

**Code No: 12285/PCI** 

## **FACULTY OF PHARMACY**

B. Pharmacy VI-Semester (PCI) (Main &Backlog) Examination, September 2021

**Subject: Medicinal Chemistry-III** 

Time: 2 Hours Max. Marks: 75

Note: Answer any seven questions from Part –A, Any one questions from Part-B and Any five questions from Part-C
PART – A (7X3 = 21 Marks)

- 1. Define & classify  $\beta$  lactam antibiotics?
- 2. Write the structure of Benzly pencillin & Chlortetracycline?
- 3. Write the structure and uses of Strptomycin?
- 4. Write the structure of Isoniazid & Para amino salicylic acid?
- 5. Write the structure and uses of Dapsone?
- 6. Define Partition coefficient, Hammet's electronic parameter?
- 7. Write the mechanism of action of Tetracyline?
- 8. Write the mechanism of action of Macrolides?
- 9. Define prodrugs?
- 10. Write the β Lactamase inhibitors?

## **PART-B (1 X 14 = 14 Marks)**

- 11. Enumerate the various classes of antitubercular drugs. Write the synthesis & mode of action of Isoniazid?
- 12. Define Beta lactam antibiotics and explain the classification, SAR and mode faction of cephalosporins?
- 13. Write the life cycle of malaria parasite and Write the synthesis and mode of action of the Chloroquine?

## PART - C (5 X 8 = 40 Marks)

- 14. Write the chemical degradation of pencillin?
- 15. Write the SAR and uses of Tetracycline?
- 16. Write the structure, synthesis, mode of action and uses of Chloramphenicol?
- 17. Write a note on Prodrugs?
- 18. Write the structure, synthesis, mode of action and uses of Nitrofurantion?
- 19. Write the mode of action and SAR of Sulphonamides?
- 20. Write a short note on combinatorial chemistry?
- 21. Write the structure, synthesis, mode of action and uses of Miconazole?
- 22. Write the structure, synthesis, mode of action and uses of Mebendazole?



#### **FACULTY OF PHARMACY**

## B. Pharmacy VI-Semester (PCI) (Main & Backlog) Examination, September 2021

Subject: Pharmacology - III

Time: 2 Hours Max. Marks: 75

Note: Answer any seven questions Part – A, any one question from Part – B and any five questions from Part – C.

## $PART - A (7 \times 3 = 21 Marks)$

- 1 What are antiemetics?
- 2 What are nasal decongestants?
- 3 Differentiate between purgatives and laxatives.
- 4 Define the following terms
  - a. Circadian rhythm
- b. circannual rhythm
- 5 What are fluoroquinolones? Give examples.
- 6 Enumerate various antidotes available.
- 7 Define Chronotherapy and write its applications.
- 8 Write a note on appetite suppressants.
- 9 What are the cholinesterase reactivators? Give examples.
- 10 How do carminatives act?

## PART- B (1 x 14 = 14 Marks)

- 11 Classify the agents used in treatment of peptic ulcer disease. Write about the pharmacological actions and therapeutic uses of Ranitidine and Omeprazole.
- 12 Write the MOA, adverse effects and therapeutic uses of Reverse transcriptase inhibitors and cisplastin.
- 13 Explain the cell cycle. What are fluoroquinolones? Explain their MOA, therapeutic uses and adverse effects.

#### PART- C (5 x 8 = 40 Marks)

- 14 Explain the MOA and adverse effects of aminoglycosides and penicillins.
- 15 Write short notes on the pharmacology of H<sub>2</sub> receptor blockers.
- 16 Discuss the symptoms and treatment of heavy metal poisoning.
- 17 Write a note on antimalarial drugs.
- 18 What are protein based drugs? Write short notes on them.
- 19 Write about urinary tract infections.
- 20 Classify antifungal drugs. Write the MOA and adverse effects of amphotericin.
- 21 Discuss the symptoms and treatment of barbiturate poisoning.
- 22 What are immunosuppressants? Classify them.

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**Code No: 12287/PCI** 

## **FACULTY OF PHARMACY**

B. Pharmacy VI-Semester (Main & Backlog) Examination, September 2021

**Subject: Herbal Drug Technology** 

Time: 2 Hours Max. Marks: 75

Note: Answer any seven questions from Part-A, any one question from Part-B and any five questions from Part-C.

