

PROGRAMME STRUCTURE

School of Pharmacy B.Pharm

Programme Code: SOP0101

Batch: 2023-2027





School of Pharmacy Programme Structure

S. No	Course Code	Course Name	Category	L	T	P	Credits
Sem	ester I				1		
1	BP101T	Human Anatomy and Physiology I— Theory	Core	3	1	-	4
2	BP102T	Pharmaceutical Analysis I—Theory	Core	3	1	-	4
3	BP103T	Pharmaceutics I— Theory	Core	3	1	-	4
4	BP104T	Pharmaceutical Inorganic Chemistry— Theory	Core	3	1	-	4
5	BP105T	Communication skills– Theory*	Skill Enhance ment course	2	-	-	2
6	BP106RBT BP106RMT	Remedial Biology/ Remedial Mathematics—Theory*	Core	2	-	-	2
7	BP107P	Human Anatomy and Physiology— Practical	Practical	-	-	4	2
8	BP108P	Pharmaceutical Analysis I–Practical	Practical	-	-	4	2
9	BP109P	Pharmaceutics I— Practical	Practical	-	-	4	2
10	BP110P	Pharmaceutical Inorganic Chemistry— Practical	Practical	-	-	4	2
11	BP111P	Communication skills–Practical*	Practical	-		2	1
12	BP112RBP	Remedial Biology– Practical*	Practical	-	-	2	1
		Total Credits:	: 27				





S.	Course Code	Course Name	Category	L	T	P	Credits
No Seme	ester II						
1	BP201T	Human Anatomy and Physiology II – Theory	Core	3	1	-	4
2	BP202T	Pharmaceutical Organic Chemistry I -Theory	Core	3	1	-	4
3	BP203T	Biochemistry– Theory	Core	3	1	-	4
4	BP204T	Pathophysiology— Theory	Core	3	1	-	4
5	BP205T	Computer Applications in Pharmacy—Theory*	Core	3	-	-	3
6	BP206T	Environmental sciences –Theory*	Skill Enhance ment course	3	-	-	3
7	BP207P	Human Anatomy and Physiology II – Practical	Core	-	_	4	2
8	BP208P	Pharmaceutical Organic Chemistry I–Practical	Core	-	-	4	2
9	BP209P	Biochemistry– Practical	Core	-	-	4	2
10	BP210P	Computer Applications in Pharmacy–Practical*	Ability Enhance ment course	-	-	2	1
		Total Cred	its: 29				





S.	Course Code	Course Name	Category	L	T	P	Credits
No							
Seme 1	ester-III	Dis aures a queti a al	Ī	l	1		
1	BP301T	Pharmaceutical Organic Chemistry II – Theory	Core	3	1	-	4
2	BP302T	Physical Pharmaceutics I— Theory	Core	3	1	-	4
3	BP303T	Pharmaceutical Microbiology– Theory	narmaceutical licrobiology- Core			-	4
4	BP304T	Pharmaceutical Engineering—Theory	Core	3	1	-	4
5.	KV301	Universal Human value & Professional Ethics	Ability enhancem ent course	3	0	0	3
5	BP305P	Pharmaceutical Organic Chemistry II– Practical	Practical	-	-	4	2
6	BP306P	Physical Pharmaceutics I— Practical	Practical	-	-	4	2
7	BP307P	Pharmaceutical Microbiology– Practical	Practical	-	-	4	2
8	BP308P	Pharmaceutical Engineering— Practical	Core	-		4	2
		Total Cred	its: 27				





S. No	Course Code	Course Name	Category	L	T	P	Credits
Sen	nester IV			•			
1	BP401T	Pharmaceutical Organic Chemistry III– Theory	Core	3	1	-	4
2	BP402T	Medicinal Chemistry I–Theory	Core	3	1	-	4
3	BP403T	Physical Pharmaceutics II— Theory	Core	3	1	-	4
4	BP404T	Pharmacology I— Theory	Core	3	1	-	4
5	BP405T	Pharmacognosy and PhytochemistryI —Theory	Core	3	1	-	4
6	BP406P	Medicinal Chemistry I– Practical	Practical	-	-	4	2
7	BP407P	Physical Pharmaceutics II— Practical	Practical	-	-	4	2
8	BP408P	Pharmacology I— Practical	Practical	-	-	4	2
9	BP409P	Pharmacognosy and Phytochemistry I —Practical	Practical	-	-	4	2
		Total Credi	its: 28				





S. No	Course Code	Course Name	Category	L	T	P	Credits		
	ester-V								
1	BP501T	Medicinal Chemistry II–Theory	Core	3	1	-	4		
2	BP502T	Industrial Pharmacy I—Theory	Core	3	1	-	4		
3	BP503T	Pharmacology II— Theory	Core	3	1	-	4		
4	BP504T	Pharmacognosy and Phytochemistry II—Theory	Core	3	1	-	4		
5	BP505T	Pharmaceutical Jurisprudence– Theory	Core	3	1	-	4		
6	BP506P	Industrial Pharmacy I— Practical	Practical	-	-	4	2		
7	BP507P	Pharmacology II— Practical	Practical	-	-	4	2		
8	BP508P	Pharmacognosy and Phytochemistry II– Practical	Practical	-	-	4	2		
9	PCC301 Community connect P1		Practical	-	-	4	2		
	Total Credits: 28								





S. No	Course Code	Course Name	Category	L	Т	P	Credits		
Semester	·-VI								
1	BP601T	Medicinal Chemistry III–Theory	Core	3	1	-	4		
2	BP602T	Pharmacology III— Theory	Core	3	1	-	4		
3	BP603T	Herbal Drug Technology—Theory	Core	3	1	-	4		
4	BP604T	Biopharmaceutics and Pharmacokinetics— Theory	s and		1	-	4		
5	BP605T	Pharmaceutical Biotechnology— Theory	Core	3	1	-	4		
6	BP606T	Quality Assurance– Theory	Core	3	1	-	4		
7	BP607P	Medicinal chemistry III–Practical	Practical	-	-	4	2		
8	BP608P	Pharmacology III— Practical	Practical			4	2		
9	BP609P	Herbal Drug Technology– Practical	Practical	-	-	4	2		
Total Credits: 30									





S.	No Course Code	Course Name	Category	L	T	P	Credits
Se	mester-VII						
1	BP701T	Instrumental Methods of Analysis—Theory	Core	3	1	-	4
2	BP702T	Industrial Pharmacy II—Theory	Core	3	1	-	4
3	BP703T	Pharmacy Practice— Theory	Core	3	1	-	4
4	BP704T	BP704T Novel Drug Delivery System–Theory		3	1	-	4
5	BP705P	Instrumental Methods of Analysis–Practical	Practical	-	-	4	2
6	BP706PS	Practice School*	Project	-	-	12	6
		Total Credits	: 24				





S. No	Course Code	Course Name	Category	L	T	P	Credits
	ester-VIII						
1	BP801T	Biostatistics and Research Methodology	Core	3	1		4
2	BP802T	Social and Preventive Pharmacy	Core	3	1	-	4
3	BP803ET	Pharma Marketing Management	Discipline specific Elective				
4	BP804ET	Pharmaceutical Regulatory Science	Elective Discipline	3+	1+1		
5	BP805ET	Pharmacovigilance					
6	BP806ET	Quality Control and Standardization Of Herbals					
7	BP807ET	Computer Aided Drug Design	Elective				4+4=8
8	BP808ET	Cell and Molecular Biology	Discipline specific Elective				
9	BP809ET	Cosmetic Science	Elective				
10	BP810ET	Experimental Pharmacology	Elective				
11	BP811ET	Advanced Instrumentation Techniques	Elective				
12	BP812ET	Dietary Supplements and Nutraceuticals	Elective				
13	BP813PW	Project Work	Project -	_	_	12	6
		Total Credits:		•			
		Programme Credits: 21	5				



Course Module





treatment of different diseases 2. Demonstrate isolation of different organs/tissues from the laborate animals by simulated experiments 3. Demonstrate the various receptor actions using isolated tiss preparation 4. Appreciate correlation of pharmacology with related medical science CO1: The students will be able to understand the structure and function of various tissues and organs of the body. CO2: The student will be able to summarize the functioning of various systems and their homeostasis. CO3: The student will be able to apply the knowledge of the anatomy aphysiology of different body parts in explaining the working patterns different body systems. CO4: The students will analyze the structures of various tissues and the origin to evaluate their damage and repair process. CO5: The students will evaluate the mechanisms of various processes which the functioning of the various body organs depend and will obsert the anatomical differentiation of different body parts. CO6: The students will be able to conclude about the mechanisms various functioning of the body organs. This subject is designed to impart fundamental knowledge on the structure and functions of the various systems of the human body. It also helps understanding both homeostatic mechanisms. The subject provides	Sch	ool:	SOP					
Course Code BP101 T	Pro	gramme:	B. Pharm					
Credits 4	Bra	nch:	Semester: 1					
3 Credits 4 4 Contact Hours (L-T-P) Course Type Compulsory 5 Course Objective 1. Understand the mechanism of drug action and its relevance in treatment of different diseases 2. Demonstrate isolation of different organs/tissues from the laborate animals by simulated experiments 3. Demonstrate the various receptor actions using isolated tiss preparation 4. Appreciate correlation of pharmacology with related medical science 6 Course Outcomes CO1: The students will be able to understand the structure and function of various tissues and organs of the body. CO2: The student will be able to summarize the functioning of various tissues and their homeostasis. CO3: The student will be able to apply the knowledge of the anatomy aphysiology of different body parts in explaining the working patterns different body systems. CO4: The students will analyze the structures of various tissues and the origin to evaluate their damage and repair process. CO5: The students will evaluate the mechanisms of various processes which the functioning of the various body organs depend and will obset the anatomical differentiation of different body parts. CO6: The students will be able to conclude about the mechanisms various functioning of the body organs. 7 Course Description This subject is designed to impart fundamental knowledge on the struct and functions of the various systems of the human body. It also helps understanding both homeostatic mechanisms. The subject provides basic knowledge required to understand the various disciplines pharmacy.	1	Course Code	BP101 T					
Contact Hours (L-T-P)	2	Course Title	Human Anatomy & Physiology I – Theory					
Course Type Compulsory								
Course Objective	4		3-1-0					
Objective 1. Understand the mechanism of drug action and its relevance in treatment of different diseases 2. Demonstrate isolation of different organs/tissues from the laborate animals by simulated experiments 3. Demonstrate the various receptor actions using isolated tiss preparation 4. Appreciate correlation of pharmacology with related medical science Outcomes CO1: The students will be able to understand the structure and function of various tissues and organs of the body. CO2: The student will be able to summarize the functioning of various yestems and their homeostasis. CO3: The student will be able to apply the knowledge of the anatomy aphysiology of different body parts in explaining the working patterns different body systems. CO4: The students will analyze the structures of various tissues and the origin to evaluate their damage and repair process. CO5: The students will evaluate the mechanisms of various processes which the functioning of the various body organs depend and will observe the anatomical differentiation of different body parts. CO6: The students will be able to conclude about the mechanisms various functioning of the body organs. This subject is designed to impart fundamental knowledge on the structure and functions of the various systems of the human body. It also helps understanding both homeostatic mechanisms. The subject provides basic knowledge required to understand the various disciplines pharmacy.		Course Type	Compulsory					
Course Outcomes CO1: The students will be able to understand the structure and function of various tissues and organs of the body. CO2: The student will be able to summarize the functioning of various to systems and their homeostasis. CO3: The student will be able to apply the knowledge of the anatomy aphysiology of different body parts in explaining the working patterns different body systems. CO4: The students will analyze the structures of various tissues and the origin to evaluate their damage and repair process. CO5: The students will evaluate the mechanisms of various processes which the functioning of the various body organs depend and will obsert the anatomical differentiation of different body parts. CO6: The students will be able to conclude about the mechanisms various functioning of the body organs. This subject is designed to impart fundamental knowledge on the structure and functions of the various systems of the human body. It also helps understanding both homeostatic mechanisms. The subject provides basic knowledge required to understand the various disciplines pharmacy.	5		 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 by simulated experiments Demonstrate the various receptor actions using isolated tissue preparation Appreciate correlation of pharmacology with related medical sciences 					
Description and functions of the various systems of the human body. It also helps understanding both homeostatic mechanisms. The subject provides basic knowledge required to understand the various disciplines pharmacy. 8 Outline syllabus CO Mapping	6		CO2: The student will be able to summarize the functioning of various body systems and their homeostasis. CO3: The student will be able to apply the knowledge of the anatomy and physiology of different body parts in explaining the working patterns of different body systems. CO4: The students will analyze the structures of various tissues and their origin to evaluate their damage and repair process. CO5: The students will evaluate the mechanisms of various processes on which the functioning of the various body organs depend and will observe the anatomical differentiation of different body parts.					
	7		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					
	8	Outline syllabus		CO Mapping				





	UNIT-I	
	A.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.	
	B.Cellular level of organization	
	Structure and functions of cell, transport across cell membrane, cell division, cell junctions. General principles ofcell communication, intracellular signaling pathway activation by extracellular signal molecule, Forms of intracellular signaling: a) Contact-dependent b) Paracrine c) Synaptic d) Endocrine C.Tissue level of organization	
	Classification of tissues, structure, location and functions of epithelial, muscular and nervous and connective tissues.	
2	UNIT-II A. Integumentary system Structure and functions of skin	
	B. 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 C. Joints	CO2
	Structural and functional classification, types of	
2	joints movements and its Articulation	
3	UNIT-III A. Body fluids and blood	
	Body fluids, composition and functions of blood, hemopoeisis, formation of hemoglobin, anemia, mechanisms of coagulation, blood grouping, Rh factors, transfusion, its significance and disorders of blood, Reticulo endothelial system.	CO3 CO6
	B. Lymphatic system Lymphatic organs and tissues, lymphatic vessels, lymph circulation and C. Functions of lymphatic system	





	4	UNIT-IV		1000000000	c.in				
	4	A. Periphera	l norvoue eve	tom:					
		_	-						
				I nervous system:					
				sympathetic and					
			etic nervous sy		CO4				
				pinal and cranial nerves.					
		C .Special ser	ises						
				eye, ear, nose and tongue					
		and their disor	rders.						
	5	UNIT-V							
		A. Cardiovas	A. Cardiovascular system						
		Heart – anat	Heart – anatomy of heart, blood circulation, blood						
		vessels, struc	ture and funct	cions of					
		artery, vein a	and capillaries	s, elements of conduction	CO5				
		system of hea	art and heart b	eat, its					
		B. Regulation	n by autonomi	c nervous system, cardiac					
		output, car	rdiac cycle.						
		C. Regulation	n ofblood	pressure, pulse,					
		_		isorders of heart.					
	Mode of	Theory	8						
	examination	Theory							
	Weightage	Continuous	Sessional	ESE					
	Distribution	Mode	Exam						
		Assessment							
		10 Marks	15	75					
	Text book/s*	10 Marks	13	13					
	Text book/s								
		1. Essentials		al Physiology by K.					
1		Sembuling	am and P.	Sembulingam. Jaypee					
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	Other References	Sembuling brother's n 2. Anatomy a by Kathl Livingston 3. Physiologi Tailor. W USA 4. Text boo C,Guyton U.S.A. 5. Principles	am and P. nedical publish and Physiolog een J.W. e, New York cal basis of M illiams & W k of Medic andJohn.E.	Sembulingam. Jaypee hers, New Delhi. sy in Health and Illness Wilson, Churchill Medical Practice-Best and Vilkins Co,Riverview,MI real Physiology- Arthur Hall. Miamisburg, OH, and Physiology by Tortora					



Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	1	2	2	1	3	1	_	1	2	3
CO2	3	2	2	2	1	3	1	-	1	2	3
CO3	3	2	1	2	2	3	1	-	1	3	3
CO4	3	2	1	2	2	-	2	-	-	2	3
CO5	3	2	2	3	2	3	2	-	1	3	3
CO6	3	1	2	-	2	-	-	-	-	3	3

- 1-Slight (Low)
- 2-Moderate (Medium)
- 3-Substantial (High)





Programme: Branch: Semester: 1	Sch	ool:	SOP					
Branch: Semester: 1								
2 Course Title Pharmaceutical Analysis I- Theory 3 Credits 4 4 Contact Hours (L-T-P) Course Type Compulsory 5 Course Objective Upon completion of this course the student should be able to 1. Understand the mechanism of drug action and its relevance in treatment of different diseases 2. Demonstrate isolation of different organs/tissues from the laborate animals by simulated experiments 3. Demonstrate the various receptor actions using isolated tiss preparation 4. Appreciate correlation of pharmacology with related medical sciences 6 Course Outcomes CO1: Students will be able to understand about the Pharmaceutical analy and its importance in Pharmacy. CO2: Students will able to evaluate about different types of analytical techniques. CO3: Students will able to explain difference between volumet quantitative and qualitative analysis. CO5: Students will be able to conclude about Electrochemical methods analysis CO6: Students will be able to generalize about Non aqueous titrations. 7 Course Description This course deals with the fundamentals of analytical chemistry a principles of electrochemical analysis of drug								
3	1	Course Code	BP102T					
Contact Hours (L-T-P)								
Hours (L-T-P) Course Type Course Objective Upon completion of this course the student should be able to 1. Understand the mechanism of drug action and its relevance in treatment of different diseases 2. Demonstrate isolation of different organs/tissues from the laborate animals by simulated experiments 3. Demonstrate the various receptor actions using isolated tiss preparation 4. Appreciate correlation of pharmacology with related medical sciences Course Outcomes CO1: Students will be able to understand about the Pharmaceutical analy and its importance in Pharmacy. CO2: Students will able to evaluate about different types of analyt techniques. CO3: Students can apply their anlaytical knowledge regarding Gravimetr CO4: Students will able to explain difference between volumet quantitative and qualitative analysis. CO5: Students will be able to conclude about Electrochemical methods analysis CO6: Students will be able to generalize about Non aqueous titrations. Course Description This course deals with the fundamentals of analytical chemistry a principles of electrochemical analysis of drug		Credits						
Course Type	4	Hours	3-1-0					
Course Objective			Compulsory					
Outcomes CO1: Students will be able to understand about the Pharmaceutical analyand its importance in Pharmacy. CO2: Students will able to evaluate about different types of analytic techniques. CO3: Students can apply their anlaytical knowledge regarding Gravimetr CO4: Students will able to explain difference between volumet quantitative and qualitative analysis. CO5: Students will be able to conclude about Electrochemical methods analysis CO6: Students will be able to generalize about Non aqueous titrations. This course deals with the fundamentals of analytical chemistry a principles of electrochemical analysis of drug	5	Course	Upon completion of this course the student should be able to 1. Understand the mechanism of drug action and its relevance in the reatment of different diseases 2. Demonstrate isolation of different organs/tissues from the laboratory animals by simulated experiments 3. Demonstrate the various receptor actions using isolated tissue preparation					
Description principles of electrochemical analysis of drug	6		and its importance in Pharmacy. CO2: Students will able to evaluate about different techniques. CO3: Students can apply their anlaytical knowledge reg CO4: Students will able to explain difference to quantitative and qualitative analysis. CO5: Students will be able to conclude about Electrod analysis CO6: Students will be able to generalize about No	types of analytical garding Gravimetry. Detween volumetric, chemical methods of				
	7			tical chemistry and				
8 Outline syllabus CO Mapping	8	Outline syllabu	1 1S	CO Mapping				





 	W B THOUGHT WE	
1	UNIT-I A. Pharmaceutical Analysis Definition and scope Definition and scope 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, C. Methods of minimizing errors, accuracy, precision and significant figures.	CO1
2	 UNIT-II A. Acid base titration & Non aqueous titration Theories of acid base indicators, classification of acid base titrations and theory involved in titrations of strong, weak, and very weak acids and bases, neutralization curves B. Non aqueous titration: Solvents, C. acidimetry and alkalimetry titration and estimation of Sodium benzoate and Ephedrine HCl 	CO2 CO6
3	 UNIT-III A. Precipitation titrations, Complexometric tration & Gravimetry, Diazotization Precipitation titrations: Mohr's method, Volhard's, Modified Volhard's, Fajans method, estimation of sodium chloride. Complexometric titration: Classification, metal ion indicators, masking and demasking reagents, estimation of Magnesium sulphate, and calcium gluconate. B. Gravimetry: Principle and steps involved in gravimetric analysis. Purity of the precipitate: co-precipitation and post precipitation, Estimation of barium sulphate. C. Define the Diazotization with their principle, methodology and their uses 	CO3





4	UNIT-IV		www.sharda	.ac.iii		
	A. Redox titrat					
	Concepts of oxida	CO4				
	B. Types of red Cerimetry C. Iodimetry,		Principles and applications) comatometry, Dichrometry,	CO4		
		potassium ioda				
5	UNIT-V	1				
	A. Electrochen Conductometry- Conductometric to B. Potentiometry working of reference electrode and cal (metal electrode determine end applications. C.Polarography and working of platinum substitution and uses of Ethano electrode, applica	CO5				
Mode of	Theory	tions				
examinatio n						
Weightage	Continuous	Sessional	ESE			
Distributio	Mode	Exam				
n	Assessment					
E	10 Marks	15	75			
Text		kett & J.B. Ste	,			
book/s*		Pharmaceutica	•			
			ss of University			
	of Londor					
	2. A.I. Voge analysis					
	Chemistry 4. Bentley and Driver's Textbook of Pharmaceutical Chemistry					
	5. John H. K	ennedy, Analy	ytical chemistry principles			
		armacopoeia.				
	l .					



Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	2	3	2	-	2	2	2	1	2	3
CO2	3	2	3	2	-	2	2	2	2	1	3
CO3	3	2	2	1	-	3	2	2	-	2	2
CO4	3		2.	2		3	2	2	2	2	3
CO4	3	_	2	2	_	3	2	2	2	2	3
CO5	3	2	2	3	-	2	2	3	2	2	3
COL	2	2		2		2	2	2	2	2	2
CO6	3	3	2	3	-	2	2	2	2	2	3

- 1-Slight (Low) 2-Moderate (Medium)
- 3-Substantial (High)





Branch: Semester: 1 Semester: 1 Semester: 1 Course Code BP103T Credits 4 Contact Hours (L-T-P) Course Type Compulsory Upon completion of this course the student should be able to 1. Understand the mechanism of drug action and its relevance in the treatment of different diseases 2. Demonstrate isolation of different organs/tissues from the laboratory animals by simulated experiments 3. Demonstrate the various receptor actions using isolated tissue preparation 4. Appreciate correlation of pharmacology with related medical sciences CO1: The students will be able to understand about general formulation, classification of different dosage forms and various Pharmacopoeias-IP, BP, USP etc. CO2: The students will be able to apply the professional way of handling the prescription, excipients used in different dosage forms, various factors affecting Posology and solubility enhancement techniques. CO3: The students will be able to distinguish between various Monophasic and biphasic liquids. CO5: Students will be able to explain about different types of semisolid dosage forms like suspension, emulsion, ointments, pastes, creams etc. CO6: The students will be able to predict stability problems in different dosage forms.	School:		SOP					
1 Course Code 2 Course Title Communication skills – Theory 3 Credits 4 4 Contact Hours (L-T-P) 5 Course Type Compulsory 6 Dijective Objective Obje	Pro	gramme:	B.Pharm					
Course Title	Bra	nch:						
Contact Hours (L-T-P)	1	Course Code	BP103T					
4 Contact Hours (L-T-P) Course Type Course Type Course Objective 5 Course Objective 1. Understand the mechanism of drug action and its relevance in the treatment of different diseases 2. Demonstrate isolation of different organs/tissues from the laboratory animals by simulated experiments 3. Demonstrate the various receptor actions using isolated tissue preparation 4. Appreciate correlation of pharmacology with related medical sciences CO1: The students will be able to understand about general formulation, classification of different dosage forms and various Pharmacopoeias-IP, BP, USP etc. CO2: The student will be able to apply the professional way of handling the prescription, excipients used in different dosage forms, various factors affecting Posology and solubility enhancement techniques. CO3: The students will be able to illustrate different methods of preparation of various semisolid dosage forms and how to calculate the dose of pediatric patients, different calculations based on the Imperial & Metric system. CO4: The students will be able to distinguish between various Monophasic and biphasic liquids. CO5: Students will be able to explain about different types of semisolid dosage forms like suspension, emulsion, ointments, pastes, creams etc. CO6: The students will be able to predict stability problems in different dosage forms. This course is designed to impart a fundamental knowledge on the preparatory pharmacy with arts and science of preparing the different conventional dosage forms.								
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2	UNIT-II A. Pharmaceutical Calculations, Powders, Liquid dosage forms Weight & Measures, Calculation involving Percentage solution etc. B. Definition of Powders, Eutectic Mixtures, Geometric Dilutions. C.Solubility enhancement techniques Advantages & disadvantage of liquid dosage forms.	





	3	UNIT-III			
		A. Monopha	sic and Bipha	sic liquids	
		gargles, syrup B. suspension overcome C. Emulsions	s, liniments, E Suspensions, a & stability s, classification	-	f
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		С.	and classification of ral incompatibilities at types, methods of base, Evaluation and	C04	
	5	CO5			
	Mode of examination	Theory			
	Weightage Distribution	Continuous Mode Assessment	Sessional Exam	ESE	
		10 Marks	15	75	





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





Pos Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1	3	2	2	2	2	2	2	-	1	1	3
CO2	3	1	2	2	1	2	2	-	2	1	3
CO3	3	1	2	1	2	3	1	-	2	2	2
CO4	3	2	1	2	1	3	2	-	1	2	3
CO5	3	3	2	3	2	2	2	-	1	1	3
CO6	3	3	2	3	1	2	2	-	1	1	3

1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sc	hool:	SOP						
Pr	ogramme:	B.Pharm						
Br	anch:	Semester: 1						
1	Course	BP104T						
	Code							
2	Course	Pharmaceutical Inorganic Chemistry -Theory						
	Title							
3	Credits	4						
4	Contact	3-1-0						
	Hours							
	(L-T-P)							
	Course	Compulsory						
	Type							
5	Course	Upon completion of this course the student should be able to						
	Objective	1. Understand the mechanism of drug action and its relevance in the treatment						
		of different diseases						
		2. Demonstrate isolation of different organs/tissues from the laboratory						
		animals by simulated experiments						
		3. Demonstrate the various receptor actions using isolated tissue preparation						
	C	4. Appreciate correlation of pharmacology with related medical sciences						
6	Course Outcomes	CO1 Students shall be able to illustrate various sources of impurities and their control in inorganic drugs and pharmaceuticals.						
	Outcomes	CO2 Students shall be able to explain concept of acids, bases and buffers						
		andmethods of calculating and adjusting isotonicity.						
		CO3 Students shall be able to discuss major intra and extracellular ions,						
		replacement therapy and physiological acid-base balance.						
		CO4 Students shall be able to evaluate various inorganic compounds, like						
		gastrointestinal agents, dentalproducts and antimicrobials.						
		CO5 Students will be able to apply knowledge about radiopharmaceutical						
		their handling, hazards and uses.						
	CO6 Students shall be able to understand importance of inorganic compou							
		which can be used as useful medicinal compounds.						
7	Course							
	Description	This subject deals with the monographs of inorganic drugs and						
		pharmaceuticals.						
8	Outline sylla	bus CO Mapping						





1	UNIT-I A. Impurities in pharmaceutical substances History of Pharmacopoeia, B.Sources and types of impurities, principle involved in the limit test for Chloride, Sulphate, Iron, Arsenic, Lead C. Heavy metals, modified limit test for Chloride and	CO1
2	UNIT-II A. Acids, Bases and Buffers Major extra and intracellular electrolytes, Dental products	
	B. Buffer equations and buffer capacity in general, buffers in pharmaceutical systems, preparation, stability, buffered isotonic solutions, measurements of tonicity, calculations and methods of adjusting tonicity.	CO2
	C. Functions of major physiological ions, Electrolytes used in the replacement therapy: Sodium chloride*, Potassium chloride, Calcium gluconate* and Oral Rehydration Salt Dentifrices, role of fluoride in the treatment of dental caries, Desensitizing agents, Calcium carbonate, Sodium fluoride, and Zinc eugenol cement.	
3	UNIT-III A.Gastrointestinal agents Acidifiers: Ammonium chloride* and Dil. HCl Antacid: Ideal properties of antacids, ombinations of antacids, Sodium 40 Bicarbonate*, Aluminum hydroxide gel, B.Magnesium hydroxide mixture Cathartics: Magnesium sulphate, Sodium orthophosphate, Kaolin and Bentonite Antimicrobials: C. Mechanism, classification, Potassium permanganate, Boric acid, Hydrogen peroxide*, Chlorinated lime*, Iodine and its preparation	CO3, CO6





		a.ac.in					
4	UNIT-IV A.Miscellaneous con Expectorants: Potassi Emetics: Copper sulp B.Sodium potassium Haematinics: Ferrous C. Ferrous glucona thiosulphate*, Acti Astringents: Zinc Sul	CO4					
5	UNIT-V A. Radiopharmaceu Radio activity, Measu B.Properties of α, β, α and C. study of radio isot conditions, precautio radioactive substance	CO5					
Mode of examinatio	Theory	Theory					
Weightage Distributio	Continuous Mode Assessment	Sessional Exam	75 ESE				
Text book/s*	Practical Chemis Press of 4th edition 2. A.I. Vo Inorgan 3. P. Gund Chemis 4. M.L Schemis 5. Bentley Pharma	gel, Text Book of (ic analysis lu Rao, Inorganic P try, 3 rd Edition hroff, Inorganic Ph					
		& Cnatwal, Inorgan ceutical Chemistry	IIC .				





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	1	3	3	ı	-	3	1	3	1	3
CO2	3	3	2	2	ı	ı	2	3	2	3	3
CO3	3	2	2	3	-	-	2	3	3	2	3
CO4	3	2	2	2	1	-	2	3	2	2	3
CO5	3	2	3	2	1	-	2	2	2	2	3
CO6	3	2	2	2	-	-	2	3	2	2	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





School:		SOP						
Prog	gramme:	B.Pharm						
Bra	nch:	Semester: 1						
1	Course Code	BP105T						
2	Course Title	Communications Theory Theory						
3	Credits	2						
4	Contact Hours (L-T-P)	2-0-0						
	Course Type	Compulsory						
5	Course Objective	Upon completion of the course the student shall be able	e to					
	3	Understand the behavioral needs for a Pharmacist to fur effectively in theareas of pharmaceutical operation	nction					
		Communicate effectively (Verbal and Non Verbal)						
		Effectively manage the team as a team player						
		Develop interview skills						
		Develop Leadership qualities and essentials						
6	Course Outcomes	CO1: students will be able to Understand the behaviora for a Pharmacist to function effectively in theareas of pharmaceutical operation	l needs					
		CO2: students will be able to plan how to Communicate and Non Verbal)	e effectively (Verbal					
		CO3: students will be able to apply Effectively to manage team player	ge the team as a					
		CO4: students will be able to Develop interview skills						
		CO5: students will be able to generalize about Leadership qualities and essentials CO6: students will be able to plan about Group discussion						
7	Course Description	This course will brepare the young bharmacy student to interact						
8	Outline syllabu	is	CO Mapping					





1	UNIT-I
	A. Communication Skills: Introduction, Definition, The Importance of Communication, The Communication Process – Source, Message, Encoding, Channel, Decoding, Receiver, Feedback, Context
	B. Barriers to communication: Physiological Barriers, Physical Barriers, Cultural Barriers, Language Barriers, Gender Barriers, Interpersonal Barriers, Psychological Barriers, Emotional barriers
	C. Perspectives in Communication: Introduction, Visual Perception, Language, Other factors affecting our perspective - Past Experiences, Prejudices, Feelings, Environment
2	UNIT-II
	A. Elements of Communication: Introduction, Face to Face Communication - Tone of Voice, Body Language (Non-verbal communication), Verbal Communication, Physical Communication
	B. Communication Styles: Introduction, The Communication Styles Matrix with example for each –
	C. Direct Communication Style, Spirited Communication Style, Systematic Communication Style, Considerate Communication Style





			THE THE PART OF TH	illection .			
3	UNIT-III						
	A. Self-A Becon Diffice						
	Introd Writte Topic,	Effective Water with the Communication of Meaning,	CO3				
	Put th	Writing Effe he Main Po nce, Organiza					
4	UNIT-IV						
	A. intervi						
	B. Fears,	CO4					
	Delive	C. Structuring YourPresentation, Delivering Your Presentation, Techniques of Delivery					
5	UNIT-V						
	Α.	Group Discu	ssion:	CO5, CO6			
	B. in gro						
	C.						
Mode of examination	Theory						
Weightage	Continuous	Sessional	ESE				
Distribution	Mode	Exam					
	Assessment	1.0					
	05 Marks	10	35				





Text book/s*	1.	Basic communication skills for	
		Technology, Andreja. J. Ruther	
		Ford, 2 nd Edition,Pearson	
		Education, 2011	
	2.	Communication skills, Sanjay Kumar,	
		Pushpalata, 1stEdition, Oxford Press,	
		2011	
	3.	Organizational Behaviour, Stephen .P. Robbins, 1 st Edition, Pearson, 2013	
	4.	Brilliant- Communication skills, Gill	
		Hasson, 1 st Edition, Pearson Life, 2011	
	5.	The Ace of Soft Skills:	
		Attitude, Communication and	
		Etiquette for success, Gopala	
		Swamy Ramesh, 5 th Edition,	
		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, 2 nd Edition, 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	
	10	Learning india pvt.ltd,2011	
	10.	Soft skills and professional	
		communication, Francis Peters	
		SJ, 1stEdition, Mc GrawHill	
	11	Education, 2011 Effective communication, John Adair,	
	11.	4 th Edition, Pan Mac Millan,2009	
	12.	Bringing out the best in people, Aubrey	
		Daniels, 2 nd Edition, Mc Graw Hill, 1999	





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	1	3	3	-	-	3	3	3	1	3
CO2	3	3	2	2	-	-	2	3	2	3	3
CO3	3	2	2	3	-	-	2	3	3	2	3
CO4	3	2	2	2	-	-	2	3	2	2	3
CO5	3	2	3	2	-	-	2	2	2	2	3
CO6	3	2	2	2	-	-	2	3	2	2	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Scl	hool:	SOP				
Pr	ogramme:	B.Pharma				
Br	anch:	Semester: 1				
1	Course Code	BP106 RBT				
2	Course Title	Remedial biology - Theory				
3	Credits	4				
4	Contact Hours	2-0-0				
	(L-T-P)					
	Course Type	Compulsory				
5	Course	Upon completion of the course, the student shall be able to)			
	Objective	-Know the classification and salient features of five kingdo	oms of life			
		-Understand the basic components of anatomy & physiolo -Know understand the basic components of anatomy & with special reference to human				
6	Course Outcomes	CO1: Students will be able to generalize would acquire kingdoms of life, morphology and anatomy of flowering physiology of plants and humans and various plant growth	plants, anatomy and			
		CO2: Students would be able to understand the anatomy plants and humans.	y and physiology of			
		CO3: Students will be able to apply the knowledge of the anatomy and physiology of different body parts in explaining the working patterns of different body systems.				
		CO4: The students will analyze the structures of vario origin. CO5: The students would evaluate the mechanisms of various control or				
		which the functioning of the various body organs and plants depend. Moreover, will observe the anatomical differentiation of different body parts of human.				
		CO6: The students will be able to predict about the med body organs and plants	chanisms of various			
7	Course	To learn and understand the components of living world, s	structure and			
	Description	functional system of plant and animal kingdom. Scope: To				
		understand the components of living world, structure and				
		plant and animal kingdom.	Signatural Systems of			
8	Outline syllab	12	CO Mapping			





	www.sharda.	uc.iii
1	UNIT-I A.Living world: Definition and characters of living organisms Diversity in the living world Binomial nomenclature Five kingdoms of life and basis of classification. Salient features of Monera, Potista, Fungi, Animalia and Plantae, Virus, B.Morphology of Flowering plants Morphology of different parts of flowering plants – Root, stem, inflorescence, flower, leaf, fruit, seed. C.General Anatomy of Root, stem, leaf of monocotyledons & Dicotylidones	CO1
2	A.Body fluids and circulation Coposition of blood, blood groups, coagulation of blood Composition and functions of lymph Human circulatory system Structure of human heart and blood vessels Cardiac cycle, cardiac output and ECG B.Digestion and Absorption Human alimentary canal and digestive glands Role of digestive enzymes Digestion, absorption and assimilation of digested food C.Breathing and respiration Human respiratory system Mechanism of breathing and its regulation Exchange of gases, transport of gases and regulation of respiratory volumes	CO2





3	UNIT-III	
	A.Excretory products and their elimination	
	Modes of excretion	
	Human excretory system- structure and function Urine	
	formation	
	Rennin angiotensin system	
	B.Neuralcontrol and coordination	
	Definition and classification of nervous system Structure	
	of a neuron	
	Generation and conduction of nerve impulse Structure of	
	brain and spinal cord	CO3, CO6
	Functions of cerebrum, cerebellum, hypothalamus and	,
	medulla oblongata	
	C.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	
4	UNIT-IV	
	A.Plants and mineral nutrition:	
	Essential mineral, macro and micronutrients	
	B. Nitrogen metabolism, Nitrogen cycle, biological	CO4
	nitrogen fixation	201
	C.Photosynthesis	
	Autotrophic nutrition, photosynthesis, Photosynthetic	
	pigments, Factors affecting photosynthesis.	
5	UNIT-V	
	A.Plant Respiration: Respiration, glycolysis,	
	fermentation (anaerobic).	
	B.Plant growth and development	
	Phases and rate of plant growth, Condition of	CO5
	growth, Introduction to plant growth regulators	CO5
	C.Cell - The unit of life	
	Structure and functions of cell and cell organelles.Cell division	
	D.Tissues	
	Definition, types of tissues, location and functions.	
Mode of	Theory	
examination	THEOLY	
CAMIIIIIauoli		





Weightage	Continuous	Sessional	ESE				
Distribution	Mode	Exam					
	Assessment						
	05 Marks	10	35				
Text	a. Text book of Bio	logy by S. B. Gol	khale				
book/s*	A Text book of B	iology by Dr. Th	ulajappa and Dr.				
	Seetaram.						
Other	a. A Text book of	Biology by B.V.	Sreenivasa Naidu				
References	b. A Text book of	b. A Text book of Biology by Naidu and Murthy					
	c. Botany for Deg						
	d. Outlines of Zoo	d. Outlines of Zoology by M. Ekambaranatha ayyer and					
	T. N. Ananthakris						
	e. A manual for pl						
	Gokhale and C. K						





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	-	2	-	-		2	2	2	-	2
						2					
CO2	3	-	2	-	1	2	2	2	2	1	2
CO3	3	-	-	-	-	2	2	1	1	-	2
CO4	3	-	2	-	-	2	1	2	2	-	2
CO5	3	-	-	-	-	2	2	2	2	-	2
CO6	3	_	2	-	2	2	2	1	1	-	2

