

### OUTCOME-BASED EDUCATION



# Institute Vision, Mission, PEO's, PO and CO

Shri Balasaheb Mane Shikshan Prasarak Mandal Ambap's

## ASHOKRAO MANE COLLEGE OF PHARMACY PETH VADGAON



### **VISION**

• Empowerment of the nation with knowledgeable pharmacist for healthy India.

### **MISSION**

<b>Mission Code</b>	Mission statements		
M1	To provide pharmaceutical education par excellence.		
M2	To promote community, institutional and industrial pharmacy.		
М3	To foster and disseminate productive research in new & emerging		
WIS	area.		
M4	To generate human resource in the profession of pharmacy.		

### PROGRAM EDUCATIONAL OBJECTIVES (PEO's)

PEO Code	Program Educational Outcomes (PEO's) Statements
PEO1	To produce Pharmacy graduate with strong fundamental concepts and high technical competence in Pharmaceutical sciences and technology who shall be able to use these tools in the field of Pharmacy.
PEO2	To train the students to contribute towards public healthcare system and counseling for prophylaxis and prevention of diseases.
PEO3	To generate the potential knowledge pool with interpersonal and collaborative skills to identify, assess and formulate problems and execute the solution in Pharmaceutical industry.
PEO4	To promote a development of trained human resources in Pharmaceutical sciences for dissemination of quality education with highly professional and ethical attitude, strong communication skills, and effective leadership skills to do work in a team with multidisciplinary approach.
PEO5	To encourage the students to participate in lifelong learning process for the highly productive career and to relate the concepts of Pharmaceutical sciences towards surveying the cause of the society.



### Program Outcomes (PO's)/ Program Specific Outcome (PSO's)

ID	Program Outcomes (PO's)/ Program Specific Outcome (PSO's) Statements
PO1	<b>Pharmacy Knowledge:</b> Possess knowledge and comprehension of the core and basic knowledge associated with the profession of pharmacy, including biomedical sciences;
	pharmaceutical sciences; behavioral, social, and administrative pharmacy sciences; and manufacturing practices.
	Planning Abilities: Demonstrate effective planning abilities including time
PO2	management, resource management, delegation skills and organizational skills.
	Develop and implement plans and organize work to meet deadlines.
	<b>Problem analysis:</b> Utilize the principles of scientific enquiry, thinking analytically,
PO3	clearly and critically, while solving problems and making decisions during daily practice. Find, analyze, evaluate and apply information systematically and shall make
	defensible decisions.
	Modern tool usage: Learn, select, and apply appropriate methods and procedures,
PO4	resources, and modern pharmacy-related computing tools with an understanding of the
	limitations.
	Leadership skills: Understand and consider the human reaction to change, motivation
	issues, leadership and team-building when planning changes required for fulfillment of
PO5	practice, professional and societal responsibilities. Assume participatory roles as
	responsible citizens or leadership roles when appropriate to facilitate improvement in
	health and wellbeing.
	Professional Identity: Understand, analyze and communicate the value of their
PO6	professional roles in society (e.g. health care professionals, promoters of health,
	educators, managers, employers, employees).
	<b>Pharmaceutical Ethics:</b> Honor personal values and apply ethical principles in professional and social contexts. Demonstrate behavior that recognizes cultural and
DO7	personal variability in values, communication and lifestyles. Use ethical frameworks;
PO7	apply ethical principles while making decisions and take responsibility for the
	outcomes associated with the decisions.
	<b>Communication:</b> Communicate effectively with the pharmacy community and with
PO8	society at large, such as, being able to comprehend and write effective reports, make
	effective presentations and documentation, and give and receive clear instructions.
	The Pharmacist and society: Apply reasoning informed by the contextual knowledge
PO9	to assess societal, health, safety and legal issues and the consequent responsibilities
	relevant to the professional pharmacy practice.
	<b>Environment and sustainability:</b> Understand the impact of the professional pharmacy
PO10	solutions in societal and environmental contexts, and demonstrate the knowledge of,
	and need for sustainable development.
	<b>Life-long learning:</b> Recognize the need for, and have the preparation and ability to
PO11	engage in independent and life-long learning in the broadest context of technological change. Self-access and use feedback effectively from others to identify learning needs
	and to satisfy these needs on an ongoing basis.
	and to satisfy these needs on an originity basis.



### PROGRAM SPECIFIC OUTCOMES (PSO's)

### **Pharmaceutics**

M. Pharmacy graduates will be able to,

ID	Program Specific Outcome (PSO's) Statements	
PSO1	Understand analytical techniques for the identification, characterization, and	
	quantification of drugs.	
PSO2	Know theoretical and practical skills of UV, IR, HPLC, and Perform structural	
	elucidation of organic compounds using spectroscopic tools.	
PSO3	Know The elements of pre-formulation studies, & acquire knowledge of novel as	
	well as conventional drug delivery systems.	
PSO4	To identify and resolve the research problems by utilizing the technical skill gain	
	through training and experimentation.	
PSO5	Know Industrial Management and GMP Considerations, Generic drug Product	
	development & utilize skills as a part of team a professional endeavor.	

### **Pharmaceutical Quality Assurance**

M. Pharmacy graduates will be able to,

ID	Program Specific Outcome (PSO's) Statements
PSO1	Understand analytical techniques for identification, characterization and
	quantification of drugs.
PSO2	Know theoretical and practical skills of UV, IR, and HPLC, and Perform Structural
	Elucidation of organic compounds using spectroscopic tools.
PSO3	To understand the applications & responsibilities of Quality assurance and Quality
	control throughout the product life cycleand appreciate the importance of
	documentation.
PSO4	To analyze the application-based importance of emerging quality-building concepts
	in product manufacturing.
PSO5	To understand and perform procedures of method validation, process validation,
	equipment/facilities/utilities qualifications & validation, documents, and records
	designing as per the regulatory standards leading to compliance of cGMP.
PSO6	To Understand the Regulatory Requirements of Pharmaceuticals.



### **COURSE OUTCOMES**

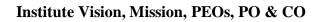
### FIRST YEAR B. PHARMACY

BP101T	Human Anatomy and Physiology I- Theory	PO
BP101T.01	Describe the gross morphology, and functions of cell and	1,3,4,6,7,8,11
	tissue.	1,5,4,0,7,0,11
BP101T.02	Describe about the gross morphology, functions of cell and	1,3,4,5,6,7,8,11
	tissue.	1,3,4,3,0,7,0,11
BP101T.03	Explain structure and functions of skeletal, muscular,	1,2,4,6,7,8,11
BF1011.03	cardiovascular system, lymphatic PNS of the human body.	
BP101T.04	Explain structure and functions skin and Special senses and	1,2,3,4,5,7,8,11
	their disorders.	1,2,3,4,3,7,0,11

BP102T	Pharmaceutical Analysis I – Theory	PO
BP102T.01	Explain the basic concepts of pharmaceutical analysis,	1,2,3,6,7,8,9,11
	impurity, their source and volumetric method used to	
	standardize various inorganic compounds.	
BP102T.02	Describe the principle of various volumetric titrations.	1,2,3,6,7,8,9,11
BP102T.03	Explain principle involved in gravimetric analysis.	1,2,3,4,6,7,8,9,11
BP102T.04	Discuss the principle and techniques of electrochemical method	1,2,3,4,6,7,8,9,11
BF 1021.04	of analysis & their applications.	

