## PES Modern College of Pharmacy, Nigdi, Pune Course outcome 2023-24 B. Pharmacy

Semeste		CO's
r	Subject Name	
	r Sem I 2019 pattern	1. To explain structure of various components, tissues and organs
BP101T	HUMAN ANATOMY AND PHYSIOLOGY I–T Div (A)	of human body systems.  2. Outline functions and Classify different components of human body systems  3. To Summarize and apply data of coordinated normal physiological pattern of different organs of each system.  4. To describe the various homeostatic mechanisms and their related disorders
	HUMAN ANATOMY AND PHYSIOLOGY I–T Div (B)	<ol> <li>To explain structure of various components, tissues and organs of human body systems.</li> <li>Outline functions and Classify different components of human body systems</li> <li>To Summarize and apply data of coordinated normal physiological pattern of different organs of each system.</li> <li>To describe the various homeostatic mechanisms and their related disorders</li> </ol>
BP102T	PHARMACEUTICAL ANALYSIS I – T (A)	<ol> <li>Assess the fundamentals of analytical chemistry and principles of electrochemical analysis of drugs</li> <li>Compare different types of titration method and expressing their concentration</li> <li>Integrate various volumetric and electrochemical titrations.</li> </ol>
		4. Select various titration method for end point detection and oxidation and reduction mechanism reaction
	PHARMACEUTICAL ANALYSIS I – T (B)	<ol> <li>Assess the fundamentals of analytical chemistry and principles of electrochemical analysis of drugs.</li> <li>Compare different types of titration methods and expressing</li> </ol>
		their concentration  3. Integrate various volumetric and electrochemical titrations.  4. Select various titration method for end point detection and oxidation and reduction mechanism reaction
BP103T	PHARMACEUTICS I – T (A)	<ol> <li>Introduction to history of pharmacy, development of pharmacy profession, industry in India, dosage form, prescription, posology, pharmaceutical calculation, powders, liquid dosage form, monophasic-biphasic liquid dosage form, semisolid dosage form and suppositories.</li> </ol>
		<ol> <li>Explain and evaluate different preparation like powders, monophasic-biphasic liquid dosage form, semisolid dosage form and suppositories</li> <li>Define &amp; classify prescription, dosage form, posology powders, liquid dosage form, monophasic-biphasic liquid</li> </ol>
		dosage form, semisolid dosage form and suppositories.  4. Summarize the Advantage, Disadvantage & Disadvantage amp; factors influencing powflers, monophasic-biphasic liquid dosage form



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	1	to the form and suppositories
	PHARMACEUTICS I – T (B)	<ol> <li>semisolid dosage form and suppositories</li> <li>Introduction to history of pharmacy, development of pharmacy profession, industry in India, dosage form, prescription, posology, pharmaceutical calculation, powders, liquid dosage form, monophasic-biphasic liquid dosage form, semisolid dosage form and suppositories.</li> <li>Explain and evaluate different preparation like powders, monophasic-biphasic liquid dosage form, semisolid dosage form and suppositories.</li> <li>Define &amp; classify prescription, dosage form, posology, powders, liquid dosage form, monophasic-biphasic liquid dosage form, semisolid dosage form and suppositories.</li> <li>Summarize the Advantage, Disadvantage &amp; factors influencing powders, monophasic-biphasic liquid dosage form, semisolid dosage form and suppositories</li> </ol>
BP104T	PHARMACEUTICAL INORGANIC CHEMISTRY T (A)	<ol> <li>Derive acquainted with the principles of limit tests.</li> <li>Integrate and analyze the different anions, cations from inorganic pharmaceuticals.</li> <li>Invent the sources of impurities and methods to determine the impurities in inorganic drugs and pharmaceuticals</li> <li>Derive acids, bases, buffers, water and different GIT agents and recall the fundamental principles of them</li> <li>Develop the medicinal used of topical agents, gases and vapors', dental products, pharmaceuticals aid and radio pharmaceuticals.</li> <li>Integrate about the major intra and extra cellular electrolytes, essential and trace elements, cationic and anionic components of inorganic drugs.</li> </ol>
	PHARMACEUTICAL INORGANIC CHEMISTRY T (B)	<ol> <li>Derive acquainted with the principles of limit tests.</li> <li>Integrate and analyze the different anions, cations from inorganic pharmaceuticals.</li> <li>Invent the sources of impurities and methods to determine the impurities in inorganic drugs and pharmaceuticals</li> <li>Derive acids, bases, buffers, water and different GIT agents and recall the fundamental principles of them</li> <li>Develop the medicinal used of topical agents, gases and vapors', dental products, pharmaceuticals aid and radio pharmaceuticals.</li> <li>Integrate about the major intra and extra cellular electrolytes, essential and trace elements, cationic and anionic components of inorganic drugs.</li> </ol>
BP105T	COMMUNICATION SKILLS – T * (A&B)	<ol> <li>Develop communication skills effectively with range of people in variety of setting, using different of modes and media</li> <li>Integrate behavioral needs of pharmacist to function effectively in the area of pharmaceutical operations</li> <li>To develop understanding and interpret subjects</li> <li>To develop ability to apply what is learned</li> <li>To focus on curricular, co-curricular and extracurricular activities</li> <li>To develop graduates with ethics, morals and social sense and decision making</li> </ol>
BP106R BT	REMEDIALBIOLOGY (A&B)	<ol> <li>To identify a given plant part based on its macroscopic and microscopic characteristics.</li> </ol>



		2. To illustrate the classification of plants, plant cell and its
		organelles.
	7-25-1	3. To describe the physiological processes in plants and humans.
		4. To explain the type of tissues present in human body.
		5. To discuss the anatomy and functions of systems of the human
		body
		6. To appraise the coordinated working pattern of different
		organs of human body.
BP107P	HUMAN ANATOMY	1. To summarize the basic knowledge of importance of Human
2	AND PHYSIOLOGY	Anatomy and Physiology in pharmacy field.
	_P	2. Apply knowledge in handling the laboratory equipment and
	Div (A)	models in order to study and compilation of data on tissues,
		blood estimations, cardiac and skeletal system
		3. To develop practical skill in Students convergent with the
		techniques for identification, counting, estimation of
		hematological studies
		4. Analyze data to investigate clinical situation based on results
	HUMAN ANATOMY	1. To summarize the basic knowledge of importance of Human
	AND PHYSIOLOGY	Anatomy and Physiology in pharmacy field.
	_P	2. Apply knowledge in handling the laboratory equipment and
	Div (B)	models in order to study and compilation of data on tissues,
		blood estimations, cardiac and skeletal system
		3. To develop practical skill in Students convergent with the
		techniques for identification, counting, estimation of
		haematological studies
		4. Analyze data to investigate clinical situation based on results
BP108P	PHARMACEUTICAL	1. Assess the fundamentals of analytical chemistry and theory of
	ANALYSIS $I - P(A)$	electrochemical analysis of drugs
		2. Evaluate various molar and normal solution in Pharmaceutical
		solution
		3. Evaluate various volumetric and electrochemical titrations.
		4. Measure end point detection in volumetric and electrochemical
		titrations
	PHARMACEUTICAL	1. Assess the fundamentals of analytical chemistry and principles
	ANALYSIS I – P (B)	of electrochemical analysis of drugs
	ANVALUE OF T (B)	2. Compare different types of titration methods and expressing
		their concentration
1		3. Integrate various volumetric and electrochemical titrations.
		4. Select various titration method for end point detection and
		oxidation and reduction mechanism reaction
DD100D	DITADIAA OPTITIOOT	Demonstrate the skill of preparation and evaluation of various
BP109P	PHARMACEUTICS I	
	- P (A)	solid liquid and semisolid dosage forms.
		2. Explain the principles of formulation and evaluation of powder
		preparations.
		3. Calculate evaluation parameters like density, specific gravity,
		angle of repose, Carr's index and Hausner's ratio of
		pharmaceutical preparations.
		4. Classify various dosage forms by using different criteria
		5. Create labels in prescribed manner for various dosage forms.
	PHARMACEUTICS I	1. Demonstrate the skill of preparation and evaluation of various
	I III II	The second of various



	– P (B)	solid liquid and semisolid dosage forms.
		2. Explain the principles of formulation and evaluation of powder
		preparations.
		3. Calculate evaluation parameters like density, specific gravity,
	34	angle of repose, Carr's index, Hausner's ratio, etc. of
		pharmaceutical preparations.
		4. Classify various dosage forms by using different criteria
		5. Create labels in prescribed manner for various dosage forms.
BP110P	PHARMACEUTICAL	1. Recall the sources of limit tests, preparation and identification
1	INORGANIC	of compounds.
	CHEMISTRY –P	2. Demonstrate the preparation of inorganic pharmaceuticals.
	(Div A)	3. Apply knowledge to perform modified limit tests.
		4. Analyze various inorganic pharmaceutical compounds
1		5. Select suitable method for the preparation of inorganic
		pharmaceuticals.
		6. Assess quality of inorganic pharmaceuticals.
	PHARMACEUTICAL	1. Derive acquainted with the principles of limit tests.
1	INORGANIC	2. Compose and Familiar with different classes of inorganic
	CHEMISTRY –P	pharmaceuticals and their analysis
	(Div B)	3. Integrate and analyze the anions, cations in different inorganic
		pharmaceuticals.
		4. Invent the sources of impurities and methods to determine the
		impurities in inorganic drugs and pharmaceuticals
BP111P	COMMUNICATION	1. Develop etiquettes, mannerism, soft skill and communication
	SKILLS – P* (A&B)	skill
	(1102)	2. Develop presentation skills, listening skills and sophisticated
		nonverbal communication
		3. Generate leadership quality, emotional intelligence and
		cognitive skills
		4. Develop good interview skills with complete professional
		etiquettes.
BP112R	REMEDIAL	To able to identify microscopy of tissues pertinent to stem,
BP 112K	BIOLOGY – P* (A&B)	root, leaf, seed, fruit and flower.
DI	BIOLOGI -I (A&B)	2. To perform blood group detection, measurement of blood
		pressure and tidal volume.
	1	3. To demonstrate different bones in human skeleton system,
T:	C 4 W	their location and significance.
	r Semester II	1. To relate the heate by aviled as about a state
BP201T	HUMAN ANATOMY	1. To relate the basic knowledge about central nervous system
	AND PHYSIOLOGY	including nervous tissue, brain and spinal cord.
	II – T (Div A)	2. To illustrate the structure and functions of gastrointestinal tract
		and to learn about ATP/CTP/BMR.
		3. To learn about structure and functions of respiratory system
	1917	and various mechanisms involved in regulation of respiration.
	Fig. 1.	4. To categorize the anatomy of urinary system and physiology
		of urine formation/ maturation.
		5. To appraise the essentiality of endocrine glands and their
		hormones
		6. To predict the physiqlogy of male and female reproductive
		organs and concepts of genetics.



	HUMAN ANATOMY	1. To relate the basic knowledge about central nervous system
	AND PHYSIOLOGY	to dealth a marriage figure brain and Spinal Cold.
	II – T (Div B)	2. To illustrate the structure and functions of gastromiestinal tract
		and to learn about ATP/CTP/BMR.
		3. To learn about structure and functions of respiratory system
		and various mechanisms involved in regulation of respiration.
		4. To categorize the anatomy of urinary system and physiology
		of urine formation/ maturation.
=		5. To appraise the essentiality of endocrine glands and their
		hormones
-		6. To predict the physiology of male and female reproductive
		organs and concepts of genetics.
BP202T	PHARMACEUTICAL	1. Derive and Understand the structure, name and the type of
	ORGANIC	isomerism of the organic compound 2. Integrate the fundamentals of atomic structure, types of bonds,
	CHEMISTRY I-T (Div	2. Integrate the fundamentals of atomic structure, types of somes,
-	(A)	hybridization and addition compounds.  3. Compose the knowledge of reagents, organic reactions and
		3. Compose the knowledge of reagents, organic reasons and
	5	electron displacement effects 4. Evaluate the rules of IUPAC nomenclature. Explain the
		structure, occurrence & stability of ions and free radicals.
		5. Derive the concept of stereochemistry and also can relate and
		value them.
		6. Understand and consider the Alkanes, Alkenes and
		Conjugated dienes and Alkyl halides
		7. Understand, analyze Alcohols, Carbonyl compounds*
		(Aldehydes and ketones)
	And the second of the second o	8. Utilize the principles of scientific Carboxylic acids, Aliphatic
		amines
	PHARMACEUTICAL	1. Derive and understand the structure, name and the type of
	ORGANIC	isomerism of the organic compound.
	CHEMISTRY I-T (Div	2. Integrate the fundamentals of atomic structure, types of bonds,
	B)	hybridization and addition compounds.
		3. Evaluate the rules of IUPAC nomenclature. Explain the
		structure, occurrence & amp; stability of ions and free radicals
		4. Understand and consider the Alkanes, Alkenes and
		Conjugated dienes and Alkyl halides
		5. Understand, analyze Alcohols, Carbonyl compounds*
		(Aldehydes and ketones)
		6. Utilize the principles of scientific Carboxylic acids, Aliphatic
		amines
BP203T	BIOCHEMISTRY – T	1. To summarize biomolecules and to conclude, evaluate and
	(Div-A)	discuss chemical nature and biological role of it with concepts
		in bioenergetics.
		2. To Summarize & Compose biosynthesis and Metabolism of
14		important biomolecules like carbohydrates, lipids, proteins and
147		Nucleic acids.
		3. Integrate, Compose, Explain & Discuss metabolism of
15		biomolecules in pathological and physiological conditions.
		4. To Compose, Assess, Explain and Discuss the genetic
		organization of manimalian genome and functions of DNA in



		in a constitution of the c
		the synthesis of RNAs and proteins.
		5. To summarize, justify and conclude catalytic role of enzymes,
		importance of enzyme in inhibitors in design of new drugs,
		therapeutic and diagnostics applications of enzymes
	BIOCHEMISTRY – T	1. Remember the qualitative analysis of carbohydrates and
	(Div-B)	proteins
		2. Understand the principle and clinical significance of blood
		glucose
		3. Examine the constituents present in Urine and their clinical
		significance
		4. Determine the effect of temperature and substrate
		concentration on salivary amylase activity
		5. Elaborate the clinical significance of creatinine, proteins and
		cholesterol in blood
BP204T	PATHOPHYSIOLOG	Explain the pathogenesis and morphology of reversible and
	Y - T (Div A)	irreversible cell injury; enumerate various lipoproteins and
		describe lipoprotein disorders
		<ol><li>Illustrate events involved in acute and chronic inflammation.</li></ol>
		3. Recognize the biological significance of various
		hypersensitivity disorders.
		4. Discuss the mechanisms involved in autoimmune diseases and
		allograft rejection
		5. Discuss the etiopathogenesis of selected diseases
		6. Describe the general biology of cancer, mechanism of shock
		and effects of radiation exposure
	PATHOPHYSIOLOG	<ol> <li>Explain the pathogenesis and morphology of reversible and</li> </ol>
	Y - T (Div B)	irreversible cell injury; enumerate various lipoproteins and
		describe lipoprotein disorders
	" []	2. Illustrate events involved in acute and chronic inflammation.
		3. Recognize the biological significance of various
		hypersensitivity disorders.
		4. Discuss the mechanisms involved in autoimmune diseases and
		allograft rejection
		5. Discuss the etiopathogenesis of selected diseases
		6. Describe the general biology of cancer, mechanism of shock
		and effects of radiation exposure
BP205T	COMPUTER	Apply the knowledge of mathematics and computing
D1 205 1	APPLICATIONS IN	fundamentals to pharmaceutical applications for any given
	PHARMACY – T *	requirement.
	(A&B)	Design and develop solutions to analyze pharmaceutical
	(Mab)	problems using computers.
	V (**)	3. Integrate and apply efficiently the contemporary IT tools to all
		Pharmaceutical related activities.
		4. Solve and work with a professional context pertaining to
		ethics, social, cultural and regulations with regard to
		Pharmacy.
		5. Student will learn relationship between ethics in clinical trials;
		computational tools etc. and their relevance to today's society
		are introduced to the students
BP206T	ENVIRONMENTAL	1. Create awareness about environmental problems among



