

| S.NO | NAME OF THE PROGRAMME | NAME OF THE SUBJECT | | COURSE OUTCOMES | | |
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| | | PHARMACEUTICS | | | | |
| 1 | I M. Pharmacy I Semester | Modern Pharmaceutical Analytical Techniques | CO 1 | Chemicals and Excipients | | |
| | | | CO 2 | The analysis of various drugs in single and combination dosage forms | | |
| | | | CO 3 | Theoretical and practical skills of the instruments | | |
| 2 | | I M. Pharmacy I Semester | Drug delivery systems | CO 1 | The various approaches for development of novel drug delivery systems. | |
| | | | | CO 2 | The criteria for selection of drugs and polymers for the development of delivering system | |
| | | | | CO 3 | The formulation and evaluation of Novel drug delivery systems | |
| 3 | | | I M. Pharmacy I Semester | Modern Pharmaceutics | CO 1 | The elements of preformulation studies. |
| | | | | | CO 2 | The Active Pharmaceutical Ingredients and Generic drug Product development |
| | | | | | CO 3 | Industrial Management and GMP Considerations. |
| | | | | | CO 4 | Optimization Techniques & Pilot Plant Scale Up Techniques |
| | | | | | CO 5 | Stability Testing, sterilization process & packaging of dosage forms |
| 4 | | | | I M. Pharmacy I Semester | Regulatory affairs | CO 1 |

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| | | | CO 2 | The Regulatory guidance's and guidelines for filing and approval process |
| | | | CO 3 | Preparation of Dossiers and their submission to regulatory agencies indifferent countries |
| | | | CO 4 | Post approval regulatory requirements for actives and drug products |
| | | | CO 5 | Submission of global documents in CTD/ eCTD formats |
| | | | CO 6 | Clinical trials requirements for approvals for conducting clinical trials |
| | | | CO 7 | Pharmacovigilance and process of monitoring in clinical trials |
| 1 | I M. Pharmacy II Semester | Molecular pharmaceuticals (Nanotechnology & Targeted DDS) | CO 1 | The various approaches for development of novel drug delivery systems. |
| | | | CO 2 | The criteria for selection of drugs and polymers for the development of NTDS |
| | | | CO 3 | The formulation and evaluation of novel drug delivery systems. |
| 2 | | Advanced Biopharmaceutics & Pharmacokinetics | CO 1 | The basic concepts in biopharmaceutics and pharmacokinetics. |
| | | | CO 2 | The use raw data and derive the pharmacokinetic models and parameters the best describe the process of drug absorption, distribution, metabolism and elimination. |
| | | | CO 3 | The critical evaluation of biopharmaceutic studies |

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| | | | | involving drug product equivalency. |
| | | | CO 4 | The design and evaluation of dosage regimens of the drugs using pharmacokinetic and biopharmaceutic parameters. |
| | | | CO 5 | The potential clinical pharmacokinetic problems and application of basics of pharmacokinetic |
| 3 | | Computer Aided Drug Delivery System | CO 1 | History of Computers in Pharmaceutical Research and Development |
| | | | CO 2 | Computational Modeling of Drug Disposition |
| | | | CO 3 | Computers in Preclinical Development |
| | | | CO 4 | Optimization Techniques in Pharmaceutical Formulation |
| | | | CO 5 | Computers in Market Analysis |
| | | | CO 6 | Computers in Clinical Development |
| | | | CO 7 | Artificial Intelligence (AI) and Robotics |
| | | | CO 8 | Computational fluid dynamics(CFD) |
| 4 | | Cosmetics And Cosmeceuticals | CO 1 | Key ingredients used in cosmetics and cosmoceuticals. |
| | | | CO 2 | Key building blocks for various formulations. |
| | | | CO 3 | Current technologies in the market |

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| | | | CO 4 | Various key ingredients and basic science to develop cosmetics and cosmeceuticals |
| | | | CO 5 | Scientific knowledge to develop cosmetics and cosmeceuticals with desired Safety, stability, and efficacy. |
| PHARMACOLOGY | | | | |
| 1 | I M. Pharmacy I Semester | Modern Pharmaceutical Analytical Techniques | CO 1 | Chemicals and Excipients |
| | | | CO 2 | The analysis of various drugs in single and combination dosage forms |
| | | | CO 3 | Theoretical and practical skills of the instruments |
| 2 | | Advanced Pharmacology - I | CO 1 | Discuss the pathophysiology and pharmacotherapy of certain diseases |
| | | | CO 2 | Explain the mechanism of drug actions at cellular and molecular level |
| | | | CO 3 | Understand the adverse effects, contraindications and clinical uses of drugs used in treatment of diseases |
| 3 | | Pharmacological And Toxicological Screening Methods - I | CO 1 | Appraise the regulations and ethical requirement for the usage of experimental animals. |
| | | | CO 2 | Describe the various animals used in the drug discovery process and good laboratory practices in maintenance and handling of experimental animals |
| | | | CO 3 | Describe the various newer screening methods involved in the drug discovery process |

