

QUESTION BANK M Pharmacy PHARMACEUTICS Semester-II



Molecular Pharmaceutics

(Nano Technology & Targeted DDS)
(NTDS)



LONG ESSAY 7.5 MARKS

UNIT I: Target Drug Delivery System

- 1. What are the events and biological process involved in drug targeting?
- 2. Explain the concepts, events and biological processes involved in drug targeting.
- 3. What do you mean by ligand mediated targeting?
- 4. Explain the blood brain barrier? What are the factors affecting transport across The blood brain barrier?
- 5. What are the ideal properties of carrier?
- 6. Write a note on tumour targeting.
- 7. Explain the cell biology and anatomy of blood brain barrier.
- 8. Describe in detail invasive methods to brain targeting.
- 9. What are the different types of targeting? Explain first order and second order targeting.
- 10. What do you mean by ligand mediated targeting?
- 11. Describe in detail different methods of tumour targeting?

UNIT II: Targeting Method

- 1. Discuss the methods for the preparation of phytosomes.
- 2. Explain the methods of preparation and evaluation of nanoparticles.
- 3. Explain about liposomal gene delivery system.
- 4. Write the methods for the preparation and applications of (a) Phytosomes (b) Electrosomes.
- 5. Explain the methods of preparation and evaluation of liposomes.
- 6. Explain the methods of preparation and evaluation of nano particles.
- 7. Write about aquasomes.
- 8. What are niosomes? Describe in detail methods of preparation and characterization of niosomes.
- 9. Explain detail about aquasomes.
- 10. Give a brief account on phytosomes.
- 11. Classify different methods of preparation of nanoparticles?
- 12. Explain in detail different types of preparation of polymeric nanoparticles.
- 13. What are niosomes? What are the differences between niosomes and liposomes.
- 14. Explain any five methods of preparation of niosomes.



UNIT III: Microcapscule / Microspheres

- 1. Describe in detail about various methods of preparation of microspheres.
- 2. Define microspheres. Write in detail preparation and evaluation methods of microspheres.
- 3. What are aerosols? Explain various propellants used in the manufacturing of aerosols.
- 4. Describe in detail evaluation methods of nasal drug delivery system.
- 5. Explain the methods for the preparation and evaluation of microspheres.

UNIT IV: Pulmonary Drug Delivery System

- 1. Write about methods of preparation and evaluation of aerosols.
- 2. Explain the factors influencing intranasal drug delivery.
- 3. Explain the nebulizers with suitable diagrams.
- 4. What are the factors influencing pulmonary drug delivery?
- 5. Discuss in detail about dry powder inhaler.
- 6. What are aerosols? Explain various propellants used in the manufacturing of Aerosols.
- 7. Explain intranasal insitu gels.
- 8. What are aerosols. What are the different types of containers used for aerosols?
- 9. Explain the various evaluation methods to evaluate aerosols.
- 10. Mention the advantages of intra nasal drug delivery system.
- 11. Explain in detail different types of intra nasal formulation and how to evaluate the same?

UNIT V: Nucelic Acid based Therapeutic DeliverySystem

- 1. Explain about bone marrow transplantation in ex-vivo gene therapy.
- 2. Explain the applications of monoclonal antibodies.
- 3. Explain about liposomal gene drug delivery.
- 4. Write various diseases treated using gene therapy.
- 5. Explain about therapeutic antisense molecules.
- 6. Describe hybridoma technology for production of monoclonal antibodies.
- 7. How will you characterize monoclonal antibodies?
- 8. Define gene therapy. Explain viral and non viral gene transfer methods.



