Legend:

L	Lecture	ESE	End Semester Examination
T	Tutorial	PC	Program Core Courses
P	Practical	FC	Foundation Courses
CT	Class Test	HS	Humanities and social science
TA	Teacher's Assessment	MC	Mandatory Course

SEMESTER- V (B. Pharmacy)

S.	Categor	Paper	Subject	L	Т	P /	Credit		Ev	aluatio	on Scho	eme
N.	y	Code				D	S		Internal Assessment		ESE	Subject Total
								CT	TA	Tota l		
	Theo	ry:					l		ı	ı	I	l
1	PC	BP 501T	Medicinal Chemistry II – Theory	3	1	-	4	15	10	25	75	100
2	PC	BP 502T	Industrial PharmacyI— Theory	3	1	-	4	15	10	25	75	100
3	PC	BP 503T	Pharmacology II – Theory	3	1	-	4	15	10	25	75	100
4	PC	BP 504T	Pharmacognosy and Phytochemistry II– Theory	3	1	-	4	15	10	25	75	100
5	PC	BP 505T	Pharmaceutical Jurisprudence – Theory	3	1	-	4	15	10	25	75	100
Labs:					•				•	•	•	
1	PC	BP 506P	Industrial PharmacyI – Practical	-	-	4	2	10	5	15	35	50
2	PC	BP 507P	Pharmacology II - Practical	-	-	4	2	10	5	15	35	50
3	PC	BP 508P	Pharmacognosy and Phytochemistry II – Practical	ı	-	4	2	10	5	15	35	50
	Total			15	5	12	26					650
	Total Work Load=32 Hr								Total	Credi	t 26	

Legend:

L	Lecture	ESE	End Semester Examination
T	Tutorial	PC	Program Core Courses
P	Practical	FC	Foundation Courses
CT	Class Test	HS	Humanities and social science
TA	Teacher's Assessment	MC	Mandatory Course



SEMESTER- VI (B. Pharmacy)

S.		Paper	Subject	L	T	P /	Credits		Eva	luatio	n Sche	me
N.		Code						Inter	rnal ssmer	nt	ESE	Subjec Total
								CT		Tota		20002
										l		
	The	ory:										
1	PC	BP 601T	Medicinal Chemistry III – Theory	3	1	-	4	15	10	25	75	100
2	PC	BP 602T	Pharmacology III – Theory	3	1	-	4	15	10	25	75	100
3	PC	BP 603T	Herbal Drug Technology – Theory	3	1	-	4	15	10	25	75	100
4	PC	BP 604T	Biopharmaceutics and Pharmacokinetics – Theory	3	1	-	4	15	10	25	75	100
5	PC	BP 605T	Pharmaceutical Biotechnology – Theory	3	1	-	4	15	10	25	75	100
6	PC	BP 606T	Quality Assurance – Theory	3	1	-	4	15	10	25	75	100
		Labs:										
1	PC	BP 607P	Medicinal chemistry III – Practical	-	-	4	2	10	5	15	35	50
2	PC	BP 608P	Pharmacology III - Practical	-	-	4	2	10	5	15	35	50
3	PC	BP 609P	Herbal Drug Technology – Practical	-	-	4	2	10	5	15	35	50
	Total	•	•	18	6	12	30					750
		ŗ	Total Work Load=36 Hrs.	<u> </u>	<u>i </u>			T	otal (Credit	30	

Note

Legend:

L	Lecture	ESE	End Semester Examination
T	Tutorial	PC	Program Core Courses
P	Practical	FC	Foundation Courses
CT	Class Test	HS	Humanities and social science
TA	Teacher's Assessment	MC	Mandatory Course

SEMESTER- VII (B. Pharmacy)

S.	Category	Paper	Subject	L	T	P/	Credits	Evaluation			n Schei	ne
N.		Code				D		Inter	rnal		ESE	Subject
								Assessment				Total
								CT	TA	Tota		
										l		
	Theo	ry:										
1	PC	BP	Instrumental Methods of	3	1	-	4	15	10	25	75	100
		701T	Analysis – Theory									
2	PC	BP	Industrial PharmacyII -	3	1	-	4	15	10	25	75	100
		702T	702T Theory									
3	PC	BP	Pharmacy Practice -	3	1	-	4	15	10	25	75	100
		703T	Theory									
4	PC	BP704	Novel Drug	3	1	-	4	15	10	25	75	100
		T	Delivery System –									
			Theory									
		Labs:										
1	PC	BP	Instrumental Methods of			4	2	10	5	15	35	50
		705P	Analysis – Practical									
2	2 PC BP Practice School*		_		12	6	20	30	50	50	100	
	706P											
	Total						24					600
		1		Total Credit 24								

^{*} Non University Examination (NUE)

SEMESTER- VIII (B. Pharmacy)

3 4	PC PC PC	BP 802T BP 803ET BP 804ET BP	Biostatistics and Research Methodology Social and Preventive Pharmacy Pharma Marketing Management Pharmaceutical Regulatory Science	3	1 1		4 4	InterAsse CT	essmer	Tota 1 25 25	75 75	Subject Total 100
3 4	PC PC PC	BP 801T BP 802T BP 803ET BP 804ET	Research Methodology Social and Preventive Pharmacy Pharma Marketing Management Pharmaceutical			-		CT 15	TA	Tota 1		100
3 4	PC PC PC	BP 801T BP 802T BP 803ET BP 804ET	Research Methodology Social and Preventive Pharmacy Pharma Marketing Management Pharmaceutical			-						
3 4	PC PC PC	BP 801T BP 802T BP 803ET BP 804ET	Research Methodology Social and Preventive Pharmacy Pharma Marketing Management Pharmaceutical			-						
3 4	PC PC	801T BP 802T BP 803ET BP 804ET	Research Methodology Social and Preventive Pharmacy Pharma Marketing Management Pharmaceutical			-						
3 4	PC PC	802T BP 803ET BP 804ET	Pharmacy Pharma Marketing Management Pharmaceutical	3	1	-	4	15	10	25	75	100
4	PC	803ET BP 804ET	Management Pharmaceutical						1			1
·		804ET						15	10	25	75	100
	PC	BP						15	10	25	75	100
5		805ET	Pharmacovigilance	-				15	10	25	75	100
6	PC	BP 806ET	Quality Control and Standardization of Herbals		1			15	10	25	75	100
7	PC	BP 807ET	Computer Aided Drug Design		+ 1		A + A = 9	15	10	25	75	100
8	PC	BP 808ET	Cell and Molecular Biology	=6	= 2	-	4+4=8	15	10	25	75	100
9	PC	BP 809ET	Cosmetic Science					15	10	25	75	100
10	PC	BP 810ET	Experimental Pharmacology	-				15	10	25	75	100
11			Advanced Instrumentation Techniques					15	10	25	75	100
12 PC BP Dietary Supplements and Nutraceuticals						15	10	25	75	100		
		Labs:										
1	PC	BP 813 PW	Project Work	-	-	12	6	-	-	-	150	150
		To	tal	12	4	12	22					550
			Total Work Load=28	1	I			<u> </u>	 Total	Credit	22	

Legend:

L	Lecture	ESE	End Semester Examination
T	Tutorial	PC	Program Core Courses
P	Practical	FC	Foundation Courses

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CT	Class Test	HS	Humanities and social science
TA	Teacher's Assessment	MC	Mandatory Course

SEMESTER WISE CREDIT 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.

BP101T: HUMAN ANATOMY AND PHYSIOLOGY-I (Theory)

Teaching and Examination Scheme:

	Teac	hing Scl	neme	Credits		Marks	Duration of End Semester	
•	L	T	P	C	Sessional	End Semester	Total	Examination
						Exam		
	3	1	0	4	25	75	100	3 hours

Scope: This subject is designed to impart fundamental knowledge on the structure and functions 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

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

COURSE CONTENT

UNIT	CONTENT	No. ofHrs.
I	Introduction to human body: Definition and scope of anatomy and physiology, levels of structural organization and body systems, basic life processes, homeostasis, basic anatomical terminology.	10
	Cellular level of organization: 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.	
	Tissue level of organization: Classification of tissues, structure, location and	

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	functions of epithelial, muscular and nervous and connective tissues.	
II	Integumentary system: Structure and functions of skin.	10
	Skeletal system: 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.	
	Joints: Structural and functional classification, types of joints movements and its	
	articulation	
III	Body fluids and blood: Body fluids, composition and functions of blood,	
	hemopoeisis, formation of hemoglobin, anemia, mechanisms of coagulation,	10
	blood grouping, Rh factors, transfusion, its significance and disorders of blood,	
	Reticulo endothelial system.	
	Lymphatic system: Lymphatic organs and tissues, lymphatic vessels, lymph	
	circulation and functions of lymphatic system	
IV	Peripheral nervous system: Classification of peripheral nervous system:	
	Structure and functions of sympathetic and parasympathetic nervous system.	08
	Origin and functions of spinal and cranial nerves.	
	origin und runedons or spiritu und erunaur nervesi	
	Special senses: Structure and functions of eye, ear, nose and tongue and their	
	disorders.	
V	Cardiovascular system: Heart – anatomy of heart, blood circulation, blood	07
•		07
	vessels, structure and functions of artery, vein and capillaries, elements of	
	conduction system of heart and heart beat, its regulation by autonomic nervous	
	system, cardiac output, cardiac cycle. Regulation of blood pressure, pulse,	
	electrocardiogram and disorders of heart.	
1		



BP107P: HUMAN ANATOMY AND PHYSIOLOGY (Practical)

Teaching and Examination Scheme:

Teac	hing Sch	neme	Credits		Marks	Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
0	0	4	2	15	35	50	3 hours

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. 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

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- 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.



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BP102T: PHARMACEUTICAL ANALYSIS (Theory)

Teaching and Examination Scheme:

Teac	hing Sch	neme	Credits	Marks I			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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

- > understand the principles of volumetric and electro chemical analysis
- > carryout various volumetric and electrochemical titrations
- develop analytical skills

COURSE CONTENT

UNIT	CONTENT	No. ofHrs.						
I	(a) Pharmaceutical analysis- Definition and scope	10						
	i) Different techniques of analysis							
	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							
	(b)Errors: Sources of errors, types of errors, methods of minimizing							
	errors,accuracy, precision and significant figures							
	(c) Pharmacopoeia, Sources of impurities in medicinal agents, limit tests.							
II	Acid base titration: Theories of acid base indicators, classification of acid base	10						
	titrations and theory involved in titrations of strong, weak, and very weak acids and							
	bases, neutralization curves							
	Non aqueous titration: Solvents, acidimetry and alkalimetry titration and							
	estimation of Sodium benzoate and Ephedrine HCl							

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III	Precipitation titrations: Mohr's method, Volhard's, Modified Volhard's, Fajans	10							
	method, estimation of sodium chloride.								
	Complexometric titration: Classification, metal ion indicators, masking and								
	demasking reagents, estimation of Magnesium sulphate, and calcium gluconate.								
	Gravimetry: Principle and steps involved in gravimetric analysis. Purity of the								
	precipitate: co-precipitation and post precipitation, Estimation of barium sulphate.								
	Basic Principles,methods and application of diazotisation titration.								
IV	Redox titrations								
	(a)) Concepts of oxidation and reduction	08							
	(b) Types of redox titrations (Principles and applications)Cerimetry,								
	Iodimetry, Iodometry, Bromatometry, Dichrometry, Titration								
	withpotassium iodate								
V	Electrochemical methods of analysis	07							
	Conductometry- Introduction, Conductivity cell, Conductometric titrations,								
	applications.								
	Potentiometry - Electrochemical cell, construction and working of 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.								
	Polarography - Principle, Ilkovic equation, construction and working of dropping								
	mercury electrode and rotating platinum electrode, applications								

BP108P: PHARMACEUTICAL ANALYSIS (Practical)

Teaching and Examination Scheme:

Teac	ching Sch	neme	Credits	Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
0	0	4	2	15	35	50	4 hours

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, StahlonePress 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.

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BP103T: PHARMACEUTICS-I (Theory)

Teaching and Examination Scheme:

Teac	ching Scl	neme	Credits	Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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:

- ➤ Know the history of profession of pharmacy
- ➤ Understand the basics of different dosage forms, pharmaceutical incompatibilities andpharmaceutical calculations
- ➤ Understand the professional way of handling the prescription
- Preparation of various conventional dosage forms

COURSE CONTENT

UNIT	CONTENT	No. ofHrs.						
I	Historical background and development of profession of pharmacy: History of							
	profession of Pharmacy in India in relation to pharmacy education, industry and	10						
	organization, Pharmacy as a career, Pharmacopoeias: Introduction to IP, BP, USP							
	and Extra Pharmacopoeia.							
	Dosage forms: Introduction to dosage forms, classification and definitions							
	Prescription: Definition, Parts of prescription, handling of Prescription and Errors							
	in prescription.							
	Posology: Definition, Factors affecting posology. Pediatric dose calculations							
	based on age, body weight and body surface area.							
II	Pharmaceutical calculations: Weights and measures – Imperial & Metric system,	10						
	Calculations involving percentage solutions, alligation, proof spirit and isotonic	10						
	solutions based on freezing point and molecular weight.							
	Powders: Definition, classification, advantages and disadvantages, Simple &							
	compound powders - official preparations, dusting powders, effervescent,							





	efflorescent and hygroscopic powders, eutectic mixtures. Geometric dilutions.	
	Liquid dosage forms: Advantages and disadvantages of liquid dosage forms.	
	Excipients used in formulation of liquid dosage forms. Solubility enhancement	
	techniques	
III	Monophasic liquids: Definitions and preparations of Gargles, Mouthwashes,	
	Throat Paint, Eardrops, Nasal drops, Enemas, Syrups, Elixirs, Liniments	08
	andLotions.	
	Biphasic liquids: Suspensions: Definition, advantages and disadvantages,	
	classifications, Preparation of suspensions; Flocculated and Deflocculated	
	suspension & stability problems and methods to overcome. Emulsions:	
	Definition, classification, emulsifying agent, test for the identification of type of	
	Emulsion, Methods of preparation & stability problems and methods to overcome.	
IV	Suppositories: Definition, types, advantages and disadvantages, types of bases,	
	methods of preparations. Displacement value & its calculations, evaluation of	08
	suppositories.	
	Pharmaceutical incompatibilities: Definition, classification, physical, chemical	
	and therapeutic incompatibilities with examples.	
V	Semisolid dosage forms: Definitions, classification, mechanisms and factors	07
	influencing dermal penetration of drugs. Preparation of ointments, pastes, creams	
	and gels. Excipients used in semi solid dosage forms. Evaluation of semi solid	
	dosages forms	

BP109P: PHARMACEUTICSI (Practical)

Teaching and Examination Scheme:

Teac	hing Sch	cheme Credits		Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
0	0	4	2	15	35	50	4 hours

- 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
- **4. Solutions**a)Iodine Throat Paint (Mandles Paint)
- b) Strong solution of ammonium acetate
 - c) Cresol with soap solution
 - d) 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)Divided powders
- **8. Suppositories**a) Glycero gelatin suppository
- b) Coca butter suppository
 - c) Zinc Oxide suppository
- **9. Semisolids**a) Sulphur ointment
- b) Non staining-iodine ointment with methyl salicylate
 - c) Carbopal gel
- 10. Gargles and Mouthwashesa) Iodine gargleb) Chlorhexidine mouthwash

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Recommended Books: (Latest Editions)

- 1. H.C. Ansel et al., Pharmaceutical Dosage Form and Drug Delivery System, LippincottWilliams andWalkins, 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, LippincottWilliams, 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)

Teaching and Examination Scheme:

Teac	hing Sch	neme	Credits	Marks		Marks Duration of End Sen		Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination	
					Exam			
3	1	0	4	25	75	100	3 hours	

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

Objectives: Upon completion of course student shall be able to

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

COURSE CONTENT

General methods of preparation, assay for the compounds superscripted with **asterisk** (*), properties and medicinal uses of inorganic compounds belonging to the following classes

UNIT	CONTENT	No. ofHrs.
I	Impurities in pharmaceutical substances: History of Pharmacopoeia, Sources and types of impurities, principle involved in the limit test for Chloride, Sulphate, Iron, Arsenic, Lead and Heavy metals, modified limit test for Chloride and	10
	Sulphate	
II	Acids, Bases and Buffers: Buffer equations and buffer capacity in general, buffers in pharmaceutical systems, preparation, stability, buffered isotonic solutions, measurements of tonicity, calculations and methods of adjusting	10
	isotonicity. Major oytro and intracellular electrolytes: Functions of major physiclogical	
	Major extra and intracellular electrolytes: Functions of major physiological ions, Electrolytes used in the replacement therapy: Sodium chloride*, Potassium chloride, Calcium gluconate* and Oral Rehydration Salt (ORS), Physiological	
	acid base balance. Dental products: Dentifrices, role of fluoride in the treatment of dental caries,	
	Desensitizing agents, Calcium carbonate, Sodium fluoride, and Zinc eugenol cement.	

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III	Gastrointestinal agents	4.0					
	Acidifiers: Ammonium chloride* and Dil. HCl	10					
	Antacid: Ideal properties of antacids, combinations of antacids,						
	SodiumBicarbonate*, Aluminum hydroxide gel, Magnesium hydroxide						
	mixture						
	Cathartics: Magnesium sulphate, Sodium orthophosphate, Kaolin						
	andBentonite						
	Antimicrobials: Mechanism, classification, Potassium						
	permanganate, Boricacid, Hydrogen peroxide*, Chlorinated						
	lime*, Iodine and its preparations						
IV	General methods of preparation, assay for the compounds superscripted with	08					
	asterisk (*), properties and medicinal uses of inorganic compounds belonging to the following classes Miscellaneous compounds						
	Expectorants: Potassium iodide, Ammonium chloride*.						
	Emetics: Copper sulphate*, Sodium potassium tartarate						
	Haematinics: Ferrous sulphate*, Ferrous gluconate						
	Poison and Antidote: Sodium thiosulphate*, Activated charcoal,						
	Sodiumnitrite						
	Astringents: Zinc Sulphate, Potash Alum						
V	Radiopharmaceuticals: Radio activity, Measurement of radioactivity, Properties	07					
	of α , β , γ radiations, Half life, radio isotopes and study of radio isotopes - Sodium						
	iodide I131, Storage conditions, precautions & pharmaceutical application of						
	radioactive substances.						

BP110P: PHARMACEUTICAL INORGANIC CHEMISTRY (Practical)

Teaching and Examination Scheme:

Teac	Teaching Scheme Credits		Marks			Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
0	0	4	2	15	35	50	4 hours

I. Limit tests for following ions

- a. Limit test for Chlorides and Sulphates
- b. Modified limit test for Chlorides and Sulphates
- c. Limit test for Iron
- d. Limit test for Heavymetals
- e. Limit test for Lead
- f. Limit test for Arsenic
- II. Identification test for 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

IV. 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. Indian Pharmacopoeia

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BP105T: COMMUNICATION SKILLS (Theory)

Teaching and Examination Scheme:

Teac	hing Sch	neme	Credits	Marks		Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
2	0	0	2	15	35	50	1.5 hours

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 teamplayer and will add value to the pharmaceutical business.

