Dr. D. Y. Patil Pratishthan's Dr. D. Y. Patil College of Pharmacy, Akurdi, Pune-411044

M. PHARM COURSE OUTCOMES

Course	Outcomes
	Department of Pharmaceutics
	Semester-I
	Semester-1
MPAT101T	MPAT101T (1) Explain principle, instrumentation of various
Modern	spectroscopic and chromatographic technique and their applications in
Pharmaceutical	Pharmaceutical research.
Analytical Techniques	MPAT101T (2) Interpret the spectrums and chromatogram of different
	methods of analysis.
	MPAT101T (3) Judge the research problems in Pharma. Analysis.
	MPAT101T (4) Examine and interpret the data obtained through
	experimentation and report the results as per regulatory requirements.
	MPAT101T (5) Utilize different analytical instruments for the assay of
MD11104/E	various APIs and formulations as per Pharmacopoeial standards.
MPH102T	MPH102T (1) Compare the concepts of development for novel drug
Drug Delivery System	delivery systems. MDH102T (2) Signify pathonbygiological conditions for development of
	MPH102T (2) Signify pathophysiological conditions for development of novel drug delivery systems.
	MPH102T (3) Justify the criteria for selection of drugs and polymers for
	the development of novel drug delivery systems.
	MPH102T (4) Evaluate novel drug delivery systems.
	MPH102T (5) Develop new technologies utilized for development of
	new dosage forms.
	MPH102T (6) Defend the recent developments for drug delivery, as per
	industry need.
MPH103T	MPH103T (1)State and perform various elements of preformulation
N.C. 1	studies.
Modern	MPH103T (2) Differentiate between the Compaction, compression and
Pharmaceutics	consolidation parameters
	MPH103T (3) Imbibe the Industrial Management and GMP
	Considerations
	MPH103T (4) Practice the optimization techniques and pilot plant Scale
	Up techniques.
	MPH103T (5) Validate and evaluate various processes, dosage forms
	and equipments.

	MPH103T (6) Estimate dissolution, diffusion and pharmacokinetic
	parameters from Pharmaceuticals point of view.
MPH104T Regulatory	MPH104T (1) Explain the importance of documentation in Pharma
Affair	industry with regulatory requirements for product approval process.
	MPH104T (2) Appraise post approval regulatory requirements for drug
	products with submission of global documents in CTD and ECTD
	formats for different countries.
	MPH104T (3) Propose the non-clinical drug development approvals for
	conducting clinical trials.
	MPH104T (4) Elaborate pharmacovigilance and process of monitoring in
	clinical trials
MPH105P	MPH105P (1) Elaborate the technique and apply skills in formulating
Pharmaceutics	dosage forms.
Practical I	MPH105P (2) Use factorial design technique to get the formulation of
	desired characteristics.
	MPH105P (3) Explain and apply the different types of analytical
	instrumental technique available for quality control of pharmaceuticals.
	(API's & Formulations).
	MPH105P (4) Interpret scientific data, represent the data in a tabular
	and/or graphical form.
	MPH105P (5) Demonstrate the effect of the physico-chemical properties
	on pharmaceutical systems
	Semester-II
MPH201T	MPH201T (1) Interpret the events, concepts and biological process in
	drug targeting.
Molecular	MPH201T (2) Differentiate, formulate and evaluate various novel
Pharmaceutics	targeted drug delivery systems.
	MPH201T (3) Summarize the process for Tumor targeting and brain
	specific delivery.
	MPH201T (4) Appraising the need of pulmonary system, differentiate,
	formulate and evaluate nasal drug delivery system.
	MPH201T (5) Summarize nucleic acid based therapeutic delivery
	system.
MPH202T	MPH202T (1) Compile the basic concepts in biopharmaceutics and
.,	pharmacokinetics with parameters that best describe process of drug
Advanced	absorption.
Biopharmaceutics &	MPH202T (2) Elaborate biopharmaceutics considerations in drug
Pharmacokinetics	product design and its performance.
	MPH202T (3) Develop pharmacokinetic models and parameters that best
	describe process of drug ADME.
	MPH202T (4) Defend use of Bioavailability and Bioequivalence studies
	for new drugs or dosage forms, as per industry need.
	for new drugs or dosage forms, as per mudstry need.