## Part – A $(7 \times 3 = 21 \text{ Marks})$

- 1 Define "IPR" and "Bioprospecting"
- 2 Write about curcumin
- 3 Explain soxhlet extraction
- 4 Differentiate conventional and organic farming
- 5 Write about any two microbial pesticides
- 6 Write composition and functions of ASUDCC
- 7 What are churnas & bhasmasWrite about guar gum and saffron
- 8 Write the underlying principle of homeopathy
- 9 What are the methods for authentication of a herb

## $Part - B (1 \times 14 = 14 Marks)$

- 10 Write a detailed account of the guidelines for stability testing of herbal drugs.
- 11 Write a note on: (a) Vitamins as antioxidants (b) Schedule Z.
- 12 Write a short note on (a) Herbal drug industry (b) Traditional Knowledge

## Part – C (5 x 8 = 40 Marks)

- 13 Write a note on the role of nutraceuticals in the prevention and management of cardiovascular diseases.
- 14 Write about pharmacokinetic herb drug interactions with examples.
- 16 Present an overview of good agricultural practices.
- 17 Write an account of plant based research institutes in India.
- 18 What is bio piracy? Discuss any three bio piracy cases in India.
- 19 Classify herbal excipients. Write in detail about naturally derived thickening agents.
- 20. Write the method of preparation and standardization of churnas.
- 21. Discuss various methods used for processing of herbal materials.
- 22. Write a note on omega fatty acids and resveratrol.

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**Code No: 12288/PCI** 

## **FACULTY OF PHARMACY**



## B. Pharmacy VI-Semester (PCI) (Main & Backlog) Examination, September 2021Subject:

## **Biopharmaceutics and Pharmacokinetics**

Time: 2Hours Max. Marks: 75

Note: Answer any Seven Questions from Part –A, Any One Questionsfrom Part-B.and Any Five Questions from Part-C

PART – A (7X3 = 21 Marks)

- 1) Define biopharmaceutics
- 2) Mention factors affecting Absorption
- 3) Differentiate Passive transport and Active transports.
- 4) Define Absolute bioavailability and relative bioavailability.
- 5) List the factors affecting elimination of drugs.
- 6) Explain Flip-flop method in Extra vascular administration.
- 7) What is apparent volume of drug distribution
- 8) Write the equation for calculating steady state drug concentration for one compartment open model.
- 9) What is proties binding. How it affects bio availability 10) Expand the terms: i. AUC, ii. Vd iii.  $t_{1/2}$  iv. Ka v. K $\theta$  vi. CLR

## Part - B (1 x 14 = 14 Marks)

- 11) Write about in-vitro drug dissolution models.
- 12) How do you estimate the pharmacokinetics parameters (K max and V max) by using Michaelis Menton equation.
- 13) Discuss about factors influencing absorption of drug in GIT

#### Part - C (5 x 8 = 40 Marks)

- 14) Write a note on Carrier mediated transport.
- 15) Describe about the physiological barriers to the distribution of drugs. Any three.
- 16) Explain the biliary excretion of drugs.
- 17) Explain the various methods for assessment of bioavailability.
- 18. Discuss in-Vitro-in-Vivo correlation
- 19. Explain kinetics of protion binding
- 20. Write in detail about compartment models.
- 21) Write a note on non-linear pharmacokinetics.
- 22)Explain methods of adjustment of dose and dosage regimen in patients withhepatic failure

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#### **FACULTY OF PHARMACY**

B. Pharmacy VI-Semester (PCI) (Main & Backlog) Examination, September 2021

Subject: Pharmaceutical Biotechnology

Time: 2 Hours Max. Marks: 75

Note: Answer any Seven Questions from Part –A, Any One Questions from Part-B. and Any Five Questions from Part-C

PART – A (7X3 = 21 Marks)

- 1. Define the following:
  - i) Biotechnology ii) Enzyme immobilization
- 2. Write the components of Biosensors
- 3. Write significance of enzyme acting on DNA
  - i) Restriction end nucleases ii) S1 nuclease
- 4. Enumerate types of cloning vectors. Add a note on COSMID as vector
- 5. What is active immunity?
- 6. Write stability tests defined for official vaccines
- 7. Give applications involved in Southern blotting technique
- 8. How will you transfer gene by conjugation method
- 9. How to control foam during fermentation?
- 10. Mention six enzymesPART- B (1 X 14 = 14 Marks)
- 11. Give the principle of rDNA technology along with significance of enzymes. Enlist and explain various methods of screening the recombinants.
- 12. Define vaccine. Write the method of preparation and quality control of bacterial vaccine
- 13. Discuss production of Penicillin by fermentation process.