Objectives:

Upon completion of the course the student shall be able to

- ➤ Understand the behavioral needs for a Pharmacist to function effectively in theareas of pharmaceutical operation
- ➤ Communicate effectively (Verbal and Non Verbal)
- > Effectivelymanage the team as a team player
- > Develop interview skills
- > Develop Leadership qualities and essentials

COURSE CONTENT

UNIT	CONTENT	No.					
		ofHrs.					
I	Communication Skills: Introduction, Definition, The Importance of						
	Communication, The Communication Process - Source, Message, Encoding,	07					
	Channel, Decoding, Receiver, Feedback, Context						
	Barriers to communication: Physiological Barriers, Physical Barriers, Cultural						
	Barriers, Language Barriers, Gender Barriers, Interpersonal Barriers,						
	Psychological Barriers, Emotional barriers						
	Perspectives in Communication: Introduction, Visual Perception, Language,						
	Other factors affecting our perspective - Past Experiences, Prejudices, Feelings,						
	Environment						
II	Elements of Communication: Introduction, Face to Face Communication - Tone						
		07					

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	of Voice, Body Language (Non-verbal communication), Verbal Communication,								
	Physical Communication								
	Communication Styles: Introduction, The Communication Styles Matrix with								
	example for each -Direct Communication Style, Spirited Communication Style,								
	Systematic Communication Style, Considerate Communication Style								
III	Basic Listening Skills: Introduction, Self-Awareness, Active Listening,	07							
	Becoming an Active Listener, Listening in Difficult Situations	07							
	Effective Written Communication: Introduction, When and When Not to Use								
	Written Communication - Complexity of the Topic, Amount of Discussion'								
	Required, Shades of Meaning, Formal Communication								
	Writing Effectively: Subject Lines, Put the Main Point First, Know Your								
	Audience, Organization of the Message								
IV	Interview Skills: Purpose of an interview, Do's and Dont's of an interview								
	Giving Presentations: Dealing with Fears, Planning your Presentation,	05							
	Structuring YourPresentation, Delivering Your Presentation, Techniques of								
	Delivery								
V	Group Discussion: Introduction, Communication skills in group discussion, Do's	04							
	and Dont's of group discussion								

BP111P: COMMUNICATION SKILLS (Practical)

Teaching and Examination Scheme:

Teac	Teaching Scheme (Credits	Marks			Duration of End
L	L T P		C	Sessional	End Semester	Total	Semester Examination
					Exam		
0	0	2	1	10	15	25	2 hours

The following learning modules are to be conducted using wordsworth® English languagelab software

Basic communication covering the following topics

Meeting People, Asking Questions, Making Friends, What did you do?, Do's and Dont's

Pronunciations covering the following topics

Pronunciation (Consonant Sounds), Pronunciation and Nouns, Pronunciation (Vowel Sounds)

Advanced Learning

Listening Comprehension / Direct and Indirect Speech, Figures of Speech, Effective Communication, Writing Skills, Effective Writing, Interview Handling Skills, E-Mail etiquette 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, GopalaSwamy Ramesh, 5thEdition, Pearson, 2013
- 6. Developing your influencing skills, Deborah Dalley, Lois Burton, Margaret, Greenhall, 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 GrawHill 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

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BP 106 RBT. REMEDIAL BIOLOGY (Theory)

Teaching and Examination Scheme:

Teac	Teaching Scheme		Credits		Marks	Duration of End Semester	
L	T P C Sessional End Semester Total		Examination				
					Exam		
2	0	0	2	15	35	50	1.5 hours

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

- ➤ know the classification and salient features of five kingdoms of life
- > understand the basic components of anatomy & physiology of plant
- > know understand the basic components of anatomy & physiology animal withspecial reference to human

COURSE CONTENT

UNIT	CONTENT	No.					
I	Living world: Definition and characters of living organisms; Diversity in the	ofHrs.					
_		07					
	living world; Binomial nomenclature ; Five kingdoms of life and basis of						
	classification. Salient features of Monera, Potista, Fungi, Animalia and Plantae,						
	Virus.						
	Morphology of Flowering plants: Morphology of different parts of flowering						
	plants - Root, stem, inflorescence, flower, leaf, fruit, seed.General Anatomy of						
	Root, stem, leaf of monocotyledons & Dicotylidones.						
II	Body fluids and circulation; Composition of blood, blood groups, coagulation						
	of blood; Composition and functions of lymph; Human circulatory system;	07					
	Structure of human heart and blood vessels; Cardiac cycle, cardiac output and						
	ECG						
	Digestion and Absorption: Human alimentary canal and digestive glands; Role						
	of digestive enzymes; Digestion, absorption and assimilation of digested food						
	Breathing and respiration: Human respiratory system; Mechanism of breathing						
	and its regulation; Exchange of gases, transport of gases and regulation of						

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	respiration; Respiratory volumes										
III	Excretory products and their elimination: Modes of excretion; Human										
	excretory system- structure and function; Urine formation; Rennin angiotensin	07									
	system										
	Neural control and coordination: 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.										
	Chemical coordination and regulation: Endocrine glands and their secretions;										
	Functions of hormones secreted by endocrine glands										
	Human reproduction: Parts of female reproductive system; Parts of male										
	reproductive system; Spermatogenesis and Oogenesis; Menstrual cycle										
IV	Plants and mineral nutrition: Essential mineral, macro and micronutrients;	05									
	Nitrogen metabolism, Nitrogen cycle, biological nitrogen fixation										
	Photosynthesis: Autotrophic nutrition, photosynthesis, Photosynthetic pigments,										
	Factors affecting photosynthesis.										
V	Plant respiration: Respiration, glycolysis, fermentation (anaerobic).	04									
	Plant growth and development: Phases and rate of plant growth, Condition of										
	growth,Introduction to plant growth regulators										
	Cell - The unit of life; Structure and functions of cell and cell organelles.Cell										
	division.										
	Tissues : Definition, types of tissues, location and functions.										

BP112 RBP. REMEDIAL BIOLOGY (Practical)

Teaching and Examination Scheme:

Teac	Teaching Scheme		Credits	Marks			Duration of End Semester
L	T	T P C Sessional End Semester Total		Examination			
					Exam		
0	0	2	1	10	15	25	2 hours

- 1. Introduction to experiments in biology
 - a) Study of Microscopeb) Section cutting techniquesc) 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

Text Books

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

Reference Books

- a. A Text book of Biology by B.V. Sreenivasa Naidu
- b. A Text book of Biology by Naidu and Murthy
- c. Botany for Degree students By A.C.Dutta.
- d.Outlines of Zoology by M. Ekambaranatha ayyer and T. N. Ananthakrishnan.
- e. A manual for pharmaceutical biology practical by S.B. Gokhale and C. K. Kokate
- f. Practical human anatomy and physiology. by S.R.Kale and R.R.Kale.
- g. A Manual of pharmaceutical biology practical by S.B.Gokhale, C.K.Kokate and S.P.Shriwastava.
- h.Biology practical manual according to National core curriculum .Biology forum of Karnataka. Prof .M.J.H.Shafi



BP 106RMT. REMEDIAL MATHEMATICS (Theory)

Teaching and Examination Scheme:

Teac	Teaching Scheme Cre		Credits	Marks			Duration of End Semester
L	T	P	C	Sessional	Sessional End Semester Total		Examination
					Exam		
2	0	0	2	15	35	50	1.5 Hours

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:-

- ➤ Know the theory and their application in Pharmacy
- > Solve the different types of problems by applying theory
- > Appreciate the important application of mathematics in Pharmacy

COURSE CONTENT

UNIT	CONTENT							
Ι	Partial fraction	ofHrs.						
	Introduction, Polynomial, Rational fractions, Proper and Improper	06						
	fractions, Partial fraction , Resolving into Partial fraction, Application of							
	PartialFraction in Chemical Kinetics and Pharmacokinetics							
	Logarithms							
	Introduction, Definition, Theorems/Properties of logarithms, Commonlogarithms,							
	Characteristic and Mantissa, worked examples, application of logarithm to solve							
	pharmaceutical problems.							
	Function: Real Valued function, Classification of real valued functions,							
	Limits and continuity							
	Introduction , Limit of a function, Definition of limit of a function (∈ - δ							
	definition), $\lim_{x \to a} \frac{x^n - a^n}{x - a} = na^{n-1}$, $\lim_{\theta \to 0} \frac{\sin \theta}{\theta} = 1$,							
II	Matrices and Determinant:	0.6						
	Introduction matrices, Types of matrices, Operation on matrices, Transpose of a	06						
	matrix, Matrix Multiplication, Determinants, Properties ofdeterminants, Product							

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	of determinants, Minors and co-Factors, Adjointor adjugate of a square matrix ,	
	Singular and non-singular matrices, Inverse of a matrix, Solution of system of	
	linear of equations using matrixmethod, Cramer's rule, Characteristic equation	
	and roots of a squarematrix, Cayley-Hamilton theorem, Application of Matrices in	
	solvingPharmacokinetic equations.	
III	Calculus	
	Differentiation : Introductions, Derivative of a function, Derivative of aconstant,	06
	Derivative of a product of a constant and a function, Derivative of the sum or	
	difference of two functions, Derivative of the product of twofunctions (product	
	formula), Derivative of the quotient of two functions(Quotient formula) -	
	Without Proof, Derivative of xn $w.r.t$ x, where n is any rational number, Derivative	
	of ex , Derivative of loge x , Derivative of ax , Derivative of trigonometric functions	
	from first principles (withoutProof), Successive Differentiation, Conditions for a	
	function to be amaximum or a minimum at a point. Application	
TT 7	1 10	
IV	Analytical Geometry	
IV	Analytical Geometry Introduction: Signs of the Coordinates, Distance formula, Straight Line: Slope	06
IV		06
IV	Introduction: Signs of the Coordinates, Distance formula, Straight Line: Slope	06
IV	Introduction: Signs of the Coordinates, Distance formula, Straight Line : Slope or gradient of a straight line, Conditions forparallelism and perpendicularity of	06
IV	Introduction: Signs of the Coordinates, Distance formula, Straight Line : Slope or gradient of a straight line, Conditions forparallelism and perpendicularity of two lines, Slope of a line joining twopoints, Slope – intercept form of a straight	06
IV	Introduction: Signs of the Coordinates, Distance formula, Straight Line: Slope or gradient of a straight line, Conditions forparallelism and perpendicularity of two lines, Slope of a line joining twopoints, Slope – intercept form of a straight line. Integration: Introduction, Definition, Standard formulae, Rules of	06
V	Introduction: Signs of the Coordinates, Distance formula, Straight Line: Slope or gradient of a straight line, Conditions forparallelism and perpendicularity of two lines, Slope of a line joining twopoints, Slope – intercept form of a straight line. Integration: Introduction, Definition, Standard formulae, Rules of integration, Method of Substitution, Method of Partial fractions, Integration by	06
	Introduction: Signs of the Coordinates, Distance formula, Straight Line: Slope or gradient of a straight line, Conditions forparallelism and perpendicularity of two lines, Slope of a line joining twopoints, Slope – intercept form of a straight line. Integration: Introduction, Definition, Standard formulae, Rules of integration, Method of substitution, Method of Partial fractions, Integration by parts, definiteintegrals, application	
	Introduction: Signs of the Coordinates, Distance formula, Straight Line: Slope or gradient of a straight line, Conditions forparallelism and perpendicularity of two lines, Slope of a line joining twopoints, Slope – intercept form of a straight line. Integration: Introduction, Definition, Standard formulae, Rules of integration, Method of substitution, Method of Partial fractions, Integration by parts, definiteintegrals, application Differential Equations: Some basic definitions, Order and degree, Equations in	
	Introduction: Signs of the Coordinates, Distance formula, Straight Line: Slope or gradient of a straight line, Conditions forparallelism and perpendicularity of two lines, Slope of a line joining twopoints, Slope – intercept form of a straight line. Integration: Introduction, Definition, Standard formulae, Rules of integration, Method of substitution, Method of Partial fractions, Integration by parts, definiteintegrals, application Differential Equations: Some basic definitions, Order and degree, Equations in separable form, Homogeneous equations, Linear Differential equations, Exact	
	Introduction: Signs of the Coordinates, Distance formula, Straight Line: Slope or gradient of a straight line, Conditions forparallelism and perpendicularity of two lines, Slope of a line joining twopoints, Slope – intercept form of a straight line. Integration: Introduction, Definition, Standard formulae, Rules of integration, Method of substitution, Method of Partial fractions, Integration by parts, definiteintegrals, application Differential Equations: Some basic definitions, Order and degree, Equations in separable form, Homogeneous equations, Linear Differential equations, Exact equations, Application in solving Pharmacokinetic equations	
	Introduction: Signs of the Coordinates, Distance formula, Straight Line: Slope or gradient of a straight line, Conditions forparallelism and perpendicularity of two lines, Slope of a line joining twopoints, Slope – intercept form of a straight line. Integration: Introduction, Definition, Standard formulae, Rules of integration, Method of substitution, Method of Partial fractions, Integration by parts, definiteintegrals, application Differential Equations: Some basic definitions, Order and degree, Equations in separable form, Homogeneous equations, Linear Differential equations, Exact equations, Application in solving Pharmacokinetic equations Laplace Transform: Introduction, Definition, Properties of Laplace transform,	
	Introduction: Signs of the Coordinates, Distance formula, Straight Line: Slope or gradient of a straight line, Conditions forparallelism and perpendicularity of two lines, Slope of a line joining twopoints, Slope – intercept form of a straight line. Integration: Introduction, Definition, Standard formulae, Rules of integration, Method of substitution, Method of Partial fractions, Integration by parts, definiteintegrals, application Differential Equations: Some basic definitions, Order and degree, Equations in separable form, Homogeneous equations, Linear Differential equations, Exact equations, Application in solving Pharmacokinetic equations Laplace Transform: Introduction, Definition, Properties of Laplace transform, Laplace Transforms of elementary functions, Inverse Laplace transforms, Laplace	

Recommended Books (Latest Edition)

- 1. Differential Calculus by Shanthinarayan
- 2. Pharmaceutical Mathematics with application to Pharmacy by PanchaksharappaGowda D.H.





- 3. Integral Calculus by Shanthinarayan
- 4. Higher Engineering Mathematics by Dr.B.S.Grewal



2ND SEMESTER

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BP 201T: HUMAN ANATOMY AND PHYSIOLOGY-II (Theory)

Teaching and Examination Scheme:

Teaching Scheme		Credits	Marks			Duration of End Semester	
L	T	T P C Sessional End Semester Total		Examination			
					Exam		
3	1	0	4	25	75	100	3 hours

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 bothhomeostatic mechanisms. The subject provides the basic knowledge required tounderstand the various disciplines of pharmacy.

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

- Explain the gross morphology, structure and functions of various organs of the human body.
- > Describe the various homeostatic mechanisms and their imbalances.
- > Identify the various tissues and organs of different systems of human body.
- ➤ 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.
- ➤ Appreciate coordinated working pattern of different organs of each system
- Appreciate the interlinked mechanisms in the maintenance of normal functioning (homeostasis) of human body.

COURSE CONTENT

UNIT	CONTENT							
		ofHrs.						
I	Nervous system: Organization of nervous system, neuron, neuroglia,							
	classification andproperties of nerve fibre, electrophysiology, action	10						
	potential,nerve impulse, receptors, synapse, neurotransmitters.							
	Central nervous system: Meninges, ventricles of brain andcerebrospinal							
	fluid.structure and functions of brain (cerebrum, brainstem, cerebellum), spinal							
	cord (gross structure, functions of afferentand efferent nerve tracts,reflex activity)							

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II	Digestive system: Anatomy of GI Tract with special reference to anatomy and	10							
	functions of stomach, (Acid production in the stomach, regulation of acid	10							
	production throughparasympathetic nervous system, pepsin role in protein								
	digestion) small intestineand large intestine, anatomy and functions of salivary								
	glands, pancreas and liver, movements of GIT, digestion and absorption of								
	nutrients and disorders of GIT.								
	Energetics: Formation and role of ATP, Creatinine Phosphate and BMR.								
III	Respiratory system: Anatomy of respiratory system with special reference to	10							
	anatomy of lungs,mechanism of respiration, regulation of respirationLung								
	Volumes and capacities transport of respiratory gases, artificial respiration, and								
	resuscitation methods.								
	Urinary system: Anatomy of urinary tract with special reference to anatomy of								
	kidney andnephrons, functions of kidney and urinary tract, physiology of urine								
	formation, micturition reflex and role of kidneys in acid base balance, role of RAS								
	in kidneyand disorders of kidney.								
IV	Endocrine system								
	Classification of hormones, mechanism of hormone action, structureand functions	10							
	of pituitary gland, thyroid gland, parathyroid gland, adrenalgland, pancreas, pineal								
	gland, thymus and their disorders.								
V	Reproductive system	09							
	Anatomy of male and female reproductive system, Functions of male and								
	femalereproductive system, sex hormones, physiology of menstruation,								
	fertilization, spermatogenesis, oogenesis, pregnancy and parturition								
	Introduction to genetics								
	Chromosomes, genes and DNA, protein synthesis, genetic pattern of inheritance								

BP 207 P: HUMAN ANATOMY AND PHYSIOLOGY (Practical)

Teaching and Examination Scheme:

Teaching Scheme			Credits		Marks	Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
0	0	2	1	10	15	25	4 hours

Practical physiology is complimentary to the theoretical discussions inphysiology. Practicals allow the verification of physiological processes discussed in theory classes through experiments on living tissue, intact animals or normalhuman 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. Jaypeebrothers medical publishers, New Delhi.

P. Technical University

Hamirpur - 177001

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- 2. Anatomy and Physiology in Health and Illness by Kathleen J.W. Wilson, ChurchillLivingstone, New York
- 3.Physiological basis of Medical Practice-Best and Tailor. Williams & WilkinsCo,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 brothers medical publishers, New Delhi.
- 7. Textbook of Practical Physiology by C.L. Ghai, Jaypee brothers medical publishers, NewDelhi.
- 8. Practical workbook of Human Physiology by K. Srinageswari and Rajeev Sharma, Jaypee brother's medical publishers, New Delhi.

