QP Code:	Reg. No.:				

# M Pharm Degree (Part I) Examinations

(Model Question Paper)

# **Modern Analytical and Research Methods**

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- 1. What is the principle and instrumentation of FTIR. Write a note on sample handling in IR spectroscopy and give its applications.. (10+10=20)
- 2. Write the principle of NMR spectroscopy. Briefly explain spin-spin coupling and decoupling techniques used. Give applications of NMR spectroscopy. Write the significance of C<sup>13</sup> NMR spectroscopy in the structural elucidation of organic compounds (15+5=20)

Short Essays: (6x10=60)

- 3. Compare GC with HPLC. Briefly discuss on derivatisation methods used in GC (5+5=10)
- 4. Write on detectors used in HPLC with neat diagram and mention the pharmaceutical applications of HPLC. (10)
- 5. Discuss the choice of solvents and solvent effects in UV spectroscopy. Factors influencing fluorescence intensity. (5+5=10)
- 6. Discuss the principle instrumentation and applications of X-ray powder diffraction technique. (10)
- 7. Give the theoretical principle of mass spectroscopy with the aid of neat diagram of double focussing mass spectrophotometer. Write a brief note on hyphenation of GC & LC with MS. (7+3=10)
- 8. Give the significance of students T- test, ANOVA, regression analysis and correlation coefficient. (10)

QP Code:	Reg. No.:
	J -

(Model Question Paper)

# Paper I - Analytical Techniques and Instrumentation

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- 1. What is the principle of flourimetry. Factors influencing fluorescence intensity. Write a note on applications of flourimetry. Explain with examples fluorescent immunoassay
- 2. Define chemical shift. Briefly explain the factors influencing chemical shift values. Write a note on 2D -NMR techniques. Enumerate various applications of NMR spectroscopy.

Short Essays: (6x10=60)

- 3. Derive Beer Lamberts law and what are the limitations of this law. Describe the applications of UV- VIS spectroscopy. (7+3=10)
- 4. Explain in detail the general fragmentation patterns for the interpretation of organic compounds in mass spectroscopy. Write a brief note on MALDI. (5+5=10)
- 5. Discuss the principle, instrumentation and applications of HPLC.
- 6. Discuss the theory, instrumentation and applications of atomic absorption spectroscopy
- 7. Write a note on factors influencing vibrational frequencies and its applications.
- 8. Explain the detectors used in GC with neat labelled diagram.

QP Code:	Reg. No.:

(Model Question Paper)

# Paper II – Advanced Pharmaceutical Analysis

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- 1. Explain about validation and calibration of a HPLC. Describe the quality control tests for tablets and parentrals.
- 2. Explain the following: (5x4=20)
  - QC of Hair care products.
- Toxicity testing of cosmetics

Particle size analysis

Assay of rabies vaccines

Short Essays: (6x10=60)

- 3. Write the methods for analyzing various carbohydrates in foods. What are the preservatives used in food products. Explain the estimation of any one of them. (5+5=10)
- 4. Explain the official methods for the determination of the following dosage forms: (2½x4=10)
  - Chloramphenicol tablets.
- Ascorbic acid tablets

Phenobarbitone tablets

- Digoxin tablets
- 5. Describe the radioimmunoassay and radiotracer techniques used in pharmaceutical analysis.
- 6. Explain the principles and procedures for the use of reagent MBTH in pharmaceutical analysis.
- 7. Write a detailed study of principle and procedure involved in various physico chemical methods of analysis of sulpha drugs.
- 8. Enlist the different applications of instrumental methods in the development and quality control of drugs.

QP Code:	Reg. No.:
	J -

(Model Question Paper)

#### Paper III - Quality Assurance

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- 1. Write the Concepts and Philosophy of TQM. Write about the Master formula records. Write note on manufacturing documents.
- 2. Classify packaging materials and tests to ensure the quality of secondary packaging materials. Explain the pharmacopoeial tests for various glass containers.

Short Essays: (6x10=60)

- 3. Write a note on Good Warehousing Practices. How the sanitation and sterile areas are maintained in pharma industry.
- 4. Write brief note on Organization and personnel in a pharma company.
- 5. Write a note on concepts and philosophy of GMP.
- 6. What are the standard operating procedures for cleaning, drying and sterilization.
- 7. What are the procedures required for evaluation of complaints and recall of distributed finished products.
- 8. Give an account on quality audits .What are the protocols to be followed in selecting vendors. Add a note on purchase, receipt, storage and release of raw materials.