	MPH202T (5) Estimate the applications of pharmacokinetics and
MDHAAT	pharmacodynamics of biotechnology drugs. MDII202T (1) Discuss and correlate studies, which is inclusive of history.
MPH203T	MPH203T (1) Discuss and correlate studies, which is inclusive of history
Computer Aided	of computers, research and development and use of computers in research.
Drug Development	MPH203T (2) Elaborate Computational Modeling of Drug Disposition.
•	MPH203T (3) Predict the role of computers in formulation development
	with optimization techniques.
	MPH203T (4) Compile Computer-aided biopharmaceutical characterization.
	MPH203T (5) Discuss with peers Artificial Intelligence (AI) and Robotics.
MDH204T	
MPH204T	MPH204T (1) Describe the regulatory provisions related to the import
Cosmetic &	and manufacture of cosmetics as per the Drugs and Cosmetics Act 1940 and the Rules 1945.
Cosmeceuticals	MPH204T (2) Discuss key building blocks for various formulations.
	MPH204T (3) Explain the various problems related to the skin and hair.
	MPH204T (4) Design cosmeceuticals for various skin, hair and dental
	problems.
	MPH204T (5) Describe the guidelines for the regulation of herbal
	cosmetics used in private bodies.
	MPH204T (6) Design cosmetics and cosmeceuticals of desired safety,
	stability and efficacy with technologies involved in their manufacturing.
MPH205P	MPH205P(1) Elaborate the technique and apply skills in formulating
Pharmaceutics	dosage and cosmetic forms.
Practical-II	MPH205P(2) Apply software to get the formulation of desired
	characteristics.
	MPH205P(3) Interpret scientific data, represent the data in a tabular
	and/or graphical form.
	MPH205P(4) Interpret Pharmacokinetic and pharmacodynamics data
	perform data analysis using computer Softwares and give conclusion.
	MPH205P(5) Perform Pharmacokinetics studies <i>in-vitro</i> .
	MPH205P(6) Compare between marketed formulations for drug release.
Depa	rtment of Pharmaceutical Quality Assurance
	Semester-I
MPAT101T	MPAT101T(1) Explain principle, instrumentation of various
	spectroscopic and chromatographic technique and their applications in
	Pharmaceutical research.

Modern	MPAT101T(2)Interpret the spectrums and chromatogram of different
Pharmaceutical	methods of analysis.
Analytical Techniques	MPAT101T(3)Judge the research problems in Pharma. Analysis.
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	MPAT101T(4)Examine and interpret the data obtained through
	experimentation and report the results as per regulatory requirements.
	MPAT101T(5)Utilize different analytical instruments for the assay of
	various APIs and formulations as per Pharmacopoeial standards.
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MQA102T	MQA102T (1) Discuss the importance of quality.
Quality Management	MQA102T (2) Identify and use tools for quality improvement.
System	MQA102T (3) Locate, Evaluate and Analyse issues in quality.
System	MQA102T (4) Establish parameters for quality evaluation of
	pharmaceuticals.
	MQA102T (5) Prepare, Summarise and build methods for stability
	testing of drug and drug substances.
MO 4 102T	MQA102T (6) Compose statistical approaches for quality.
MQA103T	MQA103T (1) Compile responsibilities of QA and QC and discuss concept of Good Laboratory Practices (GLP), QSEM and CPCSEA
Quality Control and	guidelines.
Quality Assurance	MQA103T (2) Elaborate Current Good Manufacturing Practices
	(cGMP) in Pharmaceutical Industry.
	MQA103T (3) Evaluate raw materials, finished products, packaging
	materials and in process quality control tests for different Pharmaceutical
	formulations.
	MQA103T (4) Discuss importance of documentation and the scope of
	quality certifications applicable to Pharmaceutical Industries.
	MQA103T (5) Design manufacturing operations and packaging
	operations in Pharmaceutical Industry
MQA104T	MQA104T (1) Describe new product development process and
Product Development	informational content for Investigational new drug application.
and Technology	MQA104T (2) Discuss preformulation and its impact on product
Transfer	development.
	MQA104T (3) Sketch and discuss design layout of Pilot plant for
	different dosage forms. MQA104T (4) Explain the significance of Pharmaceutical dosage form
	and their packaging requirements.
	MQA104T (5) Explain necessary information to transfer technology
	from R&D to actual manufacturing by sorting out various information
	obtained during R&D.
MQA105P	MQA105P (1)Analyze drugs and formulations by using various
Pharmaceutical	sophisticated analytical instruments
rnarmaceuticai	sopinsucated analytical instruments