Reference Books:

- 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)

Teaching and Examination Scheme:

Teaching Scheme			Credits		Marks	Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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

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

COURSE CONTENT

General methods of preparation and reactions of compounds superscripted with asterisk (*) to be ExplainedTo emphasize on definition, types, classification, principles/mechanisms, applications, examples and differences

UNIT	CONTENT	No. ofHrs.						
I	Classification, nomenclature and isomerism: Classification of Organic	0=						
	Compounds, Common and IUPAC systems of nomenclature of organic	07						
	compounds(up to 10 Carbons open chain and carbocyclic compounds).							
	Structural isomerisms in organic compounds							
II	Alkanes*, Alkenes* and Conjugated dienes*: SP3 hybridization in alkanes,							
	Halogenation of alkanes, uses of paraffins. Stabilities of alkenes, SP2							
	hybridization in alkenes							
	E1 and E2 reactions – kinetics, order of reactivity of alkyl halides,							
	rearrangement of carbocations, Saytzeffs orientation and evidences. E1 verses E2							





	reactions, Factors affecting E1	
	and E2 reactions. Ozonolysis, electrophilic addition reactions of alkenes,	
	Markownikoff'sorientation, free radical addition reactions of alkenes, Anti	
	Markownikoff's orientation.Stability of conjugated dienes, Diel-Alder,	
	electrophilic addition, free radical additionreactions of conjugated dienes, allylic	
	rearrangement	
III	Alkyl halides*: SN1 and SN2 reactions - kinetics, order of reactivity of alkyl	
	halides, stereochemistry andrearrangement of carbocations.SN1 versus SN2	10
	reactions, Factors affecting SN1 and SN2 reactions. Structure and uses of	
	ethylchloride, Chloroform, trichloroethylene,	
	tetrachloroethylene,dichloromethane, tetrachloromethane and iodoform.	
	Alcohols*- Qualitative tests, Structure and uses of Ethyl alcohol, Methyl	
	alcohol, chlorobutanol, Cetosteryl alcohol, Benzyl alcohol, Glycerol, Propylene	
	glycol	
IV	Carbonyl compounds* (Aldehydes and ketones): Nucleophilic addition,	
	Electromeric effect, aldol condensation, Crossed Aldol condensation, Cannizzaro	10
	reaction, Crossed Cannizzaro reaction, Benzoin condensation,	
	Perkincondensation, qualitative tests, Structure and uses of Formaldehyde,	
	Paraldehyde, Acetone, Chloral hydrate, Hexamine, Benzaldehyde, Vanilin,	
	Cinnamaldehyde.	
V	Carboxylic acids*: Acidity of carboxylic acids, effect of substituents on acidity,	08
	inductive effect and qualitativetests for carboxylic acids ,amide and ester.	
	Structure and Uses of Acetic acid, Lactic acid, Tartaric acid, Citric acid,	
	Succinic acid. Oxalicacid, Salicylic acid, Benzoic acid, Benzyl benzoate,	
	Dimethyl phthalate, Methyl salicylate andAcetyl salicylic acid	
	Aliphatic amines* - Basicity, effect of substituent on Basicity. Qualitative test,	
	Structure anduses of Ethanolamine, Ethylenediamine, Amphetamine	
l		

BP208P. PHARMACEUTICAL ORGANIC CHEMISTRY -I (Practical)

Teaching and Examination Scheme:

Teaching Scheme			Credits		Marks	Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
0	0	4	2	15	35	50	4 hours

Systematic qualitative analysis of unknown organic compounds

- 1. Preliminary test: Color, odour, aliphatic/aromatic compounds, saturation unsaturation, etc.
- 2. Detection of elements like Nitrogen, Sulphur and Halogen by Lassaigne's test
- 3. Solubility test
- 4. Functional group test like Phenols, Amides/ Urea, Carbohydrates, Amines, Carboxylic acids, Aldehydes and Ketones, Alcohols, Esters, Aromatic and Halogenated Hydrocarbons, Nitro compounds and Anilides.
- 5. Melting point/Boiling point of organic compounds
- 6. Identification of the unknown compound from the literature using melting point/boiling point.
- 7. Preparation of the derivatives and confirmation of the unknowncompound bymelting point/boiling point.
- 8. Minimum 5 unknown organic compounds to be analyzed systematically.
- 9. Preparation of suitable solid derivatives from organic compounds
- 10. 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.

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BP203 T. BIOCHEMISTRY (Theory)

Teaching and Examination Scheme:

Teaching Scheme			Credits		Marks	Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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

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

COURSE CONTENT

UNIT	CONTENT	No. ofHrs.					
I	Biomolecules: Introduction, classification, chemical nature and biological role	00					
	ofcarbohydrate, lipids, nucleic acids, amino acids and proteins.	08					
	Bioenergetics: Concept of free energy, endergonic and exergonic reaction,						
	Relationshipbetween free energy, enthalpy and entropy; Redox potential.Energy						
	rich compounds; classification; biological significances of ATPand cyclic AMP						
II	I 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						

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	Hormonal regulation of blood glucose level and Diabetes mellitus									
	Biological oxidation: Electron transport chain (ETC) and its									
	mechanism.Oxidative phosphorylation & its mechanism and									
	substratephosphorylation. Inhibitors ETC and oxidative									
	phosphorylation/Uncouplerslevel									
III	Lipid metabolism: β-Oxidation of saturated fatty acid (Palmitic acid)	10								
	Formation and utilization of ketone bodies; ketoacidosis									
	De novo synthesis of fatty acids (Palmitic acid)									
	Biological significance of cholesterol and conversion of cholesterol intobile acids,									
	steroid hormone and vitamin D									
	Disorders of lipid metabolism: Hypercholesterolemia, atherosclerosis,fatty liver									
	and obesity.									
	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									
IV	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									
V	Enzymes: Introduction, properties, nomenclature and IUB classification of	07								
	enzymes, Enzyme kinetics (Michaelis plot, Line Weaver Burke plot)									
	Enzyme inhibitors with examples, Regulation of enzymes: enzyme induction and									
	repression, allostericenzymes regulation, Therapeutic and diagnostic applications									
	of enzymes and isoenzymes, Coenzymes –Structure and biochemical functions									



BP 209 P. BIOCHEMISTRY (Practical)

Teaching and Examination Scheme:

Teaching Scheme			neme	Credits		Marks	Duration of End Semester	
	L	T	P	С	Sessional	End Semester	Total	Examination
						Exam		
	0	0	4	2	15	35	50	2 hours

- 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.

BP 204T.PATHOPHYSIOLOGY (THEORY)

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Teaching and Examination Scheme:

Teaching Scheme			Credits		Marks	Duration of End Semester		
L	,	T	P	C Sessional End Semester Total		Examination		
						Exam		
3		1	0	4	25	75	100	3 hours

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 itspharmacological applications, and understanding of basic pathophysiological mechanisms. Hence it will not only help to study the syllabus of pathology, but also toget baseline knowledge required to practice medicine safely, confidently, rationally and effectively.

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

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

COURSE CONTENT

UNIT	CONTENT	No.						
		ofHrs.						
I	Basic principles of Cell injury and Adaptation: 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 cellularaccumulation, Calcification, Enzyme leakage and Cell							
	Death Acidosis&Alkalosis,Electrolyte imbalance							
	Basic mechanism involved in the process of inflammation and 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 theskin,Pathophysiology of Atherosclerosis.							
II	Cardiovascular System: Hypertension, congestive heart failure, ischemic heart							
		10						





	diameter (and in the state of t	
	disease (angina,myocardialinfarction, atherosclerosis and arteriosclerosis)	
	Respiratory system: Asthma, Chronic obstructive airways diseases.	
	Renal system: Acute and chronic renal failure.	
III	Haematological Diseases: Iron deficiency, megaloblastic anemia (Vit B12 and	
	folic acid), sickle cell anemia,thalasemia, hereditary acquired anemia,	10
	hemophilia	
	Endocrine system: Diabetes, thyroid diseases, disorders of sex hormones	
	Nervous system: Epilepsy, Parkinson's disease, stroke, psychiatric	
	disorders:depression, schizophrenia and Alzheimer's disease.	
	Gastrointestinal system: Peptic Ulcer	
IV	Inflammatory bowel diseases, jaundice, hepatitis (A,B,C,D,E,F) alcoholic	
	liverdisease.	08
	Disease of bones and joints: Rheumatoid arthritis, osteoporosis and gout	
	Principles of cancer: classification, etiology and pathogenesis of cancer	
	Diseases of bones and joints: Rheumatoid Arthritis, Osteoporosis, Gout	
	Principles of Cancer: Classification, etiology and pathogenesis of Cancer	
V	Infectious diseases: Meningitis, Typhoid, Leprosy, Tuberculosis, Urinary tract	07
	infections	
	Sexually transmitted diseases: AIDS, Syphilis, Gonorrhea	

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. HarshMohan; 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 SaundersCompany; 2010.
- 8. Joseph DiPiro, Robert L. Talbert, Gary Yee, Barbara Wells, L. Michael Posey; Pharmacotherapy: A Pathophysiological Approach; 9th edition; London; McGraw-HillMedical; 2014.
- 9. V. Kumar, R. S. Cotran and S. L. Robbins; Basic Pathology; 6th edition; Philadelphia; WB Saunders Company; 1997.
- 10. 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.



BP205 T. COMPUTER APPLICATIONS IN PHARMACY (Theory)

Teaching and Examination Scheme:

Teac	ching Scheme Credits			Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	0	0	3	25	75	100	3 hours

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

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

- ➤ know the various types of application of computers in pharmacy
- > know the various types of databases
- > know the various applications of databases in pharmacy

COURSE CONTENT

UNIT	CONTENT	No. ofHrs.			
I	Number system: Binary number system, Decimal number system, Octalnumber system, Hexadecimal number systems, conversion decimal tobinary, binary to decimal, octal to binary etc, binary addition, binarysubtraction — One's complement, Two's complement method, binarymultiplication, binary division Concept of Information Systems and Software: Information gathering, requirement and feasibility analysis, data flow diagrams, processspecifications, input/output design, process life cycle, planning andmanaging the project	06			
II	Web technologies:Introduction to HTML, XML,CSS andProgramming languages, introduction to web servers and ServerProducts, Introduction to databases, MYSQL, MS ACCESS, Pharmacy Drug database				
III	Application of computers in Pharmacy – Drug information storage andretrieval, Pharmacokinetics, Mathematical model in Drug design, Hospitaland 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,	06			

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	Patient Monitoring System, Pharma Information System					
IV	Bioinformatics: Introduction, Objective of Bioinformatics,					
	BioinformaticsDatabases, Concept of Bioinformatics, Impact of Bioinformatics in					
	Vaccine					
	Discovery					
V	Computers as data analysis in Preclinical development:	06				
	Chromatographic dada analysis(CDS), Laboratory Information management					
	System (LIMS) and Text Information Management System(TIMS)					

BP210P. COMPUTER APPLICATIONS IN PHARMACY (Practical)

Teaching and Examination Scheme:

Teaching Scheme Credits			Marks	Duration of End Semester			
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
0	0	2	1	10	15	25	2 hours

- 1. Design a questionnaire using a word processing package to gather informationabout a particular disease.
- 2. Create a HTML web page to show personal information.
- 3 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 requiredfields Using access
- 6. Design a form in MS Access to view, add, delete and modify the patient record in he 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, 600South 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)
- 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

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BP 206 T. ENVIRONMENTAL SCIENCES (Theory)

Teaching and Examination Scheme:

Teac	Teaching Scheme Credits Marks				Duration of End Semester		
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	0	0	3	25	75	100	3 hours

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 studyof 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:

- > Create the awareness about environmental problems among learners.
- ➤ Impart basic knowledge about the environment and its allied problems.
- > Develop an attitude of concern for the environment.
- Motivate learner to participate in environment protection and environmentimprovement.
- Acquire skills to help the concerned individuals in identifying and solvingenvironmental problems.
- > Strive to attain harmony with Nature.

COURSE CONTENT

UNIT	CONTENT						
I	The Multidisciplinary nature of environmental studies: Natural Resources	10					
	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.						
II	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						

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	ecosystem; Aquatic ecosystems (ponds, streams, lakes, rivers, oceans, estuaries)	
III	Environmental Pollution: Air pollution; Water pollution; Soil pollution	10

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



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BP301T. PHARMACEUTICAL ORGANIC CHEMISTRY –II (Theory)

Teaching and Examination Scheme:

	Teac	hing Scl	neme	Credits	Marks			Duration of End Semester
•	L	T	P	C	Sessional	End Semester	Total	Examination
						Exam		
	3	1	0	4	25	75	100	3 hours

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

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

- write the structure, name and the type of isomerism of the organic compound
- write the reaction, name the reaction and orientation of reactions
- account for reactivity/stability of compounds,
- > prepare organic compounds

COURSE CONTENT

General methods of preparation and reactions of compounds superscripted with asterisk (*) to be ExplainedTo emphasize on definition, types, classification, principles/mechanisms, applications, examples and differences

UNIT	CONTENT	No. of							
I	Benzene and its derivatives:	10							
	A. Analytical, synthetic and other evidences in the derivation of structure of								
	benzene, Orbital picture, resonance in benzene, aromaticcharacters, Huckel's rule								
	B. Reactions of benzene - nitration, sulphonation, halogenations								
	reactivity,Friedelcrafts alkylation- reactivity, limitations,Friedelcrafts acylation.								
	C. Substituents, effect of substituents on reactivity and orientation ofmono								
	substituted benzene compounds towards electrophilicsubstitution reaction								
	D. Structure and uses of DDT, Saccharin, BHC and Chloramine								
II	Phenols* - Acidity of phenols, effect of substituents on acidity, qualitativetests,								
	Structure and uses of phenol, cresols, resorcinol, naphthols								





	Aromatic Amines* - Basicity of amines, effect of substituents on basicity, and synthetic uses of aryl diazonium salts Aromatic Acids* -Acidity, effect of substituents on acidity and important reactions of benzoic acid.	
III	Fats and Oils	10
	a. Fatty acids – reactions.	
	b. Hydrolysis, Hydrogenation, Saponification and Rancidity of oils, Dryingoils.	
	c. Analytical constants - Acid value, Saponification value, Ester value, Iodine	
	value, Acetyl value, Reichert Meissl (RM) value – significance andprinciple	
	involved in their determination.	
IV	Polynuclear hydrocarbons:	08
	a. Synthesis, reactions, structure and medicinal uses of Naphthalene,	
	Phenanthrene, Anthracene, Diphenylmethane, Triphenylmethane and their	
	derivatives	
V	Cyclo alkanes*Stabilities – Baeyer's strain theory, limitation of Baeyer's strain	07
	theory, Coulson and Moffitt's modification, Sachse Mohr's theory (Theory	
	ofstrainless rings), reactions of cyclopropane and cyclobutane only	

BP305P. PHARMACEUTICAL ORGANIC CHEMISTRY -II (Practical)

Teaching and Examination Scheme:

Teac	hing Scl	neme	Credits	Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
0	0	4	2	15	35	50	3 hours

I Experiments involving laboratory techniques

- a) Recrystallization
- b) Steam distillation

II Determination of following oil values (including standardization of reagents)

- a) Acid value
- b) Saponification value c) 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

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.

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BP302T. PHYSICAL PHARMACEUTICS-I (Theory)

Teaching and Examination Scheme:

	Teaching Scheme			Credits		Marks	Duration of End Semester	
•	L	T	P	C	Sessional	End Semester	Total	Examination
						Exam		
	3	1	0	4	25	75	100	3 hours

Scope: The course deals with the various physica 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

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

COURSE CONTENT

UNIT	CONTENT	No. of						
		Hrs.						
I	Solubility of drugs: Solubility expressions, mechanisms of solute solvent							
	interactions, ideal solubility parameters, solvation & association, quantitative							
	approach to the factorsinfluencing solubility of drugs, diffusion principles in							
	biological systems. Solubility of gas in liquids, solubility of liquids in liquids,							
	(Binary solutions, ideal solutions)Raoult's law, real solutions. Partiallymiscible							
	liquids, Critical solution temperature and applications. Distribution law, its							
	limitations and applications							
II	States of Matter and properties of matter: State of matter, changes in the state	10						
	of matter, latent heats, vapour pressure, sublimation critical point, eutectic							
	mixtures, gases, aerosols- inhalers, relative humidity, liquid complexes, liquid							
	crystals, glassy states, solidcrystalline,amorphous & polymorphism.							

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	Physicochemical properties of drug molecules: Refractive index, optical rotation, dielectric constant, dipole moment, dissociation constant, determinations and applications	
III	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.	08
IV	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	08
V	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.	07

BP306P. PHYSICAL PHARMACEUTICS – I (Practical)

Teaching and Examination Scheme:

Teaching Scheme			Credits		Marks		Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
0	0	4	2	15	35	50	3 hours

- 1. Determination the solubility of drug at room temperature
- 2. Determination of pKa value by Half Neutralization/ Henderson Hasselbalchequation.
- 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 byCST method
- 6. Determination of surface tension of given liquids by drop count and drop weightmethod
- 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-Caffeinecomplex by solubilitymethod
- 11. Determination of stability constant and donor acceptor ratio of Cupric-Glycinecomplex 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 to3, MarcelDekkar Inc.
- 6. Liberman H.A, Lachman C, Pharmaceutical Dosage forms. Dispersesystems, volume 1, 2, 3. Marcel Dekkar Inc.
- 7. Physical Pharmaceutics by Ramasamy C and ManavalanR.
- 8. LaboratoryManual of Physical Pharmaceutics, C.V.S. Subramanyam, J.Thimma settee
- 9. Physical Pharmaceutics by C.V.S. Subramanyam
- 10. Test book of Physical Phramacy, by Gaurav Jain & Roop K. Khar

BP 303 T. PHARMACEUTICAL MICROBIOLOGY (Theory)

Teaching and Examination Scheme:

Teaching Scheme			Credits		Marks	Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

Scope:Study of all categories of microorganisims especially for the production of alcholantibiotics, vaccines, vitamins enzymes etc..

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

- > Understand methods of identification, cultivation and preservation of various microorganisms
- ➤ To understand the importance and implementation of sterlization inpharmaceutical processing and industry
- ➤ Learn sterility testing of pharmaceutical products.
- ➤ Carried out microbiological standardization of Pharmaceuticals.
- ➤ Understand the cell culture technology and its applications in pharmaceuticalindustries.

COURSE CONTENT

UNIT	CONTENT	No. of						
		Hrs.						
I	Introduction, history of microbiology, its branches, scope and itsimportance.							
	Introduction to Prokaryotes and Eukaryotes, 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 methodsfor pure cultures, cultivation of anaerobes, quantitative							
	measurement ofbacterial growth (total & viable count). Study of different types of							
	phase constrast microscopy, dark fieldmicroscopy and electron microscopy.							
II	Identification of bacteria using staining techniques (simple, Gram's &Acidfast							
	staining) and biochemical tests (IMViC).Study of principle, procedure, merits,							
	demerits and applications of physical, chemical gaseous, radiation and mechanical							
	method of sterilization. Evaluation of the efficiency of sterilization							
	methods. Equipments employed in large scale sterilization. Sterility indicators.							