#### PART - C (5 X 8 = 40 Marks)

- 14. Enlist methods of immobilization of enzymes. Add a note on applications of enzyme Immobilization
- 15. Explain the applications of Biosensors
- 16. Write a brief account on production of insulin by rDNA technology
- 17. Differentiate between 'type II Hypersensitivity' the 'type III Hypersensitivity'.
- 18. Give role of HAT medium in monoclonal antibody production
- 19. Write short notes on ELISA technique
- 20. Differentiate between prokaryote and Eukaryote
- 21. Describe components and working of fermentor
- 22. Write short note on vitamin B12 Production by fermentation

## Code No:12007/PCI

#### **FACULTY OF PHARMACY**

## B. Pharmacy VI-Semester (PCI) (Suppl.) Examination, March 2021

**Subject : Medicinal Chemistry-III** 

Time: 2 Hours Max. Marks: 75

Note: Answer Any Seven Questions from Part –A, Any one Question from Part-B. and Any Five Questions from Part-C

#### PART - A (7 X 3 = 21 Marks)

- 1) Define &classify antibiotics?
- 2) Write the structure of Sulbactam & Monobactam?
- 3) Write the structure and uses of Strptomycin?
- 4) Write the structure of Chloroquine & Ethambutol?
- 5) Write the structure and uses of Dapsone?
- 6) Define Partition coefficient, Tafts steric parameter?
- 7) Write the mechanism of action of Aminoglycosides?
- 8) Write the mechanism of action of Macrolides?
- 9) Define prodrugs?
- 10) Write the mode of action of anti-fungal agents?

# Part - B (1 x 14 = 14 Marks)

- 11) Write the various classes of antitubercular drugs. Write the synthesis & mode of action of Isoniazid?
- 12) Define Beta lactam antibiotics and explain the classification, SAR and mode faction of Pencillins?
- 13) Write the mode of action and SAR of Sulphonamides and Write the synthesis of Sulfamethoxaole?

## Part - C (5x8 = 40 Marks)

- 14) Write the chemical degradation of Cephalosporins?
- 15) Write the SAR and uses of Tetracycline?
- 16) Write the structure, synthesis, mode of action and uses of Chloramphenicol?
- 17) Write a note on Prodrugs?
- 18) Write the structure, synthesis, mode of action and uses of Ciprofloxacin?
- 19) Write the structure, synthesis, mode of action and uses Trimethoprim?
- 20) Write a short note on combinatorial chemistry?
- 21) Write the structure, synthesis, mode of action and uses of Tolnaftate?
- 22) Write the structure, synthesis, mode of action and uses of Mebendazole?

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Code No: 12012/PCI

#### **FACULTY OF PHARMACY**

B. Pharmacy VI-Semester (PCI) (Suppl.) Examination, March 2021

**Subject: Quality Assurance** 

Time: 2 Hours Max. Marks: 75

Note: Answer Any Seven Questions from Part –A, Any one Question from Part-B. and Any Five Questions from Part-C

PART - A (7 X 3 = 21 Marks)

- 1 Define ISO 14000
- 2 Define Quality by design (QbD)
- 3 Define Quality Assurance.
- 4 Define GMP.
- 5 Explain out of trend (OOT)?
- 6 What is Warehousing? Explain.
- 7 Explain on out of specification.
- 8 What is Sources of impurities? Explain.
- 9 What is calibration. Why it should be bone for equipment
- 10 What is SOP.? Explain.

## Part - B (1x 14 = 14 Marks)

- 11 Write briefly about Quality control test for secondary packing materials.
- 12 Define ICH. Explain about ICH Guidelines.
- 13 Write briefly about importance, scope of validation and types of validation.

## Part - C (5x 8 = 40 Marks)

- 14 Describe SOP, Quality audit and Quality Review.
- 15 Write a short note on Total Quality Management (TQM)
- 16 List out what are the different analytical instrumentation used in the estimation of impurities.
- 17 Explain about validation master plan.
- 18 Explain about Personnel responsibilities, training, and hygiene
- 19 Write a short note on ISO 9000 series?
- 20 Explain about Equipment selection, purchase specifications and maintenance
- 21 Explain ISO certification procedure and its advantages?
- 22 Write briefly about Quality control test for containers.