\$4 · .	CCIENCES T*	learners and society at large.
	SCIENCES – T *	2. Develop the knowledge about environmental issues and its
	(A&B)	-11: -d muchlama
		3. Develop an attitude of concern about the role of an individual
	1	in a manustion of natural resources
	2.5	4. Create an urge to participate in environmental protection,
		and concernation
		5. Integrate skill to help the society in identifying and solving
		in the state of th
BP207P	HUMAN ANATOMY	1. To summarize the basic knowledge of importance of Human
BP20/P		A A A MARIO DISTRICTOR IN THE PROPERTY OF THE INC.
	AND PHYSIOLOGY	2 The send and the different mechanisms [1056 govern normal]
	II –P (Div A)	working of various organs and systems of human body with
		the half of aborta models and demonstrations.
		3. To develop practical skill in Students convergent with the
	1,200	toobniques for identification, counting, determination
		of Platelet, differential leukocyte count, Arneth index, osmotic
		Con with and hometalogical studies
		A paly knowledge to study Permanent slides and develop skills
		among students in recording of bodies various estimations and
		overminations
	HUMAN ANATOMY	1. Recall the physiology of special senses with the help of
		models charts and specimens.
	AND PHYSIOLOGY	2. Dayslan the knowledge on coordinating working of organs of
	II –P (Div B)	various systems with the help of models, charts and specificis
		3. Analyze the functions of cranial nerves by various sensory and
		motor functions
		4. Determine blood cell count, Arneth index, osmotic fragility of
		RBCs, tidal volume and vital capacity
	DILL BULL CELIEICAI	1. Integrate the reaction, Possess knowledge and synthesis of
BP208P	PHARMACEUTICAL	organic compounds
	ORGANIC	2. Compose the reactivity/stability of compounds
	CHEMISTRY I– P	3. Integrate, Analyze, identify/confirm the identification of
	(Div A)	3. Integrate, Analyze, Identifycommir are izenasta
		organic compound 4. Recall, understand, apply and analyse the unknown organic
		compound by systematic qualitative analysis that includes
	Day the last	preliminary test, detection of element , solubility test,
	1998	preliminary test, detection of element , solutinity test,
		functional group test, mp, bp, derivatives
	PHARMACEUTICAL	1. Introduce safety measures in an organic laboratory and
	ORGANIC	laboratory techniques
	CHEMISTRY I– P	2. Access Systematic qualitative analysis of unknown organic
	(Div B)	compounds
		3. Understand and Prepare of suitable solid derivatives from
	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	organic compounds
		4. Explain Building of molecular models of structures containing
	The state of the s	various functional groups
		5. Discuss the acidity of carboxylic acids and basicity of amines
		and mechanism of some named reaction
		6. Describe the classification of organic compounds and
		nomenclature, isomerism and explain structural isomerism



BP209P	BIOCHEMISTRY – P (Div-A)	1. To understand, remember and apply principles of biochemistry and to analyze, evaluate and create/conclude the biochemistry
		<ol> <li>theory.</li> <li>To evaluate, categorize, summarize and report carbohydrates, amino acids, proteins, normal and abnormal constituents of urine.</li> <li>To prepare folin wu filtrate, determine and conclude serum level of creatinine, cholesterol, reducing sugars and proteins and to measure the pH of buffer solution.</li> <li>To judge, explain, interpret the salivary amylase enzyme activity and able to summarize factors affecting it.</li> </ol>
	BIOCHEMISTRY – P (Div-B)	<ol> <li>To understand, remember and apply principles of biochemistry and to analyze, evaluate and create/conclude the biochemistry theory.</li> <li>To evaluate, categorize, summarize and report carbohydrates, amino acids, proteins, normal and abnormal constituents of</li> </ol>
	17-18-	<ul> <li>urine.</li> <li>To prepare folin wu filtrate, determine and conclude serum level of creatinine, cholesterol, reducing sugars and proteins and to measure the pH of buffer solution.</li> <li>To judge, explain, interpret the salivary amylase enzyme activity and able to summarize factors affecting it.</li> </ul>
BP210P	COMPUTER APPLICATIONS IN PHARMACY – P	<ol> <li>Know the various types of application of computers in pharmacy.</li> <li>Know various types of databases.</li> <li>Know various applications of databases in Pharmacy.</li> <li>It enables us to prepare our students to become more ethical pharmaceutical technologists.</li> <li>Know the web based tools for pharmacy practice</li> <li>Apply the knowledge to design and develop digital tools for pharmaceutical applications</li> </ol>
Second V	ear Semester III	phulmuoution upphousess
BP301T	Pharmaceutical Organic Chemistry II – Theory (Div A)	<ol> <li>Summarize and understand the structure, name, theories, Classification &amp; use of the organic compound as well as fatty acids and oils.</li> <li>Support reasons for Acidity, Basicity, Reactivity &amp; Derive methods of preparation &amp; Derive methods of preparation amp; reactions of organic compounds.</li> <li>Recommend particular chemical entities to predict the product problems.</li> <li>Summarize isomerism with types and choose proper conformation &amp; Derive methods of preparation for assigning them by building various projection formulas &amp; Derive methods of preparation for assigning them by building various projection formulas &amp; Derive methods of preparation for assigning them by building various projection formulas &amp; Derive methods of the product problems.</li> </ol>
	Pharmaceutical Organic Chemistry II – Theory (Div B)	<ol> <li>Summarize the structure, name, theories, Classification &amp; Classification &amp; Support reasons for Acidity, Basicity, Reactivity &amp; Stability of organic compounds.</li> <li>Derive methods of preparation &amp; Support reactions of organic</li> </ol>
		The Art of Significance of Sig



BP302T	Physical Pharmaceutics I  – Theory (Div A)	<ol> <li>compounds.</li> <li>Recommend particular chemical entities to predict the product problems.</li> <li>Smmarize isomerism with types and choose proper conformation &amp; projection for assigning them by building various projection formulas &amp; properties of drug and excipients in designing the dosage forms.</li> <li>Integrate and apply various physicochemical properties of drug and excipients in designing the dosage forms.</li> <li>Demonstrate basics involved in Solubility of drugs, states and properties of matter, surface and interfacial phenomenon and pH and buffer systems.</li> <li>Distinguish the principles of complexation/ protein binding &amp; properties of use them for calculations of drug release and stability constant.</li> <li>Demonstrate ability to develop critical thinking and problem solving required to address problems related to dosage form design and evaluation.</li> <li>Recognize and Apply basic rules and equations regarding</li> </ol>
	Physical Pharmaceutics I  – Theory (Div B)	physical principles essential for pharmaceutical applications  1. Integrate and apply various physicochemical properties of drug and excipients in designing the dosage forms
		<ol> <li>Demonstrate basics involved in Solubility of drugs, states and properties of matter, surface and interfacial phenomenon and pH and buffer systems</li> </ol>
		<ol> <li>Distinguish the principles of complexation/ protein binding &amp; protein binding</li> <li>&amp; amp; to use them for calculations of drug release and stability constant</li> </ol>
		<ol> <li>Demonstrate ability to develop critical thinking and problem solving required to address problems related to dosage form design and evaluation</li> </ol>
		<ol><li>Recognize and Apply basic rules and equations regarding physical principles essential for pharmaceutical applications.</li></ol>
BP303T	Pharmaceutical Microbiology – Theory (Div A)	<ol> <li>Determine the core concepts of microbiology, bacteria, culture media, identification test and microscopy with its application to pharmaceutical field.</li> </ol>
		<ol><li>Summarize and demonstrate sterilization methods, sterility testing and also acquire the knowledge about preservation and disinfection.</li></ol>
		<ol> <li>Prepare a detail scheme which involves basics of aseptic area, clean area classification</li> </ol>
		<ol> <li>Discuss the concept of cell technology and its application in pharmaceutical industry.</li> </ol>
		<ol><li>Describe the spoilage, cell culture technology with its applications in pharmaceutical industries</li></ol>
	1928	<ol> <li>Describe microbiological assay along with methods for standardization of antibiotics, vitamins and amino acids</li> </ol>
I	Pharmaceutical Microbiology – Theory Div B)	1. Determine the core concepts of microbiology, bacteria, culture media, identification test and microscopy with its application to pharmaceutical field.



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		2. Summarize and demonstrate sterilization methods, sterility testing and also acquire the knowledge about preservation and disinfection.
		3. Prepare a detail scheme which involves basics of aseptic area,
		<ul><li>clean area classification</li><li>Discuss the concept of cell technology and its application in</li></ul>
		pharmaceutical industry.
		5. Describe the spoilage, cell culture technology with its
		applications in pharmaceutical industries
		Describe microbiological assay along with methods for
BP304T	Dl. a	standardization of antibiotics, vitamins and amino acids
DP3041	Pharmaceutical Engineering – Theory	<ol> <li>Define drying, crystallization, flow of fluid, evaporation, heat transfer, distillation, corrosion, centrifugation, filtration, size</li> </ol>
	(Div A)	reduction, size separation
		2. Illustrate objective, mechanism & theories of drying, crystallization, flow of fluid, evaporation, heat transfer,
		distillation, corrosion, centrifugation, filtration, size reduction,
		size separation, material handling system
		3. Explain the types & describe the factors affecting drying,
	and the second s	crystallization, flow of fluid, evaporation, heat transfer,
		distillation, corrosion, centrifugation, filtration, size reduction,
		size separation, material handling system
		4. Explain the principle, instrumentation, working & application of drying, crystallization, flow of fluid, evaporation, heat
		transfer, distillation, corrosion, centrifugation, filtration, size
		reduction, size separation, material handling system
		5. Illustrate layout of industrial plant, industrial hazards & plant
		safety
	Pharmaceutical	1. Define drying, crystallization, flow of fluid, evaporation, heat
	Engineering – Theory	transfer, distillation, corrosion, centrifugation, filtration, size
	(Div B)	reduction, size separation
		2. Illustrate objective, mechanism & theories of drying, crystallization, flow of fluid, evaporation, heat transfer,
		distillation, corrosion, centrifugation, filtration, size reduction,
		size separation, material handling system
		3. Explain the types & describe the factors affecting drying,
		crystallization, flow of fluid, evaporation, heat transfer,
		distillation, corrosion, centrifugation, filtration, size reduction,
		size separation, material handling system
		4. Explain the principle, instrumentation, working & application
		of drying, crystallization, flow of fluid, evaporation, heat
		transfer, distillation, corrosion, centrifugation, filtration, size reduction, size separation, material handling system
		5. Illustrate layout of industrial plant, industrial hazards & plant
		safety
BP305P	Pharmaceutical Organic	Use of various equipments and take safety measures while
DF 303F	Chemistry II – Practical	working in organic chemistry laboratory.
	(Div A)	2. Demonstrate and perform laboratory techniques like
		recrystallization and steam distillation.
		3. Identify, implement the separation and identification of given



1,		binary mixture 4. Examine and evaluate the saponification value of given oil compounds
		5. Explain, apply, and justify the principle and theory behind synthesis of compound
		6. Develop and apply skill to synthesize compound and purify synthesized compounds.
	Pharmaceutical Organic Chemistry II – Practical	<ol> <li>Use of various equipments and take safety measures while working in organic chemistry laboratory.</li> </ol>
	(Div B)	<ol> <li>Demonstrate and perform laboratory techniques like recrystallization and steam distillation.</li> <li>Identify, implement the separation and identification of given</li> </ol>
		binary mixture 4. Examine and evaluate the saponification value of given oil
	9	<ul><li>compounds</li><li>5. Explain, apply, and justify the principle and theory behind synthesis of compound</li></ul>
		6. Develop and apply skill to synthesize compound and purify synthesized compounds.
BP306P	Physical Pharmaceutics I  - Practical (Div A)	1 Exercise different pharmaceutical laboratory procedures used in determining various physical properties such as solubility, pka, surface tension, HLB, CMC, adsorption, partition coefficient, stability constant etc.
		2 Design and Perform skillfully some laboratory experiments needed in pharmacy practice as determination of physical properties
		3 Acquire and use technical vocabulary to discuss pharmaceutical problems.
		4 Demonstrate knowledge about some important concepts from preformulation and formulation point of view.
14		5 Employ proper documentation system to record observations and analyze information gathered through experimentation.
	Physical Pharmaceutics I  – Practical ( Div B)	1 Exercise different pharmaceutical laboratory procedures used in determining various physical properties such as solubility, pka, surface tension, HLB, CMC, adsorption, partition
		coefficient, stability constant etc.  2 Design and Perform skillfully some laboratory experiments needed in pharmacy practice as determination of physical properties
		3 Acquire and use technical vocabulary to discuss pharmaceutical problems.
		4 Demonstrate knowledge about some important concepts from preformulation and formulation point of view.
		Employ proper documentation system to record observations and analyze information gathered through experimentation.
3P307P	Pharmaceutical Microbiology – Practical	Know the principle, construction and working of equipments and skill to handle microscope for observation of microbes
	(Div A)	2. Prepare and sterilize nutrient broth, nutrient agar, slants, stabs and plates and adopt the skills required for maintaining strictly aseptic condition & handling inoculating loop, its sterilization
	25	toop, its sterifization



	Pharmaceutical Microbiology – Practical (Div B)	and inoculation procedure  3. Practice aseptic procedures for inoculation and examine sterility testing of pharmaceuticals  4. Practice different methods of sterilization and isolate pure culture of microorganism  5. Adapt the technique involved to see motility of bacteria i.e. hanging drop technique  6. Develop skill to execute morphology of bacteria by staining and determine quality of water by most probable number test (bacteriological analysis)  7. Differentiate Gram negative intestinal bacteria by performing IMVIC test  8. Learn standardization of pharmaceutical products microbiologically  1. Demonstrate theory and practical skills to prepare and view specimens using microscopy (bright field microscope) and staining procedures.  2. Practice aseptic techniques and be able to perform routine culture handling tasks safely and effectively  3. Use appropriate experimental microbiological lab equipment and methods  4. Recognize the various methods for identification of microorganisms and able to classify the bacteria  5. Develop culture media by understanding various physical
		growth requirements of bacteria  6. Write a document and report experimental protocol, results and conclusion
308P	Pharmaceutical Engineering —Practical (Div A)	<ol> <li>Determine radiation heat constant, overall heat transfer, moisture content, loss on drying, humidity of air, uniformity index by double cone blender.</li> <li>Explain the principle, instrumentation, working &amp; application of rotary tablet machine, fluidized bed coater, fluid energy mill, dehumidifier, colloidal mill, planetary mixer, fluidized bed dryer, freeze dryer</li> <li>Illustrate construction of drying curves, various size distribution curves &amp; verify laws of size reduction by using different equation</li> <li>Study the effect of time on rate of filtration, evaporation &amp; crystallization</li> <li>Demonstration of Steam distillation, colloid mill, planetary mixer, fluidized bed dryer, freeze dryer</li> </ol>
	Pharmaceutical Engineering –Practical (Div B)	<ol> <li>Determine radiation heat constant, overall heat transfer, moisture content, loss on drying, humidity of air, uniformity index by double cone blender.</li> <li>Explain the principle, instrumentation, working &amp; application of rotary tablet machine, fluidized bed coater, fluid energy mill, dehumidifier, colloidal mill, planetary mixer, fluidized bed dryer, freeze dryer</li> </ol>



		<ol> <li>Illustrate construction of drying curves, various size distribution curves &amp; verify laws of size reduction by using different equation</li> <li>Study the effect of time on rate of filtration, evaporation &amp; crystallization</li> <li>Demonstration of Steam distillation, colloid mill, planetary mixer, fluidized bed dryer, freeze dryer</li> </ol>
Second SEMES	year 2019 pattern	λ
BP401T	Pharmaceutical Organic Chemistry III– Theory (Div A)	<ol> <li>Understand the methods of preparation and properties of organic compounds and apply them to actual experiments.</li> <li>Know the stereo chemical aspects of organic compounds and stereo chemical reactions.</li> <li>Know the medicinal uses and other applications of organic compounds</li> <li>Classify heterocyclic compounds based on various criteria, predict their common as well as IUPAC name and recall their synthetic &amp; amp; physicochemical properties</li> <li>Identify various rearrangement reactions and conclude the mechanism in the formation of a particular compound</li> </ol>
	Pharmaceutical Organic Chemistry III- Theory (Div B)	<ol> <li>mechanism in the formation of a particular compound</li> <li>Understand the methods of preparation and properties of organic compounds and apply them to actual experiments.</li> <li>Know the stereo chemical aspects of organic compounds and stereo chemical reactions.</li> <li>Know the medicinal uses and other applications of organic compounds</li> <li>Classify heterocyclic compounds based on various criteria, predict their common as well as IUPAC name and recall their synthetic &amp; amp; physicochemical properties</li> <li>Identify various rearrangement reactions and conclude the mechanism in the formation of a particular compound.</li> </ol>
BP402T	Medicinal Chemistry I – Theory (Div A)	<ol> <li>Recall, apply and illustrate SAR of different classes of drugs</li> <li>Describe, apply, categorize and justify of Physiochemical properties of drug molecules in relation to drug ADME.</li> <li>Classify, discuss and memorize different classes of drugs and receptors under ANS and CNS</li> <li>Explain and determine some routes for the chemical synthesis of some drugs.</li> <li>Describe and explain an outline of drug metabolic pathways, adverse effects and therapeutic values.</li> <li>Compare and summarize the chemistry of drugs with respective to their pharmacological activity</li> </ol>
	Medicinal Chemistry I – Theory (Div B)	<ol> <li>Recall, apply and illustrate SAR of different classes of drugs</li> <li>Describe, apply, categorize and justify of Physiochemical properties of drug molecules in relation to drug ADME.</li> <li>Classify, discuss and memorize different classes of drugs and receptors under ANS and CNS</li> <li>Explain and determine some routes for the chemical synthesis of some drugs.</li> <li>Describe and explain an outline of drug metabolic pathways,</li> </ol>