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III	Study of morphology, classification, reproduction/replication and cultivation of	10				
	Fungi and Viruses. Classification and mode of action of disinfectants Factors	10				
	influencing disinfection, antiseptics and their evaluation. Forbacteriostatic and					
	bactericidal actionsEvaluation of bactericidal & Bacteriostatic.Sterility testing of					
	products (solids, liquids, ophthalmic and other sterileproducts) according to IP,					
	BP and USP.					
IV	Designing of aseptic area, laminar flow equipments; study of differentsources of	08				
	contamination in an aseptic area and methods of prevention, clean area					
	classification.Principles and methods of different microbiological assay. Methods					
	forstandardization of antibiotics, vitamins and amino acids. Assessment of a new					
	antibiotic.					
V	Types of spoilage, factors affecting the microbial spoilage ofpharmaceutical	07				
	products, sources and types of microbial contaminants, assessment of microbial					
	contamination and spoilage.Preservation of pharmaceutical products using					
	antimicrobial agents, evaluation of microbial stability of formulations. Growth of					
	animal cells in culture, general procedure for cell culture, Primary, established					
	and transformed cell cultures. Application of cell cultures in pharmaceutical					
	industry and research.					

BP 307P. PHARMACEUTICAL MICROBIOLOGY (Practical)

Teaching and Examination Scheme:

Teaching Scheme			Credits		Marks	Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
0	0	4	2	15	35	50	3 hours

- 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 withpractical).
- 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.

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. Ananthnarayan: Text Book of Microbiology, Orient-Longman, Chennai
- 10. Edward: Fundamentals of Microbiology.
- 11. N.K.Jain: Pharmaceutical Microbiology, Vallabh Prakashan, Delhi
- 12. Bergeys manual of systematic bacteriology, Williams and Wilkins- A Waverly company

BP 304 T. PHARMACEUTICAL ENGINEERING (Theory)

Teaching and Examination Scheme:

Teaching Scheme			neme	Credits		Marks	Duration of End Semester	
L		T	P	C	Sessional	End Semester	Total	Examination
						Exam		
3	}	1	0	4	25	75	100	3 hours

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:

- To know various unit operations used in Pharmaceutical industries.
- > To understand the material handling techniques.
- > To perform various processes involved in pharmaceutical manufacturing process.
- > To carry out various test to prevent environmental pollution.
- ➤ To appreciate and comprehend significance of plant lay out design for optimumuse of resources.
- To appreciate the various preventive methods used for corrosion control inPharmaceutical industries.

COURSE CONTENT

UNIT	CONTENT	No. of Hrs.						
I	Flow of fluids: Types of manometers, Reynolds number and its	10						
	significance,Bernoulli's theorem and its applications, Energy losses, Orifice							
	meter, Venturimeter, Pitot tube and Rotometer.							
	Size Reduction: Objectives, Mechanisms & Laws governing size							
	reduction, factors affecting size reduction, principles, construction, working, uses,							
	merits anddemerits of Hammer mill, ball mill, fluid energy mill, Edge runner mill							
	& endrunner mill.							
	Size Separation: Objectives, applications & mechanism of size separation, official							
	standards of powders, sieves, size separation Principles, construction, working,							
	uses, merits and demerits of Sieve shaker, cyclone separator, Airseparator, Bag							
	filter & elutriation tank.							

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II	Heat Transfer: Objectives, applications & Heat transfer mechanisms.	10
	Fourier'slaw, Heat transfer by conduction, convection & radiation. Heat	
	interchangers &heat exchangers.	
	Evaporation: Objectives, applications and factors influencing	
	evaporation, differences between evaporation and other heat process. principles,	
	construction,working, uses, merits and demerits of Steam jacketed kettle,	
	horizontal tubeevaporator, climbing film evaporator, forced circulation evaporator,	
	multipleeffect evaporator& Economy of multiple effect evaporator.	
	Distillation: Basic Principles and methodology of simple	
	distillation, flashdistillation, fractional distillation, distillation under reduced	
	pressure, steamdistillation & molecular distillation	
III	Drying: Objectives, applications & mechanism of drying process, measurements&	08
	applications of Equilibrium Moisture content, rate of drying curve.	
	principles, construction, working, uses, merits and demerits of Tray dryer, drum	
	dryer spraydryer, fluidized bed dryer, vacuum dryer, freeze dryer.	
	Mixing: Objectives, applications & factors affecting mixing, Difference	
	betweensolid and liquid mixing, mechanism of solid mixing, liquids mixing	
	andsemisolids mixing. Principles, Construction, Working, uses, Merits and	
	Demeritsof Double cone blender, twin shell blender, ribbon blender, Sigma blade	
	mixer,planetarymixers, Propellers, Turbines, Paddles & Silverson Emulsifier	
IV	Filtration: Objectives, applications, Theories & Factors influencing filtration, filter	08
	aids, filter medias. Principle, Construction, Working, Uses, Merits anddemerits of	
	plate & frame filter, filter leaf, rotary drum filter, Meta filter &Cartridge filter,	
	membrane filters and Seidtz filter.	
	Centrifugation: Objectives, principle & applications of Centrifugation, principles,	
	construction, working, uses, merits and demerits of Perforated basketcentrifuge,	
	Non-perforated basket centrifuge, semi continuous centrifuge & supercentrifuge.	
V	Materials of pharmaceutical plant construction, Corrosion and itsprevention:	07
	Factors affecting during materials selected for Pharmaceutical plantconstruction,	
	Theories of corrosion, types of corrosion and there prevention. Ferrous and	
	nonferrous metals, inorganic and organic non metals, basic ofmaterial handling	
	systems.	



BP308P - PHARMACEUTICAL ENGINEERING (Practical)

Teaching and Examination Scheme:

Teaching Scheme			Credits		Marks		Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
0	0	4	2	15	35	50	3 hours

- 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 PharmaceuticalMachinery 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 arithmeticandlogarithmic probability plots.
- 9. Size reduction: To verify the laws of size reduction using ball mill anddetermining Kicks, Rittinger's, Bond's coefficients, power requirement and ritical speed of Ball Mill.
- 10. Demonstration of colloid mill, planetary mixer, fluidized bed dryer, freeze dryerand such othermajor equipment.
- 11. Factors affecting Rate of Filtration and Evaporation (Surface area, Concentrationand 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 ConeBlender.

Recommended Books: (Latest Editions)

- 1. Introduction to chemical engineering Walter L Badger & Julius Banchero, Latestedition.
- 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., Latestedition.
- 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.

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4th SEMESTER

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BP401T. PHARMACEUTICAL ORGANIC CHEMISTRY –III (Theory)

Teaching and Examination Scheme:

Teaching Scheme			Credits		Marks		Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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

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

- > understand the methods of preparation and properties of organic compounds
- > explain the stereo chemical aspects of organic compounds and stereo chemicalreactions
- ➤ know the medicinal uses and other applications of organic compounds

Course Content:

Note: To emphasize on definition, types, mechanisms, examples, uses/applications

UNIT	CONTENT						
I	Stereo isomerism: Optical isomerism —Optical activity, enantiomerism, diastereoisomerism, meso compounds. Elements of symmetry, chiral and achiral molecules, DL system of nomenclature of optical isomers, sequence rules, RS system ofnomenclature of optical isomers, Reactions of chiral molecules, Racemic modification and resolution of racemic mixture. Asymmetric synthesis: partial and absolute	Hrs. 10					
II	Geometrical isomerism: Nomenclature of geometrical isomers (Cis Trans, EZ, Syn Anti systems), Methods of determination of configuration of geometrical isomers. Conformational isomerism in Ethane, n-Butane and Cyclohexane. Stereo isomerism in biphenyl compounds (Atropisomerism) and conditions for optical activity. Stereospecific and stereoselective reactions	10					
III	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	10					

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IV	Synthesis, reactions and medicinal uses of following compounds/derivatives	08				
	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					
V	Reactions of synthetic importance: Metal hydride reduction (NaBH4 and	07				
	LiAlH4), Clemmensen reduction, Birch					
	reduction, Wolff Kishner reduction. Oppenauer-oxidation and Dakin reaction.					
	Beckmanns rearrangement and Schmidt rearrangement.Claisen-Schmidt					
	condensation					

Recommended Books (Latest Editions)

- 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)

Teaching and Examination Scheme:

	Teac	hing Scl	neme	Credits		Marks		Duration of End Semester
İ	L	T	P	C	Sessional	End Semester	Total	Examination
						Exam		
	3	1	0	4	25	75	100	3 hours

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

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

Course Content:

Note: Study of the development of the following classes of drugs, Classification, mechanism ofaction, 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 (*)

UNIT	CONTENT	No. of
		Hrs.
I	Introduction to Medicinal Chemistry	10
	History and development of medicinal chemistry	
	Physicochemical properties in relation to biological action	
	Ionization, Solubility, Partition Coefficient, Hydrogen bonding, Proteinbinding,	
	Chelation, Bioisosterism, Optical and Geometrical isomerism.	
	Drug metabolism	
	Drug metabolism principles- Phase I and Phase II.Factors affecting drug	
	metabolism including stereo chemical aspects.	

II	Drugs acting on Autonomic Nervous System	10							
	Adrenergic Neurotransmitters:								
	Biosynthesis and catabolism of catecholamine.Adrenergic receptors (Alpha &								
	Beta) and their distribution.								
	Sympathomimetic agents: SAR of Sympathomimetic agents								
	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.								
	Adrenergic Antagonists:								
	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.								
III	Cholinergic neurotransmitters:								
	Biosynthesis and catabolism of acetylcholine. Cholinergic receptors (Muscarinic								
	& Nicotinic) and their distribution.								
	Parasympathomimetic agents: SAR of Parasympathomimetic agents								
	Direct acting agents: Acetylcholine, Carbachol*, Bethanechol,Methacholine,								
	Pilocarpine. Indirect acting/ Cholinesterase inhibitors (Reversible & Irreversible):								
	Physostigmine, Neostigmine*, Pyridostigmine, Edrophonium chloride, Tacrine								
	hydrochloride, Ambenonium chloride, Isofluorphate, Echothiophateiodide,								
	Parathione, Malathion.								
	Cholinesterase reactivator: Pralidoxime chloride.								
	Cholinergic Blocking agents: SAR of cholinolytic agents								
	Solanaceous alkaloids and analogues: Atropine sulphate,								
	Hyoscyaminesulphate, Scopolamine hydrobromide, Homatropine								
	hydrobromide,Ipratropium bromide*.								
	Synthetic cholinergic blocking agents: Tropicamide,								



	Cyclopentolatehydrochloride, Clidinium bromide, Dicyclomine	
	hydrochloride*,Glycopyrrolate, Methantheline bromide, Propantheline	
	bromide,Benztropine mesylate, Orphenadrine citrate, Biperidine	
	hydrochloride,Procyclidine hydrochloride*, Tridihexethyl chloride, Isopropamide	
	iodide,Ethopropazine hydrochloride.	
IV	Drugs acting on Central Nervous System	08
	A. Sedatives and Hypnotics:	
	Benzodiazepines: SAR of Benzodiazepines, Chlordiazepoxide,	
	Diazepam*,Oxazepam, Chlorazepate, Lorazepam, Alprazolam, Zolpidem	
	Barbiturtes: SAR of barbiturates, Barbital*, Phenobarbital,	
	Mephobarbital, Amobarbital, Butabarbital, Pentobarbital, Secobarbital	
	Miscelleneous:	
	Amides & imides: Glutethmide.Alcohol & their carbamate derivatives:	
	Meprobomate, Ethchlorvynol.	
	Aldehyde & their derivatives: Triclofos sodium, Paraldehyde.	
	B. Antipsychotics	
	Phenothiazeines: SAR of Phenothiazeines - Promazine	
	hydrochloride,Chlorpromazine hydrochloride*, Triflupromazine,	
	Thioridazinehydrochloride, 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.	
	C. Anticonvulsants: SAR of Anticonvulsants, mechanism of anticonvulsant	
	Action. Barbiturates: Phenobarbitone, Methabarbital. Hydantoins: Phenytoin*,	
	Mephenytoin, Ethotoin Oxazolidine diones:Trimethadione, Paramethadione	
	Succinimides: Phensuximide, Methsuximide, Ethosuximide* Urea	
	andmonoacylureas: Phenacemide, Carbamazepine*Benzodiazepines:	
	Clonazepam	
	Miscellaneous: Primidone, Valproic acid, Gabapentin, Felbamate	
V	Drugs acting on Central Nervous System	07



General anesthetics:

Inhalation anesthetics: Halothane*, Methoxyflurane, Enflurane,Sevoflurane, Isoflurane, Desflurane.

Ultra short acting barbitutrates: Methohexital sodium*, Thiamylalsodium, Thiopental sodium.**Dissociative anesthetics:** Ketamine hydrochloride.*

Narcotic and non-narcotic analgesics

Morphine and related drugs: SAR of Morphine analogues, Morphinesulphate, 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.

Sodium Anti-inflammatory Mefenamic agents: salicylate, Aspirin, acid*, Meclofenamate, Indomethacin, Sulindac, Tolmetin, Zomepriac, Naproxen, Diclofenac, Ketorolac, Ibuprofen*, Piroxicam, Phenacetin, Acetaminophen, Antipyrine, Phenylbutazone.



BP406P. MEDICINAL CHEMISTRY – I (Practical)

Teaching and Examination Scheme:

Teac	hing Scl	neme	Credits	Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
0	0	4	2	15	35	50	3 hours

I Preparation of drugs/intermediates

a)1,3-pyrazole

- b) 1,3-oxazole
- c) Benzimidazole
- d) Benztriazole

- e) 2,3- diphenyl quinoxaline
- f) Benzocaine
- g) Phenytoin
- h) Phenothiazine

i) Barbiturate

II Assay of drugs: Chlorpromazine, Phenobarbitone, Atropine, Ibuprofen, Aspirin, 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.

BP 403 T. PHYSICAL PHARMACEUTICS-II (Theory)

Teaching and Examination Scheme:

Teac	Ceaching Scheme Credits		Credits	Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

Scope: The course deals with the various physica 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

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

COURSE CONTENT

UNIT	CONTENT	No. of Hrs.
I	Colloidal dispersions: Classification of dispersed systems & their generalcharacteristics, 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	07
II	Rheology: Newtonian systems, law of flow, kinematic viscosity, effect of temperature,non-Newtonian systems, pseudoplastic, dilatant, plastic, thixotropy, thixotropy informulation, determination of viscosity, capillary, falling Sphere, rotational viscometers. Deformation of solids: Plastic and elastic deformation, Heckel equation, Stress, Strain, Elastic Modulus	10

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III	Coarse dispersion: Suspension, interfacial properties of suspended particles,	10								
	settling insuspensions, formulation of flocculated and deflocculated suspensions.									
	Emulsions and theories of emulsification, microemulsion and multiple emulsions;									
	Stability of emulsions, preservation of emulsions, rheological properties of									
	emulsions and emulsionformulation by HLB method.									
IV	Micromeretics: Particle size and distribution, mean particle size, number and	10								
	weightdistribution, particle number, methods for determining particle size by									
	differentmethods, counting and separation method, particle shape, specific									
	surface, methods fordetermining surface area, permeability, adsorption, derived									
	properties of powders, porosity, packing arrangement, densities, bulkiness & flow									
	properties.									
V	Drug stability: Reaction kinetics: zero, pseudo-zero, first & second order, units									
	of basicrate constants, determination of reaction order. Physical and chemical									
	factors influencingthe 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									

BP 407P. PHYSICAL PHARMACEUTICS- II (Practical)

Teaching and Examination Scheme:

Teac	hing Scheme Credits		Marks			Duration of End Semester	
L	T	P	С	Sessional	End Semester	Total	Examination
					Exam		
0	0	4	2	15	35	50	3 hours

- 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 1,2, 3. Marcel Dekkar Inc.
- 7. Physical Pharmaceutics by Ramasamy C, and Manavalan R.



BP 404 T. PHARMACOLOGY-I (Theory)

Teaching and Examination Scheme:

Teac	hing Sch	neme	e Credits		Marks		Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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 theinformation about the drugs like, mechanism of action, physiological and biochemicaleffects (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

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

COURSE CONTENT

UNIT	CONTENT	No. of Hrs.
I	General Pharmacology	08
	Introduction to Pharmacology- Definition, historical landmarks and scope of	
	pharmacology, nature and source of drugs, essential drugs concept and routes of	
	drug administration, Agonists, antagonists(competitive and non competitive),	
	sparereceptors, addiction, tolerance, dependence, tachyphylaxis, idiosyncrasy,	
	allergy. Pharmacokinetics- Membrane transport, absorption, distribution,	
	metabolism and excretion of drugs .Enzyme induction, enzyme inhibition,	
	kinetics of elimination	

II	General Pharmacology	12						
	Pharmacodynamics- Principles and mechanisms of drug action. Receptor							
	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 bindingreceptor and receptors that regulate transcription factors, dose							
	responserelationship, therapeutic index, combined effects of drugs and factors							
	modifyingdrug action. Adverse drug reactions, Drug interactions							
	(pharmacokinetic and pharmacodynamic), Drug discovery and clinical evaluation							
	of new drugs -Drug discovery phase, preclinical evaluation phase, clinical trial							
	phase, phases of clinical trials andpharmacovigilance.							
III	Pharmacology of drugs acting on peripheral nervous system	10						
111		10						
	Organization and function of ANS.Neurohumoral transmission,co-transmission							
	and classification of							
	neurotransmitters.Parasympathomimetics,Parasympatholytics,							
	Sympathomimetics, sympatholytics.Neuromuscular blocking agents and skeletal							
	muscle relaxants (peripheral).Local anesthetic agents.Drugs used in myasthenia							
	gravis and glaucoma							
IV	Pharmacology of drugs acting on central nervous system	08						
	Neurohumoral transmission in the C.N.S.special emphasis on importance of							
	variousneurotransmitters like with GABA, Glutamate, Glycine, serotonin,							
	dopamine.General anesthetics and pre-anesthetics.Sedatives, hypnotics and							
	centrally acting muscle relaxants. Anti-epileptics, Alcohols and disulfiram							
V	Pharmacology of drugs acting on central nervous system	07						
	Psychopharmacological agents: Antipsychotics, antidepressants, anti-anxiety							
	agents,anti-manics and hallucinogens.Drugs used in Parkinsons disease and							
	Alzheimer's disease.CNS stimulants and nootropics.Opioid analgesics and							
	antagonists, Drug addiction, drug abuse, tolerance and dependence							
L								



BP 408 P.PHARMACOLOGY-I (Practical)

Teaching and Examination Scheme:

Teac	hing Scheme Credits		Marks			Duration of End Semester	
L	T	P	С	Sessional	End Semester	Total	Examination
					Exam		
0	0	4	2	15	35	50	3 hours

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

- 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, an esthetics 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 sleepingtime 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 byMES 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

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 McGraw-Hill

Technical University

Hamirpur - 177001

3. Goodman and Gilman's, The Pharmacological Basis of Therapeutics

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- 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 LippincottWilliams &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 MedicalPublishers (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,

BP 405 T.PHARMACOGNOSY AND PHYTOCHEMISTRY I (Theory)

Teaching and Examination Scheme:

Teac	hing Sch	neme	Credits	Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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

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

- ➤ know the techniques in the cultivation and production of crude drugs
- > know the crude drugs, their uses and chemical nature
- ➤ know the evaluation techniques for the herbal drugs
- > carry out the microscopic and morphological evaluation of crude drugs

Course Content

UNIT	CONTENT	No. of Hrs.
I	Introduction to Pharmacognosy:	08
	(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 andmucilages, oleoresins and oleo- gum -resins).	
	Classification of drugs:	
	Alphabetical, morphological, taxonomical, chemical, pharmacological, chemo	
	and serotaxonomical classification of drugs	
	Quality control of Drugs of Natural Origin:	
	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.	





