Code No: G-13061/PCI

# **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Make-up) Examination, November 2024
Subject: Biostatistics & Research Methodology

Time: 3 Hours Max. Marks: 75

#### PART - A

### Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Give importance of biostatistics in Pharmacy.
- 2. Hardness of 6 tablets is found as 6,8,5,7,9 and 11. Find mean and medium.
- 3. Find the mode and range for the angle of repose values of granul as give i as 13,18,13,14,13,16,14, 21 and 13.
- 4. Define the term probability.
- 5. What is meant by sample?
- 6. What is the need for Research?
- 7. Mention different types of graphs.
- 8. What is meant by Hypothesis?
- Give different statistical tools available in EXCEL
- 10. What are the advantages of factorial design?

PART - B

# Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Discuss on report writing in research methodology.
- 12. What is parametric test? Explain one way ANOVA in detail.
- 13. Give informative notes on (A) Correlation (B) Plagiarism

#### PART - C

# Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. The relative humidity values in a tablet production department of a pharmaceutical company from Monday to Saturday were recorded as 60,62,65,69,75 and 65. Calculate standard deviation.
- 15. Twenty hard gelatin capsules were examined for its physical properties. The frequency with a given number of defects per capsule is given. What is the probability of finding a capsule chosen at random contains 3 or more surface defects?

Number of Defects	0	1	2	3	4	5	6
Frequency	4	3	5	2	4	1	1

- 16. Write notes on any one non-parametric test.
- 16. What is sampling? Explain different types of sampling techniques.
- 17. Write about different types of graphs.
- 18. Explain the features of MINITAB in brief.
- 19. Write about experimental deisgn?
- 20. Obtain the line of regression of Y on Xfor the following data

Age in Yrs								63 47		
Blood Pressui	re (Y)	145	124	147	125	160	118	149 128	150	124

- 21. Explain 2<sup>2</sup> factorial design.
- 22. Give informative notes on response surface methodology.

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Code No: G-13062/PCI

# **FACULTY OF PHARMACY**

# B. Pharmacy VIII - Semester (PCI) (Make-up) Examination, November 2024 Subject: Social and Preventive Pharmacy

Time: 3 Hours Max.Marks:75

#### PART - A

#### Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Write a note on concept of public health?
- 2. What is Balanced diet?
- 3. What is drug addiction? Give few examples.
- 4. Write a note on prevention and control of hypertension?
- 5. What are the functions of PHC?
- 6. Write the objectives of pulse polio programme?
- 7. What are the strategies of national tobacco control programme?
- 8. Write a note on Impact of urbanization on health and disease?
- 9. Explain different types of diabetes mellitus?
- 10. Write a note on improvement in rural sanitation?

#### PART - B

# Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. (a) Explain various socio-cultural factors related to health and disease.
  - (b) Write a note on personal hygiene and healthcare.
- 12. (a) Write about the transmission, signs &symptoms, and treatment of Pneumonia.
  - (b) Write a note on Universal immunization programme.
- 13. (a) Write a note on Social health programme.
  - (b) Explain about the health promotion schemes in school.

#### PART - C

# Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Write a note on Vitamin deficiencies.
- 15. Explain malnutrition and its prevention.
- 16. Write a note on prevention and control of Drug addiction.
- 17. Describe the various roles of WHO in Indian national programs.
- 18. Write a note on Integrated Disease Surveillance Project (IDSP).
- 19. Write a note on National Family welfare programme.
- 20. Write a note on general principles of prevention and control of Dengue.
- 21. Write a note on objectives, functions and outcomes of TB control programme.
- 22. Write the risk factors, diagnosis and treatment of Cancer.