Code No: 12011/PCI

#### **FACULTY OF PHARMACY**

B. Pharmacy VI-Semester (PCI) (Suppl.) Examination, March 2021

**Subject : Pharmaceutical Biotechnology** 

Fime: 2 Hours Max. Marks: 75

Note: Answer Any Seven Questions from Part –A, Any one Question from Part-B. and Any Five Questions from Part-C

## PART - A (7 X 3 = 21 Marks)

- 1 What is biosensor? Write the main components used in biosensor.
- 2 What is genetic engineering?
- 3 Explain restriction endonucelases.
- 4 What are vectors? Write the ideal properties of vectors.
- 5 Write few applications of hybridoma technology.
- 6 What are toxins? Explain the method of conversion of toxin to toxoid.
- 7 Write the preparation and uses of human fibrinogen.
- 8 Write about types of aerators in Fermentor.
- 9 Write a brief note on plasmids.
- 10 Differentiate between prokaryotic and Eukaryotic organisms.

#### Part - B (1 x 14 = 14 Marks)

- 11 Discuss the structure and function of Major Histocompatability Complex.
- 12 Explain the typical structure of Immunoglobulin and types of Antibodies.
- 13 What are plasma substitutes? Explain the manufacturing of plasma substitutes and standardization.

#### Part - C (5x 8= 40 Marks)

- 14 Write a brief notes on Protein Engineering
- 15 Explain the steps involved in PCR.
- 16 Explain insertion and replacement vectors (Bacteriophage vector)
- 17 Discuss the general methods of preparation of vaccines.
- 18 Explain Southern blotting technique.
- 19 Explain generalized transduction and specialized transduction.
- 20 What are mutations? Explain the types of mutations.
- 21 Draw a neat labeled diagram of a typical fermentor.
- 22 Explain the aeration and agitation process in fermentation.

Code No. 12010 / PCI



B. Pharmacy VI-Semester (PCI) (Suppl.) Examination, March 2021

Subject: Biopharmaceutics & Pharmacokinetics

Time: 2 Hours Max. Marks: 75

Note: Answer any seven questions Part – A, any one questions from Part – B and any five question from Part – C.

## **PART – A (7x3=21 Marks)**

- 1 Define Noyes and Whitney equation and its application.
- 2 What is Lipinskis rule of five?
- 3 Differentiate between plasma protein-drug binding and tissue drug binding.
- 4 Define Microconstants and Hybrid constants and write relationship between them.
- 5 What is Flip-Flop Phenomenon and how it is useful in method of residual?
- 6 What are the different methods used to calculate the AUC?
- 7 Define orange book and objectives of bioavailability studies.
- 8 Difference between Absolute bioavailability and Relative bioavailability.
- 9 Define Creatinine and how to calculate the Creatinine Clearance
- 10 If Vd of thiopental is 2000ml. Calculate the amount of drug in the body when plasma concentration is 2µg/ml.

## **PART- B (1x14=14 Marks)**

- 11 Define absorption. Write in detail about mechanism of drug absorption with diagram.
- 12 Explain in detail about Bioequivalence study protocols.
- 13 Derive Michaelis-Menten equation and how do you estimate Km and Vmax.

## **PART- C (5x8=40 Marks)**

- 14 Write in detail about pH partition hypothesis and its limitation.
- 15 Write about Gastric emptying rate and Volume of distribution.
- 16 Significance and kinetics of protein drug binding.
- 17 What are the factor causing Non-Linearity?
- 18 How do you calculate absorption rate constant, Ka by using Wagner Nelson method?
- 19 Explain the pharmacokinetic parameters of a drug which follows one compartment open model when given by intravenous bolus with relevant mathematical equations.
- 20 What are the different methods for Assessment of Bioavailability?
- 21 Explain various cross over designs in Bioequivalence studies.
- 22 A 60 kg male received 2mg/kg of a drug orally. The following plasma concentration vs time data is obtained. Assume the drug follows one compartment open model and it is completely absorbed. Calculate all possible pharmacokinetic parameters.