- 9. Explain aptamers as drugs of future.
- 10. Describe in detail preparation of monoclonal antibodies. Mention few marketed preparations.
- 11. Write a note on: (a) Aptamers (b) Liposomal gene drug delivery system



Advanced Biopharmaceutics and Pharmacokinetics



LONG ESSAY 7.5 MARKS

UNIT I: Drug Absorption from the Gastrointestinal Tract

- 1. Explain various mechanism of drug absorption.
- 2. Explain pH partition hypothesis and its limitations.
- 3. Explain the influence of gastric emptying and intestinal transit time on absorption of drug.
- 4. Explain various pharmaceutical factors affecting drug absorption.
- 5. Discuss Noye's whiteys equation of drug dissolution process.
- 6. Define absorption. Discuss in detail the mechanism of drugabsorption.
- 7. Explain biological effect of drug absorption.
- 8. Discuss about GIT with respect to drug absorption.
- 9. In detail about active transport of drug absorption.
- 10. Discuss physico-chemical properties of a drug considered in drug product design

UNIT II: Biopharmaceutic Considerations in Drug Product Design and In Vitro Drug Product Performance

- 1. Discuss in detail in-vitro, in-vivo correlation. Explain briefly dissolution profile comparisons.
- 2. Describe the in vitro drug dissolution testing models. Explain theirmerits and limitations.
- 3. Discuss various factors to be considered in the design of a drugproduct.
- 4. Discuss the various official methods of dug dissolution.
- 5. Explain the different theories of dissolution process.
- 6. Write a detailed note on in vitro-in vivo correlation.
- 7. What are different compendial methods of dissolution.
- 8. Enumerate different categories of in vitro-in vivo correlationregarding drug product performance.
- 9. Describe compendial methods of dissolution testing.
- 10. Explain formulation factors affecting drug product performance



UNIT III: Pharmacokinetics

- 1. Explain one compartment model with extra vascular route of administration along with various pharmacokinetic parameters.
- 2. Define and classify compartment models. Discuss briefly the significance and limitations of compartment modelling.
- 3. Explain the pharmacokinetics of drug given by IV bolus, which follows one compartment open model.
- 4. Discuss in detail one compartment open model for a drug administered as IV infusion. Give the schematic representation, graphs and equations for the same.
- 5. Explain in brief extra vascular compartmental modelling.
- 6. Explain the causes of non linearity in pharmacokinetics of drug. Add a noteon estimation of Vmax and Km.
- 7. Discuss pharmacokinetic drug interaction.
- 8. Write a note on cytochrome P450 based drug interactions.
- 9. Write Michael's Menten equation. How do you estimate Vmax and Km.
- 10. Discuss about drug interactions linked to transporters.

UNIT IV: Drug Product Performance, In Vivo: Bioavailability and Bioequivalence

- 1. Explain the protocol for bioequivalence studies.
- 2. Write the objectives of bioavailability studies. Enlist the methods for measurement of bioavailability. Explain method using plasma concentration data.
- 3. What is bioequivalence? Discuss in detail various bioequivalence study designs
- 4. Define bioavailability? Write a note on various method assessing bioavailability.
- 5. What are the objective and considerations in bioavailability studies.
- 6. Discuss in detail various bioequivalence study design
- 7. Define the terms relative and absolute bioavailability. Explain any two methodsused to determine AUC.
- 8. Explain cross-over study design of bioavailability in detail. Gibe a note on measurement of bioavailability.
- 9. Enumerate clinical significance of bioequivalence studies.
- 10. Enumerate the various study designs available for carrying out bioequivalence studies and explain any one in detail



UNIT V: Application of Pharmacokinetics

- 1. Discuss Pharmacokinetic drug interactions
- 2. Discuss pharmacokinetics and Pharmacodynamics of biotechnology drugs.
- 3. Write a note on application of Pharmacokinetics in targeted drug deliverysystems.
- 4. Write a note on Pharmacokinetics and Pharmacodynamics of Protein andpeptides.
- 5. Give a note on Pharmacokinetics and Pharmacodynamics drug interactions.
- 6. Describe Pharmacokinetics and Pharmacodynamics of Vaccines.
- Compare and contrast the pharmacokinetics of conventional vs extended release dosage forms
- 8. Explain the Gene therapy with the help of any one FDA approve product
- Describe the application of pharmacokinetics in the development of modified-release drug products.
- 10. Write a detailed note on Biosimilar and generic drug products.



