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Syllabus-2023-2024

(SOP)(MPharm-PharmaCeutics)

Title of the Course	Regulatory Affairs
Course Code	MPH 104T

			Part A					
Veen	1	Compositor	4	Que dite	L	Т	Ρ	С
rear	TSL	Semester	ist	Creaits	4	0	0	4
Course Type	Theory only		•	•				
Course Category	Discipline Core							
Pre-Requisite/s				Co-Requisite/s				
Course Outcomes & Bloom's Level	CO1- Understan Understand) CO2- Know the CO3- Knowledg CO4- Know the CO5- Understan	nd the concepts of innovator and generic preparation of dossiers and their submiss e on post approval regulatory requiremer Regulatory guidance's and guidelines for nd the clinical trials requirements for appr	drugs, drug development process, re sion to regulatory agencies in differer nts for actives and drug products, sub r filing and approval process in differe ovals for conducting clinical trials, ph	gulatory guidance's and guidelines for filing nt countries (BL2-Understand) omission of global documents in CTD/ eCTD ent countries. (BL3-Apply) armacovigilance and process of monitoring	and app formats in clinica	oroval pro 6(BL2-Un al trials (B	ocess.(BL derstanc L3-Apply	2- 1) 1)
Coures Elements	Skill Developme Entrepreneurshi Employability ✓ Professsonal Ett Gender × Human Values > Environment ×	nt √ p X hics X K	SDG (Goals)	SDG4(Quality education)				

Part B

Modules	Contents	Pedagogy	Hours
UNIT-1	1. Documentation in Pharmaceutical industry: Master formula record, DMF (Drug Master File), distribution records. Generic drugs product development Introduction, Hatch- Waxman act and amendments, CFR (CODE OF FEDERAL REGULATION),drugproduct performance, in-vitro, ANDA regulatory approval process, NDA approval process, BE and drug product assessment, in –vivo, scale up process approval changes, post marketing surveillance, outsourcing BA and BE to CRO. b. Regulatory requirement for product approval: API, biologics, novel, therapies obtaining NDA, ANDA for generic drugs ways and means of US registration for foreign drugs	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	15
UNIT-2	2. CMC, post approval regulatory affairs. Regulation for combination products and medical devices. CTD and ECTD format, industry and FDA liaison. ICH - Guidelines of ICH-Q, S E, M. Regulatory requirements of EU, MHRA, TGA and ROW countries.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	15
UNIT-3	3.Nonclinical drug development: Global submission of IND, NDA, ANDA. Investigation of medicinal products dossier, dossier (IMPD) and investigator brochure (IB).	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	15
UNIT-4	4. Clinical trials: Developing clinical trial protocols. Institutional review board/ independent ethics committee Formulation and working procedures informed Consent process and procedures. HIPAA- new, requirement to clinicalstudy process, pharmacovigilance safety monitoring in clinical trials	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	15

Part C

Modules	Title	Indicative-ABCA/PBL/ Experiments/Field work/ Internships	Bloom's Level	Hours
UNIT-III	prepare regulatory guidelines for different countries	Seminar	BL3-Apply	10

Part D(Marks Distribution)

	Theory									
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation					
100	50	75	38	25	13					
Practical										
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation					
	0									

	Part E
Books	1. Generic Drug Product Development, Solid Oral Dosage forms, Leon Shargel and Isader Kaufer, Marcel Dekker series, Vol.143 2. The Pharmaceutical Regulatory Process, Second Edition Edited by Ira R. Berry and Robert P.Martin, Drugs and the Pharmaceutical Sciences, Vol.185, Informa Health care Publishers. 3. New Drug Approval Process: Accelerating Global Registrations By Richard A Guarino, MD,5th edition, Drugs and the Pharmaceutical Sciences, Vol.180, Informa Health Care Publishers. 3. New Drug Approval Process: Accelerating Global Registrations By Richard A Guarino, MD,5th edition, Drugs and the Pharmaceutical Sciences, Vol.190
Articles	https://regsci-ojs-tamu.tdl.org/regsci/
References Books	1. Guidebook fordrug regulatory submissions / Sandy Weinberg. By John Wiley & Sons.Inc. 2. FDA regulatory affairs: a guide for prescription drugs, medical devices, and biologics/edited By Douglas J. Pisano, David Mantus. 3. Clinical Trials and Human Research: A Practical Guide to Regulatory Compliance By Fay A.Rozovsky and Rodney K. Adams 4. www.ich.org/ 5. www.fda.gov/ 6. europa.eu/index_en.htm 7. https://www.tga.gov.au/tga- basics
MOOC Courses	https://onlinecourses.nptel.ac.in/
Videos	https://www.youtube.com/watch?v=xrZl8g70HoI&list=PLpGCFhhV_JSXuh8vFq4MwInuNj9f6hfRY

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COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	1	1	-	1	1	-	-	2	-	-	3	-	1	-	1
CO2	2	-	-	-	2	1	-	3	-	-	3	-	2	1	2
CO3	1	2	-	-	-	-	-	2	-	-	2	-	-	-	-
CO4	3	2	-	1	-	2	-	3	-	-	2	-	-1	2	-
CO5	2	1	-	-	1	1	-	1	-	-	2	-	-	2	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Modern Pharmaceutical Analytical Techniques
Course Code	MPH 101T

			Part A						
Voar	1et	Somostor	1st	Cradite	L	Т	Р	С	
i eai	151	Jeniestei	151	Offults	4	0	0	4	
Course Type	Theory only								
Course Category	Discipline Core								
Pre-Requisite/s				Co-Requisite/s					
Course Outcomes & Bloom's Level	CO1- Investigate CO2- Investigate CO3- Investigate CO4- Recognize CO5- Apprehene	 CO1- Investigate the pharmaceutical substance by absorption and emission techniques(BL2-Understand) CO2- Investigate the pharmaceutical substance by Nuclear Magnetic spectroscopy techniques.(BL3-Apply) CO3- Investigate the pharmaceutical substance by Mass spectroscopy techniques.(BL3-Apply) CO4- Recognize the principle, instrumentation and applications of different chromatographic techniques.(BL4-Analyze) CO5- Apprehend the fundamentals of immunological assays.(BL4-Analyze) 							
Coures Elements	Skill Developme Entrepreneurshi Employability X Professsonal Ett Gender X Human Values X Environment X	nt ✓ p X nics X K	SDG4(Quality education)						

Part B

Modules	Contents	Pedagogy	Hours
UNIT-1	a. UV-Visible spectroscopy: Introduction, Theory, Laws, Instrumentation associated with UV-Visible spectroscopy, Choice of solvents and solvent effect and Applications of UV- Visible spectroscopy. b. IR spectroscopy: Theory, Modes of Molecular vibrations, Sample handling, Instrumentation of Dispersive and Fourier - Transform IR Spectrometer, Factors affecting vibrational frequencies and Applications of IR spectroscopy c. Spectroflourimetry: Theory of Fluorescence, Factors affecting fluorescence, Quenchers, Instrumentation and Applications of fluorescence spectrophotometer	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	10
UNIT-2	NMR spectroscopy: Quantum numbers and their role in NMR, Principle, Instrumentation, Solvent requirement in NMR, Relaxation process, NMR signals in various compounds, Chemical shift, Factors influencing chemical shift, Spin-Spin coupling, Coupling constant, Nuclear magnetic double resonance, Brief outline of principles of FT-NMR and 13CNMR. Applications of NMR spectroscopy	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12
UNIT-3	Mass Spectroscopy: Principle, Theory, Instrumentation of Mass Spectroscopy, Different types of ionization like electron impact, chemical, field, FAB and MALDI, APCI, ESI, APPI Analyzers of Quadrupole and Time of Flight, Mass fragmentation and its rules, Meta stable ions, Isotopic peaks and Applications of Mass spectroscopy	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	8
UNIT-4	Chromatography: Principle, apparatus, instrumentation, chromatographic parameters, factors affecting resolution and applications of the following: a) Paper chromatography b) Thin Layer chromatography c) Ion exchange chromatography d) Column chromatography e) Gas chromatography f) High Performance Liquid chromatography g) Affinity chromatography	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12
UNIT-5	a. Electrophoresis: Principle, Instrumentation, working conditions, factors affecting separation and applications of the following: a) Paper electrophoresis b) Gel electrophoresis c) Capillary electrophoresis d) Zone electrophoresis e) Moving boundary electrophoresis f) Iso electric focusing b. X ray Crystallography: Production of X rays, Different X ray diffraction methods, Bragg's law, Rotating crystal technique, X ray powder technique, Types of crystals and applications of X- ray diffraction.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12
UNIT-6	Immunological assays: RIA (Radio immunoassay), ELISA, Bioluminescence assays.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	8

Part C

Modules	Title	Indicative-ABCA/PBL/ Experiments/Field work/ Internships	Bloom's Level	Hours
UNIT-4	High Performance Liquid chromatography	PBL	BL4-Analyze	10

Part D(Marks Distribution)

	Theory								
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation				
100	50	75	38	25	13				
Practical									
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation				
	0								

	Part E
Books	Spectrometric Identification of Organic compounds - Robert M Silverstein, Sixth edition, John Wiley & Sons, 2004. 2. Principles of Instrumental Analysis - Doglas A Skoog, F. James Holler, Timothy A. Niemar Pharmaceutical Chemistry – Beckett and Stenlake, Vol II, 4th edition, CBS Publishers, New Delhi, 1997.
Articles	https://www.sciencedirect.com/journal/journal-of-pharmaceutical-analysis
References Books	1. Organic Spectroscopy - William Kemp, 3rd edition, ELBS, 1991. 2. Quantitative Analysis of Drugs in Pharmaceutical formulation - P D Sethi, 3rd Edition, CBS Publishers, New Delhi, 1997. 3. Pharmaceutic
MOOC Courses	https://onlinecourses.nptel.ac.in/
Videos	https://www.udemy.com/course/modern-analytical-techniques/?utm_source=adwords- pmax&utm_medium=udemyads&utm_campaign=PMax_la.EN_cc.INDIA&utm_content=deal4584&utm_term=agkwadde_cdmpltili_1007795pd&gad_source=2&gclid=(

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COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	3	1	3	-	3	-	-	-	-	-	3	-	1	-	1
CO2	3	2	3	-	2	1	-	-	-	-	3	-	2	1	2
CO3	3	1	1	-	1	2	-	-	-	-	3	-	2	-	1
CO4	3	1	1	2	2	2	-	-	-	-	3	-	1	2	2
CO5	2	1	2	-	2	2	-	-	-	-	3	-	-	1	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Drug Delivery System
Course Code	MPH 102T

