

(A State University Estb. by Govt. of Punjab vide Punjab Act No. 5 of 2015 and Approved u/s 2(f) & 12 (B) of UGC; Member AIU)

Department: Pharmaceutical Sciences & Technology

Program: M. Pharmacy

COs, POs, PSOs Mapping

Subject: Modern Pharmaceutical Analytical Techniques	Subject Code: MPH101T	Semester: 1 st
Credit: <u>4</u>	LTP <u>400</u>	Duration: <u>60 Hrs.</u>

COs	Statement	PO1	PO2	PO3	PO4	PO5	PO6
CO1	Chemicals and Excipients	1			1		1
CO2	The analysis of various drugs in single and combination dosage forms		1	1			1
CO3	Theoretical and practical skills of the instruments	1		1			1



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Department: Pharmaceutical Sciences & Technology

Program: <u>M. Pharmacy</u>

Subject: <u>Drug Delivery System</u>	Subject Code: MPH102T	Semester: 1 st
Credit: 4	LTP <u>400</u>	Duration: <u>60 Hrs.</u>

COs	Statement	PO1	PO2	PO3	PO4	PO5	PO6
CO1	The various approaches for development of novel drug delivery		2	1			1
	systems.						
CO2	The criteria for selection of drugs and polymers for the development of	1	2	1			1
	delivering system						
CO3	The formulation and evaluation of Novel drug delivery systems		2	1			1



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Department: Pharmaceutical Sciences & Technology

Program: M. Pharmacy

Subject: Modern Pharmaceutics	Subject Code: MPH103T	Semester: 1 st
Credit: <u>4</u>	LTP 400	Duration: <u>60 Hrs.</u>

COs	Statement	PO1	PO2	PO3	PO4	PO5	PO6
CO1	The elements of preformulation studies			1			1
CO2	The Active Pharmaceutical Ingredients and Generic drug Product Development	1		1			1
CO3	Industrial Management and GMP Considerations.	1	1				
CO4	Optimization Techniques & Pilot Plant Scale Up Techniques	2	1				
CO5	Stability Testing, sterilization process & packaging of dosage forms.	1		1			1



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Subject: Regulatory Affairs	Subject Code: MPH104T	Semester: 1 st
Credit: 4	LTP <u>400</u>	Duration: <u>60 Hrs.</u>

Cos	Statement	PO1	PO2	PO3	PO4	PO5	PO6
CO1	The Concepts of innovator and generic drugs, drug development Process	1	1				
CO2	The Regulatory guidance's and guidelines for filing and approval Process	1	1				
CO3	Preparation of Dossiers and their submission to regulatory agencies in different countries	1	1				
CO4	Post approval regulatory requirements for actives and drug products	1	1				
CO5	Submission of global documents in CTD/ eCTD formats	1	1				
CO6	Clinical trials requirements for approvals for conducting clinical trials	1	1				
CO7	Pharmacovigilence and process of monitoring in clinical trials.	1	1				



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Program: <u>M. Pharmacy</u>

Subject: Pharmaceutics Practical I	Subject Code: MHP105P	Semester: 1 st
Credit: <u>6</u>	LTP <u>006</u>	Duration: <u>180 Hrs.</u>

COs	Statement	PO1	PO2	PO3	PO4	PO5	PO6
CO1	The elements of performulation studies	1		1			1
CO2	Optimization Techniques & Dilot plant scale up techniques		1		1		1
CO3	The various approaches for development of novel drug delivery systems	1	1				1
CO4	The criteria for selection of drugs and polymers for the development of delivering system	1				1	1
CO5	The analysis of various drugs in single and combination dosage forms	2	1				1



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Program: M. Pharmacy

Subject: Molecular Pharmaceutics (Nano Tech and Targeted DDS)	Subject Code: MPH201T	Semester: 2 nd
Credit: <u>4</u>	LTP 400	Duration: <u>60 Hrs.</u>

COs	Statement	PO1	PO2	PO3	PO4	PO5	PO6
CO1	The various approaches for development of novel drug delivery		1		1		
	Systems						
CO2	The understanding of critical variables (material and process) for the development of novel drug/		1		1		
	gene delivery systems.						
CO3	The formulation, evaluation & application of novel drug/ gene delivery systems.		1		1		



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Program: M. Pharmacy

Subject: Advanced Biopharmaceutics and Pharmacokinetics	Subject Code: MPH202T	Semester: 2 nd
Credit: <u>4</u>	LTP <u>400</u>	Duration: <u>60 Hrs.</u>

COs	Statement	PO1	PO2	PO3	PO4	PO5	PO6
CO1	O1 The basic concepts in biopharmaceutics and pharmacokinetics		1				1
CO2	CO2 The use raw data and derive the pharmacokinetic models and parameters the best describe the process of drug absorption, distribution, metabolism and elimination		1				1
CO3	CO3 The critical evaluation of biopharmaceutic studies involving drug product equivalency		1				1
CO4	The design and evaluation of dosage regimens of the drugs using pharmacokinetic and biopharmaceutic parameters	1	1				1
CO5	The potential clinical pharmacokinetic problems and application of basics of pharmacokinetic	1	1				1



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Subject: Computer Aided Drug Development	Subject Code: MPL203T	Semester: 2 ND
Credit: <u>4</u>	LTP 400	Duration: <u>60 Hrs.</u>

COs	Statement	PO1	PO2	PO3	PO4	PO5	PO6
CO1	History of Computers in Pharmaceutical Research and Development		1				
CO2	Computational Modeling of Drug Disposition		1				
CO3	Computers in Preclinical Development		1				
CO4	Optimization Techniques in Pharmaceutical Formulation		1				
CO5	Computers in Market Analysis		1				
CO6	Computers in Clinical Development		1				
CO7	Artificial Intelligence (AI) and Robotics		1				
CO8	Computational fluid dynamics(CFD)		1				



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Subject: Cosmetics and Cosmeceuticals	Subject Code: MPH204T	Semester: 2 nd
Credit: <u>4</u>	LTP 400	Duration: <u>60 Hrs.</u>

COs	Statement	PO1	PO2	PO3	PO4	PO5	PO6
CO1	Key ingredients used in cosmetics and cosmeceuticals	1	1				1
CO2	Key building blocks for various formulations	1	1				1
CO3	Current technologies in the market	1	1				1
CO4	Various key ingredients and basic science to develop cosmetics and cosmeceuticals	1	1				1
CO5	Scientific knowledge to develop cosmetics and cosmeceuticals with desired Safety, stability, and efficacy.						



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Subject: Pharmaceutics Practicals - II	Subject Code: MPH 205P	Semester: 2 nd
Credit: <u>6</u>	LTP <u>006</u>	Duration: 180 Hrs.

COs	Statement	PO1	PO2	PO3	PO4	PO5	PO6
CO1	Various key ingredients and basic science to develop Novel drug delivery system.	1		1			1
CO2	Optimization Techniques in pharmaceutical formulation using factorial design		1		1		1
CO3	The use raw data and derive the pharmacokinetic models and parameters the best describe the process of during absorption, distribution, metabolism	1	1				1
CO4	The formulation and evaluation of novel drug delivery systems	1				1	1
CO5	Drafting of various pharmaceutical Process related documentation	2	1				1