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sch	ool:	SOP								
Pro	gramme:	B.Pharm								
Bra	nch:	Semester: 1								
1	Course Code	BP106 RMT								
2	Course Title	Remedial Mathematics - Theory								
3	Credits	2								
4	4 Contact 2-0-0 Hours (L-T-P)									
	Course Type	Compulsory								
5	Course Objective	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 plantKnow understand the basic components of anatomy & physiology	t							
		special reference to human								
6	Course Outcome s	CO1: Students would acquire knowledge of partial fraction and loga CO2: Students would be able to understand about matrices and deter CO3: Students will be able to apply the knowledge of Differentiation CO4: Students will be able to analyze the knowledge of Integration CO5: Students will be able to interpret differential equations. CO6: Students will be able to apply mathematical techniques.	rminants							
7	Course Descripti on This is an introductory course in mathematics. This subject deals with the introduction to Partial fraction, Logarithm, matrices and Determinant, Analytical geometry, Calculus, differential equation and Laplace transform									
8	Outline syl	llabus	CO Mapping							





 	www.sharda.ac.in	
1	UNIT – I	
	A.Partial fraction	
	Introduction, Polynomial, Rational fractions,	
	Proper and Improper fractions, Partial fraction,	
	Resolving into Partial fraction, Application of	
	Partial Fraction in Chemical Kinetics and	
	Pharmacokinetics	
	B.Logarithms	CO1, CO6
	Introduction, Definition, Theorems/Properties of	
	logarithms, Common logarithms, Characteristic and Mantissa, worked examples, application of	
	logarithm to solve pharmaceutical problems.	
	logarithm to solve pharmaceutical problems.	
	C.Function:	
	Real Valued function, Classification of real valued	
	functions,	
	Limits and continuity:	
	Introduction, Limit of a function, Definition of limit of a function	
2	UNIT -II	
	A.Matrices and Determinant:	
	Introduction matrices, Types of matrices,	
	Operation on matrices, Transpose of a matrix,	
	Matrix Multiplication, Determinants, Properties	
	of determinants,	
	B.Product of determinants, Minors and co-	CO2
	Factors, Adjoint or adjugate of a square matrix, Singular and non-singular matrices, Inverse of a	CO2
	matrix,	
	C.Solution of system of linear of equations using	
	matrix method, Cramer's rule, Characteristic	
	equation and roots of a square matrix, Cayley-	
	Hamilton theorem, Application of Matrices in	
	solving Pharmacokinetic equations	





	www.sharda.ac.in	
3	UNIT - III	
	A. Calculus	
	Differentiation : Introductions, Derivative of a	
	function, Derivative of a constant,	
	B. Derivative of a product of a constant and a	
	function, Derivative of the sum or difference of	
	two functions, Derivative of the product of	
	two functions (product formula),	
	C. Derivative of the quotient of two functions	CO3
	(Quotient formula) – Without Proof,	
	Derivative of x^n w.r.tx,where n is any rational number,	
	Derivative of e^x , Derivative of $\log_e x$, Derivative	
	of a^x . Derivative of trigonometric functions from	
	first principles (without Proof), Successive	
	Differentiation, Conditions for a function to be a	
	maximum or a minimum at a point. Application	
4	maximum of a minimum at a point. Application	
4	UNIT – IV	
	A.Analytical Geometry	
	Introduction: Signs of the Coordinates, Distance formula,	
	B.Straight Line : Slope or gradient of a straight	
	line, Conditions for parallelism and	
	perpendicularity of two lines, Slope of a line	CO4
	joining two points, Slope – intercept form of a	201
	straight line	
	C.Integration:	
	Introduction, Definition, Standard formulae,	
	Rules of integration, Method of substitution,	
	Method of Partial fractions, Integration by parts,	
	definite integrals, application	
	definite integrals, application	





			www.sharda.ac.in							
5	UNIT-V									
	definitions, Order and separable form, Homoge Differential equations, Application in solv equations B. Laplace Transforms of Laplace Transforms of Inverse Laplace transform derivatives, C.Application to solve equations, Application	definitions, Order and degree, Equations in separable form, Homogeneous equations, Linear Differential equations, Exact equations, Application in solving Pharmacokinetic equations B. Laplace Transform: Introduction, Definition, Properties of Laplace transform, Laplace Transforms of elementary functions, Inverse Laplace transforms, Laplace transform of derivatives, C.Application to solve Linear differential equations, Application in solving Chemical kinetics and Pharmacokinetics equations								
Mode of examinat ion	Theory	Theory								
Weightag e	Continuous Mode Assessment	Sessional Exam	ESE							
Distributi on	5 Marks	10	35							
Text book/s*	2. Pharmaceuti application (Panchakshar3. Integral Cale	Differential Calculus by Shanthinarayan Pharmaceutical Mathematics with application to Pharmacy by Panchaksharappa Gowda D.H. Integral Calculus by Shanthinarayan Higher Engineering Mathematics by Dr.B.S.Grewal								
Other Referenc es										





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	-	2	-	-		2	1	2	-	2
						2					
CO2	3	-	2	-	1	2	1	1	2	1	2
CO3	3	-	1	-	-	2	1	1	1	-	2
CO4	3	-	2	-	-	2	1	1	1	-	2
CO5	3	-	2	-	-	2	2	1	1	-	2
CO6	3	-	2	_	_	2	1	1	1	-	2

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)



Sc	chool:	SOP						
Pr	ogramme:	B.Pharm						
Bı	anch:	Semester: 1						
1	Course	BP107P						
	Code							
2	Course	Human Anatomy and Physiology- Practical						
	Title							
3	Credits	2						
4	Contact	0-0-4						
	Hours							
	(L-T-P)							
	Course	Compulsory						
	Type							
5	Course	1. To understand how to handle the microscope in Human Anaton	ny &					
	Objective	Physiology lab						
		2. To calculate Hb content, WBC & RBC count and Erythrocyte						
		3. To identify axial and skeletal bones of Human skeleton						
		4.To learn and practice how to record Blood Pressure of given subject						
6	Course	CO1: Student will be able to Understand how to handle the microscope in	n Human					
	Outcomes	Anatomy & Physiology lab						
		CO2: Student will be able to analyze Hb content, WBC & RBC coun	t and					
		Erythrocyte sedimentation rate						
		CO3: Student will be able to Identify axial and skeletal bones of Human sk	celeton					
		CO4: Student will be able to interpret Blood Pressure of given subject						
		CO5: Student will be able to skeletal bones of Human skeleton						
		CO6: Students shall be able to understand and know body parts						
7	Course	Practical physiology is complimentary to the theoretical discussions in I						
	Description	Practicals allow the verification of physiological processes discussed in the						
		through experiments on living tissue, intact animals or normal human beir	ngs. This is					
		helpful for developing an insight on the subject.	,					
8	Outline syllal	bus	CO					
			Mapping					
	1	UNIT-I						
		a). Study of compound microscope	CO1					
		b). Microscopic study of epithelial and connective tissue						
	c). Microscopic study of muscular and nervous tissue							





2	UNIT-II a). Identification of axial bob). Identification of append			CO2, CO6					
3	UNIT-III a). Introduction to hemocyt (WBC) count b). Enumeration of total red c). Determination of bleeding	ometry and enumeration I blood corpuscles (RBC)	C) count	CO3,					
4	UNIT-IV a). Determination of blood b). Estimation of hemoglob	NIT-IV . Determination of blood group . Estimation of hemoglobin content . Determination of erythrocyte sedimentation rate (ESR)							
5	UNIT-V a). Determination of heart rate and pulse rate b). Recording of blood pressure								
Mode of examinatio									
Weightage Distribution	Continuous Mode Assessment	Sessional Exam	ESE						
Text book/s*	05 10 35 1. Essentials of Medical Physiologyy K.Sembulingam and P.Sembulingam. Jaypee brothers medical publishers, NewDelhi. 2.Anatomy and Physiology in Health and Illness by Kathleen W.Wilson ,Churchil lLivingstone,NewYork								
Other References	Physiological basis of Me Co,Riverview,MI USA		ailor.Williams&Wilkins						





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	2	1	3	-	-	2	1	3	-	3
CO2	3	1	3	3	-	-	1	1	2	1	3
CO3	3	2	2	3	-	-	2	2	2	-	3
CO4	3	2	3	3	-	-	1	1	3	-	3
CO5	3	1	3	3	-	-	1	1	2	1	3
CO6	3	1	3	3	_	_	1	2	2	1	3

- 1-Slight (Low) 2-Moderate (Medium)
- 3-Substantial (High)





Sch	ool:	SOP								
Prog	gramme:	B.Pharm								
Bra	nch:	Semester: 1								
1	Course Code	BP108P								
2	Course Title	Pharmaceutical Analysis – Practical								
3	Credits	2								
4	Contact	0-0-4								
	Hours									
	(L-T-P)									
	Course Type	Compulsory								
5	Course Objective	Upon completion of the course, the student shall be able								
	Objective	-Know the classification and salient features of five king								
		-Understand the basic components of anatomy & physic -Know understand the basic components of anatomy &								
6	Covera	with special reference to human	to understand about							
6	Course Outcomes	CO1: Upon completion of course student shall be able limits of impurities in a particular drug and to perform								
	Gutcomes	and determine the impurities in pharmaceuticals.	minit test to identify							
		and determine the impurities in pharmaceuticals.								
		CO2: Students shall be able to perform standardization	n and analyze given							
		sample strength of drug or pharmaceuticals.	, , , , , , , , , , , , , , , , , , ,							
		CO2. Students shall be able to know the purity to	esting of drugs and							
		CO3: Students shall be able to know the purity te pharmaceuticals. They can apply these strength tests to	_							
		the sample.								
		CO4: Students shall be able to understand about elec-	trochemical analysis							
		for pharmaceutical sample.								
		CO5: Students shall be able to perform standardization	n and analyze given							
		sample strength of drug or pharmaceuticals.								
		CO6: Students shall be able toapply purity testing of pha	armaceuticals							
7	Course	Deals with the fundamentals of analytical chemistry and								
	Description	electrochemical analysis of drugs	rrr							
8	Outline syllabi		CO Mapping							
	1	LINUT	CO1, CO6							
		UNIT-I a). Limit test for Chlorides and Sulphates								
	b). Modified limit test for Chlorides and Sulphates									
	c). Limit test for Iron									
		d). Limit test for Heavy metals								
		e). Limit test for Lead Limit test for Arsenic								





2	UNIT-II								
_	a). Sodium hy	droxide		CO2					
	b). Sulphuric								
	c). Sodium thi								
	d). Potassium								
	e). Ceric amm								
3	UNIT-III	<u> </u>							
	a). Ammoniur	n chloride by	acid base titration	CO3					
		b). Sodium Chloride by precipitation titration							
4	ÚNIT-IV	 	1						
	Conductometr	Conductometric titration of strong acid against strong							
	base								
5	UNIT-V			CO5					
	a). Sodium hy	droxide							
	b). Sulphuric								
	c). Sodium thi	osulfate							
Mode of	Theory/Jury/P	ractical/Viva							
examination									
Weightage	Continuous	Sessional	ESE						
Distribution	Mode	Exam							
	Assessment								
	05	10	35						
Text book/s*	Practical hu	ıman anatom	y and physiology.by						
	S.R.Kale and	R.R.Kale.							
	A Manual o	of pharmaceut	ical biology practical by						
		_	and S.P.Shriwastava.						
			according to National						
	. .		forum of Karnataka.						
	ProfM.J.H.Sl		Torum or Mamataka.						
	FIOIM.J.H.S	11411							



Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	2	1	3	-	-	2	-	3	-	3
CO2	3	1	3	3	-	-	1	-	2	1	3
CO3	3	2	2	3	-	-	2	-	2	-	3
CO4	3	2	3	3	-	-	2	-	3	-	3
CO5	3	1	3	3	-	-	2	-	2	1	3
CO6	3	1	3	3	-	-	2	-	2	1	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





So	chool:	SOP		
-	rogramme	B.Pharm		
:				
B	ranch:	Semester: 1		
1	Course	BP109P		
	Code			
2	Course	Pharmaceutics –I Practical		
2	Title			
3	Credits Contact	0-0-4		
4	Hours	0-0-4		
	(L-T-P)			
	Course	Compulsory		
	Type			
5	Course	This course will impart basic knowledge in the area of pharmaceutics and		
Objectiv formulation of different pharmaceutical dosage forms. The student				
	e	hands-on training in the preparation of such dosage forms in the laboratory.		
6	Course	CO1: Upon completion of course student shall be able to understand about		
	Outcome	different methods of preparation of various monophasic and biphasic liquid		
	S	dosage forms.		
		CO2: Students shall be able to apply specific types of excipients used in preparation of semisolid dosage forms.		
		CO3: Students shall be able to interpret different types of pharmaceutical dosage		
		forms like		
		syrups,elixirs,solutions,paints,gargles,mouthwashes,suspensions,emulsions,powd		
		ers,ointments,pastes etc.		
		CO4: Students shall be able to differentiate between different methods of		
		preparation of pharmaceutical dosage forms.		
		CO5: Students shall be able to interpret specific types of excipients used in preparation of semisolid dosage forms.		
		CO6: Students shall be able to understand Different types of pharmaceutical		
		formulations for the human use.		
7	Course	This course is designed to impart knowledge on preparatory pharmacy and		
	Descripti	professional way of preparing various dosage forms such as monophasic liquids,		
	on	biphasic liquids,		
8	Outline syl	semisolid dosage forms etc. llabus CO Mapping		
U	Outility syl	Taous CO Mapping		





	www.sha	ua.ac.iii
1	UNIT-I A. Syrups Syrup IP'66 Compound syrup of Ferrous Phosphate BPC'68 B. Elixirs Piperazine citrate elixir Paracetamol pediatric elixir	CO1, CO6
2	UNIT-II A. Linctus Terpin Hydrate Linctus IP'66 Iodine Throat Paint (Mandles Paint) B. Solutions Strong solution of ammonium acetate Cresol with soap solution Lugol's solution	CO2, CO6
3	UNIT-III A. Suspensions Calamine lotion Magnesium Hydroxide mixture Aluminimum Hydroxide gel B. Emulsions Turpentine Liniment Liquid paraffin emulsion	CO3
4	UNIT-IV A. Powders and Granules ORS powder (WHO) Effervescent granules Dusting powder Divided powders B. Suppositories Glycero gelatin suppository Coca butter suppository Zinc Oxide suppository	CO4, CO6





5	UNIT-V A. Semisolids Sulphur ointment Non staining-iodine o Carbopal gel B. Gargles and M Iodine gargle Chlorhexidine mouthy	CO5, CO6		
Mode of examinat ion	Practical/Viva			
Weighta ge	Continuous Mode Assessment	Sessional Exam	ESE	
Distribut ion	05Marks	10	35	
Text book/s*	H.C. Ansel et al., Phand Drug Delivery Sand Walkins, New De Carter S.J., Cooper Pharmaceutical Stude Delhi. M.E. Aulton, Phar Dosage Form Design Edinburgh. 1. Indian pharmacopo 2. British pharmacopo			
Other Referenc				
es				





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	2	1	2	-		2	-	2	-	2
						2					
CO2	3	2	1	2	_	2	1	-	2	1	2
CO3	3	2	-	2	-	2	1	-	1	-	2
CO4	3	2	-	2	-	2	1	-	1	-	2
CO5	3	2	-	2	-	2	1	-	1	-	2
CO6	3	2	-	2	-	2	1	-	1	-	2

1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





School:		SOP					
Program	me:	B.Pharm					
Branch:		Semester: I					
1	Course Code	BP110P					
2	Course Title	Pharmaceutical inorganic chemistry- Practical					
3	Credits	2					
4	Contact Hours (L-T-P)	0-0-4					
	Course Type	Compulsory					
5	Course Objective	The students will know the sources of impurities and methods impurities in inorganic drugs and pharmaceuticals.	to determine the				
6	Course Outcomes	CO1: Student shall be able to understand about the limits of	-				
	Outcomes	particular drug and to perform limit test to identify and impurities in inorganic drugs and pharmaceuticals.	determine the				
		CO2: Students shall be able to perform identification te given sample of drug or pharmaceuticals. CO3: Students shall be able to interpret the purity testing drugs and pharmaceuticals. They can apply these purity te and evaluate the sample. CO4: Students shall be able to apply methods of preparation drugs and pharmaceuticals. CO5: Students shall be able to apply methods of preparation inorganic drugs. CO6: Students shall be able to evaluate the presence	g of inorganic ests to analyze of tion of various				
		compounds in biological fluids.	of morganic				
7	Course Description	Limit test for non- toxic and toxic impurities, identification of Drugs, preparation of some drugs and purity test for some is and pharmaceuticals.					
8	Outline syllabus		CO Mapping				
	1	UNIT-I a). Limit test for Chlorides and Sulphates b). Modified limit test for Chlorides and Sulphates c). Limit test for Iron d). Limit test for Heavy metals e). Limit test for Lead Limit test for Arsenic	CO1, CO6				





			www.sharda.ac.in	
2	UNIT-II			CO2
	a). Magnesium hydr	oxide		
	b). Ferrous sulphate			
	c). Sodium bicarbon	ate		
	d). Calcium glucona	ite		
	e). Copper sulphate			
3	UNIT-III			CO3
	a). Swelling power of	of Bentonite		
	b). Neutralizing cap	acity of aluminum	hydroxide gel	
	c). Determination of	potassium iodate	and iodine in	
	potassium Iodide			
4	UNIT-IV			CO4
	a). Boric acid			
	b). Potash alum			
	c). Ferrous sulphate			
5	UNIT-V			CO5
	a). Ferrous sulphate			
	b). Sodium bicarbor	nate		
Mode of	Practical/Viva			
examination				
Weightage	Continuous	Sessional Exam	ESE	
Distribution	Mode			
	Assessment			
	05 Marks	10	35	
Text book/s*	Practical human	anatomy and p	physiology. by	
	S.R.Kale and R.R.	•		
	A Manual of n	harmaceutical bi	ology practical by	
	S.B.Gokhale, C.K.			
	,			
	Biology practical	=		
		ology forum	of Karnataka.	
	ProfM.J.H.Shafi			
Other				
References				



Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	2	3	3	_	_	2	_	2	3	3
COI	3	2	3	3			2		2	3	3
CO2	3	2	3	3	-	-	2	-	2	3	2
CO3	3	2	3	3	-	-	2	-	3	3	2
CO4	3	2	3	3	-	-	2	-	2	3	2
CO5	3	2	3	3	-	-	2	-	3	3	2
CO6	3	2	3	3	-	-	2	-	2	3	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





School	•	SOP	
Progra	mme:	B.Pharm	
Branch		Semester: 1	
1	Course Code	BP112 RBP	
2	Course Title	Remedial biology Practical	
3	Credits	2	
4	Contact Hours (L-T-P)	0-0-2	
	Course Type	Compulsory	
5	Course	1. To understand how to handle the microscope in lab.	
	Objective	2. To identify axial and skeletal bones of Human skeleton	
		3. To learn and practice how to record Blood Pressure of g	given subject.
		4. To Study morphology and microscopy of Stem, Rooffruit, flower and their modifications.5. Identification of blood group.	
6	Course	CO1: Student will be able to understand how to handle th	e microscope in
	Outcomes	lab.	· moroscopo m
		CO2: Student will be able to Identify axial and skeletal backeleton	oones of Human
		CO3: Student will be able to Record Blood Pressure of giv	en subject.
		CO4: Student will be able to understand study Mon histological characteristics of root, Stem, Leaf, Seed, Fruit a	
		CO5: Student will be able to interpret the blood group of su CO6: Students will be able to apply practical knowledge of Bones.	
7	Course Description	Practical is complimentary to the theoretical discussions read allow the verification of physiological processes discusses through experiments on living tissue, intact animhuman beings and plants. This is helpful for developing a subject.	cussed in theory mals or normal
8	Outline syllabus		CO Mapping
	1	UNIT-I a). Study of compound microscope b). Microscopic study of leaves and flowers c). Microscopic study of roots and stem	CO1





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2	· ·	UNIT-II a). Identification of axial bones b). Identification of appendicular bones						
3	3 UNIT-III a). Determination of blood group b). Estimation of hemoglobin content							
4	UNIT-IV Determination of he	art rate and pulse 1	rate	CO4				
5	UNIT-V Recording of blood	UNIT-V Recording of blood pressure						
Mode of examination	Practical/Viva							
Weightage Distribution	Continuous Mode Assessment	Sessional Exam	ESE					
Text book/s*	05 Marks Practical human S.R.Kale and R.R.J		15 hysiology. by					
	A Manual of p S.B.Gokhale, C.K.	harmaceutical bi Kokate and S.P.Sh						
		Biology practical manual according to National core urriculum. Biology forum of Karnataka. Prof I.J.H.Shafi						
Other References								





Pos	РО	РО	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos	1	2									
CO1	3	2	3	3	-	-	1	-	1	3	3
CO2	3	2	3	3	-	-	1	-	1	3	2
CO3	3	2	3	3	-	-	1	-	1	3	2
CO4	3	2	3	3	-	-	1	-	1	3	2
CO5	3	2	3	3	-	-	1	-	1	3	2
CO6	3	2	2	1	2	1	2	-	1	-	1

1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)



Scho	ool:	SOP
Programme:		B.Pharm
Bra	nch:	Semester: 2
1	Course Code	BP 201T
2	Course Title	Human Antomy & Physiology-II
3	Credits	4
4	Contact	3-1-0
	Hours	
	(L-T-P)	
	Course Type	Compulsory
5	Course Objective	1. Explain the gross morphology, structure and functions of various organs of the human body.
		2. Describe the various homeostatic mechanisms and their imbalances.
		3. Identify the various tissues and organs of different systems of human body.4. Perform the hematological tests like blood cell counts, haemoglobin estimation,
		bleeding/clotting time etc and also record blood pressure, heart rate, pulse and respiratory volume.
		5. Appreciate coordinated working pattern of different organs of each system6. Appreciate the interlinked mechanisms in the maintenance of normal functioning (homeostasis) of human body.
6	Course Outcomes	CO1 : The students will understand the structure and functions of various systems and organs of the body. Also about increase the understanding about genes and genetics.
		CO2: The student will be able to summarize the functioning of various body systems and their homeostasis.
		CO3: The student will be able to apply the knowledge of the functioning of various body systems and the structures of the organs involved in it.
		CO4: The students will analyze the correlation of various body systems and how they result in particular kind of functions.
		 CO5: The students would evaluate the processes like respiration, Excretion, digestion hormone release and reproduction by understand their mechanisms. CO6: The students would evaluate the processes and analyze their correlation with various body systems





7	Course Description	This subject is designed to impart fundamental knowledge on the struct functions of the various systems of the human body. It also helps in und both homeostatic mechanisms. The subject provides the basic knowledge to understand the various disciplines of pharmacy.	lerstanding
8	Outline syllab	ous	CO Mapping
		 UNIT-I A. Nervous system Organization of nervous system, neuron, neuroglia, classification and properties of nerve fibre, electrophysiology, action potential, nerve impulse, receptors, synapse, neurotransmitters. B. Central nervous system: Meninges, ventricles ofbrain and cerebrospinal fluid.structure and fu nctions of brain (cerebrum, brain stem, cerebellum), C. spinal cord (gross structure, functions of afferent and efferent nervetracts, reflexactivity) 	CO1, CO6
	2	 UNIT-II A. Digestive system Anatomy of GI Tract with special reference to anatomy and functions of stomach, (Acid production in the stomach, regulation of acid production through parasympathetic nervous system, pepsinrole in protein digestion) small intestine and large intestine, anatomy and functions of salivary glands, pancreas and liver, movements of GIT, digestion and absorption of nutrients and disorders of GIT. B. Energetics Formation and role of ATP, CreatininePhosphateandBMR. C. Joints Structural and functional classification, types of joints movements and its articulation 	CO2





	3	UNIT-III			
		A. Respiratory syster	n 10 hours		CO3,
		Anatomy of respiratory sylungs, mechanism of respiratory Sylungs and capac respiration, and resuscitation methods B. Urinary system			
		Anatomy of urinary tract and nephrons, functions urine formation, C. micturition reflex role of RAS in kidn	of kidney and urinary	tract, physiology of acid base balance,	
-	4	UNIT-IV			CO4
		A. Endocrine system			
		Classification of hormones			
		B.structure and functions of	of pituitary gland, thyroi	id gland, parathyroid	
		gland, adrenalgland, C.pancreas, pineal gland, the	hymus and their disords	arc	
-	5	UNIT-V	nymus and their disorde	.18.	CO5
		A. Reproductive system			
		Anatomy of male and fem and femalereproductive menstruation, fertilization spermatogenesis, oogenes B. Introduction to genetic	system, sex hormon sis, pregnancy and partu	nes, physiology of	
		C. Chromosomes, genes a	and DNA, protein synth	nesis, genetic pattern	
	N 1	of inheritance			
	Mode of examination	Theory			
	Weightage	Continuous Mode	Sessional Exam	ESE	
	Distribution	Assessment	STORIGHT LIMIN		
		10 Marks	15	75	
	Text book/s*	Practical human anator and R.R.Kale.			
		A Manual of pharmace C.K.Kokate and S.P.Shriv			
		Biology practical manual	according to National	l core curriculum	
		.Biology forum of Karnat	aka. ProfM.J.H.Shafi		





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	1	2	2	1	3	1	-	1	2	3
CO2	2	2	2	2	1	2	1		1	2	2
CO2	3	2	2	2	1	3	1	-	1	2	3
CO3	3	2	1	2		3	1	-	1	3	3
CO4	3	2	1	2	2	3	2	-	1	2	3
CO5	3	2	2	3	2	3	2	-	1	3	3
CO6	3	2	1	2	2	3	2	-	1	2	3

- 1-Slight (Low) 2-Moderate (Medium)
- 3-Substantial (High)





School:		SOP				
Progran	nme:	B.Pharma				
Branch:		Semester: 2				
1	Course Code	BP202T				
2	Course Title	Pharmaceutical organic chemistry-I Theory				
3	Credits	4				
4	Contact Hours (L-T-P)	3-1-0				
	Course Type	Compulsory				
5	Course Objective	Upon completion of the course the student shall be able to 1. Write the structure, name and the type of isomerism of the organic compound				
		2. Write the reaction, name the reaction and orientation of reactions.				
		3. Account for reactivity/stability of compounds.				
		4. Identify/ confirm the identification of organic compound.				
6	Course Outcomes	CO1: The students will have the knowledge to identify, name, and write the structure of different aliphatic compounds and their derivatives. CO2: The students will be able to understand and explain the mechanism behind the naming reactions of different aliphatic compounds and their derivatives. CO3: The students can apply the knowledge to prepare the derivatives of aliphatic compounds with different functional groups. CO4: Students will analyze the chemical reactions, stabilities of organic compounds and properties of the compounds prepared by them in the laboratory. CO5: Students would evaluate by comparing compounds prepared by them with standard compounds by chemical and physical properties				
		CO6: Students will analyze the chemical reactions and stabilities of organic compounds				
7	Course Description	This subject deals with classification and nomenclature of simple organic compounds, 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.				
8	Outline syllabus	CO Mapping				





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1	UNIT-I	CO1
	A. Classification, nomenclature and isomerism	
	Classification of Organic Compounds	
	B.Common and IUPAC systems of	
	nomenclature of organic compounds	
	C.Structural isomerisms in organic compounds	
	c.ou detaila isomerisms in organic compounds	
2	UNIT-II	CO2
	A. Alkanes*, Alkenes* and Conjugated dienes*	
	SP ³ hybridization in alkanes, Halogenation of alkanes,	
	uses of paraffins.	
	B. Stabilization of alkenes, SP2 hybridization of 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	
	C. reactions of alkenes, Markownikoff's orientation, free radical addition reactions of alkenes, Anti	
	Markownikoff's orientation.	
	Stability of conjugated dienes, Diel-Alder, electrophilic addition, free radical addition reactions of	
	conjugated dienes, allylic rearrangement	
3	UNIT-III	CO3
	A. Alkyl halides*	CO3
	SN ₁ and SN ₂ reactions - kinetics, order of reactivity of	
	alkyl halides, stereochemistry	
	and rearrangement of carbocations.	
	SN1 versus SN2 reactions, Factors affecting	
	SN1 and SN2 reactions	
	Structure and uses of ethylchloride,	
	Chloroform, trichloroethylene,	
	tetrachloroethylene, dichloromethane,	
	tetrachloromethane and iodoform.	
	B. Alcohols*- Qualitative tests, Structure and	
	uses of Ethyl alcohol,	
	C.Methyl alcohol, chlorobutanol, Cetosteryl alcohol,	
	Benzyl alcohol, Glycerol, Propylene glycol	





UNIT-IV A. Carbonyl compounds* (Aldchydes and ketones) Nucleophilic addition, Electromeric effect, aldol condensation, Crossed Aldolcondensation, Cannizzaro reaction, Crossed Cannizzaro reaction, Benzoin condensation, Perkin condensation, B. Qualitative tests of carbonyl compounds Structure and uses of Formaldehyde, Paraldehyde, Acetone, C. Chloral hydrate, Hexamine, Benzaldehyde, Vanilin, Cinnamaldehyde. 5 UNIT-V A. Carboxylic acids* Acidity of carboxylic acids, effect of substituents on acidity, inductive effect and qualitative tests for carboxylic acid, amide and ester Structure and Uses of Acetic acid, Lactic acid, Tartaric acid, Citric acid, Succinic acid. B. Oxalic acid, Salicylic acid, Benzoic acid, Benzyl benzoate, Dimethyl phthalate,Methyl salicylate and Acetyl salicylic acid Aliphatic amines. C.Basicity, effect of substituents on basicity, identification test, Structure and uses of Ethanolamine,ehthylenediamine, amphetamine. Mode of examination Weightage Distribution Weightage Distribution Sessional Exam ESE Mode Assessment 10 Marks 15 75 Text book/s* Practical human anatomy and physiology. by S.R.Kale and R.R.Kale. A Manual of pharmaceutical biology practical by S.B.Gokhale, C.K.Kokate and S.P.Shriwastava. Biology practical manual according to National core curriculum. Biology forum of Karnataka. Prof M.J.H.Shafi		1		WWW.Sharda.ac.iii	_
A. Carbonyl compounds* (Aldehydes and ketones) Nucleophilic addition, Electromeric effect, aldol condensation, Crossed Aldolcondensation, Cannizzaro reaction, Crossed Cannizzaro reaction, Benzoin condensation, Perkin condensation, B. Qualitative tests of carbonyl compounds Structure and uses of Formaldehyde, Paraldehyde, Acetone, C.Chloral hydrate, Hexamine, Benzaldehyde, Vanilin, Cinnamaldehyde. 5 UNIT-V A. Carboxylic acids* Acidity of carboxylic acids, effect of substituents on acidity, inductive effect and qualitative tests for carboxylic acids, amide and ester Structure and Uses of Acetic acid, Lactic acid, Tartaric acid, Citric acid, Succinic acid. B. Oxalic acid, Salicylic acid, Benzoic acid, Benzyl benzoate, Dimethyl phthalate,Methyl salicylate and Acetyl salicylic acid Aliphatic amines. C.Basicity, effect of substituents on basicity, identification test, Structure and uses of Ethanolamine, ethylenediamine, amphetamine. Mode of Ethanolamine, Experimentation of Ethanolamine, amphetamine. Mode Sessional Exam ESE Mode Assessment 10 Marks 15 75 Text book/s* Practical human anatomy and physiology. by S.R.Kale and R.R.Kale. A Manual of pharmaceutical biology practical by S.B.Gokhale, C.K.Kokate and S.P.Shriwastava. Biology practical manual according to National core curriculum. Biology forum of Karnataka. Prof M.J.H.Shafi	4	IINIT-IV			CO4, CO6
Nucleophilic addition, Electromeric effect, aldol condensation, Crossed Aldolcondensation, Cannizzaro reaction, Crossed Cannizzaro reaction, Benzoin condensation, Perkin condensation, B. Qualitative tests of carbonyl compounds Structure and uses of Formaldehyde, Paraldehyde, Acetone, C.Chloral hydrate, Hexamine, Benzaldehyde, Vanilin, Cinnamaldehyde. S			ounds* (Aldehyd	es and	
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Cos	1										
CO1	3	2	-	-	-	2	2	2	1	1	3
CO2	3	2	2	2	-	2	2	2	2	1	3
CO3	3	2	_	1	_	3	2	2	_	2	2
CO4	3	2	2	2	-	3	2	2	1	2	3
CO5	3	2	1	2	-	3	2	2	1	2	3
CO6	3	3	2	3	-	2	2	3	1	1	3
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- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





School:		SOP
Program	me:	B.Pharm
Branch:		Semester: 2
1	Course Code	BP203 T
2	Course Title	Biochemistry- Theory
3	Credits	4
4	Contact Hours (L-T-P)	3-1-0
	Course Type	Compulsory
5	Course	Upon completion of the course, the student shall be able to
	Objective	-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 with special reference to human
6	Course Outcomes	CO1: The students will understand the structure and functions of carbohydrate, lipids, nucleic acids, amino acids and proteins. Concept of free energy, endergonic and exergonic reaction, Relationship, between free energy. CO2:The student will be able to summarize the Citric acid cycle-Pathway, energetics and significance, HMP shunt and its significance; Glucose-6-Phosphate dehydrogenase (G6PD) deficiency CO3: The student will be able to apply the knowledge of the Amino acid and lipid metabolism. CO4: The students will analyze the correlation of Nucleic acid metabolism and genetic information transfer. CO5: The students would Introduction, properties, nomenclature and IUB classification of enzymes, Enzyme kinetics (Michaelis plot, Line Weaver Burke plot). CO6: The student will understand the importance of biochemistry.
7	Course Description	Biochemistry deals with complete understanding of the molecular levels of the chemical 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.





8	Outline syllabus	www.sharda.ac.in	CO Mapping
	1	UNIT-I	CO1
		A. Biomolecules and Bioenergetics	
		Topic1- Introduction, classification, chemical nature and biological role of carbohydrate. Topic2- Introduction, classification, chemical nature	
		and biological rolelipids, nucleic acids,amino acids and proteins.	
		B. Topic3-Concept of free energy, endergonic and exergonic reaction, Relationship between free energy, enthalpy and entropy; Redox potential.	
		C. Energy rich compounds; classification; biological significances of ATP and cyclic AMP	
	2	UNIT-II A. Carbohydrate metabolism and Biological oxidation	CO2, CO6
		Topic1- 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). B. Topic2- Gluconeogenesis- Pathway and its significance Hormonal regulation of blood glucose level and Diabetes mellitus 	
		C. Topic3- Electron transport chain (ETC) and its mechanism. Oxidative phosphorylation & its mechanism and substrate	
		Phosphorylation, Inhibitors ETC and oxidative	
	3	phosphorylation/Uncouplrslevel UNIT-III	CO3
	3	A. Lipid metabolism and Amino acid metabolism	CO3
		β-Oxidation of saturated fatty acid (Palmitic acid) 61Formation and utilization of ketone bodies; ketoacidosis De novo synthesis of fatty acids (Palmitic acid) Biological significance of cholesterol and conversion of cholesterol into bile acids, steroid hormone and vitamin D Disorders of lipid metabolism: Hypercholesterolemia, atherosclerosis, fatty liver and	
		obesity. B. General reactions of amino acid metabolism: (Phenyketonuria, Albinism, alkeptonuria, tyrosinemia) Synthesis and significance of biological substances; C.HT,melatonin,dopamine,noradrenaline,adrenaline Catabolism of heme; hyperbilirubinemia and jaundice	





			CO4				
UNIT-V A. Enzymes			CO4				
IUB classification o Enzyme kinetics (I plot) Enzyme inhibi	f enzymes Michaelis plot, L tors with examples	ine Weaver Burke					
induction and regulation C.Topic3- Therape	epression, alloste eutic and diagno	eric enzymes stic applications of					
Theory							
Continuous Mode Assessment	Sessional Exam	ESE					
10 Marks	15	75					
Practical human	anatomy and p						
=	A Manual of pharmaceutical biology practical by S.B.Gokhale, C.K.Kokate and S.P.Shriwastava.						
	=						
	A. Enzymes Topic1- Introduction of Enzyme kinetics (It plot) Enzyme inhibit B.Topic2- Regulation of Enzyme inhibit B.Topic2- Regulation C.Topic3- Theraped enzymes and isoer biochemical function. Theory Continuous Mode Assessment 10 Marks Practical human S.R.Kale and R.R.I. A Manual of p S.B.Gokhale, C.K.I. Biology practical curriculum .Biochemical function.	A. Enzymes Topic1- Introduction, properties, no IUB classification of enzymes Enzyme kinetics (Michaelis plot, Le plot) Enzyme inhibitors with examples B.Topic2- Regulation of enzymenduction and repression, allosted regulation C.Topic3- Therapeutic and diagnosed enzymes and isoenzymes, Coenzyment biochemical functions Theory Continuous Sessional Examemode Assessment 10 Marks 15 Practical human anatomy and properties of the properties of	Topic1- Introduction, properties, nomenclature and IUB classification of enzymes Enzyme kinetics (Michaelis plot, Line Weaver Burke plot) Enzyme inhibitors with examples B.Topic2- Regulation of enzymes: enzyme induction and repression, allosteric enzymes regulation C.Topic3- Therapeutic and diagnostic applications of enzymes and isoenzymes, Coenzymes —Structure and biochemical functions Theory Continuous Sessional Exam ESE Mode Assessment 10 Marks 15 75 Practical human anatomy and physiology. by S.R.Kale and R.R.Kale. A Manual of pharmaceutical biology practical by S.B.Gokhale, C.K.Kokate and S.P.Shriwastava. Biology practical manual according to National core curriculum .Biology forum of Karnataka.				

Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	-	3	-	-	2	-	2	1	1	3
CO2	3	1	3	2	-	2	-	2	2	1	3
CO3	3	1	2	1	-	3	1	2	-	2	2
CO4	3	-	2	2	-	3	2	2	1	2	3
CO5	3	2	2	3	_	2	2	3	1	1	3
CO6	3	2	2	3	-	1	2	2	1	1	3





- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High



School:		SOP
Programi	me:	B.Pharm
Branch:		Semester: 2
1	Course Code	BP204T
2	Course Title	Pathophysiology- Theory
3	Credits	4
4	Contact Hours (L-T-P)	3-1-0
	Course Type	Compulsory
5	Course Objective	 To distinguish between environmental factors, physical, psychosocial, and cognitive characteristics of various diseases and conditions. To understand basic concepts of inflammatory diseases To Demonstrate and understand mechanisms of diseases, the diagnosis of diseases, and the treatment of diseases To understand how the various organ systems are interrelated, and use this understanding to promote a holistic approach towards the evaluation and treatment of patients
6	Course Outcomes	CO1: Student will be able to Distinguish environmental factors, physical, psychosocial, and cognitive characteristics of various diseases and conditions. CO2: Student will be able to apply concepts and elements of inflammatory diseases CO3: Student will be able to Demonstrate an understanding of the mechanisms of diseases, the diagnosis of diseases, and the treatment of diseases CO4: Students will be able to understand how the various organ systems are interrelated, and use this understanding to promote a holistic approach towards the evaluation and treatment of patients CO5: Students will be able to compare and discriminate between the Infectious and sexually transmitted diseases. CO6: Students will be able to understand about infectious diseases.
7	Course Description	Pathophysiology is the study of causes of diseases and reactions of the body to such disease producing causes. This course is designed to impart a thorough knowledge of the relevant aspects of pathology of various conditions with reference to its pharmacological applications, and understanding of basic pathophysiological mechanisms. Hence it will not only help to study the syllabus of pathology, but also to get baseline knowledge required to practice medicine safely, confidently, rationally and effectively.
8	Outline syllabus	CO Mapping





	www.sharda.ac.in	1
	 UNIT-I A.Basic principles of Cell injury and Adaptation & Basic mechanism involved in the process of inflammation and repair Causes of cellular injury,Pathogenesis (Cell membrane damage, Mitochondrial damage, Ribosome damage, Nuclear damage). A. Morphology of cell injury – Adaptive changes (Atrophy, Hypertrophy, hyperplasia, Metaplasia, Dysplasia),Cell swelling, Intra cellular accumulation, Calcification, Enzyme leakage and Cell Death Acidosis& Alkalosis, Electrolyte imbalance. B. Introduction, Clinical signs of inflammation, Different types of Inflammation,Mechanism of Inflammation – Alteration in vascular permeability and blood flow, migration of WBC's, C. Mediators of inflammation,Basic principles of wound healing in the skin,Pathophysiology of Atherosclerosis 	CO1
2	UNIT-II A. Cardiovascular, Respiratory and Renal Diseases Hypertension, congestive heart failure, ischemic heart disease (angina,myocardial infarction, B.atherosclerosis and arteriosclerosis)Asthma,Chronic obstructive airways diseases. C. Acute and chronic renal failure	CO2
3	UNIT-III A.Hematological, Endocrine, Nervous and GIT diseases Iron deficiency, megaloblastic anemia (Vit B12 and folic acid), sickle cell anemia, thalasemia, hereditary acquired anemia, B.haemophilia Diabetes, thyroid diseases, disorders of sex hormones & Peptic ulcer C. Epilepsy, Parkinson's disease, stroke, psychiatric disorders: depression, schizophrenia and Alzheimer's disease.	CO3





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4	UNIT-IV A. Cancer and in Classification, etion Inflammatory bowel of alcoholic liver diseases B. Rheumatoid C. osteoporosis	CO4						
5	UNIT-V A. Infectious &Sex Meningitis,Typhoid, I B.Tuberculosis Urinary tract infection C.AIDS, Syphilis & C	CO5, CO6						
Mode of examination	Theory	Theory						
Weightage Distribution	Continuous Mode Assessment	Sessional Exam	ESE					
Text book/s*	10 Marks Practical human S.R.Kale and R.R. A Manual of pharm S.B.Gokhale, C.K.K Biology practical marks	10 Marks 15 Practical human anatomy and physiology. by S.R.Kale and R.R.Kale. A Manual of pharmaceutical biology practical by S.B.Gokhale, C.K.Kokate and S.P.Shriwastava. Biology practical manual according to National core curriculum .Biology forum of Karnataka. Prof						
Other References								





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	1	-	-	1	3	1	-	1	2	3
CO2	3	2	-	2	1	3	1	-	1	2	3
CO3	3	2	1	2		3	1	-	1	3	3
CO4	3	2	1	2	2	3	2	-	1	2	3
CO5	3	2	2	3	2	3	2	-	1	3	3
CO6	3	2	2	3	2	3	2	-	1	3	3

1-Slight (Low)

2-Moderate (Medium) 3-Substantial (High)



School:		SOP						
Program	nme:	B.Pharm						
Branch		Semester: II						
1	Course Code	BP205T						
2	Course Title	Computer applications in Pharmacy- Theory						
3	Credits	4						
4	Contact Hours (L-T-P)	3-1-0						
	Course Type	Compulsory						
5	Course Objective	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						
6	Course	Upon completion of the course, the student shall be able to						
	Outcomes	CO1: understand the Binary number system						
		CO2: interpret the web technologies						
		CO3: apply about application of computers in Pharmacy						
		CO4: the Bioinformatics Databases, Concept of Bioinformatics						
		CO5: Computers as data analysis in Preclinical development:						
		CO6: Students will understand use of Computers in Preclinical	levelopment:					
7	Course Description	This subject deals with the introduction Database, Database system, computer application in clinical studies and use of database	Management					
8	Outline syllabus		CO					
			Mapping					
	1		CO1					
		UNIT-I						
		A. Number system : Binary number system, Decimal number						
		system, Octal number system, Hexadecimal number						
		systems, conversion decimal to binary, binary to decimal,						
		octal to binary etc, binary addition, binary subtraction –						
		One's complement ,Two's complement method, binary						
		multiplication, binary division						





			www.snarua.ac.iii						
2	UNIT-II			CO2					
	A. Webtechnologies: on to HTML, XMI Programmeming introduction to we and Server Product	L,CSS and languages, eb servers		CO2					
	B. Introduction to Pharmacy Drug date		L, MS ACCESS,						
3	UNIT-III			CO3					
	Mathematical mode Pharmacy, Electronsystems, barcode dispensing of drumonitoring Diagno Patient Monitoring	ge and retrieval, el in Drug design, I onic Prescribing a medicine identifica gs, mobile technol	Pharmacokinetics, Hospital and Clinical and discharge (EP) tion and automated ogy and adherence diagnostic System,	COS					
4			Objective of abases, Concept of matics in Vaccine	CO4					
5	Chromatographic Information mana	A. Computers as data analysis in Preclinical development:							
Mode of examination	Theory/Jury/Practical/	Viva							
Weightage Distribution	Continuous Mode Assessment	Sessional Exam	ESE						
	10	15	75						





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





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	1	-	-	1	3	1	-	1	2	3
CO2	3	2	-	2	1	3	1	-	1	2	3
CO3	3	2	1	2		3	1	-	1	3	3
CO4	3	2	1	2	2	3	2	-	1	2	3
CO5	3	2	2	3	2	3	2	-	1	3	3
CO6	3	2	2	2	2	3	2		1	3	3

- 1-Slight (Low) 2-Moderate (Medium)
- 3-Substantial (High)





Scho	ol:	SOP							
Prog	ramme:	B. Pharm							
Bran		Semester: II							
1	Course Code	BP206 T							
	Course Title	Environmental Sciences (Theory)							
3	Credits	4							
4	Contact Hours (L-T-P)	3-1-0							
	Course Type	Compulsory							
5	Course Objective	Upon completion of the course the student shall be able to:							
		Create the awareness about environmental problems among lear Impart basic knowledge about the environment and its allied pro-							
		Develop an attitude of concern for the environment.							
		Motivate learner to participate in environment protection and environment improvement.							
		Acquire skills to help the concerned individuals in identifying and solving environmental problems.							
		Strive to attain harmony with Nature.							
6	Course Outcomes	CO1: Student shall be able to understand the Multidisciplinary nature of environmental studies							
		CO2: Student shall be able to apply Concept of an ecosystem. CO3: Student shall be able to analyze Structure and function of an ecosystem.							
		CO4: Student shall be able to explain Structure and function of anenvironment							
		CO5: Student shall be able to analyze computer data in Preclinical development:							
		CO6: Students will be able to understand about water and air po	ollution						
7	Course Description	Environmental Sciences is the scientific study of the environmental and the status of its inherent or induced changes on organisms. Only the study of physical and biological characters of the enalso the social and cultural factors and the impact of man on environmental Sciences is the scientific study of the the scientif	It includes not vironment but						
8	Outline syllabu	S S	СО						
			Mapping						
	1	UNIT-I	CO1						
		A. The Multidisciplinary nature of environmental studies Natura lResources Renewable and non-renewable resources:							
		B . Natural resources and associated problems							
		(a)Forest resources; b) Water resources; c) Mineral resources; C.Food resources; e) Energy resources; f) Land resources:							
		Role of an individual in conservation of natural resources.							





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2	UNIT-II				COA					
	Ecosystem				CO2					
	-	t of an ecosy								
		B. Structure and function of an ecosystem.								
		• •		ures, structure and						
		the ecosyst								
		•	-	n; Desert ecosystem;						
	_	cosystems (ponds, streams, la	ikes, rivers, oceans,						
2	estuaries									
3	UNIT-III				GO2 GO6					
		ental Pollution	on: Air pollution; \	Water pollution; Soil	CO3, CO6					
3.5.1		pollution								
Mode of examination	Theory	Theory								
Weightage	Continuous Mode		Sessional Exam	ESE						
Distribution	Assessmen	nt								
	10		15	75						
Text book/s*	1.									
		_	ational Pvt, Publish	ers,						
		Bangalore								
	2.	_		nental Biology, Nidi						
		Publ. Ltd.	Bikaner.							
	3.		Erach, The Biodiver	•						
		-	in Pu blishing Pvt.	Ltd.,						
	_		d - 380 013, India,							
	4.		C., 1989, Hazardov							
	_		n, McGraw Hill Inc							
	5.	Oxford	Marine Pollution,	Cianderson Press						
	6.	Cunninghan	n, W.P. Cooper,	T.H.						
		Gorhani,	E & Hepw							
		M.T.	· ·	nvironmental						
		Encycloped		l. House,						
	7	Mumbai, 1	-	nistry, Wiley Eastern						
	7.	Ltd.	nvironmentai Chen	msuy, whey Eastern						
	8.		arth, Centre for Sci	ence and						
	ο.	Environme		chec and						
	<u> </u>	PHAHOIIIIC	111		I					





Pos	PO	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos	1										
CO1	3	1	-	-	1	3	1	-	1	2	3
CO2	3	2	-	2	1	3	1	-	1	2	3
CO3	3	2	1	2		3	1	-	1	3	3
CO4	3	2	1	2	2	3	2	-	1	2	3
CO5	3	2	2	3	2	3	2	-	1	3	3
CO6	3	2	2	3	2	3	2	_	1	3	3

Slight (Low) 2-Moderate (Medium)

3-Substantial (High)





School:		SOP						
Program	me:	B. Pharm						
Branch:		Semester: 2						
1	Course Code	BP207 P						
2	Course Title	Human Anatomy & Physiology-II Practical						
3	Credits	2						
4	Contact Hours (L-T-P)	0-0-4						
	Course Type	Compulsory						
5	Course	Upon completion of the course, the student shall be able to						
	Objective	-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 with special reference to human						
6	Course Outcomes	CO1 : The students will understand the structure and functions of various tissues and organs of the body. Also correlate their relevance with each other.						
		CO2: The student will be able to summarize the functioning of various body systems and their homeostasis.						
		CO3: The student will be able to apply the knowledge of the anatomy and physiology of different body parts in explaining the working patterns of different body systems.						
		CO4: The students will analyze the structures of various tissues and their origin to evaluate their damage and repair process.						
		CO5: The students would evaluate the mechanisms of various processes on which the functioning of the various body organs depend. Moreover, will observe the anatomical differentiation of different body parts.						
		CO6: The students would evaluate the molecular mechanisms of various body pathways.						
7	Course Description	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.						
8	Outline syllabus	CO Mapping						





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1	UNIT-I To study the integur system, endocrine system	• •		CO1
2	UNIT-II a). To demonstrate to different types of tast c). Recording of boo	the function of olfa ste, visual acuity, r	actory nerve,	CO2
3	UNIT-III a). To demonstrate processing mechanism. b). Determination of	positive and negati		CO3
4	UNIT-IV a). Study of digestive urinary and reproduction charts and speciments b). Recording of bases	diovascular systems, the help of models,	CO4, CO6	
5	UNIT-V a). Study of family produced diagnosistest. b). Demonstration of Permanent slides of	CO5, CO6		
Mode of examination		That organis and g	onaus.	
Weightage Distribution	Continuous Mode Assessment	Sessional Exam	ESE	
	05	10	35	
Text book/s*	1. Essentials of I Sembulingam brothers medi 2. Anatomy and Illness by Kat Churchill Liv 3. Physiological Best and Taile Co,Riverview			
Other References				