Quality Assurance Practical-I	MQA105P(2) Evaluate Quality control tests for tablet, capsules, parenterals and ointment.
	MQA105P(3) Determine the process capability, stability study protocol, accelerated stability studies.
	MQA105P(4) Determine solubility of drugs using surfactant systems and co-solvency method.
	Semester-II
MQA201T Hazards and Safety	MQA201T(1) Summarize multidisciplinary nature of environmental studies and various natural resources.
Management	MQA201T(2) Discuss concept, structure and function of an Ecosystem.
	MQA201T(3) Compile sources and types of Air based hazards.
	MQA201T(4) Adopt the types of chemical based hazards and their prevention.
	MQA201T(5) Select preventive measure and management system for fire and explosion.
	MQA201T(6) Elaborate rules and guidelines for risk assessment and management.
MQA202T Pharmaceutical	MQA202T (1) Summarize concepts of calibration, qualification and validation.
Validation	MQA202T (2) Evaluate qualification of manufacturing equipments and analytical instruments.
	MQA202T (3) Compile qualification of laboratory equipments and validation of utility system.
	MQA202T (4) Design process validation of different dosage forms and validate analytical method for estimation of different drugs.
	MQA202T (5) Elaborate cleaning validation of equipments and
	computer system validation. MQA202T (6) Compile concepts of Intellectual property, patents,
MQA203T	copyright, trademark and significance of Transfer Technology. MQA203T (1) Justify the importance of auditing, parameters involved,
	departments. MQA203T (2) Explain design and develop the methodology for pre auditing, auditing and post auditing of the facility