BP103T	Pharmaceutics I – Theory	PO
BP103T.01	Explain the historical development of Pharmacy profession,	1,2,3,6,7,8,9,11
	pharmacopoeial specifications, introduction to dosage forms,	
	posology, pharmaceutical calculations and prescription.	
BP103T.02	Explain powder and liquid dosage forms	1,2,3,6,7,8,9,11
BP103T.03	Explain semisolid dosage forms and suppositories.	1,2,3,6,7,8,9,11
BP103T.04	Explain the pharmaceutical incompatibilities like physical,	1,2,3,6,7,8,9,11
	chemical and therapeutic.	

BP104T	Pharmaceutical Inorganic Chemistry – Theory	PO
BP104T.01	Summarize the sources of impurities and methods to determine	1,2,4,6,7,8,9,11
	the impurities in inorganic drugs and pharmaceuticals.	, , , , , , ,
BP104T.02	Explain acids, bases, buffers, electrolytes, dental products with	1,2,4,6,7,8,9,11
	their official monographs.	1,2,4,0,7,0,7,11
BP104T.03	Explain the medicinal and pharmaceutical importance of	1,2,3,6,7,8,9,11
	inorganic compounds.	1,2,3,0,7,0,9,11
BP104T.04	Describe and relate radiopharmaceuticals and radioisotopes with	1,2,3,4,6,7,8,9,11
	their pharmaceutical application.	1,2,3,4,0,7,0,9,11





BP105T	Communication skill	PO
BP105T.01	Understand the behavioral needs for a Pharmacist to function effectively in the areas of pharmaceutical operation.	2,5,8,11
BP105T.02	Communicate effectively Verbal and Non Verbal.	2,5,8,11
BP105T.03	Relate basic listening skill and effective written communication.	2,5,8,11
BP105T.04	Develop interview skills, Leadership qualities and essentials	2,5,8,11

BP107P	Human Anatomy and Physiology I- Practical	PO
BP107P.01	Describe different types tissue and bones.	1,3,4,6,7,8,11
BP107P.02	Illustrate the Hematological experiments.	1,3,4,5,6,7,8,11
BP107P.03	Illustrate the Hematological experiments.	1,2,4,6,7,8,11

BP108P	Pharmaceutical Analysis I – Practical	PO
BP108P.01	Illustrate the chemical compounds for their purity using limit	1,2,3,4,5,6,
	tests.	7,8,9,11
BP108P.02	Identify percentage purity of given pharmaceutical drugs by	1,2,3,4,5,6,
	titrimetric analysis along with preparation of standardization	7,8,9,11
	of secondary standard solutions.	
BP108P.03	Perform and carryout various electrochemical titrations.	1,2,3,4,5,6,
		7,8,9,11

BP109P	Pharmaceutics I – Practical	PO
BP109P .01	Illustrate the preparation, labeling aspects and evaluation of	1,2,3,4,6,7,8,9,11
	liquid dosage form.	, ,-, ,-,-,-,
BP109P .02	Illustrate the preparation, labeling aspects and evaluation of	1,2,3,4,6,7,8,9,11
BI 1071 .02	granules and powders.	1,2,3,4,0,7,0,7,11
BP109P .03	Illustrate the preparation, labeling aspects and evaluation of	1,2,3,4,6,7,8,9,11
DF 109F .03	suppositories and semisolids.	1,2,3,4,0,7,0,9,11

BP110P	Pharmaceutical Inorganic Chemistry – Practical	PO
BP110P.01	Identify impurities and carry out limit test and various tests for	1,2,3,4,6,7,8,9,11
DI 1101.01	purity for pharmaceuticals.	1,2,3,4,0,7,0,9,11
BP110P.02	Identify the inorganic compounds by undertaking various	1,2,3,4,6,7,8,9,11
DF 110F.02	identification tests.	1,2,3,4,0,7,0,9,11
BP110P.03	Justify the preparation method for inorganic pharmaceuticals.	1,2,3,4,6,7,8,9,11



BP111P	Communication skills	PO
BP111P.01	Comprehend the concept of communication and describe the basic communication skills and ways to overcome barriers.	2,5,8,11
BP111P.02	Convert the conceptual understanding of communication into everyday practice for better business communication.	2,5,8,11
BP111P.03	Apply the concept of positive thinking to keep a good stead at the time of crisis.	2,5,8,11

BP201T	Human Anatomy and Physiology II – Theory	PO
BP201T.01	Explain anatomy and physiology of nervous system.	1,3,4,6,7,8,11
BP201T.02	Explain anatomy and physiology of Digestive & respiratory system.	1,3,4,5,6,7,8,11
BP201T.03	Explain anatomy and physiology of urinary system.	1,3,4,6,7,8,11
BP201T.04	Explain anatomy and physiology Endocrine, Reproductive and genetics.	1,2,3,4,5,7,8,11

BP202T	Pharmaceutical Organic Chemistry I – Theory	PO
BP202T.01	Explain IUPAC systems, classification, nomenclature and	1,2,6,7,8,11
DI 2021.01	isomerism of organic compounds.	1,2,0,7,0,11
BP202T.02	Explain the hybridization, elimination reactions of alkanes,	1,2,6,7,8,11
DF 2021.02	alkenes and alkyl halides and stability of conjugated dienes.	1,2,0,7,0,11
BP202T.03	Describe the reaction mechanisms, qualitative tests, structure and	1,2,6,7,8,11
DF 2021.03	uses of alcohols and carbonyl compounds.	1,2,0,7,0,11
BP202T.04	Describe the reaction mechanisms, qualitative tests, structure and	1,2,6,7,8,11
DF 2021.04	uses of carboxylic acids and aliphatic amines.	1,2,0,7,0,11

BP203T	Biochemistry – Theory	PO
BP203T.01	Classify the biomolecules and explain their importance	1,2,4,6,7,8,9,11
BP203T.02	Relate the metabolism of nutrient molecules in physiological and pathological conditions.	1,2,4,6,7,8,9,11
BP203T.03	Explain the catalytic role of enzymes, importance of enzyme inhibitors in design of new drugs, therapeutic and diagnostic applications of enzymes.	1,2,4,6,7,8,9,11
BP203T.04	Illustrate the genetic organization of mammalian genome and functions of DNA in the synthesis of RNAs and proteins	1,2,4,6,7,8,9,11



BP204T	Pathophysiology – Theory	PO
BP204T.01	Describe the principles of cell injury, adaptation and mechanism involved in the inflammation and repair.	1,2,6,8,9,11
BP204T.02	Describe etiology and pathogenesis of various diseases and complications.	1,2,6,8,9,11
BP204T.03	Describe etiology and pathogenesis of liver, cancer, bone and joints.	1,2,6,8,9,11
BP204T.04	Describe etiology and pathogenesis of Infectious diseases and sexually transmitted diseases	1,2,6,8,9,11