3P405T	Pharmacognosy and	1. Summarize the case and allowed or importance and scope of
77.10.57	Discourand	dependence  1. Summarize the basic knowledge of importance and scope of import
		and to elaborate the concepts of drug addiction/abuse/tolerance/
		6. To predict the effects of drugs against neurodegenerative disorders
		5. To appraise the pharmacology of Psychopharmacological agents
		central nervous system.
.*		4. To analyze the functions of neurotransmitters and drugs acting on
		on peripheral nervous system
		3. To identify the role of neurohumoral transmission and drugs acting
		drug interactions and drug discovery
	(Div-B)	<ol> <li>To understand the basics of pharmacodynamics, adverse reactions,</li> </ol>
	Pharmacology I – Theory	pharmacokinetics.
(3=)	Dhamaaalagu I Thaam	To define the fundamental concepts of pharmacology and
		dependence
		6. To predict the effects of drugs against neurodegenerative disorders and to elaborate the concepts of drug addiction/abuse/tolerance/
	1/2	5. To appraise the pharmacology of Psychopharmacological agents
		central nervous system.
		4. To analyze the functions of neurotransmitters and drugs acting on
		on peripheral nervous system
	7 7 7	3. To identify the role of neurohumoral transmission and drugs acting
		drug interactions and drug discovery
		2. To understand the basics of pharmacodynamics, adverse reactions,
	(Div-A)	pharmacokinetics.
BP404T	Pharmacology I – Theory	To define the fundamental concepts of pharmacology and
	3/2 //	physical principles essential for pharmaceutical applications
		4. Recognize and Apply basic rules and equations regarding
		design and evaluation.
		3. Demonstrate ability to develop critical thinking and problem solving required to address problems related to dosage form
		systems, micromerities and rheological studies.
		studies, deformation, colloidal systems and coarse dispersion
		2. Demonstrate basics involved in reaction kinetics in stability
	- Theory (Div B)	drug and excipients in designing the dosage forms
	Physical Pharmaceutics II	1. Integrate and apply various physicochemical properties of
		physical principles essential for pharmaceutical applications
	Total Control	design and evaluation. 4. Recognize and Apply basic rules and equations regarding
	25/1	solving required to address problems related to dosage form
	3.21	3. Demonstrate ability to develop critical thinking and problem
		systems, micromerities and rheological studies.
		studies, deformation, colloidal systems and coarse dispersion
		2. Demonstrate basics involved in reaction kinetics in stability
	- Theory (Div A)	drug and excipients in designing the dosage forms
BP403T	Physical Pharmaceutics II	respective to their pharmacological activity  1. Integrate and apply various physicochemical properties of
	1	6. Compare and summarize the chemistry of drugs with



	Phytochemistry I– Theory	pharmacognosy and phytochemistry I in pharmacy field and its
	(Div- A)	role in various systems of medicines
		2. Apply knowledge of pharmacognostic study and quality
	5	control of crude drugs
		3. Describe biological source, chemical nature, proprties,
		identification tests and uses of crude drugs as well as plant
		products of natural origin ( Primary & secondary metabolites)
		4. Classify crude drugs and explain importance of cultivation,
		collection, processing, evaluation, preservation and storage of
	-	drugs of natural origin as well as plant tissue culture
	Pharmacognosy and Phytochemistry I— Theory	Summarize the basic knowledge of importance and scope of pharmacognosy and phytochemistry I in pharmacy field and its
	(Div-B)	role in various systems of medicines
	(BIV-B)	2. Apply knowledge of pharmacognostic study and quality
		control of crude drugs
		3. Describe biological source, chemical nature, proprties,
		identification tests and uses of crude drugs as well as plant
	: 1 -	products of natural origin (Primary & secondary metabolites)
	1.00	4. Classify crude drugs and explain importance of cultivation,
		collection, processing, evaluation, preservation and storage of
		drugs of natural origin as well as plant tissue culture
BP406P	Medicinal Chemistry I –	1. Understand, discuss and Synthesize the medicinally important
	Practical (Div A)	compounds / drug intermediates such as 1,3-pyrazole, 1,3-
		oxazole, Benzimidazole, Benztriazole, 2,3- diphenyl
		quinoxaline, Benzocaine, Phenytoin, Phenothiazine
		Barbiturate.
	47 (83.5)	2. Understand, determine and develop the recrystallization
		poceducre for organic compound and monitor reactions by
		TLC
		3. Determine, develop and apply the purification procedure for
		synthesized organic compounds by column chromatography
		4. Understand, discuss and determine and perform the procedure
	Fig.	of Partition coefficient and Ionization constants
	Medicinal Chemistry I –	1. Understand, discuss and Synthesize the medicinally important
	Practical (Div B)	compounds / drug intermediates such as 1,3-pyrazole, 1,3-
		oxazole, Benzimidazole, Benztriazole, 2,3- diphenyl
		quinoxaline, Benzocaine, Phenytoin, Phenothiazine
		Barbiturate.
		2. Understand, determine and develop the recrystallization
		poceducre for organic compound and monitor reactions by
		TLC
	x 4	
		3. Determine, develop and apply the purification procedure for
41		synthesized organic compounds by column chromatography
		4. Understand, discuss and determine and perform the procedure
		of Partition coefficient and Ionization constants
BP407P	Physical Pharmaceutics II	1. Exercise different pharmacoutical laboratory procedures used



	Prosting! (Div. A)	in 1-1 minima various physical properties such as micrometric
	- Practical (Div A)	in determining various physical properties such as micrometric
	10 - 10 - 10 - 10 - 10 - 10 - 10 - 10 -	properties, viscosity, order of reaction, etc.
	- L	2. Design and perform skillfully some laboratory experiments
		needed in pharmacy practice from preformulation and
1		formulation point of view
		3. Acquire and use technical vocabulary to discus
		pharmaceutical problems
		4. Employ proper documentation system to record observations
		and analyze information gathered through experimentation.
	Physical Pharmaceutics II	1. Exercise different pharmaceutical laboratory procedures used
	– Practical (Div B)	in determining various physical properties such as micrometric
		properties, viscosity, order of reaction, etc.
		2. Design and perform skillfully some laboratory experiments
	1 1 1 1 1 1 1 1 1 1 1	needed in pharmacy practice from preformulation and
		formulation point of view
		3. Acquire and use technical vocabulary to discuss
		pharmaceutical problems
		1. Employ proper documentation system to record observations
		and analyze information gathered through experimentation.
BP408P	Pharmacology I –	1. To learn about basic instruments, common laboratory animals
	Practical (Div A)	used in experimental pharmacology and to organize animal
		house as per the CPCSEA guidelines.
		2. To demonstrate the common laboratory techniques like routes
		of administration, blood withdrawal, anesthetics and
		euthanasia used for animal studies
		3. To interpret the effects of various drugs on rabbit eye and
		ciliary motility of frog oesophagus in correlation with humans
	372.7-	4. To analyse the effect of drugs acting as enzyme inducers,
		skeletal muscle relaxants and affecting locomotor activity in
		laboratory animal
		5. To evaluate the stereotype and anticatatonic activity of drugs
	Again the second second second	in rats/mice
	128	6. To predict various screening models for anticonvulsant
		anxiolytic and local anaesthetics activity
	Pharmacology I –	Students would have understood the different list of
	Practical (Div B)	equipment's and animals used for pharmacological actions of
		different categories of drugs
		system/cellular/ macromolecular levels.
		3. Demonstrate the use of EX- Pharma software to study effect of
		different categories of drug along with mechanism of action.
		4. To observed the effect of drugs on animals by simulated
		experiments.
	短背 : 1	5. Explain the ideas about correlation of pharmacology with
		other bio medical sciences
P409P	Pharmacognosy and	1. Understand, analyze and differentiate between the various
	Phytochemistry I –	crude drugs
	Practical (Div A)	2. Perform Quantitative analysis of the various crude drugs
		3. Design the extraction techniques by selecting suitable solvents
1		4. Analyze and differentiate between authentic and adulterated
	200	4. Thany's and differentiate between authentic and admicrated



		drugs
-	Pharmacognosy and	<ul><li>5. Develops hands on skills in micrometry</li><li>1. Understand, analyze and differentiate between the various</li></ul>
	Phytochemistry I – Practical (Div B)	crude drugs  2. Perform Quantitative analysis of the various crude drugs
		3. Design the extraction techniques by selecting suitable solvents
		<ol> <li>Analyze and differentiate between authentic and adulterated drugs</li> </ol>
Thind	2010 44	5. Develops hands on skills in micrometry
SEMEST	ar 2019 pattern CER-V	
BP501T	Medicinal Chemistry II –	1. Recall, apply and illustrate SAR of different classes of drugs
	Theory (Div A)	<ol> <li>Classify, discuss and memorize different classes of drugs and receptors under Antihistaminic, cardiovascular system, Endocrine system, antidiabetic and Local anesthetics</li> </ol>
		3. Explain and determine some routes for the chemical synthesis of some drugs.
		<ol> <li>Describe and explain an outline of drug metabolic pathways, adverse effects and therapeutic values.</li> </ol>
		5. Compare and summarize the chemistry of drugs with respective to their pharmacological activity
	Medicinal Chemistry II – Theory (Div B)	<ol> <li>Recall, apply and illustrate SAR of different classes of drugs</li> <li>Classify, discuss and memorize different classes of drugs and receptors under Antihistaminic, cardiovascular system</li> </ol>
		Endocrine system, antidiabetic and Local anesthetics 3. Explain and determine some routes for the chemical synthesis
		of some drugs.
		4. Describe and explain an outline of drug metabolic pathways adverse effects and therapeutic values.
		1. Compare and summarize the chemistry of drugs with respective to their pharmacological activity
BP502T	Industrial Pharmacy-I– Theory (Div A)	Describe the concept of pre-formulation study for the development of pharmaceutical dosage forms.
	na 1	2. Design solid and liquid dosage forms; describe their
		manufacture and evaluate them for their quality.
		3. Apply basic concepts of dosage forms; analyse techniques and equipment used in their manufacture and apply
		pharmacopoieal specifications during evaluation of solid, liquid dosage forms
		4. Construct sterile parenteral products and ophthalmic preparations; manufacture and evaluate them for their quality.
		5. Design cosmetic products, pharmaceutical aerosols and evaluate them. Describe pharmaceutical packaging materials
		required for various dosage forms and illustrate quality contro tests for them.
		6. Analyse techniques and equipment used in manufacture o sterile parenteral products, ophthalmic products, cosmetics and



	Dharmacognosy and	drugs 5. Develops hands on skills in micrometry 1. Understand, analyze and differentiate between the various
	Pharmacognosy and Phytochemistry I – Practical (Div B)	<ul> <li>crude drugs</li> <li>2. Perform Quantitative analysis of the various crude drugs</li> <li>3. Design the extraction techniques by selecting suitable solvents</li> <li>4. Analyze and differentiate between authentic and adulterated drugs</li> </ul>
		5. Develops hands on skills in micrometry
Third yes	ear 2019 pattern FER-V	2 4460
BP501T	Medicinal Chemistry II – Theory (Div A)	<ol> <li>Recall, apply and illustrate SAR of different classes of drugs</li> <li>Classify, discuss and memorize different classes of drugs and receptors under Antihistaminic, cardiovascular system, Endocrine system, antidiabetic and Local anesthetics</li> <li>Explain and determine some routes for the chemical synthesis of some drugs.</li> <li>Describe and explain an outline of drug metabolic pathways, adverse effects and therapeutic values.</li> <li>Compare and summarize the chemistry of drugs with respective to their pharmacological activity</li> </ol>
	Medicinal Chemistry II – Theory (Div B)	<ol> <li>Recall, apply and illustrate SAR of different classes of drugs</li> <li>Classify, discuss and memorize different classes of drugs and receptors under Antihistaminic, cardiovascular system, Endocrine system, antidiabetic and Local anesthetics</li> <li>Explain and determine some routes for the chemical synthesis of some drugs.</li> <li>Describe and explain an outline of drug metabolic pathways, adverse effects and therapeutic values.</li> <li>Compare and summarize the chemistry of drugs with respective to their pharmacological activity</li> </ol>
BP502T	Industrial Pharmacy-I— Theory (Div A)	<ol> <li>Describe the concept of pre-formulation study for the development of pharmaceutical dosage forms.</li> <li>Design solid and liquid dosage forms; describe their manufacture and evaluate them for their quality.</li> <li>Apply basic concepts of dosage forms; analyse techniques and equipment used in their manufacture and apply pharmacopoieal specifications during evaluation of solid, liquid dosage forms</li> <li>Construct sterile parenteral products and ophthalmic preparations; manufacture and evaluate them for their quality.</li> <li>Design cosmetic products, pharmaceutical aerosols and evaluate them. Describe pharmaceutical packaging materials required for various dosage forms and illustrate quality control tests for them.</li> <li>Analyse techniques and equipment used in manufacture of sterile parenteral products, ophthalmic products, cosmetics and</li> </ol>



		specifications during their evaluation and interpret their quality.
	Industrial Pharmacy-I– Theory (Div B)	<ol> <li>Describe the concept of pre-formulation study for the development of pharmaceutical dosage forms.</li> <li>Design solid and liquid dosage forms; describe their manufacture and evaluate them for their quality.</li> <li>Apply basic concepts of dosage forms; analyse techniques and</li> </ol>
		equipment used in their manufacture and apply pharmacopoical specifications during evaluation of solid, liquid dosage forms  4. Construct sterile parenteral products and ophthalmic
		preparations; manufacture and evaluate them for their quality.  5. Design cosmetic products, pharmaceutical aerosols and evaluate them. Describe pharmaceutical packaging materials
		required for various dosage forms and illustrate quality control tests for them.  6. Analyse techniques and equipment used in manufacture of
		sterile parenteral products, ophthalmic products, cosmetics and pharmaceutical aerosols and apply pharmacopoieal specifications during their evaluation and interpret their quality.
BP503T	Pharmacology II –	1. Describe, explain and summarize the classification,
	Theory (Div-A)	mechanism of action, pharmacological action,
		pharmacokinetic, therapeutic uses, adverse drug reaction,
•		drug interaction and contraindications of drug acting on
		Cardiovascular system, endocrine system and urinary system
		2. Describe the pharmacology of drugs used in the management
		of cardiovascular diseases.
		3. To study the pharmacology of drugs used as Diuretics and
		anti-diuretics.
		4. Describe pharmacology of drugs that affect the major
		endocrine gland and their hormones  5. To study the principles and applications of bioassay
7		6. Explain and outline correlation of pharmacology with related
		medical sciences.
	Pharmacology II –	Describe, explain and summarize the classification,
	Theory (Div-B)	mechanism of action, pharmacological action,
		pharmacokinetic, therapeutic uses, adverse drug reaction,
		drug interaction and contraindications of drug acting on
		Cardiovascular system, endocrine system and urinary system
		2. Describe the pharmacology of drugs used in the management
* *2 -		of cardiovascular diseases.
		3. To study the pharmacology of drugs used as Diuretics and
	A	anti-diuretics.  4. Describe pharmacology of drugs that affect the major
		endocrine gland and their hormones
		5. To study the principles and applications of bioassay
		6. Explain and outline correlation of pharmacology with related
		medical sciences.



BP504T	Pharmacognosy and phytochemistry II– Theory (Div A)	Sketch & relate the origin of phytoconstituents by biogenetic pathway & summarize the various techniques for their investigation.
		<ul> <li>investigation</li> <li>Develop and design the technique for extraction, isolation, purification of phytoconstituents for medicinal, agricultural,</li> </ul>
		food & healthcare industry
		3. Design & explain various methods for determination of
		authenticity of crude drugs, extracts, phytoconstituents,
		formulation & detect their adulteration
		4. Develop themselves to design a method for analysis, detection,
		purification of various extracts, phytoconstituents & herbal
		raw materials.
		5. Correlate various techniques involved in extraction, isolation,
	Pharmacognosy and	estimation & application of phytoconstituents  1. Sketch & relate the origin of phytoconstituents by biogenetic
	phytochemistry II— Theory (Div B)	pathway & summarize the various techniques for their investigation
	1, 2, 2, 3	2. Develop and design the technique for extraction, isolation,
		purification of phytoconstituents for medicinal, agricultural,
		food & healthcare industry
		3. Design & explain various methods for determination of authenticity of crude drugs, extracts, phytoconstituents,
		formulation & detect their adulteration
		4. Develop themselves to design a method for analysis, detection,
		purification of various extracts, phytoconstituents & herbal
		raw materials.
		<ol> <li>Correlate various techniques involved in extraction, isolation, estimation &amp; application of phytoconstituents</li> </ol>
BP505T	Pharmaceutical	To impart basic knowledge on important legislations related to
	Jurisprudence – Theory	the profession of pharmacy in India.
	(Div A)	2. To understand definition, committees, rules, scope,
		functioning and offences and penalties of various Indian
	Section 25	pharmaceutical Acts and Laws
		3. To understand objective and ethics of various pharmaceutical
		Acts and Laws.
		4. To impart basic knowledge of development of Pharmaceutical
		Legislation and enforcement of various Pharmaceutical Acts
		and laws.
		5. To be able to relate dynamics of pharmaceutical profession to
		the Acts and Laws applicable
		6. To be able to analyse dos and donts of pharmacy profession
	Pharmaceutical	keeping pharmaceutical Acts and Laws as the reference.  1. To impart basic knowledge on important legislations related to
	Jurisprudence – Theory	the profession of pharmacy in India.
	(Div B)	2. To understand definition, committees, rules, scope,
		functioning and offences and penalties of various Indian
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		pharmaceutical Acts and Laws 3. To understand objective and ethics of various pharmaceutical
		Acts and Laws.  4. To impart basic knowledge of development of Pharmaceutical
		Legislation and enforcement of various Pharmaceutical Acts and laws.
		5. To be able to relate dynamics of pharmaceutical profession to the Acts and Laws applicable.
		6. To understand the pharmacy profession keeping pharmaceutical Acts and Laws as the reference.
BP506P	Industrial Pharmacy-I – Practical (Div A)	Formulate solid and liquid dosage forms, manufacture and evaluate them.
		Perform calculations needed to perform evaluation tests of solid and liquid dosage forms and interpret the results
		3. Analyse the rational and use of ingredients in formulation and category of respective formulation
		<ol> <li>Select and use appropriate equipment and apparatus needed for the particular preparation</li> </ol>
		5. Formulate sterile parenteral products and ophthalmic preparations, prepare and evaluate them.
		<ol> <li>Prepare labels to suit regulatory requirements and select proper packaging (container and closure) and labeling materials for preparations</li> </ol>
	Industrial Pharmacy-I – Practical (Div B)	Formulate solid and liquid dosage forms, manufacture and evaluate them.
		<ol> <li>Perform calculations needed to perform evaluation tests of solid and liquid dosage forms and interpret the results</li> </ol>
		<ol> <li>Analyse the rational and use of ingredients in formulation and category of respective formulation</li> </ol>
		<ol> <li>Select and use appropriate equipment and apparatus needed for the particular preparation</li> </ol>
		5. Formulate sterile parenteral products and ophthalmic preparations, prepare and evaluate them.
		<ol> <li>Prepare labels to suit regulatory requirements and select proper packaging (container and closure) and labeling materials for preparations</li> </ol>
3P507P	Pharmacology II – Practical (Div A)	<ol> <li>Demonstrate the biological evaluation of drugs by in vitro and in vivo methods and to study different physiological salt</li> </ol>
		solutions  2. Analyze knowledge how to deal with experimental animals to test the potency of drugs.
		<ol> <li>Compare, select, locate and isolate different organs/tissues from the laboratory animals used in pharmacological experiments.</li> </ol>
	and the second s	4. Develop skills in handling animals and perform and evaluate



