



Course Outcomes: M.Pharm.

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ORIENTAL COLLEGE OF PHARMACY

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M.Pharm. Q.A. Course Outcomes Semester I

| | | |
|---|--|--|
| MQA 101T | After completion of this course students should be able to: | |
| MODERN PHARMACEUTICAL ANALYTICAL TECHNIQUES | MQA 101T.1 | Know the analysis of various drugs in single & combination dosage forms |
| | MQA 101T.2 | Learn Theoretical & Practical skills of instruments |
| | MQA 101T.3 | UV-Visible Spectroscopy, IR Spectroscopy, Spectrofluorometric, AAS |
| | MQA 101T.4 | NMR Spectroscopy : Quantum number, Principle, Theory, Instrumentation |
| | MQA 101T.5 | Mass Spectroscopy : Principle, Theory, Instrumentation |
| | MQA 101T.6 | Chromatography : Principle, Apparatus, Instrumentation, Types |
| | MQA 101T.7 | Electrophoresis : Principle, Theory, Factors affecting separation |
| | MQA 101T.8 | Regulatory Compliance through Quality Management & 4Hrs development of Quality |
| MQA 102T | After completion of this course students should be able to understand: | |
| QUALITY MANAGEMENT SYSTEMS | MQA 102T.1 | Importance of Quality |
| | MQA 102T.2 | Pharmaceutical Quality Management : ISO management systems |
| | MQA 102T.3 | Six System Inspection Model- Tools for quality improvement |
| | MQA 102T.4 | Analysis of issues in quality and stability testing |
| | MQA 102T.5 | Statistical approaches for quality |
| MQA 103T | After completion of this course students should be able to: | |
| QUALITY CONTROL AND QUALITY ASSURANCE | MQA 103T.1 | Understand the cGMP aspects in a Pharmaceutical industry |
| | MQA 103T.2 | appreciate the importance of documentation |
| | MQA 103T.3 | Analysis of raw materials, finished products, packaging materials |
| | MQA 103T.4 | understand the scope of quality certifications applicable to Pharmaceutical industries |
| | MQA 103T.5 | Understand the responsibilities of QA&QC departments. |

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**M.Pharm. Q.A. Course Outcomes Semester I**

| MQA 104T | After completion of this course students should be able to: | |
|---|---|---|
| PRODUCT DEVELOPMENT AND TECHNOLOGY TRANSFER | MQA 104T.1 | Understand new product development process |
| | MQA 104T.2 | Understand necessary information to transfer technology from R&D to Manufacturing |
| | MQA 104T.3 | Understand Principles of drug discovery and Development |
| | MQA 104T.4 | Understand Pre-formulation studies |
| | MQA 104T.5 | Know Pilot plant scale up |
| | MQA 104T.6 | Know Pharmaceutical Packaging |
| | MQA 104T.7 | Technology Transfer : Development of Technology by R&D, Documentation |
| MQA 105P | MQA 105P.1 | Analyse Pharmacopoeial compounds in bulk and in their formulations |
| QUALITY ASSURANCE PRACTICAL - I | MQA 105P.2 | Perform experiments based on HPLC GC, UV, AAS |
| | MQA 105P.3 | Develop stability study protocols |
| | MQA 105P.4 | Carry out preformulation studies, accelerated studies and QC tests |

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**M.Pharm. Q.A. Course Outcomes Semester II**

| MQA 201T | After completion of this course students should be able to understand: | |
|----------------------------------|---|---|
| Hazard and safety Management | MQA 201T.1 | environmental problems among learners |
| | MQA 201T.2 | Multidisciplinary nature of environmental studies |
| | MQA 201T.3 | Air based hazards |
| | MQA 201T.4 | Chemical based hazards |
| | MQA 201T.5 | Fire and Explosion |
| | MQA 201T.6 | Hazard and Risk Management |
| MQA 202T | After completion of this course students should be able to understand : | |
| Pharmaceutical Validation | MQA 202T.1 | concept of calibration and validation |
| | MQA 202T.2 | Validation of instruments |
| | MQA 202T.3 | Qualification of Manufacturing Instruments like LC-MS, HPLC, HPTLC etc |
| | MQA 202T.4 | Qualification of Laboratory Equipment: Disintegration Test |
| | MQA 202T.5 | Process Validation |
| | MQA 202T.6 | Cleaning Validation |
| | MQA 202T.7 | General Principle of Intellectual Property |
| MQA 203T | After completion of this course students should be able to: | |
| Audits and regulatory compliance | MQA 203T.1 | Understand the importance of auditing and their methods |
| | MQA 203T.2 | Know Objective Planning Process and prepare audit report |
| | MQA 203T.3 | Understand Role of quality systems and audits in pharmaceutical manufacturing environment |
| | MQA 203T.4 | Know Auditing of vendors and production department |
| | MQA 203T.5 | Know Auditing of Microbiological Laboratory |
| | MQA 203T.6 | Know Auditing of Quality Assurance Department |