BP205 T	Computer applications in Pharmacy	PO
BP205T.01	Understand the various types of application of computers	1,2,3,4,5,6,7,8,11
DI 2031.01	in pharmacy.	1,2,3,4,3,0,7,0,11
BP205T.02	Understand basic handling techniques of computers.	1,2,3,4,5,6,7,8,11
BP205T.03	Describe the various types of databases.	1,2,3,4,5,6,7,8,11
BP205T.04	Know the various applications of databases in pharmacy.	1,2,3,4,5,6,7,8,11

BP206 T	Environmental Sciences	PO
	Impart basic knowledge about the environment and	
BP206T.01	awareness about environmental problems its allied	1,2,3,4,5,6,7,8,9,10,11
	problems.	
BP206T.02	Motivate learner to participate in environment protection	1,2,3,4,5,6,7,8,9,10,11
BF 2001.02	and environment improvement.	1,2,3,4,3,0,7,6,9,10,11
BP206T.03	Acquire skills to help the concerned individuals in	1 2 2 4 5 6 7 9 0 10 11
BF2001.03	identifying and solving environmental problems.	1,2,3,4,5,6,7,8,9,10,11
BP206T.04	Strive to attain harmony with Nature.	1,2,3,4,5,6,7,8,9,10,11

BP207P	Human Anatomy and Physiology II –Practical	PO
BP207P.01	Describe and recognize different types of organ system and	
	family planning devices and pregnancy diagnosis test. With	1,3,4,6,7,8,11
	models, charts and specimen.	
BP207P.02	Identify and examination of the functions of cranial nerves.	1,3,4,5,6,7,8,11
BP207P.03	Interpret the recording of body temp and BMI total blood count.	1,2,4,6,7,8,11



BP208P	Pharmaceutical Organic Chemistry I— Practical	PO
BP208P.01	Identify unknown organic compounds by qualitative analysis.	1,2,3,4,5,6,
		7,8,9,11
BP208 P.02	Plan the synthesis of solid derivatives from organic compounds.	1,2,3,4,5,6,
		7,8,9,11
BP208 P.03	Illustrate the molecular models of compounds using atomic	1,2,3,5,6,
	models sets.	7,8,9,11

BP209P	Biochemistry – Practical	PO
BP209P.01	Interpret the biomolecules like proteins and carbohydrates	1,2,3,4,6,7,8,9,11
BP209P.02	Interpret the biochemical investigation of blood and urine	1,2,3,4,6,7,8,9,11
BP209P.03	Plan the effect of different factors on enzymatic activity & contrast the importance of buffer solution in pharmaceuticals.	1,2,3,4,6,7,8,9,11

### SECOND YEAR B. PHARMACY

BP301T	Pharmaceutical Organic Chemistry II – Theory	PO
BP301T.01	Explain IUPAC systems, classification, nomenclature and isomerism of organic compounds.	1,2,6,7,8,11
BP301T.02	Explain the hybridization, elimination reactions of alkanes, alkenes and alkyl halides and stability of conjugated dienes.	1,2,6,7,8,11
BP301T.03	Describe the reaction mechanisms, qualitative tests, structure and uses of alcohols and carbonyl compounds.	1,2,6,7,8,11
BP301T.04	Describe the reaction mechanisms, qualitative tests, structure and uses of carboxylic acids and aliphatic amines.	1,2,6,7,8,11

BP302T	Physical Pharmaceutics I – Theory	PO
DD202E 01	Relate various physicochemical properties of drug molecules	1.2.2.6.7.0.0.11
BP302T.01	in the designing the dosage forms.	1,2,3,6,7,8,9,11
BP302T.02	Explain principles of states and properties of matter.	1,2,3,4,6,7,8,9,11
BP302T.03	Describe the classification, methods and application of	1,2,3,4,6,7,8,9,11
BP3021.03	complexation and protein binding.	
DD202T 04	Describe the preparation and application of buffers and	1 2 2 4 6 7 9 9 11
BP302T.04	isotonic solutions in pharmaceutical and biological system.	1,2,3,4,6,7,8,9,11



BP303T	Pharmaceutical Microbiology – Theory	PO
BP303T.01	Explain various methods of identification, cultivation and preservation of microorganisms.	1,2,3,4,6,7, 8,9,10,11
BP303T.02	Discuss the importance and implementation of sterilization in pharmaceutical processing and industry.	1,2,3,4,6, 7,8,9,11
BP303T.03	Interpret sterility testing of pharmaceutical products.	1,2,3,4,6,7,8,9,11
BP303T.04	Explain the cell culture technology and its applications in pharmaceutical industries.	1,2,3,4,6, 7,8,9,11

BP304T	Pharmaceutical Engineering – Theory	PO
BP304T.01	Explain the principle, equipment and application for flow of	1,2,3,4,5,6,
DF 3041.01	fluids and heat transfer.	7,8,9,11
DD204T 02	Explain the various unit operations used in pharmaceutical	1,2,3,4,5,6,
BP304T.02	industries.	7,8,9,11
DD204T 02	Explain the various processes involved in pharmaceutical	1,2,3,4,5,6,
BP304T.03	manufacturing processes.	7,8,9,11
BP304T .04	Describe the various preventive methods for corrosion and	1,2,3,4,5,6,
Dr3041.04	pollution control in pharmaceutical plant construction.	7,8,9,10,11

BP305P	Pharmaceutical Organic Chemistry II – Practical	PO
DD205D 01	Plan the synthesis of various organic compounds by different	1,2,3,4,5,
BP305P.01	chemical reactions.	6,7,8,9,11
BP305P.02	Illustrate the purity of oils and fats.	1,2,3,4,5,6,
		7,8,9,11
BP305P.03	Relate the experiments involving recrystallisation and distillation techniques.	1,2,3,4,5,6 7,8,9,11

BP306P	Physical Pharmaceutics I – Practical	PO
BP306P.01	Identify the physical properties in the formulation development.	1,2,3,4,6,7,8,9,11
BP306P.02	Identify the HLB value, CMC of the surfactant for evaluation of dosage forms.	1,2,3,4,6,7,8,9,11
BP306P.03	Illustrate the stability constant by various methods.	1,2,3,4,6,7,8,9,11



BP307P	Pharmaceutical Microbiology – Practical	PO
BP303T.01	Relate various equipment and processing in conduct of experiment microbiology.	1,2,4,5,6,7, 8,9,11
BP303T.02	Illustrated microbiological assay, biochemical testing, and sterility testing standardization of Pharmaceuticals.	1,2,3,5,6, 8,9,11
BP303T.03	Illustrated the identification, cultivation and preservation of various microorganisms.	1,2,3,4,5,6, 7,8,9,11

BP 308P	Pharmaceutical Engineering –Practical	PO
DD200D 01	Illustrate the determination of heat transfer on various	1,2,3,4,5,6,
BP308P.01	techniques.	7,8,9,11
DD200D 02	Relate the various unit operations used in pharmaceutical	1,2,3,4,5,6,
BP308P.02	industries.	7,8,9,11
DD200D 02	Relate the various processes involved in pharmaceutical	1,2,3,4,5,6,
BP308P.03	manufacturing processes.	7,8,9,11