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| | | | CO 4 | Appreciate and correlate the preclinical data to humans |
| 4 | | Cellular and Molecular Pharmacology | CO 1 | Explain the receptor signal transduction processes. |
| | | | CO 2 | Explain the molecular pathways affected by drugs. |
| | | | CO 3 | Appreciate the applicability of molecular pharmacology and biomarkers in drug discovery process. |
| | | | CO 4 | Demonstrate molecular biology techniques as applicable for pharmacology |
| 1 | | Advanced Pharmacology - II | CO 1 | Explain the mechanism of drug actions at cellular and molecular level |
| | | | CO 2 | Discuss the Pathophysiology and pharmacotherapy of certain diseases |
| | | | CO 3 | Understand the adverse effects, contraindications and clinical uses of drugs used in treatment of diseases |
| 2 | I M. Pharmacy II Semester | Pharmacological and Toxicological Screening methods-II | CO 1 | Explain the various types of toxicity studies. |
| | | | CO 2 | Appreciate the importance of ethical and regulatory requirements for toxicity studies. |
| | | | CO 3 | Demonstrate the practical skills required to conduct the preclinical toxicity studies. |
| 3 | | Principles of Drug Discovery | CO 1 | Explain the various stages of drug discovery. |
| | | | CO 2 | Appreciate the importance of the role of genomics, proteomics and bioinformatics in drug discovery |

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| | | | CO 3 | Explain various targets for drug discovery. |
| | | | CO 4 | Explain various lead seeking method and lead optimization |
| | | | CO 5 | Appreciate the importance of the role of computer aided drug design in drug discovery |
| 4 | | Clinical Research And Pharmacovigilance | CO 1 | Explain the regulatory requirements for conducting clinical trial |
| | | | CO 2 | Demonstrate the types of clinical trial designs |
| | | | CO 3 | Explain the responsibilities of key players involved in clinical trials |
| | | | CO 4 | Execute safety monitoring, reporting and close-out activities |
| | | | CO 5 | Explain the principles of Pharmacovigilance |
| | | | CO 6 | Detect new adverse drug reactions and their assessment |
| | | | CO 7 | Perform the adverse drug reaction reporting systems and communication in Pharmacovigilance |
| PHARMACEUTICAL CHEMISTRY | | | | |
| 1 | I M. Pharmacy I Semester | Modern Pharmaceutical Analytical Techniques | CO 1 | The analysis of various drugs in single and combination dosage forms |
| | | | CO 2 | Theoretical and practical skills of the instruments |
| 2 | | Advanced Organic Chemistry - I | CO 1 | The principles and applications of reterosynthesis |
| | | | CO 2 | The mechanism & applications of various named reactions |

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| | | | CO 3 | The concept of disconnection to develop synthetic routes for small target molecule. |
| | | | CO 4 | The various catalysts used in organic reactions |
| | | | CO 5 | The chemistry of heterocyclic compounds |
| 3 | | Advanced Medicinal Chemistry | CO 1 | Different stages of drug discovery |
| | | | CO 2 | Role of medicinal chemistry in drug research |
| | | | CO 3 | Different techniques for drug discovery |
| | | | CO 4 | Various strategies to design and develop new drug like molecules for biological targets |
| | | | CO 5 | Peptidomimetics |
| 4 | | Chemistry of Natural Products | CO 1 | Different types of natural compounds and their chemistry and medicinal importance |
| | | | CO 2 | The importance of natural compounds as lead molecules for new drug discovery |
| | | | CO 3 | The concept of rDNA technology tool for new drug discovery |
| | | | CO 4 | General methods of structural elucidation of compounds of natural origin |
| | | | CO 5 | Isolation, purification and characterization of simple chemical constituents from natural source |
| 1 | I M. Pharmacy II Semester | Advanced Spectral Analysis | CO 1 | Interpretation of the NMR, Mass and IR spectra of various organic compounds |