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	1	-	-	1	3	1	-	1	2	3
~~~					_		_				
CO2	3	2	-	2	1	3	1	-	1	2	3
CO3	3	2	1	2		3	1	-	1	3	3
CO4	3	2	1	2	2	3	2	-	1	2	3
CO5	3	2	2	3	2	3	2	-	1	3	3
CO6	3	2	2	3	2	3	2	-	1	3	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)



*	SHAKDA
	UNIVERSITY
	Beyond Boundaries
www.shard	a.ac.in

School:		SOP
Program	me:	B. Pharm
Branch:		Semester: 2
1	Course Code	BP208 P
2	Course Title	Pharmaceutical Organic Chemistry-I Practical
3	Credits	2
4	Contact Hours (L-T-P)	4
	Course Type	Compulsory
5	Course Objective	This subject deals with classification and nomenclature of simple organic compounds, 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.
6	Course Outcomes	CO1 - Students will be able to understand practical laboratory skills and get hands-on training of systematic qualitative analysis of organic compounds.  CO2 - Students will be able to apply knowledge and understanding of systematic qualitative analysis of organic compounds and will be able to apply this knowledge in identification of organic compounds.  CO3 - Students will be able to prepare the solid derivatives of organic compounds and can apply this knowledge for the identification of drugs and pharmaceuticals also use these skills to modify various characteristics of drugs and Pharmaceuticals.  CO4 - Students will be able to analyze professional transferable skills as exemplified by problem solving and teamwork.  CO5 - Students will be able to generalize skills for the predicting the atomic structure of drugs and chemicals.  CO6 - Students will be able to analyze models of different organ system that will elaborate the learning



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7	Course Description	<ol> <li>Systematic qualitative analysis of unknown compounds like</li> <li>Preliminary test: Color, odour, aliphatic/aromat saturation and unsaturation, etc.</li> <li>Detection of elements like Nitrogen, Sulphur an Lassaigne's test.</li> <li>Functional group test like Phenols, Amides/ Urea, Amines, Carboxylic acids, Aldehydes and Keto Esters, Aromatic and Halogenated Hydrocarbons, Ni and Anilides.</li> <li>Melting point/Boiling point of organic compounds</li> <li>Identification of the unknown compound from the melting point/ boiling point</li> <li>Preparation of the derivatives and confirmation or compound by melting point/ boiling point.</li> <li>Minimum 5 unknown organic compounds to systematically.</li> <li>Preparation of suitable solid derivatives from organic</li> <li>Construction of molecular models</li> </ol>	ic compounds, d Halogen by Carbohydrates, nes, Alcohols, tro compounds literature Using f the unknown be analyzed
8	Outline syllabus		CO Mapping
	1	UNIT-I I. Experiments involving preliminary test: Color, odour, aliphatic/aromatic compounds, saturation and unsaturation, etc. Physical characteristics Flame Test Bromine Test	CO1
	2	UNIT-II Element Detection (Lassaigne's test)	CO2
	3	UNIT-III Solubility test	CO3
	4	UNIT-IV Functional group test like Phenols, Amides/ Urea, Carbohydrates, Amines, Carboxylic acids, Aldehydes and Ketones, Alcohols, Esters, Aromatic and Halogenated Hydrocarbons, Nitro compounds and Anilides.	CO4,





	www.shards.ac.in					
5	Preparation of suital compounds	Melting point/Boiling point of organic compounds Preparation of suitable solid derivatives from organic				
Mode of examination	Practical/Viva					
Weightage Distribution	Continuous Mode Assessment	Sessional Exam	ESE			
	05	10	35			
Text book/s*	<ol> <li>Organic Chen</li> <li>Textbook of Oarun Bahl.</li> <li>Organic Chen</li> <li>Practical Organic Chen</li> <li>Vogel's text bear of the N.K. Vishnoi.</li> <li>Introduction the Pavia, Lampner</li> </ol>	book of Practical Cactical organic chemosomers of the Cache of the Cac	by B.S. Bahl &  Mann and Saunders.  Organic Chemistry  mistry by  ory techniques by			
Other References						





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	2	1	3	_	_	2	-	3	-	3
CO2	3	1	3	3	-	-	1	-	2	1	3
CO3	3	2	2	3	-	-	2	-	2	-	3
CO4	3	2	3	3	-	-	1	-	3	-	3
CO5	3	2	2	3	-	-	2	-	2	-	3
CO6	3	2	2	3	-	-	2	-	3	-	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





School:		SOP
<b>Programme:</b>		B. Pharm
Branch:		Semester: 2
1	Course Code	BP 209 0P
2	Course Title	Biochemistry Practical
3	Credits	2
4	Contact Hours (L-T-P)	0-0-4
	Course Type	Compulsory
5	Course Objective	Upon completion of course student shall able to  Understand the catalytic role of enzymes, importance of enzyme inhibitors in design of new drugs, therapeutic and diagnostic applications of enzymes. Understand the metabolism of nutrient molecules in physiological and pathological conditions.  Understand the genetic organization of mammalian genome and functions of DNA in the synthesis of RNAs and proteins.
6	Course Outcomes	CO1: Students will be able to understand the Qualitative analysis of carbohydrates CO2: Students will be able to understand the Quantitatative analysis of reducing sugars. CO3: Students will be able to analyze how to determine creatinine CO4: Students will be able to determine serum cholesterol CO5: Students will be able to compare amino acids by Paper Chromatographic Technique. CO6: Students will be able to apply the practical aspect and use of biochemistry.
7	Course Description	<ol> <li>Qualitative analysis of carbohydrates (Glucose, Fructose, Lactose, Maltose, Sucrose and starch)</li> <li>Identification tests for Proteins (albumin and Casein)</li> <li>Quantitative analysis of reducing sugars (DNSA method) and Proteins (Biuret method)</li> <li>Qualitative analysis of urine for abnormal constituents</li> <li>Determination of blood creatinine</li> <li>Determination of serum total cholesterol</li> <li>Preparation of buffer solution and measurement of pH</li> <li>Study of enzymatic hydrolysis of starch</li> <li>Determination of Salivary amylase activity</li> <li>Study the effect of Temperature on Salivary amylase activity.</li> <li>Study the effect of substrate concentration on salivary amylase activity.</li> </ol>
8	Outline syllabus	CO Mapping





T	1					
1	UNIT-I			CO1, CO6		
	a). Qualitative analy					
	Fructose, Lactose, N					
	b). Identification te	sts for Proteins (al	bumin and Casein)			
2	UNIT-II	UNIT-II				
	a). Quantitative ana	alysis of reducing s	ugars (DNSA	CO2, CO6		
	method) and Proteir	ns (Biuret method)				
	b). Qualitative analy	ysis of urine for ab	normal constituents			
3	UNIT-III					
	a). Determination of	f blood creatinine		CO3		
	b). Determination of	f blood sugar				
		-				
4						
	UNIT-IV			CO1		
	a). Determination of	a). Determination of serum total cholesterol b). Preparation of buffer solution and measurement of pH UNIT-V a). Study of enzymatic hydrolysis of starch				
	b). Preparation of b					
5						
	a). Study of enzyma					
	b). Determination of	CO4				
	Chromatographic To		•			
	Practical/Viva	•				
Mode of						
examination						
Weightage	Continuous	Sessional Exam	ESE			
Distribution	Mode					
	Assessment					
	05	10	35			
Text book/s*			a and S. Bhargavan.			
Text book/s			istry by David T.			
			isay by David 1.			
		Plummer. (3rd Edition) Practical Biochemistry for Medical students by Rajagopal				
	and Ramakrishna	•	adems by Rajagopai			
	Practical Biochemis		ev			
Other	Tractical Diocitcinis	bily by Harold Vall	ic y			
References						
References						





	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Pos											
Cos											
CO1	3	2	1	2	-	2	2	-	2	2	2
CO2	3	2	1	2	1	2	1	1	2	2	2
CO3	3	2	ı	2	-	2	1	1	1	2	2
CO4	3	2	ı	2	-	2	1	1	1	2	2
CO5	3	2	-	2	_	2	1	-	1	2	2
CO6	3	2	-	2	_	2	1	- 1	2	2	2

1-Slight (Low)

2-Moderate (Medium)

3-Substantial (High)



	SHARDA UNIVERSITY Beyond Boundaries www.sharda.ac.in	
•		-

Programme:         B. Pharm           Branch:         Semester: 2           1         Course Code         BP210 P           2         Course Title         Computer applications in Pharmacy- Practical           3         Credits         2           4         Contact Hours (L-T-P)         Course Type         Compulsory           5         Course Objective         Upon completion of the course the student shall be able to know the various types of application of computers in pharmacy know the various applications of databases know the various applications of databases in pharmacy           6         Course Outcomes         Upon completion of the course, the student shall be able to CO1: understand the Binary number system CO2: the web technologies	
1 Course Code BP210 P 2 Course Title Computer applications in Pharmacy- Practical 3 Credits 2 4 Contact Hours (L-T-P) Course Type Compulsory 5 Course Objective 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 Outcomes Upon completion of the course, the student shall be able to CO1: understand the Binary number system CO2: the web technologies	
2 Course Title Computer applications in Pharmacy- Practical 3 Credits 2 4 Contact Hours (L-T-P)  Course Type Compulsory  5 Course Objective 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 Outcomes Upon completion of the course, the student shall be able to CO1: understand the Binary number system CO2: the web technologies	
3 Credits 2 4 Contact Hours (L-T-P) Course Type Compulsory  5 Course Objective Upon completion of the course the student shall be able to know the various types of application of computers in pharmacy know the various applications of databases know the various applications of databases in pharmacy  6 Course Outcomes Upon completion of the course, the student shall be able to CO1: understand the Binary number system CO2: the web technologies	
4 Contact Hours (L-T-P)  Course Type Compulsory  5 Course Objective 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 Outcomes Upon completion of the course, the student shall be able to CO1: understand the Binary number system CO2: the web technologies	
Course Type Compulsory  5 Course Objective 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 Upon completion of the course, the student shall be able to CO1: understand the Binary number system CO2: the web technologies	
5 Course Objective 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 Outcomes Upon completion of the course, the student shall be able to CO1: understand the Binary number system CO2: the web technologies	
Objective  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 Outcomes  Upon completion of the course, the student shall be able to CO1: understand the Binary number system CO2: the web technologies	
Course Upon completion of the course, the student shall be able to CO1: understand the Binary number system CO2: the web technologies	
Outcomes CO1: understand the Binary number system CO2: the web technologies	
CO3: the application of computers in Pharmacy CO4: the Bioinformatics Databases, Concept of Bioinformatics CO5: Computers as data analysis in Preclinical development	20.00
CO6: Students will overall understand the use of computers in pharm	iacy
Course Description Design a questionnaire using a word processing package to information about a particular disease.  Create a HTML web page to show personal information.	
Retrieve the information of a drug and its adverse effects using onlin	
Creating mailing labels Using Label Wizard, generating label in MS	WORD
Create a database in MS Access to store the patient information required fields Using access	with the
Design a form in MS Access to view, add, delete and modify the record in the database	e patient
Generating report and printing the report from patient database	
Creating invoice table using – MS Access	
Exporting Tables, Queries, Forms and Reports to XML pages	





Outline syllabus		CO Mapping
I	UNIT-I  A. Number system: Binary number system, Decimal number system, Octal number system, Hexadecimal number systems, conversion decimal to binary, binary to decimal, octal to binary etc, binary addition, binary subtraction – One's complement ,Two's complement method, binary multiplication, binary division	CO1, CO6
2	UNIT-II A. Webtechnologies: Introduction to HTML, XML,CSS and Programmeming languages, introduction to web servers and Server Products Introduction to databases, MYSQL, MS ACCESS, Pharmacy Drug database	CO2, CO6
3	UNIT-III  A. Application of computers in Pharmacy – Drug information storage and retrieval, Pharmacokinetics, Mathematical model in Drug design, Hospital and Clinical Pharmacy, Electronic Prescribing and discharge (EP) systems, barcode medicine identification and automated dispensing of drugs, mobile technology and adherence monitoring Diagnostic System, Lab-diagnostic System, Patient Monitoring System, Pharma Information System	CO1, CO2,
1	UNIT-IV  A. <b>Bioinformatics:</b> Introduction, Objective of Bioinformatics, Bioinformatics Databases, Concept of Bioinformatics, Impact of Bioinformatics in Vaccine Discovery	CO1, CO4
5	UNIT-V A. Computers as data analysis in Preclinical development: Chromatographic dada analysis(CDS), Laboratory Information management System (LIMS) and Text Information Management System(TIMS)	CO1, CO3
Mod	de of mination	A. Computers as data analysis in Preclinical development: Chromatographic dada analysis(CDS), Laboratory Information management System (LIMS) and Text Information Management System(TIMS)  de of Practical/Viva





Weightage	<b>Continuous Mode</b>	Sessional Exam	ESE	
Distribution	Assessment			
	05	05	15	
Text book/s*	<ol> <li>Computer Application Lea and Febiger (215) 922-1330.</li> <li>Computer Application Computer Application Computer Application, USA</li> <li>Bioinformatics S.C.Rastogi-CBS 11 Darya Gani, March 2003, Application Computer C</li></ol>	cation in Pharmacy – r, 600 South Washi ication in Pharmace lean Ekins – lee, A John Wille A (Concept, Skills a S Publishers and Dis New Delhi – 110 002 Microso ion Development I Infopath – Cary Mia ia (P) Ltd., 443	William E.Fassett – ngton Square, USA, utical Research and y and Sons, INC., nd Applications) – tributors, 4596/1- A,	





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	1	-	-	1	3	1	-	1	2	3
CO2	3	2	-	2	1	3	1	-	1	2	3
CO3	3	2	1	2		3	1	-	1	3	3
CO4	3	2	1	2	2	3	2	-	1	2	3
CO5	3	2	2	3	2	3	2	-	1	3	3
CO6	3	2	1	2	2	3	1	_	1	3	3

-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





School:		SOP						
Prograi	nme:	B. Pharm						
Branch	•	Semester: 3						
1	Course Code	BP301 T						
	Course Title	Pharmaceutical organic chemistry-II- Theory						
3	Credits	4						
4	Contact Hours (L-T-P)	3-1-0						
	Course Type	Compulsory						
5	Course Objective	Upon completion of the course the student shall be able to						
		write the structure, name and the type of isomerism of the organi	ic compound					
		write the reaction, name the reaction and orientation of reactions						
		account for reactivity/stability of compounds,						
		prepare organic compounds						
6	Course Outcomes	CO1: The students will have the knowledge to identify, name, and write the structure of different aromatic compounds and their derivatives.						
		CO2: The students will be able to understand and explain the behind the naming reactions of different aromatic compounderivatives.						
		CO3: The students can apply the knowledge to prepare the aromaic compounds with different fuctional groups.	derivatives of					
		CO4: Students will analyze the chemical reactions, stabiliti compounds and properties of the compounds prepared by laboratory.	_					
		CO5: Students would evaluate bycomparing compounds prep with standard compounds by chemical and physical properties.	ared by them					
		CO6: Students would be able to evaluate Poly nuclear hyd comparing structures.	rocarbons. by					
7	Course Description	This subject deals with general methods of preparation and reactions of some organic compounds. Reactivity of organic compounds are also studied here. The syllabus emphasizes on mechanisms and orientation of reactions. Chemistry of fats and oils are also included in the syllabus.						
8	Outline syllabus  CO  Mappi							





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1	UNIT-I	CO1
	A. Benzene and its derivatives	
	Analytical, synthetic and other evidences in the derivation of	
	structure of benzene, Orbital picture, resonance in benzene,	
	aromatic characters, Huckel's rule	
	<b>B.</b> Reactions of benzene - nitration, sulphonation,	
	halogenation- reactivity, Friedelcrafts alkylation-	
	reactivity, limitations, Friedelcrafts acylation.	
	C. Substituents, effect of substituents on reactivity	
	and orientation of mono substituted benzene compounds	
	towards electrophilic substitution reaction	
	-	
2	Structure and uses of DDT, Saccharin, BHC and Chloramine	
2	UNIT-II  A. Phenols* - Acidity of phenols, effect of substituents	CO2
	¥ 1	CO2
	on acidity, qualitative tests, Structure and uses of phenol,	
	cresols, resorcinol, naphthols	
	<b>B.</b> Aromatic Amines* - Basicity of amines, effect of	
	substituents on basicity, and synthetic uses of aryl	
	diazonium salts	
	C. Aromatic Acids* – Acidity, effect of substituents on	
2	acidity and important reactions of benzoic acid.	
3	UNIT-III	CO3
	A. Fatty acids – reactions.	CO3
	Hydrolysis, Hydrogenation, Saponification and Rancidity of	
	oils, Drying oils.	
	B. Analytical constants – Acid value, Saponification	
	value, Ester value, Iodine value, Acetyl value, Reichert	
	Meissl (RM) value – significance and principle involved in	
	their determination.	
	UNIT-IV	CO4,CO6
	A. Polynuclear hydrocarbons:	Ź
	Synthesis, reactions	
	B. Structure and medicinal uses of Naphthalene,	
	Phenanthrene, Anthracene,	
	C. Diphenylmethane, Triphenylmethane and their	
	derivatives	





	UNIT-	V			CO5,			
	Α.	A. Cyclo alkanes*						
	Stabiliti	es – Baeyer's s	train theory, limitat	tion of Baeyer's strain				
				Sachse Mohr's theory				
			rings), reactions	of cyclopropane and				
		tane only						
Mode	of Theory							
examinat	ion							
Weightag	ge Contin	uous Mode	Sessional Exam	ESE				
Distribut	ion Assessi	nent						
	10		15	75				
Text boo	k/s*	. Organic Che	mistry by Morrison	n and Bovd				
		_	mistry by I.L. Fina	•				
		_		try by B.S. Bahl &				
		Arun Bahl.	C	<b>J</b>				
	4	. Organic Che	mistry by P.L. Son	i				
	5	5. Practical Organic Chemistry by Mann and Saunders.						
	6	. Vogel's text	book of Practical (	Organic Chemistry				
	7	. Advanced	Practical organ	ic chemistry by				
		N.K.Vishnoi						





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	1	-	-	1	3	1	-	2	2	3
CO2	3	2	-	2	1	3	1	-	2	2	3
CO3	3	2	1	2		3	1	-	2	3	3
CO4	3	2	1	2	2	3	2	-	1	2	3
CO5	3	2	2	3	2	3	2	-	2	3	3
CO6	3	2	1	2		3	1	-	1	3	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





School:		SOP					
Program	me:	B. Pharm					
Branch:		Semester: 3					
1	Course Code	BP302T					
2	Course Title	Physical Pharmaceutics I- Theory					
3	Credits	4					
4	Contact Hours (L-T-P)	3-1-0					
	Course Type	Compulsory					
5	Course Objective	Upon the completion of the course student shall be able to 1. Understand various physicochemical properties of drug molecules in the designing the dosage forms 2. Demonstrate use of physicochemical properties in the formulation development and evaluation of dosage forms. 3. Apply the concept of surface tension and surfactants in formulation and development.					
6	Course Outcomes	CO1: Students would be able to understand the concept of solubility, solutions, diffusion, CST, distribution and apply them in formulation, development and biological systems.  CO2: Students would be able to explain the basics of states of matter and physical properties of drugs and use them in pharmaceutical field.  CO3: Students would be able to apply the basics of surface and interfacial tension, surface active agents, HLB and adsorption in formulation and development of pharmaceutical systems.  CO4: Students would be able to describe Complexation, protein binding and relate it with drug action.  CO5: Students would be able to compare the methods of determination of pH and demonstrate the applications of buffered isotonic solutions in pharmaceutical and biological systems.  CO6: Students would be able to evaluate the Isoonicity of solutions.					
7	Course Description	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.					





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8	Outline syllabi	us	CO Mapping
		<ul> <li>UNIT-I</li> <li>A. Solubility of drugs: Solubility expressions, mechanisms of solute solvent interactions, ideal solubility parameters, solvation &amp; association, quantitative approach to the factors influencing solubility of drugs, diffusion principles in biological systems.</li> <li>B. Solubility of gas in liquids, solubility of liquids in liquids, (Binary solutions, ideal solutions) Raoult's law, real solutions.</li> <li>C. Partially miscible liquids, Critical solution temperature and applications. Distribution law, its limitations and applications</li> </ul>	CO1
	2	UNIT-II  A. States of Matter and properties of matter: State of matter, changes in the state of matter, latent heats, vapour pressure, sublimation critical point, eutectic mixtures, gases, aerosols inhalers, relative humidity, liquid complexes, liquid crystals, glassy states, solid-crystalline, amorphous & polymorphism.	CO2
		<ul> <li>B. Physicochemical properties of drug molecules: Refractive index, optical rotation, dielectric constant, dipole moment,</li> <li>C. dissociation constant, determinations and applications</li> </ul>	
	3	UNIT-III  A. 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.	CO3





4	Classification of of analysis, <b>B.</b> protein binding, <b>C.</b> Complexation and	<ul> <li>A. Complexation and protein binding: Introduction, Classification of Complexation, Applications, methods of analysis,</li> <li>B. protein binding,</li> <li>C. Complexation and drug action, crystalline structures of complexes and thermodynamic treatment of stability</li> </ul>					
5	<ul> <li>UNIT-V</li> <li>A. buffers and Iso pH determinatio</li> <li>B. B. applications capacity,</li> <li>C. buffers in pha buffered isotonic</li> </ul>	CO5, CO6					
Mode of examination	Theory						
Weightage	Continuous	Sessional Exam	ESE				
Distribution	Mode						
	Assessment						
	10 Marks	15	75				
Text book/s*	Physical Pharmacy I Experimental Pharm Tutorial Pharmacy I Stocklosam J. Pharm Philadelphia. Liberman H.A, La Dosage forms, MarcelDekkar Inc. Liberman H.A, Pharmaceutical Disperse systems, Marcel Dekkar Inc.						



Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	-	1	-	-		2	-	2	-	2
						2					
CO2	3	-	2	-	1	2	1	-	2	2	2
CO3	3	-	-	-	-	2	1	-	1	-	2
CO4	3	-	-	-	-	2	1	-	1	1	2
CO5	3	-	-	-	-	2	2	-	1	-	2
CO6	3	-	-	-	_	2	2	-	1	-	2

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





hool:	SOP
ogramme:	B. Pharm
anch:	Semester: 3
Course Code	BP303 T
Course Title	Pharmaceutical Microbiology- Theory
Credits	4
Contact Hours (L-T-P)	3-1-0
Type	Compulsory
Course Objective	Upon completion of the course the student shall be able to tell about the history, scope of microbiology and describe the structure, morphology and cultivation of microorganism.  Student shall identify the bacteria on the basis of various staining technique and importance of sterilization in microbiology.  Upon completion of the course the student shall understand the various methods for assessment of antibiotic, test for sterility for preparation.  Student shall analyze the source of contamination and their prevention in aseptic areas and importance of cell culture technique.
Outcomes	CO1: Students shall have knowledge about history of microbiology, its scope, branches, and application of various kind of microscopy.  CO2: Students shall be able the differentiate the types of bacteria on the basis of staining technique and biochemical test and with different type of microscopic technique and their method of validation  CO3: Students shall acquire complete knowledge of microorganism (viruses, fungi) like classification reproduction pattern, disinfection and antiseptic their evaluation methods and about sterility testing of various pharmaceutical products.  CO4: Students can apply their knowledge to design the aseptic area and standardization of antibiotic, biomolecules.  CO5: Students will be able to analyze the sources of contamination and their preventions in pharmaceutical products, and their application in pharmaceutical industry and research.  CO6: Students can apply their knowledge about cell cultures.
	anch: Course Code Course Title Credits Contact Hours (L-T-P) Course Type Course Objective  Course





7	Course Description  • Study of all categories of microorganisims especially for the production of a antibiotics, vaccines, vitamins enzymes etc				
8	Outline syllab	bus	CO Mapping		
		<ul> <li>UNIT-I</li> <li>A. Introduction, history of microbiology, its branches, scope and its importance.</li> <li>B. Introduction to Prokaryotes and Eukaryotes</li> <li>Study of ultra-structure and morphological classification of bacteria, nutritional requirements, raw materials used for culture media and physical parameters for growth, growth curve, isolation and preservation methods for pure cultures, cultivation of anaerobes, quantitative measurement of bacterial growth (total &amp; viable count).</li> <li>C. Study of different types of phase constrast microscopy, dark field microscopy and electron microscopy.</li> </ul>			
	2	<ul> <li>UNIT-II</li> <li>A. Identification of bacteria using staining techniques (simple, Gram's &amp; Acid fast staining) and biochemical tests (IMViC).</li> <li>B. Study of principle, procedure, merits, demerits and applications of physical, chemical gaseous, radiation and mechanical method of sterilization.</li> <li>C. Evaluation of the efficiency of sterilization methods.</li> </ul>	CO2		
	3	<ul> <li>UNIT-III</li> <li>A. Study of morphology, classification, reproduction/replicationand cultivation of Fungi and Viruses.</li> <li>Classification and mode of action of disinfectants</li> <li>Factors influencing disinfection, antiseptics and their evaluation. For bacteriostatic and bactericidal actions</li> <li>B. Evaluation of bactericidal &amp; Bacteriostatic.</li> <li>C. Sterility testing of products (solids, liquids, ophthalmic and other sterile products) according to IP, BP and USP.</li> </ul>	CO3		





4	<ul> <li>UNIT-IV</li> <li>A. Designing of aseptic area, I sources of contamination in clean area classification.</li> <li>B. Principles and methods of d standardization of antibiotics.</li> <li>Assessment of a new antibiotic.</li> </ul>	an aseptic area and met	hods of prevention, assay. Methods for	CO4			
5 UNIT-V A. Types of spoilage, factors affecting the microbial spoilage of pharmaceutical products. B. sources and types of microbial contaminants, assessment of microbial contamination and spoilage. C. Preservation of pharmaceutical products using antimicrobial agents, evaluation of microbial stability of formulations.							
Mode of examination	Theory						
Weightage Distribution	Continuous Mode Assessment	Sessional Exam	ESE				
	10 Marks	15	75				



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





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	1	3	3	-		1	-	1	2	3
						2					
CO2	3	2	2	2	-	1	-	-	-	1	3
CO3	3	-	2	-	-	2	2	-	2	-	3
CO4	3	-	2	-	-	1	-	-	-	2	3
CO5	3	2	3	3	-	2	2	-	2	2	3
CO6	3	_	2	-	-	1	2	-	2	-	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





School:		SOP				
Program	me:	B. Pharm				
Branch:		Semester: 3				
1	Course Code	BP 304 T				
2	Course Title	Pharmaceutical Engineering - Theory				
3	Credits	4				
4	Contact Hours (L-T-P)	3-1-0				
	Course Type	Compulsory				
5	Course Objective	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 optimum use of resources.				
		To appreciate the various preventive methods used for corrosion control in Pharmaceutical industries.				
6	Course Outcomes	CO1: Students will be able to describe about various unit operations used in pharmaceutical industries. and size separation and their applications in pharmaceutical field.				
		CO2: Students will be able to understand about basic concepts and importance of various heat transfer methods involved in pharmaceutical filed. CO3: Students will be able to illustrate about the concepts, equipments and pharmaceutical applications of drying and mixing.				
		CO4: Students will be able to distinguish between different types of equipments used in various unit operations such as filtration and centrifugation. CO5: Students will be able to predict about various materials used in pharmaceutical plant construction, types of corrosion and its prevention methods and basics of material handling system CO6: Students will be able to evaluate about various mechanisms of unit operations.				





7	Course Description	This course is designed to impart a fundamental knowledge art and science of various unit operations used in pharma industry.	
8	Outline syllabus		CO Mapping
	1	UNIT-I	CO1, CO6
		A. Flow of fluids: Types of manometers, Reynolds number and its significance, Bernoulli's theorem and its applications, Energy losses, Orifice meter, Venturimeter, Pitot tube and Rotometer.	
		B. Size Reduction: Objectives, Mechanisms & Laws governing size reduction, factors affecting size reduction, principles, construction, working, uses, merits and demerits of Hammer mill, ball mill, fluid energy mill, Edge runner mill & end runner mill.	
		C. 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, Air separator, Bag filter & elutriation tank.	
	2	UNIT-II	
		A. Heat Transfer: Objectives, applications & Heat transfer mechanisms. Fourier's law, Heat transfer by conduction, convection & radiation. Heat interchangers & heat exchangers.	CO2
		<ul> <li>B. 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 tube evaporator, climbing film evaporator, forced circulation evaporator, multiple effect evaporator&amp; Economy of multiple effect evaporator.</li> <li>C. Distillation: Basic Principles and</li> </ul>	
		methodology of simple distillation, fractional distillation, distillation under reduced pressure, steam distillation & molecular distillation	





3	UNIT-III		www.sharda.ac.in		
	applications of drying curve. promerits and demodryer, fluidized by the second of the	drying process, Equilibrium Moisi rinciples, construct erits of Tray drye bed dryer, vacuum g: Objectives, app g, Difference betweenism of solid mix mixing. Principles, ruction, Working, uble cone blender Sigma blade mixe		CO3	
4	UNIT-IV			CO4	
	<b>A.Filtration:</b> Objectives, applications, Theories & Factors influencing filtration, filter aids, filter medias. Principle, Construction, Working, Uses, Merits and demerits of plate & frame filter, filter leaf, rotary drum filter, Meta filter & Cartridge filter, membrane filters and Seidtz filter.				
	<b>B. Centrifugation:</b> Centrifugation, prince		ole & applications of		
	Perforated basket centrifuge, semi con	centrifuge, Nor	its and demerits of n-perforated basket & super centrifuge.		
5	UNIT-V	36.	of pharmaceutical	G0.5	
	A. Pharm plant construct Factors affection Pharmaceutical plant B. Theori there prevention C. Nonfernon metals, basic	CO5			
Mode of	Theory				
examination	Continue	Canalanal E	ECE		
Weightage Distribution	Continuous	Sessional Exam	ESE		
Distribution	Mode Assessment				
	10 Marks	15	75		
	10 11141110	10	, .	l l	





Text book/s*	1. Total destination to all anti-states
	Introduction to chemical engineering —     Walter L Badger & Julius Banchero, Latest edition.
	2. Solid phase extraction, Principles, techniques and applications by Nigel J.K. Simpson- Latest edition.
	3. Unit operation of chemical engineering – Mcabe Smith, Latest edition.
	4. Pharmaceutical engineering principles and practices – C.V.S Subrahmanyam et al., Latest edition.
	5. Remington practice of pharmacy- Martin, Latest edition.
	6. Theory and practice of industrial pharmacy by Lachmann., Latest edition.
	7. Physical pharmaceutics- C.V.S Subrahmanyam et al., Latest edition.
	8. Cooper and Gunn's Tutorial pharmacy, S.J. Carter, Latest edition.

Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	-	1	-	-	-	2	2	2	-	2
CO2	3	-	2	-	1	-	1	2	2	2	2
CO3	3	-	2	-	-	-	2	1	1	-	2
CO4	3	-	3	2	-	-	1	1	2	-	2
CO5	3	-	-	-	-	-	2	1	1	-	2
CO6	3	-	1	-	2	-	1	2	2	1	2

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





School:		SOP				
Progran	nme:	B. Pharm				
Branch:		Semester: 3				
1	Course Code	BP305 P  Pharmacautical organic chamistry II Practical				
	Course Title	Pharmaceutical organic chemistry II – Practical				
3	Credits	2				
4	Contact Hours (L-T-P)	0-0-4				
	Course Type	Compulsory				
5	Course Objective	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 environment improvement.  Acquire skills to help the concerned individuals in identifying and solving				
		environmental problems.  Strive to attain harmony with Nature.				
		Strive to accum narmony with rvacure.				
6	Course Outcomes	CO1: The students will have the knowledge to identify, name, and write the structure of different aromatic compounds and their derivatives.				
		CO2: The students will be able to understand and explain the mechanism behind the naming reactions of different aromatic compounds and their derivatives.				
		CO3: The students can apply the knowledge to prepare the derivatives of aromaic compounds with different fuctional groups.				
		CO4: Students will analyze the chemical reactions, stabilities of organic compounds and properties of the compounds prepared by them in the laboratory.				
		CO5: Students would evaluate bycomparing compounds prepared by them with standard compounds by chemical and physical properties.				
		CO6: The students will be able to plan about new derivatives based on the above knowledge.				





7	Course	1 Experiments involving laboratory techniques					
	Description	1. Experiments involving laboratory techniques					
		a. Recrystallization					
		b. Steam distillation					
		2. Determination of following oil values (inclustandardization of reagents)	ding				
		a. Acid value					
		b. Saponification value					
		c. Iodine value  3. Preparation Of Compounds					
		Benzanilide/Phenyl benzoate/Acetanilide     Aniline/Phenol/Aniline by acylation reaction.					
	• 2,4,6-Tribromo aniline/Para bromo acetanii Aniline/						
		Acetanilide by halogenation (Bromination) reaction	•				
		• 5-Nitro salicylic acid/Meta di nitro benzene Salicylic acid / Nitro benzene by nitration reaction.	from				
		Benzoic acid from Benzyl chloride by oxid reaction.	ation				
		Benzoic acid/ Salicylic acid from alkyl benzoate/ salicylate by hydrolysis reaction.	alkyl				
		• 1-Phenyl azo-2-napthol from Aniline by diazotiz and coupling reactions.	ation				
		<ul> <li>Benzil from Benzoin by oxidation reaction.</li> </ul>					
		<ul> <li>Dibenzal acetone from Benzaldehyde by Classics</li> <li>Schmidt reaction</li> </ul>	aison				
		Cinnammic acid from Benzaldehyde by Perkin reac	tion				
		P-Iodo benzoic acid from P-amino benzoic acid					
8	Outline syllabus						
	1	Mappin CO1,	g				
		Experiments involving laboratory techniques a). Recrystallization b). Steam distillation c). Derivatives of benzene					





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2	Deterination of followi	ng oil values		CO2, CO3			
	<ul><li>b). Saponification valu</li><li>c). Iodine value</li></ul>	b). Saponification value					
3	III Preparation of compa). Benzil b. Phenyl benzoate	CO3, CO6					
	Benzoic acid Oxalic acid	,					
	Analytical constants – value, Iodine value, Ac	c. Rancidity of oils, Drying oils.  Analytical constants – Acid value, Saponification value, Ester value, Iodine value, Acetyl value, Reichert Meissl (RM) value – significance and principle involved in their determination.					
Mode of examination	Theory						
Weightage Distribution	Continuous Mode Assessment	Sessional Exam	ESE				
	0 5	10	35				
Text book/s*	<ol> <li>Organic Che</li> <li>Organic Che</li> <li>Textbook of Arun Bahl.</li> <li>Organic Che</li> <li>Practical Org</li> <li>Vogel's text</li> <li>Advanced N.K.Vishnoi</li> </ol>						





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	1	-	-	1	3	1	-	1	2	3
CO2	3	2	-	2	2	3	2	-	2	2	3
CO3	3	2	2	2		3	1	-	1	3	3
CO4	3	2	-	-	-	-	2	-	3	1	-
CO5	3	2	1	2	2	3	2	-	2	2	3
CO6	3	2	-	-	-	-	2	-	3	-	-

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





School:		SOP				
Program	me:	B. Pharm				
Branch:		Semester: 3				
1	Course Code	BP306 P				
2	Course Title	Physical pharmaceutics I-Practical				
3	Credits	4				
4	Contact Hours (L-T-P)	0-0-4				
	Course Type	Compulsory				
5	Course Objective	Upon the completion of the course student shall be able to Understand various physicochemical properties of drug molecules in the designing the dosage forms Demonstrate use of physicochemical properties in the formulation development and evaluation of dosage forms.				
6	Course Outcomes	CO1: The students would be able to describe the various methods of determination of physicochemical properties of drugs and pharmaceuticals. CO2: The students would be able to demonstrate methods for determination of HLB value and Critical Micelle concentration of surfactants.  CO3: The students would be able to calculate the value of stability constants in complexation by various methods.  CO4: The students would be able to compare various methods of determ ination of stability constants  CO5: The students would be able to determine the effect of addition of salt CST and to determine adsorption constants.  CO6: The students would be able to determine of Freundlich and Langmuir constants.				
7	Course Description	Determination of physicochemical properties of drugs and pharmaceuticals and determination of stability constants, adsorption constants, HLB and CMC values.				
8	Outline syllabus	CO Mapping				





		To determine vario	ous physicochemic	cal properties of	
		drugs and Pharma a). Determination of b). Determination of Henderson Hasselba c). Determination of in benzene and wate d). Determination of CCl4 and water e). Determination of drop count and drop	recuticals Solubility of drug f pKa value by Hal alch equation. Partition co- efficer f Partition co- efficer surface tensionof	at room temperature f Neutralization/ ient of benzoic acid ient of Iodine in	
2		To Determine impo a). Determination of saponification metho b). Determination of surfactants	CO1, CO2		
3		To determine stabi various methods a). Determination of ratio of PABA-Caffe b). Determination of ratio of Cupric-Glyc	CO2, CO3		
Mod exam	de of mination	To study the effect determine adsorptian. Determination of using phenol-water sub). Determination of using activated charman Practical/Viva	CO1, CO6		
Wei	ghtage ribution	Continuous Mode Assessment 0 5 Marks	Sessional Exam	ESE 35	





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Text book/s*	Physical Pharmacy by Alfred Martin	
	Experimental Pharmaceutics by Eugene, Parott.	
	Tutorial Pharmacy by Cooper and Gunn.	
	Stocklosam J. Pharmaceutical Calculations, Lea &Febiger,	
	Philadelphia.	
	Liberman H.A, Lachman C., Pharmaceutical Dosage	
	forms, Tablets, Volume-1 to 3, MarcelDekkar Inc.	
	Liberman H.A, Lachman C, Pharmaceutical Dosage forms.	
	Disperse systems, volume 1, 2, 3. Marcel Dekkar Inc.	
	Physical Pharmaceutics by Ramasamy C and ManavalanR.	
	Laboratory Manual of Physical Pharmaceutics, C.V.S.	
	Subramanyam, J. Thimma settee	
	Physical Pharmaceutics by C.V.S. Subramanyam	
	Test book of Physical Phramacy, by Gaurav Jain & Roop	
	K. Khar	
Other		
References		

Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	-	2	-	-		2	1	2	-	2
						2					
CO2	3	-	1	-	2	2	2	1	2	2	2
CO3	3	-	-	-	1	2	1	2	1	1	2
CO4	3	-	-	-	-	2	3	2	1	-	2
CO5	3	-	-	-	-	2	1	2	2	-	2
CO6	3	-	-	-	-	2	1	2	1	-	2

- 1-Slight (Low)
- 2-Moderate (Medium)
- 3-Substantial (High)





Sc	hool:	SOP
-	ogramme:	B. Pharm
	anch:	Semester: 3
1	Course	BP307 P
	Code	
2	Course	Pharmaceutical microbiology Practical
	Title	
3	Credits	2
4	Contact	0-0-4
	Hours	
	(L-T-P)	
	Course	Compulsory
5	Type Course	
	Objective	Upon completion of the course the student shall be able to tell about the history, scope of microbiology and describe the structure, morphology and cultivation of microorganism.  Student shall identify the bacteria on the basis of various staining technique and importance of sterilization in microbiology.  Upon completion of the course the student shall understand the various methods for assessment of antibiotic, test for sterility for preparation.  Student shall analyze the source of contamination and their prevention in aseptic areas and importance of cell culture technique.
6	Course Outcomes	CO1: Students shall have knowledge about the various equipment used in experimental microbiology and understand the principle and working of these instruments.  CO2: Students shall be able to understand the importance of sterilization in microbiology and apply this knowledge for the preparation of various media.  CO3: Students shall acquire complete knowledge of isolation procedure of microorganism (viruses, fungi) and will be able to differentiate microorganism on the basis of various staining technique  CO4: Students can apply their knowledge for the standardization of antibiotics.  CO5: Students can apply their knowledge for identification of bacteria.
		CO6: Students can apply their knowledge for nutrient and slab culture.
7	Course Description	To emphasize on definition, types, classification, principles/mechanisms, applications, examples and differences
8	Outline sylla	bus CO Mapping





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1	a). To study various equipme b). To perform the sterilization			CO1, CO2				
2		a). Preparation of sterile nutrient broth b). Preparation of sterile nutrient agar media						
3	a). Study of environmental microflora of various region							
4	a). Standardization of antibiotic by cup and plate method Identification of bacteria by gram staining technique identification of bacteria by acid fast staining technique Preparation of nutrient slant and stab culture							
Mode of examination	Practical/Viva							
Weightage Distribution	tribution Assessment							
	05	10	35					
book/s*	Blackwell Scientific publication Prescott and Dunn.,	W.B. Hugo and A.D. Russel: Pharmaceutical Microbiology, Blackwell Scientific publications, Oxford London.  Prescott and Dunn., Industrial Microbiology, 4 th edition, CBS Publishers & Distributors, Delhi.						
	Publishers & Distributors, Delhi.  Pelczar, Chan Kreig, Microbiology, Tata McGraw Hill edn.  Malcolm Harris, Balliere Tindall and Cox: Pharmaceutical Microbiology.  Rose: Industrial Microbiology.  Probisher, Hinsdill et al: Fundamentals of Microbiology, 9th ed.  Japan  Cooper and Gunn's: Tutorial Pharmacy, CBS Publisher and Distribution.  Peppler: Microbial Technology.  I.P., B.P., U.S.P latest editions.  Ananthnarayan: Text Book of Microbiology, Orient-Longman, Chennai  Edward: Fundamentals of Microbiology, Vallabh Prakashan, Delhi  Bergeys manual of systematic bacteriology, Williams and Wilkins-A Waverly company							
Other References								





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	1	3	3	3	2	2	1	2	1	3
CO2	3	2	2	2	2	2	2	3	2	2	3
CO3	3	3	2	-	2	2	2	3	2	2	3
CO4	3	2	2	-	2	1	2	3	2	2	3
CO5	3	2	3	3	3	2	1	1	2	1	3
CO6	3	2	2	2	2	1	2	3	2	2	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





School:		SOP
Program	me:	B. Pharm
Branch:		Semester: 3
1	Course Code	BP 308 P
2	Course Title	Pharmaceutical Engineering Practical
3	Credits	4
4	Contact Hours (L-T-P)	0-0-4
	Course Type	Compulsory
5	Course Objective	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 optimum use of resources.
		To appreciate the various preventive methods used for corrosion control in Pharmaceutical industries.
6	Course Outcomes	CO1: Student shall be able to Understand the different factors effecting rate of filtration, evaporation and overall heat transfer coefficient etc.
		CO2: Students shall be able to predict humidity of air, effect of time on crystallization rate and laws of size reduction.
		CO3: Students shall be able to calculate uniformity index of given sample, efficiency of steam distillation and construct various size frequency curves, drying curves etc. CO4: Students shall be able to evaluate size distribution of tablet granulations.
		CO5: Students shall be able to calculate time of crystallization
		CO6: Students shall be able to study the effect of time on rate of crystallization.





_											
7	Course Description	science of various u	nit operations used	ndamental knowledge I in pharmaceutical in pharmaceutical unit is	dustryand their						
8	Outline syllabus				CO Mapping						
	1	<u> </u>			Contrapping						
		transfer coefficient efficiency of steam b). Students would calcium carbonate a	<ul><li>a). Students would be able to determine the overall heat transfer coefficient by heat exchanger and calculate the efficiency of steam distillation.</li><li>b). Students would be able to construct drying curves (for calcium carbonate and starch) and determine moisture content and loss on drying.</li></ul>								
	2	<ul><li>a). Students would I From wet and dry b method.</li><li>b). Students would tablet granulations I</li></ul>	CO1, CO3								
	3	a). Students would be reduction using ball b). Students would of Evaporation and c). Students would be major equipment us	mill using Ball M be able to relate fact Filtration. be able to understan	ill etors affecting Rate	CO2 CO3						
	4	the Rate of Crystall b). Students would	a). Students would be able to study the effect of time on the Rate of Crystallization b). Students would be able to calculate the uniformity Index for given sample by using Double Cone Blender								
	Mode of examination	Practical/Viva									
	Weightage Distribution	Continuous Mode Assessment	Sessional Exam	ESE							
		0 5 Marks	10	35							





Text book/s*	1. Introduction to chemical engineering – Walter Badger & Julius Banchero, Latest edition.	L
	<ol><li>Solid phase extraction, Principles, techniques an applications by Nigel J.K. Simpson- Latest edition.</li></ol>	d
	3. Unit operation of chemical engineering – Mcabe Smith Latest edition.	1,
	4. Pharmaceutical engineering principles and practices C.V.S Subrahmanyam et al., Latest edition.	_