Code No: G-13063/PCI

# **FACULTY OF PHARMACY**

# B. Pharmacy VIII - Semester (PCI) (Make-up) Examination, November 2024 Subject: Pharma Marketing Management (Elective –I)

Time: 3 Hours Max. Marks: 75

#### PART - A

# Note: Answer all the question.

 $(10 \times 2 = 20 \text{ Marks})$ 

- Distinguish between marketing and selling.
- 2. Give an overview of Drug Price Control Order.
- 3. What is market segmentation and targeting?
- 4. What is sampling in promotion?
- Define consumerism.
- 6. What is product branding?
- 7. What is Physical distribution management?
- 8. What are the objectives of Pricing?
- 9. What are the future prospects of Professional sales representative?
- 10. Write about product life cycle.

#### PART - B

# Note: Answer any two question.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Write in detail about packaging and labeling decisions.
- 12. Discuss about Vertical and Horizontal Marketing.
- 13. Write a note on Pharmaceutical marketing channels.

#### PART - C

# Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. How a business management person maintains public relations?
- 15. What are the online promotional techniques for OTC products?
- 16. What are the duties of Professional sales representative (PSR)?
- 17. Write a note on Product decision
- 18. Explain about the patient's choice of physician and retail pharmacist.
- 19. Write about emerging concepts in marketing.
- 20. Give an overview of personal selling and advertising.
- 21. Write about global marketing.
- 22. What are the different components in marketing environment?

Code No: G-13064/PCI

# **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Make-up) Examination, November 2024 Subject: Pharmaceutical Regulatory Science (Elective - I)

Time: 3 Hours Max. Marks: 75

#### PART - A

Note: Answer all the questions.  $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Define (a) DMF (b) CTD.
- 2. Explain pre-clinical studies.
- 3. Define concept of generics.
- 4. Explain orange book, federal register.
- 5. What are Exclusion criteria in clinical trials.
- 6. Define Organogram of CDSCO.
- 7. What is clinical trial. Define pharmacovigilance?
- 8. Describe Objectives of regulatory affairs.
- 9. Write about Regulatory authorities of Canada.
- 10. What is good clinical practice

#### PART - B

### Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Write about drug development process.
- 12. Write a note on ANDA.
- 13. Explain in detail stages of drug delivery.

#### PART - C

# Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Briefly discuss CTD and eCTD.
- 15. Explain the technical documentation required for regulation of Indian drug product.
- 16. What is Orange book? Explain.
- 17. Write a note on 21 CFR.
- 18. Explain the GCP obligations of investigators.
- 19. Describe in detail new drug approval process along with its documentation requirements as per USFDA.
- 20. Describe formation and working procedure of independent ethics committee.
- 21. Briefly explain about India regulatory authority
- 22. What is electronic common technical document?

Code No: G-13065/PCI

### **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Make-up) Examination, November 2024 Subject: Pharmacovigilance (Elective - I)

Time: 3 Hours Max. Marks: 75

PART - A

Note: Answer all the questions.  $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Define the adverse drug reaction.
- 2. What are CIOMS working groups?
- 3. What is phase IV of clinical trials?
- 4. Write a short note on Harmonization.
- 5. Discuss the PSUR.
- 6. What is eudravigilance.
- 7. Explain ICH Steering Committee.
- 8. What is teratogenicity? Give examples.
- 9. Illustrate the importance of Pharmacogenomics.
- 10. Describe common technical document.

PART - B

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Differentiate between adverse drug reactions and adverse events with suitable examples. Explain the mechanisms of Type-A and Type-B ADRs.
- 12.(a) Write a note on MedDRA.
  - (b) Write a note on Pharmacovigilance program of India (PvPI).
- 13. Explain the criteria for Drug safety evaluation in Pediatric population.

PART - C

Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Describe in details CDSCO in India.
- 15. Explain the Schedule Y of Drugs and Cosmetics Act in brief.
- 16. Write about CROs in pharmacovigilance.
- 17. Discuss Naranjo's and WHO causality scales.
- 18. Write a note on post approval expedited reporting.
- 19. What is CIOMS? Enlist various CIOMS working groups and give their functions.
- 20. Discuss in brief the objectives of ICH guidelines.
- 21. Explore the Pre- marketing and Post marketing clinical trials.
- 22. Explain about eudravigilance medicinal product dictionary.