	experiments on them  5. Calculate and compare experimental observations and results statistically
Pharmacology II – Practical (Div B)	Demonstrate the biological evaluation of drugs by in vitro and in vivo methods and to study different physiological salt solutions
	<ol> <li>Analyze knowledge how to deal with experimental animals to test the potency of drugs.</li> <li>Compare, select, locate and isolate different organs/tissues</li> </ol>
	from the laboratory animals used in pharmacological experiments.  4. Develop skills in handling animals and perform and
	<ul> <li>evaluate experiments on them</li> <li>5. Calculate and compare experimental observations and results statistically</li> </ul>
Pharmacognosy and Phytochemistry II – Practical (Div A)	<ol> <li>Modify the techniques for isolation &amp; Develop &amp; modify the techniques involved in analysis of crude drugs &amp; their extracts</li> <li>Design various analytical parameters required for the</li> </ol>
	<ul> <li>authentication of crude drugs, Extracts, Oils etc.</li> <li>Judge various crude drugs on the basis of their morphological &amp; microscopical Studies.</li> <li>Evaluate the purity &amp;/or stability of various unorganized crude</li> </ul>
Pharmacognosy and Phytochemistry II – Practical (Div B)	drugs & oils.  1. Modify the techniques for isolation & Develop & modify the techniques involved in analysis of crude drugs & their extracts
	<ol> <li>Design various analytical parameters required for the authentication of crude drugs, Extracts, Oils etc.</li> <li>Judge various crude drugs on the basis of their morphological &amp; microscopical Studies.</li> <li>Evaluate the purity &amp;/or stability of various unorganized crude drugs &amp; oils.</li> </ol>
2019 pattern	
Medicinal Chemistry III – Theory (Div A)	<ol> <li>Recall, apply and illustrate SAR of different classes of drugs</li> <li>Understand and apply the chemistry of drugs with respective to their biological activity.</li> <li>Describe and explain an outline of drug metabolic pathways, adverse effects and therapeutic values.</li> <li>Explain and determine some routes for the chemical synthesis of some drugs.</li> <li>Understand and apply importance of drug design and different techniques of drug design.</li> </ol>
	Pharmacognosy and Phytochemistry II – Practical (Div A)  Pharmacognosy and Phytochemistry II – Practical (Div B)  2019 pattern R-VI Medicinal Chemistry III –



		6. Classify, discuss and memorize different classes of drugs under Antibiotics, anti-infective, antimalarial, antimycobaterial, antiviral, antifungal, anttiprotozoal,
		anthalmintic and antineonlastic agents.
	Medicinal Chemistry III – Theory (Div B)	
		<ul> <li>4. Explain and determine some routes for the chemical synthesis of some drugs.</li> <li>5. Understand and apply importance of drug design and different</li> </ul>
		techniques of drug design.  6. Classify, discuss and memorize different classes of drugs under Antibiotics, anti-infective, antimalarial,
		anthelmintic and antineoplastic agents.
BP602T	Pharmacology III – Theory (Div A)	Understand the pharmacological concepts of drugs acting on respiratory system.
		2. Understand pharmacological concepts of drugs acting on Gastrointestinal tract.
		<ul><li>3. Clarify the pharmacological features of common and important drugs as antibiotics and chemotherapeutic agents.</li><li>4. Describe the pharmacology of immunosuppressant and</li></ul>
		immunostimulants. 5. Explain principles of toxicology and treatment of various
		poisonings. 6. Correlate toxicology and its pharmacology with different types of disease and poisoning.
	Pharmacology III – Theory (Div B)	Understand the pharmacological concepts of drugs acting on respiratory system.
		2. Understand pharmacological concepts of drugs acting on Gastrointestinal tract.
		3. Clarify the pharmacological features of common and important drugs as antibiotics and chemotherapeutic agents.
		4. Describe the pharmacology of immunosuppressant and immunostimulants.
		5. Explain principles of toxicology and treatment of various poisonings.
		<ol> <li>Correlate toxicology and its pharmacology with different types of disease and poisoning.</li> </ol>
BP603T	Herbal Drug Technology  – Theory (Div A)	1. Describe and explain herbs as a raw materials source from the cultivation of herbal drug products.
		2. Discuss and apply various guidelines issued by WHO in relation to cultivation, collection, storage, etc. in order to
		ethically develop pharmaceutical dosage forms.  3. Describe and explain the concept of health and pathogenesis
		philosophical basis, diagnosis & treatment aspects of Ayurveda, Unani, Siddha & Homeopathic system of medicine
	-12	understand & explain the method of preparation of Ayurvedic

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	Herbal Drug Technology	delivery of natural products like liposomes, phytosomes; herbs used in cosmetic preparation & the method of their formulations.  4. Understand health benefits and potentials of Nutraceuticals, Describe and explain Herbal-Drug and Herb-food interactions.  5. Describe and explain Good Manufacturing Practices (GMP), patenting and regulatory issues of herbal drugs
	Herbal Drug Technology	Describe and explain Herbal-Drug and Herb-food interactions.  5. Describe and explain Good Manufacturing Practices (GMP).
1.	Herbal Drug Technology	patenting and regulatory issues of herbal drugs
	Herbal Drug Technology	terming and regulatory issues of fictival dides
	- Theory (Div B)	<ol> <li>Describe and explain herbs as a raw materials source from the cultivation of herbal drug products.</li> <li>Discuss and apply various guidelines issued by WHO in relation to cultivation, collection, storage, etc. in order to ethically develop pharmaceutical dosage forms.</li> <li>Describe and explain the concept of health and pathogenesis, philosophical basis, diagnosis &amp; treatment aspects of Ayurveda, Unani, Siddha &amp; Homeopathic system of medicine; understand &amp; explain the method of preparation of Ayurvedic</li> </ol>
		dosage forms; understand the importance of novel drug delivery of natural products like liposomes, phytosomes; herbs used in cosmetic preparation & the method of their formulations.  7. Understand health benefits and potentials of Nutraceuticals, Describe and explain Herbal-Drug and Herb-food interactions. Describe and explain Good Manufacturing Practices (GMP), patenting and regulatory issues of herbal drugs
BP604T	Biopharmaceutics and	Understand the basic concepts in biopharmaceutics and
	Pharmacokinetics – Theory (Div A)	pharmacokinetics and their significance
. "	, (= = : = )	2. Calculate the plasma drug concentration-time data & the
		<ul> <li>pharmacokinetic parameters</li> <li>3. Describe the processes &amp; kinetics of drug absorption, distribution, metabolism, excretion, elimination.</li> <li>4. Discuss &amp; Understand the concepts of bioavailability and bioequivalence of drug products and their significance</li> <li>5. Collaborate the concept of dissolution and application of in vitro in vivo correlation in drug product development</li> </ul>
F	Biopharmaceutics and Pharmacokinetics – Theory (Div B)	Understand the basic concepts in biopharmaceutics and pharmacokinetics and their significance     Calculate the plasma drug concentration-time data & the pharmacokinetic parameters
		<ol> <li>Describe the processes &amp; kinetics of drug absorption, distribution, metabolism, excretion, elimination.</li> <li>Discuss &amp; Understand the concepts of bioavailability and bioequivalence of drug products and their significance</li> <li>Collaborate the concept of dissolution and application of in</li> </ol>
В	Pharmaceutical Biotechnology – Theory Div A)	vitro in vivo correlation in drug product development.  1. Summarize the basic knowledge of Biotechnology and its scope in pharmacy  2. To assess understanding about use of advanced biotechnological terms, principles and methods to solve



	<ul> <li>biotechnological tasks.</li> <li>3. To apply knowledge of immunology in healthcare and diagnostic purposes</li> <li>4. To clarify equipments and the steps involved in production of biotechnologically derived product as well as methods to develop modern techniques in biotechnology field helpful to</li> </ul>
The state of the s	society
Biotechnology – Theory (Div B)	<ol> <li>Summarize the basic knowledge of Biotechnology and its scope in pharmacy</li> <li>To assess understanding about use of advanced biotechnological terms, principles and methods to solve biotechnological tasks.</li> <li>To apply knowledge of immunology in healthcare and diagnostic purposes</li> <li>To clarify equipments and the steps involved in production of biotechnologically derived product as well as methods to develop modern techniques in biotechnology field helpful to society</li> </ol>
Quality Assurance  -Theory (Div A)	<ol> <li>Understand and analyze cGMP aspects in a pharmaceutical industry</li> <li>Explain the concept of quality management and elaborate quality responsibilities of QA &amp; QC departments</li> <li>Compile and discuss quality certifications applicable to pharmaceutical industries and their scope</li> <li>Understand document maintenance in pharmaceutical industry and judge the importance of documentation.</li> <li>Explain quality control of packaging materials; understand good laboratory practices and identify its importance.</li> <li>Outline calibration and validation principles and good warehouse practices</li> </ol>
Quality Assurance  -Theory (Div B)	<ol> <li>Understand and analyze cGMP aspects in a pharmaceutical industry</li> <li>Explain the concept of quality management and elaborate quality responsibilities of QA &amp; QC departments</li> <li>Compile and discuss quality certifications applicable to pharmaceutical industries and their scope</li> <li>Understand document maintenance in pharmaceutical industry and judge the importance of documentation.</li> <li>Explain quality control of packaging materials; understand good laboratory practices and identify its importance.         <ul> <li>Outline calibration and validation principles and good warehouse practices</li> </ul> </li> </ol>
Medicinal chemistry III – Practical (Div A)	<ol> <li>Use of Various equipment &amp; and take safety measures while working in medical chemistry laboratory</li> <li>Explain, apply, and justify the principle and theory behind synthesis of compound of our interest</li> <li>Develop and apply skill to synthesize compound and purify synthesized compounds.</li> <li>Develop and apply skill to synthesize by microwave assisted</li> </ol>
	Quality Assurance -Theory (Div A)  Quality Assurance -Theory (Div B)



		synthesis as green chemistry approach.
		5. Predict and derive physicochemical properties of synthesized
		compound.
		6. Develop virtual learning and skill of drawing structures and
		determining physicochemical parameters using online free
		software.
	Medicinal chemistry III -	1. Use of Various equipment & and take safety measures while
	Practical (Div B)	working in medical chemistry laboratory
		2. Explain, apply, and justify the principle and theory behind
		synthesis of compound of our interest
		3. Develop and apply skill to synthesize compound and purify
	2,000	synthesized compounds.
	100	4. Develop and apply skill to synthesize by microwave assisted
		synthesis as green chemistry approach.
		5. Predict and derive physicochemical properties of synthesized
		compound.
	5= <sub>0</sub>	6. Develop virtual learning and skill of drawing structures and
	7	determining physicochemical parameters using online free
DDCOOD	DI	software.
BP608P	Pharmacology III –	1. Understand the effect of drugs on various muscles and tissue
	Practical (Div A)	on animal.
	12 22	2. Study toxicity of drug using given data.
		3. Evaluate corrosion/irritation of test substances on animals.
	Pharmacology III –	4. Study Biostatistics methods in experimental pharmacology.
	Practical (Div B)	<ol> <li>Understand the effect of drugs on various muscles and tissues on animal.</li> </ol>
	Tractical (BIV B)	
		<ol> <li>Study toxicity of drug using given data.</li> <li>Evaluate corrosion/irritation of test substances on animals.</li> </ol>
		Study Biostatistics methods in experimental pharmacology
BP609P	Herbal Drug Technology	Classify and analyze various herbal raw materials required for
	- Practical (Div A)	formulation of herbal products.
	1100.001 (21.11)	2. Develop skills for evaluation of various Ayurvedic
		formulations such as Asava, Arista. Design and evaluate
	15.4	various excipients of natural origin.
		3. Design, evaluate, optimize & develop cosmetic
	Water the second	formulations like creams, lotions and shampoos.
		4. Design, evaluate, optimize & develop standardized extract in
		formulations like syrups, mixtures and tablets.
		5. Discuss about the analysis of a Monograph of herbal drugs
		from recent Pharmacopoeias.
	81.55	6. Design, optimize analysis methods for determination of
		aldehydes, phenols and alkaloids in medicinal herbs, herbal
	, a	products.
	Herbal Drug Technology	1. Classify and analyze various herbal raw materials required for
	- Practical (Div B)	formulation of herbal products.
		2. Develop skills for evaluation of various Avurvedic
	A A A A A A A A A A A A A A A A A A A	formulations such as Asava, Arista. Design and evaluate
		various excipients of natural origin.
		3. Design, evaluate, optimize & amp; develop cosmetic
- Nega		A

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		formulations like creams, lotions and shampoos.  4. Design, evaluate, optimize & develop standardized extract in
		Ioinulations like syrups, mixtures and tablets
		5. Discuss about the analysis of a Monograph of herbal drug from recent Pharmacopoeias.
		6. Design, optimize analysis methods for determination o
		aldehydes, phenols and alkaloids in medicinal herbs, herba
Final yea SEMEST	nr 2019 pattern	products
BP701		
T	Instrumental Methods of Analysis- Theory	1. To explain, discuss and integrate theory and principle o
	(Div A)	various analytical techniques like UV/Visible Spectroscopy
		FTIR spectroscopy, Fluorimetry, Flame Photometry, Atomic
		absorption spectroscopy, Nepheloturbidimetry, Adsorption and partition column chromatography, Paper chromatography
		Thin layer chromatography, HPTLC, Theory o
		Chromatography, Gas chromatography, High performance
		ilquid chromatography (HPLC), Ion exchange
		chromatography, Gel chromatography.
		2. To explain, discuss, compose, demonstrate and evaluate instrumentation in UV/Visible Spectroscopy, FTIR
		instrumentation in UV/Visible Spectroscopy, FTIR spectroscopy, Fluorimetry, Flame Photometry, Atomic
	2 -3 -	absorption spectroscopy, Nepheloturbidimetry, Adsorption and
	* " "	partition column chromatography, Paper chromatography
		Thin layer chromatography, HPTLC, Theory of
		Chromatography, Gas chromatography, High performance
		liquid chromatography (HPLC), Ion exchange chromatography, Gel chromatography.
		3. To apply, compare, rank and justify the analytical tools
		UV/Visible Spectroscopy, FTIR spectroscopy, Fluorimetry
	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	Flame Photometry, Atomic absorption spectroscopy
		Nepheloturbidimetry, Adsorption and partition column
		chromatography, Paper chromatography, Thin layer
		chromatography, HPTLC, Theory of Chromatography, Gaschromatography, High performance liquid chromatography
	N.	(HPLC), Ion exchange chromatography, Gel chromatography
		4. To summarize, integrate, analyze, critique, compare and
		discuss UV/Visible Spectroscopy, FTIR spectroscopy
		Fluorimetry, Flame Photometry, Atomic absorption
		spectroscopy, Nepheloturbidimetry, Adsorption and partition column chromatography, Paper chromatography, Thin layer
		chromatography, HPTLC, Theory of Chromatography, Gas
	To Carlo	chromatography, High performance liquid chromatography
	Instrumental Methods	(HPLC), Ion exchange chromatography, Gel chromatography
	of Analysis- Theory	To describe, discuss and summarize theory and principle of analytical techniques like UV Visible spectroscopy.
	(Div B)	analytical techniques like UV Visible spectroscopy, Fluorimetry, IR spectroscopy, Flame Photometry, Atomic
	` ′	absorption spectroscopy, Nepheloturbidometry, Adsorption
		and partition column chromatography Thin layer
		chromatography, Paper chromatography, Electrophoresis, Gas



		chromatography, High performance liquid chromatography
		(HPLC), Ion exchange chromatography, Gel chromatography,
		Affinity chromatography
		2. Explain, discuss, and demonstrate in details the above
1		instruments.
		3. Understand, discuss and compare the applications of
		instruments.
		4. Understand, discuss, interpret and elucidate the data obtained
		from above spectroscopic instruments.
BP702	Industrial pharmacy-II-	1. Understand and analyse the process of pilot plant and scale up
T	Theory (Div A)	of pharmaceutical dosage forms
		2. Explain and elaborate the process of technology transfer from
		lab scale to commercial batch
		3. Compile and discuss different Laws and Acts that regulate
-		pharmaceutical industry
	2 (1)	4. Compile and explain different quality management systems in
		pharmaceutical industry
		5. Understand the approval process and regulatory requirements
	1	for drug products
	Industrial pharmacy-II-	1. Understand and analyse the process of pilot plant and scale up
	Theory (Div B)	of pharmaceutical dosage forms
İ		2. Explain and elaborate the process of technology transfer from
		lab scale to commercial batch
		3. Compile and discuss different Laws and Acts that regulate
		pharmaceutical industry
		4. Compile and explain different quality management systems in
		pharmaceutical industry
		5. Understand the approval process and regulatory requirements
BP703	Pharmacy Practice –	for drug products
T 703	Theory (Div A)	1. Apply the basic knowledge of various drug distribution methods, pharmacy stores management and inventory control
•	Theory (DIV A)	procedures.
		•
		<ol> <li>Classify the drug interactions and adverse drug reactions.</li> <li>Describe and justify the concept of hospital formulary,</li> </ol>
		therapeutic drug monitoring and Patient counselling.
		4. Understand the ethical concerns in clinical pharmacy.
		5. Apply the concepts of professional ethics by producing safe
		and appropriate medication use throughout society.
		6. Interpret and justify the use of various clinical laboratory tests.
	Pharmacy Practice –	Understand and apply all the basic knowledge related to the
	Theory (Div B)	hospital administration, inventory procedures and Pharmacy
1		management as well as drug interactions.
		2. Classify and understand the drug stores in community
		pharmacy and its management.
		3. Describe and justify the concept of hospital formulary,
		therapeutic drug monitoring and Patient counselling.
	7 24 - 1	4. Understand the ethical concerns in clinical pharmacy.
		5. Apply the concepts of professional ethics by producing safe
		and appropriate medication use throughout society.
	· · ·	6. Interpret and justify the use of various glinical laboratory tests.
		A / many many me and of various difficult faboratory tests.