BP401T	Pharmaceutical Organic Chemistry III- Theory	PO
BP401T.01	To explain the stereo chemical aspects of organic compounds	1 2 4 10 11
BP4011.01	and stereo chemical reactions with respect to optical isomerism.	1,3,4,10,11
	To explain the stereo chemical aspects of organic compounds	
BP401T.02	and stereo chemical reactions with respect to geometrical	1,3,4,10,11
	isomerism.	
DD401T 02	To summarize the synthesis, chemical reactions, and medicinal	1 2 4 10 11
BP401T.03	uses of some heterocyclic compounds.	1,3,4,10,11
DD401T 04	To illustrate statement, mechanism, and applications of some	1 2 4 10 11
BP401T.04	named reactions.	1,3,4,10,11

BP402T	Medicinal Chemistry I – Theory	PO
BP402T.01	Explain the physicochemical properties and drug metabolism for various categories of drugs.	1,2,3,5,6, 7,8,9,11
BP402T.02	Explain the classification, mechanism of action and uses of different classes of drugs.	1,2,3,5,6, 7,8,9,11
BP402T.03	Relate the Structure activity relationship of different classes of drugs.	1,2,3,5,6, 7,8,9,11
BP402T.04	Plan the synthesis of different class of selective medicinal drugs.	1,2,3,4,5,6, 7,8,9,11



BP403T	Physical Pharmaceutics II – Theory	PO
BP403T.01	Explain the general characteristics and classification of	1 2 6 7 9 0 11
BP4031.01	colloidal dispersion.	1,2,6,7,8,9,11
BP403T.02	Describe principles involved in the rheological characteristics	124679011
BP4031.02	and coarse dispersion.	1,2,4,6,7,8,9,11
BP403T.03	Explain the micromeritic of the powder and their application in	124679011
BP4031.03	solid dosage forms.	1,2,4,6,7,8,9,11
BP403T.04	Describe the kinetics study and accelerated stability testing in	124679011
Dr4031.04	pharmaceutical dosage forms.	1,2,4,6,7,8,9,11

BP404T	Pharmacology I – Theory	PO
BP404T.01	Explain the general pharmacology of the drugs regarding pharmacokinetics & pharmacodynamics.	1,3,4,6,9,10, 11
BP404T.02	Relate the drug discovery & clinical trials for a new drug.	1,2,3,4,9,10,11
BP404T.03	Describe the Pharmacology of drugs acting on peripheral nervous system.	1,3,4,9,10,11
BP404T.04	Describe the pharmacology of drugs acting on central nervous system.	1,3,4,6,9,10,11

BP405T	Pharmacognosy and Phytochemistry I– Theory	PO
BP405T.01	Explain the Pharmacognostic scheme of the crude drug of natural origin.	1,2,3,4,6, 7,9,11
BP405 T.02	Explain the role of pharmacognosy in various systems of medicines and study the Pharmacognostic scheme of secondary metabolites.	1,2,3,4,6, 7,8,9,1
BP405 T.03	Describe the cultivation, collection, processing and storage of natural origin of medicinal drugs and development and application of tissue culture.	1,2,3,4,6, 7,8,9,11
BP405 T.04	Explain the Pharmacognostic scheme of drugs of natural origin containing plant products, primary metabolites and marine drugs.	1,2,3,4,6, 7,8,9,11

BP406P	Medicinal Chemistry I – Practical	PO
BP406P.01	Synthesize drugs and drug intermediates of selective medicinal	1,2,3,4,5,6,
D1 4001 .01	compounds.	7,8,9,11
BP406P.02	Assess the percentage purity of selective drugs.	1,2,3,4,5,6,
DF 400F .02		7,8,9,11
BP406P.03	Assess the partition coefficient of drugs.	1,2,3,4,5,6,
DF400P.03		7,8,9,11



BP407P	Physical Pharmaceutics II – Practical	PO
BP407P.01	Identify the micromeritic properties of the powders.	1,2,3,4,6,7,8,9,11
BP407P .02	Identify the physical properties in the formulation development.	1,2,3,4,6,7,8,9,11
BP407P.03	Relate the principle of chemical kinetics and apply them in stability testing of formulation.	1,2,3,4,6,7,8,9,11

BP408P	Pharmacology I – Practical	PO
BP408P.01	Summarize about the construction and working the about basic instruments, common laboratory animals and maintenance as per CPCSEA guidelines used in experimental pharmacology.	1,4,10,11
BP408P.02	Describe and demonstrate the common laboratory techniques like routes of drug administration, blood withdrawal and plasma separation, anesthetics and euthanasia used for animal studies.	1,3,4,10,11
BP408P.03	Interpret and observe the effect of different drugs on animals by simulated experiments.	1,3,4,9,10,11

BP409P	Pharmacognosy and Phytochemistry I – Practical	PO
BP 409P.1	Illustrate the analysis of crude drugs by chemical tests.	1,2,3,4,6,7,8,9,11
BP 409P.2	Determine the characteristics of crude drugs by physical method.	1,2,3,4,6,7,8,9,11
BP 409P.3	Describe the physical characteristics of crude drug by microscopy method.	1,2,3,4,6,7,8,9,11

### THIRD YEAR B. PHARMACY

BP501T	Medicinal Chemistry II – Theory	PO
BP501T.01	Explain the classification and uses of different classes of drugs.	1,2,3,5,6,7,8,9,11
BP501T.02	Relate the Mechanism of action of different classes of drugs.	1,2,3,5,6,7,8,9,11
BP501T.03	Relate the Structure activity relationship of different classes of	1,2,3,5,6,7,8,9,11
<b>D1</b> 3011.03	drugs.	1,2,3,3,0,7,0,7,11
BP501T.04	Plan the synthesis of different class of selective medicinal	12245679011
DF 3011.04	drugs.	1,2,3,4,5,6,7,8,9,11



BP502T	Industrial Pharmacy I– Theory	PO
BP502T.01	Explain the preformulation studies and relate the packaging materials in the development of various dosage form.	1,2,3,4,5,6, 7,8, 9,11
BP502T.02	Describe the formulation and quality control test for solid and liquid dosage forms such as tablets, pellets, capsules and liquid orals.	1,2,3,4,5,6, 7,8,9,11
BP502T.03	Describe the production condition, formulation and quality control test of parentrals, ophthalmic products and aerosols.	1,2,3,4,5,6, 7,8,9,11
BP502T.04	Explain the formulation considerations for the various cosmetic preparations.	1,2,3,4,5,6, 7,8,9,11

BP503T	Pharmacology II – Theory	PO
DD502T 01	Explain the pharmacological action and its relevance of drugs	1,2,6,7,
BP503T.01	acting on CVS and urinary system.	8,9,11
BP503T.02	Explain the pharmacological action and its relevance of drugs	1,2,6,7,
BP3031.02	acting on autacoids and drugs.	8,9,11
DD502T 02	Explain the pharmacological action and its relevance of drugs	1,2,6,7,
BP503T.03	acting on endocrine system.	8,9,11
DD502T 04	Describe the principles, types and application of bioassay of	1,2,4,6,7,
BP503T.04	drugs.	8,9,11