Time(hr)	1	2	3	4	5	6	8	10	12	14
Plasma Conc. (µg/ml)	3.2	7.3	9.1	9.7	9.7	9.2	7.1	5.3	4.0	3.0

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Code No: 12009/PCI

## **FACULTY OF PHARMACY**

B. Pharmacy VI-Semester (PCI) (Suppl.) Examination, March 2021

Subject: Herbal Drug Technology

Time: 2 Hours Max. Marks: 75

Note: Answer Any Seven Questions from Part –A, Any one Question from Part-B. and Any Five Questions from Part-C

PART - A (7 X 3 = 21 Marks)

- 1 What is Tridosha
- 2 What are Biopesticides
- 3 Define the term nutraceuticals
- 4 Write the health benefits of Amla.
- 5 What are advantages of Herbal Excipients.
- 6 Define the term Cosmetics.
- 7 Define the term patent and farmers right
- 8 What do you mean by evaluation of drugs?
- 9 List plant based government research institutes in India.
- 10 Write a note on Biopiracy.

#### Part - B (1x 14 = 14 Marks)

- 11 Briefly explain good agricultural practices in cultivation of medicinal plants.
- 12 What is traditional knowledge. Explain patenting aspects of Traditional knowledge and natural products.
- 13 Briefly explain the objectives and components of Schedule-T.

## Part - C (5x 8 = 40 Marks)

- 14 List the Ayurvedic formulations and write the preparation of Bhasma.
- 15 Explain the principles of Siddha system of medicine.
- 16 Discuss the future prospects of Herbal drug industry.
- 17 Write a detailed account of case study of Neem and Curcuma.
- 18 Write the methods of stability testing of herbal drugs.
- 19 Write note on herbal binders and diluents.
- 20 Write the possible side effects and interactions of garlic and pepper.
- 21 Write the role of Honey and Alfa alfa as health food.
- 22 Write about pest management in medicinal plants.



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## **FACULTY OF PHARMACY**

# B. Pharmacy VI-Semester (PCI) (Main) Examination, December 2020 Subject: Quality Assurance

Time: 2 Hours Max. Marks: 75

#### PART - A

#### Note: Answer any Seven questions.

 $(7 \times 3=21 Marks)$ 

- 1. Define TQM
- 2. Give difference between Quality Assurance & Quality Control.
- 3. State the purpose of ICH.
- 4. Name Quality Control tests for glass containers.
- 5. Name different parameters of Analytical method validation.
- 6. Name any four responsibilities of Quality control people.
- 7. Mention classification of Recall.
- 8. What is qualification and validation.
- 9. Enlist the scope for validation.
- 10. Give the principles of NABL accreditation.

## PART - B

#### Note: Answer One question.

(1 x14=14 Marks)

- 11.a) Define Quality by Design.
  - b) Write in detail note on QbD.
- 12. Write a short note on plant layout with example.
- 13. Explain Good Warehousing practices.

#### PART - C

#### Note: Answer any Five questions.

(5x8=40 Marks)

- 14. Write in detail Equipment Validation.
- 15. Draw cause and effect diagram for tablet manufacturing process.
- 16. Write in detail parameters to be checked in Quality Audit.
- 17. Write short note on ISO 9000.
- 18. Explain in short Good Laboratory practices.
- 19. Explain steps involved in complain handling.
- 20. Explain the term "validation Master Plan".
- 21. What is forced degradation stability study? Explain in short.
- 22. Write a note on Quality Management System.



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Code No: 6225/PCI

## **FACULTY OF PHARMACY**

## B.Pharmacy VI-Semester (PCI) (Main) Examination, November 2020

Subject : Pharmaceutical Biotechnology

Time: 2 Hours Max. Marks: 75

## PART - A

## Note: Answer any Seven questions.

(7 x3=21 Marks)

- 1. Enlist applications of biotechnology to pharmaceutical industry.
- 2. Describe the terms biosensor and bioreactor.
- 3. Write significance of enzyme acting on DNA.
  - i) Polymerase
- ii) Ligase
- 4. Describe the importance linkers and adapters.
- 5. What is toxoid. Give examples
- 6. What are plasma substitutes?
- 7. Define the following:
  - i) Immunoblotting
- ii) Immuno suppression.
- 8. How will you transfer gene by transduction method?
- 9. Define fermentation.
- 10. Write six enzymes.