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BP704	N 15	
T T	Novel Drug Delivery System- Theory (Div A)	<ol> <li>Illustrate the basic knowledge of novel drug delivery system concept with respect to classification and definition of various novel drug delivery systems.</li> <li>Illustrate the basic knowledge for selection of drug and various excipients for designing and formulation of novel drug delivery system</li> <li>Discuss the application, advantages and disadvantages of novel drug delivery system over conventional dosage forms.</li> <li>Discuss the various approaches for development of NDDS.</li> </ol>
	Novel Drug Delivery System- Theory (Div B)	<ol> <li>Illustrate the basic knowledge of novel drug delivery system concept with respect to classification and definition of various novel drug delivery systems.</li> <li>Illustrate the basic knowledge for selection of drug and various excipients for designing and formulation of novel drug delivery system</li> <li>Discuss the application, advantages and disadvantages of novel drug delivery system over conventional dosage forms.</li> </ol>
BP705 P	Instrumental Methods of Analysis- Practical (Div A)	<ol> <li>Discuss the various approaches for development of NDDS.</li> <li>To understand, discuss and determine working principle of instruments UV/Visible spectrophotometer, Flame photometer, fluorescence spectrophotometer, colorimeter, HPLC, FTIR, TLC and paper chromatography.</li> </ol>
		<ol> <li>To explain, discuss, demonstrate and perform the practical procedure for quantitative and qualitative analysis of drugs or ions.</li> <li>To interpret and summarize the data obtained from UV/Visible spectrophotometer, Flame photometer, fluorescence spectrophotometer, colorimeter, HPLC, FTIR, TLC and paper chromatography.</li> <li>To calculate and estimate the results from observations and summarized data.</li> </ol>
	Instrumental Methods of Analysis- Practical (Div B)	<ol> <li>Understand, discuss and determine the working principles of instruments such as UV, colorimeter, fluorimetry, flame photometry, HPLC, FTIR, TLC and Paper Chromatography.</li> <li>Explain, discuss, and demonstrate the practical procedure for quantitative and qualitative analysis of drugs or ions</li> <li>Interpret and summarize the data obtained from instruments</li> <li>Calculate and estimate the results from summarized data.</li> </ol>
BP706PS	Practice School* (Div A)	<ol> <li>To develop and illustrate the techniques for quantification of API and Formulation by Modern Analytical instruments</li> <li>To Plan various quality control parameters, understand the Good Laboratory Practices and Document maintenance required in a pharmaceutical industry.</li> <li>Develop the hands-on expertise in animal handling as well as on the advanced instruments such as auto analyzer and hematology analyzer.</li> </ol>
college	Practice School* (Div B)	<ul> <li>4. Develop basic skills, demonstrate use of modern equipments in formulation development and documentation practices.</li> <li>1. To develop and illustrate the techniques for quantification of API and Formulation by Modern Analytical instruments</li> </ul>



?806ET	Quality Control and Standardization of Herbals (Div A)	<ol> <li>development.</li> <li>Evaluate alternative ways of solving problems related to healt and pharmaceutical issues.</li> <li>Describe the Objectives, functioning and importance of national programmes for prevention and control of diseases.</li> <li>Outline community services offered in urban and rural areas with respect to prevention and control of communicable and non communicable disease.</li> <li>Illustrate the general measures of prevention and control of infections and diseases.</li> <li>Select and compare various test protocols and parameters for the standardization of herbal drugs and formulations</li> <li>Link and hypothesize the parameters for evaluation and to</li> </ol>
		<ol> <li>Evaluate alternative ways of solving problems related to healt and pharmaceutical issues.</li> <li>Describe the Objectives, functioning and importance of national programmes for prevention and control of diseases.</li> <li>Outline community services offered in urban and rural areas with respect to prevention and control of communicable and non communicable disease.</li> <li>Illustrate the general measures of prevention and control of infections and diseases.</li> </ol>
		<ol> <li>Evaluate alternative ways of solving problems related to healt and pharmaceutical issues.</li> <li>Describe the Objectives, functioning and importance of national programmes for prevention and control of diseases.</li> <li>Outline community services offered in urban and rural areas</li> </ol>
		3. Evaluate alternative ways of solving problems related to healt and pharmaceutical issues.
	Social and Preventive Pharmacy (Div B)	<ol> <li>Understand and solve the current issues related to health and pharmaceutical problems within the country and worldwide.</li> <li>Develop a critical way of thinking based on current healthcare</li> </ol>
, ,	4	<ul> <li>national programmes for prevention and control of diseases.</li> <li>Outline community services offered in urban and rural areas.</li> <li>Illustrate the general measures of prevention and control of infections and diseases</li> </ul>
		<ol> <li>Evaluate alternative ways of solving problems related to healt and pharmaceutical issues.</li> <li>Describe the Objectives, functioning and importance of</li> </ol>
	Pharmacy	<ol><li>Develop a critical way of thinking based on current healthcare development.</li></ol>
BP802T	Social and Preventive Pharmacy	<ol> <li>Understand and solve the current issues related to health and pharmaceutical problems within the country and worldwide.</li> </ol>
Final y 2019 Pa BP801T	Biostatistics and Research Methodology	required in a pharmaceutical industry.  3. Develop the hands-on expertise in animal handling as well on the advanced instruments such as auto analyzer are hematology analyzer.  4. Develop basic skills, demonstrate use of modern equipment in formulation development and documentation practices.  1. Demonstrate knowledge about basic concepts about research and the methodology to conduct industrial and clinic research.  2. Explain the basic statistical concepts, terminology, tool Regression calculation and parametric test.  3. Describe the appropriate statistical methods required for particular research design Non parametric test, Introduction research, Graphs and designing the methodology.  4. Demonstrate the use of softwares like M.S. Excel, SPSS, and MINITAB, Hypothesis testing and clinical trial.  5. Demonstrate use of tools for design, analysis and optimization of experiments like factorial designs, RSM methodology are response surface methodology.



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		apply various techniques for the standardisation of Herbal
		products.
		3. Differentiate between the raw material, finished products
		based upon assessing their quality
		4. Explain, Correlate and implement various guidelines for
		sustaining the quality, safety and efficacy of neroal materials
		and also for their registration in Indian and International
		markets
	Quality Control and	1 Select and compare various test protocols and parameters
	Standardization of	for the standardization of herbal drugs and formulations
	Herbals ( Div B )	2 Link and hypothesize the parameters for evaluation and to
		apply various techniques for the standardisation of Herbal
		products.
	7-1	3. Differentiate between the raw material, finished products
		based upon assessing their quality
		4. Explain, Correlate and implement various guidelines for
		sustaining the quality, safety and efficacy of herbal
		materials and also for their registration in Indian and
		International markets
BP808ET	Cell and Molecular	Understand cell and molecular biology history, cellular
Di cocbi	Biology- Theory (Div A)	composition and functioning of cell biology.
	Dielogy Theory (Bit It)	2. Learn chemical foundations of cell biology.
		3. To explain cellular membrane structure and function
	- 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	properties and functions of DNA, Cell Cycle.
	1622 64,	properties and functions of DNA, cell cycle.
		4. Depict basic molecular genetics mechanisms.
		5. Comprehend the cell signaling pathways with their regulations
	Cell and Molecular	1. Understand cell and molecular biology history, cellular
	Biology- Theory (Div B)	composition and functioning of cell biology.
		2. Learn chemical foundations of cell biology.
		3. To explain cellular membrane structure and function
		properties and functions of DNA, Cell Cycle.
		4. Depict basic molecular genetics mechanisms.
		5. Comprehend the cell signaling pathways with their regulations
BP809ET	Cosmetic Science-	1. Discuss the fundamental & scope in cosmetic science &
	Theory (Div A)	explain the concepts and summarize general additives, water,
		perfume, preservatives colors, humectants, binders used in
	V.	cosmetics
		2. Explain theoretical aspects in formulation and evaluation of
		skin care products, hair care products, shaving preparation,
		baby care products & dental preparation.
		3. Develop and evaluate skin care products, hair care products,
		shaving preparation, baby care products, nail care product &
		dental preparation.
		4. Design and develop cosmetics formulation for hair, eyes, nail,
		dental and baby care products
		5. Illustrate the concept of cosmetics, cosmeceuticals &
		cosmeceutical agents
	Cosmetic Science-	1. Define and classify cosmetic & cosmecutical product,
	-3	Cosmetic excipients, skin care product, Hair care product, sun
	Theory (Div B)	
А		protection product.
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		<ol> <li>Describe Advantage, disadvantage and application of cosmetic &amp; cosmecutical product, Cosmetic excipients, skin care product, Hair care product, sun protection product.</li> <li>Illustrate the evaluation parameters and analytical method of Cosmetic product, cosmetic excipients, skin care product, Hair care product, sun protection product.</li> <li>Explain the problem associated with oral cavity, skin and hair.</li> <li>Understand the principle, mechanism of action and formulation aspect of skin, hair, oral cavity, antiperspants and deodorants</li> </ol>
BP811ET	Advanced Instrumentation Techniques (Div A& B)	<ol> <li>Understand, discuss and determine in detail the theory and principle of H- NMR, C- NMR spectroscopy, mass spectrometry, thermal methods, X-Ray diffraction method, calibration, of radio immune assay, electrophoresis, extraction techniques and hyphenated techniques.</li> <li>Explain, discuss, and demonstrate in details the above instruments.</li> </ol>
		<ol> <li>Understand, discuss and compare the applications of instruments.</li> <li>Understand, discuss, interpret and elucidate the data obtained from above spectroscopic instruments.</li> </ol>



## PES Modern College of Pharmacy, Nigdi, Pune 44

## ACADEMIC YEAR 2023-2024

## COURSE OUTCOMES

Course: M. Pharm

SEMESTER: I

Subject Code	Subject Name	CO
	Dej	partment: Pharmacology
MPAT 101T	Modern	MPAT101T CO1: Integrate and apply theoretical
	Pharmaceutical	knowledge of various spectroscopic techniques available
	Analytical	for analysis of drugs.
	Techniques	MPAT101T CO2: Understand and apply basic Knowledge of chromatographic and electrophoretic separation
		techniques.
		MPAT101T CO3: Summarize the instrumentation of
		modern analytical techniques.
		MPAT101T CO4: Elucidate a structure based on UV, IR,
		NMR and Mass data.
		MPAT101T CO5: Explain, Compare and appraise X-ray
		crystallographic and Thermal Techniques.
MPL 102T	Advanced	MPL102T CO1: Discuss the pathophysiology and
	pharmacology I	pharmacotherapy of certain diseases.
		MPL102T CO2: Explain the mechanism of drug actions at
		cellular and molecular level.
		MPL102T CO3: Understand the adverse effects,
		contraindications and clinical uses of drugs used in
		treatment of diseases.
		MPL102T CO4: Summarize, classify and apply
		epidemiology, etiology, pathophysiology sign and
		symptoms, diagnosis, complications, treatment and
	Augen L	management of disease affecting CVS, CNS.
		MPL102T CO5: Understand and summarize different
		autocoids and their role in physiological functioning.
MPL 103T	Pharmacological	Upon completion of this course, students will be able to-
	and Toxicological	MPL103T CO1: Illustrate the regulations and ethical
	Screening	requirements for use of experimental animals and be able
	Methods-I	to explain various good laboratory practices for their
		maintenance and handling.
		MPL103T CO2: Describe the various laboratory animals
		used in experimental pharmacology and their applications.
	and the same of th	MPL103T CO3: Classify the various preclinical screening
	College or	methods involved in expensionental pharmacology.
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		MPL103T CO4: Describe the various preclinical screening
		methods involved in experimental pharmacology.
		MPL103T CO5: Summarize the general principles and
		methods of immunoassay.
		MPL103T CO6: Outline the limitations of animal
		experimentations justifying the use of alternative methods
		for animal studies.
MPL104T	Cellular and	MPL104T CO1: Understand the basics of cell biology,
	Molecular	recombinant DNA technology and transfer of genes to
	Pharmacology	mammalian cells.
		MPL104T CO2: Apprehend the genetic elements of DNA,
		fingerprint analysis and various molecular techniques
		applicable in drug discovery.
		MPL104T CO3: Apply the knowledge of molecular
		MPL1041 CO3: Apply the knowledge of molecular
		pharmacology and biomarkers in drug discovery process.
		MPL104T CO4: Demonstrate molecular biology
		techniques as applicable for drug discovery.
		MPL104T CO5: Explain the molecular pathways affected
		by drugs.
MPL 105P	Pharmacological	MPL105P CO1: To demonstrate the biological evaluation
2 1001	Practical I	of drugs by in vitro and in vivo methods.
	Tractical I	MPL105P CO2: To analyze knowledge how to deal with
	4.75.1.3	experimental animals to test the potency of drugs.
		MPL105P CO3: To develop skills in handling animals and
		perform and evaluate experiments on them.
		MPL105P CO4: To calculate and compare experimental
		observations and results statistically. Student can
		differentiate between actual experimental results or if these
		results are due to chance.
		MPL105P CO5: Manage her / his time effectively by
		performing pharmacological activity on experimental
		animals.
		ent: Pharmaceutical Chemistry
MPAT 101T	Modern	MPAT101T CO1: Integrate and apply theoretical
	Pharmaceutical	knowledge of various spectroscopic techniques available
	Analytical	for analysis of drugs.
	Techniques	MPAT101T CO2: Understand and apply basic Knowledge
	1 1 25	of chromatographic and electrophoretic separation
		techniques.
		MPAT101T CO3: Summarize the instrumentation of
		modern analytical techniques.
		MPAT101T CO4: Elucidate a structure based on UV, IR,
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		NMR and Mass data
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		<u>-</u>

	T	MPAT101T CO5: Explain, Compare and appraise X-ray
		crystallographic and Thermal Techniques.
MPC102T	Advanced Organic	MPC102T CO1: Recall, apply and interpret basic aspects
	Chemistry I	of organic intermediates and different types of reactions.
		MPC102T CO2: Understand and appraise mechanism and
		applications of various named reactions.
	4-	MPC102T CO3: Recognize and apply different synthetic
		reagent and protecting groups used in organic synthesis.
		MPC102T CO4: Explain and understand name reactions
		including heterocyclic compounds.
		MPC102T CO5: Understand and apply the concept of
		disconnection to develop synthetic routs for small target
		molecule.
		MPC102T CO6: Recall and apply retrosynthetic pathways
		in organic synthesis.
MPC 103T	Advanced	MPC103T CO1: Plan strategies and prepare new chemical
WI C 1031	Medicinal	entities as potential drugs by following the process of drug
	Chemistry	discovery knowledge.
	Chemistry	MPC103T CO2: Synthesize new generation of molecules
	A STATE OF THE STA	of various classes, and understand role of stereochemistry
	The second second second	and drug action and apply the knowledge of chirality for
	A STATE OF THE STA	better drug action.
	73.77.6.	MPC103T CO3: Understand the concept of & design of
		peptidomimetics including the chemistry of
		prostaglandins, leukotrienes and thromboxanes.
		MPC103T CO4: Rationally design the enzyme inhibitors
		in medicine.
		MPC103T CO5: Understand the concept of design
		resistance.
MPC 104T	Chemistry of	MPC104T CO1: Understand, discuss and determine the
	Natural Products	natural products as leads for new pharmaceuticals
		MPC104T CO2: Explain, discuss, and demonstrate the introduction, classification, isolation, purification,
		initiodadvious,
		molecular modification and biological activity of natural
		occurring organic substances
		MPC104T CO3: Understand, discuss the recombinant
		DNA technology and crude drugs used in Diabetic, liver
		dysfunction and antitumor therapy
		MPC104T CO4: Understand, discuss and determine the
		structural characterization of natural compounds
	College	