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(Model Question Paper)

# Paper IV – Clinical Chemistry and Toxicological Analysis

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

1. Explain the determination of the following:

(4x5=20)

Blood glucose

SGOT & SGPT

Serum cholesterol

- Serum alkaline phosphatase
- 2. Write in detail on DOPE tests and methods for the estimation of anabolic steroids and drugs of abuse from biological samples.

Short Essays: (6x10=60)

- 3. Write a note on TDM and its application. Estimation of Carbamazepine from blood
- 4. Write methods for extraction of drugs from biological samples.
- 5. Enumerate screening procedure –spot tests and the use of TLC, GC and HPTLC in the identification of poisons.
- 6. Explain the analytical procedures for the estimation of the following:
  - Theophylline

- Phenytoin
- 7. Clinical correlation and significance of abnormal values of biochemical constituents
- 8. Write a note on:
  - Detoxification pathway of Phenobarbitone.
  - Detection of organophosphorus poisoning.
  - Name the antidotes for Heavy metal poisoning
  - Biochemical role of Choline esterase.

QP Code:	Reg. No.:

(Model Question Paper)

#### Paper I - Drug Design

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- 1. What is meant by Analog deign. Explain analog approach of drug design. what is Hansch analysis. How it is useful in drug design. Monte Caro method of conformational analysis.
- 2. Explain methods of parallel and mixed combinatorial synthesis. Write applications of combinatorial chemistry in drug discovery. Explain different types of molecular graphics in molecular modeling.

Short Essays: (6x10=60)

- 3. Discuss energy components for inter molecular non covalent interactions with suitable examples.
- 4. Explain applications of recombinant DNA technology in medicinal chemistry.
- 5. What is topless decision tree analysis. Explain peptidomimetics in drug design.
- 6. Define antisense technology. How antisense oligonucleotides are used in drug design.
- 7. Explain in detail about phase I and phase II metabolic reactions.
- 8. Explain about conventional methods of drug design

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QP Code:	Reg. No.:
	J -

(Model Question Paper)

# Paper II – Advanced Medicinal Chemistry

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- 1. Discuss recent advances in cancer therapy. Classify anti viral drugs. Explain the mechanism of action and synthesis of one drug from two different classes.
- 2. Give classification mechanism of action and SAR of anti hypertensive agents. Drugs used in neuro degenerative disorders.

Short Essays: (6x10=60)

- 3. Explain how radio sensitizers are used in drug therapy
- 4. Discuss the agents used in management of tuberculosis. Explain the concept of multiple drug resistance in tuberculosis.
- 5. Discuss the chemistry of β lactam antibiotics. Steroidal anti inflammatory agents
- 6. Explain about anti hyperlipidemic agents. Give synthesis of any two drugs.
- 7. What are tranquillizers . Give SAR of phenothiazine derivatives. give an account of thiazide diuretics.
- 8. Write the important classes of anti malaraial agents and give synthesis of amodiaquine and chloroquine

QP Code:	Reg. No.:

(Model Question Paper)

# Paper III – Advanced Organic Chemistry

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- 1. Explain localized and delocalized bonds with examples. Explain generation fate and biological significance of free radicals.
- 2. Write notes on (4X5=20)
  - Oppeneaur oxidation
- Wolf Kishner reduction

Birch reduction

Meerwin Pondroff's reduction

Short Essays: (6x10=60)

- 3. Discuss various methods of determining organic reaction mechanisms.
- 4. Discuss in detail the various mechanisms involved in the addition to carbon carbon multiple bonds.
- 5. Explain in detail about Retro synthetic analysis. Explain hyper conjugation with examples.
- 6. Discuss the phase transfer catalysis and its applications in reduction reactions.
- 7. Give a detail account of carbocations and carboanions.
- 8. Explain the mechanism of aromatic electrophilic substitution reaction. Write the basic theory of photochemical reactions and mention its applications.