#### PART - B

#### Note: Answer One question.

(1 x14=14 Marks)

- 11. Explain benefits of recombinant DNA products. Write a detailed account on human insulin production by rDNA technology
- 12. What is Hybridoma technology? Explain the steps involved in the production of monoclonal antibodies and applications.
- 13. Describe Microbial biotransformation and its pharmaceutical applications.

## PART - C

#### Note: Answer any Five questions.

(5x8=40 Marks)

- 14. Explain the concept of enzyme immobilization. Comment on its applicability with suitable examples.
- 15. Write short notes on production of amylase.
- 16. Write short notes on interferon production by rDNA technology.



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- 17. Write a short note on PCR.
  - 18. Differentiate between humoral mediated immunity and cell mediated immunity.
  - 19. Give an account of collection, processing & storage of whole human blood.
  - 20. What is solthern blotting? Give details of southern blotting and application.
  - 21. Enlist various criteria to be considered in designing of a fermentor, Draw a neat schematic labelled diagram of fermentor.
  - 22. Write short notes on antibiotic production by fermentation with suitable example.



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## **FACULTY OF PHARMACY**

## B.Pharmacy VI-Semester (PCI) (Main) Examination, November 2020

**Subject: Medicinal Chemistry - III** 

Time: 2 Hours Max. Marks: 75

#### PART - A

#### Note: Answer Seven Questions.

(7 X 3 = 21 Marks)

- 1. Write the general synthesis of sulfonamides.
- 2. What are folate reductase inhibitors?
- 3. Give the mechanism of action of Trimethoprim.
- 4. Mention any six quinolone drugs.
- 5. What are Monobactams?
- 6. Classify antitubercular agents with examples.
- 7. Mention any six sulfonamide drugs
- 8. Mention any six antifungal agents
- 9. Mention any six antiviral drugs.
- 10. Mention any six antiprotozoal agents?

#### PART - B

#### Note: Answer One Question.

(1X14 = 14 Marks)

- 11.a) Write a note on B-lactam antibiotics
  - b) Write a note on tetracyclines.
- 12.a) Write the classification of antifungal agents
  - b) Give the synthesis, mechanism of action and uses of any one antifungal drug.
- 13. a) Write a note on Tetracyclines.
  - b) Write a note on Anti-protozoal agents.

#### PART - C

#### Note: Answer Five Question.

(5X8 = 40 Marks)

- 14. Discuss the SAR of semi-synthetic Penicillins.
- 15. What are prodrugs? Write the classification of Prodrugs based on functional groups.
- 16. Write the synthesis and mechanism of any two sulfa drugs.
- 17. Give a note on Artemisinin derivatives.



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- 18. Write the synthesis, mode of action and therapeutic uses of Isoniazid and Para amino salicylic acid
- 19. Write a note on Anti-HIV drugs.
- 20. Write the synthesis and mechanism of Diethylcarbamazine citrate and Metronidazole.
- 21. Write about Quinoline antibiotics.
- 22. What are  $\beta$  lactam antibiotics? Write their mechanism of action.



Code No: 6224/PCI

## **FACULTY OF PHARMACY**

## B. Pharmacy VI-Semester (PCI) (Main) Examination, November 2020

**Subject: Biopharmaceutics and Pharmacokinetics** 

Time: 2 Hours Max. Marks: 75

#### PART - A

Note: Answer Seven Questions.

(7 X 3 = 21 Marks)

- 1) Mention the factors effecting elimination of drugs
- 2) List the factors influencing absorption of drugs through GIT
- 3) Differentiate tissue binding and protein binding.
- 4) Write the markers used in renal clearance.
- 5) Define Bioavailability.
- 6) Expand the terms  $i.A \cup C$  ii.  $t_{\frac{1}{2}}$ , iii. Vd iv. IV v. Ka vi.  $E_{E}$
- 7) What is  $t_{\frac{1}{2}}$  what is its importance
- 8) Write the equation for calculating loading dose.
- 9) What is apparent volume of distribution and its importance
- 10) What are the factors for cause of non-linear kinetics.

#### PART - B

Note: Answer One Question.

(1X14 = 14 Marks)

- 11) Write about in vitro drug dissolution models
- 12) Derive mathematical equations used to calculate Pharmaco-Kinetic parameters following IV bolus administration blood data, assuming that the drug follows two compartment open model.
- 13) Discuss about protein binding and various factors affecting drug-protein binding.