	Chemistry Practical -I	importance and characterize the synthesized compounds using physicochemical and spectroscopic methods.  MPC105P CO2: Purify organic solvents using column chromatographic techniques.  MPC105P CO3: Isolate and characterize the islolated compounds using physicochemical and spectroscopic techniques.  MPC105P CO4: Operate, demonstrate, apply and record UV Vis spectrophotometer, column chromatography, HPLC, GC, fluorimetry and flame photometry.  MPC105P CO5: Perform degradation reactions on selected plant constituents.  MPC105P CO6: Isolate, characterize melting point, mixed melting point, molecular weight determination, functional group analysis, co-chromatographic technique for identification of isolated compounds and interpretation of UV and IR data.
	Department	Pharmaceutical Quality Assurance
MPAT 101T	Modern Pharmaceutical Analytical Techniques	MPAT101T CO1: Integrate and apply theoretical knowledge of various spectroscopic techniques available for analysis of drugs.  MPAT101T CO2: Understand and apply basic Knowledge of chromatographic and electrophoretic separation techniques.  MPAT101T CO3: Summarize the instrumentation of modern analytical techniques  MPAT101T CO4: Elucidate a structure based on UV, IR, NMR and Mass data.  MPAT101T CO5: Explain, Compare and appraise X-ray crystallographic and Thermal Techniques.
MQA 102T	Quality Management Systems (2019 pattern)	The learner will be able to MQA 102T -CO1: Explain the basic concepts, terminology and factors affecting the framework of quality, quality control and quality management systems and implement the same.  MQA 102T -CO2: Explain the scope of quality certifications applicable to Pharmaceutical industries.  MQA 102T -CO3: Demonstrate the effective utilization of different QMS tools for monitoring and improving product quality and process performance within the regulatory framework.

MPC105P CO1: Synthesize compounds of medicinal

MPC 105P

Pharmaceutical

		MQA 102T -CO4: Suggest suitable guidelines for quality
		management which can be applied to pharmaceutical
		industry.
MQA103T	Quality Control and Quality	MQA103T CO1: Explain significance of quality in pharmaceutical manufacturing and understand the
	Assurance	responsibilities of QA & QC departments.
	Assurance	MQA103T CO2: Explain role of national and international
		regulatory agencies in deciding quality standards.
		MQA103T CO3: Follow cGMP while working in
		Pharmaceutical industry.
		MQA103T CO4: Describe various aspects of
		documentation.
		MQA103T CO5: Perform analysis of raw materials, IPQC
		and FPQC of drug products and other manufacturing
		operations and controls.
		MQA103T CO6: Explain different guidelines related to
		Pharmaceutical industry.
MQA104T	Product	Upon completion of this course, the student should be able
	Development and	to:
	Technology	MQA104T CO1: describe the regulatory principles and
	Transfer	requirements of drug discovery and product
		Development.
		MQA104T CO2: relate and execute the concept of pre-
		formulation studies for various formulation.
		MQA104T CO3: explain the detailed knowledge about
		pilot plant scale-up.  MOA 104T CO4: categorize and use various
		MQATOTI COT. categorize
		pharmaceutical packaging systems.
		MQA104T CO5: implement the concept of technology
		transfer from R&D to production plant.
MQA105P	Pharmaceutical	MQA105P CO1: Perform qualitative and quantitative analysis of pharmacopoeial compounds in bulk and in their
	Quality Assurance	formulations
	Practical I	MQA105P CO2: Perform experiments on various
	4 1 86° A	analytical instruments such as UV Vis spectrophotometer,
		HPLC, GC etc.
		MQA105P CO3: Demonstrate use of tools for quality
		management.
		MQA105P CO4: Perform quality control tests for drugs,
		raw materials, dosage forms and primary and secondary
		packaging materials.
	College	MQA105P CO5: Perior experiments on pre-formulation
	The second of th	studies.
	(1) (1) (1) (1) (1) (1) (1) (1) (1) (1)	P. E. S. Modern (Street of Pharman

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		epartment: Pharmaceutics
MPAT101T	Modern	MPAT101T CO1: Integrate and apply theoretical
	Pharmaceutical	knowledge of various spectroscopic techniques available
	Analytical	for analysis of drugs.
	Techniques	MPAT101T CO2: Understand and apply basic Knowledge
		of chromatographic and electrophoretic separation
	0	techniques.
	100	MPAT101T CO3: Summarize the instrumentation of
		modern analytical techniques.
		MPAT101T CO4: Elucidate a structure based on UV, IR,
		NMR and Mass data.
		MPAT101T CO5: Explain, Compare and appraise X-ray
		crystallographic and Thermal Techniques.
MPH102T	Drug Delivery	MPH102T CO1: To formulate and evaluate novel drug
WII 111021	System	delivery systems.
	bystem	MPH102T CO2: To justify selection criteria of drug and
		polymers for the development of novel drug delivery
	57	system.
		MPH102T CO3: To understand and construct rate
		controlled drug delivery systems.
		MPH102T CO4: To construct, justify and evaluate
		gastroretentive drug delivery systems.
		MPH102T CO5: To construct, justify and evaluate ocular
		drug delivery systems and transdermal drug delivery
		systems.
		MPH102T CO6: To construct, justify and evaluate protein
) (D111000	Sylver Say	and peptide delivery systems and vaccine delivery systems.
MPH103T	Modern	On successful completion of following theory topics, a
	Pharmaceutics	learner should be able to;
		MPH103T CO1: Understand and analyze the concept of
		preformulation studies in oral solid dosage form
		development.
		MPH103T CO2: Compile and integrate preformulation
		data and apply this knowledge in development of disperse
	the way the same	systems and parenterals.
		MPH103T CO3: Analyze and apply optimization and scale
		up techniques in formulation development and
		manufacturing.
		MPH103T CO4: Demonstrate and understand the concept
		of validation of pharmaceutical processes, equipment's and
		methods.
	10,32-0	MPH103T CO5: Outline and analyze the industrial
		management techniques and GMP considerations.
	College	

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		MPH 103T CO6: Evaluate and understand stability, compression, diffusion and dissolution processes in drug product development.
MPH104T	Regulatory Affair	MPH104T CO1: Discuss the basic regulatory Documentation in pharmaceutical industry. MPH104T CO2: Discuss the preparation and submission of CTD,e-CTD. MPH104T CO3: Explain the chemistry, manufacturing controls and their regulatory importance in Anda,Nda Submission & Approval process by different regulatory agencies. MPH104T CO4: Describe the process of Preparation of Dossiers and their submission to regulatory agencies in different countries. MPH104T CO5: What are Clinical trials requirements &
MPH105P	Pharmaceutics Practical I	different approval procedures for conducting clinical trials.  On successful completion of following theory topics, a learner should be able to;  MPH105P CO1: Understand and apply the analytical methods for pharmaceutical compounds.  MPH105P CO2: Compile and integrate various analytical techniques for estimation of drug alone or in combination.  MPH105P CO3: Analyze and evaluate formulation aspects of different drug delivery systems.  MPH105P CO4: Demonstrate and understand in vitro dissolution studies and effect of various parameters on dissolution of drug.  MPH105P CO5: Outline and perform preformulation studies as a significant step in drug product development.  MPH105P CO6: Demonstrate and apply the concept of preformulation studies in product life cycle.

Course: M. Pharm

SEMESTER: II

Subject Code	Subject Name	CO
	De	partment: Pharmacology
MPL 201T	Advanced Pharmacology II (SEM 2)	Upon completion of this course, students will be able to-MPL201T CO1: Recall the pathophysiological aspects of different endocrine disorders, be able to classify different drugs used in endocrine disorders and be able to describe the pharmacological aspects associated with them.  Principal  P.S. Modern College of Pharmacy

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		MPL201T CO2: Identify the etiological aspects of various infective disorders, be able to classify different drugs used in infective disorders and be able to summarize their pharmacology in relevance to the management of different microbial infections.  MPL201T CO3: Recall the pathophysiological aspects of different GIT disorders, be able to classify different drugs used in GIT disorders and be able to describe their pharmacological aspects.  MPL201T CO4: Illustrate the current scenario of tuberculosis, HIV, cancer and respiratory disorders in society, be able to classify the drugs used in management of these diseases and be able justify the use of these drugs for management of these diseases.  MPL201T CO5: Outline the different aspects of free radical pharmacology and be able to summarize the role of various antioxidants in management of free radical induced diseases.
MPL 202T	Pharmacological and toxicological screening methods II (SEM 2)	MPL202T CO1: Explain the various types of toxicity studies.  MPL202T CO2.Appreciate the importance of ethical and regulatory requirements for toxicity studies.  MPL202T CO3: Demonstrate the practical skills require conducting the preclinical toxicity studies.  MPL202T CO4: To study Importance and applications of toxicokinetic studies.
MPL203T	Principles of Drug Discovery	MPL202T CO5: To study and apply good laboratory practices and its importance in drug development.  MPL203T CO1: Outline the various stages of drug discovery.  MPL203T CO2: Justify the importance of the role of
		genomics, proteomics and bioinformatics in drug discovery.  MPL203T CO3: Apply the knowledge of various targets, biomarkers and in vitro screening techniques for drug discovery.  MPL203T CO4: Summarize various lead seeking and lead optimization methods.  MPL203T CO5: Integrate the concepts of computer aided drug design in drug discovery.
MPL 204T	Clinical research and pharmacovigilance	MPL204T CO1: Explain the regulatory requirements for conducting clinical trial.  P. K. Modern College of Pharmacy

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MPL 205P	Pharmacological practical II (SEM 2)	MPL204T CO2: Demonstrate the types of clinical trial designs.  MPL204T CO3: Explain the responsibilities of key players involved in clinical trials.  MPL204T CO4: Execute safety monitoring, reporting and close-out activities.  MPL204T CO5: Explain the principles of Pharmacovigilance.  MPL204 CO6: Detect new adverse drug reaction and their assessment. Perform the adverse drug reaction reporting systems and communication in Pharmacovigilance.  MPL205P CO1: To demonstrate the biological evaluation of drugs by in vitro and in vivo methods  MPL205P CO2: To analyze knowledge how to deal with experimental animals to test the potency of drugs.  MPL205P CO3: Compare, select, locate and isolate different organs/tissues from the laboratory animals used in pharmacological experiments as well as by simulated experiments which give an idea on the recent methods of bioassay.  MPL205P CO4: To develop skills in handling animals and perform and evaluate experiments on them.  MPL205P CO5: To calculate and compare experimental observations and results statistically. Student can differentiate between actual experimental results or if these results are due to chance.  MPL205P CO6: Manage her / his time effectively by performing pharmacological activity on experimental
		animals.
	Departme	ent: Pharmaceutical Chemistry
MPC201T	Advanced Spectral Analysis	MPC201T CO1: To explain, discuss and integrate theory and principle of various analytical techniques like UV and
		IR spectroscopy, NMR spectroscopy, Mass Spectroscopy, Chromatography, Thermal methods of analysis MPC201T CO2: To apply the knowledge of UV and IR spectroscopy, NMR spectroscopy, Mass Spectroscopy, Chromatography, Thermal methods of analysis for identification, characterization and quantification of drugs. MPC201T CO3: To understand, explain and interpret UV, IR, NMR, and Mass spectra-of various organic compounds

		like GC MS, GCAAS, LC MS, LC FTIR, LC NMR, CE
		MS, and LC MS MS
MPC 202T	Advanced Organic Chemistry - II	MPC202T CO1: Apply the principles and techniques of Green chemistry in synthesis of organic compounds.  MPC202T CO2: Synthesize chiral compounds using various methods of asymmetric synthesis based on the stereochemistry.  MPC202T CO3: Apply the techniques of peptide synthesis.  MPC202T CO4: Apply the principles of photochemical and pericyclic reactions.  MPC202T CO5: Use different methods of catalysis and
		catalysts in synthesis of organic compounds.
MPC 203T	Computer Aided Drug Design	MPC 203T CO1 To utilize various molecular modeling softwares in the design of novel drug-like molecules.  MPC 203T CO2 To apply the various softwares for physicochemical property prediction.  MPC 203T CO3 To understand how current drugs were developed by using pharmacophores modeling and docking technique.  MPC 203T CO4 To study Case studies in Molecular Modeling to apply those in development of novel entities by the use of computer aided drug design.  MPC 203T CO5 Discuss History, different techniques and applications of Computer aided drug design & apply them to determine route for designing new molecules.  MPC 203T CO6 To know history, physicochemical parameters, applications and statistical methods used to develop QSAR & apply these concepts in developing best QSAR model.
MPC 204T	Pharmaceutical Process Chemistry	MPC204T CO1: Assess Synthetic strategy Stages of process in Bench, pilot and large scale.  MPC204T CO2: Outline Industrial Safety MSDS (Material Safety Data Sheet and Personal Protection Equipment (PPE) Fire hazards, Occupational Health & Safety Assessment Series 1800 (OHSAS-1800) and ISO-14001.  MPC204T CO3: Classify Extraction Filtration Distillation Evaporation Crystallization operations in Pharmaceutical process chemistry.  MPC204T CO4: Summarize Nitration, Halogenation and Oxidation and Reduction operations in Pharmaceutical process chemistry and Case study on industrial reduction
- marie		process.

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		MPC204T CO5: Select Fermentation used in production of
		medicine and Reaction progress kinetic analysis useful for
		scale-up.
MPC 205P	Pharmaceutical	MPC205P CO1: Compare synthesis of API's /
WII C 2031	Chemistry	intermediates by different synthetic routes and use various
	Practical -II	approaches for synthesis of organic compounds.
		1
		MPC205P CO2: Synthesize organic compounds using
		microwave technique.
		MPC205P CO3: Design drug molecules using computer
	2	nided drug design.
		MPC205P CO4: Interpret and characterize synthesized
		organic compounds using FT-IR, NMR, CNMR and Mass
		spectra.
		MPC205P CO5: Describe, discuss and compare study of
		synthesis of APIs/intermediates by different synthetic
		routes and assignments on regulatory requirements in API.
		MPC205P CO6: Describe, discuss, explain and
		demonstrate calculation of ADMET properties
		of drug molecules and its analysis using softwares
		Pharmacophore modeling, 2D-QSAR based experiments,
		3D-QSAR based experiments, docking study based
		experiment, virtual screening based experiment.
	Department:	Pharmaceutical Quality Assurance
MOAZOIT	Harand and Cafata	MOA201T COL. Understand the basic concert behind
MQA201T	Hazard and Safety Management	MQA201T CO1: Understand the basic concept behind
	Wanagement	studies of multidisciplinary nature of environment and
		empower ideas to clear mechanism and management in
	Mar 2811	different kinds of hazard management system like
		ecosystem, air, chemical & fire based hazards, fire
	-10 W.p.	protection system.
	Y 3.	MQA201T CO2: Impart basic knowledge about the
		environment and its allied problems related to air,
		chemical, fire-based hazards, and fire protection systems.
		MQA201T CO3: Illustrate the different sources and types
		of environmental studies, air, chemical & fire-based
		hazards.
		MQA201T CO4: Describe and Illustrate safety guidelines,
		rules, regulation to prevent environmental hazards like air,
		chemical & fire based hazards and its risk management
1/01/2007	71	system.
MQA 202T	Pharmaceutical	The learner will be able to
	Validation	MQA 202T-CO1: Explain the concepts, importance, scope
College		types, In ethodology and application of calibration,
18/ 100		qualification and validation activities in pharma industry.
2/		than the variety of a curvines in pharma industry.