Computer Aided Drug Delivery System



UNIT I: Computers in Pharmaceutical Research and Development

- 1. Outline the quality by design concept in pharmaceutical product development with respect to International conference on harmonization guidelines.
- 2. Describe the different levels of *In-vitro* and *In-vivo* correlation
- 3. Discuss the role of computers in pharmaceutical formulation.
- 4. Write the benefits of pharmaceutical automation in packaging.
- 5. Explain the computational modeling concept with respect to drug absorption and solubility.
- 6. Enumerate the history of computers in pharmaceutical research & development.
- 7. Describe the use of computers in market analysis.
- 8. Enumerate the history of computers in pharmaceutical research & development.
- 9. What are the bio waiver considerations to be considered to get the exception for *In vivo* studies?
- 10. Computers in Market analysis.
- 11. The various statistical modeling in Pharmaceutical Research and Development
- 12. Applications of computer aided techniques in development of Pharmaceutical emulsion.

UNIT II: Computational Modeling of Drug Disposition

- 1. What is artificial intelligence? Mention its application.
- Explain the computational modeling concept concerning drug absorption and solubility.
 - a. Discuss the optimization parameters and different optimization techniques in formulation development.
- 3. Discuss QbD.
- 4. Discuss ICHQ8 Guideline
- 5. Biomedical simulations (Pharmacokinetic & Pharmaco dynamic).
 - a. Applications of scientifically based QbD.
- 6. The experimental design in optimization of Pharmaceutical formulations.



- 7. Comparison of the traditional and QbD approach (ICH Q8 guidelines) in Pharmaceutical development.
- 8. The benefits of QbD in Industry and regulation bodies.
- 9. Discuss in detail about ICH Q 8 guidelines.
- 10. What is modeling? Elaborate computational modeling of drug disposition

UNIT III: Computer-aided formulation development

- 1. Computers in Clinical development.
- 2. Explain the role of computers in clinical data collection and management
- 3. What are the differences between non clinical, preclinical and clinical studies write short notes on clinical data collection?
- 4. Write the role of computers in clinical data collection and management for clinical development.
- 5. Write about computers in clinical development as clinical data collection and management.
- 6. The various statistical modeling in Pharmaceutical Research and Development

UNIT IV: Computer-aided biopharmaceutical characterization

- 1. Explain the role of computers in clinical data collection and management.
- 2. What is Modeling? Explain Computational Modeling of Drug Disposition.
- 3. Explain in detail about computer aided formulation development.
- 7. The applications of computer in various Intellectual Property Rights of Pharmaceutical R & D.
- 8. The various statistical modeling in Pharmaceutical Research and Development.
- 9. The history of computers in Pharmaceutical Research and Development.
- 10. The experimental design in optimization of Pharmaceutical formulations.



UNIT V: Artificial Intelligence (AI), Robotics and Computational fluid dynamics:

- 1. Write history and role of computers in pharmaceutical research and development.
- 2. Write a note on Artificial intelligence and robotics in pharmaceutical automation
- 3. Write a note on Pharmaceutical applications, advantages and challenges of robotics in pharmaceutical product development.
- 4. Write a note on the Artificial intelligence and robotics in pharmaceutical automation.
- 5. Write a note on the Pharmaceutical applications, advantages and challenges of robotics in pharmaceutical product development.
- 6. Discuss about artificial intelligence and robotics in pharmaceutical automation and write their application, advantages and disadvantages.
- 7. Write current challenges and future directions of AI.
- 8. Mention and explain the various applications of Artificial intelligence in pharmaceutical automation.
- 9. Write brief notes about Robotics in Pharmacy



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LONG ESSAY 7.5 MARKS

UNIT I: Cosmetics-Regulatory

- 1. Explain in detail plant layout and factory requirements for manufacturing of cosmetics.
- 2. Discuss on salient features of international nomenclature cosmetic ingredients. Add a note on the labelling requirements for cosmetic products.
- 3. Discuss the conditions to be fulfilled for manufacturing of cosmetics according to Indian regulatory provisions.
- 4. Discuss the regulatory provisions for import of cosmetics
- 5. Explain the regulatory requirements for labelling of cosmetics
- 6. Explain the conditions for obtaining license
- 7. Write a note on prohibition of manufacture and sale of certain cosmetics.
- 8. Explain about the offences and penalties of cosmetics
- 9. Write a note on Indian regulatory requirements for labelling of cosmetics.
- 10. Explain about cosmetic- regulatory aspects