BP504T	Pharmacognosy and Phytochemistry II – Theory	PO
DD504T 01	Explain the basics of metabolic biosynthesis and biogenesis	1 2 2 4 6 7 9 0 11
BP504T.01	pathways and their determinations.	1,2,3,4,6,7,8,9,11
DD504T 02	Describe the various chemical categories of secondary	1 2 2 4 6 7 9 0 11
BP504T.02	metabolites and its Pharmacognostic studies.	1,2,3,4,6,7,8,9,11
BP504T.03	Illustrate the evaluation of Phytoconstituents by isolation,	1 2 2 4 6 7 9 0 11
DP3041.03	identification, estimation, production and utilization.	1,2,3,4,6,7,8,9,11
BP504T.04	Relate the basics in Phytochemistry, its extraction and	1 2 2 4 6 7 9 0 11
DP3041.04	application of various techniques of screening.	1,2,3,4,6,7,8,9,11



BP505T	Pharmaceutical Jurisprudence – Theory	PO
BP505T.01	Explain the import, manufacture, sale of drugs as per Drugs & cosmetics act 1940 and rules1945.	1,2,4,5,6,7, 8,9,10,11
BP505T.02	Summarize the administration of the Drug & Cosmetics act and rules.	1,2,3,5,6,7, 8,9,10,11
BP505T.03	Describe the various pharmaceutical acts & laws M & T P Act 1955,N D &P S Act 1985,D& M R Act, Prevention of Cruelty to animalsAct-1960, DPCO act, MTP Act, RTIACT, IPR.	1,2,3,5,6,7, 8,9,10,11
BP505T.04	Describe the development of pharmaceutical legislation of India &pharmacist in relation to ethics practice.	1,2,3,5,6,7, 8,9,10,11

BP506P	Industrial Pharmacy I – Practical	PO
BP 506P.01	Illustrate the preformulation and evaluation of packaging materials.	1,2,3,4,6,7,8,9,11
BP 506P.02	Relate the formulation and evaluation of tablets, capsules and compare with marketed products.	1,2,3,4,6,7,8,9,11
BP 506P.03	Relate the preparation of parentral and ophthalmic dosage forms and cosmetic creams.	1,2,3,4,6,7,8,9,11

BP507P	Pharmacology II – Practical	PO
BP507P.01	Summarize the preparation of physiological salt solutions in the study of effect of drugs on various receptor & animal model.	1,3,4,6,9,10,11
BP507P.02	Compare the bioassay of drugs using animal models by various methods.	1,3,4,6,9,10,11
BP507P.03	Explain the analgesic & Anti-inflammatory activity of drugs by using various methods.	1,3,4,6,9,10,11

BP508P	Pharmacognosy and Phytochemistry II – Practical	PO
BP508P.01	Assess the morphology, histology, powder characteristics and	1,2,3,4,5,6,7,8,9,11
DF 306F.01	extraction, isolation and detection of crude drugs.	1,2,3,4,3,0,7,6,9,11
BP508P.02	Plan the separation and detection of Phytoconstituents by	1,2,3,4,5,6,7,8,9,11
DP308P.02	different chromatographic methods.	1,2,5,4,5,0,7,8,9,11
BP508P.03	Assess the analysis of crude drugs by chemical test.	1,2,3,4,5,6,7,8,9,11



BP601T	Medicinal Chemistry III – Theory	PO
BP601T.01	Explain the classification and uses of different classes of drugs.	1,2,3,5,6,7,8,9,11
BP601T.02	Relate the Mechanism of action and Structure activity relationship of different classes of drugs.	1,2,3,5,6,7,8,9,11
BP601T.03	Plan the synthesis of different class of selective medicinal drugs.	1,2,3,4,5,6,7,8,9,11
BP601T.04	Explain the approaches in drug design, concept of QSAR and combinatorial chemistry.	1,2,3,4,5,6,7,8,9,11

BP602T	Pharmacology III – Theory	PO
BP602T.01	Explain the pharmacological action and its relevance of drugs acting on respiratory system and gastrointestinal	1,2,6,7,8,9,11
	system.	
	Explain general principle of chemotherapy of disease,	
BP602T.02	mechanism of bacterial resistance and pharmacology of	1,2,6,7,8,9,11
	antimicrobial and antineoplastic agent.	
	Explain the pharmacological action and its relevance of	
BP602T.03	drugs acting on immunopharmacological drugs and	1,2,6,7,8,9,11
	chronopharmacology.	
BP602T.04	Summarize principle of toxicology and treatment of various	1,2,6,7,8,9,11
D1 0021.04	acute and chronic poisoning.	1,2,0,7,0,9,11

BP603T	Herbal Drug Technology – Theory	PO
BP603T.01	Explain the source of herbs, identification, authentication, cultivation and processing of herbal raw materials in various systems of medicines.	1,2,3,6,7,8,9,11
BP603T.02	Explain the general aspects benefits and role of neutraceuticals and herbal cosmetics.	1,2,3,6,7,8,9,11
BP603T.03	Relate the evaluation and assessment of herbal drugs by WHO and ICH guidelines.	1,2,3,6,7,8,9,11
BP603T.04	Use of schedule T in the manufacturing of herbal drugs and Indian system of medicine.	1,2,3,6,7,8,9,11

BP604T	Bio pharmaceutics and Pharmacokinetics – Theory	PO
BP604T.01	Describe the basic concept of absorption, distribution,	1,2,3,6,7,8,9,11
DI 0041.01	metabolism and excretion of drugs.	1,2,3,0,7,6,7,11
BP604T.02	Explain the principles and methods of bioavailability and	1,2,3,4, 6,7,8,9,11
DF 0041.02	bioequivalence studies.	1,2,3,4, 0,7,0,9,11
BP604T.03	Describe the primary, secondary and tertiary	1,2,3,6,7,8,9,11
DI 0041.03	pharmacokinetic parameters in clinical setting.	1,2,3,0,7,6,9,11
BP604T .04	Explain the factors and parameters of nonlinear	1,2,3,6,7,8,9,11
DF0041.04	pharmacokinetics.	1,2,3,0,7,0,9,11



BP605T	Pharmaceutical Biotechnology – Theory	PO
BP605T.01	Summarize the importance of Immobilized enzymes in Pharmaceutical Industries.	1,2,4,6,7,8, 9,11
BP605T.02	Describe genetic engineering applications in relation to production of pharmaceuticals.	1,2,4,7,8,9,11
BP605T.03	Explain of Monoclonal antibodies in Industries.	1,2,4,5,7,8,11
BP605T.04	Illustrated the use of microorganisms in fermentation technology.	1,2,4,5,6,7,8,9,11

BP606T	Quality Assurance –Theory	PO
BP606T.01	Explain the concept and control of quality management systems.	1,2,3,5,6,7,8,9,11
BP606T.02	Describe the quality control tests for raw materials, containers and secondary packaging materials.	1,2,3,5,6,7,8,9,11
BP606T.03	Describe the documentation in pharmaceutical industry.	1,2,3,5,6,7,8,9,11
BP606T.04	Explain the general principles of calibration, validation process and equipment qualification.	1,2,3,4,5,6,7,8,9,11

BP607P	Medicinal chemistry III – Practical	PO
BP607P.01	Synthesize drugs and drug intermediates of selective	1,2,3,4,5,6,7,
DI 00/1.01	medicinal compounds.	8,9,11
BP607P.02	Assess the percentage purity and physichochemical	1,2,3,4,5,6,7,
BF00/F.02	properties of selective drugs.	8,9,11
BP607P.03	Illustrate the structures and reactions using chem draw.	1,2,3,4,5,6,7,
DF00/F.03		8,9,11

BP608P	Pharmacology III – Practical	PO
BP608P.01	Demonstrate the screening of different category of drugs.	1,5,6,7,8,11
BP608P.02	Assess the dose calculation, acute oral toxicity study and	1,9,10,11
	acute skin and eye irritation study.	
BP608P.03	Estimation of serum biochemical parameters and	1,2,3,9,11
	biostatistics methods in experimental pharmacology.	