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validation of instruments, facilities and processes as per guidelines.  MQA 202T-CO3: Explain the importance of patent and intellectual property rights  MQA 202T-CO4: Suggest methodology for qualification of laboratory, analytical and manufacturing equipments.  MQA 202T-CO5: Suggest methodology for validation of utilities, analytical methods, cleaning methods, computerized systems and manufacturing processes of various dosage forms.  MQA203T  Audits and  Regulatory Compliance  MQA203T Explain the different methods of auditing, MQA203T Explain the different methods of auditing MQA203T Understand the process of carrying out audits in departments like production, quality assurance, and different laboratories of the pharmaceutical industry.  MQA203T Design and carry out the audit process and prepare an audit report.  MQA203T Prepare audit checklist for audits.  MQA204T CO1: Understand and discuss common practices in manufacturing of sterile and non-sterile dosage forms in pharmaceutical industry.  MQA204T CO2: Understand, analyze and describe principles and processes in production planning, manufacturing of sterile and non-sterile dosage forms, in selecting containers and closures and in quality monitoring tools.  MQA204T CO3: Elaborate legal requirements for pharmaceutical industry development, plant layout and compile steps of production planning.  MQA204T CO3: Elaborate legal requirements for pharmaceutical industry development, plant layout and compile steps of production planning.  MQA204T CO3: Construct sterile product manufacturing and non-sterile product manufacturing technology and differentiate them.  MQA204T CO5: Compile, classify and compare different containers and closures for pharmaceuticals and to correlate stability aspects of packaging materials.  MQA204T CO6: Understand and elaborate principles of			LANDA 2027 CO2. Deserve sectionals for qualification and
guidelines.  MQA 202T-CO3: Explain the importance of patent and intellectual property rights  MQA 202T-CO4: Suggest methodology for qualification of laboratory, analytical and manufacturing equipments, MQA 202T-CO5: Suggest methodology for validation of utilities, analytical methods, cleaning methods, computerized systems and manufacturing processes of various dosage forms.  MQA203T  Audits and Regulatory Compliance MQA203T Explain the different methods of auditing MQA203T Understand the process of carrying out audits in departments like production, quality assurance, and different laboratories of the pharmaceutical industry.  MQA203T Design and carry out the audit process and prepare an audit report.  MQA203T Prepare audit checklist for audits.  MQA203T Prepare audit checklist for audits.  MQA203T Prepare audit of the pharmaceutical industry.  MQA203T Prepare audit of the pharmaceutical industry.  MQA203T Prepare audit of the pharmaceutical industry.  MQA204T CO1: Understand and discuss common practices in manufacturing of sterile and non-sterile dosage forms in pharmaceutical industry.  MQA204T CO2: Understand, analyze and describe principles and processes in production planning, manufacturing of sterile and non-sterile dosage forms, in selecting containers and closures and in quality monitoring tools.  MQA204T CO3: Elaborate legal requirements for pharmaceutical industry development, plant layout and compile steps of production planning.  MQA204T CO4: Construct sterile product manufacturing and non-sterile product manufacturing technology and differentiate them.  MQA204T CO5: Compile, classify and compare different containers and closures for pharmaceuticals and to correlate stability aspects of packaging materials.  MQA204T CO6: Understand and elaborate principles of Quality by Design (QbD) and Process Analytical Technology (PAT) in pharmaceutical manufacturing.  MQA205P CO1: Perform qualitative and quantitative analysis of certain pharmaceutical contaminants and drugs.  MQA205P CO2: Analyze and validate ph			MQA 202T-CO2: Prepare protocols for qualification and
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		MQA205P CO3: Study qualification of pharmaceutical
		equipment.
		MQA205P CO4: Create checklist for pharmaceutical
		facilities.
		MQA205P CO5: Design plant layout for sterile and
		nonsterile products.
		MQA205P CO6: Perform case study on certain quality
		management tools.
		management tools.
	De	partment: Pharmaceutics
MPH201T	Molecular	MPH201T CO1: To integrate the knowledge of various
	Pharmaceutics	approaches for development of TDDS.
	(Nano Tech and	MPH201T CO2: To discuss and elaborate the criteria for
* 3	Targeted DDS)	selection of drugs and polymers for the development of
		TTDS.
		MPH201T CO3: To demonstrate basics involved in the
		formulation and evaluation of TDDS.
		MPH201T CO4:To integrate and apply various concepts
		and strategies for improving targeting and absorption in the
MINION	A 1 1	design of TDDS.
MPH202T	Advanced	On successful completion of following theory topics, a
	Biopharmaceutics &	learner should be able to;
	Pharmacokinetics	MPH202T CO1: Understand and analyze the concept of
		biopharmaceutics and pharmacokinetic.
		MPH202T CO2: Compile and integrate pharmacokinetic
		data and derive the pharmacokinetic models and
	- No. 200	parameters that best describe the process of drug
		absorption, distribution and metabolism.
		MPH202T CO3: Analyze and apply critical evaluation of
		biopharmaceutic studies involving drug product
		equivalency.
		MPH202T CO4: Demonstrate and understand the design
		and evaluation of dosage regimens of the drugs using
		pharmacokinetic and biopharmaceutic parameters.
		MPH202T CO5: Outline and analyze the potential clinical
		pharmacokinetic problems and application of basics of
		pharmacokinetic.
		MPH202T CO6: Demonstrate and apply the applications
		of biopharmaceutics and pharmacokinetic in drug delivery
		development.
MPH203T	Computer Aided	MPH203T CO1 Discuss the role of Computer Aided Drug
	Drug	Delivery in drug discovery.
	Development	MPH203T CO2 Explain different Computer Aided Drug
	1/1/	Delivery techniques and their applications.
1/2	Wincipa	

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		MPH203T CO3 Demonstrate various strategies to design
		and develop new drug-like molecules.
		MPH203T CO4 Explain in detail the working with
		molecular modeling software to design new drug
		molecules.
	49.11	MPH203T CO5 Discuss various in silico virtual screening
		protocols.
MPH204T	Cosmetic &	Upon completion of the course, the students shall be able
WII 112041	Cosmeceuticals	to understand:
	Cosmeccuticals	
		MPH204T CO1: Using knowledge of regulatory and
		biological aspects is a fundamental need for developing
		cosmetics and cosmeccuticals.
		MPH204T CO2: Discuss the various building blocks used
		in cosmetics and cosmeceuticals formulations.
		MPH204T CO3: Develop and evaluate various cosmetics
		and cosmeceutical formulations with the desired safety,
		stability, and efficacy.
	1 (3)(4)	MPH204T CO4: Discuss the current technologies in the
		market.
		MPH204T CO5: Explain guidelines and challenges in
		formulating herbal cosmetics
MPH205P	Pharmaceutics	On successful completion of following theory topics, a
	Practical II	learner should be able to;
		MPH205P CO1: Understand and evaluate the effect of
		various factors on development of formulation of drug.
		MPH205P CO2: Compile and apply different types of in
	( )	
		vitro dissolution studies of drug formulation and its
		comparison with marketed product.
		MPH205P CO3: Analyze and understand the techniques in
		ex vivo and in situ evaluation of the formulation for its
		performance.
		MPH205P CO4: Demonstrate and understand various
		software's in optimization of the formulation
		MPH205P CO5: Outline and learn various in silico
		methods used in formulation development and
		optimization.
		MPH205P CO6: Demonstrate and apply the formulation
		knowledge and skills in development of herbal
		formulations.
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## **ACADEMIC YEAR 2023-2024**

## **COURSE OUTCOME**

Course: Pharm D

	First Year		
Subject Code	Subject Name	СО	
1.1T	Human Anatomy and Physiology	1.1T CO1: To recall the terminologies in the human anatomy and physiology, along with learn the functions of human cell 1.1T CO2: To summarize the functions of tissue, bones and joints in the skeleton 1.1T CO3: To explain the functions of formed elements in the blood along with lymph and its role in immunity 1.1T CO4: To compare the anatomical features of heart, lungs and GIT and to analyze their physiology 1.1T CO5: To assess the structure and function of brain, spinal cord and cranial nerves and to interpret the physiology of urinary system 1.1T CO6: To elaborate the physiology of endocrine glands, reproductive organs, sensory organs and to discuss the physiology skeletal muscles	
1.1P	Human Anatomy and Physiology	1.2P CO1: Illustrate different types of Tissues and explain various Anatomical models 1.2P CO2: Identify the bones of skeletal system 1.2P CO3: Determine Blood cell count, Hemoglobin, Blood grouping, ESR, Bleeding time and Clotting time 1.2P CO4: Record Blood Pressure, Pulse rate, Body temperature 1.2P CO5: Choose family planning devices and conduct Pregnancy diagnosis test 1.2P CO6: construct and record simple curves using frog gastroenemius sciatic nerve	
1.2T	Pharmaceutics	1.2T CO1: Integrate the knowledge related to introduction and classification of dosage form, prescription and posology 1.2T CO2: Build knowledge about the historical background and development of Pharmacy profession including different pharmacopoeias 1.2T CO3: Develop and demonstrate solid dosage form like powders and granules and pharmaceutical calculations related to weights and measures	



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	The state of the s	
		1.2T CO4: Build knowledge of liquid dosage form including both monophasic and biphasic dosage form 1.2T CO5: Investigate information related to suppositories and surgical aids 1.2T CO6: Develop knowledge related to different galenical products and its extraction processes along with incompatibilities associated with dosage form
1.2P	Pharmaceutics	1.2P CO1: Demonstrate the skill of preparation and evaluation of various solid and liquid dosage forms 1.2P CO2: Explain principles of formulation and evaluation of dosage forms 1.2P CO3: Calculate evaluation parameters like density, specific gravity, angle of repose, carr's index and hausner's ratio of pharmaceutical preparation 1.2P CO4: Classify various dosage forms by using different criteria 1.2P CO5: Create labels in prescribed manner for various dosage forms 1.2P CO6: Build knowledge regarding different types of incompatibilities for safety, efficacy and therapeutic effect of dosage forms
1.3T	Medicinal Biochemistry	1.3T CO1: Define the basic concepts in medicinal biochemistry and clinical chemistry 1.3T CO2: Apply concepts and knowledge of medicinal biochemistry to clinical scenarios 1.3T CO3: Critically interpret how the biomolecules acts on the body and its mechanisms 1.3T CO4: Link the biochemical reactions and pathways of several diseases 1.3T CO5: Explain the common laboratory values in clinical chemistry 1.3T CO6: Use the scientific laboratory equipment in order to gather and analyze data on medicinal biochemistry
1.3P	Medicinal Biochemistry	1.3P CO1: Evaluate and analyze presence of various biomolecules/ normal and abnormal constituents in body fluids using qualitative and quantitative tests 1.3P CO2: Understand and analyze the catalytic activity of enzymes and the importance of isoenzymes in the diagnosis of diseases 1.3P CO3: Apply the biochemical principles of organ function tests of the kidney, liver, and endocrine gland 1.3P CO4: Understand the importance of levels of various biomolecules in body fluids
1.4T	Pharmaceutical Organic Chemistry	1.4T CO1: Explain the physical properties of organic compounds 1.4T CO2: Identify the structures of a given organic compound and give the nomenclature



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		Transport of the state of the s
		1.4T CO3: Draw the mechanisms involved in various named organic reactions 1.4T CO4: Analyse the reactivity, orientation and stability of organic reactions
	101 (S)	1.4T CO5: Predict the products obtained through simple organic reactions
		1.4T CO6: To study on some important official organic
		compounds
1.4P	Pharmaceutical Organic Chemistry	1.4P CO1: Compose different classes of organic pharmaceuticals using some named organic reactions with mechanisms
		1.4P CO2: Apply stereo models and explain the structural aspects of organic compounds
		1.4P CO3: Integrate and Analyze the elements in different organic pharmaceuticals by Performing organic qualitative analysis
		1.4P CO4: Identify various classes of organic compounds by systematic qualitative analysis
1.600		1.4P CO5: Analyse various organic pharmaceuticals 1.5T CO1: To discuss and explain the principles and
1.5T	Pharmaceutical Inorganic Chemistry	procedures of analysis of drugs
	morganic Chemistry	1.5T CO2: To apply and determine the applications of
		inorganic pharmaceuticals in analysis of drug
		1.5T CO3: To discuss the fundamentals of analytical chemistry and examine inorganic pharmaceuticals
		regarding their monograph
	and the second	1.5T CO4: To justify the importance of inorganic
	The state of the s	pharmaceuticals in preventing and curing disease 1.5T CO5: Knowledge about the sources of impurities
		and methods to determine the impurities in inorganic
		drugs and pharmaceuticals 1.5T CO6: To have been introduced to a variety of
		inorganic drug classes
1.5P	Pharmaceutical Inorganic Chemistry	1.5P CO1: Perform the limit test for certain impurities like chloride, sulphate, iron, arsenic, lead and heavy
	mergame enemaly	metals as per the Indian Pharmacopoeia
		1.5P CO2: Determine percentage purity of given pharmaceutical drugs by titrimetric analysis
		1.5P CO3: Perform qualitative analysis of given
		inorganic mixtures
		1.5P CO4: Identify the Inorganic compounds through various chemical tests
		1.5P CO5: Know the sources of impurities and methods
		to determine the impuritles in inorganic drugs and pharmaceuticals
		1.5P CO6: Use different methods for preparation of
	The second secon	Inorganic substances



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Second Year		
Subject Name	CO	
Pathophysiology	2.1T CO1: Describe the etiology and pathogenesis of selected diseases 2.1T CO2: Knowledge the signs and symptoms of disease 2.1T CO3: Identify complications of diseases 2.1T CO4: Know most commonly encountered pathophysiological state and/or disease mechanism as well as any clinical testing requirements 2.1T CO5: Students will make correlation between pathophysiology and clinical skills they are learning in their allied health science programs 2.1T CO6: Students will understand how the various organ systems are interrelated, and use this understanding to promote a holistic approach towards the application and treatment of nations.	
Pharmaceutical Microbiology	the evaluation and treatment of patients  2.2T CO1: To list the branches, scope of microbiology and morphology of microbes  2.2T CO2: To explain the methods of identification, cultivation and preservation of various microorganisms  2.2T CO3: To apply the principles of sterilization in pharmaceutical processing and sterility testing  2.2T CO4: To compare different types of immunological reactions, antigens, vaccines and their role in immunity	
	2.2T CO5: To evaluate microbiological standards of pharmaceuticals and presence of pathogens 2.2T CO6: To elaborate the characteristics, mode of infection, diagnosis, prophylaxis and treatment of bacterial, fungal and viral infectious agents	
Pharmaceutical Microbiology	2.2P CO1: To recal different techniques of sterilization and equipment used in microbiology laboratory 2.2P CO2: To interpret characteristics of microbes using staining techniques, Isolation methods and quantitative estimation 2.2P CO3: To construct standard graphs for estimating antibiotics and vitamins using microbes 2.2P CO4: To test for possible microbial contamination in a given sample 2.2P CO5: To estimate qualitatively and quantitatively the amount of microbes in a sample 2.2P CO6: To choose the correct method for evaluating	
	Pharmaceutical Microbiology  Pharmaceutical	



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2.27	DI	2.2T GOL D
2.3T	Pharmacognosy & Phytopharmaceuticals	2.3T CO1: Develop and design agricultural & amp; storage requirement of crude drugs & amp; explain detailed pharmacognostic account of medicinal plants 2.3T CO2: Develop and design the technique for extraction, isolation, purification of phytoconstituents for medicinal, agricultural, food & amp; healthcare
		industry 2.3T CO3: Design & De
		2.3T CO4: Develop themselves to design a method for analysis, detection, purification of various extracts, phytoconstituents & amp; herbal raw materials. 2.3T CO5: Correlate various herbal crude drugs & amp; their method of extraction, chemistry & amp; pharmacognostic details including their medicinal applications
2.3P	Pharmacognosy & Phytopharmaceuticals	2.3P CO1: Judge various crude drugs on the basis of their morphological & mp; microscopical studies
		2.3P CO2: Design various analytical parameters required for the authentication of crude drugs, extracts, oils etc.
		2.3P CO3: Illustrate the anatomical architecture of various crude drugs & amp; its significance for the plants & amp; for its analysis
		2.3P CO4: Evaluate the purity & amp;/or stability of various unorganized crude drugs & amp; oils 2.3P CO5: Correlate the various staining reagents required for the authentication of crude drugs during
- Am	-	microscopy
2.4T	Pharmacology-I	2.4T CO1: Relate pharmacokinetics and pharmacodynamics of a drug with drug action 2.4T CO2: Identify the factors modifying drug action 2.4T CO3: Assess drug interactions and detect adverse
		drug reactions 2.4T CO4: Classify and explain the pharmacology of drugs acting on various systems
		2.4T CO5: Correlate and apply the knowledge of pharmacology as therapeutically
2.4P	Pharmacology-I	2.4P CO1: Utilize and handle the Experimental animals 2.4P CO2: Assess and handle the computerized
		simulated software programme such as COLPHARM 2.4P CO3: Compare the effects of various drugs on
		animals



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		2.4P CO5: recommend physiological salt solutions for
		different isolated tissues
		2.4P CO6: Apply different routes of drug
		administration and the techniques of Euthanasia and
		analgesia in laboratory animals
2.5T	Community Pharmacy	2.5T CO1: Know the pharmaceutical care services
		2.5T CO2: Understand the business and professional
		practice management skills in community pharmacies
		2.5T CO3: Patient counselling and provide health
1		screening services to public in community pharmacy
		2.5T CO4: Respond to minor ailments and provide
		appropriate medication
		2.5T CO5: Encourage patients for rational drug therapy
		2.5T CO6: Show empathy and sympathy to patients
2.6T	Pharmacotherapeutics-	2.6T CO1: To recall the pathophysiology of
	I	cardiovascular disorders and relate their etiology with
}	2	the therapeutic approach including treatment
		controversies.
		2.6T CO2: To outline the concept of essential drugs use
		and rational drug therapy and summarize the choice of
		drugs with justification in various disease conditions
	4	2.6T CO3: To identify various types of respiratory and
	-516 2	endocrine disorders with respect to clinical features and
		laboratory investigations, list their complications along
		with replacement in their management
	1 34 16 2	2.6T CO4: To distinguish between various disease
		conditions and analyse the results with drug induced
		disorders
		2.6T CO5: To select the patient-specific parameters
l	to the second second	relevant in initiating drug therapy, and monitoring
		therapy among pediatric, geriatric, pregnant and
		lactating women.
	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	2.6T CO6: To develop competency to design individual
		care plan for cardiovascular, respiratory, ocular and
		hormonal disorders
2.6P	Pharmacotherapeutics-	
2.01	I	2.6P CO1: To recall the pathophysiology of
	1	cardiovascular disorders and relate their etiology with
		the therapeutic approach including treatment
	ay	controversies
		2.6P CO2: To outline the concept of essential drugs use
		and rational drug therapy and summarize the choice of
	Landon Company	drugs with justification in various disease conditions
		2.6P CO3: To identify various types of respiratory and
	2 2 2 1	endocrine disorders with respect to clinical features and
		laboratory investigations, list their complications along
	7.55	with replacement in their management