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ſ	Voar	1et	Somostor	1et	Credite	L	Т	Р	С
	16di	131	Cemester	131	oreans	4	0	0	4
	Course Type	Theory only							
	Course Category	Discipline Cor	e						
	Pre-Requisite/s				Co-Requisite/s				
	Course Outcomes & Bloom's Level	CO1- Explain therapeutic eff its desired the CO2- Know va CO3- Ability to administration CO4- Apply la CO5- Create r	 C01- Explain drug delivery systems which give detailed information on transporting a pharmaceutical compound in the body as needed to safely achieve its desired herapeutic effect. Also about the approaches, formulations, technologies, and systems for transporting a pharmaceutical compound in the body as needed to safely achieve ts desired therapeutic effect with suitable drug delivery (BL2-Understand) C02- Know vaccine delivery and different mode of application approach for clinical use. (BL2-Understand) C03- Ability to communicate different types of Drug carrier used in the process of drug delivery which serves to improve the selectivity, effectiveness, and/or safety of drug administration(BL3-Apply) C04- Apply latest drug delivery knowledge and think to develop new formulation based on the individual requirement(BL3-Apply) C05- Create recent developments in protein and peptide for parenteral delivery approaches will give new dimension of drug deliver for antibiotics, insulin, etc(BL4-Analyze) 						
	Coures Elements	Skill Developn Entrepreneurs Employability Professsonal I Gender X Human Values Environment X	nent ✓ hip X ✓ Ethics X S X K	SDG (Goals)	SDG1(No poverty) SDG3(Good health and well-being) SDG4(Quality education) SDG17(Partnerships for the goals)				

Part B

Modules	Contents	Pedagogy	Hours
UNIT-1	1.Sustained Release (SR) and Controlled Release (CR) formulations: Introduction & basic concepts, advantages/ disadvantages, factors influencing, Physicochemical & biological approaches for SR/CR formulation, Mechanism of Drug Delivery from SR/CR formulation. Polymers: introduction, definition, classification, properties and application Dosage Forms for Personalized Medicine: Introduction, Definition, Pharmacogenetics, Categories of Patients for Personalized Medicines: Customized drug delivery systems, Bioelectronic Medicines, 3D printing of pharmaceuticals, Telepharmacy	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12
UNIT-2	2. Rate Controlled Drug Delivery Systems: Principles & Fundamentals, Types, Activation; Modulated Drug Delivery Systems; Mechanically activated, pH activated, Enzyme activated, and Osmotic activated Drug Delivery Systems Feedback regulated Drug Delivery Systems; Principles & Fundamentals.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	10
UNIT-3	3. Gastro-Retentive Drug Delivery Systems: Principle, concepts advantages and disadvantages, Modulation of GI transit time approaches to extend GI transit. Buccal Drug Delivery Systems: Principle of muco-adhesion, advantages and disadvantages, Mechanism of drug permeation, Methods of formulation and its evaluations	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	6
UNIT-4	4. Ocular Drug Delivery Systems: Barriers of drug permeation, Methods to overcome barriers. Rate Controlled Drug Delivery Systems: Principles & Fundamentals, Types, Activation; Modulated Drug Delivery Systems; Mechanically activated, pH activated, Enzyme activated, and Osmotic activated Drug Delivery Systems Feedback regulated Drug Delivery Systems; Principles & Fundamentals.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	6
UNIT-5	5, Gastro-Retentive Drug Delivery Systems: Principle, concepts advantages and disadvantages, Modulation of GI transit time approaches to extend GI transit	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	6
UNIT-6	6. Buccal Drug Delivery Systems: Principle of muco-adhesion, advantages and disadvantages, Mechanism of drug permeation, Methods of formulation and its evaluations	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	6
UNIT-7	7.Ocular Drug Delivery Systems: Barriers of drug permeation, Methods to overcome barriers	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board/PPT	4
UNIT-8	8. Transdermal Drug Delivery Systems: Structure of skin and barriers, Penetration enhancers, Transdermal Drug Delivery Systems, Formulation and evaluation	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	3

Modules	Title	Indicative-ABCA/PBL/ Experiments/Field work/ Internships	Bloom's Level	Hours
UNIT-1	3D PRINTING TECHNOLOGY	PBL	BL4-Analyze	10

	Part D(Marks Distribution)							
	Theory							
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation			
100	50	75	38	25	13			
			Practical					
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation			
	0							

Part E

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Books	1.N.K. Jain, Controlled and Novel Drug Delivery, CBS Publishers & Distributors, New Delhi, First edition 1997 (reprint in 2001). 2. S. P. Vya s and R. K. Khar, Controlled Drug Delivery- conceptsand advances, Vallabh Prakashan, New Delhi, First edition 2002
Articles	2.Indian Journal of Pharmaceutical Sciences (IPA) 3. Indian drugs (IDMA) 4. Journal of controlled release (Elsevier Sciences) desirable 4. Drug Development and Industrial Pharmacy (Marcel & Decker) desirable
References Books	5. Y W. Chien, Novel Drug Delivery Systems, 2nd edition, revised and expanded, Marcel Dekker, Inc., New York, 1992. 2. Robinson, J. R., Lee V. H. L, Controlled Drug Delivery Systems, Marcel Dekker, Inc., New York, 1992. 3. Encyclopedia of controlled delivery, Editor- Edith Mathiowitz, Published by Wiley Interscience Publication, John Wiley and Sons, Inc, New York! Chichester/Weinheim
MOOC Courses	https://onlinecourses.nptel.ac.in/
Videos	https://www.youtube.com/watch?v=1Jk08tf1Gh8

	Course Articulation Matrix														
COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	3	-	3	-	3	1	-	-	-	-	3	-	1	-	1
CO2	3	-	2	-	1	2	-	-	-	-	3	-	2	1	-
CO3	3	1	3	-	2	-	-	-	-	-	3	-	-	-	2
CO4	3	-	1	-	2	1	-	-	-	-	3	-	-	1	2
CO5	2	2	3	1	1	-	-	-	-	-	3	-	1	2	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Modern Pharmaceutics
Course Code	MPH 103T

			Part A						
Yoar	1-t		1st	Cradita	L	т	Р	С	
Tear	150	Semester	150	Cleans	4	0	0	4	
Course Type	Theory only								
Course Category	Discipline Cor	sipline Core							
Pre-Requisite/s				Co-Requisite/s					
Course Outcomes & Bloom's Level	CO1- Apply th CO2- Recogni CO3- Describe CO4- Analyze CO5- Understa	 CO1- Apply the concepts of preformulation and optimization techniques during formulation development. (BL3-Apply) CO2- Recognize the importance of validation of methods, equipments and processes during pharmaceutical manufacturing (BL2-Understand) CO3- Describe current good manufacturing practices guidelines and industrial management (BL2-Understand) CO4- Analyze the importance of tablet compression and compaction studies. (BL4-Analyze) CO5- Understand different consolidation parameters employed in formulation development and evaluation. (BL5-Evaluate) 							
Coures Elements	Skill Developn Entrepreneurs Employability Professsonal I Gender X Human Values Environment X	nent ✓ hip × ✓ Ethics ×	SDG (Goals)	SDG1(No poverty) SDG3(Good health and well-being) SDG4(Quality education) SDG17(Partnerships for the goals)					

Part B

Modules	Contents	Pedagogy	Hours
UNIT-I	a. Preformation Concepts – Drug Excipient interactions - different methods, kinetics of stability, Stability testing. Theories of dispersion and pharmaceutical Dispersion (Emulsion and Suspension, SMEDDS) preparation and stability Large and small volume parental – physiological and formulation consideration, Manufacturing and evaluation. b. Optimization techniques in Pharmaceutical Formulation: Concept and parameters of optimization, Optimization techniques in pharmaceutical formulation and processing. Statistical design, Response surface method, Contour designs, Factorial designs and application in formulation	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	14
UNIT-II	2. Validation: Introduction to Pharmaceutical Validation, Scope & merits of Validation, Validation and calibration of Master plan, ICH & WHO guidelines for calibration and validation of equipments, Validation of specific dosage form, Types of validation. Government regulation, Manufacturing Process Model, URS, DQ, IQ, OQ & P.Q. of facilities.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	14
UNIT-III	3. cGMP & Industrial Management: Objectives and policies of current good manufacturing practices, layout of buildings, services, equipments and their maintenance Production management: Production organization, , materials management, handling and transportation, inventory management and control, production and planning control, Sales forecasting, budget and cost control, industrial and personal relationship. Concept of Total Quality Management.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12
UNIT-IV	4. Compression and compaction: Physics of tablet compression, compression, consolidation, effect of friction, distribution of forces, compaction profiles. Solubility	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	8
UNIT-V	5. Study of consolidation parameters; Diffusion parameters, Dissolution parameters and Pharmacokinetic parameters, Heckel plots, Similarity factors – f2 and f1, Higuchi and Peppas plot, Linearity Concept of significance, Standard deviation, Chi square test, students T-test, ANOVA test.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12

Part C

Modules	Title	Indicative-ABCA/PBL/ Experiments/Field work/ Internships	Bloom's Level	Hours
UNIT-V	study of diffussion and dissolution parameters	PBL	BL4-Analyze	10

Part D(Marks Distribution)								
Theory								
Total Marks	Total Marks Minimum Passing Marks External Evaluation Min. External Evaluation Internal Evaluation Min. Internal Evaluation							
100	50	75	38	25	13			
			Practical					
Total Marks Minimum Passing Marks		External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation			

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Books	1. Theory and Practice of Industrial Pharmacy By Lachmann and Libermann 2. Pharmaceutical dosage forms: Tablets Vol. 1-3 by Leon Lachmann. 3. Pharmaceutical Dosage forms: Disperse systems, Vol, 1-2; By Leon Lachmann. 4. Pharmaceutical Dosage forms: Parenteral medications Vol. 1-2; By Leon Lachmann. 5. Modern Pharmaceutics; By Gillbert and S. Banker. 6. Remington's Pharmaceutical Sciences
Articles	https://www.ijmpronline.com/
References Books	7.Advances in Pharmaceutical Sciences Vol. 1-5; By H.S. Bean & A.H. Beckett. 8. Physical Pharmacy; By Alfred martin 9. Bentley's Textbook of Pharmaceutics – by Rawlins. 10. Good manufacturing practices for Pharmaceuticals: Aplan for total quality control, Second edition; By Sidney H. Willig. 11. Quality Assurance Guide; By Organization of Pharmaceutical producers of India. 12.Drug formulation manual; By D.P.S. Kohli and D.H.Shah. Eastern publishers, New Delhi. 13.How to practice GMPs; By P. P. Sharma. Vandhana Publications, Agra. 12. Pharmaceutical Process Validation; By Fra. R. Berry and Robert A. Nash. 13.Pharmaceutical Preformulations; By J.J. Wells. 14. Applied production and operations management; By Evans, Anderson, Sweeney and Williams. 15. Encyclopedia of Pharmaceutical technology, Vol I – III
MOOC Courses	https://swayam.gov.in/nc_details/NPTEL
Videos	https://www.youtube.com/watch?v=mRJvss9bMVc&list=PL0o-kamDFTumhseOKF-6OCrHdhBySNkAA