Audits and Regulatory Compliance MQA203T (3) Formulate the audit process, constitute the team requite to complete the process and assign the role to each member. MQA203T (4) Develop the auditing report, authorise the reprince investigate NCs, suggest and follow up compliance audited MQA203T (5) Prepare the check list for auditing the department signifying the importance of each component. MQA203T (6) Discuss and appraise a premaudited list which shall be base for auditing for GAMP. MQA204T (1) Describe the common practice in the pharmaceutic industry developments, plant layout and production planning. MQA204T(2) Discuss principles and practices of aseptic process technology. MQA204T(3) Explain production principles and practices of non stemanufacturing process technology. MQA204T(4) Elaborate the process of selection of Containers and process of selection o	ed
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manufacturing process technology.	ile
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Manual (4) Elaborate the process of selection of Containers	nd
closures for pharmaceuticals.	
MQA204T(5) Explain principles and implementation of Quality	hv
design (QbD) and process analytical technology (PAT) in pharmaceuti	•
manufacturing.	Jai
manuracturing.	
MQA205P (1) Analyse drugs and their formulations by varie	us
analytical instruments.	
Pharmaceutical MQA205P (2) Validate sterile and non-sterile dosage form by Proc	ess
Quality Assurance validation	
Practical II MQA205P (3) Measure qualification of Pharmaceutical equipments a	nd
Analytical instruments	
MQA205P (4) Validate equipment by cleaning validation method.	
Department of Pharmaceutical Chemistry	
Semester-I	
Defitester-1	
MPAT101T (1) Explain principle, instrumentation of variety	us
spectroscopic and chromatographic technique and their applications	in
Modern Pharmaceutical research.	
Pharmaceutical Applicational Teacheries and MPAT101T (2) Interpret the spectrums and chromatogram of different forms.	ent
Analytical Techniques methods of analysis.	
MPAT101T (3) Judge the research problems in Pharma. Analysis.	
MPAT101T (4) Examine and interpret the data obtained through	gh
experimentation and report the results as per regulatory requirements	
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	MPAT101T (5) Utilize different analytical instruments for the assay of
N CO	various APIs and formulations as per Pharmacopoeial standards.
MPC102T	MPC102T (1) Explain the different organic intermediates involved in
Advanced Organic	determining the reaction mechanism.
Chemistry – I	MPC102T (2) Elaborate the mechanism and applications of SN1, SN2,
	E1, E2 and various named reactions
	MPC102T (3) Discuss the applications of various synthetic reagents
	MPC102T (4) Explain the various protecting and de-protecting groups
	MPC102T (5) Describe the chemistry, synthesis and mechanism of
	reactions in heterocyclic compounds
	MPC102T (6) Compile the principle, process and applications of
MDC102T	Synthon approach and retrosynthesis.
MPC103T	MPC103T (1) Discuss various stages and techniques of drug discovery and their role in drug research
Advanced Medicinal	
Chemistry	MPC103T (2) Appraise the structural activity relationship and MOA of the important class of drugs and role of stereochemistry on drug action
	MPC103T (3) Apply peptidomimetics approach and applications
	MPC103T (4) Explain types of Enzyme inhibition and its application in
	medicine
	MPC103T (5) Elaborate prodrug and Analog development along with
	its applications
	MPC103T (6) Interpret techniques of combating drug resistance
MPC104T	MPC104T (1) Discuss the different types of natural compounds, their
112 01011	chemistry and medicinal importance.
Chemistry of Natural	MPC104T (2) Explain the importance of natural compounds as lead
Products	molecules for new drug discovery.
	MPC104T (3) Illustrate rDNA technology tool for new drug discovery.
	MPC104T (4) Elaborate chemistry and physiological significance of
	vitamins.
	MPC104T (5) Summarise general and advanced methods of structural
	elucidation of compounds of natural origin.
	MPC104T (6) Describe isolation, purification and characterization of
	simple chemical constituents from the natural source
MPC105P	MPC105P (1) Analyze Pharmacopoeial compounds by various
Dhawa a autia a	instrumental techniques
Pharmaceutical Chamistry Proctical I	MPC105P (2) Estimation of components by fluorimetry and flame
Chemistry Practical I	photometry
	MPC105P (3) Synthesize compounds based on rearrangement reaction
	MPC105P (4) Perform purification and characterization of medicinally
	important synthesized compounds
	MPC105P (5) Estimation of elements and functional groups in organic
	natural compounds

	MPC105P (6) Perform isolation and typical degradation reactions on plant constituents
	Semester-II
MPC201T	MPC201T (1) Correlate different analytical data using discriminate
Advanced Spectral	instruments.
Analysis	MPC201T (2) Annalyze and conclude the data of unknown structures.
•	MPC201T (3) Evaluate data of hyphenated instruments.MPC201T (4) Discuss structural elucidation of organic and natural
	compounds by IR, NMR and MASS spectral data
MPC202T	MPC202T (1) Describe the principles and applications of Green
WII C2021	chemistry
Advanced Organic	MPC202T (2) Illustrate the concept, principle and applications of
Chemistry –II	stereochemistry and asymmetric synthesis
	MPC202T (3) Explain the chemistry of peptide.
	MPC202T (4) Elaborate the concept of Photochemical reactions
	MPC202T (5) Discuss the principles of different types of pericyclic
	reactions.
	MPC202T (6) Explain the applications of various catalysis used in the
	reaction.
MPC203T	MPC203T (1) Predict and analyzed molecular properties of new
Computer Aided	molecules and explain various drug design methods
Drug Design	MPC203T (2) Elaborate the concept of pharmacophore mapping and
	Virtual Screening
	MPC203T (3) Discuss the Molecular Modeling and Docking technique.
	MPC203T (4) Assess the role of computer aided drug design in drug
	discovery MPC203T (5) Discuss history and development of QSAR
	MPC203T (6) Apply statistically QSAR based applications.
MPC204T	MPC204T(1) Develop synthetic routes that is safe, cost-effective,
Pharmaceutical	environmentally friendly, and efficient.
Process Chemistry	0.1.1.1.0.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1
J	MPC204T(2) Impart knowledge on the development and optimization
	of a synthetic route/s.
	MPC204T(3) Discuss pilot plant procedure for the manufacture of
	Active Pharmaceutical Ingredients and new chemical entities for the drug
	development phase.
	MPC204T(4) Create and carry out work up and separation procedure.
	MPC204T(5) Predict the outcome of organic reactions using a basic understanding of the general reactivity of functional groups and mechanism.