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	2.6P CO4: To distinguish between various disease conditions and analyze the results with drug induced disorders 2.6P CO5: To select the patient-specific parameters relevant in initiating drug therapy, and monitoring therapy among pediatric, geriatric, pregnant and lactating women 2.6P CO6: To develop competency to design individual care plan for cardiovascular, respiratory, ocular and hormonal disorders
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Third Year		
Subject Code	Subject Name	СО
3.1T	Pharmacology-II	3.1T CO1: Explicate the pharmacological actions of different categories of drugs 3.1T CO2: Understand the mechanism of drug action at organ system/ sub cellular/ macromolecular level 3.1T CO3: Illustrate the application of basic pharmacological knowledge in the prevention and treatment of various diseases 3.1T CO4: Enlighten the effect of drugs on animal by simulated experiments 3.1T CO5: Explain the correlation of pharmacology with other biomedical science 3.1T CO6: Elucidate the signal transduction mechanism of various mechanism
3.1P	Pharmacology-II	3.1P CO1: Illustrate the experimental pharmacology, instrumental pharmacology, CPCSEA: guidelines and common laboratory animals 3.1P CO2: Demonstrate of Common laboratory techniques. Blood withdrawal, serum and plasma separation, anesthetics and euthanasia used for animal studies 3.1P CO3: Explain different route of drug administration in rat & amp; mice, effect of hepatic microsomal enzyme inducers, Anticonvulsant effect of drugs by MES and PTZ method 3.1P CO4: Explain in details effect of drugs on ciliary motility of frog oesophagus, on rabbit eye, skeletal muscle relaxants using rota-rod apparatus, on locomotor activity using actophotometer, of stereotype and anti-catatonic activity of drugs on rats/mice, anxiolytic activity of drugs using rats/mice, local anesthetics by different methods



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		3.1P CO5: Judge various terms and factors that influence the pharmacokinetics and Pharmacodynamics and Build the concept of discovery and development of New drugs-Preclinical and clinical studies 3.1P CO6: They would have understood the pharmacological aspects of drugs used to treat ailment of different organ systems of the body. 3.1P CO7: They would appreciate the importance of drug discovery by preclinical and clinical trials 3.1P CO8: They would have finally learnt to apply the knowledge of drugs practically using simulated pharmacological experiments
3.2T	Pharmaceutical Analysis	3.2T CO1: Assess quality Assurance and validation methods concept with guidelines and their regulations and Understand the concepts of calibration and validation various pharmaceutical analytical instruments 3.2T CO2: Explain Introduction, theory, instrumentation and application in the different types of chromatography and spectroscopic technique 3.2T CO3: Categorize the electrometric methods, theoretical aspects, Instrumentation, Interpretation of data and analytical applications 3.2T CO4: Understand the principles of analytical techniques and its application in analysis of drugs
3.2P	Pharmaceutical Analysis	3.2P CO1: Develop mobile phase for separation and identification of drugs by Chromatography 3.2P CO2: Operate various instruments to develop practical skill 3.2P CO3: Interpret the data obtained through UV, IR, NMR, spectra and report the result 3.2P CO4: Summarize theoretical knowledge on various instrumental technique
3.3T	Pharmacotherapeutics-II	3.3T CO1: To remember and recall the pathophysiology of selected diseases and rationale for drug therapy 3.3T CO2: To identify various therapeutic approaches for the management of selected diseases 3.3T CO3: To apply the concepts of various drug therapies and identify the controversies in drug therapy 3.3T CO4: To assess the drug therapy by preparing individual therapeutic plan based on diagnosis 3.3T CO5: To evaluate the patient specific parameters relevant in initiating drug therapy and monitoring therapy



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		3.3T CO6: To create a pharmaceutical care plan, design a list of patient counselling points on the
		specific illness
3.3P	Pharmacotherapeutics-	3.3P CO1: To remember and recall the pathophysiology and management of cardiovascular,
		respiratory, endocrine diseases and viral infections 3.3P CO2: To identify various drug interactions and
		rationalize the prescription 3.3P CO3: To plan the quality use of medicines
		surrounding the therapeutic agents in the treatment of selected diseases
		3.3P CO4: To analyze the clinical skills in the
		therapeutic management of selected disease conditions 3.3P CO5: To prioritize the treatment strategies for
		better patient outcome and discuss the controversies in treatment
3.4T	Pharmaceutical Jurisprudence	3.4T CO1: Illustrate pharmaceutical legislation in India concepts, principle and significance of
		pharmaceutical ethics drafted by PCI 3.4T CO2: Compile and describe objectives, legal
		definitions, do's and don'ts, penalties in case of breach
		of act mentioned under various pharmaceutical acts and rules
		thereunder 3.4T CO3: Practice legal provisions for import,
		manufacture, sale of drugs and cosmetics and schedules thereunder and similarly, provisions under
		Pharmacy Act 3.4T CO4: Organize and describe provisions under
		medicinal and toilet preparation act and narcotic drug and psychotropic substances act
		3.4T CO5: Analyse and describe salient features of
		drugs and magic remedies act and essential commodities act relevant to drugs price control order
		3.4T CO6: Examine provisions made under drugs price control order, prevention of cruelty to animal act,
		patent and design act and prescription and non- prescription product
3.5T	Medicinal Chemistry	3.5T CO1: Understand Modern concepts of rational drug design
		3.5T CO2: Derive and understand the Structural
		influences on mechanism of pharmacologic action (structure-activity relationship)
		3.5T CO3: Evaluate the chemistry of drugs with
		respect to their pharmacological activity 3.5T CO4: Summarize the mechanism pathways of
		different class of medicinal compounds



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		t '-lalatura brand
		3.5T CO5: Illustrate chemical nomenclature, brand
	37 000	names of important marketed products and their side
		effects
		3.5T CO6: Analyse and apply Diagnostic agents
3.5P	Medicinal Chemistry	3.5P CO1: Access the fundamentals of medicinal
3.31	Wiedlemar Chemistry	chemistry and theory of analysis of drugs
	2.0	2.50 CO2: Perform synthesis of medicinal compounds
		3.5P CO2: Perform purification and evaluation of
		3.51 COS: Periorit purification and oversely
		synthesized compounds
		3.5P CO4: Analyse pharmaceutical drugs using
	the second second	appropriate assay method
		3.5P CO5: Study and Analyse monographs of
,		important drugs
		3.5P CO6: Determine physical properties of organic
		compounds with respect to OSAR analysis
3.6T	Pharmaceutical	3.6T COI: Integrate the knowledge related to different
3.01	Formulations	pharmaceutical dosage forms
	Formulations	3.6T CO2: Develop and prepare various
		pharmaceutical dosage forms
		3.6T CO3: Analyse different pharmaceutical dosage
		3.61 CO3: Analyse different pharmaceurion assignments
		forms according to their evaluation parameters
	13 1 1 1	3.6T CO4: Integrate knowledge regarding different
		terms like bioavailability and bioequivalence
		3.6T CO5: Study role of various pharmaceutical
		dosage forms
3.6P	Pharmaceutical	3.6P CO1: Develop different types of dosage form
3.01	Formulations	according to route of administration
	1 officiations	3.6P CO2: Develop and prepare various
		pharmaceutical dosage forms
	1.3	3.6P CO3: Analyse different dosage forms according
		to their evaluation parameters
		3.6P CO4: Design labels according to different types
	7,31	3.0P CO4: Design labels according to 2
	12 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	of pharmaceutical dosage forms
		3.6P CO5: Integrate knowledge regarding principle involved in formulation and role of various
		Involved in termination
		pharmaceutical dosage forms
		A CONTRACTOR OF THE PROPERTY O
		Fourth Year
Subject Code	Subject Name	CO
4.17	Pharmacotherapeutics-	4.1T CO1: To explain the etiopathogenesis of selected
4.1T		gastrointestinal, hematological, neurological and
	III	psychiatric diseases
		psychiatric discuss the principles of evidence based
		4.1T CO2: To discuss the principles of evidence based
		therapy and pain management



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		4.1T CO3: To identify the patient-specific parameters relevant in initiating and monitoring drug therapy and adverse effects 4.1T CO4: To discuss the therapeutic approach in the management of selected diseases and controversies in drug therapy 4.1T CO5: To prepare individualized therapeutic plans based on diagnosis 4.1T CO6: To recognize the role of pharmacist in essential and rational drug use
4.1P	Pharmacotherapeutics-	4.1P CO1: To identify drug interactions and rationalize the prescription 4.1P CO2: To discuss the therapeutic approach to management of selected diseases 4.1P CO3: To prepare individualized therapeutic plans based on diagnosis 4.1P CO4: To conduct patient counselling 4.1P CO5: To conduct planned experiments and
4.2T	Hospital Pharmacy	prepare laboratory report in a standard format  4.2T CO1: To apply and analyze structure,
	of bullet	organisation and functions of hospital and hospital pharmacist 4.2T CO2: To understand, Prepare and implement budget, inventory control in various drug policies 4.2T CO3: To analyze and evaluate various hospital drug policies to develop hospital pharmacy news letters hospital formulary 4.2T CO4: To define, classify and prepare various drug distribution methods for inpatients and outpatients including narcotic and controlled drugs 4.2T CO5: To understand and explain the procurement, manufacturing and storage process of various formulations and handling of radio pharmaceuticals 4.2T CO6: To develop programmes for professional upraising continuously and to build inter professional relations in the hospitals
4.2P	Hospital Pharmacy	4.2P CO1: To understand various drug distribution systems in hospital 4.2P CO2: To extend the professional practice management skills in hospital pharmacy 4.2P CO3: To utilize various methods for the preparation and labeling of pharmaceutical products such as powders and intravenous solutions 4.2P CO4: To recommend the solutions to overcome the drug interaction and adverse drug reactions 4.2P CO5: To appreciate various store management and inventory control



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		4.2P CO6: To solve drug related problems through
		prescription analysis and individualized dose
4.2T	Clinical Pharmany	4.3T CO1: Assess, Detect and monitor ADRs
4.3T	Clinical Pharmacy	4.3T CO2: Design, analyze, interpret and formulate the
		drug or medicine information
		4.3T CO3: Interview Medication history and Drug
		utilization review and Provide clinical pharmacy
		services
		4.3T CO4: Solve Drug and Poison information Queries
		and provide disease related counseling to patient
		4.3T CO5: Relate common medical abbreviations,
		terminologies and lab results used in clinical practice
		and interpret the data obtained through laboratory tests
		4.3T CO6: Inspect the potential sources of Medication
		error and act for its prevention
4.3P	Clinical Pharmacy	4.3P CO1: To create awareness in patients by
		counselling them on various diseases and drugs using
	2 4 2	clinical knowledge and communication skills
		4.3P CO2: To conduct comprehensive and meticulous
		medication history interview for the preparation of
		individualized pharmaceutical care plan
		4.3P CO3: To interpret laboratory results of specific
		disease states mentioned and correlating with patient
		drug therapy while monitoring disease progression
		4.3P CO4: To provide response to a drug and poison
		information queries using modified systemic approach
		by critically appraising the blomedical literature
		4.3P CO5: To report and assess causality of adverse
		drug reactions to establish acausal relation between an
4 4T	Biostatistics &	ADR and a drug  4.4T CO1: To Demonstrate knowledge about basic
4.4T	Research Methodology	concepts about research and the methodology to
	Research Methodology	conduct research
		4.4T CO2: To simplify the understanding of statistical
		methods in epidemiology and be conscious about its
		relative, attributable risks
		4.4T CO3: To learn to utilize the computer applications
		and their advantages in both hospital, community
	4 4 7	pharmacy
	* 5	4.4T CO4: To Develop an appropriate framework for
		research studies including study design, method of data
		collection, sampling techniques, report writing
4.5T	Biopharmaceutics &	4.5T CO1: To recall basic concepts of absorption,
	Pharmacokinetics	distribution, metabolism and excretion of drugs
		4.5T CO2: To understand the mechanisms, interpret
		various factors affecting drug absorption, distribution,
	1	metabolism and excretion of drugs



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		4.5T CO3: To apply the pharmacokinetic models for the determination of Pharmacokinetic parameters 4.5T CO4: To examine multiple dosage regimens based on pharmacokinetic Parameters for maximizing therapeutic effectiveness and patient compliance 4.5T CO5: To evaluate various pharmacokinetic parameters for the drugs exhibiting saturation kinetics 4.5T CO6: To design the bioavailability testing protocol of a drug and compare the bioequivalence between marketed products
4.5P	Biopharmaceutics & Pharmacokinetics	
		4.5P CO2: To interpret the effect of surfactant, diluents, lubricant and polymorphism on rate of drug dissolution 4.5P CO3: To solve bioavailability parameters of drugs by using plasma data and methods to improve
		bioavailability 4.5P CO4: To analyze absorption rate constant, KE, biological half-life, mean residence time and mean
	10	absorption time for the given data 4.5P CO5: To estimate the extent of protein biding by equilibrium dialysis or dynamic dialysis methods 4.5P CO6: To predict the pharmacokinetic parameters for the given data as per one compartment and two
4.6T	Clinical Toxicology	compartment models  4.6T CO1: To understand the general principles
		involved in the management of poisoning with toxicokinetics parameters 4.6T CO2: To Evaluate the role of antidotes, supportive care, gut decontamination and elimination
		enhancement in poisoning 4.6T CO3: To distinguish the clinical symptoms and to plan various managements of pesticides, drugs acting on CNS, hydrocarbons, caustics and radiation poisoning
		4.6T CO4: To categorize the toxic symptoms and management of venomous snake bites, toxicity of plants and contaminated foods and heavy metals 4.6T CO5: To compare the characteristics and specific standard treatment guideline for the treatment of
		various toxins 4.6T CO6: To design several preventive approaches to reduce unintended poisoning
4.7T	Pharmacotherapeutics I & II	4.7T CO1: Identify drug interactions, controversies and adverse effects of drug therapy



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		4.7T CO2: Discuss the various methods involved in the diagnosis of selected disease state 4.7T CO3: Identify Goals of therapy and therapeutic approach to management of diseases
		4.7T CO4: Describe the pathophysiology of selected disease states and explain the rationale for drug therapy 4.7T CO5: Describe the individualized therapeutic plan based upon the diagnosis
4.7P	Pharmacotherapeutics I & 11	4.7P CO1: Propose the therapeutic management of selected diseases 4.7P CO2: Plan individualized therapeutic plans based on diagnosis 4.7P CO3: Identify patient specific parameters relevant in initiating drug therapy and monitoring therapy 4.7P CO4: Develop evidence based medicine therapeutic plan according to patient 4.7P CO5: Modify the controversies in drug therapy
		Fifth Year
Subject Code	Subject Name	CO
5.1T	Clinical Research	5.1T CO1: Discuss the Pharmacological and Toxicological considerations in process of development of new drugs 5.1T CO2: Discuss the principles and phases in clinical trial of drug 5.1T CO3: Explain the guidelines for ethics and safe monitoring in clinical trial of a drug 5.1T CO4: Design the documents of clinical trial 5.1T CO5: Distinguish the guidelines of national and international regulatory bodies for clinical trial 5.1T CO6: Recognize differing roles and obligations of the Investigator, Sponsor and Institutional Review Board
5.2T	Pharmacoepidemiology and Pharmacoeconomics	5.2T CO1: Define Pharmacoepidemiology and Pharmacoeconomics & Define Pharmacoepidemiology and application 5.2T CO2: Calculate various outcome and drug use measures involved in pharmacoepidemiology 5.2T CO3: Demonstrate competency in the design, conduct and evaluation of Pharmacoepidemiology studies 5.2T CO4: Summarize the methods used in Pharmacoepidemiology and Pharmacoeconomic analysis 5.2T CO5: Solve various case studies by applying the concepts of pharmacoepidemiology and Pharmacoeconomics in designing a good outcome



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5.3T	Clinical Pharmacokinetics & Pharmacotherapeutic Drug Monitoring	5.3T CO1: To Discuss the pharmacokinetic principles to individualize drug therapy in patient care situations 5.3T CO2: To Determine dose, dosing intervals and dosage adjustments of a drug for a given patient 5.3T CO3: To Apply the principles of pharmacokinetics to analyse and predict drug interactions
	Drug Monitoring	5.3T CO3: To Apply the principles of
		interactions
		5.3T CO4: To Prepare protocol for TDM of drugs for selected diseases
		5.3T CO5: To Discuss the concept of genetic polymorphism in metabolism, transport and target of a
		drug



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