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QP Code:	Reg. No.:
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(Model Question Paper)

#### Paper IV – Chemistry of Natural Products

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- 1. Define alkaloids . Give the methods of isolation of alakaloids. Elucidate the structure of quinine
- 2. Write the applications of IR, NMR and MASS Spectroscopy in the structural elucidation of natural products. Explain the importance of GLC and HPLC in separation

Short Essays: (6x10=60)

- 3. Explain the chemistry of :
  - Rutin
     Carotenes
- 4. Elucidate the structure of Cholestrol
- 5. Outline the synthesis:
  - Progesterone
     Reserpine
- 6. Write a note on role of natural products in new drug development. Explain the constitution of vitamin A.
- 7. Write in detail the role of recombinant DNA technology. Write about the isolation and characterization of important neutraceuticals.
- 8. Define terpenoids and elucidate the structure of camphor

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QP Code:	eg. No.:
M Pharm Degree (Part I) Examinations – Pharma	acognosy & Phytochemistry
(Model Question Paper)	
Paper I –Phytochemis	try
Time: 3 hrs	Maximum marks: 100
<ul> <li>Answer all questions</li> <li>Draw diagrams wherever necessary</li> </ul> Essays:	(2x20=40)
1. What are the steps involved in drug discovery process fro terms "Lead "," Hit " and " Activity Guided fractionation"	m natural products. Explain the
2. Explain the chemistry, methods of extraction, sources, ide flavanoids	ntification tests and uses of
Short Essays:	(6x10=60)
3. What is the extraction and evaluation technique of artimisis	in.
4. Give the methods of separation and identification of protein	ins.
5. What are the applications of docking studies with respect	to natural products Z.
6. What are the uses of NMR and Mass Spectroscopic meth	ods in identification of
phytochemicals.	
7. Explain the significance of alkaloids, fatty acids and volatil	le oils in chemotaxonomic studies.
8. Phytoestrogens	
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Q	P Code:	Reg. No.:		
N	M Pharm Degree (Part I) Examinations – Pharmacognosy & Phytochemistry			
		(Model Question Paper)		
		Paper II –Cultivation and Collection of Drugs		
	Time: 3 hrs	Maximum ma	rks: 100	
Es	• • • •	Answer all questions Draw diagrams wherever necessary	(2x20=40)	
1.	Explain the culti of cinchona.	vation methods including soil, fertilizers, irrigation and post har	vest treatment	
2.	What re Good A	agricultural Practices as guided by WHO.		
Sł	nort Essays:		(6x10=60)	
3.	Auxins and Gibb	perellins		
4.	Effect of Climate	e and soil moisture on the yield of phytoconstituents		
5.	Germplasm con	servation		
6.	Macro and micro	onutrients needed for cultivation of medicinal plants		
7.	Biopesticides			
8.	Different types of	of soil		
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Q	Reg. No.:	
M	M Pharm Degree (Part I) Examinations – Pharmacognosy & Phytoc	hemistry
	(Model Question Paper)	
	Paper III –Applied Pharmacognosy	
	Time: 3 hrs Maximum mark	s: 100
Es	<ul> <li>Answer all questions</li> <li>Draw diagrams wherever necessary</li> <li>ssays:</li> </ul>	(2x20=40)
1.	What re the problems encountered in the development and evaluation of herbal formulations.	
2.	What are the standards for asavas, arishtas, churnas and lehyas. Write about an medicinal plants used in Ayurveda.	y 5
Sł	hort Essays:	(6x10=60)
3.	Invitro screening for anti-inflammatory activity	
4.	Cardioactive toxicity of herbal drugs	
5.	DNA Fingerprinting	
6.	Methods of preparation and uses of phytosomes	
7.	Biomarkers in HPTLC standardization of herbals	
8.	Determination of aflotoxins in samples of crude drugs	
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Q	QP Code: Reg. No.:				
N	M Pharm Degree (Part I) Examinations – Pharmacognosy & Phytochemistry				
	(Model Question Paper)				
Paper IV –Medicinal Plant Biotechnology					
	Time: 3 hrs Maximum mark	s: 100			
Es	<ul> <li>Answer all questions</li> <li>Draw diagrams wherever necessary</li> <li>Essays:</li> </ul>	(2x20=40)			
1.	What are the different types of plant tissue culture methods				
2.	<ol><li>What are the genetic factors affecting the production of phytochemicals. Explain t gene mutation and gene transfer.</li></ol>	he terms			
SI	Short Essays: (6x10=60)				
3.	3. Role of elicitors and precursors in the production of phytochemicals				
4.	4. Biotransformation				
5.	5. Role of transgenic plants in production of medicines				
6.	6. DNA Recombinant technology				
7.	7. Bioreactors				
3.	8. Uses of PCR in gene mapping				
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#### M Pharm Degree (Part I) Examinations – Pharmaceutics

(Model Question Paper)

# Paper I – Formulation Technology

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- 1. Explain the design and formulation of sustained release tablets. Write the film coating process and the defects of film coated tablets.
- 2. Explain the Polymers used in controlled drug delivery. What are the various techniques used for microencapsulation. Explain phase separation Coacervation technique.