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COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	1	-	3	-	1	-	-	-	-	-	3	-	1	-	1
CO2	3	1	1	-	2	-	-	-	-	-	3	-	-	1	-
CO3	2	2	2	1	2	-	-	-	-	-	3	-	2	2	2
CO4	2	1	1	-	2	-	-	-	-	-	3	-	1	-	2
CO5	2	2	2	2	-	1	-	-	-	-	3	-	1	-	2
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Pharmaceutics Practical I
Course Code	MPH 105P

Part A Т Ρ С L Credits Year 1st Semester 1st 0 0 6 6 Course Type Lab only Discipline Core **Course Category** Co-Requisite/s Pre-Requisite/s CO1- Chemicals and Excipients(BL4-Analyze) CO2- The analysis of various drugs in single and combination dosage forms(BL5-Evaluate) CO3- Theoretical and practical skills of the instruments(BL3-Apply) **Course Outcomes** & Bloom's Level Skill Development 🗸 Entrepreneurship \checkmark SDG1(No poverty) SDG3(Good health and well-being) SDG4(Quality education) SDG17(Partnerships for the goals) Employability 🗸 **Coures Elements** Professsonal Ethics \mathbf{X} SDG (Goals) Gender \mathbf{X} Human Values old XEnvironment old X

Part B

Modules	Contents	Pedagogy	Hours
UNIT-1	1.Sustained Release (SR) and Controlled Release (CR) formulations: Introduction & basic concepts, advantages/ disadvantages, factors influencing, Physicochemical & biological approaches for SR/CR formulation, Mechanism of Drug Delivery from SR/CR formulation. Polymers: introduction, definition, classification, properties and application Dosage Forms for Personalized Medicine: Introduction, Definition, Pharmacogenetics, Categories of Patients for Personalized Medicines: Customized drug delivery systems, Bioelectronic Medicines, 3D printing of pharmaceuticals, Telepharmacy	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12

	Par	t C		
Modules	Title	Indicative-ABCA/PBL/ Experiments/Field work/ Internships	Bloom's Level	Hours
1	1. Analysis of pharmacopoeial compounds and their formulations by UV Vis spectrophotometer 2. Simultaneousestimation ofmulti component containing formulations by UV spectrophotometry	Experiments	BL4-Analyze	8
2	3. Experiments based on HPLC 4. Experiments based on Gas Chromatography	Experiments	BL3-Apply	8
3	5. Estimation of riboflavin/quinine sulphate by fluorimetry 6. Estimation of sodium/potassium by flame photometry	Experiments	BL4-Analyze	8
4	7. To perform In-vitro dissolution profile of CR/ SR marketed formulation 8. Formulation and evaluation of sustained release matrix tablets	Experiments	BL5-Evaluate	8
5	9. Formulation and evaluation osmotically controlledDDS 10. Preparation and evaluation of Floating DDS- hydro dynamically balanced DDS	Experiments	BL6-Create	8
6	11. Formulation and evaluation of Muco adhesive tablets. 12. Formulation and evaluation of trans dermal patches	Experiments	BL6-Create	8
7	13. To carry out preformulation studies of tablets. 14. To study the effect of compressional force on tablets disintegration time.	Experiments	BL3-Apply	8
8	15. To study Micromeritic properties of powders and granulation. 16. To study the effect of particle size on dissolution of a tablet	Experiments	BL3-Apply	8

Part D(Marks Distribution)

			Theory					
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation			
	Practical							
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation			
150	0	100	50	50	25			

	Part E						
Books	PRACTICAL MANUAL						
Articles	JOURNALS						
References Books	LAB MANUAL						
MOOC Courses	SWAYAM NPTEL						
Videos	YOUTUBE						

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COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	1	1	3	1	1	-	-	-	-	-	3	-	1	-	1
CO2	3	1	2	2	2	-	-	-	-	-	3	-	2	1	2
CO3	3	2	3	-	1	-	-	-	-	-	3	-	3	-	2
CO4	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO5	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Assignments
Course Code	MPH 106A

Part A									
Voar	1ct Semanter		1et	Cradite	L	Т	Р	С	
i eai	150	Semester		oreans	7	0	0	7	
Course Type	Project								
Course Category	Discipline Core								
Pre-Requisite/s				Co-Requisite/s					
Course Outcomes & Bloom's Level	CO1- To understand modern pharmaceutical techniques(BL4-Analyze) CO2- To understand various pharmaceutical formulations() CO3- To understand the IPR in pharmacy()								
Coures Elements	Skill Developme Entrepreneurshi Employability ✓ Professsonal Ett Gender × Human Values > Environment ×	nt √ p √ hics X K	SDG (Goals)	SDG4(Quality education)					

Part	В
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Modules	Contents	Pedagogy	Hours
1	Different Analytical techniques	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	7
2	Various aprroaches for development of NDDS	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	7
3	Optimization techniques and pilot plant scale up	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	7
4	write about various Pharmaceutical regulatory bodies across the globe	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	

	Part C										
Modules	Title	Indicative-ABCA/PBL/ Experiments/Field work/ Internships	Bloom's Level	Hours							
1	1. Analysis of pharmacopoeial compounds and their formulations by UV Vis spectrophotometer 2. Simultaneousestimation ofmulti component containing formulations by UV spectrophotometry	Experiments	BL4-Analyze	8							

Theory									
Total Marks	Total Marks Minimum Passing Marks External Evaluation Min. External Evaluation Internal Evaluation Min. Internal Evaluation								
			Practical						
Total Marks Minimum Passing Marks External Evaluation Min. External Evaluation Internal Evaluation Min. Internal Evaluation									
100	50	75	38	25	13				

Part E						
Books	REFERENCE BOOKS					
Articles	INTERNET SOURCES					
References Books	REFERENCE BOOKS					
MOOC Courses	SWAYAM					
Videos	YOUTUBE					

	Course Articulation Matrix														
COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	1	1	1	-	-	-	-	-	-	-	1	-	1	-	1

Part F

CO2	2	2	2	-	-	-	-	-	-	-	1	-	-	2	-
CO3	1	1	-	-	-	-	1	-	-	-	1	-	-	-	1
CO4	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO5	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Online Certificate Course
Course Code	MPH 107ET

			Part A						
Yoar	1et	Semester	1et	Cradits	L	Т	Р	С	
Teal	150	Semester	151	Oreans	1	0	0	1	
Course Type	Online course								
Course Category	Skill Enhanceme	xill Enhancement Courses							
Pre-Requisite/s	ELECTIVES			Co-Requisite/s					
Course Outcomes & Bloom's Level	CO1- TO INCRE	CO1- TO INCREASE CREATIVE AND TECHNICAL SKILLS(BL4-Analyze)							
Coures Elements	Skill Developme Entrepreneurshi Employability ✓ Professsonal Ett Gender X Human Values > Environment X	nt ✔ p X hics X K	SDG (Goals)	SDG4(Quality education)					

Part B								
Modules Contents Pedagogy Ho								
UNIT	REGULATORY AFFAIRS	ONLINE COURSES	10					

Part D(Marks Distribution)								
Theory								
Total Marks Minimum Passing Marks External Evaluation Min. External Evaluation Internal Evaluation Min. Internal Evaluation								
50	25	25	12	25	13			
			Practical					
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation			
	0							

Part E							
Books	NA						
Articles	NA						
References Books	NA						
MOOC Courses	1.https://www.coursera.org/courses?query=regulatory%20affairs 2. https://www.igmpi.ac.in/RAprograms.html						
Videos	NA						

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	3	1	1	-	-	-	-	3	-	-	3	-	3	1	2
CO2	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO3	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO4	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO5	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-

Course Articulation Matrix



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Good Manufacturing in Pharma
Course Code	MPH 108ET

			Part A	Λ				
Voar	1 ct	Somostor	1 st	Cradita	L	Т	Р	С
Tear	151	Semester		3	1	0	4	
Course Type	Theory only	,						
Course Category	Discipline S	pecific Elective						
Pre-Requisite/s				Co-Requisite/s				
Course Outcomes & Bloom's Level	CO1- Mean CO2- Gene CO3- The ro CO4- To ap CO5- To lea	CO1- Meaning and importance of GMP in the manufacturing and pharmaceutical industries.(BL1-Remember) CO2- General and specific requirements for documentation and records(BL2-Understand) CO3- The role of Production, Quality Control (QC), Quality Assurance (QA) and the Qualified Person (QP) in GMP(BL3-Apply) CO4- To apply the GMP certification in industry (BL3-Apply) CO5- To learn the documentation and GMP SOPs (BL3-Apply)						
Coures Elements	Skill Develo Entreprene Employabili Professson Gender ✓ Human Valu Environmer	opment ✓ urship ✓ ty ✓ al Ethics ✓ ues ✓ tt ✓	SDG (Goals)	SDG1(No poverty) SDG3(Good health and well-being) SDG4(Quality education) SDG6(Clean water and sanitation) SDG8(Decent work and economic growth) SDG12(Responsible consuption and produ SDG17(Partnerships for the goals)	uction)			

Part B

Modules	Contents	Pedagogy	Hours
UNIT 1	Introduction What is Good Manufacturing Practice? Why is GMP important? Official GMP Directives. the basic requirements of Good Manufacturing Practice., Pharmaceutical Quality System Principle and overview of the Pharmaceutical Quality System. Major updates. Development, content and implementation of PQS.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	8
UNIT 2	Personnel Key personnel. Background and duties of the Qualified person. Duties of the Head of production department. Duties of the Head of quality control. Person releasing the batch. Consultants. Personnel training and hygiene, Premises and Equipment Production area. Storage area. Quality control areas. Ancillary areas. Equipment.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	8
UNIT 3	Documentation Premises. Generation and control of documentation. Types of documents and specifications. Manufacturing formula and processing instructions. Packaging instructions. Procedures and records., Production General principles. Prevention of cross-contamination in production. Guidelines for starting materials. Processing operations. Packaging materials and operations. Guidelines for finished products.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	8
UNIT4	Quality Control General principles. Main tasks of the Quality control department. Technical transfer of testing methods. Transfer protocol., Complaints and Recalls GMP Guidelines related to complaints. Classification of defects. Product Recalls.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	8

Part C

Modules	Title	Indicative-ABCA/PBL/ Experiments/Field work/ Internships	Bloom's Level	Hours
1	GMP	Seminar	BL3-Apply	2