II	Cultivation, Collection, Processing and storage of drugs of natural origin:	10
	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, Conservation of	
	medicinal plants	
III	Plant tissue culture: Historical development of plant tissue culture, types of	07
	cultures, Nutritional requirements, growth and their maintenance. Applications of	
	plant tissue culture in pharmacognosy.Edible vaccines	
IV	Pharmacognosy in various systems of medicine: Role of Pharmacognosy in	10
	allopathy and traditional systems of medicine namely, Ayurveda, Unani, Siddha,	
	Homeopathy and Chinese systems of medicine.	
	Introduction to secondary metabolites: Definition, classification, properties	
	and test for identification of Alkaloids, Glycosides, Flavonoids, Tannins, Volatile	
	oil and Resins	
V	Study of biological source, chemical nature and uses of drugs of natural origin	08
	containingfollowing drugs	
	Plant Products:	
	Fibers - Cotton, Jute, Hemp	
	Hallucinogens, Teratogens, Natural allergens	
	Primary metabolites:	
	General introduction, detailed study with respect to chemistry, sources,	
	preparation, evaluation, preservation, storage, therapeutic used and commercial	
	utility as PharmaceuticalAids and/or Medicines for the following	
	Primarymetabolites:	
	Carbohydrates: Acacia, Agar, Tragacanth, Honey	
	Proteins and Enzymes: Gelatin, casein, proteolytic enzymes (Papain,	
	bromelain, serratiopeptidase, urokinase, streptokinase, pepsin).	
	Lipids(Waxes, fats, fixed oils): Castor oil, Chaulmoogra oil, Wool Fat, Bees	
	Wax	
	Marine Drugs:	
	Novel medicinal agents from marine sources	



BP408 P. PHARMACOGNOSY AND PHYTOCHEMISTRY I (Practical)

Teaching and Examination Scheme:

Teac	hing Sch	neme	Credits	Marks			Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination	
					Exam			
0	0	4	2	15	35	50	3 hours	

- 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 piecemicrometer
- 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, NewDelhi.
- 7. Essentials of Pharmacognosy, Dr.SH.Ansari, IInd edition, Birla publications, NewDelhi, 2007
- 8. Practical Pharmacognosy: C.K. Kokate, Purohit, Gokhlae
- 9. Anatomy of Crude Drugs by M.A. Iyengar

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5th SEMESTER

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BP501T. MEDICINAL CHEMISTRY – II (Theory)

Teaching and Examination Scheme:

Teac	ching Scl	neme	Credits	Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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

- ➤ Understand the chemistry of drugs with respect to their pharmacological activity
- ➤ Understand the drug metabolic pathways, adverse effect and therapeutic value ofdrugs
- ➤ Know the Structural Activity Relationship of different class of drugs
- > Study the chemical synthesis of selected drugs

COURSE CONTENT

Note: 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 (*)

UNIT	CONTENT								
		Hrs.							
I	Antihistaminic agents: Histamine, receptors and their distribution in	10							
	thehumanbody. H1-antagonists: Diphenhydramine hydrochloride*,								
	Dimenhydrinate, Doxylamines cuccinate, Clemastine fumarate,								
	Diphenylphyraline hydrochloride,								
	Tripelenamine hydrochloride, Chlorcyclizine hydrochloride,								
	Meclizinehydrochloride, Buclizine hydrochloride, Chlorpheniramine maleate,								
	Triprolidine								
	hydrochloride*, Phenidamine tartarate, Promethazine								
	hydrochloride*, Trimeprazine tartrate, Cyproheptadine hydrochloride, Azatidine								
	maleate, Astemizole, Loratadine, Cetirizine, Levocetrazine Cromolyn sodium H2-								

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antagonists: Cimetidine*, Famotidine, Ranitidin. Gastric **Proton** pump inhibitors: Omeprazole, Lansoprazole, Rabeprazole, Pantoprazole **Anti-neoplastic agents:** Alkylating agents: 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. II 10 **Anti-anginal**: Vasodilators: Amyl nitrite. Nitroglycerin*, Pentaerythritol tetranitrate. Isosorbidedinitrite*, Dipyridamole. Calcium channel Verapamil, Bepridil hydrochloride, blockers: Diltiazemhydrochloride, Nifedipine, Amlodipine, Felodipine, Nicardipine, Nimodipine. **Diuretics:** Carbonic anhydrase inhibitors: Acetazolamide*, Methazolamide, Dichlorphenamide. Thiazides: Chlorthiazide*, Hydrochlorothiazide, Hydroflumethiazide, Cyclothiazide. Loop diuretics: Furosemide*, Bumetanide, Ethacrynic acid.Potassium sparing Diuretics: Spironolactone, Triamterene, Amiloride. Osmotic Diuretics: Mannitol Anti-hypertensive Agents: Timolol, Captopril, Lisinopril, Enalapril, Benazepril hydrochloride, Quinapril hydrochloride, Methyldopate hydrochloride,* Clonidine hydrochloride, Guanethidine monosulphate, Guanabenz acetate. Sodiumnitroprusside, Reserpine, Hydralazine Diazoxide, Minoxidil, hydrochloride. Ш Quinidine 10 **Anti-arrhythmic Drugs**: sulphate, Procainamide hydrochloride, Disopyramide phosphate*, Phenytoin sodium, Lidocaine hydrochloride, Tocainide hydrochloride, Mexiletine hydrochloride, Lorcainide hydrochloride,



	Amiodarone, Sotalol.							
	Anti-hyperlipidemic agents: Clofibrate, Lovastatin, Cholesteramine							
	andCholestipol							
	Coagulant & Anticoagulants: Menadione, Acetomenadione,							
	Warfarin*, Anisindione, clopidogrel							
	Drugs used in Congestive Heart Failure: Digoxin, Digitoxin,							
	Nesiritide,Bosentan, Tezosentan.							
IV	Drugs acting on Endocrine system	08						
	Nomenclature, Stereochemistry and metabolism of steroids							
	Sex hormones: Testosterone, Nandralone, Progestrones, Oestriol,							
	Oestradiol,Oestrione, Diethyl stilbestrol.							
	Drugs for erectile dysfunction: Sildenafil, Tadalafil.							
	Oral contraceptives: Mifepristone, Norgestril, Levonorgestrol							
	Corticosteroids: Cortisone, Hydrocortisone, Prednisolone,							
	Betamethasone,Dexamethasone							
	Thyroid and antithyroid drugs: L-Thyroxine, L-Thyronine,							
	Propylthiouracil, Methimazole.							
V	Antidiabetic agents:	07						
	Insulin and its preparations, Sulfonyl ureas: Tolbutamide*, Chlorpropamide,							
	Glipizide, Glimepiride., Biguanides: Metformin., Thiazolidinediones:							
	Pioglitazone, Rosiglitazone., Meglitinides: Repaglinide, Nateglinide.Glucosidase							
	inhibitors: Acrabose, Voglibose.							
	Local Anesthetics: SAR of Local anesthetics							
	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.*							

Recommended Books (Latest Editions)

1. Wilson and Giswold's Organic medicinal and Pharmaceutical Chemistry.

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- 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.



BP 502 T. Industrial PharmacyI (Theory)

Teaching and Examination Scheme:

Teac	ching Scl	neme	Credits	Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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

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

COURSE CONTENT

UNIT	CONTENT	No. of Hrs.							
I	Preformulation Studies: Introduction to preformulation, goals and objectives,								
	study ofphysicochemical characteristics of drug substances.								
	a. Physical properties: Physical form (crystal & amorphous), particle size, shape,								
	flowproperties, solubility profile (pKa, pH, partition coefficient), polymorphism								
	b. Chemical Properties: Hydrolysis, oxidation, reduction, racemisation,								
	polymerization, BCS classification of drugs & its significant, Application of								
	preformulation considerations in the development of solid, liquid oral								
	andparenteral dosage forms and its impact on stability of dosage forms.								
II	Tablets:	10							
	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								
	coatingcomposition, methods of coating, equipment employed and defects in								
	coating.								
	c. Quality control tests: In process and finished product tests								

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	Liquid orals: Formulation and manufacturing consideration of syrups and elixirs							
	suspensions and emulsions; Filling and packaging; evaluation of liquid orals							
	official in pharmacopoeia							
III	Capsules:	08						
	a. <i>Hard gelatin capsules:</i> Introduction, Production of hard gelatin capsule shells.							
	Sizeof capsules, Filling, finishing and special techniques of formulation of hard							
	gelatincapsules, manufacturing defects. In process and final product quality							
	control testsfor capsules.							
	b. Soft gelatin capsules: Nature of shell and capsule content, size							
	ofcapsules,importance of base adsorption and minim/gram factors, production, in							
	process and final product quality control tests. Packing, storage and stability							
	testingof soft gelatin capsules and their applications.							
	Pellets: Introduction, formulation requirements, pelletization process,							
	equipments formanufacture of pellets							
IV	Parenteral Products:	10						
	a. Definition, types, advantages and limitations. Preformulation factors and							
	essentialrequirements, vehicles, additives, importance of isotonicity							
	b. Production procedure, production facilities and controls, aseptic processing							
	c. Formulation of injections, sterile powders, large volume parenterals and							
	lyophilized products.							
	d. Containers and closures selection, filling and sealing of ampoules, vials and							
	infusionfluids. Quality control tests of parenteral products.							
	Ophthalmic Preparations: Introduction, formulation considerations;							
	formulation of eyedrops, eye ointments and eye lotions; methods of preparation;							
	labeling, containers; evaluation of ophthalmic preparations							
V	Cosmetics: Formulation and preparation of the following cosmetic preparations:	10						
	lipsticks, shampoos, cold cream and vanishing cream, tooth pastes, hair dyes and							
	sunscreens.							
	Pharmaceutical Aerosols: Definition, propellants, containers, valves, types of							
	aerosolsystems; formulation and manufacture of aerosols; Evaluation of aerosols;							
	Qualitycontrol and stability studies.							
	Packaging Materials Science: Materials used for packaging of pharmaceutical							



products, factors influencing choice of containers, legal and official requirements for containers, stability aspects of packaging materials, quality control tests.

BP 506 P. INDUSTRIAL PHARMACYI (PRACTICAL)

Teaching and Examination Scheme:

Teac	ching Scl	neme	Credits	Marks			Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination	
					Exam			
0	0	4	2	15	35	50	3 hours	

- 1. Preformulation studies on paracetamol/asparin/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. Qulaity 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 PharmaceuticalScience (RPS)
- 6. Theory and Practice of Industrial Pharmacy by Liberman & Lachman
- 7. Pharmaceutics- The science of dosage form design by M.E.Aulton, Churchilllivingstone, 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.

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BP503.T. PHARMACOLOGY-II (Theory)

Teaching and Examination Scheme:

	Teac	hing Scl	neme	Credits	Marks			Duration of End Semester
•	L	T	P	C	Sessional	End Semester	Total	Examination
						Exam		
	3	1	0	4	25	75	100	3 hours

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, emphasison the basic concepts of bioassay.

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

- > Understand the mechanism of drug action and its relevance in the treatment of different diseases
- Demonstrate isolation of different organs/tissues from the laboratory animals bysimulated experiments
- > Demonstrate the various receptor actions using isolated tissue preparation
- ➤ Appreciate correlation of pharmacology with related medical sciences

COURSE CONTENT

UNIT	CONTENT	No. of
I	Pharmacology of drugs acting on cardio vascular system	Hrs. 10
	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.	
II	Pharmacology of drugs acting on cardio vascular system	10
	a. Drug used in the therapy of shock.	
	b. Hematinics, coagulants and anticoagulants.	
	c. Fibrinolytics and anti-platelet drugs	

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	d. Plasma volume expanders	
	Pharmacology of drugs acting on urinary system	
	a. Diuretics	
	b. Anti-diuretics.	
III	Autocoids and related drugs	10
	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	
IV	Pharmacology of drugs acting on endocrine system	08
	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.	
	d. Insulin, Oral Hypoglycemic agents and glucagon.	
	e. ACTH and corticosteroids.	
V	Pharmacology of drugs acting on endocrine system	07
	a. Androgens and Anabolic steroids.	
	b. Estrogens, progesterone and oral contraceptives.	
	c. Drugs acting on the uterus.	
	Bioassay	
	a. Principles and applications of bioassay.	
	b.Types of bioassay	
	c. Bioassay of insulin, oxytocin, vasopressin, ACTH,d-tubocurarine,digitalis,	
	histamineand 5-HT	
		1



BP 507 P. PHARMACOLOGY-II (Practical)

Teaching and Examination Scheme:

Teac	ching Scl	neme	Credits	Marks			Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination	
					Exam			
0	0	4	2	15	35	50	3 hours	

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

- 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 (bySchilds 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-edemamodel.
- 15. Analgesic activity of drug using central and peripheral methods

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 McGraw-Hill.
- 3. Goodman and Gilman's, The Pharmacological Basis of Therapeutics

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- 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 PointLippincott 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 MedicalPublishers (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.



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BP504 T. PHARMACOGNOSY AND PHYTOCHEMISTRY II (Theory)

Teaching and Examination Scheme:

Teac	hing Sch	neme	Credits	Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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 to:

- know the modern extraction techniques, characterization and identification of theherbal drugs and phytoconstituents
- > understand the preparation and development of herbal formulation.
- > understand the herbal drug interactions
- carryout isolation and identification of phytoconstituents

COURSE CONTENT

UNIT	CONTENT	No. of
		Hrs.
I	Metabolic pathways in higher plants and their determination	07
	a) Brief study of basic metabolic pathways and formation of different secondary	
	metabolitesthrough these pathways- Shikimic acid pathway, Acetate pathways	
	and Amino acid pathway.	
	b) Study of utilization of radioactive isotopes in the investigation of Biogenetic	
	studies.	
II	General introduction, composition, chemistry & chemical classes, biosources,	14
	therapeuticuses and commercial applications of followingsecondary metabolites:	
	Alkaloids: Vinca, Rauwolfia, Belladonna, Opium,	
	Phenylpropanoids and Flavonoids: Lignans, Tea, Ruta	
	Steroids, Cardiac Glycosides & Triterpenoids: Liquorice, Dioscorea, Digitalis	

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	Volatile oils: Mentha, Clove, Cinnamon, Fennel, Coriander,								
	Tannins: Catechu, Pterocarpus								
	Resins: Benzoin, Guggul, Ginger, Asafoetida, Myrrh, Colophony								
	Glycosides: Senna, Aloes, Bitter Almond								
	Iridoids, Other terpenoids & Naphthaquinones: Gentian, Artemisia, taxus,								
	carotenoids								
III	Isolation, Identification and Analysis of Phytoconstituents	06							
	a) Terpenoids: Menthol, Citral, Artemisin								
	b) Glycosides: Glycyrhetinic acid & Rutin								
	c) Alkaloids: Atropine, Quinine, Reserpine, Caffeine								
	d) Resins: Podophyllotoxin, Curcumin								
IV	Industrial production, estimation and utilization of the following	10							
	phytoconstituents:Forskolin, Sennoside, Artemisinin, Diosgenin, Digoxin,								
	Atropine, Podophyllotoxin, Caffeine, Taxol, Vincristine and Vinblastine								
V	Basics of Phytochemistry	08							
	Modern methods of extraction, application of latest techniques like Spectroscopy,								
	chromatography and electrophoresis in the isolation, purification and								
	identification of crudedrugs.								

BP 508 P. PHARMACOGNOSY AND PHYTOCHEMISTRY II (Practical)

Teaching and Examination Scheme:

Teac	hing Sch	neme	Credits	Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
0	0	4	2	15	35	50	3 hours

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 Belladonnad. 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, NewDelhi.
- 5. Essentials of Pharmacognosy, Dr.SH.Ansari, IInd edition, Birla publications, NewDelhi, 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.

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BP 505 T. PHARMACEUTICAL JURISPRUDENCE (Theory)

Teaching and Examination Scheme:

Teac	eaching Scheme Credits			Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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:

- ➤ The Pharmaceutical legislations and their implications in the development andmarketing of pharmaceuticals.
- ➤ Various Indian pharmaceutical Acts and Laws
- > The regulatory authorities and agencies governing the manufacture and sale ofpharmaceuticals
- > The code of ethics during the pharmaceutical practice

COURSE CONTENT

UNIT	CONTENT	No. of
		Hrs.
I	Drugs and Cosmetics Act, 1940 and its rules 1945:	10
	Objectives, Definitions, Legal definitions of schedules to the Act andRules	
	Import of drugs – Classes of drugs and cosmetics prohibited from import, Import	
	underlicense or permit. Offences and penalties.	
	Manufacture of drugs – Prohibition of manufacture and sale of certain drugs,	
	Conditions for grant of license and conditions of license for manufacture of	
	drugs, Manufacture of drugs for test, examination and analysis, manufacture of	
	new drug, loanlicense and repacking license.	
II	Drugs and Cosmetics Act, 1940 and its rules 1945.	10
	Detailed study of Schedule G, H, M, N, P,T,U, V, X, Y, Part XII B, Sch F &	
	DMR (OA)Sale of Drugs - Wholesale, Retail sale and Restricted license.	
	Offences and penalties, Labeling & Packing of drugs- General labeling	

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	requirements and specimen labels fordrugs and cosmetics, List of permitted	
	colors. Offences and penalties.	
	Administration of the Act and Rules – Drugs Technical Advisory Board, Central	
	drugsLaboratory, Drugs Consultative Committee, Government drug analysts,	
	Licensingauthorities, controlling authorities, Drugs Inspectors	
III	Pharmacy Act –1948: Objectives, Definitions, Pharmacy Council of India;	10
	itsconstitution and functions, Education Regulations, State and Joint state	
	pharmacy	
	councils; constitution and functions, Registration of Pharmacists, Offences and	
	Penalties	
	Medicinal and Toilet Preparation Act -1955: Objectives, Definitions,	
	Licensing, Manufacture In bond and Outside bond, Export of alcoholic	
	preparations, Manufacture of Ayurvedic, Homeopathic, Patent & Proprietary	
	Preparations.Offences and Penalties.	
	Narcotic Drugs and Psychotropic substances Act-1985 and Rules: 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	
IV	Study of Salient Features of Drugs and Magic Remedies Act and its	08
	rules: Objectives, Definitions, Prohibition of certain advertisements, Classes of	
	Exempted advertisements, Offences and Penalties	
	Prevention of Cruelty to animals Act-1960: Objectives, Definitions,	
	InstitutionalAnimal Ethics Committee, CPCSEA guidelines for Breeding and	
	Stocking of Animals, Performance of Experiments, Transfer and acquisition of	
	animals forexperiment, Records, Power to suspend or revoke registration,	
	Offences and Penalties	
	National Pharmaceutical Pricing Authority: Drugs Price Control Order	
	(DPCO)-2013. Objectives, Definitions, Sale prices of bulk drugs, Retail price of	
	formulations,Retail price and ceiling price of scheduled formulations, National	
	List of EssentialMedicines (NLEM)	