Short Essays: (6x10=60)

- 3. Explain Oral Osmotic pumps and Floating drug delivery systems
- 4. Explain the various mechanism of drug distribution in pulmonary drug delivery. Explain the design and development of inhalation drug delivery systems
- 5. Write an essay on the delivery of pharmaceutical peptides and proteins. Explain the problems in the delivery of peptides and proteins in conventional form.
- 6. Give a brief account on the following: Applications of nanoparticles and Concepts of physical drug targeting.
- 7. Explain any two types of parenteral controlled drug delivery system. Explain the factors that affect the release of drug from its delivery system
- 8. Explain the design and various types of transdermal drug delivery systems. Explain the evaluation of transdermal delivery system.

QP Code:	Reg. No.:
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(Model Question Paper)

# Paper II – Biopharmaceutics and Pharmacokinetics

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- 1. Define the terms bioavailability and bioequivalence. Discuss selection criteria of volunteers in bioavailability and bioequivalence studies. Why the bioavailability studies are carried out in healthy subjects. Explain any one method of determination of bioavailability
- 2. Explain flip –flop model. What are the various methods for estimation of absorption rate constant. Explain the method of residuals.

Short Essays: (6x10=60)

- 3. What is meant by first -pass metabolism. Explain its clinical significance. Explain the Biopharmaceutics of intramuscular injection.
- 4. Discuss the methods of dose adjustment in renal impairment. Explain the need of short term i.v infusions.
- 5. Explain Wagner Nelson method. Write about its merits and demerits.
- 6. Explain pharmacokinetic model and their objectives. Discuss assumptions of 'one compartment model'.
- 7. What is capacity limited kinetics. Explain the causes of non-linearity. Discus the application of Michaelis-Menton equation in non-linearity.
- 8. Discuss various biopharmaceutical factors affecting drug absorption from an injectable. Explain sigma –minus method and its limitation.

QP Code:	Reg. No.:
	J -

(Model Question Paper)

# Paper III - Industrial Microbiology and Biotechnology

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- 1. Describe the production, harvest and recovery of lactic acid by fermentation
- 2. Give examples of toxoids . Explain the production and standardization of Diphtheria toxoid.

Short Essays: (6x10=60)

- 3. What are gene libraries. Give examples of pharmaceuticals produced by recombinant DNA technology
- 4. What are monoclonal antibodies. Explain the term 'Cloning'
- 5. Give examples of viral vaccines. Explain the production of a viral vaccine by tissue culture method
- 6. Define the term fermentation. Describe the production of ethanol by fermentation
- 7. How aseptic processes are validated. Explain how the efficiency of air filters for sterilization is tested.
- 8. What is an allergy. List out foods, which are recognized to produce allergic reactions.

# M Pharm Degree (Part I) Examinations – Pharmaceutics

(Model Question Paper)

# Paper IV – Industrial Pharmacy

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- 1. Write a note on preformulation studies with special reference to polymorphism and solubility.. Write a note on stability of drugs in preformulation study.
- 2. What is called as CGMP. Explain Good manufacturing practices (GMP) in the quality control of parenteral products

Short Essays: (6x10=60)

- 3. Write notes on:. Optimization parameters, Simplex method
- 4. Explain the various types of industrial hazards. Explain industrial effluent testing procedures.
- 5. Write in detail about the elements of cost. Write in detail about the Revocation of patents.
- 6. Write notes on: cost control, ISO 9000 series
- 7. Explain the factors to be considered for pilot plant scale up. Explain the requirements of New Drug Application (NDA).
- 8. Explain the following terms: Inventory control, Materials management.