	Part D(Marks Distribution)						
			Theory				
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation		
100	50	75	38		13		
			Practical				
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation		

Books	Karmacharya JB. Good manufacturing practices (GMP) for medicinal products. Promising Pharmaceuticals. 2014;101.
Articles	Patel KT, Chotai NP. Pharmaceutical GMP: past, present, and future-a review. Die Pharmazie-An International Journal of Pharmaceutical Sciences. 2008 Apr 1;63(4):251-5.
References Books	Durivage MA, editor. The Certified Pharmaceutical GMP Professional Handbook. Quality Press; 2016 May 23.
MOOC Courses	UDEMY, COURSERA, PHARMASTATE ACADEMY
Videos	You tube

Part F

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	3	-	-	-	1	-	1	1	2	-	3	-	1	1	1
CO2	1	-	-	1	1	-	1	1	2	-	3	-	1	1	1
CO3	1	-	-	-	1	-	2	1	2	-	3	-	2	1	1
CO4	2	-	-	1	1	-	1	2	2	-	2	-	1	-	3
CO5	1	-	-	1	1	-	2	2	2	-	2	-	-	-	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-

Course Articulation Matrix

6/13/24, 2:09 PM

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Syllabus-2023-2024

(SOP)(MPharm-PharmaCeutics)

Title of the Course	Molecular Pharmaceutics (Nano Tech and Targeted DDS)
Course Code	MPH 201T

		Part A						
Year	1st	Semester	2nd	Credits	L	Т	Ρ	С
					4	0	0	4
Course Type	Theory only							
Course Category	Discipline Core							
Pre-Requisite/s	Upon completion of novel drug delivery The formulation and	the course student shall be able to understand T systems. • The criteria for selection of drugs and d evaluation of novel drug delivery systems	he various approaches for development of polymers for the development of NTDS	Co-Requisite/s				
Course Outcomes & Bloom's Level	CO1- Recall the bas CO2- Describe met CO3- Outline the pr CO4- Discuss the d CO5- Choose comp	sic aspects and approaches of targeted drug deliv hods for the preparation and evaluation of polyme eparation methods and applications of monoclona ifferent aspects of pulmonary drug delivery syster ponents of nucleic acid based therapeutic delivery	very systems (BL2-Understand) eric nanoparticles and liposome's (BL3-Appi al antibodies and vesicular nanocarriers. (BL ns (BL4-Analyze) systems (BL5-Evaluate)	y) 3-Apply)				
Coures Elements	Skill Development Entrepreneurship Employability Professsonal Ethics Gender Human Values Environment	×	SDG (Goals) SDG3(Good health and well-being)					

Part B

Modules	Contents	Pedagogy	Hours
UNIT-1	Targeted Drug Delivery Systems: Concepts, Events and biological process involved in drug targeting. Tumor targeting and Brain specific delivery.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12
UNIT-2	Targeting Methods: introduction preparation and evaluation. Nano Particles & Liposomes: Types, preparation and evaluation.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12
UNIT-3	Micro Capsules / Micro Spheres: Types, preparation and evaluation, Monoclonal Antibodies; preparation and application, preparation and application of Niosomes, Aquasomes, Phytosomes, Electrosomes	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12
UNIT-4	Pulmonary Drug Delivery Systems: Aerosols, propellants, Containers Types, preparation and evaluation, Intra Nasal Route Delivery systems; Types, preparation and evaluation	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12
UNIT-5	5 Nucleic acid based therapeutic delivery system: Gene therapy, introduction (ex-vivo & in- vivo gene therapy). Potential target diseases for gene therapy (inherited disorder and cancer). Gene expression systems (viral and nonviral gene transfer). Liposomal gene delivery systems. Biodistribution and Pharmacokinetics. knowledge of therapeutic antisense molecules and aptamers as drugs of future.	white board/ppt	12

	Part C								
Modules	Title	Indicative-ABCA/PBL/ Experiments/Field work/ Internships	Bloom's Level	Hours					
UNIT-2	PREPARATION OF NANOSOMES	PBL	BL4-Analyze	8					

	Part D(Marks Distribution)								
	Theory								
Total Marks	Total Marks Minimum Passing Marks External Evaluation Min. External Evaluation Internal Evaluation Min. Internal Evaluation								
100	50 75		38 25		13				
			Practical						
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation				
	0								

Books	Y W. Chien, Novel Drug Delivery Systems, 2nd edition, revised and expanded, Marcel Dekker, Inc., New York, 1992. 2. S. P. Vyas and R. K. Khar, Controlled Drug Delivery - concepts and advances, Vallabh Prakashan, New Delhi, First edition 2002. N.K. Jain, Controlled and Novel Drug Delivery, CBS Publishers & Distributors, New Delhi, First edition 1997 (reprint in 2001).
Articles	https://pubs.acs.org/journal/mpohbp
References Books	Y W. Chien, Novel Drug Delivery Systems, 2nd edition, revised and expanded, Marcel Dekker, Inc., New York, 1992. 2. S. P. Vyas and R. K. Khar, Controlled Drug Delivery - concepts and advances, Vallabh Prakashan, New Delhi, First edition 2002. N.K. Jain, Controlled and Novel Drug Delivery, CBS Publishers & Distributors, New Delhi, First edition 1997 (reprint in 2001).
MOOC Courses	https://onlinecourses.nptel.ac.in/
Videos	https://www.youtube.com/watch?v=rGP7KZOTkzE

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COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	3	1	3	3	3	-	-	-	-	-	3	-	3	-	3
CO2	2	2	2	3	2	-	-	-	-	-	3	-	2	1	2
CO3	2	1	3	2	1	-	-	-	-	-	3	-	2	-	1
CO4	3	2	3	2	3	-	-	-	-	-	3	-	3	2	2
CO5	1	1	3	2	2	-	-	-	-	-	2	-	2	2	3
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Advanced Biopharmaceutics & Pharmacokinetics
Course Code	MPH 202T

			Part A						
Voar	1et	Somostor	and	Credits	L	Т	Р	С	
	130	Gemester	210	oreans	4	0	0	4	
Course Type	Theory only	ery only							
Course Category	Discipline Core	e							
Pre-Requisite/s				Co-Requisite/s					
Course Outcomes & Bloom's Level	CO1- Understa CO2- Explain CO3- Analyze elimination(BL CO4- Know cr CO5- Identify	CO1- Understand basic concepts in biopharmaceutics and pharmacokinetics (BL2-Understand) CO2- Explain design and evaluation of dosage regimens of the drugs using pharmacokinetic and biopharmaceutics parameters(BL2-Understand) CO3- Analyze raw data and derive the pharmacokinetic models and parameters the best describe the process of drug absorption, distribution, metabolism and elimination(BL3-Apply) CO4- Know critical evaluation of biopharmaceutics studies involving drug product equivalency(BL5-Evaluate) CO5- Identify potential clinical pharmacokinetic problems and apply basic pharmacokinetic principles to solve them(BL5-Evaluate)							
Coures Elements	Skill Development ✓ Entrepreneurship × Employability ✓ Professsonal Ethics × Gender × Human Values × Environment ×								

Part B

Modules	Contents	Pedagogy	Hours
UNIT-I	drug absorption, Factors affecting drug absorption, pH–partition theory of drug absorption. Formuulation and physicochemical factors: Dissolution rate, Dissolution process, Noyes– Whitney equation and drug dissolution, Factors affecting the dissolution rate. Gastrointestinal absorption: role of the dosage form: Solution (elixir, syrup and solution) as a dosage form , Suspension as a dosage form, Capsule as a dosage form, Tablet as a dosage form ,Dissolution methods , Formulation and processing factors, Correlation of in vivo data with in vitro dissolution data. Transport model: Permeability-Solubility-Charge State and the pH Partition Hypothesis, Properties of the Gastrointestinal Tract (GIT), pH Microclimate Intracellular pH Environment, Tight-Junction Complex.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12
UNIT-II	Biopharmaceutic considerations in drug product design and In Vitro Drug Product Performance: Introduction, biopharmaceutic factors affecting drug bioavailability, rate- limiting steps in drug absorption, physicochemical nature of the drug formulation factors affecting drug product performance, in vitro: dissolution and drug release testing, compendial methods of dissolution, alternative methods of dissolution testing, meeting dissolution requirements, problems of variable control in dissolution testing performance of drug products. In vitro-in vivo correlation, dissolution profile comparisons, drug product stability, considerations in the design of a drug product.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12
UNIT-III	Pharmacokinetics: Basic considerations, pharmacokinetic models, compartment modeling: one compartment model- IV bolus, IV infusion, extra-vascular. Multi compartment model: two compartment - model in brief, non-linear pharmacokinetics: cause of non-linearity, Michaelis – Menten equation, estimation of kmax and vmax. Drug interactions: introduction, the effect of protein- binding interactions, the effect of tissue-binding interactions, cytochrome p450- based drug interactions, drug interactions linked to transporters	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	10
UNIT-IV	Drug Product Performance, in vivo: Bioavailability and Bioequivalence: drug product performance, purpose of bioavailability studies, relative and absolute availability. methods for assessing bioavailability, bioequivalence studies, design and evaluation of bioequivalence studies, study designs, crossover study designs, evaluation of the data, bioequivalence example, study submission and drug review process. biopharmaceutics classification system, methods. Permeability: In-vitro, in-situ and In-vivo methods. Generic biologics (biosimilar drug products),clinical significance of bioequivalence studies, special concerns in bioavailability and bioequivalence studies, generic substitution.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	14
UNIT-V	Application of Pharmacokinetics: Modified-Release Drug Products, Targeted Drug Delivery Systems and Biotechnological Products. Introduction to Pharmacokinetics and pharmacodynamic, drug interactions. Pharmacokinetics and pharmacodynamics of biotechnology drugs. Introduction, Proteins and peptides, Monoclonal antibodies, Oligonucleotides, Vaccines (immunotherapy), Gene therapies.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12

Part C

Modules

Title

Indicative-ABCA/PBL/ Experiments/Field work/ Internships

EXPERIMENT	PREPARATIONS OF TARGETED DRUG DELIEVERY SYSTEM	PBL	BL3-Apply	10

Theory Minimum Passing Marks External Evaluation Min. External Evaluation Internal Evaluation Min. Internal Evaluation **Total Marks** 100 50 75 38 25 13 Practical Total Marks Minimum Passing Marks External Evaluation Min. External Evaluation Internal Evaluation Min. Internal Evaluation 0

Part D(Marks Distribution)