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M.Pharm. Q.A. Course Outcomes Semester II

| MQA 204T | After completion of this course students should be able to understand: | |
|---|--|---|
| Pharmaceutical Manufacturing Technology | MQA 204T.1 | Pharmaceutical Industry Developments |
| | MQA 204T.2 | Aseptic process technology |
| | MQA 204T.3 | Non Sterile manufacturing process technology |
| | MQA 204T.4 | Containers and closures for pharmaceuticals, packaging technology |
| | MQA 204T.5 | Quality by design (QbD) and process analytical technology |


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**M.Pharm. Pharmacology Course Outcomes Semester I**

| MPL 101T | After completion of this course students should be able to learn: | |
|--|--|---|
| MODERN PHARMACEUTICAL ANALYTICAL TECHNIQUES | MPL 101T.1 | Theoretical & Practical skills of instruments |
| | MPL 101T.2 | UV-Visible Spectroscopy, IR Spectroscopy, Spectrofluorimetry, AAS |
| | MPL 101T.3 | NMR Spectroscopy : Quantum number, Principle, Theory, Instrumentation |
| | MPL 101T.4 | Mass Spectroscopy : Principle, Theory, Instrumentation |
| | MPL 101T.5 | Chromatography : Principle, Apparatus, Instrumentation, Types |
| | MPL 101T.6 | Electrophoresis : Principle, Theory, Factors affecting separation |
| | MPL 101T.7 | Potentiometry |
| MPL 102T | After completion of this course students should be able to understand: | |
| ADVANCED PHARMACOLOGY | MPL 102T.1 | General Pharmacology, Pharmacokinetics, Pharmacodynamics |
| | MPL 102T.2 | Neurotransmission, Systematic Pharmacology, Autonomic Pharmacology |
| | MPL 102T.3 | Central Nervous System Pharmacology |
| | MPL 102T.4 | Cardiovascular Pharmacology |
| | MPL 102T.5 | Autocoid Pharmacology |
| | MPL 102T.5 | |
| MPL 103T | After completion of this course students should be able to: | |
| PHARMACOLOGICAL AND TOXOLOGICAL SCREENING | MPL 103T.1 | Appraise the regulations and ethical requirement for the usage of experimental animals. |
| | MPL 103T.2 | Describe the various animals used in the drug discovery process and good laboratory practices in maintenance and handling of experimental animals |
| | MPL 103T.3 | Describe the various newer screening methods involved in the drug discovery process |
| | MPL 103T.4 | Appreciate and correlate the preclinical data to humans |
| | MPL 103T.5 | Understand Preclinical Screening of new substances for the pharmacological activity |

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M.Pharm. Pharmacology Course Outcomes Semester I

| MPL 104T | After completion of this course students should be able to: | |
|-------------------------------------|---|---|
| CELLULAR AND MOLECULAR PHARMACOLOGY | MPL 104T.1 | Explain the receptor signal transduction processes |
| | MPL 104T.2 | Explain the molecular pathways affected by drugs. |
| | MPL 104T.3 | Appreciate the applicability of molecular pharmacology and biomarkers in drug discovery process |
| | MPL 104T.4 | Demonstrate molecular biology techniques as applicable for pharmacology |

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**M.Pharm. Pharmacology Course Outcomes Semester II**

| MPL 201T | After completion of this course students should be able to: | |
|--|---|--|
| ADVANCED PHARMACOLOGY II | MPL201T.1 | Explain the mechanism of drug actions at cellular and molecular level |
| | MPL201T.2 | Understand Endocrine Pharmacology : Molecular and cellular mechanism of actions of hormones Chemotherapy : |
| | MPL201T.3 | Understand Chemotherapy : Drugs used in protozoal Infections |
| | MPL201T.4 | Understand GIT Pharmacology : Chemo pharmacology, Cardiovascular disease |
| | MPL201T.5 | Understand Free Radical Pharmacology: Generation of free radicals, recent advances in treatment |
| MPL 202T | After completion of this course students should be able to: | |
| PHARMACOLOGICAL AND TOXOLOGICAL SCREENING II | MPL 202T.1 | Explain the various types of toxicity studies |
| | MPL 202T.2 | Understand Acute, Subacute and chronic oral ,Acute eye irritation |
| | MPL 202T.3 | Understand Reproductive toxicology studies |
| | MPL 202T.4 | Understand IND enabling studies : Definition if IND Industry Perspective |
| | MPL 202T.5 | Understand Toxicokinetics : Evaluation in Preclinical Studies, Alternative method of Animal toxicity testing |
| MPL 203T | After completion of this course students should be able to Explain: | |
| PRINCIPLES OF DRUG DISCOVERY | MPL 203T.1 | various stages of drug discovery |
| | MPL 203T.2 | Lead Identification, Protein Structure |
| | MPL 203T.3 | Rational drug Design : Traditional Vs rational drug design |
| | MPL 203T.4 | Molecular docking : rigid docking, flexible docking |
| | MPL 203T.5 | QSAR Stastical Method : regression analysis, Prodrug design basic concept |