# M Pharm Degree (Part I) Examinations – Pharmacology

(Model Question Paper)

# Paper I–Pharmacological Screening Methods & Clinical Evaluation Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- 1. Discuss the pharmacological models employed in the screening of new drugs belonging to the following categories (In-vivo & In- vitro models) (10+10=20)
  - Nootropics
- Anxiolytics
- 2. Discuss in detail about the cell culture & cell line techniques about the (10+6+4=20)
  - Types, propagation & preservation of cultures
     Design, equipments for cell culture
     Application of cell cultures

Short Essays: (6x10=60)

- 3. Briefly explain the CPCSEA guidelines for laboratory animal facility.
- 4. Explain the principle, methods and applications of ELISA
- 5. Discuss the different models of anti-inflammatory agents screening procedure.
- 6. Explain the applicability of Analysis of Variance (ANOVA) with reference to biomedical research.
- 7. Write down the procedure involved in the following
  - Streptozotocin induced hyperglycemia
     NSAIDs induced ulcer model
- 8. Explain the drug discovery approaches in

(5+5=10)

- High throughput screening
- Combinatorial chemistry

# M Pharm Degree (Part I) Examinations – Pharmacology

(Model Question Paper)

# Paper II-Biochemical & Molecular Pharmacology

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

**Essays:** (2x20=40)

- 1. Explain the following in molecular approach (10+10=20)
  - Acute & Chronic inflammation Apoptosis
- 2. Discuss in detail of the following: (10+0=20)
  - Replication & transcription of DNA & RNA
     Gene therapy of genetic disorders

(6x10=60)**Short Essays:** 

- 3. Explain the biological functions & pharmacological implications of Nitric oxide(NO)
- 4. Write short notes on the following: (3+3+4=10)
  - Neuropeptide Y (NPY)
- Multidrug resistance (mtr) proteins
- Tissue Necrosis factor (TNF)
- 5. Write notes on the following techniques.

(4+3+3=10)

- Polymerase chain reaction (PCR)
   Southern blotting

- Northern blotting
- 6. What do you mean by gene cloning. Write short notes on transgenic animals & their (3+5+2=10)application.
- 7. Write down the biochemical pathway of reactive oxygen species & anti- oxidant defense mechanisms. (5+5=10)
- 8. Explain the role of pharmacogenetics with respect to drug action. What do you mean by chronopharmacology. (8+2=10)

QP Code:	Reg. No.:			
M Pharm Deg	ree (Part I) Examinations – Pharmacology			
(Model Question Paper)  Paper III – Recent Advances in Pharmacology				
<ul><li>Answer al</li><li>Draw diag</li><li>Essays:</li></ul>	questions rams wherever necessary (2x20=40)			
	hannel Receptors. Classify it . Explain the transduction pathway el with neat diagram. Write down their therapeutic importance.			
Recent approac	(10+7+3=20) I & humoral mediated immunity with flow diagram nes in drug discovery of AIDS ors of Indigenous origin			
Short Essays:	(6x10=60)			
3. Write short notes on the following short notes of the following short notes on the following short notes of the following short n	,			
	tem cells and their potential in various disorders. (5+5=10) chanism of anti-microbial resistance. What do you mean by			
<ul><li>6. Write short notes on the fo</li><li>Forces involved in</li></ul>	lowing: (5+5=10) D-R binding • Purinoceptors			
<ul><li>7. Write down the methods of target for gene therapy.</li><li>8. Write short notes on the following Nutraceuticals</li></ul>	gene transfer technologies (Viral & non Viral). What are disease (7+3=10) owing:  • Anti sense agents			

#### M Pharm Degree (Part I) Examinations – Pharmacology

(Model Question Paper)

# Paper IV – Clinical Pharmacology & Toxicology

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- 1. Discuss in detail about clinical research in the following: (10+10=20)
  - Design and ethical guidelines in clinical research
  - Phases of clinical research as per ICMR guidelines.
- 2. Discuss the types , methods & importance of the: (10+10=20)
  - Pharmacoeconomics
     Rational drug use

Short Essays: (6x10=60)

- 3. Discuss the Pathophysiology and pharmacotherapy of diabetes.
- 4. Write short notes on the following:
  - Helsinki declaration
     Stakeholders in clinical research
- 5. What do you mean by Pharmacovigilance. Write down the protocol followed in it and mention the importance of Pharmacovigilance in clinical practice. (2+6+2=10)
- 6. Write short notes on the following poison management: (5+5=10)
  - Organophosphorous compounds
     Methanol
- 7. Describe the pathophysiology and pharmacotherapy of Rheumatoid arthritis.
- 8. Discuss the importance of dose adjustments in renal & hepatic failure.