Code No: G-13067/PCI

### **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Make-up) Examination, November 2024
Subject: Computer-Aided Drug Design (Elective-I)

Time: 3 Hours Max. Marks: 75

PART - A

Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Differentiate among lead, drug candidate & drug.
- 2. Explain metabolism-based lead discovery with an example.
- 3. What is 3D QSAR technique? Give examples.
- 4. Explain the significance of the partition coefficient.
- 5. What is pharmacophore mapping?
- 6. Define de novo drug design.
- 7. Write the applications of cheminformatics tools in drug design.
- 8. What is the importance of ADME databases? Give a few examples.
- 9. Define the terms molecular mechanics and quantum mechanics.
- 10. What is conformational analysis?

PART - B

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Explain the methodology involved in Hansch QSAR analysis with its advantages & disadvantages. Highlight its role in predicting biological activity with a model QSAR equation.
- 12. Describe the concept of docking-based virtual screening in drug design.
- 13. Explain various stages involved in drug discovery and development.

PART - C

Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Describe the serendipitous discovery of drugs.
- 15. Classify bioisosteres with examples.
- 16. Describe Free-Wilson QSAR analysis with its advantages and disadvantages.
- 17. Discuss the methodology involved in CoMFA.
- 18. Explain Lipinski Rule of Five. How does it help in Drug design?
- 19. Explain chemical databases with suitable examples. Give their importance in drug design.
- 20. Describe the applications of bioinformatics tools in drug design.
- 21. Write about various energy minimization methods.
- 22. Write the applications of quantum mechanics in drug design.

#### **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Make-up) Examination, November 2024 Subject: Quality Control and Standardization of Herbals (Elective-I)

Time: 3 Hours Max. Marks: 75

PART - A

Note: Answer all the questions.  $(10 \times 2 = 20 \text{ Marks})$ 

1. Write about traditional system of medicine.

- 2. Write the biological method of crude drug evaluation with examples.
- 3. Explain the significance of ICH guidelines.
- 4. Write the test procedure of predinisolone phosphate.
- 5. How do you authenticate medicinal plants.
- 6. Write a note on cGMP.
- 7. Explain Lycopodium Spore Method Formula.
- 8. Write a note on long term toxicity test.
- 9. What are weedicides? Mention two examples.
- 10. Explain the stability testing of herbal medicine.

PART - B

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Explain different requirements to follow Good manufacturing practice? Why it is followed in herbal drug industry.
- 12. Discuss the guidelines given by Europian medicine agency on quality of traditional herbal medicine.
- 13. Write the comparison of various famous herbal pharmacopoeias.

PART - C

Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Write a note on good collection practices for herbal drugs.
- 15. Write the GMP requirement for herbal medicine.
- 16. Explain the preparation of document for new drug application.
- 17. Discuss the protocol for clinical guidelines in herbal medicine.
- 18. Enumerate various aspects of GLP.
- 19. Write the applications of chromatography technique in the standardization of herbal drugs.
- 20. Write briefly about stability studies of herbal medicinal products.
- 21. Describe the basic test for medicinal plant materials.
- 22. Write a note on research guidelines for evaluating safety and efficacy of herbal medicine.

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Code No: G-13068/PCI

### **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Make-up) Examination, November 2024 Subject: Cell & Molecular Biology (Elective-II)

Time: 3 Hours Max. Marks: 75

PART - A

Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. What are the basic features of cell theory?
- 2. Describe prophase-I of meiosis in brief.
- 3. Write about physical properties of DNA.
- 4. Write the functions of rRNA and mRNA.
- 5. What are essential and non-essential amino acids? Give examples.
- 6. Give a brief classification of proteins.
- 7. What are 'Restriction endonucleases' and write their function.
- 8. Enlist various enzymes used in genetic engineering.
- 9. Define ligands and receptors.
- 10. Enlist important intracellular signalling pathways.

PART - B

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Explain in detail about the properties of cell membrane. Differentiate between Prokaryotic and Eukaryotic cell.
- 12. Describe in detail about regularities in Protein pathway.
- 13. Describe Southern blotting technique and its applications.