Part E

Books	1. Biopharmaceutics and Clinical Pharmacokinetics, An Introduction, 4th edition, revised and expande by Robert. E. Notari, Marcel Dekker Inc, New York and Basel, 1987. 2. Biopharmaceutics and Relevant Pharmacokinetics by John. G Wagner and M.Pemarowski, 1st edition, Drug Intelligence Publications, Hamilton, Illinois, 1971. 3. Encyclopedia of Pharmaceutical Technology, Vol 13, James Swarbrick, James. G. Boylan, Marcel Dekker Inc, New York, 1996. 12. Basic Pharmacokinetics, 1st edition, Sunil S Jambhekar and Philip J Breen, pharmaceutical press, RPS Publishing,2009. 13. Absorption and Drug Development- Solubility, Permeability, and Charge State, Alex Avdeef, John Wiley & Sons, Inc,2003. 9. Biopharmaceutics and Clinical Pharmacokinetics, An Introduction, 4th edition, revised and expande by Robert. E. Notari, Marcel Dekker Inc, New York and Basel,1987. 10. Biopharmaceutics and Relevant Pharmacokinetics by John. G Wagner and M.Pemarowski, 1st edition, Drug Intelligence Publications, Hamilton, Illinois, 1971. 11. Encyclopedia of Pharmaceutical Technology, Vol 13, James Swarbrick, James. G. Boylan, Marcel Dekker Inc, New York, 1996. 12. Basic Pharmacokinetics, 1st edition, Sunil S Jambhekar and Philip J Breen, pharmaceutical press, RPS Publishing,2009. 13. Absorption and Drug Development- Solubility, Permeability, and Charge State, Alex Avdeef, John Wiley & Sons, Inc,2003.
Articles	JOURNAL OF PHARMACEUTICS JOURNAL OF MOLECULAR PHARMACEUTICS
References Books	1. Biopharmaceutics and Clinical Pharmacokinetics by Milo Gibaldi, 4th edition, Philadelphia, Lea and Febiger, 1991 2. Biopharmaceutics and Pharmacokinetics, A. Treatise, D .M. Brahmankar and Sunil B. Jaiswal., VallabPrakashan, Pitampura, Delhi 3. Applied Biopharmaceutics and Pharmacokinetics by Shargel. Land YuABC, 2ndedition, Connecticut Appleton Century Crofts, 1985 4. Textbook of Biopharmaceutics and Pharmacokinetics, Dr. Shobha Rani R. Hiremath, Prism Book 5. Pharmacokinetics by Milo Gibaldi and D. Perrier, 2nd edition, Marcel Dekker Inc.,New York, 1982 6. Current Concepts in Pharmaceutical Sciences: Biopharmaceutics, Swarbrick. J, Leaand Febiger, Philadelphia, 1970 7. Clinical Pharmacokinetics, Concepts and Applications 3rd edition by MalcolmRowland and Thom~ N. Tozer, Lea and Febiger, Philadelphia, 1995 8. Dissolution, Bioavailability and Bioequivalence, Abdou. H.M, Mack Publishing Company, Pennsylvania 1989
MOOC Courses	NPTEL
Videos	https://www.youtube.com/watch?v=hXrniai9ZR0

Course Articulation Matrix

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	3	1	1	3	1	3	-	-	-	-	3	-	1	1	1
CO2	2	1	2	2	2	2	-	-	-	-	2	-	2	2	2
CO3	2	2	3	1	2	1	-	-	-	-	2	-	1	2	2
CO4	3	-	1	2	2	2	-	-	-	-	3	-	2	-	2
CO5	3	1	3	2	3	2	-	-	-	-	2	-	1	1	3
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Computer Aided Drug Delivery System
Course Code	MPH 203T

			Part A					
Veer	1 of	Somostor	and	Cradita	L	т	Р	С
Tear	151	Semester	2110	Creuits	4	0	0	4
Course Type	Theory only							
Course Category	Discipline C	ore						
Pre-Requisite/s		Co-Requisite/s						
Course Outcomes & Bloom's Level	C01- Describe statistical modeling and quality by design in pharmaceutical research and development (BL2-Understand) C02- Discuss the descriptors of drug disposition utilized in computational modeling(BL2-Understand) C03- Understand the ethics and legal protection of computing in pharmaceutical research(BL2-Understand) C04- Defend in silico approaches for biopharmaceutical characterization(BL3-Apply) C05- Recognize the importance of automation in pharmaceutical development(BL4-Analyze)							
Coures Elements	nts Skill Development ✓ Entrepreneurship × Employability ✓ Professsonal Ethics × SDG (Goals) Gender × Human Values × Environment ×			SDG3(Good health and well-being) SDG4(Quality education) SDG7(Affordable and clean energy) SDG9(Industry Innovation and Infrastructu	re)			

Part B

Modules	Contents	Pedagogy	Hours
UNIT-I	a. Computers in Pharmaceutical Research and Development: A General Overview: History of Computers in Pharmaceutical Research and Development. Statistical modeling in Pharmaceutical research and development: Descriptive versus Mechanistic Modeling, Statistical Parameters, Estimation, Confidence Regions, Nonlinearity at the Optimum, Sensitivity Analysis, Optimal Design, Population Modeling b. Quality-by-Design In Pharmaceutical Development: Introduction, ICH Q8 guideline, Regulatory and industry views on QbD, Scientifically based QbD - examples of application.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12
UNIT-II	2 Computational Modeling Of Drug Disposition: Introduction ,Modeling Techniques: Drug Absorption, Solubility, Intestinal Permeation, Drug Distribution ,Drug Excretion, Active Transport; P-gp, BCRP, Nucleoside Transporters, hPEPT1, ASBT, OCT, OATP, BBB-Choline Transporter	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12
UNIT-III	3 Computer-aided formulation development: Concept of optimization, Optimization parameters, Factorial design, Optimization technology & Screening design. Computers in Pharmaceutical Formulation: Development of pharmaceutical emulsions, microemulsion drug carriers Legal Protection of Innovative Uses of Computers in R&D, The Ethics of Computing in Pharmaceutical Research, Computers in Market analysis	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	15
UNIT-IV	4 a. Computer-aided biopharmaceutical characterization: Gastrointestinal absorption simulation. Introduction, Theoretical background, Model construction, Parameter sensitivity analysis, Virtual trial, Fed vs. fasted state, Invitro dissolution and in vitro- in vivo correlation, Biowaiver considerations b. Computer Simulations in Pharmacokinetics and Pharmacodynamics: Introduction, Computer Simulation: Whole Organism, Isolated Tissues, Organs, Cell, Proteins and Genes. c. Computers in Clinical Development: Clinical Data Collection and Management, Regulation of Computer Systems	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12
UNIT-V	5 Artificial Intelligence (AI), Robotics and Computational fluid dynamics: General overview, Pharmaceutical Automation, Pharmaceutical applications, Advantages and Disadvantages. Current Challenges and Future Directions	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	9

Part C											
Modules	Title	Indicative-ABCA/PBL/ Experiments/Field work/ Internships	Bloom's Level	Hours							
1	ARTIFICIAL INTELLIGENCE IN HEALTH CARE	Seminar	BL4-Analyze	2							

Part D(Marks Distribution)

	Ineory												
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation								
100	50	75	38	25	13								
			Practical										
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation Min. Internal Evaluation									

	Part E
Books	1. Computer Applications in Pharmaceutical Research and Development, Sean Ekins, 2006, John Wiley & Sons. 2. Computer-Aided Applications in Pharmaceutical Technology, 1st Edition, Jelena Djuris, Woodhead Publishing 3. Encyclopediaof Pharmaceutical Technology, Vol 13, James Swarbrick, James. G. Boylan, Marcel Dekker Inc, New York, 1996.
Articles	1. Computer Applications in Pharmaceutical Research and Development, Sean Ekins, 2006, John Wiley & Sons. 2. Computer-Aided Applications in Pharmaceutical Technology, 1st Edition, Jelena Djuris, Woodhead Publishing 3. Encyclopediaof Pharmaceutical Technology, Vol 13, James Swarbrick, James. G. Boylan, Marcel Dekker Inc, New York, 1996.
References Books	1. Computer Applications in Pharmaceutical Research and Development, Sean Ekins, 2006, John Wiley & Sons. 2. Computer-Aided Applications in Pharmaceutical Technology, 1st Edition, Jelena Djuris, Woodhead Publishing 3. Encyclopediaof Pharmaceutical Technology, Vol 13, James Swarbrick, James. G. Boylan, Marcel Dekker Inc, New York, 1996.
MOOC Courses	nptel
Videos	pharmawins

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COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	1	1	3	-	3	-	1	-	-	-	3	-	1	-	1
CO2	2	2	3	-	3	-	-	-	-	-	3	-	-	1	2
CO3	1	-	2	-	2	-	-	-	-	-	3	-	2	2	2
CO4	3	1	2	-	3	-	1	-	-	-	2	-	1	-	-
CO5	3	-	1	-	3	-	1	-	-	-	2	-	-	2	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Cosmetics and Cosmeceuticals
Course Code	MPH 204T

			Part A					
Voar	1st Somostor		and	Cradita	L	т	Р	С
Tear	151	Semester		Oreans	4	0	0	4
Course Type	Theory only							
Course Category	Discipline C	ore						
Pre-Requisite/s				Co-Requisite/s				
Course Outcomes & Bloom's Level	Outcomes CO1- Gain information on key ingredients used in cosmetics and cosmeceuticals (BL1-Remember) CO2- Understand key building blocks of cosmetics for various formulations (BL2-Understand) CO3- Know the current technologies in the market(BL2-Understand) CO4- Understand the scientific principles to develop cosmetics and cosmeceuticals with desired safety(BL2-Understand) CO5- Understand the regulatory aspects in cosmetics(BL4-Analyze)							
Coures Elements	Skill Develop Entrepreneu Employabilit Professsona Gender X Human Valu Environmen	orment ✓ irship ✓ y ✓ il Ethics X es X t X	SDG (Goals)	SDG3(Good health and well-being) SDG4(Quality education) SDG9(Industry Innovation and Infrastructure)				