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(5+5=10)

QP Code:	Reg. No.:
	J -

#### M Pharm Degree (Part I) Examinations – Pharmacy Practice

(Model Question Paper)

#### Paper I – Clinical Pharmacy Practice

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- What are various markers of ischemic heart injury. How their levels change during ischemic injury. What factors are considered while analysing the patient data for case presentation.
   What are beneficial drug-drug interactions. Give atleast two examples with mechanism and benefits offered
- 2. What is Drug Use Evaluation. How is it carried out. Develop a protocol for carrying out a DUE for antimicrobial use in a tertiary care hospital.

Short Essays: (6x10=60)

- 3. Discuss the setting up of Drug information centre
- 4. Discuss the importance of medication history in therapeutic management of patients.
- 5. What are the types of medication errors, explain with suitable examples from each category
- 6. What is Non-compliance. Explain the causes, outcome and suggestions for improving the compliance
- 7. What are drug induced diseases. Explain their mechanism and management
- 8. What are the various methods of pharmacoeconomic evaluations, explain cost benefit analysis

QP Code:	Reg. No.:

#### M Pharm Degree (Part I) Examinations – Pharmacy Practice

(Model Question Paper)

# Paper II – Pathophysiology and Pharmacotherapeutics

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- 1. Explain the etiology and pathophysiology of Diabetes mellitus. Discuss the diabetes related complications. Discuss the therapeutic management options of diabetes mellitus.
- 2. Describe the various etiological factors responsible for acute renal failure. How to differentiate acute renal failure and chronic renal failure diagnostically. Discuss the supportive care and pharmacotherapeutic management of Acute renal failure.

Short Essays: (6x10=60)

- 3. Write the pharmacotherapeutic management of epilepsy in adult
- 4. Chart out and explain an algorithm for therapeutic management of rheumatoid arthritis
- 5. Discuss Cell cycle and the principles of cancer chemotherapy
- 6. Explain pain pathway. Discuss management of various types of pain using WHO pain ladder.
- 7. Present pathophysiology of Tuberculosis .Explain the latest antitubercular regimen for Tuberculosis
- 8. Write the JNC-7 classification of hypertension and explain with drug of choice in each stage

# M Pharm Degree (Part I) Examinations – Pharmacy Practice

(Model Question Paper)

# Paper III – Clinical Toxicology and Pharmacokinetics

Time: 3 hrs Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- 1. Describe the protocol, procedure and statistical interpretation of bioequivalence studies
- 2. "The Investigator's Brochure is important and constantly evolving document." Justify this statement. What is role and responsibilities of principal investigator as per ICH GCP guidelines. Describe briefly the content and format of NDA.

Short Essays: (6x10=60)

- 3. Discuss the importance of therapeutic drug monitoring with particular reference to anticonvulsants.
- 4. What are the signs and symptoms observed in organophosphorus poisoning. Write its management.
- 5. Describe design, conduct and outcome of Phase I and Phase II of clinical trials
- 6. What in Institution Ethics Committee (IEC). Give its composition and responsibilities.
- 7. Discuss the principles of sampling in clinical trials. Differentiate parametric and non-parametric tests.
- 8. What are the main factors that influence drug dosing in renal failure. Explain one method for adjusting drug dose in renal disease.

# M Pharm Degree (Part I) Examinations – Pharmacy Practice

(Model Question Paper)

Paper IV – Hospital and Community Pharmacy & Drug Store Management
Time: 3 hrs

Maximum marks: 100

- Answer all questions
- Draw diagrams wherever necessary

Essays: (2x20=40)

- 1. Discuss the term inventory control from the point of view of a hospital pharmacist. What are the various methods of drug distribution used in modern hospital pharmacies.
- 2. What is the role of Community Pharmacy practice in family planning and first aid. Write a note on code of ethics for community pharmacists. What are the barriers a community pharmacist encounters during patient counselling.

Short Essays: (6x10=60)

- 3. Write notes on drug information sources Explain the term Polypharmacy and what are its consequences
- 4. What are OTC drugs, write patient counselling notes for any one OTC drug. IV additive service and role of pharmacist
- 5. Discuss the feasibility of manufacturing Large volume parenterals in district head quarter hospital
- 6. Explain the role of computers in hospital and clinical pharmacy
- 7. Write a note on organization of Hospital pharmacy. Discuss procurement procedures for various materials used in Hospital.
- 8. Define CSSD. Discuss the typical plan of central supply for medium sized hospital.