BP609P	Herbal Drug Technology – Practical	PO
BP609P.01	Assess the evaluation of crude drugs by preliminary phytochemical screening.	1,2,3,4,6,7,8,9,11
BP609P.02	Assess the preparation and standardization of Herbal drugs and herbal cosmetics by evaluation parameters.	1,2,3,4,6,7,8,9,11
BP609P.03	Illustrate the Monograph analysis of herbal drugs from recent Pharmacopoeias.	1,2,3,4,6,7,8,9,11



### FINAL YEAR B. PHARMACY

BP701T	Instrumental Methods of Analysis – Theory	PO
BP701T.01	To summarize the interaction of matter with electromagnetic radiations and instrumentation in various spectroscopic techniques.	1,2,3,4,6,7, 8,9,10,11
BP701T.02	To illustrate the principle and technique behind various chromatographic separations.	1,2,3,4,6,7, 9,10,11
BP701T.03	To emphasize qualitative and quantitative analysis of drugs using various chromatographic techniques.	1,2,3,4,6, 8,10,11
BP701T.04	To articulate the qualitative and quantitative applications of various spectroscopic techniques in the analysis of drugs.	1,2,3,4,6,8,10,11

BP702T	Industrial Pharmacy II – Theory	PO
BP702T.01	Describe the process of pilot plant scale up of	1,2,3,4,5,6,7,8,9,11
DI 7021.01	pharmaceutical dosage forms.	1,2,3,4,3,0,7,0,9,11
BP702T.02	Develop the practice and the process of technology transfer	1,2,3,4,5,6,7,8,9,11
DF /021.02	from lab scale to production.	1,2,3,4,3,0,7,0,9,11
BP702T.03	Explain the different laws, approval process, role and	1,2,3,4,5,6,7,8,9,11
DF 7021.03	responsibility of Regulatory agencies.	1,2,3,4,3,0,7,0,9,11
BP702T.04	Explain the different Quality Management systems and their	1,2,3,4,5,6,7,8,9,11
DF /021.04	role.	1,4,5,4,5,0,7,0,9,11

BP703T	Pharmacy Practice – Theory	PO
BP703T.01	Describe the knowledge on organization of hospital, community pharmacy, various methods of distribution and hospital formulary in hospitals and apply it in in the practice of pharmacy.	1,2,5,6,7,8,9,11
BP703T.02	Categorize the role of hospital pharmacist in pharmacy, therapeutic committee, drug information services, patient counseling, education and training programme in hospitals.	1,2,5,6,7,8,9,11
BP703T.03	Explain concept, function and responsibility of clinical pharmacist.	1,2,5,6,7,8,9,11
BP703T.04	Explain organization of drug store management and inventory control.	1,2,5,6,7,8,9,11

<b>BP704T</b>	Novel Drug Delivery System – Theory	PO
BP704T.01	Explain the various approaches for development of novel drug delivery system.	1,2,3,4,6,7,8,9,11
BP704T.02	Identify the criteria for selection of drugs and polymer for the development of novel drug delivery system.	1,2,3,4,6,7,8,9,11
BP704T.03	Explain the concept, methodology and their applications by various drug delivery systems.	1,2,3,4,6,7,8,9,11
BP704T.04	Explain the concept, methodology and their applications by various targeted drug delivery system.	1,2,3,4,6,7,8,9,11



BP705P	Instrumental Methods of Analysis – Practical	PO
BP705P.01	To interpret the absorption maxima, assay by Colorimeter and	1,2,3,4,5,6,
BP/03P.01	UV Visible Spectrophotometer.	8,9,10,11
BP705P.02	To relate the estimation of concentration of ions by Flame	1,2,3,4,5,
DP/03P.02	Photometer and turbidance by Nepheloturbidimeter.	6,9,10,11
BP705P.03	To relate the purity of the drugs by various chromatographic techniques such as TLC, PC, Column chromatography, and HPLC.	1,2,3,4,5, 8,9,11

BP801T	Biostatistics and Research Methodology	PO
BP801T.01	Relate the basic terminologies involved in Statistics and measures of central tendency, dispersion and correlation.	1,3,4,6,7,8,11
BP801T.02	Compare and measure the various parametric and non-parametric statistical techniques.	1,3,4,5,6,7,8,11
BP801T.03	Explain the concept of research and various methodologies involved in research.	1,3,4,6,7,8,11
BP801T.04	Explain the measure of different statistical software's using design of experiments and clinical trial study.	1,2,3,4,5,7,8,11

BP802T	Social and Preventive Pharmacy	PO
BP802T.01	Illustrate the concept and evaluation of public health.	1,5,6,7,8,9,11
BP802T.02	Explain the principle on prevention and control of	1,5,6,7,8,9,11
DF 8021.02	communicable and non communicable diseases.	
BP802T.03	Identify current issues related with various diseases in	15679011
BF 8021.03	related to the prevention and control within the country.	1,5,6,7,8,9,11
	Role play of the community services in improvement of	
BP802T.04	ruler sanitation, urban health care and promotion of school	1,5,6,7,8,9,11
	health.	

BP803ET	Pharma Marketing Management	PO
BP803ET.01	Explain the general concepts and scope of pharmaceutical	1,2,5,6,7,8,9,11
DI 603E1.01	marketing.	1,2,3,0,7,0,9,11
BP803ET.02	Describe the product decision and management in	1,2,5,6,7,8,9,11
DF 603E1.02	pharmaceutical industry.	1,2,3,0,7,0,9,11
BP803ET.03	Describe the methods of promotion, role of PSR and	1,2,5,6,7,8,9,11
DF 603E1.03	various applications of marketing channels.	1,2,3,0,7,0,9,11
BP803ET.03	Relate the emerging concepts in marketing and price	1,2,5,6,7,8,9,11
DI 003E1.03	management as per BPCO and NPPA.	1,2,3,0,7,0,9,11



BP805ET	Pharmacovigilance	PO
BP805ET.01	Discuss the importance of drug safety monitoring and the development of pharmacovigilance programme.	1,2,6,7,8,9,11
BP805ET.02	Identify methods and management of adverse drug reaction.	1,2,3,4,6,7,8,9,11
BP805ET.03	Assess international standards for classification of diseases and drugs.	1,2,3,4,6,7,8,9,11
BP805ET.04	Explain various methods of programmes and terminologies in drug safety surveillance and communication in pharmacovigilance programme.	1,2,3,4,6,7,8,9,11