	5. Remington practice of pharmacy- Martin, Late edition.	st
	<ol><li>Theory and practice of industrial pharmacy b Lachmann., Latest edition.</li></ol>	у
	<ol> <li>Physical pharmaceutics- C.V.S Subrahmanyam et al Latest edition.</li> </ol>	.,
	8. Cooper and Gunn's Tutorial pharmacy, S.J. Carte Latest edition.	r,

Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	-	1	-	-		2	_	2	-	2
						2					
CO2	3	-	2	-	2	2	1	-	2	2	2
CO3	3	-	-	-	-	2	1	-	1	-	2
CO4	3	-	_	_	_	2	2	_	1	2	2
CO5	3	-	-	-	-	2	2	-	2	-	2
CO6	3	-	-	-	-	2	2	-	1	2	2

Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





So	chool:	SOP	
_	rogramme:	B.Pharm	
	ranch:	Semester: IV	
1	Course	BP401T	
	Code		
2	Course	Pharmaceutical Organic Chemistry III - Theory	
	Title		
3	Credits	4	
4	Contact	3-1-0	
	Hours		
	(L-T-P)		
	Course	Compulsory	
	Type		
5	Course	Upon completion of the course the student shall be able to	
	Objective	1. 1. understand the methods of preparation and prop	erties of organic
		compounds	
		2. explain the stereo chemical aspects of organic compo	ounds and stereo
		chemical reactions	
		3. 3. know the medicinal uses and other applicat	ions of organic
_	Carrage	compounds	on to outlool and
6	Course Outcomes	CO1: Students shall be able to understand the configuration	-
	Outcomes	geometrical isomers. They also get the knowledge of propertical and geometrical isomers and diasteriomers.	es of enantiomers
		CO2: Students shall acquire the knowledge of separation of	different isomers
		and on the basis of this knowledge students can separate the	
		form.	desired isometic
		CO3: Students shall be able to do nomenclature of heteroc	velic compounds
		and draw the structure of heterocyclic compounds.	J
		CO4: students shall gain the knowledge of various heterocyc	lic compounds in
		terms of their synthesis, chemical reactions and their application	
		CO5: The students will be able to understand and explain	n the mechanism
		behind various naming reactions and acquire the know	wledge of their
		applications in preparation of various drugs and intermediates.	
		CO6: Students shall be able to study the SAR of compounds	•
7	Course Description	This subject imparts knowledge on stereo-chemical asp	ects of organic
		compounds and organic reactions, important named reaction	ons, chemistry of
		important hetero cyclic compounds. It also emphasizes on me	edicinal and other
		uses of organic compounds.	
8	Outline syllab	ous	CO Mapping





1	UNIT-I	CO1
	A. Stereo isomerism	
	Optical isomerism —Optical activity, enantiomerism, diastereoisomerism, meso compounds Elements of symmetry, chiral and achiral molecules  B. DL system of nomenclature of optical isomers, sequence rules, RS system of nomenclature of optical isomersReactions of chiral molecules	
	Racemic modification and resolution of racemic mixture.	
	C. Asymmetric synthesis: partial and absolute	
2	UNIT-II	CO2, CO6
	A. Geometrical isomerism	
	Nomenclature of geometrical isomers (Cis Trans, EZ, Syn Anti systems)	
	Methods of determination of configuration of geometrical isomers.	
	<b>B.</b> Conformational isomerism in Ethane, n-Butane and Cyclohexane.	
	Stereo isomerism in biphenyl compounds (Atropisomerism) and conditions for optical activity.  Stereospecific and stereoselective reactions	
3	Unit III	CO3
	A. Heterocyclic compounds:	
	Nomenclature and classification	
	B. Synthesis, reactions and medicinal uses of following	
	compounds/derivatives Pyrrole, Furan, and Thiophene	
	C. Relative aromaticity and reactivity of Pyrrole, Furan and Thiophene	





4 UNIT-IV A. Synthesis, reactions and medicinal uses of following compounds/derivatives Pyrazole, B. Imidazole, Oxazole and Thiazole. Pyridine, Quinoline, Isoquinoline, Acridine and Indole. Basicity of pyridine Synthesis C. medicinal uses of Pyrimidine, Purine, azepines and their derivatives  5 UNIT-V CO5 A. Reactions of synthetic importance	
compounds/derivatives Pyrazole,  B. Imidazole, Oxazole and Thiazole. Pyridine, Quinoline, Isoquinoline, Acridine and Indole. Basicity of pyridine Synthesis  C. medicinal uses of Pyrimidine, Purine, azepines and their derivatives  5 UNIT-V  CO5	
B. Imidazole, Oxazole and Thiazole. Pyridine, Quinoline, Isoquinoline, Acridine and Indole. Basicity of pyridine Synthesis C. medicinal uses of Pyrimidine, Purine, azepines and their derivatives  5 UNIT-V  CO5	
Isoquinoline, Acridine and Indole. Basicity of pyridine Synthesis C. medicinal uses of Pyrimidine, Purine, azepines and their derivatives  5 UNIT-V  CO5	
Synthesis C. medicinal uses of Pyrimidine, Purine, azepines and their derivatives  5 UNIT-V  CO5	
C. medicinal uses of Pyrimidine, Purine, azepines and their derivatives  5 UNIT-V CO5	
derivatives  5 UNIT-V CO5	
5 UNIT-V CO5	
UNIT-V	
A. Reactions of synthetic importance	
Metal hydride reduction (NaBH ₄ and LiAlH ₄ ), Clemmensen reduction, Birch reduction, Wolff Kishner reduction.	
<b>B.</b> Oppenauer-oxidation and Dakin reaction.	
C. Beckmanns rearrangement and Schmidt rearrangement.	
Claisen-Schmidt condensation	
Mode of Theory examinat ion	
Weighta Continuous Mode Sessional Exam ESE	
ge Assessment	
Distribut ion 10 Marks 15 75	
Text book/s*  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	
Other Referenc	



## COURSE ARTICULATION MATRIX

Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	2	2	3	3	1	-	2	2	3
CO2	3	3	-	1	3	3	2	-	3	2	3
CO3	3	3	2	2	3	2	2	-	3	3	3
CO4	3	2	-	1	3	3	2	-	2	2	3
CO5	3	3	1	2	3	3	1	-	2	1	3
CO6	3	3	2	-	3	2	-	-	3	3	3

- 1. Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





School:		SOP					
Progr	amme:	B.Pharm					
Branc		Semester: IV					
1	Course Code	BP402T					
2	Course Title	Medicinal chemistry I - Theory					
3	Credits	4					
4	Contact Hours (L-T-P)	3-1-0					
	Course Type	Compulsory					
5	Course Objective	Upon completion of the course the student shall be able to 1. understand the chemistry of drugs with respect to their pharactivity 2. understand the drug metabolic pathways, adverse effect and value of drugs know the Structural Activity Relationship (SAR) of different cl write the chemical synthesis of some drugs	therapeutic				
6	Course Outcomes	CO1: The students will be able to apply knowledge to identify classify the different categories of drugs with respect to their pharactivities.  CO2: The students will be able to understand and explain the structure relationship, drug metabolic pathways, adverse effects and their activity of different categories of drugs.  CO3: The students will be able to apply the knowledge to echemical synthesis of some drugs.  CO4: The students will be able to analyze chemical reactions, a compounds and properties of the compounds prepared by the laboratory.  CO5: The students would be able to illustrate different drug Nervous system.  CO6: The students would be able to evaluate various transmitters.	eture activity therapeutic construct the stabilities of hem in the				
7	Course Description	This subject is designed to impart fundamental knowledge on the chemistry and therapeutic value of drugs. The subject emphasizes activity relationships of drugs, importance of physicochemical primetabolism of drugs. The syllabus also emphasizes on chemical important drugs under each class.	on structure operties and				
8	Outline syllabu	IS STATE OF THE PROPERTY OF TH	CO Mapping				





1	UNIT- I	CO1
	UNII-I	
	A. Introduction to Medicinal Chemistry History and development of medicinal chemistry Physicochemical properties in relation to biological action Ionization, Solubility, Partition Coefficient, Hydrogen bonding, Protein binding, Chelation, Bioisosterism, Optical and Geometrical isomerism.	
	B. Drug metabolism	
	a). Drug metabolism principles- Phase I and Phase II.	
	b). Factors affecting drug metabolism including stereo chemical aspects.	
2	UNIT- II	CO2, CO6
	A. Drugs acting on Autonomic Nervous System Adrenergic Neurotransmitters:	
	a). Biosynthesis and catabolism of catecholamine.	
	b). Adrenergic receptors (Alpha & Beta) and their distribution.	
	B. Sympathomimetic agents: SAR of Sympathomimetic agents	
	a). Direct acting: Nor-epinephrine, Epinephrine, Phenylephrine*, Dopamine, Methyldopa, Clonidine, Dobutamine, Isoproterenol, Terbutaline, Salbutamol*, Bitolterol, Naphazoline, Oxymetazoline and Xylometazoline.	
	Indirect acting agents: Hydroxyamphetamine,	
	Agents with mixed mechanism: Ephedrine, Metaraminol.	
	C. Adrenergic Antagonists:	
	Alpha adrenergic blockers: Tolazoline*, Phentolamine, Phenoxybenzamine, Prazosin, Dihydroergotamine, Methysergide.	
	<b>D. Beta adrenergic blockers:</b> SAR of beta blockers, Propranolol*, Metibranolol, Atenolol, Betazolol, Bisoprolol, Esmolol, Metoprolol, Labetolol, Carvedilol.	





3 CO3 **UNIT-III** A. Cholinergic neurotransmitters: a). Biosynthesis and catabolism of acetylcholine. b). Cholinergic receptors (Muscarinic & Nicotinic) and their distribution. c). Parasympathomimetic agents: SAR of Parasympathomimetic agents B. Direct acting agents: Acetylcholine, Carbachol*, Bethanechol, Methacholine, Pilocarpine. C. Indirect acting/ Cholinesterase inhibitors (Reversible & Irreversible): Physostigmine, Neostigmine*, Pyridostigmine, Edrophonium chloride, Tacrine hydrochloride, Ambenonium chloride, Isofluorphate, Echothiophate iodide, Parathione, Malathion. Cholinesterase reactivator: Pralidoxime chloride. Cholinergic Blocking agents: SAR of cholinolytic agents Solanaceous alkaloids and analogues: Atropine Hyoscyamine sulphate, Scopolamine hydrobromide, Homatropine hydrobromide, Ipratropium bromide*. cholinergic Tropicamide, **Synthetic** blocking agents: Cyclopentolate hydrochloride, Clidinium bromide, Dicyclomine hydrochloride*, Glycopyrrolate, Methantheline bromide,

Propantheline bromide, Benztropine mesylate, Orphenadrine citrate,

Procyclidine

Isopropamide iodide,

hydrochloride*,

Ethopropazine

hydrochloride,

chloride,

Biperidine

Tridihexethyl

hydrochloride.





4 UNIT- IV

CO₄

### **Drugs acting on Central Nervous System**

### A. Sedatives and Hypnotics:

**Benzodiazepines:** SAR of Benzodiazepines, Phenobarbital, Mephobarbital, Amobarbital, Butabarbital, Pentobarbital, Secobarbital

#### **B.** Miscelleneous:

Amides & imides: Glutethmide.

Alcohol & their carbamate derivatives: Meprobomate, Ethchlorvynol. Aldehyde & their derivatives: Triclofos sodium, Paraldehyde.

#### C. Antipsychotics

**Phenothiazeines:** SAR of Phenothiazeines - Promazine hydrochloride, Chlorpromazine hydrochloride*, Triflupromazine, Thioridazine hydrochloride, Piperacetazine hydrochloride, Prochlorperazine maleate, Trifluoperazine hydrochloride.

**Ring Analogues of Phenothiazeines:** Chlorprothixene, Thiothixene, Loxapine succinate, Clozapine.

Fluro buterophenones: Haloperidol, Droperidol, Risperidone.

Beta amino ketones: Molindone hydrochloride.

Benzamides: Sulpieride.

Chlordiazepoxide, Diazepam*, Oxazepam, Chlorazepate, Lorazepam, Alprazolam, Zolpidem

Barbiturtes: SAR of barbiturates, Barbital*,





5	UNIT – V			CO5
	Drugs acting on Central N	ervous System		
	A. General anesthetics:			
	<b>a). Inhalation anesthetics:</b> Sevoflurane, Isoflurane, Des		hoxyflurane, Enflurane,	
	<b>b). Ultra short acting</b> Thiamylal sodium, Thiopen		Methohexital sodium*,	
	c). Dissociative anesthetics	: Ketamine hydro	chloride.*	
	B. Narcotic and non-narc	otic analgesics		
	a). Morphine and related Morphine sulphate, Codeine hydrochloride, Diphenos hydrochloride, Fentanyl Propoxyphene hydrochloride	e, Meperidine hy xylate hydroc citrate*, Metha	drochloride, Anilerdine hloride, Loperamide adone hydrochloride*,	
	b). Narcotic antagonists: tartarate, Naloxone hydrochi			
	c). Anti-inflammatory Mefenamic acid*, Mecinological Tolmetin, Zomepriac, Diclo Piroxicam, Phenacetin Phenylbutazone.	lofenamate, Inc ofenac, Ketorolac,	lomethacin, Sulindac, Ibuprofen*, Naproxen,	
	C.Actions of synthetic import	ance		
	a). Metal hydride reduction, veduction, Birch reduction,	*	* *	
	b). Oppenauer-oxidation and	d Dakin reaction.		
	c). Beckmanns rearrangeme	ent and Schmidt	rearrangement. Claisen-	
	Schmidt condensation			
Mode of examinati on	Theory			
Weightag	Continuous Mode	Sessional	ESE	
e Distributi	Assessment	Exam		
Distributi on	10 Marks	15	75	





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



### **COURSE ARTICULATION MATRIX**

Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	1	2	3	3	2	-	2	2	3
CO2	3	3	-	2	3	3	1	-	3	1	3
CO3	3	3	2	2	3	2	2	-	3	3	3
CO4	3	2	-	2	3	3	2	-	2	1	3
CO5	3	3	-	1	3	3	1	_	3	2	3
CO6	3	3	2	2	3	2	1	-	3	3	3

Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sc	hool:	SOP
	ogramme:	B. Pharm
	anch:	Semester: IV
1	Course Code	BP403T
2	Course Title	Physical Pharmaceutics II - Theory
3	Credits	4
4	Contact	3-1-0
	Hours	
	(L-T-P)	
	Course Type	Compulsory
5	Course	Upon the completion of the course student shall be able to
	Objective	1. Understand various physicochemical properties of drug molecules in the
		designing the dosage form  2. Know the principles of chemical kinetics & to use them in assigning expiry
		2. Know the principles of chemical kinetics & to use them in assigning expiry date for Formulation
		3. Demonstrate use of physicochemical properties in evaluation of dosage forms.
		Appreciate physicochemical properties of drug molecules in formulation research
		and Development
6	Course	CO1: Students would be able to understand the concept of reaction kinetics,
	Outcomes	degradation pathways, factor effects stability of drugs.
		CO2: Students would be able to understand flow of liquid, law of flow,
		determination of viscosity of liquid by viscometer.
		CO3: Students would be able to apply the basics of surface and interfacial tension,
		surface active agents.
		CO4: Students would be able to describe properties of powder like particle size and
		distribution, determining particle size by different methods.  CO5: Students would be able to evaluate about the colloidal dispersion, role of
		particle size and shape in colloidal dispersion.
		CO6: Students would be able to illustrate the Stabilization of medicinal agents.
7	Course	200. Statement would be use to madrate the Statement of medicinal agents.
	Description	The course deals with the various physica and physicochemical properties, and
	r ·	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.
8	Outline syllabu	CO Mapping





1	UNIT-I	CO1
	A. Colloidal dispersions: Classification of dispersed systems & their general characteristics, size & shapes of colloidal particles, classification of colloids & comparative account of their general properties. Optical, kinetic	
	<b>B.</b> B. electrical properties.	
	C. Effect of electrolytes, coacervation, peptization& protective action.	
2		CO2
2	UNIT-II	CO2
	<b>A. Rheology:</b> Newtonian systems, law of flow, kinematic viscosity, effect of temperature, non-Newtonian systems, pseudoplastic, dilatant, plastic, thixotropy, thixotropy in formulation, determination of viscosity, capillary, falling Sphere, rotational viscometers	
	B. Deformation of solids: Plastic and elastic deformation,	
	C. Heckel equation, Stress, Strain, Elastic Modulus	
3	UNIT-III	CO3
	<b>A. Coarse dispersion:</b> Suspension, interfacial properties of suspended particles, settling in suspensions, formulation of flocculated and deflocculated suspensions.	
	<b>B.</b> B. Emulsions and theories of emulsification, microemulsion and multiple emulsions;	
	<b>C.</b> Stability of emulsions, preservation of emulsions, rheological properties of emulsions and emulsion formulation by HLB method.	





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4	UNIT IV			CO4
	C04			
5	UNIT-V			CO5, CO6
	<ul><li>A. Drug stabi second ordereaction ordereaction ordered desolvent, ion acid base ca</li><li>B. Stabilization like hydroly</li></ul>			
	C. Accelerated pharmaceut prevention			
Mode of examination	Theory			
Weightage Distribution	Continuous S Mode Assessment	Sessional Exam	ESE	
	10 Marks	15	75	
Text book/s*	<ol> <li>Physical Pharm</li> <li>Experimental p</li> </ol>			
	3. Tutorial pharm	nacy by Cooper and Gunn		
	4. Stocklosam J. Philadelphia.			
	5. Liberman H.A. Tablets, Volum			
		A, Lachman C, Pharm ms, volume 1, 2, 3. Marce	aceutical dosage forms. el Dekkar Inc.	
	7. Physical Pharm	naceutics by Ramasamy C	C, and Manavalan R.	



## COURSE ARTICULATION MATRIX

Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	2	2	3	3	2	2	2	1	3
CO2	3	3	-	1	3	3	2	1	3	2	3
CO3	3	3	2	2	3	2	1	3	3	3	3
CO4	3	2	-	2	3	3	2	2	2	2	3
CO5	3	3	2	2	3	2	1	3	3	3	3
CO6	3	2	-	1	3	3	2	2	2	2	3

Slight (Low) 2-Moderate (Medium) 3-Substantial (High)

		* SHARDA
School:		SOP UNIVERSITY
	ogramme:	B. Pharm
	anch:	Semester: IV
1	Course Code	BP404T
2	Course Title	Pharmacology I - Theory
3	Credits	4
4	Contact Hours (L-T-P)	3-1-0
	Course Type	Compulsory
5	Course Objective	Upon completion of this course the student should be able to
	J	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
6	Course Outcomes	CO1: Students will able to understand the pharmacological actions of different categories of drugs.  CO2: Students will able to explain the mechanism of drug action at organ system/sub cellular/macromolecular levels.  CO3: Students will able to apply the basic pharmacological knowledge in the prevention and treatment of various diseases.  CO4: Students will able to illustrate the effect of drugs on animals by simulated experiments.  CO5: Students will able to apply the correlation of pharmacology with other biomedical Sciences.  CO5: Students will able to evaluate the mechanisms and action of drugs under CNS CO6: Students will able to evaluate the mechanisms and action of drugs.
7	Course Descriptio n	The main purpose of the subject is to understand what drugs do to the living organisms and how their effects can be applied to therapeutics. The subject covers the information about the drugs like, mechanism of action, physiological and biochemical effects (pharmacodynamics) as well as absorption, distribution, metabolism and excretion (pharmacokinetics) along with the adverse effects, clinical uses, interactions, doses, contraindications and routes of administration of different classes of drugs.
8	Outline syll	abus CO Mapping





	www.sharda.ac.in	GO.1
1	UNIT-I	CO1
	A. General Pharmacology	
	<ul> <li>a. Introduction to Pharmacology- Definition, historical landmarks and scope of pharmacology, nature and source of drugs, essential drugs concept and routes of drug administration, receptors, addiction, tolerance, dependence, tachyphylaxis, idiosyncrasy, allergy.</li> <li>b. Pharmacokinetics- Membrane transport, absorption, distribution, metabolism and excretion of drugs .Enzyme induction, enzyme inhibition, kinetics of elimination</li> <li>c.Agonists, antagonists( competitive and non competitive), spare</li> </ul>	
	receptors, addiction, tolerance, dependence, tachyphylaxis,	
2	UNIT-II	CO2, CO6
	A. General Pharmacology 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 binding receptor and receptors that regulate transcription factors, dose response relationship, therapeutic index, combined effects of drugs and factors modifying drug action. a. Adverse drug reactions. b. Drug interactions (pharmacokinetic and pharmacodynamic)  Drug discovery and clinical evaluation of new drugs -Drug discovery phase, preclinical evaluation phase,	





UNIT-III Pharmacology of drugs acting on peripheral nervous system  A. 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).  B. Local anesthetic agents.  C. Drugs used in myasthenia gravis and glaucoma  UNIT-IV Pharmacology of drugs acting on central nervous system  a. Neurohumoral transmission in the C.N.S. special emphasis on importance of various neurotransmitters like with GABA, Glutamate, Glycine, serotonin, dopamine.  b. General anesthetics and pre-anesthetics.  c. Sedatives, hypnotics and centrally acting muscle relaxants. Anti-epileptics, Alcohols and disulfiram  5  UNIT-V Pharmacology of drugs acting on central nervous system  a. Psychopharmacological agents: Antipsychotics, antidepressants, anti-anxiety agents, anti-manics and hallucinogens.  b. Drugs used in Parkinsons disease and Alzheimer's disease.  c. CNS stimulants and nootropics.  Opioid analgesics and antagonists  Drug addiction, drug abuse, tolerance and dependence.  Mode of Examination  Weightage Distribution  Weightage Distribution  Mode Assessment  10 Marks 15 75				www.sharda.ac.in	
A. 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).  B. Local anesthetic agents.  C. Drugs used in myasthenia gravis and glaucoma  UNIT-IV  Pharmacology of drugs acting on central nervous system  a. Neurohumoral transmission in the C.N.S. special emphasis on importance of various neurotransmitters like with GABA, Glutamate, Glycine, serotonin, dopamine.  b. General anesthetics and pre-anesthetics.  c. Sedatives, hypnotics and centrally acting muscle relaxants. Anti-epileptics, Alcohols and disulfiram  5  UNIT-V  Pharmacology of drugs acting on central nervous system  a. Psychopharmacological agents: Antipsychotics, antidepressants, anti-anxiety agents, anti-manics and hallucinogens.  b. Drugs used in Parkinsons disease and Alzheimer's disease.  c. CNS stimulants and nootropics.  Opioid analgesics and antagonists  Drug addiction, drug abuse, tolerance and dependence.  Mode of examination  Weightage  Distribution  Mode  Assessment  A. Ocotinuous  Mode  Assessment  A. ESE  Mode  Continuous  Mode  Assessment	3	UNIT-III			CO3
Neurohumoral transmission, co-transmission and classification of neurotransmitters.  Parasympathomimetics, Parasympatholytics, Sympathomimetics, sympatholytics.  Neuromuscular blocking agents and skeletal muscle relaxants (peripheral).  B. Local anesthetic agents.  C. Drugs used in myasthenia gravis and glaucoma  4  UNIT-IV Pharmacology of drugs acting on central nervous system  a. Neurohumoral transmission in the C.N.S. special emphasis on importance of various neurotransmitters like with GABA, Glutamate, Glycine, serotonin, dopamine.  b. General anesthetics and pre-anesthetics.  c. Sedatives, hypnotics and centrally acting muscle relaxants. Anti-epileptics, Alcohols and disulfiram  5  UNIT-V Pharmacology of drugs acting on central nervous system  a. Psychopharmacological agents: Antipsychotics, antidepressants, anti-anxiety agents, anti-manics and hallucinogens.  b. Drugs used in Parkinsons disease and Alzheimer's disease.  c. CNS stimulants and nootropics.  Opioid analgesics and antagonists  Drug addiction, drug abuse, tolerance and dependence.  Mode of examination  Weightage Distribution  Mode Assessment  Neurohumoral ransmission and skeletal muscle relaxants  CO4  CO5  CO5  CO5  CO5  CO5  CO5  CO5		Pharmacology of	drugs acting on periph	eral nervous system	
neurotransmitters. Parasympathomimetics, Parasympatholytics, Sympathomimetics, sympatholytics. Neuromuscular blocking agents and skeletal muscle relaxants (peripheral).  B. Local anesthetic agents. C. Drugs used in myasthenia gravis and glaucoma  4 UNIT-IV Pharmacology of drugs acting on central nervous system  a. Neurohumoral transmission in the C.N.S.special emphasis on importance of various neurotransmitters like with GABA, Glutamate, Glycine, serotonin, dopamine.  b. General anesthetics and pre-anesthetics. c. Sedatives, hypnotics and centrally acting muscle relaxants. Anti-epileptics, Alcohols and disulfiram  5 UNIT-V Pharmacology of drugs acting on central nervous system  a. Psychopharmacological agents: Antipsychotics, antidepressants, anti-anxiety agents, anti-manics and hallucinogens. b. Drugs used in Parkinsons disease and Alzheimer's disease. c. CNS stimulants and nootropics. Opioid analgesics and antagonists Drug addiction, drug abuse, tolerance and dependence.  Mode of examination  Weightage Distribution  Mode Assessment  CO4  CO4  CO5  CO5  CO5  CO5  CO5  CO5		A. Organiz	zation and function of A	NS.	
sympatholytics.  Neuromuscular blocking agents and skeletal muscle relaxants (peripheral).  B. Local anesthetic agents.  C. Drugs used in myasthenia gravis and glaucoma  4 UNIT-IV Pharmacology of drugs acting on central nervous system  a. Neurohumoral transmission in the C.N.S.special emphasis on importance of various neurotransmitters like with GABA, Glutamate, Glycine, serotonin, dopamine.  b. General anesthetics and pre-anesthetics.  c. Sedatives, hypnotics and centrally acting muscle relaxants. Anti-epileptics, Alcohols and disulfiram  5 UNIT-V Pharmacology of drugs acting on central nervous system  a. Psychopharmacological agents: Antipsychotics, antidepressants, anti-anxiety agents, anti-manics and hallucinogens.  b. Drugs used in Parkinsons disease and Alzheimer's disease.  c. CNS stimulants and nootropics.  Opioid analgesics and antagonists  Drug addiction, drug abuse, tolerance and dependence.  Mode of Theory  ESE  Mode Assessment  ESE			nsmission, co-transmiss	sion and classification of	
(peripheral).  B. Local anesthetic agents. C. Drugs used in myasthenia gravis and glaucoma  4  UNIT-IV Pharmacology of drugs acting on central nervous system a. Neurohumoral transmission in the C.N.S.special emphasis on importance of various neurotransmitters like with GABA, Glutamate, Glycine, serotonin, dopamine. b. General anesthetics and pre-anesthetics. c. Sedatives, hypnotics and centrally acting muscle relaxants. Anti-epileptics, Alcohols and disulfiram  5  UNIT-V Pharmacology of drugs acting on central nervous system a. Psychopharmacological agents: Antipsychotics, antidepressants, anti-anxiety agents, anti-manics and hallucinogens. b. Drugs used in Parkinsons disease and Alzheimer's disease. c. CNS stimulants and nootropics. Opioid analgesics and antagonists Drug addiction, drug abuse, tolerance and dependence.  Mode of examination  Mode Sessional Exam ESE  Mode Assessment  CO4  Assessment			etics, Parasympatholy	tics, Sympathomimetics,	
C. Drugs used in myasthenia gravis and glaucoma  4  UNIT-IV Pharmacology of drugs acting on central nervous system  a. Neurohumoral transmission in the C.N.S. special emphasis on importance of various neurotransmitters like with GABA, Glutamate, Glycine, serotonin, dopamine.  b. General anesthetics and pre-anesthetics.  c. Sedatives, hypnotics and centrally acting muscle relaxants. Anti-epileptics, Alcohols and disulfiram  5  UNIT-V Pharmacology of drugs acting on central nervous system  a. Psychopharmacological agents: Antipsychotics, antidepressants, anti-anxiety agents, anti-manics and hallucinogens.  b. Drugs used in Parkinsons disease and Alzheimer's disease.  c. CNS stimulants and nootropics.  Opioid analgesics and antagonists Drug addiction, drug abuse, tolerance and dependence.  Mode of examination  Weightage Distribution  Weightage Distribution  CO5			locking agents and s	skeletal muscle relaxants	
4 UNIT-IV Pharmacology of drugs acting on central nervous system a. Neurohumoral transmission in the C.N.S. special emphasis on importance of various neurotransmitters like with GABA, Glutamate, Glycine, serotonin, dopamine. b. General anesthetics and pre-anesthetics. c. Sedatives, hypnotics and centrally acting muscle relaxants. Anti-epileptics, Alcohols and disulfiram  CO5  UNIT-V Pharmacology of drugs acting on central nervous system a. Psychopharmacological agents: Antipsychotics, antidepressants, anti-anxiety agents, anti-manics and hallucinogens. b. Drugs used in Parkinsons disease and Alzheimer's disease. c. CNS stimulants and nootropics. Opioid analgesics and antagonists Drug addiction, drug abuse, tolerance and dependence.  Mode of examination  Weightage Distribution  CO5  CO5  CO5  CO5  CO5  CO5  CO5  CO		<b>B.</b> Local a	nesthetic agents.		
UNIT-IV Pharmacology of drugs acting on central nervous system  a. Neurohumoral transmission in the C.N.S.special emphasis on importance of various neurotransmitters like with GABA, Glutamate, Glycine, serotonin, dopamine.  b. General anesthetics and pre-anesthetics.  c. Sedatives, hypnotics and centrally acting muscle relaxants. Anti-epileptics, Alcohols and disulfiram  5  UNIT-V Pharmacology of drugs acting on central nervous system  a. Psychopharmacological agents: Antipsychotics, antidepressants, anti-anxiety agents, anti-manics and hallucinogens.  b. Drugs used in Parkinsons disease and Alzheimer's disease.  c. CNS stimulants and nootropics.  Opioid analgesics and antagonists  Drug addiction, drug abuse, tolerance and dependence.  Mode of examination  Weightage Distribution  Mode Assessment  CO5		C. Drugs v	ised in myasthenia gravi	s and glaucoma	
importance of various neurotransmitters like with GABA, Glutamate, Glycine, serotonin, dopamine.  b. General anesthetics and pre-anesthetics.  c. Sedatives, hypnotics and centrally acting muscle relaxants. Anti-epileptics, Alcohols and disulfiram  CO5  UNIT-V Pharmacology of drugs acting on central nervous system  a. Psychopharmacological agents: Antipsychotics, antidepressants, anti-anxiety agents, anti-manics and hallucinogens.  b. Drugs used in Parkinsons disease and Alzheimer's disease.  c. CNS stimulants and nootropics.  Opioid analgesics and antagonists  Drug addiction, drug abuse, tolerance and dependence.  Mode of Examination  Weightage Distribution  Mode Assessment  ESE	4		drugs acting on centra	l nervous system	CO4
c. Sedatives, hypnotics and centrally acting muscle relaxants. Anti-epileptics, Alcohols and disulfiram  CO5  UNIT-V Pharmacology of drugs acting on central nervous system  a. Psychopharmacological agents: Antipsychotics, antidepressants, anti-anxiety agents, anti-manics and hallucinogens.  b. Drugs used in Parkinsons disease and Alzheimer's disease.  c. CNS stimulants and nootropics.  Opioid analgesics and antagonists  Drug addiction, drug abuse, tolerance and dependence.  Mode of examination  Weightage Distribution  Mode Assessment  CO5  CO5  CO5  CO5  CO5  CO5  CO5  CO		importance of vari			
Anti-epileptics, Alcohols and disulfiram  UNIT-V Pharmacology of drugs acting on central nervous system  a. Psychopharmacological agents: Antipsychotics, antidepressants, anti-anxiety agents, anti-manics and hallucinogens.  b. Drugs used in Parkinsons disease and Alzheimer's disease.  c. CNS stimulants and nootropics.  Opioid analgesics and antagonists  Drug addiction, drug abuse, tolerance and dependence.  Mode of examination  Weightage Distribution  Mode Assessment  CO5  CO5  CO5  CO5  CO5  CO5  CO5  CNS  Antipsychotics, antidepressants, anti-manics and hallucinogens.  b. Drugs used in Parkinsons disease and Alzheimer's disease.  c. CNS stimulants and nootropics.  Opioid analgesics and antagonists  Drug addiction, drug abuse, tolerance and dependence.		<b>b.</b> General an			
Weightage Distribution  UNIT-V Pharmacology of drugs acting on central nervous system  a. Psychopharmacological agents: Antipsychotics, antidepressants, anti-anxiety agents, anti-manics and hallucinogens.  b. Drugs used in Parkinsons disease and Alzheimer's disease.  c. CNS stimulants and nootropics.  Opioid analgesics and antagonists  Drug addiction, drug abuse, tolerance and dependence.  Mode Assessment  ESE  Mode Assessment			· ·	y acting muscle relaxants.	
a. Psychopharmacological agents: Antipsychotics, antidepressants, anti-anxiety agents, anti-manics and hallucinogens.  b. Drugs used in Parkinsons disease and Alzheimer's disease.  c. CNS stimulants and nootropics.  Opioid analgesics and antagonists  Drug addiction, drug abuse, tolerance and dependence.  Mode of examination  Weightage Distribution  Mode Assessment  ESE	5		CO5		
c. CNS stimulants and nootropics. Opioid analgesics and antagonists Drug addiction, drug abuse, tolerance and dependence.  Mode of examination Weightage Distribution Mode Assessment  Continuous Mode Assessment  ESE		a. Psychopha			
c. CNS stimulants and nootropics. Opioid analgesics and antagonists Drug addiction, drug abuse, tolerance and dependence.  Mode of examination Weightage Distribution Mode Assessment  Continuous Mode Assessment  ESE		<b>b.</b> Drugs use	d in Parkinsons disease	and Alzheimer's disease.	
Drug addiction, drug abuse, tolerance and dependence.  Mode of examination  Weightage Distribution  Mode Assessment  Drug addiction, drug abuse, tolerance and dependence.  ESE		c. CNS stime	alants and nootropics.		
Drug addiction, drug abuse, tolerance and dependence.  Mode of examination  Weightage Distribution  Mode Assessment  Drug addiction, drug abuse, tolerance and dependence.  ESE		Opioid analgesics	and antagonists		
Mode of examination  Weightage Distribution  Mode Assessment  Mode Assessment  Mode Assessment			_	dependence.	
examination     Continuous     Sessional Exam     ESE       Distribution     Mode     Assessment					
Distribution Mode Assessment		Theory			
Assessment	Weightage		Sessional Exam	ESE	
	Distribution				
			15	75	



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



## COURSE ARTICULATION MATRIX

Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	1	2	3	-	2	-	2	2	3
CO2	3	3	-	2	3	-	1	-	3	1	3
CO3	3	3	2	2	3	-	1	-	3	3	3
CO4	3	2	-	2	3	-	2	-	2	2	3
CO5	3	3	-	1	3	_	2	_	3	2	3
CO6	3	2	2	-	-	-	1	-	3	1	2

- Slight (Low)
   Moderate (Medium)
   Substantial (High)



Sc	hool:	SOP						
Pr	ogramme:	B.Pharm						
Br	anch:	Semester: IV						
1	Course Code	BP405T						
2	Course Title	Pharmacognosy and Phytochemistry I - Theory						
3	Credits	4						
4	Contact	3-1-0						
	Hours							
	(L-T-P)							
	Course Type	Compulsory						
5	Course	Upon the completion of the course student shall be able to						
	Objective	Understand the techniques in the cultivation and production of crude drugs.						
		Identify the crude drugs, their uses and chemical nature.						
		Understand the evaluation techniques for the herbal drugs.						
		Carry out the microscopic and morphological evaluation of crude drugs						
6	Course	CO1: Students shall be able to define pharmacognosy, identify the sources of crude						
	Outcomes	drugs, and describe type of adulteration, evaluation of crude drugs, cultivation techniques, various medicine systems and plant tissue culture.						
		CO2: Students will be able to classify the crude drugs, understand their properties,						
		chemical nature and uses and are able to distinguish drugs with the help of chemical tests and describe various cultivation techniques.						
		CO3: Students will be able to apply their knowledge in identification, cultivation,						
		evaluation of drugs, and prescribing the crude drug for various health issues.						
		CO4: Students will analyze the crude drugs and its chemical nature and their						
		activities.						
		CO5: Students would be able to compare two drugs with the help of chemical and						
		physical properties, and evaluate them for their quality.						
		CO6: Students will be able to study plant tissue culture techniques.						
7	Course	The subject involves the fundamentals of Pharmacognosy like scope, classification of						
	Description	crude drugs, their identification and evaluation, phytochemicals present in them and						
		their medicinal properties.						
8	Outline syllab	us CO Mapping						





1	UNIT-I	CO1
	A. Introduction to Pharmacognosy:	
	a. Definition, history, scope and development of	
	Pharmacognosy	
	<ul> <li>b. Sources of Drugs – Plants, Animals, Marine &amp; Tissue culture</li> <li>c. Organized drugs, unorganized drugs (dried latex, dried juices, dried extracts, gums and mucilages, oleoresins and oleogum -resins).</li> </ul>	
	<b>B. Classification of drugs:</b> Alphabetical, morphological, taxonomical, chemical, pharmacological, chemo and sero taxonomical classification of drugs	
	<ul> <li>C. Quality control of Drugs of Natural Origin:</li> <li>a. Adulteration of drugs of natural origin. Evaluation by organoleptic, microscopic, physical, chemical and biological methods and properties.</li> </ul>	
	b. Quantitative microscopy of crude drugs including lycopodium spore method, leafconstants, camera lucida and diagrams of microscopic objects to scale with camera lucida.	
2	UNIT-II	CO2
	A. Cultivation, Collection, Processing and storage of drugs of	
	natural origin:	
	a. Cultivation and Collection of drugs of natural origin Factors influencing cultivation of medicinal plants. Plant hormones and	
	their applications.	
	b. Polyploidy, mutation and hybridization with reference to medicinal plants	
	B. Conservation of medicinal plants	
3	UNIT-III	CO3, CO6
	A. Plant tissue culture:	
	Historical development of plant tissue culture,	
	<b>B.</b> types of cultures, Nutritional requirements, growth and their maintenance.	
	C. Applications of plant tissue culture in pharmacognosy. Edible vaccines	
 i	, 44 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4	





4	UNIT V A. Study of biologi natural origin con B. Plant Products: Fibers - Cotton, Jute Hallucinogens, Terat	CO4						
5	UNIT V Study of biological natural origin contain A. Plant Products: Fibers - Cotton, Jute Hallucinogens, Terat	CO5						
	B. Primary metabo	olites:						
	General introduction	, detailed study with r	respect to chemistry,					
	sources, preparation	n, evaluation, preser I utility as Pharmaceu	with respect to chemistry, vation, storage, therapeutic atical Aids and/or Medicines					
	C. Carbohydrates:	Acacia, Agar, Tragac	eanth, Honey					
	(Papain, bromelain pepsin).	serratiopeptidase,	asein, proteolytic enzymes urokinase, streptokinase, oil, Chaulmoogra oil, Wool					
	Marine Drugs:							
		nts from marine source	ces					
Mode of examinatio	Theory	Theory						
Weightage	<b>Continuous Mode</b>	Sessional Exam	ESE					
Distributio	Assessment							
n	10 Marks	15	75					





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

### **COURSE ARTICULATION MATRIX**

Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	1	2	3	3	2	-	2	2	3
CO2	3	2	-	2	3	2	1	-	3	1	3
CO3	3	3	2	2	3	2	2	-	3	3	3
CO4	3	2	2	2	2	3	2	-	2	2	3
CO5	3	3	3	2	3	3	1	-	3	2	3
CO6	3	3	3	1	3	3	2	-	3	2	3

Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





<b>School:</b>		SOP						
Prograi	mme:	B. Pharm						
Branch	:	Semester: 4						
1	Course Code	BP406 P						
2	Course Title	Medicinal chemistry-i ( practial)						
3	Credits	2						
4	Contact Hours (L-T-P)	0-0-4						
	Course Type	Compulsory						
5	Course	Upon completion of the course, the student shall be able to						
	Objective	-Know the classification and salient features of five kingdon	ms of life					
		-Understand the basic components of anatomy & physiology of plantage. Know understand the basic components of anatomy & physiology with special reference to human						
6	Course							
	Outcomes	CO1: Students will be able to understand about practical land get hands-on experience of modern scientific instrumethodology, particularly in relation to the chemistry of phase	umentation and					
		CO2: Students will be able to apply knowledge and understanding of the fundamental principles of chemistry and their applications to pharmaceuticals.						
		CO3: Students will be able to use and apply their skills ar to a range of techniques used in pharmaceutical chemistry.	nd methodology					
		CO4: Students will be able to generalize professional trans exemplified by problem solving and teamwork.	ferable skills as					
		CO5: Students will be able to predict the skills to make sy for certain reactions involved in synthesis of drugs.	ynthetic scheme					
		CO6: Students will be able to apply knowledge about different methods to establish qualitative as well as quantitative rechemical entity.	-					
7	Course Description	Preparation of drugs/ intermediates, assay of drugs and dete Partition coefficient.	rmination of					
8	Outline syllabus		CO Mapping					
0	1	I Preparation of drugs/ intermediates	CO1					
		1,3-pyrazole 1,3-oxazole Benzimidazole	CO2					





	<b>5</b>		www.sharda.ac.in	G04
2	Benztriazole			CO1
	2,3- diphenyl quinox	CO6		
	Benzocaine			
	Phenytoin			
	Phenothiazine			
3	Assay of drugs			CO2
				CO3
	Barbiturate			CO6
	II Assay of drugs			
	Chlorpromazine			
	Phenobarbitone			
	Atropine			
	Ibuprofen			
4	Determination of P	Partition coefficie	nt for any two	CO1
	drugs			CO6
N 1 2	D (* 1/57*			
Mode of examination	Practical/Viva			
Weightage	Continuous	Sessional	ESE	
Distribution	Mode	Exam		
	Assessment			
	05	10	35	
Text book/s*		and Giswold's O	rganic medicinal and	
	2. Foye's F	Principles of Medi	cinal Chemistry.	
	3. Burger's	s Medicinal Chem	istry, Vol I to IV.	
	4. Introduction Smith and Williams			
	5. Remingt Martindale's extra	ton's Pharmaceuti pharmacopoeia	cal Sciences.	
Other				
References				





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	-	2	-	-		2	-	2	-	2
						2					
CO2	3	-	2	-	2	2	1	-	2	2	2
CO3	3	-	-	-	-	2	2	-	2	-	2
CO4	3	-	-	-	_	2	1	-	1	-	2
CO5	3	-	-	-	_	2	2	-	2	-	2
CO6	3	-	-	-		1	1	-	-	2	2

- 1-Slight (Low)
- 2-Moderate (Medium)
- 3-Substantial (High)





School	•	SOP					
Progra	mme:	B. Pharm					
Branch		Semester: 4					
1	Course Code	BP407P					
2	Course Title	Physical pharmaceutics II (Practical)					
3	Credits	2					
4	Contact Hours (L-T-P)	0-0-4					
	Course Type	Compulsory					
5	Course	Upon completion of the course, the student shall be able to					
	Objective	-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 with special reference to human					
6	Course Outcomes	CO1: Thestudents would be able to describe the derived properties of powder like angle of repose, bulk density, true density and porosity.					
		CO2: The students would be able to demonstrate methods for determination of HLB value and Critical Micelle concentration of surfactants.					
		CO3: The students would be able to analyze the particle size, particle size distribution by using methods like Sieving and Microscopic.					
		CO4: The students would be able to describe the viscosity, effect of sedimentation on suspension					
		CO5: The students would be able to plan about rate of reaction and accelerated stability studies according to ICH guidelines.					
		CO6: The students would be able to compare different types of stability parameters.					
7	Course Description	Determination of particle size, particle size distribution, derived properties of powder, viscosity, effect of suspending agent on sedimentation volume, factors affecting viscosity, viscosity determination and various stability studies as per ICH guidelines					
8	Outline syllabus	CO Mapping					





	1	1. Determination of particle size, particle size	CO1
		distribution using sieving method.	CO5
		2. Determination of particle size, particle size	CO6
		distribution using microscopic method.	
		3. Determination of bulk density, true density and	
		porosity.	
		4. Determine 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 of 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	
		Accelerated stability studies	
	2	Determination of particle size, particle size distribution	CO2
		using sieving	CO3
		method& microscopic method	
		I. Determination of particle size	
		II. Determination of particle size distribution	
		III. Using the sieving method	
	3	To determine the derived properties of powder	
		Determine the bulk density, true density and	CO3
		porosity of the powder	CO4
		Determine the flow properties of powder	
		Determine the effect of glidants on flow properties	
		of powder	
		or portaon	
	4	Determination of viscosity of liquids & semi solids	CO2
		Determination of viscosity of liquid by using	CO4
		Ostwald's viscometer	CO5
		Determination of viscosity of different	
		concentration of glycerineby using Ostwald's	
		viscometer	
		VISCOILICICI	
1			





5	Determination of	<b>Determination of sedimentation volume of suspension</b>					
	I. Determination	I. Determination sedimentation volume with effect of					
	different	suspending agent					
	II. Determination	on of sedimentation	n volume with				
Mode of examination	Practical/Viva						
Weightage	Continuous	Sessional Exam	ESE				
Distribution	Mode						
	Assessment						
	05	10	35				
Text book/s*	1. Physical Phar	macy by Alfred M	artin, Sixth edition				
	2. Experimental	pharmaceutics by	Eugene, Parott.				
	3. Tutorial pharm	macy by Cooper ar	nd Gunn.				
		4. Stocklosam J. Pharmaceutical calculations, Lea & Febiger, Philadelphia.					
Other							
References							

Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1		-	3	2	-		2	2	2	-	2
	2					2					
CO2	2	-	2	2	1	2	1	2	2	1	2
CO3	2	-	-	2	-	2	1	2	2	-	3
CO4	2	-	-	2	-	2	1	2	1	-	2
CO5	2	-	-	2	-	2	2	2	1	-	2
CO6	2	_	2	2	-	2	2	2	1	-	2

- 1-Slight (Low) 2-Moderate (Medium)
- 3-Substantial (High)



School:		SOP				
Program	ime:	B. Pharm				
Branch:		Semester: 4				
1	Course Code	BP408 P				
2	Course Title	Pharmacology I Practical				
3	Credits	2				
4	Contact Hours (L-T-P)	0-0-4				
	Course Type	Compulsory				
5	Course	Objectives:				
	Objective	1. Upon the completion of the course student shall be able to				
		2. Understand the pharmacological actions of different categories of drugs.				
		3.Observe the effect of drugs on animals by simulated experiments 4.Appreciate correlation of pharmacology with other bio medical sciences				
6	Course Outcomes	CO1: Thestudents would be able to explain the pharmacological aspects of drugs.				
		CO2: Thestudents would be able to understand and carry out the animal experiments				
		CO3: Thestudents would be able to appreciate the importance of Pharmacology subject as a basis of therapeutics.				
		CO4: The students would be able to Correlate and apply the knowledge therapeutically. CO5: The students would be able to apply experimental concepts. CO6: The students will be able to explore CNS action on experimental models.				





7	Course Description	Introduction to experimental pharmacology.					
		Commonly used instruments in experimental pharmacol	logy. Study of				
		common laboratory animals.					
		Maintenance of laboratory animals as per CPCSEA guideli	nes				
		Common laboratory techniques. Blood withdrawal, seru separation, anesthetics and euthanasia used for animal stud					
		Study of different routes of drugs administration in mice/ra	ts.				
		Study of effect of hepatic microsomal enzyme inducers on the phenobarbitone sleeping time in mice.					
		Effect of drugs on ciliary motility of frog oesophagus					
		Effect of drugs on rabbit eye.					
		Effects of skeletal muscle relaxants using rota-rod apparatu	IS.				
		Effect of drugs on locomotor activity using actophotometer	:.				
		Anticonvulsant effect of drugs by MES and PTZ method.					