Part B

Modules	Contents	Pedagogy	Hours
UNIT-I	Cosmetics – Regulatory: Definition of cosmetic products as per Indian regulation. Indian regulatory requirements for labeling of cosmetics Regulatory provisions relating to import of cosmetics., Misbranded and spurious cosmetics. Regulatory provisions relating to manufacture of cosmetics – Conditions for obtaining license, prohibition of manufacture and sale of certain cosmetics, loan license, offences and penalties.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12
UNIT-II	Cosmetics - Biological aspects: Structure of skin relating to problems like dry skin, acne, pigmentation, prickly heat, wrinkles and body odor. Structure of hair and hair growth cycle. Common problems associated with oral cavity. Cleansing andcare needs for face, eye lids, lips, hands, feet, nail, scalp, neck, body and under-arm.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12
UNIT-III	Formulation Building blocks: Building blocks for different product formulations of cosmetics/cosmeceuticals. Surfactants – Classification and application. Emollients, rheological additives: classification and application. Antimicrobial used as preservatives, their merits and demerits. Factors affecting microbial preservative efficacy. Building blocks for formulation of a moisturizing cream, vanishing cream, cold cream, shampoo and toothpaste. Soaps and syndetbars. Perfumes; Classification of perfumes. Perfume ingredients listed as allergens in EU regulation	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	14
UNIT-IV	Design of cosmeceutical products: Sun protection, sunscreens classification and regulatory aspects. Addressing dry skin, acne, sun-protection, pigmentation, prickly heat, wrinkles, body odor., dandruff, dental cavities, bleeding gums, mouth odor and sensitive teeth through cosmeceutical formulations	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12
UNIT-V	Herbal Cosmetics: Herbal ingredients used in Hair care, skin care and oral care. Review of guidelines for herbal cosmetics by private bodies like cosmos with respect to preservatives, emollients, foaming agents, emulsifiers and rheology modifiers. Challenges in formulating herbal cosmetics.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	10

Part C

Modules	Title	Indicative-ABCA/PBL/ Experiments/Field work/ Internships	Bloom's Level	Hours
UNIT-4	COSMETIC PREPARATIONS	PBL	BL3-Apply	12

	Part D(Marks Distribution)											
Theory												
Total Marks	Total Marks Minimum Passing Marks External Evaluation Min. External Evaluation Internal Evaluation Min. Internal Evaluation											
100	50 75 38 25 13											
			Practical									
Total Marks	Total Marks Minimum Passing Marks External Evaluation Min. External Evaluation Internal Evaluation Min. Internal Evaluation											
	0											

Part E

Books	Harry's Cosmeticology. 8th edition. Poucher'sperfumecosmeticsandSoaps,10th edition. 3. Cosmetics - Formulation, Manufacture and quality control, PP.Sharma,4th edition Handbook of cosmetic science and Technology A. O. Barel, M. Payeand H.I. Maibach. 3rd edition 5. Cosmetic and Toiletries recent suppliers catalogue. CTFA directory.
Articles	https://onlinelibrary.wiley.com/journal/14682494
References Books	Harry's Cosmeticology. 8th edition. Poucher'sperfumecosmeticsandSoaps,10th edition. Cosmetics - Formulation, Manufacture and quality control, PP.Sharma,4th edition Handbook of cosmetic science and Technology A. O. Barel, M. Payeand H.I. Maibach. 3rd edition Cosmetic and Toiletries recent suppliers catalogue. CTFA directory.
MOOC Courses	https://nptel.ac.in/
Videos	https://www.youtube.com/watch?v=bcCkQ1liaKA

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COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	1	-	-	-	2	-	-	-	-	-	3	-	1	1	1
CO2	3	1	-	-	2	-	-	-	-	-	3	-	2	2	2
CO3	2	2	-	-	2	-	-	-	-	-	1	-	1	-	2
CO4	1	2	-	-	1	-	-	-	-	-	2	-	1	-	3
CO5	3	-	-	-	-	-	-	-	-	-	1	-	-	2	2
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Pharmaceutics Practical II
Course Code	MPH 205P

			Part A					
Voar	1et	Somostor	and	Cradits	L	Т	Р	С
i eai	150	Jennester	210	Credits	7	0	0	7
Course Type	Lab only							
Course Category	Discipline Core)						
Pre-Requisite/s	TO GAIN EXP	TO GAIN EXPERIMEMTAL KNOWLEDGE Co-Requisite/s						
Course Outcomes & Bloom's Level	CO1- To understand and apply the practical knowledge(BL2-Understand) CO2- To understand the concepts of bioavailability and bioequivalence of drug products and their significance(BL2-Understand) CO3- To prepare several herbal and cosmetic formulations(BL3-Apply) CO4- Hands training on new deug devlopment softwares(BL3-Apply)							
Coures Elements	Skill Developm Entrepreneursl Employability Professsonal E Gender X Human Values Environment X	nent √ hip X / Ethics X X	SDG (Goals)	SDG3(Good health and well-being) SDG4(Quality education)	education)			

Part B

Modules	Contents	Pedagogy	Hours

Part	C

Modules	Title	Title Indicative-ABCA/PBL/ Title Experiments/Field work/ Internships			
1	1. To study the effect of temperature change, non-solvent addition, incompatible polymer addition in microcapsules preparation	Experiments	BL4-Analyze	4	
2	2. Preparation and evaluation of Alginate beads	PBL	BL6-Create	4	
3	3. Formulation and evaluation of gelatin /albumin microspheres	PBL	BL6-Create	4	
4	4. Formulation and evaluation of liposomes/niosomes	PBL	BL6-Create	4	
5	5. Formulation and evaluation of spherules	PBL	BL6-Create	4	
6	6. Improvement of dissolution characteristics of slightly soluble drug by Solid dispersion technique.	PBL	BL4-Analyze	4	
7	7. Comparison of dissolution of two different marketed products /brands	PBL	BL5-Evaluate	4	
8	8. Protein binding studies of a highly protein bound drug & poorly protein bound drug	PBL	BL4-Analyze	4	

Part D(Marks Distribution)

Theory							
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation		
Practical							
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation		
150	75	100	50	50	25		

Books	7. Pharmaceutics- The science of dosage form design by M.E.Aulton, Churchill Livingstone, Latest edition				
Articles	https://www.ipinnovative.com/journal-name/JPBS				
References Books	1 Modern Pharmaceutics by Gilbert S. Banker & C.T. Rhodes, 3rd Edition 2. Remington: The Science and Practice of Pharmacy, 20th edition Pharmaceutical Science (RPS) 3. Theory and Practice of Industrial Pharmacy by Liberman & Lachman				
MOOC Courses	https://nptel.ac.in/				
Videos	NA				

Part E

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	3	-	-	-	-	-	-	-	-	-	3	-	1	-	1
CO2	3	-	1	1	-	-	-	-	-	-	3	-	-	1	2
CO3	2	-	2	2	-	-	-	-	-	-	3	-	2	-	3
CO4	3	-	-	1	-	-	-	-	-	-	3	-	-	-	2
CO5	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-

Course Articulation Matrix



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Seminars
Course Code	MPH 206S

			Part A						
Voar	1et	Somostor	2nd	Credite	L	Т	Р	С	
i eai	151	Semester	2110	Credits	0	0	7	7	
Course Type	Lab only	only							
Course Category	Discipline Cor	scipline Core							
Pre-Requisite/s				Co-Requisite/s					
Course Outcomes & Bloom's Level	CO1- The pur CO2- To Focu CO3- To impre	O1- The purpose of a seminar is to enable students to improve their knowledge and understanding of a topic by engaging with key issues(BL3-Apply) O2- To Focus on sharing the knowledge(BL3-Apply) O3- To improve the communicative ,presentation and understanding skills in students(BL3-Apply)							
Coures Elements	Skill Developr Entrepreneurs Employability Professsonal Gender X Human Value Environment	nent ✓ ship ✓ ✓ Ethics X s X X	SDG (Goals)	SDG3(Good health and well-being) SDG4(Quality education) SDG7(Affordable and clean energy)					

Part B

Modules	Contents	Pedagogy	Hours
UNIT-I	a. Computers in Pharmaceutical Research and Development: A General Overview: History of Computers in Pharmaceutical Research and Development. Statistical modeling in Pharmaceutical research and development: Descriptive versus Mechanistic Modeling, Statistical Parameters, Estimation, Confidence Regions, Nonlinearity at the Optimum, Sensitivity Analysis, Optimal Design, Population Modeling b. Quality-by-Design In Pharmaceutical Development: Introduction, ICH Q8 guideline, Regulatory and industry views on QbD, Scientifically based QbD - examples of application.	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	12

	Part C									
Modules	ules Title Indicative-ABCA/PBL/ Internships		Bloom's Level	Hours						
1	Polymers in NDDS	Seminar	BL2-Understand	12						
2	Gene theray	Seminar	BL2-Understand	10						
3	PKPD studies	Research Paper Presentation	BL4-Analyze	12						
4	Insilico drug designing	Simulation	BL3-Apply	8						

Part D(Marks Distribution)									
	Theory								
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation				
			Practical						
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation				
100	50	75	38	25	13				

	Par E							
Books	NA							
Articles	NA							
References Books	NA							
MOOC Courses	NA							
Videos	NA							

Part F

Course Articulation Matrix

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	1	1	1-	-	-	-	-	-	-	3	3	1	1	-	3
CO2	-	2	2	-	-	-	-	-	-	3	3	2	-	-	1
CO3	2	-	1	-	-	-	-	-	-	3	3	1	2	-	2
CO4	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO5	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Online Certificate Course
Course Code	MPH 207ET

			Part A						
Voar	1 ct	Somostor	and	Credite	L	Т	Р	С	
Teal			2110	Oreans	2	0	0	2	
Course Type	Online cours	line course							
Course Category	Skill Enhand	Il Enhancement Courses							
Pre-Requisite/s	Pharmaceu	tical background		Co-Requisite/s					
Course Outcomes & Bloom's Level	CO1- Why (CO2- QbD a CO3- Deter	CO1- Why QbD, what is QbD? FDA objectives, the QbD approach() CO2- QbD and continuous processing: FDA perspective, advantages, challenges() CO3- Determining CPPs and CMAs, CQA,QTPP,RISK ASSESMENT()							
Coures Elements	Skill Develo Entrepreneu Employabili Professsona Gender X Human Valu Environmen	pment ✓ urship × ty ✓ al Ethics × tes × t ×	SDG (Goals)	SDG3(Good health and well-being) SDG4(Quality education) SDG9(Industry Innovation and Infrastructure)					

Part B

Modules	Contents	Pedagogy	Hours
UNIT-I	As per recommended Course provider	PEER TUTORIAL	60

	Theory								
Total Marks	ks Minimum Passing Marks External Evaluation Min. External Evaluation Internal Evaluation Min. Internal Evaluation								
50	25	35	18	15	8				
			Practical						
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation Min. Internal Evalua					
	0								

Ρ	а	r	t	E

Books	NA
Articles	ΝΑ
References Books	ΝΑ
MOOC Courses	1. https://ispe.org/training/course/qbd 2. https://www.udemy.com/course/quality-by-design-qbd-in-pharmaceutical-development/? =&utm_source=adwords&utm_medium=udemyads&utm_campaign=LongTail_la.EN_cc.INDIA&utm_content=deal4584&utm_term=ag_118445032537ad_618853564450kwde_cdmpl 1212271230479li_1007795pd&matchtype=&gad_source=1&gclid=Cj0KCQjwztOwBhD7ARIsAPDKnkBXWZvoPEBACFjanUEyfigFRV8AJSdYMY2AH1tTLjXCBcpyt7DSYX8aAsvOEALw_wcB&cou 3. https://www.6sigma.us/quality-by-design/
Videos	NA