### M. Pharmacy (Pharmaceutics)

### Sem I

MPH 101T	Modern pharmaceutical analytical techniques	PO
MPH101T.01	Illustrate assay of single and multiple component	1 2 2 4 5 6 7 9 0 11
MPH1011.01	Pharmaceuticals by using various analytical instruments.	1,2,3,4,5,6,7,8,9,11
MPH101T.02	Describe basic practical skills using Instrumentation	1 2 2 4 5 6 7 9 0 11
MPH1011.02	techniques.	1,2,3,4,5,6,7,8,9,11
MDII101T 02	Identify the theoretical knowledge on various instrumental	1 2 2 4 5 6 7 9 0 11
MPH101T.03	techniques for analysis of organic substances.	1,2,3,4,5,6,7,8,9,11
MDII101T 04	Explain the knowledge in developing new procedures for	1 2 2 4 5 6 7 9 0 11
MPH101T.04	analysis.	1,2,3,4,5,6,7,8,9,11

MPH 102T	Drug Delivery System	PO
	Describe concept, principle involved, formulation and	
MPH102T.01	evaluation methods for sustained release and	1,2,3,4,6,7,
	controlled formulations.	8,9,11
	Explain the principle and fundamentals involved in	
MPH102T.02	personalized medicines and rate controlled drug	1,2,3,4,6,7,8,9,11
	delivery systems.	
	Explain the principle, formulation and evaluation of	1 2 2 4 6 7 9 0 11
MPH102T.03	ocular and transdermal drug delivery systems.	1,2,3,4,6,7,8,9,11
	Describe fundamentals, formulation and evaluation of	
MPH102T.04	drug delivery system of proteins, macromolecules	1,2,3,4,6,7,8,9,11
	and vaccines.	

MPH 103T	Modern Pharmaceutics	PO
MPH103T.01	Describe preformulation concepts, formulation considerations of pharmaceutical Dispersion parenterals and application of optimization technique in formulation design.	1,2,3,4,6,7,9,10,11
MPH103T.02	Explain the elements, methods of equipment and process parameters validation.	1,2,3,4,6,7,9,10,11
MPH103T.03	Explain the industrial management and cGMP considerations.	1,2,3,4,6,7,9,10,11
MPH103T.04	Explain the principles involved in statistics and kinetic models in formulation consideration.	1,2,3,4,6,7,9,10,11



MPH 104T	Regulatory affairs	PO
MPH104T.01	Describe the chemistry, manufacturing controls and	1,2,3,4,6,
MPH1041.01	various regulatory agencies involved.	7,8,9,10,11
MDII104T 02	Explain the regulatory requirements for drug approval	1,2,3,4,6,7,
MPH104T.02	process.	8,9,10,11
MPH 104T.03	Explain the non clinical drug development process.	1,2,3,4,6,
WII 11 1041.03		7,8,9,10,11
MPH 104T.04	Illustrate the clinical trial requirements and protocols.	1,2,3,4,6,7,
WII II 1041.04		8,9,10,11

MPH 105P	Pharmaceutics Practical I	PO
MDII 105D 01	Illustrate the analysis of pharmacopoeial compounds and	1 2 2 4 5 6 7 9 9 11
MPH 105P.01	formulation using various instrumental techniques.	1,2,3,4,5,6,7,8,9,11
MDII 105D 02	Design the formulation and evaluation methods for	1 2 2 4 5 6 7 9 0 11
MPH 105P.02	various sustained and controlled release formulation.	1,2,3,4,5,6,7,8,9,11
	Summarize the pre formulation concept, micromeritic	
MPH 105P.03	properties and application of pharmacokinetic models	1,2,3,4,5,6,7,8,9,11
	for various dosage forms.	

### Sem II

MPH 201T	Molecular Pharmaceutics	PO
MPH201T.01	Explain concepts, preparations and evaluation of various targeted drug delivery systems.	1,2,3,4,6,7,8,9,11
MPH201T.02	Explain preparation, evaluation and applications of micro particulate drug delivery systems.	1,2,3,4,6,7,8,9,11
MPH201T.03	Explain the principle, preparation and evaluation involved in pulmonary drug delivery systems.	1,2,3,4,6,7,8,9,11
MPH201T.04	Describe fundamentals and applications of nucleic acid based therapeutic delivery system.	1,2,3,4,6,7,8,9,11



MPH 202T	Advanced Bio pharmaceutics and pharmacokinetics	PO
MPH202T.01	Explain the principles of absorption, distribution, metabolism and excretion of the drug from various dosage forms.	1,2,3,4,5, 6,7,8,11
MPH202T.02	Explain the various biopharmaceutical considerations in drug product design and in vitro drug product performance.	1,2,3,4,5, 6,7,8,11
MPH202T.03	Describe the basic consideration and pharmacokinetic model and application of pharmacokinetics in conventional and modified drug delivery system.	1,2,3,4,5, 6,7,8,11
MPH202T.04	Explain the drug product performance in vivo Bioavailability and bioequivalence for generic drugs and biologics.	1,2,3,4,5, 6,7,8,11

MPH203T	Computer Aided Drug Delivery	PO
MPH203T.01	Explain the QbD in formulation development by computer modeling and statistical application.	1,2,3,4,5,6,7,8,11
MPH203T.02	Explain the computational modeling of drug disposition.	1,2,3,4,5,6,7,8,11
MPH203T.03	Explain the computer-aided biopharmaceutical characterization, simulations in pharmacokinetics and pharmacodynamics, and clinical development.	1,2,3,4,5,6,7,8,11
MPH203T.04	Explain artificial intelligence, robotics and Computational fluid dynamics in pharmaceuticals.	1,2,3,4,5,6,7,8,11

MPH204T	Cosmetics & Cosmeceuticals	PO
MPH204T.01	Describe the regulatory provisions related to the import	1,2,3,4,5,6,7,8,11
	and manufacture of cosmetics as per the Drugs and	
	Cosmetics Act 1940 and the Rules 1945.	
MPH204T.02	Explain various formulation considerations in the various	1,2,3,4,5,6,7,8,11
	cosmeticeutical products.	
MPH204T.03	Explain the biological aspects and various problems	1,2,3,4,5,6,7,8,11
	related to the skin, hair and oral hygienic products.	
MPH204T.04	Explain herbal ingredients used in formulating various	1,2,3,4,5,6,7,8,11
	cosmeticeutical products.	