		Study of stereotype and anti-catatonic activity					
8	Outline syllabus		CO Mapping				
	1	Basic Pharmacology Experiment	CO1				
		a). Introduction to experimental pharmacology.	CO5				
		b). Commonly used instruments in experimental					
		pharmacology.					
		c). Study of common laboratory animals.					
		d). Maintenance of laboratory animals as per CPCSEA					
		guidelines.					





1 _		T	_				
2			•	lab techniques an	d study the		CO2
		effect	s of Drugs				CO3
		a.	Common	laboratory	techniques.	Blo	CO6
			withdrawal,	serum and plasma	separation,		
		anesth	etics and euth	anasi			
		b.	Study of	different r	outes of	drugs	
		admin	istration in mi	ice/rats.			
		c.	Study of effe	ect of hepatic micro	osomal enzy	me	
		induce	ers on the phei	nobarbitone sleepir	ng time		
		in mic	ee.	•			
		d.	Effect of dru	gs on ciliary motil	ity of frog		
		oesop	hagus	•			
		_	_	gs on rabbit eye.			
				eletal muscle relax	ants using ro	ota-	
			paratus.		C		
			-	gs on locomotor a	ctivity using		
		_	notometer.		, ,		
		_		ant effect of drugs	by MES an		
					<i>y</i>		
N	Mode o	f Practi	cal/Viva				
e	xamination						
W	Veightage	Conti	nuous	Sessional Exam	ESE		
	Distribution	Mode					
			sment				
		05	<u></u>	10	35		
		0.5		10	55		





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

Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	2	3	3	-	2	1	-	2	2	3
CO2	3	2	2	2	-	2	-	-	-	1	3
CO3	3	-	2	-	-	2	2	-	2	-	3
CO4	3	-	2	-	-	2	-	-	-	2	3
CO5	3	-	2	-	-	2	2	-	2	-	3
CO6	1	2	2	3	_	3	2	_	1	2	-

1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





School:		SOP
Progran	nme:	B. Pharm
Branch:		Semester: 4
1	Course Code	BP409 P
2	Course Title	Pharmacognosy and Phytochemistry Practical
3	Credits	2
4	Contact Hours (L-T-P)	0-0-4
	Course Type	Compulsory
5	Course Objective	Upon completion of the course the student shall be able to understand different methods for analysis of crude drugs by various evaluation parameters i.e. Physical, chemical and organoleptic and anatomical parameters.
6	Course Outcomes	CO1: Students will discover practical laboratory skills and get hands- on experience of modern scientific instrumentation in relation to the Pharmacognosy.
		CO2: Students will receive knowledge and understanding of the fundamental principles of pharmacognosy and their applications to pharmaceuticals.
		CO3: Students will be able to use and apply their skills to a range of techniques used in pharmacognosy.
		CO4: Students will analyze various crude drugs by various methods.
		CO5: Students would be able to evaluate various crude drugs for their quality.
		CO6: Students would be able to study about chemical and quantitative analysis.
7	Course Description	Analysis of crude drugs by chemical tests: (i)Tragaccanth (ii) Acacia (iii)Agar (iv) Gelatin (v) starch (vi) Honey (vii) Castor oil Determination of stomatal number and index
		Determination of stomatal number and mack  Determination of vein islet number, vein islet termination and palisade ratio.  Determination of size of starch grains, calcium oxalate crystals by eye piece
		micrometer
		Determination of Fiber length and width
		Determination of number of starch grains by Lycopodium spore method
		Determination of Ash value
		Determination of Extractive values of crude drugs Determination of moisture
		content of crude drugs Determination of swelling index and foaming index.
8	Outline syllabus	CO Mapping
υ	Junine Synabus	CO iviapping





			www.sharda.ac.in				
 1				CO1			
	I Experiments invo	•	techniques	CO4			
	Chemical an	nalysis					
	<ul> <li>Macroscopio</li> </ul>	cal and microscopi	cal analysis				
	<ul> <li>Use of micro</li> </ul>	oscope, camera luc	ida, eye piece				
	micrometer	etc.					
2	II Determination of	II Determination of physical evaluation parameters					
	<ul> <li>Ash values</li> </ul>			CO5			
	<ul> <li>Extractive v</li> </ul>	alues					
	Moisture co	ntent					
	Swelling and	d foaming index					
3	III Evaluation of c	rude drugs by		CO1			
	anatomical/micros	copical evaluation	1	CO3			
	Stomatal nu	mber and Stomatal	Index	CO6			
	<ul> <li>Vein islet, v</li> </ul>	ein termination and	d Palisade ratio				
	<ul> <li>Fiber length</li> </ul>	and width					
	Size of starc	h grains and calciu	ım oxalate crystals				
4		Chemical and Quantitative analysis					
	•	crude drugs by che		CO2			
	Quantitative	analysis by lycop	odium spore method	CO3			
Mode of examination	Practical/Viva						
Weightage	Continuous	Sessional Exam	ESE				
Distribution	Mode						
	Assessment						
	05	10	35				
Text book/s*	1. W.C.Evans, Treedition, W.B. Sound	ease and Evans	Pharmacognosy, 16th				
		Brady, L.R. a					
		•	Febiger, Philadelphia,				
	1988.	i Lan., Lea and i	corger, rimaderpina,				
		<ul><li>1988.</li><li>3. Text Book of Pharmacognosy by T.E. Wallis</li></ul>					
			and Phytochemistry,				
	CBS Publishers & D	•	•				
		·	C.K. Kokate, Purohit,				
	Gokhlae (2007), 37th	~					
	1 ' ' '		dhary (1996), Ist Edn,				
	Eastern Publisher, N	• •	<i>y</i> ( : : = 0), -== = 011,				
Other	,						
References							





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	2	1	3	-	-	2	-	3	-	3
CO2	3	2	3	3	-	-	2	-	2	2	3
CO3	3	2	2	3	-	-	2	-	2	1	3
CO4	3	2	3	3	-	-	1	-	3	-	3
CO5	3	1	2	2	-	-	2	-	3	-	2
CO6	3	2	2	3	-	-	2	-	2	-	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





_	ogramme:	B.Pharm							
Br	anch:	Semester: V							
1	Course	BP501T							
	Code								
2	Course	Medicinal Chemistry-II - Theory							
	Title								
3	Credits	4							
4	Contact	3-1-0							
	Hours								
_	(L-T-P)								
	Course	Compulsory							
	Type								
	Course	Upon completion of the course the student shall be able to							
	Objective	1. understand the chemistry of drugs with respect to their pharmacological							
		activity							
		2. understand the drug metabolic pathways, adverse effect and therapeutic							
		value of drugs							
		3. know the Structural Activity Relationship (SAR) of different class of drugs							
6	Course	write the chemical synthesis of some drugs CO1: The students will be able to understand different categories of drugs							
	Outcomes	with respect to their pharmacological activities.							
	Outcomes	CO2: The students will apply and explain the structure activity relationship,							
		drug metabolic pathways, adverse effects and their therapeutic activity of							
		different categories of drugs.							
		CO3: The student's will be able to plan the knowledge to construct the							
		chemical synthesis of some drugs.							
		CO4: The students will analyse chemical reactions, stabilities of compounds							
		and properties of the compounds prepared by them in the laboratory.							
		CO5: The students will be able to modify and design new chemical							
		compounds with therapeutic activity.							
		CO6: Students will be able to anlyze chemical compounds with therapeutic							
		activity in different diseases.							
	C								
	Course	This subject is designed to impart fundamental knowledge on the structure,							
	Description	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.							
8	Outline syllab	bus CO Mapping							





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 (*)

CO1 CO6

#### Unit I

- A. **Antihistaminic agents:** Histamine, receptors and their distribution in the human body.
- B. **H1–antagonists:** Diphenhydramine hydrochloride*,
  Dimenhydrinate, Doxylamines cuccinate, Clemastine
  fumarate, Diphenylphyraline hydrochloride, Tripelenamine
  hydrochloride, Chlorcyclizine hydrochloride, Meclizine
  hydrochloride, Buclizine hydrochloride, Chlorpheniramine
  maleate, Triprolidine hydrochloride*, Phenidamine
  tartarate, Promethazine hydrochloride*, Trimeprazine
  tartrate, Cyproheptadine hydrochloride, Azatidine maleate,
  Astemizole, Loratadine, Cetirizine, Levocetrazine
  Cromolyn sodium
  - a. **H₂-antagonists:** Cimetidine*, Famotidine, Ranitidin.
  - b. **Gastric Proton pump inhibitors**: Omeprazole, Lansoprazole, Rabeprazole, Pantoprazole

### C. Anti-neoplastic agents:

- **a).** Alkylating agents: Meclorethamine*, Cyclophosphamide, Melphalan, Chlorambucil, Busulfan, Thiotepa
- **b).** Antimetabolites: Mercaptopurine*, Thioguanine, Fluorouracil, Floxuridine, Cytarabine, Methotrexate*, Azathioprine
- c). Antibiotics:Dactinomycin, Daunorubicin,Doxorubicin Bleomycin

**Plant products**: Etoposide, Vinblastin sulphate, Vincristin sulphate Miscellaneous: Cisplatin, Mitotane.





2		CO2
	Unit II	
	<ul> <li>A. Anti-anginal:</li> <li>B. Vasodilators: Amyl nitrite, Nitroglycerin*, Pentaerythritol tetranitrate, Isosorbide dinitrite*, Dipyridamole.</li> <li>C. Calciumchannel blockers: Verapamil, Bepridil hydrochloride, Diltiazem hydrochloride, Nifedipine, Amlodipine, Felodipine, Nicardipine, Nimodipine.</li> <li>Diuretics:</li> </ul>	
	Carbonicanhydrase inhibitors: Acetazolamide*,	
	Thiazides:Chlorthiazide*,Hydrochlorothiazide, 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, Sodium nitroprusside, Diazoxide, Minoxidil, Reserpine, Hydralazine hydrochloride.	
3	<ul> <li>Unit III         A. Anti-arrhythmic Drugs: Quinidine sulphate, Procainamide hydrochloride, Disopyramide phosphate*, Phenytoin sodium, Lidocaine hydrochloride, Tocainide hydrochloride, Mexiletine hydrochloride, Lorcainide hydrochloride, Amiodarone, Sotalol.     </li> <li>B. Anti-hyperlipidemicagents: Clofibrate, Lovastatin, Cholesteramineand Cholestipol</li> <li>C. Coagulant&amp;Anticoagulants: Menadione, Acetomenadione, Warfarin*, Anisindione, clopidogrel</li> <li>Drugs used in Congestive Heart Failure: Digoxin, Digitoxin, Nesiritide, Bosentan, Tezosentan.</li> </ul>	CO3





4	UNIT- IV			CO4			
	A. Drugs acting on	Endocrine system	n				
	Nomenclature, Stere	eochemistry and me	etabolism of steroids				
	<ul> <li>B. Sex hormones: Testosterone, Nandralone, Progestrones, Oestriol, Oestradiol, Oestrione, Diethyl stilbestrol.</li> <li>C. 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.</li> </ul>						
5	UNIT – V  A. Antidiabetic ag	ents:		CO5			
	Insulin and its prepa	rations					
	Sulfonyl ureas Tolbi	utamide*, Chlorpro	ppamide,Glipizide,				
	Glimepiride. Biguan	ides: Metformin.					
	ThiazolidinedionPio	glitazone, Rosiglit	azone. Meglitinides:				
	Repaglinide, Nategli	inide.					
	Glucosidase inhibito	ors: Acrabose, Vog	libose.				
	B. Local Anestheti	cs: SAR of Local	anesthetics				
	C. <b>Benzoic Acid de</b> Meprylcaine, Cy	erivatives; Cocaino clomethycaine, Pi	•				
			Benzocaine*, Butamben, e, Tetracaine, Benoxinate.				
	<b>Lidocaine/Anil</b> Prilocaine, Etido		Lignocaine, Mepivacaine,				
	Miscellaneous:						
Mode of examinat ion	Theory						
Weighta ge	Continuous Mode Assessment						
Distribut ion	10 Marks	15	75				





	www.snarda.ac.in	
Text	1. Wilson and Giswold's Organic medicinal and	
book/s	* 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.	
	<ul><li>8. The Organic Chemistry of Drug Synthesis by Lednicer,</li><li>Vol. 1to 5.</li><li>9. Indian Pharmacopoeia.</li></ul>	
	10. Text book of practical organic chemistry- A.I.Vogel.	
Other		
Referen	nc	
es		

## **COURSE ARTICULATION MATRIX**

Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	1	1	3	3	1	-	2	1	3
CO2	3	3	-	1	3	3	1	-	3	1	3
CO3	3	3	2	2	3	2	1	-	3	3	3
CO4	3	2	-	1	3	3	2	_	2	1	3
CO5	3	3	1	1	3	3	1	-	2	1	3
CO6	3	3	-	1	3	3	1	-	3	1	3

- 1-Slight (Low) 2-Moderate (Medium)
- 3-Substantial (High)





Scho	ool:	SOP								
Programme:		B. Pharm								
Bra		Semester: V								
1	Course Code	BP502 T								
2	Course Title	Industrial Pharmacy I - Theory								
3	Credits									
4	Contact	3-1-0								
	Hours									
	(L-T-P)									
	Course Type	Compulsory								
5	Course Objective	Upon completion of the course the student shall be able to  1. Know about the various pharmaceutical dosage forms and their manufacturing techniques (large scale equipment's etc)								
		2. Understand the various considerations in development of pharmaceutical dosage forms								
		3. Develop solid, liquid dosage forms and evaluate them for their quality Know about containers, closures and packaging material used for different type of dosage forms.								
6	Course Outcomes CO1: Students would be able to understand the concept of preformula studies for the development of safe and effective dosage form.									
		CO2: Students would be able to apply the knowledge various types of dosage form. (tablet, capsule, parenteral, liquid orals, pellets cosmetic preparations etc)								
		CO3: Students would be able to understand the formulation component and manufacturing procedures for different dosage form on Laboratory scale.								
		CO4: Students would be able to formulate ophthalmic Preparations.								
		CO5: Students would be able to formulate and evaluate the formulations for their quality.								
		CO6: Students shall be able to interpret regarding various packaging material for pharmaceutical products and evaluate them for quality.								
7	Course Description	Course enables the student to understand and appreciate the influence of pharmaceutical additives and various pharmaceutical dosage forms on the performance of the drug product.								
8	Outline syllabı	IS CO Mapping								





1	Unit I	CO1
	Preformulation studies-I Introduction to Preformulation, Goals and Objective, study of physicochemical characteristics of drug substances.  A. Physical properties: Physical form (crystal & amorphous), particle size, shape, flow properties, 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 and parenteral dosage forms and its impact on stability of dosage forms.	
2	<ul> <li>Unit II <ul> <li>A. Tablets: a. Introduction, ideal characteristics of tablets, classification of tablets. Excipients,</li> <li>Formulation of tablets, granulation methods,</li> <li>compression and processing problems. Equipments and tablet tooling.</li> <li>B. Tablet coating: Types of coating, coating materials, formulation of coating composition, methods of coating, equipment employed and defects in coating.</li> <li>C. Quality control tests: In process and finished product tests</li> <li>Liquid orals: Formulation and manufacturing consideration of syrups and elixirs suspensions and emulsions; Filling and packaging; evaluation of liquid orals official in pharmacopoeia</li> </ul> </li> </ul>	CO2





		27 TO TO THE THE TO THE	and the same of th
	3	Unit III	
		Capsules:	CO3
		A. Hard gelatin capsules: Introduction,	
		Production of hard gelatin capsule shells. size of	
		capsules, Filling, finishing and special techniques	
		of formulation of hard gelatin capsules,	
		manufacturing defects. In process and final	
		product quality control tests for capsules.	
		<b>B</b> . Soft gelatin capsules: Nature of shell and	
		capsule content, size of capsules,importance of	
		base adsorption and minim/gram factors,	
		production, in process and final product quality	
		control tests. Packing, storage and stability	
		testing of soft gelatin capsules and their	
		applications.	
		C. Pellets: Introduction, formulation requirements,	
		pelletization process, equipments for manufacture	
		of pellets	
	4	Unit IV	CO4
		<b>A.</b> Definition, types, advantages and limitations.	
		Preformulation factors and essential requirements,	
		vehicles, additives, importance of isotonicity	
		<b>B.</b> Production procedure, production facilities and	
		controls, aseptic processing c. Formulation of	
		injections, sterile powders, large volume	
		parenterals and lyophilized products.	
		C. Containers and closures selection, filling and	
		sealing of ampoules, vials and infusion fluids.	
		Quality control tests of parenteral products.	
		Ophthalmic Preparations: Introduction, formulation	
		considerations; formulation of eye drops, eye	
		ointments and eye lotions; methods of preparation;	
		labeling, containers; evaluation of ophthalmic	
1		preparations	





E .	TT24 T7			
5	following shampoos tooth paste  B. Pharmacer containers formulation Evaluation stability st  C. Packaging packaging influencin requireme	cosmetic , cold cream es, hair dyes ar utical Aerosols , valves, typ on and man n of aerosols cudies. ; Materials Sc of pharmac g choice of co nts for contai	a and preparation of the preparations: lipsticks, and vanishing cream, and sunscreens is: Definition, propellants, are of aerosol systems; aufacture of aerosols; is; Quality control and definition in the products of actions and are in the products of actions and in the products of actions are stability aspects of ality control tests	CO5 CO6
Mode of examination	Theory			
Weightage	Continuous	Sessional	ESE	
Distribution	Mode	Exam		
213410441011				
	Assessment			
	10 Marks	15	75	





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



### **COURSE ARTUCULATION MATRIX**

Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	2	1	3	-	2	_	2	2	3
CO2	3	2	3	2	2	-	3	-	3	3	3
CO3	3	3	2	2	3	-	1	-	3	3	3
CO4	3	2	-	1	3	-	2	-	2	1	3
CO5	3	3	1	1	2	-	1	-	2	2	3
CO6	3	3	-	1	2	-	1	-	2	2	3

1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sch	ool:	SOP
	gramme:	B. Pharm
	nch:	Semester: V
1	Course Code	BP503 T
2	Course Title	Pharmacology II-Theory
3	Credits	4
4	Contact	3-1-0
	Hours	
	(L-T-P)	
	Course Type	Compulsory
5	Course Objective	Upon completion of this course the student should be able to 1. Understand the mechanism of drug action and its relevance in the treatment of different diseases 2. Demonstrate isolation of different organs/tissues from the laboratory animals by simulated experiments 3. Demonstrate the various receptor actions using isolated tissue preparation 4. Appreciate correlation of pharmacology with related medical sciences
6	Course Outcomes	CO1: Students would be able to define and describe various categories of drugs to be used in the treatment of cardiovascular, haematological, endocrine and inflammatory disorders.  CO2: Students would be able to understand and explain the mechanisms, pharmacokinetic profile, adverse effects and uses of various drugs.  CO3: Students would be able to demonstrate the use of various categories of drugs and their bioassays.  CO4: Students would be able to analyze and explain the pathology of the cardiovascular, blood related and endocrine disorders.  CO5: Students would be able to evaluate and discriminate amongst the normal and abnormal physiological processes, and various drugs that can e employed for different treatment protocols.  CO6: Students would be able to plan about mechanism and therapeutic actions of drugs related to CVS, hormones.
7	Course Description	This subject is intended to impart the fundamental knowledge on various aspects (classification, mechanism of action, therapeutic effects, clinical uses, side effects and contraindications) of drugs acting on different systems of body and in addition, emphasis on the basic concepts of bioassay.





	0 11 11 1	www.sharda						
8	Outline syllabu		CO Mapping					
		Unit-1 A. Pharmacology of drugs acting on cardio vascular system a. Introduction to hemodynamic and electrophysiology of heart. b. Drugs used in congestive heart failure c. Anti-hypertensive drugs. d. Anti-anginal drugs. e. Anti-arrhythmic drugs. f. Anti-hyperlipidemic drugs.	CO1					
	2	Unit-2 A. Pharmacology of drugs acting on cardio vascular system a. Drug used in the therapy of shock. b. Hematinics, coagulants and anticoagulants. c. Fibrinolytics and anti-platelet drugs d. Plasma volume expanders B. Pharmacology of drugs acting on urinary system a. Diuretics b. Anti-diuretics.						
	3	Unit-3  A. Autocoids and related drugs  a. Introduction to autacoids and classification b. Histamine, 5-HT and their antagonists. c. Prostaglandins, Thromboxanes and Leukotrienes. d. Angiotensin, Bradykinin and Substance P. e. Non-steroidal anti-inflammatory agents f. Anti-gout drugs g. Antirheumatic drugs.	CO3					





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4	Unit-4 A. Pharmaco system B. Basic conce Anterior Pitu inhibitors. C.Thyroid hor Hormones Parathormone d. Insulin, Ora ACTH and co	CO4 CO6							
5	Unit-5 A. Pharmacolo a. Androgens b. Estrogens, pi c. Drugs actin B. Bioassay a. Principles a b. Types of bio c. Bioassay ACTH,d-tubo and 5-HT	CO5							
Mode of	Theory								
examination	<b>~</b> .•		Lege						
Weightage	Continuous	Sessional	ESE						
Distribution	Mode	Exam							
	Assessment								
Text book/s*			arma K. K., Principles of						
	<ul><li>2. Moo App</li><li>3. Gho of Phan</li></ul>	Pharmacology, Paras medical publisher  2. Modern Pharmacology with clinical Applications, by Charles R.Craig& Robert.  3. Ghosh MN. Fundamentals of Experimental Pharmacology. Hilton & Company, Kolkata.							
			andbook of experimental allabh Prakashan.						



## COURSE ARTUCULATION MATRIX

Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	2	1	1	3	-	1	-	2	1	3
CO2	3	2	2	1	3	-	1	-	2	1	3
CO3	3	3	2	1	3	-	1	-	3	3	3
CO4	3	2	2	1	3	-	2	-	2	1	3
CO5	3	3	2	1	3	-	1	-	2	1	3
C06	3	2	2	2	2	-	1	-	2	2	2

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





School:		SOP				
Programme:		B. Pharm				
Branch:		Semester: V				
1	Course	BP504T				
	Code					
2	Course	Pharmacognosy – II Theory				
	Title	·				
3	Credits	4				
4	Contact	3-1-0				
	Hours					
	(L-T-P)					
	Course	Compulsory				
	Type					
5	Course	Upon completion of the course, the student shall be able				
	Objective	1. to know the modern extraction techniques, characterization and				
		identification of the herbal drugs and phytoconstituents				
		2. to understand the preparation and development of herbal formulation.				
		3. to understand the herbal drug interactions				
6	Course	to carryout isolation and identification of phytoconstituents  CO1: Students would be able to define and describe various metabolic				
	Outcomes	pathways, various secondary metabolites like alkaloids glycosides by spectroscopic techniques and chromatography and various extraction methods				
		CO2: Students would be able to explain applications of phytoconstituents and their industrial production, isolation process and extraction methods				
		CO3: Students would be able to apply and demonstrate various identification process and latest technique of phytoconstituents				
		CO4: Students would be able to separate and analyse various phytoconstituents				
		CO5: Students would be able to evaluate various phytoconstituent				
		CO6: Students would be able to understand about metabolic pathways.				
7	Course	The main purpose of subject is to impart the students the				
	Description	knowledge of how the secondary 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				





8	Outline Syllabus		CO
	1	<ul> <li>UNIT-I</li> <li>Metabolic pathways in higher plants and their determination</li> <li>a. Brief study of basic metabolic pathways and formation of different secondary metabolites through these pathways-Shikimic acid pathway,</li> <li>b. Acetate pathways and Amino acid pathway.</li> <li>c. Study of utilization of radioactive isotopes in the investigation of Biogenetic studies.</li> </ul>	Mapping CO1 CO6
	2	UNIT-II General introduction, composition, chemistry & chemical classes, biosources, therapeutic uses and commercial applications of following  a. secondary metabolites:	CO2
		Alkaloids: Vinca, Rauwolfia, Belladonna, Opium, Phenylpropanoids and Flavonoids: Lignans, Tea, Ruta Steroids, Cardiac Glycosides & Triterpenoids: Liquorice, Dioscorea, Digitalis Volatile oils: Mentha, Clove, Cinnamon, Fennel, Coriander, Tannins: Catechu, Pterocarpus b. Resins: Benzoin, Guggul, Ginger, Asafoetida, Myrrh, Colophony c. Glycosides: Senna, Aloes, Bitter Almond Iridoids, Other terpenoids & Naphthaquinones: Gentian, Artemisia, taxus, carotenoids	
	3	UNIT-III Isolation, Identification and Analysis of Phytoconstituents ATerpenoids: Menthol, Citral, Artemisin B. Glycosides: Glycyrhetinic acid & Rutin C. Alkaloids: Atropine, Quinine, Reserpine, Caffeine Resins: Podophyllotoxin, Curcumin	CO3
	4	<ul> <li>UNIT-IV</li> <li>A. Industrial production, estimation and utilization of the following phytoconstituents:</li> <li>B. Forskolin, Sennoside, Artemisinin, Diosgenin, Digoxin, Atropine, Podophyllotoxin,</li> <li>C. Caffeine, Taxol, Vincristine and Vinblastine</li> </ul>	CO4





		-	THE TRANSPORT OF THE PARTY OF T		
Mode of examinati on	a. Basics of Phytochemistry b. Modern methods of extraction, application of latest techniques like Spectroscopy, c. chromatography and electrophoresis in the isolation, purification and identification of crude drugs.  Theory				
Weightag e	Continuous Mode Assessment	Sessional Exam	ESE		
Distributi on	10 Marks	15	75		
Text book/s*	W.C. Evans, Trease and Evans Pharmacognosy, 16th edition, W.B. Sounders & Co., London, 2009.  Mohammad Ali. Pharmacognosy and Phytochemistry, CBS Publishers & Distribution, New Delhi.  Text book of Pharmacognosy by C.K. Kokate, Purohit, Gokhlae (2007), 37th Edition, Nirali Prakashan, New Delhi.  Herbal drug industry by R.D. Choudhary (1996), Ist Edn, Eastern Publisher, New Delhi.  Essentials of Pharmacognosy, Dr.SH.Ansari, IInd edition, Birla publications, New Delhi, 2007  Herbal Cosmetics by H.Pande, Asia Pacific Business press, Inc, New Delhi.  A.N. Kalia, Textbook of Industrial Pharmacognosy, CBS Publishers, New Delhi, 2005.  R Endress, Plant cell Biotechnology, Springer-Verlag, Berlin, 1994.  Pharmacognosy & Pharmacobiotechnology. James Bobbers, Marilyn KS, VE Tylor.				
Other Reference s					





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	2	1	3	-	1	-	2	1	3
CO2	3	2	-	1	3	-	1	-	3	1	3
CO3	3	3	2	2	3	-	1	-	3	3	3
CO4	3	2	-	1	3	-	2	-	2	1	3
CO5	3	3	1	1	3	-	1	-	2	1	3
CO6	3	3	2	2	3	-	1	_	3	3	3

1-Slight (Low) 2-Moderate (Medium)

3-Substantial (High)





Sc	chool:	SOP	
Pr	ogramme:	B.Pharm	
Bı	ranch:	Semester: V	
1	Course	BP505T	
	Code		
2	Course	Pharmaceutical Jurisprudence - Theory	
	Title		
3	Credits	4	
4	Contact	3-1-0	
	Hours		
	(L-T-P)		
	Course	Compulsory	
	Type		
5	Course	Upon completion of the course, the student shall be able to un	
	Objective	1. The Pharmaceutical legislations and their implications in	the development
		and marketing of pharmaceuticals.	
		Various Indian pharmaceutical Acts and Laws	C . 1 1
		3. The regulatory authorities and agencies governing the mar	nufacture and sale
-	Carrage	of PharmaceuticalsCode of ethics	4111 - 1 C
6	Course Outcomes	CO1: Students would be able to identify and understand	_
	Outcomes	Legal definitions of schedules to the Act and Rules, license f and sale of drugs.	of manufacturing
		and sale of drugs.	
		CO2: Students would be able to explain various schedul	es labelling and
		packaging of drugs and various acts and rules.	ies labelling and
		packaging of drags and various acts and rules.	
		CO3: Students would be able to differentiate various acts and	d rules Schedules
		sale of drugs and various Acts and Rules and apply Rules.	<b>3 1010</b>
		CO4: Students would be able to interpret various Acts and I	Rules and how to
		apply various acts and Rules.	
		CO5: Students would be able to summarize the acts and code	and conduct and
		also would explain Intellectual Proprietary Rights	
		CO6: Students would be able to plan about various schedules	of D&C Act.
7	Course	This course is designed to impart basic knowledge	on important
	Description	legislations related to the profession of pharmacy in India	on important
$\vdash$		registations related to the profession of pharmacy in mula	
	Outline syllab	MIC.	CO Manning
8	Outime synat	ous	CO Mapping
О			





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1	UNIT-I	CO1
	Drugs and Cosmetics Act, 1940 and its rules 1945:	
	Objectives, Definitions, Legal definitions of schedules to the	
	Act and Rules	
	a. Import of drugs – Classes of drugs and cosmetics	
	prohibited from import, Import under license or permit.	
	Offences and penalties.	
	b. Manufacture of drugs – Prohibition of manufacture and sale of certain	
2	UNIT-II	CO2
		CO2 CO6
	A. Drugs and Cosmetics Act, 1940 and its rules 1945.	C00
	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	
	B. Labeling & Packing of drugs- General labeling requirements and specimen labels for drugs and cosmetics, List of permitted colors. Offences and penalties.	
	C. Administration of the Act and Rules – Drugs Technical Advisory Board, Central drugs Laboratory, Drugs Consultative Committee, Government drug analysts, Licensing authorities, controlling authorities, Drugs Inspectors	





3 **UNIT-III** CO₃ **A. Pharmacy Act –1948**: Objectives, Definitions, Pharmacy Council of India; its constitution and functions, Education Regulations, State and Joint state pharmacy councils; constitution and functions, Registration of Pharmacists, Offences and Penalties B. 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. C. 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 **UNIT-IV** 4 CO₄ A. Study of Salient Features of Drugs and Magic Remedies and its rules: Objectives, Definitions, Prohibition of certain advertisements, Classes of Exempted advertisements, Offences and Penalties B. Prevention of Cruelty to animals Act-1960: Objectives, Definitions, Institutional Animal Ethics Committee, CPCSEA guidelines for Breeding and Stocking of Animals, Performance of Experiments, Transfer and acquisition of animals for experiment, Records, Power to suspend or revoke registration, Offences and Penalties C. 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 Essential Medicines (NLEM)





5	UNIT-V	CO5				
	A. Pharmaceutic Introduction, Study and development c committee					
			ession and his profession,			
	C.Medical Terminat	tion of Pregnancy	Act			
	Right to I	nformation Act				
	Introduct	ion to Intellectual	Property Rights (IPR)			
Mode of examina tion	Theory					
Weighta ge	Continuous Mode Assessment	Sessional Exam	ESE			
Distribut ion	10 Marks	15	75			
Text	1. Text book o	of Forensic Pharma	cy by B.M. Mithal			
book/s*	2. Hand book					
	3. A text book	of Forensic Pharm	nacy by N.K. Jain			
	4. Drugs and publications					
	5. Medicinal a India public		ions act 1955 by Govt. of			
	6. Narcotic drugs and psychotropic substances act by Govt. of India publications					
	7. Drugs and publication	_	act by Govt. of India			
		of the said laws pu ooks (Theory)	iblished by Government.			



Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	2	1	2	3	3	1	-	2	1	3
CO2	3	2	2	1	3	3	1	-	3	1	3
CO3	3	3	2	2	3	2	1	-	3	3	3
CO4	3	2	2	2	3	3	2	-	2	1	3
CO5	3	3	1	1	3	3	1	-	2	1	3
CO6	3	2	2	2	3	3	2	-	2	1	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sc	hool:	SOP
Pr	ogramme:	B.Pharm
	anch:	Semester: V
1	Course	BP506P
	Code	
2	Course	Industrial Pharmacy- I Practical
	Title	
3	Credits	
4	Contact	0-0-4
	Hours	
	(L-T-P)	Commulación
	Course Type	Compulsory
5	Course	Upon completion of the course the student shall be able to
	Objective	opon completion of the course the student shan be able to
	<b></b>	1. Know about the various pharmaceutical dosage forms and their
		manufacturing techniques( large scale equipment's etc)
		2. Understand the various considerations in development of pharmaceutical
		dosage forms
		3. Develop solid, liquid dosage forms and evaluate them for their quality
		Know about containers, closures and packaging material used for different
		type of dosage forms.
6	Course	CO1: Students would be able to understand various types of dosage
	Outcomes	form. (tablet, capsule, Parenterals, creams etc)
		CO2: Students would be able to plan about the manufacturing procedures for
		different dosage form on laboratory scale. (tablet, capsule, Parenterals,
		creams etc)
		CO3: Students would be able to evaluate the Aerosols.
		CO3. Students would be able to evaluate the Acrosols.
		CO4: Students would be able to plan various packaging material for
		pharmaceutical products and evaluate them for quality
		CO5: Students shall be able to formulate Cosmetics.
		CO6: Students shall be able to evaluate the different types of Descar forms
		CO6: Students shall be able to evaluate the different types of Dosage forms.





	~						
7	Course Description	To emphasize on definition, types, classification, principles/mechanisms, applications, examples and differences Formulation and evaluation of the following dosage forms containing drugs mentioned in pharmacopoeia.  1. Capsules. 2. Microcapsules/microspheres 3. Tablets by dry and wet granulation methods					
		4. Film coated tablets/ Enteric coated tablets, Ear drops					
8	Outline Syllal	bus	CO Mapping				
	1	To study the various instruments used in evaluation of tablet.	CO1				
	2	Evaluation of tablet as per IP.	CO2 CO6				
	3	Prepare and evaluate granules of Calcium lactate 50 Tablets.	CO3 CO6				
	4	To compress the prepared granules Of Acetyl salicylic acid by using Tablet Making Machine and determine their Disintegration Time and hardness of prepared tablets.	CO4				
	5	To study the effect of coating on disintegration of tablets.	CO5				
	6	To prepare effervescent granules by hot and wet method.	CO1				
	7	To prepare microcapsules by using phase separation & coacervation technique brought about by polymer polymer interaction.	CO2				
	8	To prepare and submit cold cream	CO3				
	9	Preparation of injection	CO4				
	10	To prepare and submit vanishing cream	CO5				
	Mode of examination	Practical/Viva					





Weightage	Continuous	Sessional Exam	ESE				
Distribution	Mode						
	Assessment						
	05Marks	10	35				
Text book/s*	H.A. Liberman, Le Pharmaceutical dos	Pharmaceutical dosage forms - Tablets, volume 1 -3 by H.A. Liberman, Leon Lachman & J.B.Schwartz. Pharmaceutical dosage form - Parenteral medication vol-&2 by Liberman & Lachman					
	Pharmaceutical dos & Lachman	sage form disperse	system VOL-1 by Liberman				
Other							
References							





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	2	2	2	3	3	1	-	2	1	3
CO2	3	3	-	1	3	2	1	-	3	1	3
CO3	3	3	2	2	3	2	1	-	3	3	3
CO4	3	2	-	1	2	3	2	-	2	1	3
CO5	3	3	1	1	3	3	1	-	2	1	3
CO6	3	3	-	1	3	3	1	-	2	1	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sc	hool:	SOP
Pr	ogramme:	B. Pharm
Br	anch:	Semester: V
1	Course	BP507P
	Code	
2	Course	Pharmacology- II Practical
	Title	
3	Credits	2
4	Contact	0-0-4
	Hours	
	(L-T-P)	
	Course	Compulsory
	Type	
5	Course	Upon completion of this course the student should be able to
	Objective	1. Understand the mechanism of drug action and its relevance in the
		treatment ofdifferent diseases
		2. Demonstrate isolation of different organs/tissues from the laboratory
		animals bysimulated experiments
		3. Demonstrate the various receptor actions using isolated tissue
		preparation
		4. Appreciate correlation of pharmacology with related medical
6	Course	sciences  CO1: Students would be able to define and describe various instruments and
U	Outcomes	methods used in the evaluation of <i>in vitro</i> and <i>in vivo</i> evaluation of various
	outcomes	drugs.
		CO2: Students would be able to understand and explain the working
		principles of the instruments used and actions of various drugs on biological
		systems.
		CO3: Students would be able to demonstrate the effects of various categories
		of drugs and bioassays of physiological substances.
		CO4: Students would be able to analyze and explain the outcomes of
		experiments through simulation studies.
		COS. Students mould be able to confer and 1 1 1 1 1 1
		CO5: Students would be able to evaluate and discriminate amongst the
		normal and abnormal physiological processes, and various drugs that can be employed for different treatment protocols.
		employed for different treatment protocols.
		CO6: Students would be able to plan about the effects of various bioassays
		of physiological substance and its application.
		or physiological substance and no application.
		I





		27 27 27	
7	Course Description	<ol> <li>Introduction to <i>in-vitro</i> pharmacology and physiological salt solu</li> <li>Effect of drugs on isolated frog heart.</li> <li>Effect of drugs on blood pressure and heart rate of dog.</li> <li>Study of diuretic activity of drugs using rats/mice.</li> <li>DRC of acetylcholine using frog rectus abdominis muscle.</li> <li>Effect of physostigmine and atropine on DRC of acetylcholine rectus abdominis muscle and rat ileum respectively.</li> </ol>	using frog
		7. Bioassay of histamine using guinea pig ileum by matching methe 8. Bioassay of oxytocin using rat uterine horn by interpolation meth 9. Bioassay of serotonin using rat fundus strip by three point bioass 10. Bioassay of acetylcholine using rat ileum/colon by four point be	nod. say. ioassay.
		<ul> <li>11. Determination of PA2 value of prazosin using rat anococcygeu (by Schilds plot method).</li> <li>12. Determination of PD2 value using guinea pig ileum.</li> <li>13. Effect of spasmogens and spasmolytics using rabbit jejunum.</li> </ul>	
		<ul><li>14. Anti-inflammatory activity of drugs using carrageenan induced edema model.</li><li>Analgesic activity of drug using central and peripheral methods</li></ul>	ı paw-
8	Outline Syall		CO Mapping
	1	Introduction to <i>in-vitro</i> pharmacology and physiological salt solutions.	CO1
	2	Effect of drugs on isolated frog heart.	CO2
	3	Effect of drugs on blood pressure and heart rate of dog.	CO3
	4	Study of diuretic activity of drugs using rats/mice.	CO4
	5	DRC of acetylcholine using frog rectus abdominis muscle.	CO5
	6	Effect of physostigmine and atropine on DRC of acetylcholine using frog rectus abdominis muscle and rat ileum respectively.	CO1
	7	Bioassay of histamine using guinea pig ileum by matching method.	CO2 CO6
	8	Bioassay of oxytocin using rat uterine horn by interpolation method.	CO3 CO6





9	Bioassay of seroto bioassay.	Bioassay of serotonin using rat fundus strip by three point bioassay.						
10	Bioassay of acety bioassay.	Bioassay of acetylcholine using rat ileum/colon by four point bioassay.						
Mode of examination	Practical/Viva							
Weightage Distribution	Continuous Mode Assessment	Mode						
Text book/s*	medical publisher Modern Pharmaco R.Craig& Robert. Ghosh MN. Fu Pharmacology. Hill	Sharma H. L., Sharma K. K., Principles of Pharmacology, Paras medical publisher Modern Pharmacology with clinical Applications, by Charles R.Craig& Robert. Shosh MN. Fundamentals of Experimental Pharmacology. Hilton & Company, Kolkata. Kulkarni SK. Handbook of experimental pharmacology. Vallabh						
Other References								



Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	1	1	3	3	1	-	2	1	3
CO2	3	3	2	1	3	3	1	-	3	1	3
CO3	3	3	2	2	3	2	1	-	3	3	3
CO4	3	2	2	1	3	3	2	-	2	1	3
CO5	3	3	1	1	3	3	1	-	2	1	3
C06	2	2	2	1	1	3	3	-	3	3	1

1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sc	hool:	SOP
Pr	ogramme:	B. Pharm
	anch:	Semester: V
1	Course	BP508P
	Code	
2	Course	Pharmacognosy - II Practical
	Title	
3	Credits	2
4	Contact	0-0-4
	Hours	
	(L-T-P)	
	Course	Compulsory
	Type	
5	Course Objective	1. Explain correct use of various equipments in Pharmacognosy laboratory.
		2. Handle simple/compound microscope in technically correct way.
		3. Expain and understand the Morphology, histology and powder characteristics
		4. Demonstrate skill of plant material sectioning, staining, mounting & focusing.
		5. Decide on staining reagents required for specific part of plant.
		6. Demonstrate Isolation and detection methods
		7. Separate phytoconstituents by TLC
6	Course	CO1: Students would able to identify and describe the morphology and
	Outcomes	chemical test of crude drugs
		CO2: Students would be able to explain and compare the microscopy of crude
		drugs and powder
		CO3: Students would be able to calculate the Rf value of phytoconstituents
		CO4: Students would be able to separate and analyze the phytoconstituents
		CO5: Students would be able to apply knowledge to isolate the compounds.
		CO6: Students would be able to evaluate the Phytoconstituents.





7	Course Description	Morphology, histology and powder characteristics & extraction & detection of: Cinchona, Cinnamon, Senna, Clove, Ephedra, Fennel and Coriander Exercise involving isolation & detection of active principles a. Caffeine - from tea dust. b. Diosgenin from Dioscorea c. Atropine from Belladonna d. Sennosides from Senna							
		<ol> <li>Separation of sugars by Paper chromatography</li> <li>TLC of herbal extract</li> </ol>	titutanta by						
		<ul> <li>4. Distillation of volatile oils and detection of phytocons TLC</li> <li>5. Analysis of crude drugs by chemical tests:</li> </ul>	intutents by						
		(i) Asafoetida (ii) Benzoin (iii) Colophony (iv) Aloes (v) Myrrh							
8	Outline Sylla	bus	CO Mapping						
	1	To study the morphological and microscopy of Cinchona bark.	CO1						
	2	To study the morphological and microscopy of Fennel fruits.	CO2						
	3	To study the morphological characteristics of Senna leaves, Cinnamon bark and Ephedra stem	CO3						
	4	To study the powder characteristics of clove buds and Cinnamon bark	CO4						
	5	To study the morphological, and histological characteristics of clove bud	CO5						
	6	To study the morphological, microscopy and powder characteristics of Ephedra stem	CO1						
	7	To extract Caffeine from tea powder and identify by Thin Layer chromatography  CO2 CO6							
	8	To perform separation of sugars by Paper chromatography CO3							
	9	To perform Thin Layer chromatography of the given herbal extract.	CO4 CO6						





10	To isolate volate Clavengers apparatiquid using Ostw	CO5				
Mode of examination	Practical/Viva					
Weightage Distribution	Continuous Mode Assessment	Mode				
	05 Marks	10	35			
Text book/s*	Pharmodelling Ph	n, 1994. nacognosy & Phers, Marilyn KS, VI formulation and ances and flavours. ngton's Pharmaceut Book of Biotechnol	Publishers, New otechnology, Springer-Verlag, narmacobiotechnology. James E Tylor. preparation of cosmetic,			
Other References						



Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	2	2	1	2	3	1	_	2	2	3
CO2	3	3	-	1	3	3	1	-	3	1	3
CO3	3	3	1	2	3	2	1	-	3	3	3
CO4	3	2	-	1	3	3	2	-	2	1	3
CO5	3	3	1	1	3	3	1	-	2	1	3
CO6	3	2	-	1	3	3	2	-	2	1	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sc	hool:	SOP							
	ogramm	B.Pharm							
e:									
Branch:		Semester: VI							
1	Course	BP601T							
	Code								
2 Course		Medicinal Chemistry III – Theory							
	Title								
3	Credits	4							
4	Contact	3-1-0							
	Hours								
	(L-T-P)								
	Course	Compulsory							
_	Type	The accomplation of the course student shall be able to							
5	Course	Upon completion of the course student shall be able to-							
	Objectiv e	1. Understand the importance of drug design and different techniques of drug							
	C	design.  2. Understand the chemistry of drugs with respect to their biological							
		activity.							
		3. Know the metabolism, adverse effects and therapeutic value of drugs.							
		Know the importance of SAR of drugs.							
6	Course	CO1: Student will be able to describe get fundamental knowledge of the							
	Outcom	structure, chemistry and its correlation with the therapeutic value of drugs.							
	es								
		CO2: Students will be able to apply conceptual knowledge and background of							
		drugs and ensure their rational use.							
		CO3: Students will be able to plan about the synthesis and Structure Activity							
		Relationships (SAR) associated with the drugs structure.							
		CO4: Students will also conclude about the chemistry, mechanism of action,							
		metabolism, adverse effects, and therapeutic uses of important drugs.							
		memoonism, adverse effects, and merapeutic uses of important drugs.							
		CO5: Students will be able to the modern techniques of rational drug design like							
		quantitative structure activity relationship							
		•							
		CO6: Students will be able to construct the Drug design							
7	Course	This subject is designed to impart fundamental knowledge on the structure,							
	Descript	chemistry and therapeutic value of drugs. The subject emphasis on modern							
	ion	techniques of rational drug design like quantitative structure activity relationship							
		(QSAR), Prodrug concept, combinatorial chemistry and Computer aided drug							
		design (CADD). The subject also emphasizes on the chemistry, mechanism of							
		action, metabolism, adverse effects, Structure Activity Relationships (SAR),							
		therapeutic uses and synthesis of important drugs.							





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8	Outline sy	flabus	CO Mapping
	1	<ul> <li>UNIT – I         <ul> <li>A. Antibiotics</li> </ul> </li> <li>Historical background, Nomenclature, Stereochemistry,         <ul> <li>Structure activity relationship, Chemical degradation                 classification and important products of the following classes.</li> </ul> </li> <li>β-Lactam antibiotics: Penicillin, Cepholosporins, β- Lactamase inhibitors, Monobactams         <ul> <li>Aminoglycosides: Streptomycin, Neomycin, Kanamycin</li> <li>Tetracyclines: Tetracycline, Oxytetracycline,</li></ul></li></ul>	CO1
	2	<ul> <li>UNIT - II</li> <li>A. Antibiotics</li> <li>Historical background, Nomenclature, Stereochemistry, Structure activity relationship, Chemical degradation classification and important products of the following classes.</li> <li>B. Macrolide: Erythromycin Clarithromycin, Azithromycin.</li> <li>C. Miscellaneous: Chloramphenicol*, Clindamycin.</li> <li>Prodrugs: Basic concepts and application of prodrugs design.</li> </ul>	CO2
		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	





3 UNIT - III CO₃ A. Anti-tubercular Agents B. Synthetic anti tubercular agents: Isoniozid*, Ethionamide, Ethambutol, Pyrazinamide, Para amino salicylic acid.* C. Anti tubercular antibiotics: Rifampicin, Rifabutin, Cycloserine Streptomycine, Capreomycin sulphate. Urinary tract anti-infective agents Quinolones: SAR of quinolones, Nalidixic Acid, Norfloxacin, Ciprofloxacin*, Enoxacin, Ofloxacin, Lomefloxacin, Sparfloxacin, **Miscellaneous:** Furazolidine, Nitrofurantoin*, Methanamine. **Antiviral agents:** Amantadine hydrochloride, Rimantadine hydrochloride, Idoxuridine trifluoride, Acyclovir*, Gancyclovir, Zidovudine, Didanosine, Zalcitabine, Lamivudine, Loviride, Delavirding,

Ribavirin, Saquinavir, Indinavir,





4 UNIT - IV CO4

- A. Antifungal agents:
- B. **Antifungal antibiotics:** Amphotericin-B, Nystatin, Natamycin, Griseofulvin.
- C. **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.