Pharmaceutical Legislations – A brief review, Introduction, Study of drugs enquirycommittee, Health survey and development committee, Hathi committee andMudaliar committee

Code of Pharmaceutical ethics D efinition, Pharmacist in relation to his job, trade,medical profession and his profession, Pharmacist's oath

Medical Termination of Pregnancy Act

Right to Information Act

Introduction to Intellectual Property Rights (IPR)

Recommended books: (Latest Edition)

- 1. Forensic Pharmacy by B. Suresh
- 2. Text book of Forensic Pharmacy by B.M. Mithal
- 3. Hand book of drug law-byM.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)



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BP601T. MEDICINAL CHEMISTRY – III (Theory)

Teaching and Examination Scheme:

Teac	ching Scl	neme	Credits	Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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 actional drug design like quantitative structure activity relationship (QSAR), Prodrugconcept, combinatorial chemistry and Computer aided drug design (CADD). The subjectalso 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

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

COURSE CONTENT

Note: Study of the development of the following classes of drugs, Classification, mechanismofaction, 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 (*)

UNIT	CONTENT	No. of Hrs.							
Ι	Antibiotics	10							
	Historical background, Nomenclature, Stereochemistry, Structure								
	activityrelationship, Chemical degradation classification and important products								
	ofthe following classes.								
	β-Lactam antibiotics: Penicillin, Cepholosporins, $β$ - Lactamase								
	inhibitors, Monobactams								
	Aminoglycosides: Streptomycin, Neomycin, Kanamycin								
	Tetracyclines: Tetracycline, Oxytetracycline, Chlortetracycline, Minocycline,								

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	Doxycycline									
II	Antibiotics	10								
	Historical background, Nomenclature, Stereochemistry, Structure									
	activityrelationship, Chemical degradation classification and important products									
	of									
	the following classes.									
	Macrolide: Erythromycin Clarithromycin, Azithromycin.									
	Miscellaneous: Chloramphenicol*, Clindamycin.									
	Prodrugs: Basic concepts and application of prodrugs design.									
	Antimalarials: Etiology of malaria.									
	Quinolines: SAR, Quinine sulphate, Chloroquine*, Amodiaquine,Primaquine									
	phosphate, Pamaquine*, Quinacrine hydrochloride, Mefloquine.									
	Biguanides and dihydro triazines: Cycloguanil pamoate, Proguanil.									
	Miscellaneous: Pyrimethamine, Artesunete, Artemether, Atovoquone.									
III	Anti-tubercular Agents	10								
	Synthetic anti tubercular agents: Isoniozid*, Ethionamide,									
	Ethambutol, Pyrazinamide, Para amino salicylic acid.*									
	Anti tubercular antibiotics: Rifampicin, Rifabutin, Cycloserine, Streptomycine,									
	Capreomycin sulphate.									
	Urinary tract anti-infective agents									
	Quinolones: SAR of quinolones, Nalidixic Acid,Norfloxacin,									
	Enoxacin, Ciprofloxacin*, Ofloxacin, Lomefloxacin, Sparfloxacin,									
	Gatifloxacin, Moxifloxacin									
	Miscellaneous: Furazolidine, Nitrofurantoin*, Methanamine.									
	Antiviral agents:									
	Amantadine hydrochloride, Rimantadine hydrochloride, Idoxuridine, trifluoride,									
	Acyclovir*, Gancyclovir, Zidovudine, Didanosine, Zalcitabine,Lamivudine,									
	Loviride, Delavirding, Ribavirin, Saquinavir, Indinavir, Ritonavir.									
IV	Antifungal agents:	08								
	Antifungal antibiotics: Amphotericin-B, Nystatin, Natamycin, Griseofulvin.									
	Synthetic Antifungal agents: Clotrimazole, Econazole,									
	Butoconazole,Oxiconazole Tioconozole, Miconazole*, Ketoconazole,									



Terconazole, Itraconazole, Fluconazole, Naftifine hydrochloride, Tolnaftate*. **Anti-protozoal Agents:** Metronidazole*, Tinidazole, Ornidazole, Diloxanide, Iodoquinol, Pentamidine Isethionate, Atovaquone, Eflornithine. citrate*, **Anthelmintics:** Diethylcarbamazine Thiabendazole, Mebendazole*, Albendazole, Niclosamide, Oxamniquine, Praziquantal, Ivermectin. **Sulphonamides and Sulfones** 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*. V **Introduction to Drug Design** 07 Various approaches used in drug design. Physicochemical parameters used in quantitative structure activityrelationship (QSAR) such as partition coefficient, Hammet's electronic parameter, Tafts steric parameter and Hansch analysis. Pharmacophore modeling and docking techniques. Combinatorial Chemistry: Concept and applicationschemistry: solid phase and solution phase synthesis of combinatorial



BP607P. MEDICINAL CHEMISTRY- III (Practical)

Teaching and Examination Scheme:

Teac	hing Sch	g Scheme Credits		Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
0	0	4	2	15	35	50	3 hours

I Preparation of drugs and intermediates

Sulphanilamide, 7-Hydroxy, 4-methyl coumarin, Chlorobutanol, Triphenyl imidazole, Tolbutamide, Hexamine

II Assay of drugs

Isonicotinic acid hydrazide, Chloroquine, Metronidazole, Dapsone, Chlorpheniramine maleate Benzyl penicillin

III Preparation of medicinally important compounds or intermediates by Microwaveirradiation technique

IV Drawing structures and reactions using chem draw®

V Determination of physicochemical properties such as logP, clogP, MR, Molecularweight, Hydrogen bond donors and acceptors for class of drugs course contentusing 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.

BP602 T. PHARMACOLOGY-III (Theory)

Teaching and Examination Scheme:

Teaching Scheme Credit			Credits		Marks	Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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:

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

COURSE CONTENT

UNIT	CONTENT	No. of Hrs.
I	Pharmacology of drugs acting on Respiratory system	10
	a. Anti -asthmatic drugs	
	b. Drugs used in the management of COPD	
	c. Expectorants and antitussives	
	d. Nasal decongestants	
	e. Respiratory stimulants	
	Pharmacology of drugs acting on the Gastrointestinal Tract	
	a. Antiulcer agents.	
	b. Drugs for constipation and diarrhoea.	
	c. Appetite stimulants and suppressants.	
	d. Digestants and carminatives.	

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	e. Emetics and anti-emetics.	
II	Chemotherapy	10
	a. General principles of chemotherapy.	
	b. Sulfonamides and cotrimoxazole.	
	c. Antibiotics- Penicillins, cephalosporins, chloramphenicol,	
	macrolides,quinolones and fluoroquinolins, tetracycline and aminoglycosides	
III	Chemotherapy	10
	a. Antitubercular agents	
	b. Antileprotic agents	
	c. Antifungal agents	
	d. Antiviral drugs	
	e.Anthelmintics	
	f. Antimalarial drugs	
	g. Antiamoebic agents	
IV	Chemotherapy: Urinary tract infections and sexually transmitted diseases.	08
	Chemotherapy of malignancy	
	Immunopharmacology	
	a. Immunostimulants	
	b. Immunosuppressant	
	Protein drugs, monoclonal antibodies, target drugs to antigen, biosimilars	
V	Principles of toxicology	07
	a. Definition and basic knowledge of acute, subacute and chronic toxicity.	
	b. Definition and basic knowledge of genotoxicity, carcinogenicity,	
	teratogenicityand mutagenicity	
	c. General principles of treatment of poisoning	
	d. Clinical symptoms and management of barbiturates, morphine,	
	organophosphorus compound and lead, mercury and arsenic poisoning.	
	Chronopharmacology	
	a. Definition of rhythm and cycles.	
	b. Biological clock and their significance leading to chronotherapy.	



BP 608 P. PHARMACOLOGY-III (Practical)

Teaching and Examination Scheme:

Teaching Scheme C			Credits		Marks	Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
0	0	4	2	15	35	50	3 hours

^{*}Experiments are demonstrated by simulated experiments/videos

- 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, WilcoxonSigned Rank test)

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 McGraw-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 LippincottWilliams &Wilkins
- 5. Mycek M.J, Gelnet S.B and Perper M.M. Lippincott's Illustrated Reviews-Pharmacology

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- 6. K.D.Tripathi. Essentials of Medical Pharmacology, , JAYPEE Brothers MedicalPublishers (P) Ltd, New Delhi.
- 7. Sharma H. L., Sharma K. K., Principles of Pharmacology, Paras medical publisherModern 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.



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BP 603 T. HERBAL DRUG TECHNOLOGY (Theory)

Teaching and Examination Scheme:

Teaching Scheme			Credits		Duration of End Semester		
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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 ManufacturingPractices (GMP), patenting and regulatory issues of herbal drugs.

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

- > understand raw material as source of herbal drugs from cultivation to herbal drugproduct
- ➤ know the WHO and ICH guidelines for evaluation of herbal drugs
- know the herbal cosmetics, natural sweeteners, nutraceuticals
- > appreciate patenting of herbal drugs, GMP.

COURSE CONTENT

UNIT	CONTENT	No. of							
		Hrs.							
I	Herbs as raw materials								
	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								
	Biodynamic Agriculture								
	Good agricultural practices in cultivation of medicinal plants including Organic								
	farming.Pest and Pest management in medicinal plants:								
	Biopesticides/Bioinsecticides.								
	Indian Systems of Medicine								
	a) Basic principles involved in Ayurveda, Siddha, Unani and Homeopathy								
	b) Preparation and standardization of Ayurvedic formulations viz Aristas and								
	Asawas,Ghutika,Churna, Lehya and Bhasma.								

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II	Nutraceuticals	07
	General aspects, Market, growth, scope and types of products available in the	
	market. Healthbenefits and role of Nutraceuticals in ailments like Diabetes, CVS	
	diseases, Cancer, Irritablebowel syndrome and various Gastro intestinal diseases.	
	Study of following herbs as health food: Alfaalfa, Chicory, Ginger, Fenugreek,	
	Garlic, Honey, Amla, Ginseng, Ashwagandha, Spirulina	
	Herbal-Drug and Herb-Food Interactions: General introduction to interaction	
	and classification. Study of following drugs and their possible side effects and	
	interactions:Hypercium, kava-kava, Ginkobiloba, Ginseng, Garlic, Pepper &	
	Ephedra.	
III	Herbal Cosmetics	10
	Sources and description of raw materials of herbal origin used via, fixed oils,	
	waxes, gumscolours, perfumes, protective agents, bleaching agents, antioxidants	
	in products such as skincare, hair care and oral hygiene products.	
	Herbal excipients:	
	Herbal Excipients – Significance of substances of natural origin as excipients –	
	colorants, sweeteners, binders, diluents, viscosity builders, disintegrants, flavors	
	& perfumes.	
	Herbal formulations:	
	Conventional herbal formulations like syrups, mixtures and tablets and Novel	
	dosage formslike phytosomes	
IV	Evaluation of Drugs WHO & ICH guidelines for the assessment of herbal drugs	10
	Stability testing of herbal drugs.	
	Patenting and Regulatory requirements of natural products:	
	a) Definition of the terms: Patent, IPR, Farmers right, Breeder's right,	
	Bioprospecting andBiopiracy	
	b) Patenting aspects of Traditional Knowledge and Natural Products. Case study	
	of Curcuma& Neem.	
	· · · · · · · · · · · · · · · · · · ·	
	Regulatory Issues - Regulations in India (ASU DTAB, ASU DCC), Regulation	
	Regulatory Issues - Regulations in India (ASU DTAB, ASU DCC), Regulation ofmanufacture of ASU drugs - Schedule Z of Drugs & Cosmetics Act for ASU	



Herbal drugs industry: Present scope and future prospects.

A brief account of plant based industries and institutions involved in work on medicinal andaromatic plants in India.

Schedule T – GoodManufacturing Practice of Indian systems of medicine

Components of GMP (Schedule – T) and its objectivesInfrastructural requirements, working space, storage area, machinery and equipments, standard operating procedures, health and hygiene, documentation and records.



BP 609 P. HERBAL DRUG TECHNOLOGY (Practical)

Teaching and Examination Scheme:

Teaching Scheme C			Credits		Marks	Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
0	0	4	2	15	35	50	3 hours

- 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 inIndian Medicine & Homeopathy)
- 7. Mukherjee, P.W. Quality Control of Herbal Drugs: An Approach to Evaluation ofBotanicals. Business Horizons Publishers, New Delhi, India, 2002.



BP 604 T. BIOPHARMACEUTICS AND PHARMACOKINETICS (Theory)

Teaching and Examination Scheme:

Teaching Scheme Credits			Credits		Marks	Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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:

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

COURSECONTENT

UNIT	CONTENT	No. of						
I	Introduction to Biopharmaceutics							
	Absorption; Mechanisms of drug absorption through GIT, factors influencing							
	drugabsorption though GIT, absorption of drug from Non per oral extra-vascular							
	routes, Distribution Tissue permeability of drugs, binding of drugs, apparent,							
	volumeof drug distribution, plasma and tissue protein binding of drugs, factors							
	affectingprotein-drug binding. Kinetics of protein binding, Clinical significance							
	of proteinbinding of drugs							
II	Elimination: Drug metabolism and basic understanding metabolic pathways	10						
	renalexcretion of drugs, factors affecting renal excretion of drugs, renal							
	clearance, Non renalroutes of drug excretion of drugs							

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	Bioavailability and Bioequivalence: Definition and Objectives of								
	bioavailability, absolute and relative bioavailability, measurement of								
	bioavailability, in-vitro drugdissolution models, in-vitro-in-vivo correlations,								
	bioequivalence studies, methods toenhance the dissolution rates and								
	bioavailability of poorly soluble drugs.								
III	Pharmacokinetics: Definition and introduction to Pharmacokinetics,	10							
	Compartment models, Physiological models, One								
	compartment openmodel. (a). Intravenous Injection (Bolus) (b). Intravenous								
	infusion and (c) Extravascular administrations. Pharmacokinetics parameters -								
	KE ,t1/2,Vd,AUC,Ka, Clt andCLR- definitions methods of eliminations,								
	understanding of their significance and application								
IV	Multicompartment models: Two compartment open model. IV bolusKinetics of								
	multiple dosing, steady state drug levels, calculation of loading andmainetnance								
	doses and their significance in clinical settins.								
V	Nonlinear Pharmacokinetics: a. Introduction, b. Factors causing Non-linearity.	07							
	c. Michaelis-menton method of estimating parameters, Explanation with example								
	ofdrugs.								

Recommended Books: (Latest Editions)

- 1. Biopharmaceutics and Clinical Pharmacokinetics by, Milo Gibaldi.
- 2. Biopharmaceutics and Pharmacokinetics; By Robert F Notari
- 3. Applied biopharmaceutics and pharmacokinetics, Leon Shargel and AndrewB.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: ByMilo Glbaldi Donald, R. Mercel Dekker Inc.
- Hand Book of Clinical Pharmacokinetics, ByMilo Gibaldi and Laurie Prescott byADIS Health Science Press.
- 7. Biopharmaceutics; By Swarbrick
- 8. Clinical Pharmacokinetics, Concepts and Applications: ByMalcolm 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 editionRevised and expanded by Rebort F Notari Marcel Dekker Inn, New York and Basel, 1987.
- 12. Remington's Pharmaceutical Sciences, ByMack Publishing Company, Pennsylvnia

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BP 605 T. PHARMACEUTICAL BIOTECHNOLOGY (Theory)

Teaching and Examination Scheme:

Teaching Scheme Credit			Credits		Marks	Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

Scope:Biotechnology has a long promise to revolutionize the biological sciences andtechnology. Scientific application of biotechnology in the field of genetic engineering, medicine and fermentation technologymakes the subject interesting. Biotechnology is leading to new biological revolutions in diagnosis, preventionand 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;

- ➤ Understanding the importance of Immobilized enzymes in PharmaceuticalIndustries
- ➤ Genetic engineering applications in relation to production of pharmaceuticals
- ➤ Importance of Monoclonal antibodies in Industries
- Appreciate the use of microorganisms in fermentation technology

UNIT	CONTENT	No. of						
		Hrs.						
I	a) Brief introduction to Biotechnology with reference to Pharmaceutical Sciences.	10						
	b) Enzyme Biotechnology- Methods of enzyme immobilization and applications.							
	c) Biosensors- Working and applications of biosensors in Pharmaceutical Industries.							
	d) Brief introduction to Protein Engineering.							
	e) Use of microbes in industry. Production of Enzymes- General consideration -							
	Amylase, Catalase, Peroxidase, Lipase, Protease, Penicillinase.							
	f) Basic principles of genetic engineering.							
II	a) Study of cloning vectors, restriction endonucleases and DNA ligase.	10						
	b) Recombinant DNA technology. Application of genetic engineering in medicine.							
	c) Application of r DNA technology and genetic engineering in the production of:i)							
	Interferon ii) Vaccines- hepatitis- B iii) Hormones-Insulin.							
	d) Brief introduction to PCR							

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III	Types of immunity- humoral immunity, cellular immunity	10					
	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 Substituties.						
IV	IV a) Immuno blotting techniques- ELISA, Western blotting, Southern blotting.						
	b) Genetic organization of Eukaryotes and Prokaryotes						
	c) Microbial genetics including transformation, transduction, conjugation, plasmids						
	andtransposons.d) Introduction to Microbial biotransformation and applications.e)						
	Mutation: Types of mutation/mutants.						
V	a) Fermentation methods and general requirements, study of media, equipments,	07					
	sterilization methods, aeration process, stirring.						
	b) Large scale production fermenter design and its various controls.c) Study of the						
	production of - penicillins, citric acid, Vitamin B12, Glutamic acid, Griseofulvin,						
	d) Blood Products: Collection, Processing and Storage of whole human blood,						
	driedhuman plasma, plasma Substituties.						

Recommended Books (Latest edition):

- 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 RoyalSociety 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)

Teaching and Examination Scheme:

	Teaching Scheme			Credits	Marks			Duration of End Semester
•	L	T	P	C	Sessional	End Semester	Total	Examination
						Exam		
	3	1	0	4	25	75	100	3 hours

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

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

- > understand the cGMP aspects in a pharmaceutical industry
- > appreciate the importance of documentation
- > understand the scope of quality certifications applicable to pharmaceuticalindustries
- > understand the responsibilities of QA & QC departments

COURSE CONTENT

UNIT	CONTENT	No. of Hrs.								
I	Quality Assurance and Quality Management concepts: Definition and	10								
	concept of Qualitycontrol, Quality assurance and GMP									
	Total Quality Management (TQM): Definition, elements, philosophies									
	ICH Guidelines: purpose, participants, process of harmonization, Brief overview									
	of QSEM, with special emphasis on Q-series guidelines, ICH stability testing									
	guidelines									
	Quality by design (QbD): Definition, overview, elements of QbD program, tools									
	ISO 9000 & ISO14000: Overview, Benefits, Elements, steps for registration									
	NABL accreditation: Principles and procedures									
II	Organization and personnel: Personnel responsibilities, training, hygiene and	10								
	personal records.									
	Premises: Design, construction and plant layout, maintenance, sanitation,									
	environmentalcontrol, utilities and maintenance of sterile areas, control of									
	contamination.									