MPH205P	Pharmaceutics Practical II	PO
MPH205P.01	Design the formulation and evaluation methods for various NDDS.	1,2,3,4,5,6,7,8,9,11
MPH205P.02	Illustrate dissolution, protein binding, bioavailability studies and application of pharmamacokinetics <i>in-vivo in-vitro</i> correlation by software.	1,2,3,4,5,6,7,8,9,11
MPH205P.03	Design the formulation and evaluation methods for various cosmetics product and herbal cosmetics.	1,2,3,4,5,6,7,8,9,11

### Sem III

MRM301T	Research Methodology and Biostatistics	PO
MRM301T.01	Explain the concept of general &Medical Research and various methodologies and guidelines involved in research.	1,2,3,4,5,6,7,8,11
MRM301T.02	Apply Statistics and Statistical tests for significance and measure the various parametric and non-parametric tests.	1,2,3,4,5,6,7,8,11
MRM301T.03	Relate to the CPCSEA guidelines for laboratory animal facility and handling.	1,2,3,4,5,6,7,8,11
MRM301T.04	Articulate the basic principles of Helsinki in medical research and medical care.	1,2,3,4,5,6,7,8,11



### M. Pharmacy (Pharmaceutical Quality Assurance)

### Sem I

MQA101T	Modern pharmaceutical analytical technique	PO
MQA101T.01	Illustrate assay of single and multiple component	1 2 2 4 5 6 7 9 0 11
WiQiiioii.oi	Pharmaceuticals by using various analytical instruments.	1,2,3,4,5,6,7,8,9,11
MQA101T.02	Describe basic practical skills using Instrumentation	1,2,3,4,5,6,7,8,9,11
	techniques.	1,2,3,4,3,0,7,6,9,11
	Identify the theoretical knowledge on various	
MQA101T.03	instrumental techniques for analysis of organic	1,2,3,4,5,6,7,8,9,11
	substances.	
MQA101T.04	Explain the knowledge in developing new procedures for	1 2 2 4 5 6 7 9 0 11
	analysis.	1,2,3,4,5,6,7,8,9,11

MQA102T	Quality Management System	PO
MQA102T.01	Explain the importance of quality, tools for quality improvement and analysis of issues in quality.	1,2,3,4,5,6,7,8,9,11
MQA102T.02	Describe the quality evaluation of pharmaceuticals by different Quality Management systems.	1,2,3,4,5,6,7,8,9,11
MQA102T.03	Explain the stability testing of drug and drug substances and Statistical approaches for quality.	1,2,3,4,5,6,7,8,9,11
MQA102T.04	Explain the Regulatory compliance and Benchmarking for the Quality Management.	1,2,3,4,5,6,7,8,9,11

MQA103T	<b>Quality Control and Quality Assurance</b>	PO
MQA103T.01	Explain the concept of cGMP, GLP, ICH guidelines and non-clinical aspects in a pharmaceutical industry.	1,2,3,4,5,6,7,8,9,11
MQA103T.02	Explain the role of different regulatory affairs and the scope of quality certifications applicable to Pharmaceutical industries.	1,2,3,4,5,6,7,8,9,11
MQA103T.03	Relate the importance of documentation as per different Regulatory guidelines.	1,2,3,4,5,6,7,8,9,11
MQA103T.04	Explain the conditions of Manufacturing operations and controls for API and finished products.	1,2,3,4,5,6,7,8,9,11



<b>MQA 104T</b>	Product Development and Technology Transfer	PO
MQA104T.01	Describe the principles of drug discovery and drug approval process as per the regulatory guidelines.	1,2,3,4,5,6,7,8,9,11
MQA104T.02	Explain the preformulation studies in product development and application of packaging materials for the same.	1,2,3,4,5,6,7,8,9,11
MQA104T.03	Describe the significance design and layout of pilot plant scale up for various dosage forms.	1,2,3,4,5,6,7,8,9,11
MQA104T.04	Describe the technology transfer process from R & D to production.	1,2,3,4,5,6,7,8,9,11

MQA105P	Pharmaceutical Quality Assurance I Practical	PO
MQA105P.01	Illustrate the analysis of pharmacopoeial compounds and formulation using various instrumental techniques.	1,2,3,4,5,6,7,8,9,11
MQA105P.02	Assess case studies related to quality management system in pharmaceutical practices.	1,2,3,4,5,6,7,8,9,11
MQA105P.03	Summarize preformulation, In process quality control, stability, and packaging for different dosage forms.	1,2,3,4,5,6,7,8,9,11

### SEM II

MQA201T	Safety and hazards	PO
MQA201T.01	Explain the multidisciplinary nature of environmental	1,2,3,4,5,6,
	studies and concept, structure and function of an	7,8,9,10,11
	ecosystem.	
MQA201T.02	Illustrate the critical hazard management systems,	1,2,3,4,5,6,
	sources, types and prevention of air and fire.	7,8,9,10, 11
MQA201T.03	To plan the management and prevention of fire	1,2,3,4,5,6,
	explosion and types of chemical based hazards.	7,8,9,10, 11
MQA201T.04	To infer the rules and guidelines on risk assessment and	1,2,3,4,5,6,
	management.	7,8,9,10,11

MQA 202T	Pharmaceutical Vlidation	PO
MQA202T.01	Summarize the concepts of calibration, qualification specifications and validation types.	1,2,3,4,5,6,7,
		8,9,10, 11
MQA202T.02	Explain the Qualification of manufacturing, Laboratory equipments and Analytical instruments.	1,2,3,4,5,6,7,
		8,9,10,11
MQA202T.03	Explain the Concept, Process and documentation of Process validation and cleaning validation.	1,2,3,4,5,6,7,
		8,9,10,11
MQA202T.04	Illustrate the General Principles of Intellectual Property and Significance of transfer technology (TOT).	1,2,3,4,5,6,7,
		8,9,10,11



MQA203T	Audits & Regulatory Compliance	PO
MQA203T.01	Explain the roles and responsibility of audit system in pharmaceutical manufacturing.	1,2,5,6,7,8,9,11
MQA203T.02	Explain the auditing of vendors and production department.	1,2,5,6,7,8,9,11
MQA203T.03	Explain the auditing of microbiological laboratory.	1,2,5,6,7,8,9,11
MQA203T.04	Explain auditing of quality assurance and engineering department.	1,2,5,6,7,8,9,11

MQA204T	Pharmaceutical Manufacturing Technology	PO
MQA204T.01	Describe the pharmaceutical industry developments,	1,2,3,4,5,6,7,8,9,11
	plant layout and production planning.	
MQA204T.02	Explain the principles and practices of aseptic process	1,2,3,4,5,6,7,8,9,11
	technology.	
MQA204T.03	Explain the principles and practices of non sterile	1 2 2 4 5 6 7 9 0 11
	manufacturing technology and packaging technology.	1,2,3,4,5,6,7,8,9,11
MQA204T.04	Illustrate the principles and implementation of Quality	12245679011
	by design (QbD) and PAT.	1,2,3,4,5,6,7,8,9,11

MQA205P	Pharmaceutical Quality Assurance II Practical	PO
MQA205P.01	Assess different elements by using various instrumental analytical techniques.	1,2,3,4,5,6,7,8,9,11
MQA205P.02	Illustrate the validation of equipments and process.	1,2,3,4,5,6,7,8,9,11
MQA205P.03	Summarize case studies and check lists related to quality management system in pharmaceutical practices.	1,2,3,4,5,6,7,8,9,11

### Sem III

MRM301T	Research Methodology and Biostatistics	PO
MRM301T.01	Explain the concept of general &Medical Research and various methodologies and guidelines involved in research.	1,2,3,4,5,6,7,8,11
MRM301T.02	Apply Statistics and Statistical tests for significance and measure the various parametric and non-parametric tests.	1,2,3,4,5,6,7,8,11
MRM301T.03	Relate to the CPCSEA guidelines for laboratory animal facility and handling.	1,2,3,4,5,6,7,8,11
MRM301T.04	Articulate the basic principles of Helsinki in medical research and medical care.	1,2,3,4,5,6,7,8,11