#### PART - C

**Note: Answer Five Question.** 

(5X8 = 40 Marks)

- 14) Discuss the mechanism of Active diffusion in absorption of drugs.
- 15) How the organ size and perfusion rate influence the drug distribution?
- 16) Explain briefly about Kinetics of protein binding.



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- 17) Explain factors affecting the renal excretion of drugs.
- 18) Discuss about in vitro-in vivo correlations
- 19) A drug has a volume of distribution of 12Lts and elimination rate constant of  $0.18hr^{-1}$  A steady state concentration of  $12\mu g/ml$  is desred. Assuming one compartment kinetics, calculate time required to reach 99% of Css and infusion rate to achieve desired steady state.
- 20) Write the significance of different volumes of distribution in two compartment model.
- 21) Write a note on non-linear pharmacokinetics and Michaelis Mention equation.
- 22) How do you determine absorption rate constant, Ka by Wagner nelson method.

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## **FACULTY OF PHARMACY**

B. Pharmacy VI-Semester (PCI) (Main) Examination, November 2020

Subject: Herbal Drug Technology

Time: 2 Hours Max. Marks: 75

## PART - A

#### **Note: Answer Seven Questions.**

(7 X 3 = 21 Marks)

- 1) What is Organic farming.
- 2) Define the term Herbal medicine as per WHO.
- 3) Mention any six names of Aycervedic preparations (formulations)
- 4) What is significance of Herbal excipients
- 5) Write the health benefits of herbal medicines.
- 6) Define the term Nutraceuticals
- 7) List the parameters for evaluation of herbal tablets.
- 8) Define the term patent and IPR.
- 9) What is schedule T
- 10) What are antioxidants and give examples.

#### PART - B

#### Note: Answer One Question.

(1X14 = 14 Marks)

- 11) Briefly explain the objectives and components of Schedule-T
- 12) List the Ayurvedic formulations and write the preparation of any three.
- 13) Explain the WHO guidelines for the assessment of herbal drugs.

#### PART - C

#### Note: Answer Five Question.

(5X8 = 40 Marks)

- 14) How will you perform selection and identification of herbal materials?
- 15) Briefly explain the principles of Homeopathic system of Medicine.
- 16) Write a note on Functional foods and Dietary supplements.
- 17) Give informative note on Health benefits of nutraceuticals in management of diabetes.
- 18) What are excipients and give its classification with examples.
- 19) What are phytosomes? Give its method of preparation.
- 20) Give a detailed account of case study of neem and curcumin.
- 21) Explain the objectives and functions of ASU and DCC.
- 22) Give an informative note on future prospects of herbal drug industry.



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## **FACULTY OF PHARMACY**

## B. Pharmacy VI-Semester (PCI) (Main) Examination, November 2020 Subject : Pharamacology - III

Time: 2 Hours Max. Marks: 75

#### PART - A

#### Note: Answer Seven Questions.

(7 X 3 = 21 Marks)

- 1) What is asthma. Give four examples of drugs used in Asthma
- 2) What is ulcer. Give four examples of drugs used in ulcer.
- 3) What is the treatment for organophosphorus poisoning?
- 4) What is teratogenicity and give examples of drugs causing teratogenic effects.
- 5) Define Chronopharmacology.
- 6) What are the uses of sulfa drugs mention any four sulfa drugs.
- 7) What is amoebiasis Give any four examples of drugs.
- 8) What is BCG? What for it is used
- 9) Give two examples for Bronchiodilators and explain how they work?
- 10) Define Expectorant. Give two examples.

#### PART - B

#### Note: Answer One Question.

(1X14 = 14 Marks)

- 11) Classify anticancer agents. Add a note on antimetabolites.
- 12) Write the symptoms and management of Heavy metal poisoning.
- 13) Explain the pharmacological role of H<sub>1</sub> and H<sub>2</sub> antihistaminics.

#### PART - C

#### Note: Answer Five Question.

(5X8 = 40 Marks)

- 14) Write a note on sulfanamides.
- 15) Explain about Proton pump Inhibitors.
- 16) Write a note on Immunosuppressant's.
- 17) Explain the chemotherapy of Anti-TB drug.
- 18) Write a note on Penicillins.
- 19) Write a note on ant tubercular agents.
- 20) Write a note on antimalarial drugs.
- 21) Write the pharmacology of respirations stimulants
- 22) What are the different types of rhythms. Explain about circardian rhythm with example.



31.10h.01.