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	Equipments and raw materials: Equipment selection, purchase specifications,									
	maintenance, purchase specifications and maintenance of stores for raw materials.									
III	Quality Control: Quality control test for containers, rubber closures and	10								
	secondary packingmaterials.									
	Good Laboratory Practices: General Provisions, Organization and Personnel,									
	Facilities, Equipment, Testing Facilities Operation, Test and Control Articles,									
	Protocol for Conduct of aNonclinical Laboratory Study, Records and Reports,									
	Disqualification of Testing Facilities									
IV	Complaints: Complaints and evaluation of complaints, Handling of return good,									
	recalling andwaste disposal.									
	Document maintenance in pharmaceutical industry: Batch Formula Record,									
	Master FormulaRecord, SOP, Quality audit, Quality Review and Quality									
	documentation, Reports anddocuments, distribution records.									
V	Calibration and Validation: Introduction, definition and general principles of	07								
	calibration, qualification and validation, importance and scope of validation, types									
	of validation, validationmaster plan. Calibration of pH meter, Qualification of									
	UV-Visible spectrophotometer, Generalprinciples of Analytical method									
	Validation. Warehousing: Good warehousing practice, materials management									

Recommended Books: (Latest Edition)

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

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7th SEMESTER

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BP701T. INSTRUMENTAL METHODS OF ANALYSIS (Theory)

Teaching and Examination Scheme:

Teac	Teaching Scheme Credits			Marks	Duration of End Semester		
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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

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

- Understand the interaction of matter with electromagnetic radiations and itsapplications in drug analysis
- ➤ Understand the chromatographic separation and analysis of drugs.
- Perform quantitative & qualitative analysis of drugs using various analyticalinstruments.

COURSE CONTENT

UNIT	CONTENT	No. of						
		Hrs.						
I	UV Visible spectroscopy	10						
	Electronic transitions, chromophores, auxochromes, spectral shifts, solvent effect							
	onabsorption spectra, Beer and Lambert's law, Derivation and deviations.							
	Instrumentation - Sources of radiation, wavelength selectors, sample cells,							
	detectors-Photo tube, Photomultiplier tube, Photo voltaic cell, Silicon							
	Photodiode.Applications - Spectrophotometric titrations, Single component and							
	multi componentanalysis							
	Fluorimetry							
	Theory, Concepts of singlet, doublet and triplet electronic states, internal and							
	externalconversions, factors affecting fluorescence, quenching, instrumentation							
	andapplications							
II	IR spectroscopy: Introduction, fundamental modes of vibrations in poly atomic	10						

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10
08
07



BP705P. INSTRUMENTAL METHODS OF ANALYSIS (Practical)

Teaching and Examination Scheme:

Teac	ching Scl	neme	Credits		Marks		Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
0	0	4	2	15	35	50	3 hours

- 1 Determination of absorption maxima and effect of solvents on absorptionmaxima 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
- 10 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



BP 702 T. INDUSTRIAL PHARMACYII (Theory)

Teaching and Examination Scheme:

Teac	hing Sch	neme	Credits		Marks		Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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:

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

Course Content:

UNIT	CONTENT									
I	Pilot plant scale up techniques: General considerations - including significance	10								
	ofpersonnel requirements, space requirements, raw materials, Pilot plant scale up									
	considerations for solids, liquid orals, semi solids and relevant documentation,									
	SUPACguidelines, Introduction to platform technology									
II	Technology development and transfer: WHO guidelines for Technology									
	Transfer(TT):Terminology, Technology transfer protocol, Quality risk									
	management, Transfer from R& D to production (Process, packaging and									
	cleaning), Granularity of TT Process (API, excipients, finished products,									
	packaging materials) Documentation, Premises andequipments, qualification and									
	validation, quality control, analytical method transfer, Approved regulatory									
	bodies and agencies, Commercialization - practical aspects and problems (case									
	studies), TT agencies in India - APCTD, NRDC, TIFAC, BCIL, TBSE /SIDBI;									
	TT related documentation - confidentiality agreement, licensing, MoUs,legal									
	issues									
III	Regulatory affairs: Introduction, Historical overview of Regulatory Affairs,	10								

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	Regulatoryauthorities, Role of Regulatory affairs department, Responsibility of									
	Regulatory AffairsProfessionals									
	Regulatory requirements for drug approval: Drug Development Teams, Non-									
	ClinicalDrug Development, Pharmacology, Drug Metabolism and Toxicology,									
	Generalconsiderations of Investigational New Drug (IND) Application,									
	Investigator's Brochure(IB) and New Drug Application (NDA), Clinical research									
	/ BE studies, Clinical ResearchProtocols, Biostatistics in Pharmaceutical Product									
	Development, Data Presentation forFDA Submissions, Management of Clinical									
	Studies.									
IV	Quality management systems: Quality management & Certifications: Concept									
	ofQuality, Total Quality Management, Quality by Design (QbD), Six Sigma									
	concept, Outof Specifications (OOS), Change control, Introduction to ISO 9000									
	series of qualitysystems standards, ISO 14000, NABL, GLP									
V	Indian Regulatory Requirements: Central Drug Standard Control Organization									
	(CDSCO) and State Licensing Authority: Organization, Responsibilities,									
	Certificate ofPharmaceutical Product (COPP), Regulatory requirements and									
	approval procedures forNew Drugs.									

Recommended Books: (Latest Editions)

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



BP 703T. PHARMACY PRACTICE (Theory)

Teaching and Examination Scheme:

Teac	Teaching Scheme Credits			Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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. Incommunity 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

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

UNIT	CONTENT						
		Hrs.					
I	a) Hospital and it's organization: Definition, Classification of hospital-	10					
	Primary, Secondary and Tertiary hospitals,						
	Classification based on clinical and non- clinical basis, Organization Structure of						
	aHospital, and Medical staffs involved in the hospital and their functions.						
	b) Hospital pharmacy and its organization: Definition, functions of hospital						
	pharmacy, Organization structure, Location, Layoutand staff requirements, and						

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	Responsibilities and functions of hospital pharmacists.	
	c) Adverse drug reaction: Classifications - Excessive pharmacological effects,	
	secondary pharmacological	
	effects, idiosyncrasy, allergic drug reactions, genetically determined toxicity,	
	toxicityfollowing sudden withdrawal of drugs, Drug interaction- beneficial	
	interactions, adverse interactions, and pharmacokinetic drug interactions, Methods	
	for detectingdrug interactions, spontaneous case reports and record linkage	
	studies, and Adversedrug reaction reporting and management.	
	d) Community Pharmacy: Organization and structure of retail and wholesale	
	drug store, types and design, Legalrequirements for establishment and	
	maintenance of a drug store, Dispensing ofproprietary products, maintenance of	
	records of retail and wholesale drug store.	
II	a) Drug distribution system in a hospital: Dispensing of drugs to inpatients,	10
	types of drug distribution systems, charging policyand labelling, Dispensing of	
	drugs to ambulatory patients, and Dispensing of controlled drugs.	
	b) Hospital formulary: Definition, contents of hospital formulary,	
	Differentiation of hospital formulary andDrug list, preparation and revision, and	
	addition and deletion of drug from hospitalformulary.	
	c) Therapeutic drug monitoring: Need for Therapeutic Drug Monitoring,	
	Factors to be considered during the Therapeutic Drug Monitoring, and Indian	
	scenario for Therapeutic Drug Monitoring.	
	d) Medication adherence: Causes of medication non-adherence, pharmacist role	
	in the medication adherence, and monitoring of patient medication adherence.	
	e) Patient medication history interview: Need for the patient medication	
	history interview, medication interview forms.	
	f) Community pharmacy management: Financial, materials, staff, and	
	infrastructure requirements.	
III	a) Pharmacy and therapeutic committee: Organization, functions, Policies of	10
	the pharmacy and therapeutic committee in	
	including drugs into formulary, inpatient and outpatient prescription, automatic	
	stoporder, and emergency drug list preparation.	
	1	



	b) Druginformation services: Drug and Poison information centre, Sources of	
	drug information, Computerised	
	services, and storage and retrieval of information.	
	c) Patientcounseling: Definition of patient counseling; steps involved in patient	
	counseling, and Special	
	cases that require the pharmacist	
	d) Education and training program in the hospital: Role of pharmacist in the	
	education and training program, Internal and external	
	training program, Services to the nursing homes/clinics, Code of ethics for	
	communitypharmacy, and Role of pharmacist in the interdepartmental	
	communication and community health education.	
	e) Prescribed medication order and communication skills: Prescribed	
	medication order- interpretation and legal requirements, andCommunication	
	skills- communication with prescribers and patients.	
IV	a) Budgetpreparation and implementation: Budget preparation and	08
	implementation	
	b) Clinical Pharmacy: Introduction to Clinical Pharmacy, Concept of clinical	
	pharmacy, functions and responsibilities of clinical pharmacist, Drug therapy	
	monitoring - medication chartreview, clinical review, pharmacist intervention,	
	Ward round participation, Medicationhistory and Pharmaceutical care.Dosing	
	pattern and drug therapy based on Pharmacokinetic & disease pattern.	
	c) Over the counter (OTC) sales: Introduction and sale of over the counter, and	
	Rational use of common over the	
	counter medications.	
V	a) Drug store management and inventory control: Organisation of drug store,	07
	types of materials stocked and storage conditions, Purchaseand inventory control:	
	principles, purchase procedure, purchase order, procurementand stocking,	
	Economic order quantity, Reorder quantity level, and Methods used forthe	
	analysis of the drug expenditure	
	b) Investigational use of drugs: Description, principles involved, classification,	
	control, identification, role of hospitalpharmacist, advisory committee.	
	c) Interpretation of Clinical Laboratory Tests: Blood chemistry, hematology,	
	1	



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*, 1st ed. 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 ofHealth System Pharmacists Inc; 2009.
- 6. Parmar N.S. *Health Education and Community Pharmacy*, 18th ed. India: CBSPublishers & 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)



BP 704T: NOVEL DRUG DELIVERY SYSTEMS (Theory)

Teaching and Examination Scheme:

Teac	Teaching Scheme Credits			Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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

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

COURSE CONTENT

UNIT	CONTENT	No. of Hrs.						
I	Controlled drug delivery systems: Introduction, terminology/definitions and	10						
	rationale,advantages, disadvantages, selection of drug candidates. Approaches to							
	design controlledrelease formulations based on diffusion, dissolution and ion							
	exchange principles.Physicochemical and biological properties of drugs relevant							
	to controlled releaseformulations							
	Polymers: Introduction, classification, properties, advantages and application of							
	polymers in formulation of controlled release drug delivery systems.							
II	Microencapsulation: Definition, advantages and disadvantages,	10						
	microspheres/microcapsules, microparticles, methods of microencapsulation,							
	applications Mucosal Drug Delivery system: Introduction, Principles of							
	bioadhesion /mucoadhesion, concepts, advantages and disadvantages,							
	transmucosal permeability andformulation considerations of buccal delivery							
	systems							
	Implantable Drug Delivery Systems: Introduction, advantages and							
	disadvantages,concept of implantsand osmotic pump							
III	Transdermal Drug Delivery Systems: Introduction, Permeation through skin,	10						

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	factorsaffecting permeation, permeation enhancers, basic components of TDDS,	
	formulationapproaches	
	Gastroretentive drug delivery systems: Introduction, advantages,	
	disadvantages,approaches for GRDDS - Floating, high density systems,	
	inflatable and gastroadhesivesystems and their applications	
	Nasopulmonary drug delivery system: Introduction to Nasal and Pulmonary	
	routes ofdrug delivery, Formulation of Inhalers (dry powder and metered dose),	
	nasal sprays,nebulizers	
IV	Targeted drug Delivery: Concepts and approaches advantages and	08
	disadvantages, introduction to liposomes, niosomes, nanoparticles, monoclonal	
	antibodies and theirapplications	
V	Ocular Drug Delivery Systems: Introduction, intra ocular barriers and methods	07
	toovercome –Preliminary study, ocular formulations and ocuserts	
	Intrauterine Drug Delivery Systems: Introduction, advantages and	
	disadvantages, development of intra uterine devices (IUDs) and applications	

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 WileyInterscience 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)



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BP801T. BIOSTATISITCS AND RESEARCH METHODOLOGY (Theory)

Teaching and Examination Scheme:

Teaching Scheme			Credits		Marks	Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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

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

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

COURSE CONTENT

UNIT	CONTENT	No. of					
		Hrs.					
I	Introduction: Statistics, Biostatistics, Frequency distribution	10					
	Measures of central tendency: Mean, Median, Mode- Pharmaceutical examples						
	Measures of dispersion: Dispersion, Range, standard deviation, Pharmaceutical						
	problems						
	Correlation: Definition, Karl Pearson's coefficient of correlation, Multiple						
	correlation -Pharmaceuticals examples						
II	Regression: Curve fitting by the method of least squares, fitting the lines y= a +	10					
	bx and x= a + by, Multiple regression, standard error of regression-						
	Pharmaceutical Examples						
	Probability: Definition of probability, Binomial distribution, Normal distribution,						
	Poisson's distribution, properties – problems						

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	Sample, Population, large sample, small sample, Null hypothesis, alternative	
	hypothesis, sampling, essence of sampling, types of sampling, Error-I type, Error-	
	II type, Standarderror of mean (SEM) - Pharmaceutical examples	
	Parametric test: t-test(Sample, Pooled or Unpaired and Paired), ANOVA, (One	
	wayand Two way), Least Significance difference	
III	Non Parametric tests: Wilcoxon Rank Sum Test, Mann-Whitney U test,	10
	Kruskal-Wallistest, Friedman Test	
	Introduction to Research: Need for research, Need for design of Experiments,	
	Experiential Design Technique, plagiarism	
	Graphs: Histogram, Pie Chart, Cubic Graph, response surface plot, Counter Plot	
	graph	
	Designing the methodology: Sample size determination and Power of a study,	
	Reportwriting and presentation of data, Protocol, Cohorts studies, Observational	
	studies, Experimental studies, Designing clinical trial, various phases.	
IV	Blocking and confounding system for Two-level factorials	08
	Regression modeling: Hypothesis testing in Simple and Multiple	
	regressionmodels	
	Introduction to Practical components of Industrial and Clinical Trials	
	Problems:Statistical Analysis Using Excel, SPSS, MINITAB®, DESIGN OF	
	EXPERIMENTS, R -Online Statistical Software's to Industrial and Clinical trial	
	approach	
V	Design and Analysis of experiments:	07
	Factorial Design: Definition, 22, 23design. Advantage of factorial design	
	Response Surface methodology: Central composite design, Historical design,	
	Optimization Techniques	
1		1

Recommended Books (Latest edition):

- 1. Pharmaceutical statistics- Practical and clinical applications, Sanford Bolton, publisher Marcel Dekker Inc. New York.
- 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



BP 802T: SOCIAL AND PREVENTIVE PHARMACY

Teaching and Examination Scheme:

Teac	Teaching Scheme Credits			Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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. Theroles of the pharmacist in these contexts are also discussed.

Objectives:

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

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

COURSE CONTENT

UNIT	CONTENT	No. of
I	Concept of health and disease: Definition, concepts and evaluation of public	Hrs. 10
	health.Understanding the concept of prevention and control of disease, social	
	causes of diseasesand social problems of the sick.	
	Social and health education: Food in relation to nutrition and health, Balanced	
	diet, Nutritional deficiencies, Vitamin deficiencies, Malnutrition and its	
	prevention.	
	Sociology and health: Socio cultural factors related to health and disease,	
	Impact of urbanization on health and disease, Poverty and health	
	Hygiene and health: personal hygiene and health care; avoidable habits	
II	Preventive medicine: General principles of prevention and control of diseases	10
	such ascholera, SARS, Ebola virus, influenza, acute respiratory infections,	
	malaria, chickenguinea, dengue, lymphatic filariasis, pneumonia, hypertension,	

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	diabetes mellitus, cancer,drug addiction-drug substance abuse								
III	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, Nationalprogramme for prevention and control of								
	deafness, Universal immunization programme, National programme for control of								
	blindness, Pulse polio programme.								
IV	National health intervention programme for mother and child, National family								
	welfareprogramme, National tobacco control programme, National Malaria								
	Prevention Program, National programme for the health care for the elderly,								
	Social health programme; role of WHO in Indian national program								
V	Community services in rural, urban and school health: Functions of PHC,	07							
	Improvementin rural sanitation, national urban health mission, Health promotion								
	and education inschool.								

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 RoyRabindra Nath, Saha Indranil, 4th Edition, 2013, ISBN: 9789350901878, JAYPEEPublications
- 3. Review of Preventive and Social Medicine (Including Biostatistics), Jain Vivek, 6thEdition, 2014, ISBN: 9789351522331, JAYPEE Publications
- 4. Essentials of Community Medicine—A Practical Approach, Hiremath Lalita D,Hiremath Dhananjaya A, 2nd Edition, 2012, ISBN: 9789350250440, JAYPEEPublications
- 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. PHARMA MARKETING MANAGEMENT (Theory)

Teaching and Examination Scheme:

Teac	hing Sch	neme	Credits		Marks		Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

Scope: The pharmaceutical industry not only needs highly qualified researchers, chemists and, technical people, but also requires skilled managers who can take the industry forwardby 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.

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

COURSE CONTENT

UNIT	CONTENT	No. of Hrs.							
I	Marketing: Definition, general concepts and scope of marketing; Distinction	10							
	between marketing &selling Marketing environment; Industry and competitive								
	analysis; Analyzing consumerbuying behavior; industrial buying behavior.								
	Pharmaceutical market: Quantitative and qualitative aspects; size and								
	composition of the market; demographicdescriptions and socio-psychological								
	characteristics of the consumer; marketsegmentation& targeting.Consumer								
	profile; Motivation and prescribing habits of thephysician; patients' choice of								
	physician and retail pharmacist. Analyzing the Market; Roleof market research.								
II	Product decision: Classification, product line and product mix decisions, product lifecycle, product portfolio analysis; product positioning; New product decisions; Productbranding, packaging and labeling decisions, Product management in pharmaceuticalindustry.	10							
III	Promotion: Methods, determinants of promotional mix, promotional budget; An	10							

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	overview of								
	personal selling, advertising, direct mail, journals, sampling, retailing,								
	medicalexhibition, public relations, online promotional techniques for OTC								
	Products.								
IV	Pharmaceutical marketing channels: Designing channel, channel members,	10							
	selecting the appropriate channel, conflict inchannels, physical distribution								
	management: Strategic importance, tasks in physical distribution management.								
	Professional sales representative (PSR): Duties of PSR, purpose of detailing,								
	selection and training, supervising, norms forcustomer calls, motivating,								
	evaluating, compensation and future prospects of the PSR.								
V	Pricing: Meaning, importance, objectives, determinants of price; pricing	10							
	methods and strategies, issues in price management in pharmaceutical industry.								
	An overview of DPCO(Drug Price Control Order)and NPPA (National								
	Pharmaceutical Pricing Authority).								
	Emerging concepts in marketing: Vertical & Horizontal Marketing;								
	RuralMarketing; Consumerism; Industrial Marketing; Global Marketing.								