	MPC204T (6) Explain principles and applications of modern chemical instrumentation, experimental design, and data analysis.
MPC205P	MPC205P (1) Synthesize organic compounds by adapting different
WII C2031	
Pharmaceutical	approaches
Chemistry Practical II	MPC205P (2) Interpretation and identification of organic compounds
Chemistry Tractical II	by various analytical techniques
	MPC205P (3) Perform synthesis of organic compounds by various
	synthetic route as well as techniques
	MPC205P (4) Apply softwares for physicochemical and ADMET
	properties of drug molecules
	MPC205P (5) Perform experiments based on various computer aided
	drug design techniques
	MPC205P (6) Reflect the requirement of API through case studies as
	per regulatory guidelines
	Department of Pharmacology
	Semester-I
MPAT101T	MPAT101T (1) Explain principle, instrumentation of various
	spectroscopic and chromatographic technique and their applications in
Modern	Pharmaceutical research.
Pharmaceutical	MPAT101T (2) Interpret the spectrums and chromatogram of different
Analytical Techniques	methods of analysis.
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	MPAT101T (3) Judge the research problems in Pharma. Analysis.
	MPAT101T (4) Examine and interpret the data obtained through
	experimentation and report the results as per regulatory requirements.
	MPAT101T (5) Utilize different analytical instruments for the assay of
	various APIs and formulations as per Pharmacopoeial standards.
MPL102T	MPL102T (1) Predict Pharmacokinetic and Pharmacodynamics process
	of lipophilic and hydrophilic drugs.
Advanced	
Pharmacology-I	MPL102T (2) Differentiate pharmacological actions of drug acting on
	autonomic nervous system.
	MPL102T (3) Relate concept of drug action on central nervous system
	with its receptors.
	with its receptors.
	MPL102T (4) Describe mechanism and pharmacology of prototype
	drugs acting on CVS disorders and explain their clinical use.
	MPL102T (5) Illustrate feedback mechanism using mechanism and
	pharmacological action of hormones, autocoids and their antagonists.
MPL103T	MPL103T (1) Describe the various animals used in the drug discovery
1122 22200 1	process.
	process.