UNIT II: Cosmetics-Biological aspects

- 1. Discuss about structure of hair and hair growth cycle. List the different hair care products
- 2. Define cosmetic products as per European union guidelines. Discuss basic raw materials used in the formulations of creams.
- 3. Describe the formulations used for foot care and hygiene.
- 4. Discuss the cleaning and care needs of under arms. Discuss in detail about cosmeceutical products used to address body odour.
- 5. Define cosmetics and discuss the structure of hair and the hair growth cycle. Explain the reasons and treatment for dandruff.
- 6. Explain the biological aspects of skin relating problem like dry skin, acne and pigmentation
- 7. Write a note on common problems associated with oral cavity
- 8. Write a note on cleansing and care needs for face, eye lids
- 9. Explain about neck, body and under arm care need
- 10. Write a note on biological aspects of cosmetics.



UNIT III: Formulation Building blocks

- 1. What are rheology modifiers and classify rheology modifiers.
- 2. Classify surfactants and mention the applications.
- 3. Define cosmetics. Give the building block for formulation of cream.
- 4. Write the classification and applications of surfactants.
- 5. Write a note on factors influencing effectiveness of preservatives.
- 6. Discuss on different controversial ingredients used in cosmetic products
- 7. Classify perfumes used in industry with examples.
- 8. Explain on the formulation of amine based hair dyes.
- 9. Describe the different ingredients used in the formulation of toothpaste.
- 10. Write the difference between perfume and flavouring agents and give the list of allergens in perfume.

UNIT IV: Design of cosmeceutical products

- 1. Write in detail on cosmeceutical products for mouth odor and sensitive teeth.
- 2. Cosmeceutical products for dandruff control.
- 3. What are ideal properties of sun screening agents. How are they classified?
- 4. What is antiperspirant. Add a note on its formulation
- 5. Explain the skin condition relating to prickly heat and how to address the same.
- 6. Explain the cosmetic preparations used in treating bleeding gum and sensitive teeth.
- 7. Explain on the sunscreen for skin protection.
- 8. Classify perfumes
- 9. Write a note on parabens, formaldehyde liberators
- 10. Explain about the building blocks for formulation of a moisturizing cream, vanishing cream



UNIT V: Herbal cosmetics

- 1. Write briefly on challenges in formulating herbal cosmetics
- 2. Write a note on guidelines for herbal cosmetics by private bodies.
- 3. Discuss about herbal ingredients used in oral care.
- 4. Explain hair colorants. Add brief note on herbal hair colorants.
- 5. What are herbal cosmetics? Discuss in detail herbal ingredients used in the formulation of cosmetics.
- 6. Discuss in detail various colorants. Add a note on herbal hair colorant.
- 7. Discuss the formulation of herbal toothpaste.
- 8. Discuss the role of emollients rheology modifiers in cosmetic products including their classification.
- 9. What are the challenges in formulating herbal cosmetics.
- 10. Explain the factors influencing performance of preservatives in cosmetic products.
- 11. Explain the role of herbal ingredients for oral care.
- 12. Write a note on syndet bars used in personal care.
- 13. Explain guidelines for herbal cosmetics with respect to preservatives.



Vision and Mission of the Institution

Vision

The East Point College of Pharmacy aspires to be a globally acclaimed institution, recognized for excellence in pharmaceutical education, research and nurturing students for holistic development.

Mission

- M1 Create pharmacy graduates through quality education
- M2 Promote innovation, **creativity**, and excellence **in teaching**, learning, and **research**
- M3 Inspire integrity, teamwork, critical thinking, personal development, and ethics in students and lay the foundation for lifelong learning
- M4 Serve the healthcare, technological, scientific, and economic needs of then society.