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| | | | CO 2 | Theoretical and practical skills of the hyphenated instruments |
| | | | CO 3 | Identification of organic compounds |
| 2 | | Advanced Organic Chemistry - II | CO 1 | The principles and applications of Green chemistry |
| | | | CO 2 | The concept of peptide chemistry. |
| | | | CO 3 | The various catalysts used in organic reactions |
| | | | CO 4 | The concept of stereochemistry and asymmetric synthesis. |
| 3 | | Computer Aided Drug Design | CO 1 | Role of CADD in drug discovery |
| | | | CO 2 | Different CADD techniques and their applications |
| | | | CO 3 | Various strategies to design and develop new drug like molecules. |
| | | | CO 4 | Working with molecular modeling software's to design new drug molecules |
| 4 | | Pharmaceutical Process Chemistry | CO 1 | The strategies of scale up process of APIs and intermediates |
| | | | CO 2 | The various unit operations and various reactions in process chemistry |
| PHARMACEUTICAL ANALYSIS & QUALITY ASSURANCE | | | | |
| 1 | I M. Pharmacy I Semester | Modern Pharmaceutical Analytical Techniques | CO 1 | Chemicals and Excipients |
| | | | CO 2 | The analysis of various drugs in single and combination dosage forms |
| | | | CO 3 | Theoretical and practical skills of the instruments |
| 2 | | | CO 1 | The importance of quality |

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| | | Quality Management Systems | CO 2 | ISO management systems |
| | | | CO 3 | Tools for quality improvement |
| | | | CO 4 | Analysis of issues in quality |
| | | | CO 5 | Quality evaluation of pharmaceuticals |
| | | | CO 6 | Stability testing of drug and drug substances |
| | | | CO 7 | Statistical approaches for quality |
| 3 | | | Quality Control and Quality Assurance | CO 1 |
| | | CO 2 | | To appreciate the importance of documentation |
| | | CO 3 | | To understand the scope of quality certifications applicable to Pharmaceutical industries |
| | | CO 4 | | To understand the responsibilities of QA & QC departments. |
| 4 | | Audits And Regulatory Compliance | CO 1 | To understand the importance of auditing |
| | | | CO 2 | To understand the methodology of auditing |
| | | | CO 3 | To carry out the audit process |
| | | | CO 4 | To prepare the auditing report |
| | | | CO 5 | To prepare the check list for auditing |
| 1 | I M. Pharmacy II Semester | Hazards and Safety Management | CO 1 | Understand about environmental problems among learners. |
| | | | CO 2 | Impart basic knowledge about the environment and its allied problems. |
| | | | CO 3 | Develop an attitude of concern for the industry environment. |

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| | | | CO 4 | Ensure safety standards in pharmaceutical industry |
| | | | CO 5 | Provide comprehensive knowledge on the safety management |
| | | | CO 6 | Empower an ideas to clear mechanism and management in different kinds of hazard management system |
| | | | CO 7 | Teach the method of Hazard assessment, procedure, methodology for provide safe industrial atmosphere. |
| 2 | | Pharmaceutical Validation | CO 1 | The concepts of calibration, qualification and validation |
| | | | CO 2 | The qualification of various equipments and instruments |
| | | | CO 3 | Process validation of different dosage forms |
| | | | CO 4 | Validation of analytical method for estimation of drugs |
| | | | CO 5 | Cleaning validation of equipments employed in the manufacture of pharmaceuticals |
| 3 | | Advanced Pharmaceutical Analysis | CO 1 | Appropriate analytical skills required for the analytical method development. |
| | | | CO 2 | Principles of various reagents used in functional group analysis that renders necessary support in research methodology and demonstrates its application in the practical related problems. |
| | | | CO 3 | Analysis of impurities in drugs, residual solvents and stability |

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| | | | | studies of drugs and biological products |
| 4 | | Modern Bio-Analytical Techniques | CO 1 | Extraction of drugs from biological samples |
| | | | CO 2 | Separation of drugs from biological samples using different techniques |
| | | | CO 3 | Guidelines for BA/BE studies. |
| PHARMACEUTICAL ANALYSIS | | | | |
| 1 | I M. Pharmacy I Semester | Modern Pharmaceutical Analytical Techniques | CO 1 | The analysis of various drugs in single and combination dosage forms |
| | | | CO 2 | Theoretical and practical skills of the instruments |
| 2 | | Advanced Pharmaceutical Analysis | CO 1 | Appropriate analytical skills required for the analytical method development. |
| | | | CO 2 | Principles of various reagents used in functional group analysis that renders necessary support in research methodology and demonstrates its application in the practical related problems. |
| | | | CO 3 | Analysis of impurities in drugs, residual solvents and stability studies of drugs and biological products |
| 3 | | Pharmaceutical Validation | CO 1 | Explain the aspect of validation |
| | | | CO 2 | Carryout validation of manufacturing processes |
| | | | CO 3 | Apply the knowledge of validation to instruments and equipments |