Pharmacological and	MPL103T (2) Explain good laboratory practices in maintenance and			
Toxicological	handling of experimental animals.			
Screening Methods-I	MPL103T (3) Appraise the regulations and ethical requirement for the			
	usage of experimental animals.			
	MPL103T (4) Discuss the various preclinical <i>in-vitro</i> , <i>in-vivo</i> and other			
	possible animal alternative models for the screening various			
	pharmacological activities.			
	MPL103T (5) Elaborate general principles and evaluation of			
	Immunoassay methods.			
MPL104T	MPL104T (1) Analyze the receptor signal transduction processes.			
Cellular and	MPL104T (2) Construct the molecular pathways affected by drugs.			
Molecular	MPL104T (3) Explain mechanisms and applicability of molecular			
Pharmacology	pharmacology, genomic and proteomic tools.			
i marmaeorogy	MPL104T (4) Distinguish the process of Pharmacogenomics.			
	MPL104T (5) Interpret the concept of Immunotherapeutic.			
	MPL104T (6) Interpret various Cell culture techniques.			
MPL105P	MPL105P (1) Demonstrate route of drug administration, blood			
Pharmacology	withdrawal techniques.			
Practical-I	MPL105P (2) Evaluate of effect of drug on CNS activity, analgesic			
	activity, Anti-diabetic activity, Anti-inflammatory activity, diuretic			
	activity, antiulcer activity etc.			
	MPL105P (3) Estimate DNA/RNA isolated from biological sample			
	using various techniques.			
	MPL105P (4) Demonstrate MTT assay, gel electrophoresis, gene			
	amplification and Protein quantification. MDI 105D (5) Explain the principle precedure and applications of			
	MPL105P (5) Explain the principle, procedure and applications of enzyme inhibiton activity, pharmacokinetics studies, apoptosis			
	enzyme inhibiton activity, pharmacokinetics studies, apoptosis determination etc.			
	MPL105P (6) Exhibit the extraction techniques of drug from biological			
	samples and their estimation using various analytical techniques.			
	Semester-II			
MPL201T	MPL201T (1) Illustrate feedback mechanism using mechanism and			
Advanced	pharmacological action of drug acting on endocrine systems.			
Pharmacology II	MPL201T (2) Relate concept of mechanism and resistance of drugs			
Thurmacology II	acting microbes, fungus, virus and tuberculosis.			
	acting interests, rangus, virus and tubercuresis.			
	MPL201T (3) Discuss the pharmacotherapy of COPD, Asthma,			
	constipation, diarrhea, Ulcer, inflammation, Rheumatoid Arthritis,			
	immune disorders etc.			
	MPL201T (4) Relate significance of rhythm, cycles and biological			
	clock for application of chronotherapy in various disease conditions.			
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	MPL201T (5) Explain antioxidant scavenging effects on free radicals				
	to cure diabetes, neurodegenerative diseases and cancer etc.				
	MPL201T (6) Conclude the recent advances in the treatment of				
	Alzhemier's disease, Parkinson's disease, cancer and diabetes mellitus.				
MPL202T	MPL202T (1) Summarise the various types of toxicity studies.				
Pharmacological and	MPL202T (2) Discuss the importance, ethical and regulatory				
Toxicological	requirements for various type of toxicity studies.				
Screening Methods–II	MPL202T (3) Outline the significance of reproductive toxicity,				
	teratogenicity, Genotoxicity, carcinogenicity studies.				
	MPL202T (4) Describe the significance of IND enabling studies and				
	safety pharmacology studies.				
	MPL202T (5) Recognize the importance and applications of				
	toxicokinetics studies and alternative methods to animal toxicity testing.				
	MPL202T (6) Demonstrate the practical skills require for conducting the				
1577 2027	preclinical toxicity studies.				
MPL203T	MPL203T (1) Illustrate various stages in modern drug discovery				
Principles of Drug	process.				
Discovery	MPL203T (2) Appraise role of genomics, proteomics and				
•	bioinformatics in drug discovery. MDI 202T (2) Discovery the different months do for lead identification.				
	MPL203T (3) Discuss the different methods for lead identification. MPL203T (4) Explain different approaches for rational drug design				
	MPL203T (4) Explain different approaches for rational drug design. MPL203T (5) Elaborate role of classical target and biomarker in drug				
	MPL203T (5) Elaborate role of classical target and biomarker in drudiscovery.				
	MPL203T (6) Illustrate role of <i>in-vitro</i> screening technique in drug				
	discovery				
MPL204T	MPL204T (1) Elaborate the basic concept of clinical research.				
Clinical Research and	MPL204T (2) Discuss regulatory requirements for conducting clinical				
Pharmacovigilance	trial.				
	MPL204T (3) Summarise the types of clinical trial designs.				
	MPL204T (4) Explain the responsibilities of key players involved in				
	clinical trials				
	MPL204T (5) Describe in detail about safety monitoring, reporting and				
	close-out activities.				
	MDI 204T (C) Illustrate the main duly of Direction				
	MPL204T (6) Illustrate the principles of Pharmacovigilance.				
MPL205P	MPL205P (1) Demonstrate the determination of unknown concentration				
	of sample by Bioassay method using chicken ilium preparation.				
	MPL205P (2) Illustrate the drug effect on rat BP, heart rate and ECG				
	using computer simulation techniques.				

Pharmacology	MPL205P (3) Study the acute oral toxicity, dermal toxicity, repeated
Practical II	dose toxicity studies and drug mutagenicity study as per OECD
	guidelines.
	MPL205P (4) Assess the ADR reporting and monitoring protocol.
	MPL205P (5) Design the protocol for Clinical trial studies.
	MPL205P (6) Evaluate the efficacy of drugs using In-silico studies like
	docking studies, Pharmacophore based screening and QSAR studies.