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, TataMC 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) ExcelPublications.



BP804 ET: PHARMACEUTICAL REGULATORY SCIENCE (Theory)

Teaching and Examination Scheme:

Teac	Teaching Scheme		Credits	Marks			Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination	
					Exam			
3	1	0	4	25	75	100	3 hours	

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

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

- ➤ Know about the process of drug discovery and development
- ➤ Know the regulatory authorities and agencies governing the manufacture and saleof pharmaceuticals
- ➤ Know the regulatory approval process and their registration in Indian andinternational markets

COURSE CONTENT

UNIT	CONTENT	No. of Hrs.
I	New Drug Discovery and development	10
	Stages of drug discovery, Drug development process, pre-clinical studies, non-	
	clinicalactivities, clinical studies, Innovator and generics, Concept of generics,	
	Generic drugproduct development.	
II	Regulatory Approval Process	10
	Approval processes and timelines involved in Investigational New Drug (IND),	
	NewDrug Application (NDA), Abbreviated New Drug Application (ANDA).	
	Changes to anapproved NDA / ANDA.	
	Regulatory authorities and agencies	
	Overview of regulatory authorities of India, United States, European Union,	

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	Australia, Japan, Canada (Organization structure and types of applications)							
III	Registration of Indian drug product in overseas market							
	Procedure for export of pharmaceutical products, Technical documentation, Drug							
	MasterFiles (DMF), Common Technical Document (CTD), electronic Common							
	TechnicalDocument (eCTD), ASEAN Common Technical Document							
	(ACTD)research.							
IV	Clinical trials	08						
	Developing clinical trial protocols, Institutional Review Board / Independent							
	Ethicscommittee - formation and working procedures, Informed consent process							
	andprocedures, GCP obligations of Investigators, sponsors & Monitors,							
	Managing andMonitoring clinical trials, Pharmacovigilance - safetymonitoring in							
	clinical trials							
V	Regulatory Concepts	07						
	Basic terminology, guidance, guidelines, regulations, Laws and Acts, Orange							
	book, Federal Register, Code of Federal Regulatory, Purple book							

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 Healthcare Publishers.
- 3. New Drug Approval Process: Accelerating Global Registrations By Richard AGuarino, MD, 5th edition, Drugs and the Pharmaceutical Sciences, Vol. 190.
- 4. Guidebook for drug regulatory submissions / SandyWeinberg. By John Wiley &Sons. Inc.
- 5. FDA Regulatory Affairs: a guide for prescription drugs, medical devices, andbiologics /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 ComplianceBy 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

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BP 805T: PHARMACOVIGILANCE (Theory)

Teaching and Examination Scheme:

Teac	eaching Scheme Credits			Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

Scope: This paper will provide an opportunity for the student to learn about development ofpharmacovigilance as a science, basic terminologies used in pharmacovigilance, global scenarioof Pharmacovigilance, train students on establishing pharmacovigilance programme in anorganization, 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, andappreciate):

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

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COURSE CONTENT

UNIT	CONTENT	No. of Hrs.								
I	Introduction to Pharmacovigilance: History and development of	10								
	Pharmacovigilance; Importance of safety monitoring of Medicine; WHO									
	international drug monitoring programme; Pharmacovigilance Program of									
	India(PvPI)									
	Introduction to adverse drug reactions: 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									
	Basic terminologies used in pharmacovigilance: Terminologies of adverse									
	medication related events; Regulatory terminologies									
II	Drug and disease classification: Anatomical, therapeutic and chemical	10								
	classification of drugs; International classification of diseases; Daily defined									
	doses; International Non proprietary Names for drugs									
	Drug dictionaries and coding in pharmacovigilance: WHO adverse reaction									
	terminologies; MedDRA and Standardised MedDRA queries; WHO drug									
	dictionary; Eudravigilance medicinal product dictionary									
	Information resources in pharmacovigilance: Basic drug information									
	resources; Specialised resources for ADRs									
	Establishing pharmacovigilance programme: Establishing in a hospital,									
	Establishment & operation of drug safety department in industry; Contract									
	Research Organisations (CROs); Establishing a national programme									
III	Vaccine safety surveillance: Vaccine Pharmacovigilance; Vaccination failure;	10								
	Adverse events following immunization									
	Pharmacovigilance methods: 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									
	Communication in pharmacovigilance: Effective communication in									
	Pharmacovigilance; Communication in Drug Safety Crisis management;									





	Communicating with Regulatory Agencies, Business Partners, Healthcare									
	facilities &Media									
IV	Safety data generation: Pre clinical phase; Clinical phase; Post approval phase	08								
	(PMS)									
	ICH Guidelines for Pharmacovigilance: 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									
V	Pharmacogenomics of adverse drug reactions: Genetics related ADR with									
	example focusing PK parameters.									
	Drug safety evaluation in special population: Paediatrics; Pregnancy and									
	lactation; Geriatrics									
	CIOMS: CIOMS Working Groups; CIOMS Form									
1	CDSCO (India) and Pharmacovigilance: D&C Act and Schedule Y;									
	, ,	I								
	Drug safety evaluation in special population: Paediatrics; Pregnancy and lactation; Geriatrics CIOMS: CIOMS Working Groups; CIOMS Form									

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 andBartlett 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 PKManna
- 12. http://www.whoumc.org/DynPage.aspx?id=105825&mn1=7347&mn2=7259&mn3=7297
- 13. http://www.ich.org/

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- 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

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BP 806 ET. QUALITY CONTROL AND STANDARDIZATION OF HERBALS(Theory)

Teaching and Examination Scheme:

Teac	ching Scheme Credits			Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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;

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

COURSE CONTENT

UNIT	CONTENT	No. of Hrs.						
I	Basic tests for drugs – Pharmaceutical substances, Medicinal plants materials and	10						
	dosageforms, WHO guidelines for quality control of herbal drugs.							
	Evaluation of commercial crude drugs intended for use							
II	Quality assurance in herbal drug industry of cGMP, GAP, GMP and GLP in							
	traditional system of medicine.WHO Guidelines on current good manufacturing							
	Practices (cGMP) for Herbal Medicines. WHO Guidelines on GACP for							
	Medicinal Plants.							
III	EU and ICH guidelines for quality control of herbal drugs.	10						
	Research Guidelines for Evaluating the Safety and Efficacy of Herbal Medicines							
IV	Stability testing of herbal medicines. Application of various chromatographic	08						
	techniquesin standardization of herbal products. Preparation of documents for							
	new drug application and export registrationGMP requirements and Drugs &							
	Cosmetics Act provisions.							
V	Regulatory requirements for herbal medicines.							
	WHO guidelines on safety monitoring of herbal medicines in pharmacovigilance							

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systems. Comparison of various Herbal Pharmacopoeias.

Role of chemical and biological markers in standardization of herbal products

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, CarrierPub., 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 ofBotanicals. Business Horizons Publishers, New Delhi, India, 2002.
- 7. Shinde M.V., Dhalwal K., Potdar K., Mahadik K. Application of quality controlprinciples to herbal drugs. International Journal of Phytomedicine 1(2009); p. 4-8.
- 8. WHO. Quality Control Methods for Medicinal Plant Materials, World HealthOrganization, Geneva, 1998. WHO. Guidelines for the Appropriate Use of HerbalMedicines. WHO Regional Publications, Western Pacific Series No 3, WHORegional office for the Western Pacific, Manila, 1998.
- 9. WHO. The International Pharmacopeia, Vol. 2: Quality Specifications, 3rd edn.World Health Organization, Geneva, 1981.
- WHO. Quality Control Methods for Medicinal Plant Materials. World HealthOrganization, 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.



BP 807 ET. COMPUTER AIDED DRUG DESIGN (Theory)

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Teaching and Examination Scheme:

Teac	hing Scl	neme	Credits	Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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

- > Design and discovery of lead molecules
- > The role of drug design in drug discovery process
- > The concept of QSAR and docking
- ➤ Various strategies to develop new drug like molecules.
- > The design of new drug molecules using molecular modeling software

COURSE CONTENT

UNIT	CONTENT									
	Introduction to Dwg Diggsyawy and Dayslanmants Stages of dwg diggsyawy									
I	Introduction to Drug Discovery and Development: Stages of drug discovery	10								
	and development									
	Lead discovery and Analog Based Drug Design: 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 onclinical observation.									
	Analog Based Drug Design: Bioisosterism, Classification,									
	Bioisostericreplacement. Any three case studies									
II	Quantitative Structure Activity Relationship (QSAR): SAR versus QSAR,	10								
	History and development of QSAR, Types ofphysicochemical parameters,									
	experimental and theoretical approaches forthe determination of physicochemical									
	parameters such as Partitioncoefficient, Hammet's substituent constant and Tafts									
	steric constant. Hansch analysis, Free Wilson analysis, 3D-QSAR approaches									
	likeCOMFA and COMSIA.									
III	Molecular Modeling and virtual screening techniques:	10								

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	Virtual Screening techniques: Drug likeness screening, Concept										
	ofpharmacophore mapping and pharmacophore based Screening,										
	Molecular docking: Rigid docking, flexible docking, manual docking, Docking										
	based screening. De novo drug design.										
IV	Informatics & Methods in drug design: Introduction to Bioinformatics,										
	chemoinformatics. ADME databases, chemical, biochemical and pharmaceutical										
	databases.										
V	Molecular Modeling: Introduction to molecular mechanics and	07									
	quantummechanics.Energy Minimization methods and Conformational										
	Analysis, global conformational minima determination.										

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 OrganicMedicinal & Pharmaceutical Chemistry Lippincott, New York.
- 4. Foye WO Principles of Medicinal chemistry _ Lea & Febiger.
- 5. Koro lkovas A, Burckhalter JH. —Essentials of Medicinal Chemistry WileyInterscience.
- 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 UniversityPress.
- 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 (Elective subject)

Dean

H.P. Technical Univer

Hamirpur - 177001

Teaching and Examination Scheme:

Teac	Teaching Scheme Credits			Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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;

- > Summarize cell and molecular biology history.
- > Summarize cellular functioning and composition.
- > Describe the chemical foundations of cell biology.
- ➤ Summarize the DNA properties of cell biology.
- > Describe protein structure and function.
- > Describe cellular membrane structure and function.
- Describe basic molecular genetic mechanisms.
- > Summarize the Cell Cycle

COURSE CONTENT

UNIT	CONTENT	No. of Hrs.				
I	a) Cell and Molecular Biology: Definitions theory and basics and Applications.	10				
	b) Cell and Molecular Biology: History and Summation.					
	c) Properties of cells and cell membrane.					
	d) Prokaryotic versus Eukaryotic					
	e) Cellular Reproduction					
	f) Chemical Foundations – an Introduction and Reactions (Types)					
II	a) DNA and the Flow of Molecular Information	10				
	b) DNA Functioning					
	c) DNA and RNA					

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	d) Types of RNA							
	e) Transcription and Translation							
III	a) Proteins: Defined and Amino Acids	10						
	b) Protein Structure							
	c) Regularities in Protein Pathways							
	d) Cellular Processes							
	e) Positive Control and significance of Protein Synthesis							
IV	a) Science of Genetics	08						
	b) Transgenics and Genomic Analysis							
	c) Cell Cycle analysis							
	d) Mitosis and Meiosis							
	e) Cellular Activities and Checkpoints							
V	a) Cell Signals: Introduction	07						
	b) Receptors for Cell Signals							
	c) Signaling Pathways: Overview							
	d) Misregulation of Signaling Pathways							
	e) Protein-Kinases: Functioning							

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 Waverlycompany



- 12. B.R. Glick and J.J. Pasternak: Molecular Biotechnology: Principles and Applications of Recombinant DNA: ASM Press Washington D.C.
- 13. RA Goldshy et. al., : Kuby Immunology.



BP809ET. COSMETIC SCIENCE(Theory)

Teaching and Examination Scheme:

Teac	hing Sch	ng Scheme Credits		Marks			Duration of End Semester
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

Scope:This subject is designed to impart the basic knowledge of cosmetic sciences

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

- > Appreciate the applications of various commonly used cosmetic excipients.
- > Appreciate and demonstrate the various formulation methods used in cosmetic formulations
- > Appreciate and demonstrate the importance of analytical methods used in cosmetics.

COURSE CONTENT

UNIT	CONTENT							
I	Classification of cosmetic and cosmeceutical products; Definition of cosmetics as per							
	Indian and EU regulations, Evolution of cosmeceuticalsfrom cosmetics, cosmetics as							
	quasi and OTC drugs. Cosmetic excipients: Surfactants, rheologymodifiers,							
	humectants, 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.							
II	Principles of formulation and building blocks of skin care products:							
	Face wash, Moisturizing cream, Cold Cream, Vanishing cream and their advantages							
	anddisadvantages. Application of these products in formulation of							
	cosmecuticals. Antiperspants & deodorants- Actives & mechanism of action.							
	Principles of formulation and building blocks of Hair care products:							
	Conditioning shampoo, Hair conditioner, anti-dandruff 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.							
III	Sun protection, Classification of Sunscreens and SPF.	10						

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	Role of herbs in cosmetics: Skin Care: Aloe and turmeric; Hair care: Henna and amla								
	Oral care: Neem and clove. Analytical cosmetics: BIS specification and analytical								
	methods for shampoo, skincreamand toothpaste.								
IV	Principles of Cosmetic Evaluation:Principles of sebumeter, corneometer.								
	Measurementof TEWL, Skin Color, Hair tensile strength, Hair combing								
	propertiesSoaps,and syndet bars. Evolution and skin benfits.								
V	Oily and dry skin, causes leading to dry skin, skin moisturisation. Basic understanding								
	ofthe terms Comedogenic, dermatitis.Cosmetic problems associated with Hair and								
	scalp: Dandruff, Hair fall causesCosmetic problems associated with skin: blemishes,								
	wrinkles, acne, prickly heat andbody odor.								
	Antiperspirants and Deodorants- Actives and mechanism of action								

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.



BP810 ET. PHARMACOLOGICAL SCREENINGMETHODS

Teaching and Examination Scheme:

Teac	Teaching Scheme		Credits	Marks		Duration of End Semester	
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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,

- Appreciate the applications of various commonly used laboratory animals.
- Appreciate and demonstrate the various screening methods used in preclinical Research
- ➤ Appreciate and demonstrate the importance of biostatistics and researchmethodology
- ➤ Design and execute a research hypothesis independently

COURSE CONTENT

UNIT	CONTENT						
		Hrs.					
Ι	Laboratory Animals: Study of CPCSEA and OECD guidelines for	08					
	maintenance, breedingand conduct of experiments on laboratory animals,						
	Common labanimals: Description and applications of different species and						
	strainsof animals. Popular transgenic and mutant animals. Techniques for						
	collection of blood and common routes of drug						
	administration in laboratory animals, Techniques of blood collectionand						
	euthanasia.						
II	Preclinical screening models:						
	a. Introduction: Dose selection, calculation and conversions,preparation of drug						
	solution/suspensions, grouping of animals andimportance 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						
	Preclinical screening models: for CNS activity- analgesic,antipyretic,anti-						

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	inflammatory, general anaesthetics, sedative andhypnotics, antipsychotic,							
	antidepressant, antiepileptic, antiparkinsonism, alzheimer's disease							
III	Preclinical screening models: for ANS activity,	10						
	sympathomimetics, sympatholytics, parasympathomimetics, parasympatholytics,							
	skeletalmuscle relaxants, drugs acting on eye, local anaethetics							
IV	Preclinical screening models: for CVS activity- antihypertensives, diuretics,							
	antiarrhythmic, antidyslepidemic, anti aggregatory, coagulants, and							
	anticoagulantsPreclinical screening models for other important drugs like							
	antiulcer, antidiabetic, anticancer and antiasthmatics.							
V	Research methodology and Bio-statistics							
	Selection of research topic, review of literature, research hypothesisand study							
	designPre-clinical data analysis and interpretation using Students _t' testand One-							
	way ANOVA. Graphical representation of data							

Recommended Books (latest edition):

- 1. Fundamentals of experimental Pharmacology-byM.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 JRichard



BP 811 ET. ADVANCED INSTRUMENTATION TECHNIQUES

Teaching and Examination Scheme:

Teaching Scheme Credits		Marks		Duration of End Semester			
L	T	P	C	Sessional	End Semester	Total	Examination
					Exam		
3	1	0	4	25	75	100	3 hours

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

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

- > understand the advanced instruments used and its applications in drug analysis
- understand the chromatographic separation and analysis of drugs.
- > understand the calibration of various analytical instruments
- ➤ know analysis of drugs using various analytical instruments.

COURSE CONTENT

UNIT	CONTENT	No. of Hrs.						
I	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							
	Mass Spectrometry- Principles, Fragmentation, Ionization techniques –Electron							
	impact, chemical ionization, MALDI, FAB, Analyzers-Time offlight and							
	Quadrupole, instrumentation, applications							
II	Thermal Methods of Analysis: Principles, instrumentation and applications	10						
	of ThermogravimetricAnalysis (TGA), Differential Thermal Analysis (DTA),							
	Differential Scanning Calorimetry (DSC)							
	X-Ray Diffraction Methods: Origin of X-rays, basic aspects of crystals, Xray							
	Crystallography, rotating crystal technique, single crystal diffraction,powder							

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	diffraction, structural elucidation and applications.							
III	Calibration and validation-as per ICH and USFDA guidelines							
	Calibration of following Instruments							
	Electronic balance, UV-Visible spectrophotometer, IR							
	spectrophotometer, Fluorimeter, Flame Photometer, HPLC and GC							
IV	Radio immune assay:Importance, various components, Principle, different							
	methods, Limitation and Applications of Radio immuno assay							
	Extraction techniques:General principle and procedure involved in the solid							
	phase extraction and liquid-liquid extraction							
V	Hyphenated techniques-LC-MS/MS, GC-MS/MS, HPTLC-MS.	07						

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



BP 812 ET. DIETARY SUPPLEMENTS AND NUTRACEUTICALS

No. of hours: 3 Tutorial: 1 Credit point: 4

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.

UNIT I 07 hours

- 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 benefits of following used as nutraceuticals/functional foods: Spirulina, Soyabean, Ginseng, Garlic, Broccoli, Gingko, Flaxseeds

UNIT II 15 hours

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.



UNIT III 07 hours

- 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..

UNIT IV 10 hours

- 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

UNIT V 06 hours

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

edition. Lea and Febiger