**Anthelmintics:** Diethylcarbamazine citrate*, 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*.





5	UNIT - V  A. Introduction	CO5 CO6		
	Various approaches	used in drug desigr	ı <b>.</b>	
	activity relationship	(QSAR) such	n quantitative structure as partition coefficient, ts steric parameter and	
	Pharmacophore mod	eling and docking	techniques.	
	B. Combinator chemistry:	ial Chemistry: C	oncept and applications	
	C. solid phase a	nd solution phase s	ynthesis.	
Mode of examina tion	Theory			
Weighta ge	Continuous Mode Assessment	Sessional Exam	ESE	
Distribut ion	10 Marks	15	75	
Text book/s*	Wilson and Giswo Chemistry.     Foye's Principles of Burger's Medicina Introduction to principle.			
Other Referenc es	Text book of practice			



Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	2	1	3	-	1	-	2	1	3
CO2	3	3	1	1	3	-	1	-	3	1	3
CO3	3	3	3	2	3	-	1	-	3	3	3
CO4	3	2	-	1	3	-	2	-	2	1	3
CO5	3	3	1	1	3	-	1	-	2	1	3
CO6	3	3	2	2	3	-	1	-	3	3	3

1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





So	chool:	SOP				
Pı	ogramme:	B.Pharm				
Bı	anch:	Semester: VI				
1	Course Code	BP602T				
2	Course Title	Pharmacology - III – Theory				
3	Credits	4				
4	Contact Hours (L-T-P)	3-1-0				
	Course Type	Compulsory				
5	Course Objective	Upon completion of this course the student should be able to:				
	·	<ol> <li>understand the mechanism of drug action and its relevance in the treatment of different infectious diseases</li> <li>comprehend the principles of toxicology and treatment of various poisonings</li> <li>appreciate correlation of pharmacology with related medical</li> </ol>	sciences.			
6	Course Outcomes	CO1: Students would be able to define and describe various category drugs to be used in the treatment of respiratory, gastrointestinal, in and malignant disorders.  CO2: Students would be able to understand and explain the repharmacokinetic profile, adverse effects and uses of various drugs.  CO3: Students would be able to demonstrate the use of various care drugs and their bioassays.  CO4: Students would be able to analyze and explain the pathology infectious, respiratory and gastrointestinal diseases.  CO5: Students would be able to evaluate and discriminate amongs and abnormal physiological processes, and various drugs that can be for different treatment protocols.  CO6: The student would be able to conclude the rational use of anti-	gories of affectious mechanisms, tegories of a cancer, the normal per employed			
7	Course  Descriptio  n  aspects (classification, mechanism of action, therapeutic effects, clinical use side effects and contraindications) of drugs acting on respiratory are gastrointestinal system, infectious diseases, immuno-pharmacology and addition, emphasis on the principles of toxicology and chronopharmacology.					
8	8 Outline syllabus CO Mapp					





1		CO1					
1	UNIT-I						
	A. Pharmacology of drugs acting on Respiratory system						
	a. Anti -asthmatic drugs						
	b. Drugs used in the management of COPD						
	c. Expectorants and antitussives						
	d. Nasal decongestants						
	B. 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.						
_	e. Emetics and anti-emetics.						
2	UNIT-II						
	A. Chemotherapy	CO2					
	a. General principles of chemotherapy.	CO6					
	b. Sulfonamides and cotrimoxazole.						
	c.Antibiotics- Penicillins,						
	cephalosporins,						
	ablamamphanical magnelidae animalanee and						
	chloramphenicol, macrolides, quinolones and						
	fluoroquinolins, tetracycline and						
	aminoglycosides						
3	UNIT-III						
	A. Chemotherapy	CO3					
	a. Antitubercular agents						
	b. Antileprotic agents						
	c. Antifungal agents						
	d. Antiviral agents						
	e. Anthelmintic agents						
	f. Antimalarial drugs						
	g. Antiamoebic agents						





			THE THE TAXABLE PARTY OF TAXABLE PARTY O	
4	UNIT-IV A. Chemotherapy 1. Urinary tract infection m. Chemotherapy of m B. Immunopharmacole a. Immunostimulants b. Immunosuppressant Protein drugs, monochiosimilars	alignancy. <b>ogy</b>	mitted diseases.	CO4
5	<ul><li>c. Clinical symptoms organophosphosphopoisoning.</li><li>B. Chronopharmacola. Definition of rhymacola.</li></ul>	knowledge of acute, sion and basic knowledge of the enicity and mutagenion of treatment of poison and management of borus compound and leadings.	edge of genotoxicity, icity ning	CO5
Mode of examinat ion	Theory			
Weighta ge	Continuous Mode Assessment	Sessional Exam	ESE	
Distribut ion	10 Marks	15	75	



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



Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	2	1	3	-	2	-	2	2	3
CO2	3	3	1	1	3	-	1	-	3	1	3
CO3	3	3	1	2	3	-	1	-	3	1	3
CO4	3	2	2	2	3	-	2	_	2	2	3
CO5	3	3	1	1	3	_	1	-	2	1	3
CO6	3	3	2	2	3	-	1	-	3	3	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sc	hool:	SOP	
-	ogramme:	B.Pharm	
	anch:	Semester: VI	
1	Course Code	BP603T	
2	Course Title	Herbal Drug Technology – Theory	
3	Credits	4	
4	Contact Hours	3-1-0	
	(L-T-P)		
_	Course Type	Compulsory	
5	Course Objective	<ul> <li>Upon completion of the course, the student shall be able to</li> <li>1. Understand raw material as source of herbal drugs from cultivation therbal drug product.</li> <li>2. Know the WHO and ICH guidelines for evaluation of herbal drugs.</li> <li>know the herbal cosmetics, natural sweeteners, neutraceuticals.</li> </ul>	0.0
6	Course Outcomes	CO1: Students would be able to define herbal medicine, identify and authentication of herbal materials, describe neutraceuticals and herbal drug interactions CO2: Students would be able to differentiate Indian system of medicine and would be able to describe Stability testing of herbal drugs and explain patenting CO3: Students would be able apply and various identification process and latest technique of phytoconstituents CO4: Students would be able to demonstrate evaluation of drugs according to W.H.O. guidelines. Patenting and regulatory requirements of natural and analyse various phytoconstituents CO5: Students would be able to evaluate various phytoconstituents Herb drug Industry Schedule T-Good manufacturing practices of Indian system medicine CO6: Students would be able to formulate herbs or natural origin drugs raw materials for preparation of cosmetics, excipients, conventional herb formulation and novel dosage forms like Phytosomes	of as
7	Course Description	This subject gives the student the knowledge of basic understanding of herb drug industry, the quality of raw material, guidelines for quality of herb drugs, herbal cosmetics, natural sweeteners, nutraceutical etc. The subject all emphasizes on Good Manufacturing Practices (GMP), patenting ar regulatory issues of herbal drugs	oal so
8	Outline syllab	us CO Mapping	5





		***************************************							
	1	UNIT-I	CO1						
		A. Herbs as raw materials	CO6						
		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							
		B. Biodynamic Agriculture							
		Good agricultural practices in cultivation of medicinal plants							
		including Organic farming. Pest and Pest management in							
		medicinal plants: Biopesticides/Bioinsecticides.							
		C. Indian Systems of Medicine							
		a) Basic principles involved in Ayurveda, Siddha, Unani and							
		Homeopathy							
		a) Preparation and standardization of Ayurvedic formulations viz							
		Aristas and Asawas, Ghutika, Churna, Lehya and Bhasma.							
	2	UNIT-II							
		A. Nutraceuticals	CO2						
		a. General aspects, Market, growth, scope and types of products							
		available in the market. Health benefits and role of							
		Nutraceuticals in ailments like Diabetes, CVS diseases,							
		Cancer, Irritable bowel syndrome and various Gastro							
		intestinal diseases.							
		b. Study of following herbs as health food: Alfaalfa, Chicory,							
		Ginger, Fenugreek, Garlic, Honey, Amla, Ginseng,							
		Ashwagandha, Spirulina							
		B. Herbal-Drug and Herb-Food Interactions: General							
		introduction to interaction and classification. Study of following							
		drugs and their possible side effects and interactions:							
		C. Hypercium, kava-kava, Ginkobiloba, Ginseng, Garlic, Pepper							
1		& Ephedra.							





3	Unit III	CO3
	A. Herbal Cosmetics	
	Sources and description of raw materials of herbal origin used via, fixed oils, waxes, gums colours, perfumes, protective agents, bleaching agents, antioxidants in products such as skin care, hair care and oral hygiene products.	CO6
	B. Herbal excipients:  Herbal Excipients – Significance of substances of natural origin as excipients – colorants, sweeteners, binders, diluents, viscosity builders, disintegrants, flavors & perfumes.	
	C. Herbal formulations :	
	Conventional herbal formulations like syrups, mixtures and tablets and Novel dosage forms like phytosomes	
4	UNIT- IV	
	<b>A.</b> Evaluation of Drugs WHO & ICH guidelines for the assessment of herbal drugs Stability testing of herbal drugs.	CO4
	<ul> <li>B. Patenting and Regulatory requirements of natural products:</li> <li>a. Definition of the terms: Patent, IPR, Farmers right, Breeder's right, Bioprospecting and Biopiracy</li> <li>b. Patenting aspects of Traditional Knowledge and Natural Products. Case study of Curcuma &amp; Neem.</li> </ul>	
	C. Regulatory Issues - Regulations in India (ASU DTAB, ASU DCC), Regulation of manufacture of ASU drugs - Schedule Z of Drugs & Cosmetics Act for ASU drugs.	
5		
	A. General Introduction to Herbal Industry  Herbal drugs industry: Present scope and future prospects.  A brief account of plant based industries and institutions involved in work on medicinal and aromatic plants in India.	CO5
	B. Schedule T – Good Manufacturing Practice of Indian systems of medicine  a. Components of GMP (Schedule – T) and its objectives Infrastructural requirements, working space, storage area, machinery and equipments, standard operating procedures, health and hygiene, documentation and records.	
Mode of	Theory	
examinat		
ion		





Weighta	<b>Continuous Mod</b>	e Sessional Exam	ESE
ge	Assessment		
Distribut ion	10 Marks	15	75
Text	1. Text	book of Pharmacognosy b	y Trease & Evans.
book/s*	2. Text Rob	book of Pharmacognosy ber.	y by Tyler, Brady &
	3. Phai	macognosy by Kokate, Pu	rohit and Gokhale
	4. Esse	ntial of Pharmacognosy by	y Dr.S.H.Ansari
	5. Phai	macognosy & Phytochem	istry by V.D.Rangari
	6. Phar	macopoeal standards f	for Ayurvedic
	Form	nulation (Council of Rese	earch in Indian
	Med	icine & Homeopathy)	
	7. Muk	herjee, P.W. Quality Con	ntrol of Herbal
	Drug	gs: An Approach to	Evaluation of
	Bota	nicals. Business Horizo	ons Publishers,
	New	Delhi, India, 2002.	
_			
Other			
Referenc es			





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	2	2	3	3	1	-	2	1	3
CO2	3	3	-	1	3	3	1	-	3	1	3
CO3	3	3	2	2	3	2	1	-	3	3	3
CO4	3	2	-	1	3	3	2	-	2	1	3
CO5	3	3	1	1	3	3	1	-	2	1	3
CO6	3	3	1	1	2	3	1	-	2	1	3

1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





So	chool:	SOP						
Pı	ogramme:	B.Pharm						
Bı	ranch:	Semester: VI						
1	CourseCode	BP604T						
2	Course Title	Biopharmaceutics & Pharmacokinetics – Theory						
3	Credits	4						
4	Contact	3-1-0						
	Hours							
	(L-T-P)							
	Course Type	Compulsory						
5	Course	Upon completion of the course student shall be able to:						
	Objective	1. Understand the basic concepts in biopharmaceutics and						
		pharmacokinetics and						
		their significance.						
		2. Use of plasma drug concentration-time data to calculate the						
		pharmacokinetic parameters to describe the kinetics of drug absorption,						
		distribution, metabolism, excretion, elimination.						
		3. To understand the concepts of bioavailability and bioequivalence of drug						
		products and their significance.						
		Understand various pharmacokinetic parameters, their significance & applications.						
6	Course	CO1: Students will be able to define and differentiate the meaning of						
U	Outcomes	Biopharmaceutics and Pharmacokinetics						
	Outcomes							
		CO2: Students will be able to plan about basic concepts and importance of plasma						
		drug concentration-time data to calculate the pharmacokinetic parameters to						
		describe the kinetics of drug absorption, distribution, metabolism, excretion,						
		elimination.						
		CO3: Students will be able to categorize, sketch and relate various compartment						
		models and their orientation while learning the parameters involved in the						
		biopharmaceutical expression and infer the findings from such studies.						
		CO4: Students will be able to correlate a study and interpret basic concepts,						
		measurement and calculation of zero order and first order absorption rate constant						
		involved in various biopharmaceutical and pharmacokinetics measurements.						
	CO5: Students will be able to interpret various constraints in developing of							
		for individuals in diseased conditions and compare with the functioning of normal						
		person while incorporating the concept of pharmacokinetic study.						
		CO6: Students will be able to analyze Non linear Pharmacokinetics.						





7	Course Description	armaceutics evelopment, ised therein.				
8	Outline syl	Outline syllabus				
	1	<ul> <li>Unit I</li> <li>Introduction to Biopharmaceutics</li> <li>a. Absorption; Mechanisms of drug absorption through GIT, factors influencing drug absorption though GIT, absorption of drug from Non per oral extra-vascular routes,</li> <li>b. Distribution Tissue permeability of drugs, binding of drugs, apparent, volume of drug distribution, plasma and tissue protein binding of drugs, factors affecting protein-drug binding.</li> <li>c. Kinetics of protein binding, Clinical significance of protein binding of drugs</li> </ul>	Mapping CO1			
	2	<ul> <li>UNIT-II Hours</li> <li>A. Elimination: Drug metabolism and basic understanding metabolic pathways renal excretion of drugs, factors affecting renal excretion of drugs, renal clearance, Non renal routes of drug excretion of drugs</li> <li>B. Bioavailability and Bioequivalence: Definition and Objectives of bioavailability, absolute and relative bioavailability, measurement of bioavailability, <i>in-vitro</i> drug dissolution models, <i>in-vitro-in-vivo</i> correlations,</li> <li>C. bioequivalence studies, methods to enhance the dissolution rates and bioavailability of poorly soluble drugs.</li> </ul>	CO2			
	3	<ul> <li>UNIT- III 10 Hours</li> <li>A. Pharmacokinetics: Definition and introduction to</li> <li>Pharmacokinetics, Compartment models, Non compartment models, physiological models, One compartment open model.</li> <li>(a). Intravenous Injection (Bolus)</li> <li>(b). Intravenous infusion and</li> <li>(c) Extra vascular administrations.</li> <li>B. Pharmacokinetics parameters - KE ,t1/2,Vd,AUC,Ka, Clt and CLR- definitions methods of eliminations,</li> <li>C. understanding of their significance and application</li> </ul>	CO3			





4	ompartment open model. tate drug levels, tnance doses and their	CO4		
5	UNIT- V Nonlinear Pharmacokinetics:  a. Introduction, b. Factors causing Non-linearity. a. Michaelis-menton method of estimating paramet c.Explanation with example of drugs.			
Mode of examinat ion	Theory			
Weighta	<b>Continuous Mode</b>	Sessional Exam	ESE	
ge	Assessment			
Distribut ion	10 Marks	15	75	



*	<b>SHARDA</b>
	UNIVERSITY
www.sharda	Beyond Boundaries

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





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	2	2	3	3	2	-	2	2	3
CO2	3	3	1	1	3	3	1	_	3	1	3
CO3	3	3	2	2	3	2	1	-	3	3	3
CO4	3	2	2	1	3	3	2	-	2	2	3
CO5	3	3	2	2	3	3	1	-	2	3	3
CO6	3	2	1	1	3	3	2	-	2	2	3

1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sc	chool:	SOP			
Pr	ogramme:	B.Pharm			
Bı	ranch:	Semester: VI			
1	Course Code	BP605T			
2	Course Title	Pharmaceutical Biotechnology – Theory			
3	Credits	4			
4	Contact	3-1-0			
	Hours				
	(L-T-P)				
	Course Type	Compulsory			
5	Course Objective	Upon completion of the subject student shall be able to;			
	Objective	1. Understanding the importance of Immobilized			
		enzymes in Pharmaceutical Industries			
		2. Genetic engineering applications in relation to production of			
		pharmaceuticals			
		3. Importance of Monoclonal antibodies in Industries			
		4. Appreciate the use of microorganisms in fermentation technology			
6	Course	CO1: Students will be able to understand the importance of			
	Outcomes	Immobilized enzymes in Pharmaceutical Industries.			
		CO2: Students will be able to describe genetic engineering applications in relation to production of Pharmaceuticals			
		CO3: Students will be able to conclude about importance of Monoclonal antibodies in Industries			
		CO4: Students will be able to understand the use of microorganisms in fermentation technology			
		CO5: Students will be able to plan various fermentation methods			
		CO6: Students will be able to explore recent developments in fermentation technology.			





7	Course Description	Biotechnology has a long promise to revolutionize the biological sciences and technology.				
		• Scientific application of biotechnology in the field of genetic engineering, medicine and fermentation technology makes the subject interesting.				
		• Biotechnology is leading to new biological revolutions in diagnosis, prevention and cure of diseases, new and cheaper pharmaceutical drugs.				
		• Biotechnology has already produced transgenic crops and animals and the future promises lot more.				
		• It is basically a research-based subject.				
8	Outline syll	abus	CO Mapping			
	1	Unit I	CO1			
		a) Brief introduction to Biotechnology with reference to Pharmaceutical Sciences.				
		b) Enzyme Biotechnology- Methods of enzyme immobilization and applications.				
		e) Biosensors- Working and applications of biosensors in Pharmaceutical Industries.				
	Brief introduction to Protein Engineering.					
		Use of microbes in industry. Production of Enzymes- General consideration - Amylase, Catalase, Peroxidase, Lipase, Protease, Penicillinase.				
		Basic principles of genetic engineering.				





2	Unit II	CO2
	Types of immunity- humoral immunity, cellular immunity	CO2
	a). Structure of Immunoglobulins	
	b). Structure and Function of MHC	
	c). Hypersensitivity reactions, Immune stimulation and Immune suppressions.	
	General method of the preparation of bacterial vaccines, toxoids, viral vaccine, antitoxins, serum-immune blood derivatives and other products relative to immunity. Storage conditions and stability of official vaccines	
	Hybridoma technology- Production, Purification and Applications	
	Blood products and Plasma Substituties.	
3	Unit III  a. Types of immunity- humoral immunity, cellular immunity	CO3
	<ul> <li>b. Structure of Immunoglobulins</li> <li>Structure and Function of MHC</li> <li>Hypersensitivity reactions, Immune stimulation and Immune suppressions.</li> <li>c. General method of the preparation of bacterial vaccines, toxoids, viral vaccine, antitoxins, serum-immune blood derivatives and other products relative to immunity.</li> <li>Storage conditions and stability of official vaccines</li> <li>Hybridoma technology- Production, Purification and Applications</li> <li>Blood products and Plasma Substituties.</li> </ul>	
4	Unit IV  a) Immuno blotting techniques- ELISA, Western blotting, Southern blotting.  b) Genetic organization of Eukaryotas and Prokaryotas	CO4
	<ul><li>b) Genetic organization of Eukaryotes and Prokaryotes</li><li>c) Microbial genetics including transformation, transduction, conjugation, plasmids and transposons.</li></ul>	
	Introduction to Microbial biotransformation and applications.	
	Mutation: Types of mutation/mutants.	





5	Unit V								
		quirements, study of ods, aeration process,	CO5 CO6						
	b. Large scale p controls.	<ul> <li>Large scale production fermenter design and its various controls.</li> </ul>							
		production of - penic Glutamic acid, Griseof							
	<b>Blood Products:</b> Col whole human blood Substituties.		<u> </u>						
Mode of examinat ion	Theory								
Weighta ge	Continuous Mode Assessment	Sessional Exam	ESE						
Distribut ion	10 Marks	15	75						
Text book/s*	<ol> <li>B.R. Glick and J.J. Pasternak: Molecular Biotechnology: Principles and Applications of RecombinantDNA: ASM Press Washington D.C.</li> <li>RA Goldshy et. al., : Kuby Immunology.</li> <li>J.W. Goding: Monoclonal Antibodies. Zaborsky: Immobilized Enzymes, CRC Press, Degraland, Ohio.</li> <li>S.B. Primrose: Molecular Biotechnology (Second Edition) Blackwell Scientific Publication.</li> <li>Stanbury F., P., Whitakar A., and Hall J., S., Principles of fermentation technology, 2nd edition, Aditya books Ltd., New Delhi</li> </ol>								





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	2	2	3	3	1	-	2	1	3
CO2	3	2	-	1	3	3	1	-	3	1	3
CO3	3	3	2	2	3	2	1	-	3	3	3
CO4	3	2	-	1	3	3	2	-	2	1	3
CO5	3	3	1	1	3	3	1	-	2	1	3
CO6	3	3	2	2	3	2	1	-	3	3	3

- 1-Slight (Low)
  2-Moderate (Medium)
  3-Substantial (High)





	hool:	SOP					
Pr	ogramme:	B.Pharm					
	anch:	Semester: VI					
1	Course Code	BP606T					
2	Course Title	Pharmaceutical Quality Assurance – Theory					
3	Credits	4					
4	Contact	3-1-0					
	Hours						
	(L-T-P)						
	Course Type	Compulsory					
5	Course	Upon completion, students will be familiar with various aspects of quality					
	Objective	control and quality assurance aspects of pharmaceutical industries. It deals					
		with the important aspects like cGMP, QC tests, documentation, quality					
		certifications and					
		regulatory affairs					
6	Course	CO1: Students will be able to describe about upon completion of the course student					
	Outcomes	shall be able to understand the cGMP aspects in a pharmaceutical industry					
		CO2: Students will be able to associate basic concepts and importance of appreciate the importance of documentation					
		CO3: Students will be able to interpret andunderstand basic concepts on quality certifications applicable to pharmaceutical industries					
		CO4: Students will be able to summarize and understand the responsibilities of QA					
		& QC departments along with GLP and validation aspects					
		CO5: Students will be able to analyze Complaints and evaluation of complaints in addition to Handling of return goods					
		addition to Handing of feturi goods					
		CO6: Students will be able to compare about all the documents used in the Pharmaceutical industry.					
	Course	This course deals with the various aspects of quality control and quality					
7	Description	assurance aspects of pharmaceutical industries. It deals with the important					
	2 coorphon	aspects like cGMP, QC tests, documentation, quality certifications and					
		regulatory					
8	Outline syllabi						
	<b>3</b>	Mapping					





1	THE TAY AND REAL PROPERTY OF THE PROPERTY OF T	
1	UNIT – I	CO1
	A. Quality Assurance and Quality Management concepts:	
	Definition and concept of Quality control, Quality assurance and GMP	
	B. Total Quality Management (TQM): Definition, elements,	
	philosophies	
	<b>C.</b> 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 Programme, tools ISO 9000 & ISO14000:	
	Overview, Benefits, Elements, steps for registration NABL accreditation: Principles and procedures	
	accreditation. Frinciples and procedures	
2	UNIT - II	CO2
	A. Organization and personnel: Personnel responsibilities,	CO2
	training, hygiene and personal records. <b>Premises:</b> Design,	
	construction and plant layout, maintenance, sanitation, environmental control, utilities and maintenance of sterile	
	areas, control of contamination.	
	B. Equipments and raw materials: Equipment selection,	
	purchase specifications, maintenance,	
	C. purchase specifications and maintenance of stores for	
3	raw materials.	
3	UNIT – III	CO3
	<b>A. Quality Control:</b> Quality control test for containers, rubber closures and secondary packingmaterials.	
	B. Good Laboratory Practices: General Provisions,	
	Organization and Personnel, Facilities, Equipment, Testing	
	Facilities Operation, Test and Control Articles, C. Protocol for Conduct of a Nonclinical Laboratory Study,	
	Records and Reports, Disqualification of Testing Facilities	
4	UNIT - IV	CO4
	A. Complaints: Complaints and evaluation of complaints,	CO4 CO6
	Handling of return good, recalling and waste disposal.	
	B. Document maintenance in pharmaceutical industry:	
	Batch Formula Record, Master Formula Record, SOP,	
	Quality audit,  C. Quality Review and Quality documentation, Reports and	
	documents, distribution records.	





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5	general princivalidation, imposes of validation pH meter, Qual General princip  C. Warehousing: management	CO5			
examinat	Theory				
		T	T		
-		Sessional Exam	ESE		
_	Assessment				
	10 Marks	15	75		
			<u> </u>		
	•		•		
			egulations, 2 nd Edition,		
	•	· ·	maceuticals. A		
	compend	ium of Guidelines	and Related		
	5. How to P	Practice GMP's – PP	Sharma.		
	6. ISO 9000	and Total Quality M	lanagement – Sadhank G		
	Ghosh				
	Quality	specification for	Pharmaceutical		
		<u>-</u>	_		
		•			
	Mode of	Weighta ge Assessment  1. Quality Pharmace 2. Good La Sandy W 3. Quality compend materials 4. A guide and Sedh 5. How to F 6. ISO 9000 Ghosh 7. The Inte III, IV-Quality Substance 8. Good lab	A. Calibration and Validation: Introdegeneral principles of calibration validation, importance and scope of validation, validation master pH meter, Qualification of UV-Visil General principles of Analytical methods of the composition of the careful principles of Analytical methods of the composition of the careful principles of Analytical methods of the composition of the careful principles of Analytical methods of the composition of the composit	A. Calibration and Validation: Introduction, definition and general principles of calibration, qualification and validation, importance and scope of validation, and validation, importance and scope of validation.  B. types of validation, validation master plan. Calibration of pH meter, Qualification of UV-Visible spectrophotometer, General principles of Analytical method Validation.  C. Warehousing: Good warehousing practice, materials management  Mode of examination  Mode of examination  Continuous Mode Sessional Exam ESE  Assessment  1. Quality Assurance Guide by organization of Pharmaceutical Products of India.  2. Good Laboratory Practice Regulations, 2nd Edition, Sandy Weinberg Vol. 69.  3. Quality Assurance of Pharmaceuticals- A compendium of Guidelines and Related materials Vol I WHO Publications.  4. A guide to Total Quality Management- Kushik Maitra and Sedhan K Ghosh  5. How to Practice GMP's – P P Sharma.  6. ISO 9000 and Total Quality Management – Sadhank G Ghosh  7. The International Pharmacopoeia – Vol I, II, III, IV- General Methods of Analysis and Quality specification for Pharmaceutical Substances, Excipients and Dosage forms  8. Good laboratory Practices – Marcel Deckker Series	



Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
	3	3	1	2	3	3	1	2	2	1	3
CO1											
	3	3	2	2	3	3	1	1	3	1	3
CO2											
	3	3	2	2	3	2	1	3	3	3	3
CO3											
	3	2	2	1	3	3	2	2	2	1	3
CO4											
	3	3	2	1	3	3	2	3	2	1	3
CO5											
	3	3	2	1	3	3	2	3	2	1	3
CO6											

1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sc	chool:	SOP
Pr	ogramme:	B. Pharm
Bı	ranch:	Semester: V
1	Course	BP607P
	Code	
2	Course	Medicinal Chemistry III - Practical
	Title	
3	Credits	2
4	Contact	0-0-4
	Hours	
	(L-T-P)	
	Course	Compulsory
5	Type Course	
)	Objective	Upon completion of the course student shall be able to
	-	<ol> <li>Understand the importance of drug design and different techniques of drug design.</li> </ol>
		2. Understand the chemistry of drugs with respect to their biological activity.
		3. Know the metabolism, adverse effects and therapeutic value of drugs.
		4. Know the importance of SAR of drugs.
6	Course Outcomes	CO1: Student will be able to understand the structure, chemistry and its correlation with the therapeutic value of drugs.
		CO2: Student will be able to formulate of drugs and preparation of drugs.
		CO3: Student will be able to apply the basic knowledge about the synthesis of sulphanilamide.
		CO4: Student will be able to generalize about the synthesis of Chlorobutanol, metronidazole.
		CO5: Student will be able to conclude about the synthesis of chloroquine, dapsone CO6: Student will be able to plan about the synthesis of Tolbutamide.





7	Course	I	I Preparation of drugs and intermediates									
	Description	1	Sulpha	nilamide								
		2	7-Hydı	roxy, 4-methyl cou	marin							
		3	Chloro	butanol								
		4	Triphe	nyl imidazole								
		6	Hexam	nine								
		II Assay of drugs										
		1	Isonico	otinic acid hydrazid	le							
	2 Chloroquine											
		3	Metror	nidazole, Dapsone								
		4	Chlorp	heniramine maleat	e, Benzyl penicillin							
8	Outline Sylla	bus				СО						
	•					Mapping						
	1	Preparation	of Sulp	hanilamide		CO1						
	2	_		orpheniramine male	eate	CO2						
	3	-		ydroxy, 4-methyl c		CO3						
	4	Preparation			<del></del>	CO4						
	5	Preparation				CO5						
	6	•		henyl imidazole		CO2						
	7			zyl penicillin		CO1						
	8	Preparation	of Prep	paration of Preparat	ion of Tolbutamide	CO3						
						CO6						
	9	Preparation				CO1						
	10	Preparation	of Dap	sone		CO2						
	Mode of	Practical/Vi	Practical/Viva									
	examination											
	Weightage	Continuous	tinuous Sessional Exam ESE									
	Distribution	Mode										
		Assessment										
		05 Marks		10	35							





Text book/s*	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.





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	1	1	3	3	1	2	2	1	3
CO2	3	3	-	1	3	3	1	1	3	1	3
CO3	3	3	2	2	3	2	1	3	3	3	3
CO4	3	2	-	1	3	3	2	2	2	1	3
CO5	3	3	1	1	3	3	1	3	2	1	3
CO6	3	3	2	2	3	2	1	3	3	3	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sc	hool:	SOP						
Pr	ogramme:	B. Pharm						
Br	anch:	Semester: VI						
1	Course	BP608P						
	Code							
2	Course	Pharmacology III- Practical						
	Title							
3	Credits	2						
4	Contact	0-0-4						
	Hours							
	(L-T-P)							
	Course	Compulsory						
_	Туре							
5	Course	Upon completion of this course the student should be able to						
	Objective	1. Understand the mechanism of drug action and its relevance in the						
		treatment of different diseases						
		2. Demonstrate isolation of different organs/tissues from the laboratory						
		animals by simulated experiments 3. Demonstrate the various receptor actions using isolated tissue						
		preparation						
		Appreciate correlation of pharmacology with related medical sciences						
6	Course	CO1: Students would be able to define various instruments and						
	Outcomes	methods used in the evaluation of <i>in vitro</i> and <i>in vivo</i> evaluation of						
		various drugs.						
		CO2: Students would be able to understand the working principles of the						
		instruments used and actions of various drugs on biological systems.						
		CO3: Students would be able to demonstrate the effects of various						
		categories of drugs and bioassays of physiological substances.						
		CO4: Students would be able to analyze the outcomes of experiments						
		through simulation studies.						
		CO5: Students would be able to evaluate the normal and abnormal						
		physiological processes, and various drugs that can be employed for						
		different treatment protocols.						
		CO6: The student will be able to perform bioassays of various drugs.						





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7	Course								
	Description	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							
		<ul><li>6. Estimation of serum biochemical parameters by using autoanalyser</li><li>8. Effect of saline purgative on frog intestine</li></ul>							
		hypoglycemic effect in rabbit	ne Insulin						
		9. Test for pyrogens (rabbit method)							
		10. Determination of acute oral toxicity (LD50) of a drugiven data	g from a						
	11. Determination of acute skin irritation / corrosion of a substance								
		12. Determination of acute eye irritation / corrosion of a	test						
	substance								
		13. Calculation of pharmacokinetic parameters from a given	ven data						
		14. Biostatistics methods in experimental pharmacology( student's t test, ANOVA)							
		15. Biostatistics methods in experimental pharmaco square test, Wilcoxon Signed Rank test)	nogy (Cm						
8	Outline Sylla	l hus	СО						
			Mapping						
	1	Dose calculation in pharmacological experiments	CO1						
	2	Study of anti-ulcer activity of a drug using pylorus ligand (SHAY) rat model and NSAIDS induced ulcer model.	CO2						
	3	Study of effect of drugs on gastrointestinal motility	CO3						
	4	Effect of agonist and antagonists on guinea pig ileum	CO4						
	5	Estimation of serum biochemical parameters by using semi-	CO5						
		autoanalyser	CO6						
	6	Estimation of serum biochemical parameters by using semi-	CO1						
		autoanalyser	CO6						
	7	Effect of saline purgative on frog intestine	CO2						
	8	Insulin hypoglycemic effect in rabbit	CO3						
	9	Test for pyrogens ( rabbit method)	CO4						
	10	Determination of acute oral toxicity (LD50) of a drug from a	005						
		given data	CO5						





Mode of examination	Practical/Viva								
Weightage	Continuous	Sessional Exam	ESE						
Distribution	Mode								
	Assessment								
	05 Marks	10	35						
'Text book/s*	Pharmacolog Pharmacolog Charles R.Cr 2. Ghosh M Pharmacolog 3. Kulkarni VallabhP 4. N.Udupa	y, Paras medical y with clinical raig& Robert, MN. Fundamental y. Hilton & Compa	experimental pharmacology.						
Other									
References									



Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	2	1	3	3	3	2	2	3	3
CO2	3	3	3	1	3	3	3	1	3	3	3
CO3	3	3	3	2	3	2	1	3	3	3	3
CO4	3	2	3	1	3	3	2	2	2	2	3
CO5	3	3	2	1	3	3	1	3	2	2	3
CO6	3	2	-	1	3	3	2	2	2	1	3

- 1. Slight (Low) 2-Moderate (Medium) 3-Substantial (High)



Sc	hool:	SOP
Pr	ogramme:	B. Pharm
	anch:	Semester: VI
1	Course	BP609P
	Code	
2	Course	Herbal Drug Technology - Practical
	Title	S Si
3	Credits	2
4	Contact	0-0-4
	Hours	
	(L-T-P)	
	Course	Compulsory
	Type	
5	Course	1. Explain correct use of various equipments in Pharmacognosy
	Objective	laboratory.
		2. Evaluation of drugs and various formulations
		3. Demonstrate various herbal preparations
		4. Formulations of various herbal cosmetics
		5. Analyse the phytoconstituents
6	Course Outcomes	CO1: Students would able to prepare crude drug extract, identify extract through phytochemical screening and also identify herbal drugs through chemical test CO2: Students would be able to distinguish excipients of natural origin and estimate alcohol content in alcoholic formulations
		CO3: Students would be able to formulate herbal creams, shampoo CO4: Students would be able to analyze evaluation parameters for herbal shampoo and creams CO5: Students would be able to formulate and evaluate syrups. CO6: Students will be able to understand about herbal drug preparations.
		Coo. Students will be able to understand about herbal drug preparations.
7	Course Description	<ol> <li>To perform preliminary phytochemical screening of crude drugs.</li> <li>Determination of the alcohol content of Asava and Arista</li> <li>Evaluation of excipients of natural origin</li> <li>Incorporation of prepared and standardized extract in cosmetic formulations like creams, lotions and shampoos and their evaluation.</li> <li>Incorporation of prepared and standardized extract in formulations like syrups, mixtures and tablets and their evaluation as per Pharmacopoeial requirements.</li> <li>Monograph analysis of herbal drugs from recent Pharmacopoeias</li> <li>Determination of Aldehyde content</li> </ol>
		8. Determination of Phenol content
		Determination of total alkaloids





8	Outline Sylla	bus			СО				
					Mapping				
	1	To determine phen	ol content in clove	oil.	CO1				
	2	To determine total	To determine total alkaloids in crude drug sample.						
	3	Evaluation of excip	pients of natural ori	gin	CO3				
	4	Incorporation of pr	Incorporation of prepared and standardized extract in cosmetic formulations like creams, lotions and shampoos and their evaluation.						
	5	To perform Prelim	inary phytochemica	al screening of crude drugs.	CO5				
	6	To determine the a	lcohol content of A	sava and Arishta.	CO1				
	7	To formulate and e longa.	CO2 CO6						
	8	To formulate and e	CO3 CO6						
	9	To formulate and e	To formulate and evaluate herbal cough syrup						
	10	To perform Prelim	inary phytochemica	al screening of crude drugs.	CO2				
	Mode of examination	Practical/Viva	V 1 V						
	Weightage Distribution	Continuous Mode Assessment	Sessional Exam	ESE					
		05 Marks	10	35					
	Text		ook of Pharmacog	nosy by Trease & Evans.					
	book/s*	2. Textbook of Pharmacognosy by Tyler, Brady & Robber.							
		3. Phari	nacognosy by Kok	ate, Purohit and Gokhale					
		4. Esser	ntial of Pharmacogn	nosy by Dr.S.H.Ansari					
		5. Pharr	nacognosy & Phyto	ochemistry by V.D.Rangari					



Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	1	1	3	3	1	2	2	1	3
CO2	3	3	2	1	3	3	1	1	3	1	3
CO2	3	3	2	1	3	3	1	1	3	1	3
CO3	3	3	2	2	3	2	1	3	3	3	3
CO4	3	2	2	1	3	3	2	2	2	1	3
CO5	3	3	1	1	3	3	1	3	2	1	3
CO6	3	2	2	1	3	3	2	2	2	1	3

-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sch	ool:	SOP					
Pro	gramme:	B.Pharm					
Bra	nch:	Semester: VII					
1	Course Code	BP701T					
2	Course Title	Instrumental Methods of Analysis					
3	Credits	4					
4	Contact Hours (L-T-P)	3-1-0					
	Course Type	Compulsory					
5	Course Objective	Upon completion of this course the student should be able to 1. Understand the interaction of matter with electromagnetic radiations and its applications in drug analysis. 2. Understand various techniques in Analysis of various Pharmaceuticals. 3. Study the applications of various Instruments in analysis of Pharmaceuticals. 4. Perform quantitative & qualitative analysis of drugs using various analytical instruments.					
6	Course	CO1: Students will be able to understand about Analytical techniques in					
	Outcomes	Instrumental Methods of Analysis.  CO2 Students will be able to apply the principle of Modern instruments using in Analysis of Pharmaceuticals  CO3: Students will be able to compare about applications of Modern instruments using in Analysis of Pharmaceuticals  CO4: Students will be able to identify how to operate the Modern instruments in analysis of Pharmaceuticals.  CO5: Students will be able to interpret the chromatographic separation and analysis of drugs.  CO6: Students will be able to analyze the chromatographic separation and analysis of drugs					
7	Course Description	This subject deals with the application of instrumental methods in qualitative and quantitative analysis of drugs. This subject is designed to impart a fundamental knowledge on the principles and instrumentation of spectroscopic and chromatographic technique. This also emphasizes on theoretical and practical knowledge on modern analytical instruments that are used for drug testing.					
8	Outline syllabus	S CO Mapping					





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	1	A. UV Visible spectroscopy	G 0.4
		Electronic transitions, chromophores, auxochromes,	CO1
		spectral shifts, solvent effect on absorption spectra,	
		Beer and Lambert's law, Derivation and deviations.	
		<b>B.</b> Instrumentation - Sources of radiation,	
		wavelength selectors, sample cells, detectors- Photo	
		tube, Photomultiplier tube, Photo voltaic cell, Silicon	
		Photodiode.	
		<b>C. Applications -</b> Spectrophotometric titrations,	
		Single component and multi component analysis	
		D.Fluorimetry	
	2	A. IR spectroscopy	CO2
		Introduction, fundamental modes of vibrations in	
		poly atomic molecules, sample handling, factors	
		affecting vibrations	
		Instrumentation - Sources of radiation, wavelength	
		selectors, detectors - Golay cell, Bolometer,	
		Thermocouple, Thermister, Pyroelectric detector and	
		applications	
		<b>B. Flame Photometry-</b> Principle, interferences,	
		instrumentation and applications	
		C. Atomic absorption spectroscopy- Principle,	
		interferences, instrumentation and applications	
		Nepheloturbidometry- Principle,	
	2	instrumentation and applications	G02
	3	A. Introduction to chromatography	CO3
		Adsorption and partition column	
		chromatography-Methodology,	
		advantages, disadvantages and applications.	
		B. Thin layer chromatography- Introduction,	
		Principle, Methodology, Rf values, advantages,	
		disadvantages and applications.	
		C. Paper chromatography-Introduction,	
		methodology, development techniques,	
		advantages, disadvantages and applications	
		Electrophoresis— Introduction, factors affecting	
		electrophoretic mobility, Techniques of paper,	
		gel, capillary electrophoresis, applications	
	4	A. Gas chromatography - Introduction, theory,	
		instrumentation, derivatization,	CO4
		temperature Programmeming, advantages,	
		disadvantages and applications	
		B. High performance liquid chromatography	
		(HPLC)-Introduction, theory,	
		<b>C.</b> instrumentation, advantages and applications.	





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Mode 5	classificat mechanism affecting application B. Gel chroinstrumen C. chromato	<ul> <li>A. Ion exchange chromatography- Introduction, classification, ion exchange resins, properties, mechanism of ion exchange process, factors affecting ion exchange, methodology and applications</li> <li>B. Gel chromatography- Introduction, theory, instrumentation and applications Affinity</li> <li>C. chromatography- Introduction, theory, instrumentation and applications</li> <li>Theory</li> </ul>					
examination							
Weightage Distribution	Continuous Mode Assessment	Sessional Exam	ESE				
	10 Marks	15	75				
Text book/s*			<u>I</u>				
Other	Recommende	ed Books (Lat	test Editions)				
References		nental Metho	ds of Chemical Analysis				
	2. Organi	ic spectroscop	y by Y.R Sharma				
		book of Phar A. Connors	rmaceutical Analysis by				
	_	's Text book by A.I. Vogel	of Quantitative Chemical				
		cal Pharmaceu nd J.B. Stenlak	tical Chemistry by A.H.				
	6. Organi	ic Chemistry b	y I. L. Finar				
	7. Organi	ic spectroscop	y by William Kemp				
	8. Quanti Garrett						
	`	9. Quantitative Analysis of Drugs in Pharmaceutical Formulations by P. D. Sethi					
	_	ophotometric ds by Silverste	identification of Organic ein				





Pos Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1	3	2	3	3	1	2	2	2	2	2	1
CO2	3	2	3	3	1	2	2	2	1	2	2
CO3	3	2	3	3	2	2	2	2	2	2	2
CO4	3	1	3	3	1	1	2	1	1	2	2
CO5	3	2	3	3	2	1	2	1	2	1	1
CO6	3	1	2	3	2	3	2	3	2	1	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sch	ool:	SOP					
Pro	gramme:	B.Pharm					
Bra	nch:	Semester: VII					
1	Course Code	BP-702T					
2	Course Title	Industrial Pharmacy-II Theory					
3	Credits	4					
4	Contact Hours (L-T-P)	3-1-0					
	Course Type Course	Compulsory					
5	Objective Objective	Upon completion of the course, the student shall be able 1. Know the process of pilot plant and scale up of phart forms 2. Understand the process of technology transfer from 1 commercial batch 3. Know different Laws and Acts that regulate pharmac 4. Understand the approval process and regulatory required products	maceutical dosage lab scale to leutical industry				
6	Course Outcomes	and scale up of pharmaceutical dosage forms  CO2: Students shall be able to describe the process of from lab scale to commercial batch.  CO3: Students shall be able to plan about stepwise process from NDA filing to final FDA submission  CO4 Students shall be to able to analyze the different regulate pharmaceutical industry in India and US  CO5: Students shall be to able to Develop the comanagement and knowledge of required certifications  CO6: Students will be able to design about all the processor.	O2: Students shall be able to describe the process of technology transfer om lab scale to commercial batch.  O3: Students shall be able to plan about stepwise product development ocess from NDA filing to final FDA submission  O4 Students shall be to able to analyze the different laws and acts that gulate pharmaceutical industry in India and US  O5: Students shall be to able to Develop the concept of quality anagement and knowledge of required certifications				
7	Course	requirements for drug product.  This course is designed to impart fundamental knowled	las on pharmacoutical				
/		product development and translation from laboratory to i					
8	Outline syllabi	ls .	CO Mapping				
	Unit 1	A. Pilot plant scale up techniques:					





B. Pilot plant scale up techniques: General considerations - including	CO1
significance of personnel requirements, space requirements, raw	COI
materials, Pilot plant scale up considerations for solids, liquid	
orals, semi solids	
C. relevant documentation, SUPAC guidelines, Introduction to	
platform technology.	
Unit 2 Technology development and transfer	
A. Technology development and transfer: WHO guidelines for	CO2
Technology Transfer(TT): Terminology, Technology transfer	
protocol,	
B. 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 and equipments, qualification and	
validation, quality control, analytical method transfer,	
C. 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	
Unit 3 Regulatory affairs	
	CO2
A. Regulatory affairs: Introduction, Historical overview of	CO3
Regulatory Affairs, Regulatory authorities, Role of Regulatory	
affairs department,	
B. Responsibility of Regulatory Affairs Professionals Regulatory	
requirements for drug approval: Drug Development Teams, Non-	
Clinical Drug Development, Pharmacology, Drug Metabolism	
and Toxicology, General considerations of Investigational New	
Drug (IND) Application,	
C. Investigator's Brochure (IB) and New Drug Application (NDA),	
Clinical research / BE studies, Clinical Research Protocols,	
Biostatistics in Pharmaceutical Product Development, Data	
Presentation for FDA Submissions, Management of Clinical	
Studies	
Unit 4 Quality management systems	
A. Quality management systems: Quality management &	CO4
Certifications: Concept of Quality,	
B. Total Quality Management, Quality by Design (QbD), Six Sigma	
concept, Out of Specifications (OOS),	
C. Change control, Introduction to ISO 9000 series of quality	