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| | | | CO 4 | Validate the manufacturing facilities |
| 4 | | Food Analysis | CO 1 | Food constituents |
| | | | CO 2 | Food additives |
| | | | CO 3 | Finished food products |
| | | | CO 4 | Pesticides in food |
| | | | CO 5 | And also student shall have the knowledge on food regulations and legislations |
| 1 | | Advanced Instrumental Analysis | CO 1 | Interpretation of the NMR, Mass and IR spectra of various organic compounds |
| | | | CO 2 | Theoretical and practical skills of the hyphenated instruments |
| | | | CO 3 | Identification of organic compounds |
| 2 | I M. Pharmacy II Semester | Modern Bio-Analytical Techniques | CO 1 | Extraction of drugs from biological samples |
| | | | CO 2 | Separation of drugs from biological samples using different techniques |
| | | | CO 3 | Guidelines for BA/BE studies. |
| 3 | | Quality Control And Quality Assurance | CO 1 | The cGMP aspects in a pharmaceutical industry |
| | | | CO 2 | To appreciate the importance of documentation |
| | | | CO 3 | To understand the scope of quality certifications applicable to Pharmaceutical industries |
| | | | CO 4 | To understand the responsibilities of QA & QC departments |

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| 4 | | Herbal And Cosmetic Analysis | CO 1 | Determination of herbal remedies and regulations |
| | | | CO 2 | Analysis of natural products and monographs |
| | | | CO 3 | Determination of Herbal drug-drug interaction |
| | | | CO 4 | Principles of performance evaluation of cosmetic products. |
| PHARMACEUTICAL REGULATORY AFFAIRS | | | | |
| 1 | I M. Pharmacy I Semester | Good Regulatory Practices | CO 1 | The key regulatory and compliance elements with respect to Good Manufacturing Practices, Good Laboratory Practices, Good Automated Laboratory Practices and Good Documentation Practices. |
| | | | CO 2 | Prepare and implement the check lists and SOPs for various Good Regulatory Practices |
| | | | CO 3 | Implement Good Regulatory Practices in the Healthcare and related Industries |
| | | | CO 4 | Prepare for the readiness and conduct of audits and inspections. |
| 2 | | Documentation And Regulatory Writing | CO 1 | Know the various documents pertaining to drugs in pharmaceutical industry |
| | | | CO 2 | Understand the basics of regulatory compilation |
| | | | CO 3 | Create and assemble the regulation submission as per the requirements of agencies |

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| | | | CO 4 | Follow up the submissions and post approval document requirements |
| 3 | | Clinical Research Regulations | CO 1 | History, origin and ethics of clinical and biomedical research and evaluation |
| | | | CO 2 | Clinical drug, medical device development process and different types and phases of clinical trials |
| | | | CO 3 | Regulatory requirements and guidance for conduct of clinical trials and research |
| 4 | | Regulations And Legislation For Drugs & Cosmetics, Medical Devices, Biologicals & Herbals, And Food & Nutraceuticals In India And Intellectual Propertyrights | CO 1 | Know different Acts and guidelines that regulate Drugs & Cosmetics |
| | | | CO 2 | Understand the approval process and regulatory requirements for Drugs & Cosmetics |
| 1 | I M. Pharmacy II Semester | Regulatory Aspects Of Drugs & Cosmetics | CO 1 | Process of drug discovery and development and generic product development |
| | | | CO 2 | Regulatory approval process and registration procedures for API and drug products in US, EU |
| | | | CO 3 | Cosmetics regulations in regulated and semi-regulated countries |
| | | | CO 4 | A comparative study of India with other global regulated markets |
| 2 | | Regulatory Aspects Of Herbal And Biologicals | CO 1 | Know the regulatory Requirements for Biologics and Vaccines |

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| | | | CO 2 | Understand the regulation for newly developed biologics and biosimilars |
| | | | CO 3 | Know the pre-clinical and clinical development considerations of biologics |
| | | | CO 4 | Understand the Regulatory Requirements of Blood and/or Its Components Including Blood Products and label requirements |
| 3 | | Regulatory Aspects Of Medical Devices | CO 1 | Basics of medical devices and IVDs, process of development, ethical and quality considerations |
| | | | CO 2 | Harmonization initiatives for approval and marketing of medical devices and IVDs |
| | | | CO 3 | Regulatory approval process for medical devices and IVDs in India, US, Canada, EU, Japan and ASEAN |
| | | | CO 4 | Clinical evaluation and investigation of medical devices and IVDs |
| 4 | | Regulatory Aspects Of Food & Nutraceuticals | CO 1 | Know the regulatory Requirements for nutraceuticals |
| | | | CO 2 | Understand the regulation for registration and labeling of nutraceuticals and food supplements in India, USA and Europe. |