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	3	1	1	-	-	-	-	-	-	-	3	-	1	1	1
CO2	2	2	2	-	-	-	-	-	-	-	3	-	2	2	2
CO3	3	-	-	-	-	-	-	-	-	-	3	-	1	1	1
CO4	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO5	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Introduction to intellectual property rights
Course Code	MPH 208ET

			Part A						
Voar	1et	Somostor	and	Cradita	L	т	Р	С	
i eai	130	Semester	2114	oreuts	3	1	0	4	
Course Type	Theory only	eory only							
Course Category	Skill Enhance	ement Courses							
Pre-Requisite/s		Co-Requisite/s							
Course Outcomes & Bloom's Level	CO1- To crea CO2- To mal CO3- Develo CO4- To kno CO5- To app	ate awareness of IPR among ph ke the pharmacy students aware op the understanding of the Intel w the database of intellectual pr ly the Knowledge of IPR in draft	armacy students.(BL2-Under about the pharmaceutical R lectual Property Rights neces operty and TKDL(BL2-Under ing and fillng of IPR(BL3-App	rstand) & D and the activities therein. (BL2-Underst sary for research activities in the pharmaceu rstand) bly)	and) Itical industry	/.(BL3-Apply)			
Coures Elements	Skill Develop Entrepreneu Employability Professsona Gender X Human Value Environment	oment ✓ rship ✓ y ✓ I Ethics ✓ es √ : ×	SDG (Goals)	SDG1(No poverty) SDG3(Good health and well-being) SDG4(Quality education) SDG6(Clean water and sanitation) SDG8(Decent work and economic growth) SDG17(Partnerships for the goals)					

Part B

				1
N	Modules	Contents	Pedagogy	Hours
U	NIT 1	The pharmaceutical business and The pharmaceutical R & D	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	10
U	NIT 2	Module 3 – Intellectual Property Rights: Introduction about patents, copyright, trademark, Industrial Designs,Geographical Indications, Trade Secrets, Module 4 – IPR: With specific reference to pharma	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	10
U	NIT 3	IPR: Indian patent scenario and Patent commercialization and licensing	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	10
U	NIT 4	Patent drafting and Patent searches, patent filing, registration, granting World Intellectual Property Organization (WIPO) and its functions	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	08
U	NIT 5	IP in Traditional Knowledge, TKDL database in medicinal plants, INDIAN WEB- PORTALS FOR PATENTS AND TECHNOLOGIES	Lecture based learning, interactive class, Peer tutorial, Class using ICT tool/PPT/white board	07

Part C

Modules	Title	Indicative-ABCA/PBL/ Experiments/Field work/ Internships	Bloom's Level	Hours
1	patent drafting and filing	Case Study	BL3-Apply	5

	Part D(Marks Distribution)					
	Theory					
Total Marks	Total Marks Minimum Passing Marks External Evaluation Min. External Evaluation Internal Evaluation Min. Internal Evaluation					
100 50		75	38	25	13	
	Practical					
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation	

Books	Cockburn IM. Intellectual property rights and pharmaceuticals: challenges and opportunities for economic research. The economics of intellectual property. 2009 Jan: 150.			
Articles	Savale SK, Savale VK. Intellectual property rights (IPR). World J Pharm Pharm Sci. 2016 Apr 22;5:2559-92.			
References Books	Prabu SL, Tnk S, editors. Intellectual property rights. BoD–Books on Demand; 2017 Jun 21.			
MOOC Courses	NEPTEL			
Videos	NA			

Part E

Course Articulation Matrix

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	-	2	-	1	-	-	-	3	-	2	3	-	1	1	1
CO2	-	1	-	-	-	-	-	3	-	-	2	-	1	-	1
CO3	-	1	-	-	-	-	-	2	-	-	-	-	-	-	-
CO4	-	-	-	-	-	-	-	2	-	-	2	-	-	-	-
CO5	-	-	-	-	-	-	-	-	-	-	1	-	-	-	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-

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(SOP)(MPharm-PharmaCeutics)

Title of the Course	Research Methodology and Biostatistics
Course Code	MPH 301T

Part A

Voar	2nd	Somostor	3rd	Cradits	L	Т	Р	С
l C ai	2110	Jemester	510	erouno	4	0	0	4
Course Type	Theory on	heory only						
Course Category	Discipline	Core						
Pre-Requisite/s				Co-Requisite/s				
Course Outcomes & Bloom's Level	CO1- Dev CO2- Des CO3- Cho project(BL CO4- Dev	 CO1- Develop the ability to apply the methods while working on a research project work(BL2-Understand) CO2- Describe the appropriate statistical methods required for a particular research design (BL2-Understand) CO3- Choose the appropriate research design and develop appropriate research hypothesis for a research project(BL3-Apply) CO4- Develop an appropriate framework for research studies(BL4-Analyze) 					nd)	
Coures Elements	Coures ElementsSkill Development ✓ Entrepreneurship × Employability ✓ Professsonal Ethics × Gender × Human Values × Environment ×		SDG (Goals)	SDG3(Good health and wel SDG4(Quality education)	l-being)			

.

Modules	Contents	Pedagogy	Hours
UNIT-I	General Research Methodology: Research, objective, requirements, practical difficulties, review of literature, study design, types of studies, strategies to eliminate errors/bias, controls, randomization, crossover design, placebo, blinding techniques	WHITEBOARD	12
UNIT-II	Biostatistics: Definition, application, sample size, importance of sample size, factors influencing sample size, dropouts, statistical tests of significance, type of significance tests, parametric tests (students "t" test, ANOVA, Correlation coefficient, regression), non- parametric tests (wilcoxan rank tests, analysis of variance, correlation, chi square test), null hypothesis, P values, degree of freedom, interpretation of P values.	WHITE BOARD	12
UNIT-III	Medical Research: History, values in medical ethics, autonomy, beneficence, non- maleficence, double effect, conflicts between autonomy and beneficence/non- maleficence, euthanasia, informed consent, confidentiality, criticisms of orthodox medical ethics, importance of communication, control resolution, guidelines, ethics committees, cultural concerns, truth telling, online business practices, conflicts of interest, referral, vendor relationships, treatment of family members, sexual relationships, fatality.	WHITE BOARD	12
UNIT-IV	CPCSEA guidelines for laboratory animal facility: Goals, veterinary care, quarantine, surveillance, diagnosis, treatment and control of disease, personal hygiene, location of animal facilities to laboratories, anesthesia, euthanasia, physical facilities, environment, animal husbandry, record keeping, SOPs, personnel and training, transport of lab animals	WHITE BOARD	12
UNIT-V	Declaration of Helsinki: History, introduction, basic principles for all medical research, and additional principles for medical research combined with medical care	WHITE BOARD	12

Part D(Marks Distribution)

	Theory					
Total Minimum Passing Marks Marks		External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation	
100	50	75	37	25	12	
			Practical			
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation	
	0					

Part E	
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1 Pharmaceutical statistics, Practical and clinical applications, Sanford Bolton, publisher Marcel Dekker Inc.	

Books	NewYork. 2. Fundamental of Statistics – Himalaya Publishing House- S.C.Guptha
Articles	NA
References Books	1. Design and Analysis of Experiments –PHI Learning Private Limited, R. Pannerselvam. 2. Design and Analysis of Experiments – Wiley Students Edition, Douglas and C. Montgomery
MOOC Courses	https://www.coursera.org/search?query=biostatistics%20in%20public%20health
Videos	https://www.youtube.com/watch?v=UtivXLO7c9A

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COs	PO1	PO2	PO3	PO4	PO5	PO6	P07	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	-	3	3	3	-	-	-	-	-	-	3	-	1	1	1
CO2	-	2	2	1	-	-	-	-	-	-	3	-	2	2	2
CO3	-	3	1	2	-	-	-	-	-	-	2	-	1	2	1
CO4	-	3	3	3	-	-	-	-	-	-	2	-	2	-	2
CO5	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Journal Club
Course Code	MPH 302P

Part A

Voar	2nd Semester		3rd	Cradits	L	Т	Р	С
Teal	2110	Jemester		Credits	0	0	1	1
Course Type	Lab only			-	-	-		-
Course Category	Discipline	Core						
Pre-Requisite/s				Co-Requisite/s				
Course Outcomes & Bloom's Level	CO1- To a Understa CO2- o cri	ccquire communicatior nd) tically evaluate recent a	n skills,group discussion articles in the academ	on, team work and to gain nev ic literature (BL3-Apply)	v knowl	edge (B l	L 2 -	
Coures Elements	Skill Deve Entrepren Employab Professso Gender X Human Va Environme	lopment ✓ eurship X ility ✓ nal Ethics ✓ ilues X ent X	SDG (Goals)	SDG4(Quality education) SDG17(Partnerships for the	DG4(Quality education) DG17(Partnerships for the goals)			

Part B

Modules Contents Pedagogy Hours

Part C

Modules	Title	Indicative-ABCA/PBL/ Experiments/Field work/ Internships	Bloom's Level	Hours
1	To understand the recent literature in Pharmaceutical sciences	Research Paper Presentation	BL3-Apply	12
2	To prepare review articles	Case Study	BL5-Evaluate	5
3	To present review articles	Research Paper Presentation	BL3-Apply	2

Part D(Marks Distribution)

	пеогу								
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation				
	Practical								
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation				
25	13	25	12	0	0				

Part E

.

Books	NA
Articles	
References Books	NA
MOOC Courses	NA
Videos	NA

Course Articulation Matrix

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	1	1	-	1	-	-	-	-	-	-	3	-	1	2	3
CO2	1	2	-	2	-	-	-	-	-	-	3	-	-	3	1
CO3	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO4	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO5	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Discussion Presentation (Proposal Presentation)
Course Code	MPH 303P

Part A

Voar	2nd Semester		3rd	Cradits	L	Т	Р	С
Ισαι	2110	Semester	510	Credits	0	0	2	2
Course Type	Lab only				•	•	-	
Course Category	Projects ar	nd Internship						
Pre-Requisite/s	Selection a	and discussion of projec	ct proposal	Co-Requisite/s				
Course Outcomes & Bloom's Level	CO1- To know how to frame a research problem () CO2- to know how to set a hypothesis() CO3- can able to perform literature survey, and prepare a proposal of thesis()							
Coures Elements	Skill Devel Entreprene Employabi Professsor Gender X Human Va Environme	opment ✓ eurship × lity × nal Ethics × lues × ent ×	SDG (Goals)	SDG3(Good health and wel SDG4(Quality education) SDG17(Partnerships for the	(Good health and well-being) (Quality education) 7(Partnerships for the goals)			

Part B

Modules	Contents	Pedagogy	Hours
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Part C

Modules	Title	Indicative-ABCA/PBL/ Experiments/Field work/ Internships	Bloom's Level	Hours
1	General Research Methodology: Research, objective, requirements, practical difficulties, review of literature, study design, types of studies, strategies to eliminate errors/bias, controls, randomization, crossover design, placebo, blinding techniques	Research Paper Presentation	BL3-Apply	12

	Theory							
Total Marks	Total MarksMinimum Passing MarksExternal EvaluationMin. External 							
			Practical					
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation			

Part D(Marke Distribution)

Part E

.