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| MPL 204T | After completion of this course students should be able to understand: | |
|---|--|--|
| CLINICAL RESEARCH AND PHARMACOVIGILANCE | MPL 204T.1 | Regulatory perspective of clinical trial |
| | MPL 204T.2 | Clinical trials |
| | MPL 204T.3 | Clinical trials documentation |
| | MPL 204T.4 | Basic aspects and terminologies and establishment of pharmacovigilance |
| | MPL 204T.5 | Methods, ADR reporting and tools used in pharmacovigilance |
| | MPL 204T.6 | Pharmacoeconomics, safety pharmacology |

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**M. Pharm Pharmaceutics Course Outcomes Semester I**

| | | |
|-----------------------|--|--|
| MPH 101T | After completion of this course students should be able to : | |
| MPAT | MPH 101T.1 | Have knowledge of Chemicals and Excipients |
| | MPH 101T.2 | The analysis of various drugs in single and combination dosage forms |
| | MPH 101T.3 | Theoretical and practical skills of the instruments |
| MPH 102T | After completion of this course students should be able to understand | |
| DRUG DELIVERY SYSTEMS | MPH 102T.1 | The various approaches for development of novel drug delivery systems. |
| | MPH 102T.2 | The criteria for selection of drugs and polymers for the development of delivering system |
| | MPH 102T.3 | The formulation and evaluation of Novel drug delivery systems |
| MPH 103T | After completion of this course students should be able to understand: | |
| MODERN PHARMACEUTICS | MPH 103T.1 | The elements of pre formulation studies. |
| | MPH 103T.2 | The Active Pharmaceutical Ingredients and Generic drug Product development |
| | MPH 103T.3 | Industrial Management and GMP Considerations. |
| | MPH 103T.4 | Optimization Techniques & Pilot Plant Scale Up Technique |
| | MPH 103T.5 | Stability Testing, sterilization process & packaging of dosage forms. |
| MPH 104T | After completion of this course students should be able to understand: | |
| REGULATORY AFFAIRS | MPH 104T.1 | The Concepts of innovator and generic drugs, drug development process |
| | MPH 104T.2 | The Regulatory guidance's and guidelines for filing and approval process |
| | MPH 104T.3 | Preparation of Dossiers and their submission to regulatory agencies in different countries |
| | MPH 104T.4 | Post approval regulatory requirements for actives and drug products Submission of global documents in CTD/ eCTD formats |
| | MPH 104T.5 | Clinical trials requirements for approvals for conducting clinical trials |
| | MPH 104T.6 | Pharmacovigilance and process of monitoring in clinical trials. |

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**M.Pharm. Pharmaceutics Course Outcomes Semester II**

| | | |
|-------------------------|---|---|
| MPH 201T | After completion of this course students should be able to understand | |
| MOLECULAR PHARMACEUTICS | MPH 201T.1 | The various approaches for development of novel drug delivery systems. |
| | MPH 201T.2 | The criteria for selection of drugs and polymers for the development of NTDS |
| | MPH 201T.3 | The formulation and evaluation of novel drug delivery systems |
| MPH 202T | After completion of this course students should be able to understand | |
| ADV. BPPK | MPH 202T.1 | The basic concepts in biopharmaceutics and pharmacokinetics |
| | MPH 202T.2 | The use raw data and derive the pharmacokinetic models and parameters the best describe the process of drug absorption, distribution, metabolism and elimination. |
| | MPH 202T.3 | The critical evaluation of biopharmaceutic studies involving drug product equivalency |
| | MPH 202T.4 | The design and evaluation of dosage regimens of the drugs using pharmacokinetic and biopharmaceutic parameters |
| | MPH 202T.5 | The potential clinical pharmacokinetic problems and application of basics of pharmacokinetic |
| MPH 203T | After completion of this course students should be able to understand | |
| CADD | MPH 203T.1 | Use of Computers in Pharmaceutical Research and Development |
| | MPH 203T.2 | Computational Modeling of Drug Disposition |
| | MPH 203T.3 | Computers in Preclinical Development |
| | MPH 203T.4 | Optimization Techniques in Pharmaceutical Formulation |
| | MPH 203T.5 | Computers in Market Analysis |
| | MPH 203T.6 | Computers in Clinical Development |
| | MPH 203T.7 | Artificial Intelligence (AI) and Robotics |
| | MPH 203T.8 | Computational fluid dynamics(CFD) |

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M.Pharm. Pharmaceutics Course Outcomes Semester II

| MPH 204T | After completion of this course students should be able to know: | |
|------------------------------|--|---|
| COSMETICS AND COSMECEUTICALS | MPH 204T.1 | Key ingredients used in cosmetics and cosmeceuticals |
| | MPH 204T.2 | Key building blocks for various formulations |
| | MPH 204T.3 | Current technologies in the market |
| | MPH 204T.4 | Various key ingredients and basic science to develop cosmetics and cosmeceuticals |
| | MPH 204T.5 | Scientific knowledge to develop cosmetics and cosmeceuticals with desired Safety, stability, and efficacy |

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