systems standards, ISO 14000, NABL, GLP	
Unit 5 Indian Regulatory Requirements	





 www.shatua.ac.iii								
A. Indian Reg	CO5							
Organizatio	CO6							
		ilities, Certi	ficate of Pharmaceutical					
Product (C	, .							
		approval pro	ocedures for New Drugs					
Mode of	Theory							
examination		<b>,</b>						
Weightage	Continuous	Sessiona	ESE					
Distribution	Mode	1 Exam						
	Assessment							
	10 Marks	15	75					
Text	1. Regulatory A	ffairs from	Wikipedia, the free					
book/s	encyclopedia mo	dified on 7th	h April available at					
	http,//en.wikiped	ia.org/wiki/F	Regulatory_					
	Affairs.							
		•	Affairs Updates,					
	2005.	availal						
	http://www.iraup		<del></del>					
	_		Pavid S. Mantus. Text					
		•	Affairs A Guide for					
	<u> </u>	gs, Medical l	Devices, and Biologics'					
	Second Edition.		1.1.1.1					
		`	ght by learning plus,					
	inc. available at l	nttp.//www.c	gmp.com/ra.htm.					





CO/PO	PO1	PO 2	PO 3	PO 4	PO 5	PO 6	PO 7	P0 8	PO 9	PO1 0	PO1 1
CO1	3	2	2	2	2	3	2	_	2	2	2
CO2	3	2	1	2	2	3	2	2	3	1	2
CO3	3	2	2	2	2	2	3	2	2	1	2
CO4	3	1	1	1	1	2	3	_	3	2	2
CO5	3	1	2	1	1	3	2	-	3	2	3
CO6	3	2	2	2	2	3	2	_	3	2	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sch	ool:	SOP
Pro	gramme:	B. Pharm
	nch:	Semester:VII
1	Course Code	BP703T
2	Course Title	Pharmacy Practice Theory
3	Credits	4
4	Contact Hours (L-T-P)	3-1-0
	Course Type	Compulsory
5	Course Objective	This course has been designed to impart the fundamental knowledge of pharmacy practice and ethics along with the aspects of hospital organization.  Objectives: Upon the completion of this course the students shall be able to  1. know various drug distribution methods in a hospital
		appreciate the pharmacy stores management and inventory control
		appreciate the pharmacy stores management and inventory control     monitor drug therapy of patient through medication chart review and clinical review
		4. obtain medication history interview and counsel the patients
		5. identify drug related problems
		6. detect and assess adverse drug reactions
		7. interpret selected laboratory results (as monitoring parameters in therapeutics) of specific disease states
		8. know pharmaceutical care services
		9. do patient counselling in community pharmacy;
		10. Appreciate the concept of rational drug therapy.
6	Course	CO1: Student will be able to understand Ability to discuss the controversies
	Outcomes	in drug therapy CO2: Student will be able to apply the therapeutic approach to management of hospital CO3: Student will be able to identify the patient specific parameters relevant in monitoring therapy CO4: Student will be able to conclude the importance of individualized therapeutic plans based on diagnosis CO5: Student will be able to analyze data collected at their research work. CO6: Students shall be able to generalize the role of a Pharmacist in a community.





7	Course	In the changing scenario of pharmacy practice in	India for successful
,	Description	practice of Hospital Pharmacy, the students are requ	
	Description	skills like drug distribution, drug information, a	
		monitoring for improved patient care. In community pl	
			• .
		be learning various skills such as dispensing of drugs	
		ailments by providing suitable safe medication, pa	itient counselling for
	0 11 11 1	improved patient care in the community set up.	COM :
8	Outline syllabu		CO Mapping
	Unit 1	Hospital and it's organization	
	<b>A.</b> Definition	on, Classification of hospital- Primary, Secondary and	
	Tertia	ary hospitals, Classification based on clinical and non-	CO1
	clinic	al basis, Organization	
	Structure of a	Hospital, and Medical staffs involved in the hospital	
	and their functi	ions.	
	<b>B.</b> Definition, f	functions of hospital pharmacy, Organization structure,	
	Location, La	ayout and staff requirements, and Responsibilities and	
	functions o	f hospital pharmacists Classifications - Excessive	
		gical effects, secondary pharmacological effects,	
		, allergic drug reactions, genetically determined	
		xicity following sudden withdrawal of drugs, Drug	
		beneficial interactions, adverse interactions, and	
		netic drug interactions, Methods for detecting drug	
		spontaneous case reports and record linkage studies,	
		e drug reaction reporting and management.	
		oraș reaction reporting and management.	
	<b>D.</b> Organization	and structure of retail and wholesale drug store, types	
		Legal requirements for establishment and maintenance	
		ore, Dispensing of proprietary products, maintenance of	
		etail and wholesale drug store.	
	Unit 2	Drug distribution system in a hospital	
	l .	v i	





pati	i Commun ients.	nication skills- communication with prescribers and	
and	Prescribed 1		
cou pha exte Coo the	inseling, ararmacist in ernal training de of ethics interdepartment.		
info	ormation, (ormation.	Orug and Poison information centre, Sources of drug Computerised services, and storage and retrieval of on of patient counseling; steps involved in patient	
con	nmittee in i scription,	on, functions, Policies of the pharmacy and therapeutic including drugs into formulary, inpatient and outpatient automatic stop order, and emergency drug list	CO3
Uni	it 3	Pharmacy and therapeutic committee	CO2
Fin	medica ancial, mate		
C.		for the patient medication history interview,	
		ence, and monitoring of patient medication adherence.	
		scenario for Therapeutic Drug Monitoring. Causes of ation non-adherence, pharmacist role in the medication	
В.	consid	for Therapeutic Drug Monitoring, Factors to be lered during the Therapeutic Drug Monitoring, and	
fori	finition, con mulary and etion of dru		
A.	system ambula	sing of drugs to inpatients, types of drug distribution as, charging policy and labelling, Dispensing of drugs to atory patients, and Dispensing of controlled drugs.	CO2





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	A. Budget prep	paration and imp	lementation.		CO4			
	functions and monitoring -	responsibilities medication char Ward round p	ept of clinical pharmacy, narmacist, Drug therapy ical review, pharmacist Medication history and					
	Dosing pattern pattern.	and drug therap	oy based on Ph	armacokinetic & disease				
		on and sale of the counter medi		ter, and Rational use of				
1	Unit 5	Drug store ma	nagement and	l inventory control				
	conditions, Purprocedure, puro quantity, Reord the drug expendenticon, identicon committee.  B. Description identification, in  C. Blood chem	rchase and inchase order, productive description in the control of	ventory controcurement and sel, and Methodation, principles of hospital as involved, pharmacist, adv	visory committee.	CO5 CO6			
	Mode of examination	Theory	Theory					
	Weightage	Continuous	Sessional	ESE				
	Distribution	Mode	Exam	ESE				
	2 15410 441011	Assessment						
		10 Marks	15	75				
,	Text book/s*							



	Other	1. Merchant S.H. and Dr. J.S.Quadry. A
	References	textbook of hospital pharmacy, 4th ed.
		Ahmadabad: B.S. Shah Prakakshan; 2001.
		2. Parthasarathi G, Karin Nyfort-Hansen,
		Milap C Nahata. A textbook of Clinical
		Pharmacy Practice- essential concepts and
		· · · · · · · · · · · · · · · · · · ·
		skills, 1 st ed. Chennai: Orient Longman
		Private Limited; 2004.
		3. William E. Hassan. <i>Hospital pharmacy</i> , 5th
		ed. Philadelphia: Lea & Febiger; 1986.
		4. Tipnis Bajaj. <i>Hospital Pharmacy</i> , 1 St ed.
		Maharashtra:
		Career Publications; 2008.
		5. Scott LT. Basic skills in interpreting laboratory
		data, 4thed. American Society of Health System
		Pharmacists Inc; 2009.
		5. Parmar N.S. Health Education and Community
		Pharmacy, 18th ed. India: CBS Publishers &
		Distributers; 2008.
		Distributors, 2000.
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СО/РО	PO1	PO2	PO 3	PO4	PO 5	P O 6	PO7	P0 8	PO 9	PO10	PO11
CO1	3	2	2	2	2	-	2	-	2	2	2
CO2	3	2	1	2	2	-	2	-	3	1	2
CO3	3	2	2	2	2	-	2	-	2	1	1
CO4	3	1	1	2	1	-	3	-	3	2	1
CO5	3	1	2	2	2	-	2	-	3	2	3
CO6	3	1	2	1	1	-	2	-	2	2	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





School: SOP							
Pro	gramme:	B. Pharm					
Bra	nch:	Semester: VII					
1	Course Code	BP 704T					
2	Course Title	Novel drug delivery systems- Theory					
3	Credits	4					
4	Contact Hours (L-T-P)	3-1-0					
	Course Type	Compulsory					
5	Course Objective	<ol> <li>After the successful completion of this course, the students.</li> <li>The course aims to provide an understanding of the area of novel drug delivery systems.</li> <li>To understand various approaches for developments.</li> <li>To understand the criteria for selection of drugs the development of Novel drug delivery systems, the evaluation</li> </ol>	nent of novel drug s and polymers for				
6	Course Outcomes	CO1: The students will understand the concepts and Novel Drug Delivery Systems and to study various sustained and controlled drug delivery systems.  CO2: The student will be able to apply knowledge in novel formulations as per requirements and to Implantable drug delivery.  CO3: The student will be able to analyze var parameters for oral, parenteral, topical etc. drug deliver CO4: The students will be able to formulate indust effective strategy for development of new dosage form CO5: The students will be able to plan about site specific CO6: The Students will be able to develop ocular drug and challenges, drug selection.	developing various learn mucosal and rious evaluation ry systems. trially feasible, cost as fic drug delivery.				
7	Course Description	This subject is designed to impart basic knowledge of drug delivery systems.  1. To understand various approaches for development of understand the criteria for selection of design for the development of Novel drug delivery formulation and evaluation	elopment of novel				
8	Outline syllabus	<u></u>	CO Mapping				





	WWW.лицио.	
	A. Controlled drug delivery systems: Introduction, terminology/definitions and rationale, advantages, disadvantages, selection of drug candidates. Approaches to design controlled release formulations based on diffusion, dissolution and ion exchange principles. Physicochemical and biological properties of drugs relevant to controlled release formulations	CO1
	<b>B. Polymers:</b> Introduction, classification, properties,	
	<b>C.</b> advantages and application of polymers in formulation of controlled release drug delivery systems.	
	<ul> <li>A. Microencapsulation: Definition, advantages and disadvantages, microspheres /microcapsules, microparticles, methods of microencapsulation, applications</li> <li>B. Mucosal Drug Delivery system: Introduction, Principles of bioadhesion / mucoadhesion, concepts, advantages and disadvantages, transmucosal permeability and formulation considerations of buccal delivery systems</li> </ul>	CO2
	C. Implantable Drug Delivery Systems:Introduction, advantages and disadvantages, concept of implantsand osmotic pump	
3	<ul> <li>A. Transdermal Drug Delivery Systems:         <ul> <li>Introduction, Permeation through skin, factors affecting permeation, permeation enhancers, basic components of TDDS, formulation approaches</li> </ul> </li> <li>B. Gastroretentive drug delivery systems:         <ul> <li>Introduction, advantages, disadvantages</li> </ul> </li> <li>C. approaches for GRDDS – Floating, high</li> </ul>	CO3
	density systems, inflatable and gastroadhesive systems and their applications	





		1			I-Macazonici)			
	4	approac <b>B.</b> introduction	approaches advantages and disadvantages, <b>B.</b> introduction to liposomes, niosomes, nanoparticles,					
	5	intra ocu -Prelimir ocuserts <b>B. Intrauter</b> Introducti <b>C.</b> developm	<ul> <li>A. Ocular Drug Delivery Systems: Introduction, intra ocular barriers and methods to overcome —Preliminary study, ocular formulations and ocuserts</li> <li>B. Intrauterine Drug Delivery Systems: Introduction, advantages and disadvantages,</li> <li>C. development of intra uterine devices (IUDs) and applications</li> </ul>					
	Mode of examination	Theory						
1	Weightage Distribution	Continuous Mode Assessment	Sessional Exam	ESE 75				
	Text book/s*	TOWAND	10	1.0				
	Other References	1. Y W System expand 2. Marce 3. Robins Contro Marce 4. Inc., N 5. Encycl Delive Publis 6. Interse Analys	Recommended Books (Latest Editions)  1. Y W. Chien, Novel Drug Delivery Systems, 2nd edition, revised and expanded,  2. Marcel Dekker, Inc., New York, 1992.  3. Robinson, J. R., Lee V. H. L, Controlled Drug Delivery Systems, Marcel Dekker,  4. Inc., New York, 1992.  5. Encyclopedia of Controlled Delivery. Edith Mathiowitz, Published by Wiley  6. Interscience Publication, John Quantitative Analysis of Drugs in Pharmaceutical Formulations by P. D. Sethi					



Pos Cos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
CO1	3	2	3	3	1	2	2	-	2	2	1
CO2	3	2	3	3	1	2	2	-	1	2	2
CO3	3	2	3	3	2	2	2	-	2	2	1
CO4	3	2	3	3	1	2	2	-	1	2	1
CO5	3	1	3	3	1	1	2	_	2	1	1
CO6	3	1	3	3	1	1	2	_	2	1	1

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sch	ool:	SOP						
<b>Programme:</b>		B.Pharm						
Bra	nch:	Semester: VII						
1	Course Code	BP705P						
2	Course Title	Instrumental Methods of Analysis- theory						
3	Credits	2						
4	Contact Hours (L-T-P)	0-0-4						
	Course Type	Compulsory						
5	Course Objective	Upon completion of this course the student should be able to 1. Understand applications of instruments in drug analysis 2. Understand operation of instruments in Analysis of various Pharmaceuticals. 3. Study the preparation of various analytes 4. Perform quantitative & qualitative analysis of drugs using various analytical instruments.						
6	Course Outcomes	CO1: Students will be able to understand Analytical techniques in Instrumental Methods of Analysis.  CO2: Students will be able to describe conceptual knowledge about operation of Modern instruments using in Analysis of Pharmaceuticals  CO3: Students will be able to apply the basic knowledge about applications of Modern instruments using in Analysis of Pharmaceuticals  CO4: Student will be able to plan about how to operate the Modern instruments in analysis of Pharmaceuticals.  CO5: Students will be able to understand the chromatographic separation and analysis of drugs.  CO6 Students will be able to interpret the chromatographic separation and analysis of drugs.						





7	Course Description	Subject covers operation of various modern instrument Pharmaceuticals  1. Determination of absorption maxima and ef absorption maxima of organic compounds.  2. Estimation of dextrose by colorimetry  3. Estimation of sulfanilamide by colorimetry  4. Simultaneous estimation of ibuprofen and paspectroscopy  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 neg  11 Separation of amino acids by paper chromatogra  12 Separation of sugars by thin layer chromatogra  13 Separation of plant pigments by column chroma  14 Demonstration experiment on HPLC  15 Demonstration experiment on Gas Chromatogra	s used in analysis of fect of solvents on paracetamol by UV to helo turbidometry aphy why
8	Outline syllabi	us	CO Mapping
	1	Determination of absorption maxima and effect of solvents on absorptionmaxima of organic compounds.	CO1 CO3
	2	Estimation of dextrose by colorimetry	CO2 CO6
	3	Estimation of sulfanilamide by colorimetry	CO1 CO2
	4	Simultaneous estimation of ibuprofen and paracetamol by UV spectroscopy.	CO2 CO4
	5	Assay of paracetamol by UV- Spectrophotometry	CO1 CO3
	6	Estimation of quinine sulfate by fluorimetry.	CO1 CO5
	7	Study of quenching of fluorescence	CO2
	8	Determination of sodium by flame photometry	CO3
	9	Determination of potassium by flame photometry	CO2 CO5
	10	Determination of chlorides and sulphates by nephelo turbidometry	CO1





11	Separation of	CO3 CO4				
12	Separation of	sugars by this	n layer chromatography	CO2 CO4		
Mode of examination	Practical					
Weightage Distribution	Continuous Mode Assessment	Sessional Exam	ESE			
Text book/s*	05 Marks	10	35			
Text book/s**						
Other References	1. Instruments by B.F.					
	2. Organ	2. Organic spectroscopy by Y.R Sharma				
	3. Text Kenne					
	4. Vogel Analy					
		5. Practical Pharmaceutical Chemistry by A.H. Beckett and J.B. Stenlake				
	6. Organ	ic Chemistry	by I. L. Finar			
	7. Organ	ic spectrosco	py by William Kemp			
	8. Quant Garret	•	vsis of Drugs by D. C.			
	9. Quant Pharm Spectrophotor Compounds b	naceutical For metric ider	alysis of Drugs in mulations by P. D. Sethi ntification of Organic			





Pos Cos	PO1	PO2	PO	PO	PO5	PO6	PO7	PO8	PO9	PO10	PO11
			3	4							
CO1	3	1	3	2	2	1	2	-	2	2	2
CO2	3	2	3	3	2	1	2	-	2	2	2
CO3	3	1	3	3	2	1	2	-	2	2	2
CO4	3	2	3	2	2	2	2	-	2	2	2
CO5	3	1	3	3	2	1	1	-	1	2	1
CO6	2	2	1	3	1	3	2	-	2	2	2

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)



Sch	ool:	SOP					
	gramme:	B.Pharm					
	nch:	Semester: VIII					
1	Course Code	BP801T					
	C mid						
2	Course Title	Biostatistics & Research Methodology- Theory					
3	Credits	4					
4	Contact Hours	3-1-0					
	(L-T-P)	Commulator					
5	Course Type Course	Compulsory	®				
3	Objective	1. Know the operation of M.S. Excel, SPSS, I	R and MINITAB $^{\otimes}$ ,				
	Objective	DoE (Design of Experiment)					
		2. Know the various statistical techniques t	o solve statistical				
		problems					
		3. Appreciate statistical techniques in solving the	problems				
6	Course	CO1:Students will be able to Describe statistics,	Biostatistics and				
	Outcomes	interpretation of frequency distribution table, and their pharmaceutical					
		examples					
		CO2: Students will be able to Calculate the measures					
		and dispersion of a data and describe the method					
		including a discussion of advantages, disadvantage	ges, and necessary				
		assumptions.	CD :				
		CO3: Students will be able to Describe the proper					
		Curve fitting, Multiple regression and their pharmaceutical examples  CO4: Students will be able to Calculate and interpret the correlation					
		between two variables.	pret the correlation				
		CO5: Students will be able to Understand the concep	t of Probability and				
		different types of distributions, Poisson's distribution a	•				
		CO6: Students will be able to plan about the concep					
		different types of distributions, Poisson's distribution a					
7	Course	To understand the applications of Biostatics in Phar					
	Description	deals with descriptive statistics, Graphics, Corre	•				
		logistic regression Probability theory, Sampling tec					
		tests, Non Parametric tests, ANOVA, Introduction	ion to Design of				
		Experiments, Phases of Clinical trials and					
		Experimental studies, SPSS, R and MINITAB sta	atistical software's,				
		analyzing the statistical data using Excel.					
8	Outline syllabus	S	CO Mapping				





1	Unit-I	CO1
	<b>A. Introduction:</b> Statistics, Biostatistics,	
	Frequency distribution	
	B. Measures of central tendency: Mean,	
	Median, Mode- Pharmaceutical examples	
	C. Measures of dispersion: Dispersion, Range,	
	standard deviation, Pharmaceutical problems	
	<b>D. Correlation:</b> Definition, Karl Pearson's	
	coefficient of correlation,	
	Multiple correlation - Pharmaceuticals examples	
2	Unit-II	CO2
	<b>A. Regression:</b> Curve fitting by the method of	
	least squares, fitting the lines $y=a + bx$ and $x$	
	= a + by, Multiple regression, standard error	
	of regression–Pharmaceutical Examples	
	<b>B. Probability:</b> Definition of probability,	
	Binomial distribution, Normal distribution,	
	Poisson's distribution, properties - problems	
	Sample, Population, large sample, small	
	sample, Null hypothesis, alternative	
	hypothesis, sampling, essence of sampling,	
	types of sampling, Error-I type, Error-II type,	
	Standard error of mean (SEM) -	
	Pharmaceutical examples	
	C. Parametric test: t-test(Sample, Pooled or	
	Unpaired and Paired), ANOVA, (One way	
	and Two way), Least Significance difference	
3	Unit-III 10 Hours	CO3
	A. Non Parametric tests: Wilcoxon Rank Sum	
	Test, Mann-Whitney U test, Kruskal-Wallis test,	
	Friedman Test	
	<b>B.Introduction to Research:</b> Need for research,	
	Need for design of Experiments, Experiential Design	
	Technique, plagiarism	
	C. Graphs: Histogram, Pie Chart, Cubic Graph,	
	response surface plot, Counter Plot graph	
	<b>Designing the methodology:</b> Sample size	
	determination and Power of a study, Report writing	
	and presentation of data, Protocol, Cohorts studies,	
	Observational studies, Experimental studies,	
	Designing clinical trial, various phases.	





4 Unit-IV CO	$\Omega$ 4
A. Blocking and confounding system for Two-	04
level factorials Regression modeling:	
Hypothesis testing in Simple and Multiple	
regression models	
B. Introduction to Practical components of	
Industrial and Clinical Trials Problems:	
®Statistical Analysis Using Excel, SPSS,	
MINITAB DESIGN OF EXPERIMENTS, R	
C. Online Statistical Software's to Industrial and	
Clinical trial approach	
5 Unit-V CC	O5
A. Design and Analysis of experiments: CO	06
Factorial Design: Definition, 2, 2 design.	
Advantage of factorial design	
B. Response Surface methodology: Central	
composite design,	
C. Historical design, Optimization Techniques	
Mode of Theory	
examination	
Weightage Continuous Sessional ESE	
Distribution Mode Exam	
Assessment	
Text book/s*	
Other Recommended Books (Latest edition):	
References 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,	
Statems Edition,	





CO/PO	PO1	PO2	PO3	P O4	PO 5	PO 6	PO7	P0 8	PO9	PO10	PO11
CO1	2	2	2	2	2	ı	2	-	2	2	2
CO2	3	2	1	2	1	•	2	-	3	1	2
CO3	3	2	2	-	2	•	3	-	2	1	1
CO4	3	1	1	-	-	-	3	-	3	2	1
CO5	3	1	2	-	1	-	2	-	3	2	3
CO6	3	2	2	1	3	-	2	-	1	2	1

- 1-Slight (Low)
- 2-Moderate (Medium)
  3-Substantial (High)





Sch	ool:	SOP	
	gramme:	B.Pharm	
Branch:		Semester: VIII	
1	Course Code	BP802T	
2	Course Title	Social & Preventive Pharmacy- Theory	
3	Credits	4	
4	Contact Hours (L-T-P)	3-1-0	
	Course Type	Compulsory	
5	Course Objective	<ol> <li>Upon completion of this course the student should be a         <ol> <li>Acquire high consciousness/realization of curr health and pharmaceutical problems within worldwide.</li> <li>Have a critical way of thinking based on development.</li> </ol> </li> <li>Evaluate alternative ways of solving problem and pharmaceutical issues</li> </ol>	ent issues related to n the country and current healthcare
6	Course Outcomes	CO1: The students will understand the issues related health. CO2: The student will be able to summarize the impolicies run for various health issues. CO3: The student will be able to apply the knowledge the health issues of the society on finding the effect eradication of diseases. CO4: The students will compare the correlation affecting the health status of common people and we plans to combat the health issues. CO5: The students would evaluate the processes Programmes related to social health and prevention of CO6: The students would elaborate the processes of of diseases	pact of govt. health ge of understanding fective solution for of various factors ill assess the action of various national various diseases.
7	Course Description	The purpose of this course is to introduce to students issues and their challenges. This course also introduce national health Programmemes. The roles of the properties are also discussed.	duced a number of
8	Outline syllabus		CO Mapping





1	UnitI:	CO1
	A. Concept of health and disease: Definition,	
	concepts and evaluation of public health.	
	Understanding the concept of prevention and control	
	of disease, social causes of diseases and social	
	problems of the sick.	
	<b>D. Social and health education:</b> 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	
2	Unit II:	CO2
	<b>A. Preventive medicine:</b> General principles of	
	prevention and control of diseases such as	
	cholera, SARS,	
	<b>B.</b> Ebola virus, influenza, acute respiratory	
	infections, malaria, chicken guinea, dengue,	
	lymphatic	
	C. filariasis, pneumonia, hypertension, diabetes	
	mellitus, cancer, drug addiction-drug	
_	substance abuse	
3	Unit III:	CO3
	A. National health Programmes, its	
	objectives, functioning and outcome of the	
	following: HIV AND AIDS control	
	Programmeme, TB, Integrated disease	
	surveillance Programme (IDSP),	
	B. National leprosy control Programmeme, National mental health Programme, National	
	Programmeme for prevention and control of	
	deafness,	
	C. Universal immunization Programmeme,	
	National Programmeme for control of	
	blindness, Pulse polio Programmeme.	





4	Unit IV:			CO4
	<b>A.</b> Nation			
	for mo			
	Progra	mmeme, Natio	onal tobacco control	
	Progra	mmeme,		
	<b>B.</b> Nation	al Malaria Pre	evention Programme,	
	Nation	al Programme	me for the health care	
		elderly,		
		_	mmeme; role of WHO in	
		national Progr	ramme	
5	Unit V:			CO5
		•	in rural, urban and	CO6
		health: Functi	· ·	
	-		l sanitation, national	
			, Health promotion and	
		ion in school.		
Mode of	Theory			
examination		Sessional	ESE	
Weightage	Continuous			
Distribution	Mode			
	Assessment			
	10 Marks	15	75	
Text book/s*				





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





CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	PO7	P08	PO9	PO10	PO11
CO1	3	2	2	2	2	3	1	1	2	2	2
CO2	3	2	1	2	2	3	1	2	3	1	2
CO3	3	2	2	2	2	2	3	2	2	1	1
CO4	3	1	1	2	2	2	3	2	3	2	1
CO5	3	1	2	2	1	3	2	2	3	2	3
CO6	2	1	2	2	3	1	2	1	2	3	1

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sch	nool:	SOP							
Pro	gramme:	B.Pharm							
	nch:	Semester: VIII							
1	Course Code	BP803ET							
2	Course Title	Pharma marketing management- Theory							
3	Credits	4							
4	Contact Hours (L-T-P)	3-1-0							
	Course Type	Compulsory							
5	Course Objective	After the successful completion of this course, the student shall be able to:							
		<ol> <li>The course aims to provide an understanding of marketing concepts and techniques and their applications in the pharmaceutical industry</li> <li>Have a critical way of thinking based on different marketing strategy for product development.</li> </ol>							
		3. The aim here is to develop a community around the brand whereby audiences can interact with certain content. In the pharmaceutical industry, more so big pharma, and just like most other consumer-facing industries,							
		there are more products and more messages subsequently meaning more noise.							
6	Course Outcomes	CO1: The students will understand the Marketing concepts and techniques and the application of the same in the pharmaceutical industry.							
		CO2: The student will be able to summarize the Market research and distribution channels along with their implementation in the pharmaceutical industry.  CO3: The student will be able to apply the knowledge regarding the Concepts of product line and product mix decisions, branding and product management.  CO4: The students will analyze the Theories on promotion techniques for OTC Products, sales and pricing of a product.  CO5: The students would evaluate the processes of issues in price management in the pharmaceutical industry.  CO6: The students will be able to conclude about the Emerging concepts in marketing and Global Marketing concept.							
7	Course Description	The pharmaceutical industry not only needs highly qualified researchers, chemists and, technical people, but also requires skilled managers who can take the industry forward by managing and taking the complex decisions which are imperative for the growth of the industry. The Knowledge and Know-how of marketing management groom the people for taking a challenging role in Sales and Product management.							





8	Outline syllabus	www.sharda.	CO Mapping
0			
		Unit A.Marketing: Definition, general concepts and scope of marketing; Distinction between marketing & selling; Marketing environment; Industry and competitive analysis; Analyzing consumer buying behavior; industrial buying behavior. A. Pharmaceutical market: Quantitative and qualitative aspects; size and composition of the market; demographic descriptions and socio-psychological characteristics of the consumer; market segmentation& targeting.Consumer profile; C.Motivation and prescribing habits of the physician; patients' choice of physician and retail pharmacist.Analyzing the Market;Role of market research.	CO1
	2	Unit II:  A. Productdecision:  Classification, product line and product mix decisions, product life cycle, product portfolio analysis; product positioning;  B. New product decisions; Product branding, packaging and labeling decisions,  C. Product management in pharmaceutical	CO2
	2	industry.	CO2
	3	<ul> <li>Unit III:</li> <li>A. Promotion:</li> <li>Methods, determinants of promotional mix, promotional budget;</li> <li>B. An overview of personal selling, advertising, direct mail, journals, sampling, retailing,</li> <li>C. medical exhibition, public relations, online promotional techniques for OTC Products.</li> </ul>	CO3





4	4	Unit IV:			CO4					
		A. Pharma	ceutical n	narketing channels:						
			Designing channel, channel members, selecting							
		the appr								
				management: Strategic						
				n physical distribution						
		manager	nent.							
		B. Profes	sional sales r	epresentative (PSR):						
		Duties of PSI	R, purpose of	detailing, selection and						
		training, supe	ervising, nor	ms for customer calls,						
		motivating, ev	•							
			nsation and	future prospects of the						
		PSR.								
	5	Unit V:			CO5					
		A. Pricin	0		CO6					
		•		jectives, determinants of						
			_	and strategies, issues in						
		-	_	rmaceutical industry. An						
				ug Price Control Order)						
			A (National	Pharmaceutical Pricing						
		Authority).	•	• • •						
				in marketing:						
		Vertical		rizontal Marketing;						
		RuralMarketin	•	; Global Marketing.						
<del>                                      </del>	Mode of	Theory	nai waikeung	, Olovai Maikellig.						
	examination	THEOLY								
	Weightage	Continuous								
	Distribution	Mode	Sessional Exam	ESE						
		Assessment								
		10 Marks	15	75						
	Text book/s*									





Other	Recommended Books: (Latest Editions)
References	1. Philip Kotler and Kevin Lane Keller:
References	Marketing Management, Prentice Hall of
	India, New Delhi
	2. Walker, Boyd and Larreche: Marketing
	Strategy- Planning and Implementation, Tata
	MC GrawHill, New Delhi.
	3. Dhruv Grewal and Michael Levy: Marketing,
	Tata MC Graw Hill
	4. Arun Kumar and N Menakshi: Marketing
	Management, Vikas Publishing, India
	5. Rajan Saxena: Marketing Management; Tata
	MC Graw-Hill (India Edition)
	6. Ramaswamy, U.S & Nanakamari, S:
	Marketing Managemnt:Global Perspective,
	IndianContext,Macmilan India, New Delhi.
	7. Shanker, Ravi: Service Marketing, Excell
	Books, New Delhi
	8. Subba Rao Changanti, Pharmaceutical
	Marketing in India (GIFT – Excel series)
	Excel
	7. Rabindra Nath, Saha Indranil, 4
	Publications
	Review of Preventive and Social Medicine
	(Including Biostatistics), Jain Vivek, 6
	Edition, 2014, ISBN: 9789351522331,
	JAYPEE Publications
	8. Essentials of Community Medicine—A
	Practical Approach, Hiremath Lalita D, nd
	9. Hiremath Dhananjaya A, 2 Edition, 2012,
	ISBN: 9789350250440, JAYPEE
	Publications Park Textbook of Preventive
	and Social Medicine, K Park, 21 Edition,
	2011,
	10. ISBN-14: 9788190128285, BANARSIDAS
	BHANOT PUBLISHERS. Community
	Pharmacy Practice, Ramesh Adepu, BSP
	publishers, Hyderabad
	, , , , , , , , , , , , , , , , , , ,





CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	PO7	P08	PO9	PO10	PO11
CO1	3	1	2	1	1	3	1	2	1	2	3
CO2	3	2	1	2	1	3	1	2	2	2	3
CO3	3	2	2	2	2	3	1	2	1	3	3
CO4	3	2	1	2	2	3	2	2	2	2	3
CO5	3	2	2	3	2	3	2	3	1	3	3
CO6	3	2	1	2	2	3	2	2	2	2	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Ç.	hool:	SOP	d.dC.IN					
	ogramme: canch:	B.Pharm Semester: V						
		BP804ET						
1	Course Code	BP804E1						
2	Course	Dharmagautical Dagulatory Science Theory						
	Title	Pharmaceutical Regulatory Science - Theory						
3	Credits	4						
4	Contact	3-1-0						
	Hours							
	(L-T-P)							
	Course	Compulsory						
	Type							
5	Course Objective	Upon completion of the subject student shall be able to;						
		Know about the process of drug discovery and development	t					
		Know the regulatory authorities and agencies government manufacture and sale of pharmaceuticals	erning the					
		Know the regulatory approval process and their registration Indian and international markets	in					
6	Course Outcomes	CO1: Students would be able to understand about the proceand development	ess of drug discovery					
		CO2: Students would be able to apply the regulatory auth agencies governing the manufacture and sale of pharmaceur CO3: Students would be able to explain the regulatory appr CO4: Students would be able to analyze the registration of and international market.  CO5 Students would be able to plan about clinical trials, exprotocol designing.	ticals oval process f drug in Indian					
7	Course Descriptio n	This course is designed to impart the fundamental knowled regulatory requirements for approval of new drugs, and drain regulated markets of India & other countries like US, Australia, UK etc. It prepares the students to learn in design regulatory requirements, documentation requirements, and procedures for marketing the drug products.	ug products EU, Japan, etail on the					
8	Outline sylla	abus	CO Mapping					





1		
1	Unit I	CO1
	A. New Drug Discovery and development	
	Stages of drug discovery,	
	B.Drug development process, pre-clinical studies, non-	
	clinical activities, clinical studies,	
	C.Innovator and generics, Concept of generics, Generic	
	drug product development.	
2	Unit II	
		CO2
	A. Regulatory Approval Process	
	Approval processes and timelines involved in	
	Investigational New Drug (IND), New Drug Application	
	(NDA), Abbreviated New Drug Application (ANDA).	
	Changes to an approved NDA / ANDA.	
	B. Regulatory authorities and agencies	
	Overview of regulatory authorities of India, United	
	States, European Union,	
	C.Australia, Japan, Canada (Organization structure and	
	types of applications)	
3	Unit III	CO3
	A. Registration of Indian drug product in overseas	
	market	
	Procedure for export of pharmaceutical products,	
	Technical documentation, Drug Master Files (DMF),	
	Common Technical Document (CTD), electronic	
	Common Technical	
	<b>B.</b> Document (eCTD), ASEAN Common Technical	
	Document (ACTD) research.	
	pelletization process,	
	C.equipments for manufacture of pellets	
1	Unit IV	
4	A. Clinical trials	CO4
	Developing clinical trial protocols, Institutional Review	
	Board / Independent Ethics committee - formation and	
	working procedures, Informed consent process and	
	procedures, <b>B.</b> GCP obligations of Investigators, sponsors & Monitors,	
	C.Managing and Monitoring clinical trials,	
	Pharmacovigilance - safety monitoring in clinical trials	
5	Unit V	CO5
	A. Regulatory Concepts	CO6
	Basic terminology, guidance, guidelines, regulations,	
	Laws and Acts, Orange book, Federal Register, Code of	
	Federal Regulatory, Purple book	





			ever vv. silai u	3.40.111
Mode of examination	Theory			
Weightage	<b>Continuous Mode</b>	Sessional Exam	ESE	
Distributio	Assessment			
n	10 Marks	15	75	
Text book/s*	N.S. Vyawahare, The Pharmac Edition Edited by Drugs and the Informa Health ca  New Drug Global Registratic edition, Drugs Sciences, Vol. 190  Guidebook Sandy Weinberg. FDA Regular drugs, medical of Douglas J. Pisano Generic Drug Dosage forms, Marcel Dekker se Clinical Trial Guide to Regular Rozovsky and Ro  Principles an Second Edition Frederick P. Ogni	reutical Regulatory Ira R. Berry and I Pharmaceutical S Pharmaceutical S Pharmaceutical S Pharmaceutical S Pharmaceutical S Propose Proces Pro	Process, Second Robert P. Martin, Sciences, Vol. 185.  ss: Accelerating Guarino, MD, 5 th Pharmaceutical ry submissions / Sons. Inc. te for prescription ogics /edited by ment, Solid Oral I Isader Kaufer, earch: A Practical re By Fay A.  linical Research, I. Gallin and	





Pos	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
Cos											
CO1	3	3	1	1	3	3	1	_	2	1	3
CO2	3	3	-	1	3	3	1	-	3	1	3
CO3	3	3	2	2	3	2	1	-	3	3	3
CO4	3	2	_	1	3	3	2	_	2	1	3
									_	_	
CO5	3	3	1	1	3	3	1	-	2	1	3
CO6	3	2	1		3	3	1	-	2	1	3

1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)





Sch	ool:	SOP						
Pro	gramme:	B.Pharm						
	nch:	Semester: VIII						
1	Course Code	BP805ET						
2	Course Title	Pharmacovigilance Theory						
3	Credits	4						
4	Contact Hours (L-T-P)	3-1-0						
	Course Type	Elective						
5	Course Objective	This course has been designed to impart the fundamental knowledge of pharmacy practice and ethics along with the aspects of hospital organization.  Objectives: Upon the completion of this course the students shall be able toTell 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 Methods to generate safety data during pre clinical, clinical and post approval phases of drugs' life cycle  Drug safety evaluation in paediatrics, geriatrics, pregnancy and lactation Pharmacovigilance Programme of India (PvPI) requirement for ADR reporting in India  ICH guidelines for ICSR, PSUR, expedited reporting, pharmacovigilance planning  CIOMS requirements for ADR reporting						
6	Course	Writing case narratives of adverse events and their quality.  Student will be able to						
U	Outcomes	CO1: Understant the history of Pharmacovigilance						
		CO2: Apply about the Detection of new adverse drug reactions and their assessment CO3: Analyze the suspected drug events CO4: Interpret the importance of PV PI CO5: Understand reports like ICSR, PSUR CO6: Apply the ADR for vaccines						





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7	Course	This paper will provide an opportunity for the student to learn about										
	Description	development of pharmacovigilance as a science, basic terminologies used										
		in pharmacovigilance, global scenario of Pharmacovigilance, train										
		students on establishing pharmacovigilance Programmeme in an										
		organization, various methods that can be used to generate safety data and										
		signal detection. This paper also develops the skills of	of classifying drugs,									
		diseases and adverse drug reactions.										
8	Outline syllabus		CO Mapping									
	1	A. Introduction to Pharmacovigilance										
		C	CO1									
		a. History and development of										
		Pharmacovigilance										
		b. Importance of safety monitoring of										
		Medicine										
		c. WHO international drug monitoring										
		Programmeme										
		B. Pharmacovigilance Programme of										
		India(PvPI)										
		C. Introduction to adverse drug reactions										
		Definitions and classification of ADR										
		Detection and reportin										
		Methods in Causality assessment										
		Severity and seriousness assessment										
		Predictability and preventability assessment										
		prevention.										
		Management of adverse drug reactions										
		Basic terminologies used in										
		pharmacovigilance used in										
		Terminologies of adverse medication										
		related events										
		Regulatory terminologies										
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Drug and disease classification CO₂ 2 A. Drug dictionaries and coding in pharmacovigilance Anatomical, therapeutic and chemical classification of drugs International classification of diseases Daily defined doses International Non proprietary Names for drugs **B.** Information resources in pharmacovigilance Basic drug information resources Specialised resources for ADRs WHO adverse reaction terminologies MedDRA and Standardised MedDRA queries WHO drug dictionary Eudravigilance medicinal product dictionary C. Establishing pharmacovigilance **Programmeme** Establishing in a hospital Establishment & operation of drug safety department in industry Contract Research Organisations (CROs) Establishing a national Programmeme





3	A. Vaccine safety surveillance	CO3
	Vaccine Pharmacovigilance	
	Vaccination failureAdverse events following	
	immunization	
	B. <b>Pharmacovigilance</b> methodsPassive	
	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  C. Communication in pharmacovigilance	
	Effective communication in Pharmacovigilance	
	Communication in Drug Safety Crisis management Communicating with Regulatory Agencies,	
	Business Partners, Healthcare facilities & Media	
	Business Furthers, Heartheare Fuelines & Media	
4	A. Safety data generation	CO4
4	1. Pre clinical phase	CO4
	2. Clinical phase	
	3. Post approval phase (PMS)	
	ost approvar phase (1 1/16)	
	B. ICH Guidelines for Pharmacovigilance	
	1. Organization and objectives of ICH	
	2. Expedited reporting	
	3. Individual case safety reports	
	4. Periodic safety update reports	
	5. Post approval expedited reporting	
	6. Pharmacovigilance planning	
	7. Good clinical practice in	
	pharmacovigilance studies	





			THE TRIBUTURE	
5	A. Pharn reaction examp B. Drug so population a. Page	CO5 CO6		
	b. Preg	gnancy and lac	ctation	
	c. Geri			
	C. CIOM	-		
		S Working Gr	oups	
	CIOMS			
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	□□ D&C Act			
	pharmacovigi		_	
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Mode of examination	Theory			
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	Assessment			
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Text book/s*	D 1	1D 1 (T	4 4 1949	
Other	Recommende			
References	Brothers, Med	_	lance: S K Gupta, Jaypee	
	· ·		A to Z By Barton Cobert,	
			lett Publishers.	
			e:Elizabeth B. Andrews,	
	Nicholas, Wil	-		
	Stephens' D	etection of	New Adverse Drug	
		ohn Talbot	, Patrick Walle, Wiley	
	Publishers.			
			rmacovigilance: Patrick	
	Waller, Wiley Cobert's M	Publishers. Ianual of	Drug Safety and	
			n Cobert,Jones&Bartlett	
	Publishers.	iance. Dano	ii Coort, ones & Dartiett	



Cos/POs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11
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CO3	3	2	3	2	2	1	1	2	2	2	2
CO4	2	1	2	2	2	-	1	2	2	2	3
CO5	2	2	2	2	2	-	1	2	2	2	3
CO6	1	2	3	1	1	_	2	1	2	3	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High





Sch	ool:	SOP
Pro	gramme:	B.Pharm
Bra	nch:	Semester: VIII
1	Course Code	BP808ET
2	Course Title	Cell and molecular biology - Theory
3	Credits	4
4	Contact Hours (L-T-P)	3-1-0
	Course Type	Elective
5	Course	Upon the completion of the course student shall be able to:
	Objective	1. Summarize cell and molecular biology history.
		2. Summarize cellular functioning and composition.
		3. Describe the chemical foundations of cell biology.
		a. Summarize the DNA properties of cell biology.
		b. Describe protein structure and function.
		4. Describe cellular membrane structure and function.
		5. Describe basic molecular genetic mechanisms.
6	Course Outcomes	CO1: Students would be able to understand the concept of cell theory and basics of cellular structure and the mechanism of immune system.
		CO2: Students would be able to apply cellular reproduction in eukaryotic cells and increase their knowledge about nucleic acids
		CO3: Students would be able to interpret DNA, RNA and their role in central dogma of life, Protein synthesis and types of RNA.
		CO4: Students would be able to apply Transgenics and Genomic Analysis, Cell Cycle analysis, role of genetics, mitosis and meiosis.
		CO5: Students would be able to explain Cell Signals, Receptors for Cell Signals, Signaling Pathways, Misregulation of Signaling Pathways and Protein- Kinases: Functioning
		CO6: Students will be able to analyze different aspect of cell function which will help them to think about research involving cell biology.





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7	Course									
	Description	• Cell biology is a branch of biology that studies cel	ls – 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	_							
		specialized cells in multi-cellular organisms such as humans,								
			piants, and							
8	Outline avillation	sponges.	СО							
0	Outline syllabus									
	1	TT *4 T	Mapping							
	1	Unit I	CO1							
		a) Cell and Molecular Biology: Definitions theory and								
		basics and Applications.								
		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)								
	2	Unit II								
		a). DNA and the Flow of Molecular Information	CO2							
		b). DNA Functioning								
		c). DNA and RNA								
		d). Types of RNA								
		e). Transcription and Translation								
	3	Unit III	CO3							
		a). Proteins: Defined <b>and</b> Amino Acids								
		b). Protein Structure								
		c). c)Regularities in Protein Pathways								
		d). Cellular Processes								
		e). Positive Control and significance of Protein Synthesis								
	4	Unit IV								
	¬	a). Science of Genetics	CO4							
		b). Transgenics and Genomic Analysis	004							
		c). Cell Cycle analysis								
		d). Mitosis and Meiosis a) Collular Activities and Chackpoints								
		e). Cellular Activities and Checkpoints	COF							
	5	Unit V	CO5							
		a). Cell Signals: Introduction	CO6							
		b). Receptors for Cell Signals								
		c). Signaling Pathways: Overview								
		d). Misregulation of Signaling								
		e). Pathways Protein-Kinases: Functioning								





M	lode of	Theory							
ex	kamination	•							
W	/eightage	Continuous	Sessional	ESE					
D	istribution	Mode	Exam						
		Assessment							
			15	75					
		10 Marks							
Te	ext book/s*								
O	ther		ed Books (late						
Ro	eferences	Micro	-	A.D. Russel: Pharmaceutical ckwell Scientific publications,					
			outors, Delhi. e	nn., Industrial Microbiology, edition, CBS Publishers & Microbiology, Tata McGraw Hill					
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		Micro 7. Coope	her, Hinsdil biology, 9th ec r and Gum her and Distrik	l. Japan 1's: Tutorial Pharmacy, CBS					
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		11. Berger and W	Prakashan, Delhi 11. Bergeys manual of systematic bacteriology, Williams and Wilkins- A Waverly a. company						
		12. 12.B.F Biotec	R. Glick ar hnology: Princ	*					
		Washi		RecombinantDNA: ASM Press 3. RA Goldshy et. al., : Kuby					





CO/PO	PO1	PO2	PO3	PO4	PO5	PO6	PO7	P08	PO9	PO10	PO11
CO1	3	1	3	2	2	1	2	ı	2	3	3
CO2	3	1	3	2	2	2	2	1	2	1	3
CO3	3	2	2	2	2	3	2	-	2	2	3
CO4	3	1	1	3	3	3	3	-	2	2	3
CO5	3	2	2	3	2	2	2	-	2	1	3
CO6	3	1	2	2	2	2	2	-	2	1	3

- 1-Slight (Low) 2-Moderate (Medium) 3-Substantial (High)