Books	NA
Articles	NA
References Books	NA
MOOC Courses	NA
Videos	NA

Course Articulation Matrix

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	3	1	2	2	-	-	-	-	-	-	3	-	1	2	2
CO2	3	1	1	1	-	-	-	-	-	-	3	-	2	2	1
CO3	1	2	2	3	-	-	-	-	-	-	2	-	1	1	1
CO4	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO5	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Research Work
Course Code	MPH 304P

			Part A						
Year	2nd	Somostor	3rd	Cradits	L	Т	Ρ	С	
i cui	Zilu Semester			oreans	0	0	14	14	
Course Type	Lab only								
Course Category	Projects ar	d Internship							
Pre-Requisite/s		Co-Requisite/s							
Course Outcomes & Bloom's Level	CO1- To know how to conduct a research(BL2-Understand) CO2- To know about different research methodolgies(BL2-Understand) CO3- Apllication of research principles and stastical principles(BL3-Apply)								
Coures Elements	Skill Develo Entreprene Employabil Professsor Gender X Human Val Environme	opment ✓ eurship ✓ ity ✓ nal Ethics ✓ ues √ nt X	nt ✓ o ✓ nics ✓ SDG (Goals) SDG4(Quality education)						

Part B

Modules	Contents	Pedagogy	Hours
UNIT-I			

Part D(Marks Distribution)

	Theory							
Total Marks	Minimum Passing MarksExternal EvaluationMin. External EvaluationInternal 							
	175							
			Practical					
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation			
350	175	200	100	150	75			

Part E

Books	NA
Articles	NA
References Books	NA
MOOC Courses	NA
Videos	NA

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COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	3	3	3	3	2	2	-	-	-	-	3	-	1	3	3
CO2	3	2	3	2	2	1	-	-	-	-	3	-	2	1	3
CO3	3	3	3	2	2	1	-	-	-	-	3	-	3	1	3
CO4	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO5	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Research Presentation in Seminar/ Conference/Symposium
Course Code	MPH 305P

			Part A					
Year	2nd	Somostor	3rd	Credits	L	Т	Ρ	С
i c ai	2110	Gemester	514	Oredits	0	0	2	2
Course Type	Lab only				-	-	-	-
Course Category	Projects and Internship							
Pre-Requisite/s	know about preparation of presentation Co-Requisite/s							
Course Outcomes & Bloom's Level	CO1- devel CO2- devel	op the communication skil op presentation skills (BL3		-				
Coures Elements	Skill Develo Entreprene Employabili Professson Gender X Human Valu Environmer	opment ✓ urship × ity × al Ethics × ues × nt ×	SDG (Goals)	SDG4(Quality education)				

Part B

Modules	Contents	Pedagogy	Hours

Part D(Marks Distribution)

	Theory									
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation					
			Practical							
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation					
50	25	50	25	0	0					

Books	NA

Articles	NA
References Books	NA
MOOC Courses	NA
Videos	NA

.

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	1	-	-	-	-	-	-	-	-	3	3	3	1	2	3
CO2	1	1	1	-	-	-	-	-	-	3	3	3	2	2	2
CO3	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO4	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO5	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-

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(SOP)(MPharm-PharmaCeutics)

Title of the Course	Journal Club
Course Code	MPH 401P

Part A

Vear	2nd	Somostor	Ath	Credits Co-Requisite/s nd to gain new knowledge(E L3-Apply)	L	Т	Ρ	С
i c ai	ZIIU	Jemester	401	oroano		0	1	1
Course Type	Lab only							
Course Category	Projects and I	nternship						
Pre-Requisite/s	In a journal clu or research in purpose. The appropriatene conclusions d the research t allow pharmad make evidence in education a	n a journal club, a group of participants who have common practice or research interests meet regularly for a defined pedagogical ourpose. The club often discusses current research articles and the appropriateness of the study design, the data analysis, the conclusions drawn, and the potential applications or implications of he research to practice and patient care. In pharmacy, these clubs allow pharmacists to understand the current drug research to help nake evidence-based recommendations. The goals of journal clubs n education and research.						
Course Outcomes & Bloom's Level	CO1- To accq Understand) CO2- To critic	uire communication skills,grou ally evaluate recent articles in	p discussion, team work ar the academic literature (BL	nd to gain new knowledge(BL _3-Apply)	.2-			
Coures Elements	Skill Developr Entrepreneurs Employability Professsonal Gender X Human Values Environment 2	nent √ ship X X Ethics X s X X	SDG (Goals)	SDG4(Quality education)				

Modules Contents Pedagogy Hours

Part C

Modules	Title	Indicative-ABCA/PBL/ Experiments/Field work/ Internships	Bloom's Level	Hours
1	To understand the recent literature in Pharmaceutical sciences	Internships	BL3-Apply	8

Part D(Marks Distribution)

	Theory									
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation					
			Practical							
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation					
25	12	25	13	0	0					

Part E

.

Books	NA
Articles	NA
References Books	NA
MOOC Courses	NA
Videos	NA

Course Articulation Matrix

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	1	1	-	-	-	-	-	-	-	-	3	-	1	-	3
CO2	3	-	1	-	-	-	-	-	-	-	3	-	-	-	1
CO3	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO4	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO5	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Discussion/Presentation (Proposal Presentation)
Course Code	MPH 402P

Part A

Voar	2nd Semester		Ath	Cradits	L T		Р	С
Teal			401	Credits	0	0	4	4
Course Type	Lab only							
Course Category	Projects and	d Internship						
Pre-Requisite/s	Selection ar	nd Discussion of project pro	Co-Requisite/s					
Course Outcomes & Bloom's Level	CO1- 1.To k CO2- 2. To k CO3- 3. can	CO1- 1.To know how to frame a research problem(BL2-Understand) CO2- 2. To know how to set a hypothesis (BL3-Apply) CO3- 3. can able to perform literature survey and prepare project proposal(BL4-Analyze)						
Coures Elements	Skill Develo Entrepreneu Employabilit Professsona Gender X Human Valu Environmen	pment ✓ urship × ty × al Ethics × nes × t ×	SDG (Goals)	SDG4(Quality education)				

Part B

Modules	Contents	Pedagogy	Hours
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Part D(Marks Distribution)

	Theory									
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation					
			Practical							
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation					
75	38	75	38	0	0					

Books	NA
Articles	NA
References Books	NA
MOOC Courses	NA
Videos	NA

•

COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	3	2	3	3	-	-	-	-	-	-	3	-	1	-	2
CO2	3	3	1	3	-	-	-	-	-	-	3	-	2	-	1
CO3	3	3	2	2	-	-	-	-	-	-	2	-	2	2	2
CO4	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO5	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Discussion/Final Presentation
Course Code	MPH 403P

Part A

Voar	Year 2nd Semester 4th Credits	L	Т	Р	С			
Tear	2110	Semester	401	Creans	0	0	3	3
Course Type	Lab only	-	•	•	-			-
Course Category	Projects an	d Internship						
Pre-Requisite/s				Co-Requisite/s				
Course Outcomes & Bloom's Level	CO1- To kn CO2- To kn CO3- Apllic	CO1- To know how to conduct a research (BL2-Understand) CO2- To know about different research methodolgies(BL2-Understand) CO3- Apllication of research principles and stastical principles(BL3-Apply)						
Coures Elements	Skill Development ✓ Entrepreneurship × Employability × Professsonal Ethics × Gender × Human Values × Environment ×							

Part B

Modules Contents Pedagogy Hours	
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	Part D(Marks Distribution)								
	Theory								
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation				
	200								
			Practical						
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation				
400	200	200	100	200	100				

Books	NA
Articles	NA
References Books	NA
MOOC Courses	NA
Videos	NA

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COs	PO1	PO2	PO3	PO4	PO5	PO6	P07	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	-	3	3	3	1	-	-	-	-	-	3	-	1	-	3
CO2	-	2	3	3	1	-	-	-	-	-	3	-	-	-	1
CO3	-	3	3	3	-	-	-	-	-	-	2	-	1	-	3
CO4	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO5	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-



(SOP)(MPharm-PharmaCeutics)

Title of the Course	Research Publication /Report
Course Code	MPH 404T

Part A

Voar	and	Somostor	Ath	Credits	L	Т	Р	С
Teal	2110	Semester	401	Credits	0	0	2	2
Course Type	Lab only							
Course Category	Projects an	d Internship						
Pre-Requisite/s				Co-Requisite/s				
Course Outcomes & Bloom's Level	CO1- stude CO2- Impro CO3- shoul	CO1- students should improve thier publication writing skills(BL2-Understand) CO2- Improve their communication skills (BL3-Apply) CO3- should get sound knowledge of various databases ,referencing styles and softwares(BL3-Apply)						
Coures Elements	Skill Develo Entreprene Employabil Professson Gender X Human Val Environmer	pment × urship × ty × al Ethics ✓ SDG (Goals) SDG4(Quality education) ues √ ot ×						

Part B

Modules	Contents	Pedagogy	Hours
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Part C

Modules	Title	Indicative-ABCA/PBL/ Experiments/Field work/ Internships	Bloom's Level	Hours
1	presenting research/review papers in various seminars ,conferences or symposiums	PBL		

Part D(Marks Distribution)

Theory										
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation					
	25									

Flactical										
Total Marks	Minimum Passing Marks	External Evaluation	Min. External Evaluation	Internal Evaluation	Min. Internal Evaluation					
50	0	50	25	0	0					

Part I	=
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Books	NA
Articles	NA
References Books	NA
MOOC Courses	NA
Videos	NA

Course Articulation Matrix

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COs	PO1	PO2	PO3	PO4	PO5	PO6	PO7	PO8	PO9	PO10	PO11	PO12	PSO1	PSO2	PSO3
CO1	1	3	2	2	1	-	-	-	-	-	3	-	1	1	3
CO2	1	1	2	2	-	-	-	-	-	-	2	-	2	2	3
CO3	1	3	3	2	1	-	-	-	-	-	3	-	1	2	1
CO4	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO5	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
CO6	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-

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