PART - C

Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Briefly describe various components of a typical prokaryotic cell.
- 15. Describe Transcription and Translation.
- 16. Write the structure and functioning of DNA.
- 17. Describe types of RNA and flow of molecular information.
- 18. Describe any five colour reactions of proteins.
- 19. Explain positive control and significance of protein synthesis.
- 20. Write a note on vectors used in recombinant DNA technology.
- 21. Write short notes on genomic analysis.
- 22. Write about cell signalling and explain receptors for cell signals.

Code No: G-13069/PCI

# FACULTY OF PHARMACY B. Pharmacy VIII - Semester (PCI) (Makeup) Examination, November 2024 Subject: Cosmetic Science (Elective-II)

Time: 3 Hours Max.Marks:75

#### PART - A

#### Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Write a note on cosmetics as Quasi and OTC drugs.
- 2. Write a note on hair growth cycle.
- 3. Write a note on preservatives used in cosmetics.
- 4. Write a note on hair oils in hair care cosmetics.
- 5. Write a note on turmeric in skin care.
- 6. Write a note on mouthwashes.
- 7. Write the role of clove in oral care.
- 8. Write a note determination of skin colour.
- 9. What are the reasons for dry skin and how to prevent it?
- 10. Write a note on reasons and prevention of hair fall.

#### PART - B

### Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Write a note on following excipients with examples.
  - (i) Surfactants (ii) Humectants (iii) Rheology modifiers (iv) Emollients
- 12. Write a note on Sebumeter, Corneometer, Tewameter (TEWL) in cosmetic evaluation.
- 13. Write the causes and prevention of Blemishes, Wrinkles, Acne, Prickly heat and Body Odour.

#### PART - C

# Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Classify cosmetics and cosmeceuticals with examples.
- 15. Write a note on basic structure of skin and formulation of Moisturizing cream.
- 16. Write the principle involved in formulation of cold cream and vanishing cream.
- 17. Write the formulation of toothpaste for bleeding gums and sensitive teeth.
- 18. What is SPF? Classify the sunscreen formulations with examples.
- 19. Write a note on conditioning shampoo, antidandruff shampoo in hair care.
- 20. Write a note on Henna and Amla in hair care. Write about soaps and syndet bars.
- 21. Write formulation and mechanism of action of Antiperspirants & deodorants.
- 22. Write about soaps and syndet bars.

Code No: G-13072/PCI

# **FACULTY OF PHARMACY**

# B. Pharmacy VIII-Semester (PCI) (Make-up) Examination, November 2024

Subject: Experimental Pharmacology
Paper: (Pharmacological screening methods) (Elective-II)

Time: 3 Hours Max.Marks:75

#### PART - A

# Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. What are the methods used for preparation of drugs suspension?
- 2. What are the applications of mutant animals?
- 3. Explain the significance of sham negative group.
- 4. Write the different techniques for Euthanasia.
- 5. Explain the study designs involved in preclinical experiment.
- 6. Define a. Sedatives b. Hypnotics.
- 7. Write a short note on review of literature.
- 8. Give the examples of a. Mydriatics b. Miotics.
- 9. Mention the composition of IAEC.
- 10. Write the objectives of OECD guidelines.

#### PART - B

#### Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Describe any three screening methods for parasympatholytic drugs.
- 12. Explain the various preclinical screening methods for anti-inflammatory drugs. Give any two in-vivo methods.
- 13. Describe any three preclinical screening methods of anti-cancer drugs.

#### PART - C

#### Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Explain the different methods for collection of blood in laboratory animals.
- 15. What is the significance of statistical analysis of student t test?
- 16. Mention the objectives of CPCSEA. Write the composition and responsibilities of IAEC.
- 17. Describe on in vivo screening model for anti-epileptic drugs.
- 18. Explain any one preclinical screening method for diuretic activity.
- 19. Explain students 't' tort and one way ANOVA.
- 20. Mention briefly about any two preclinical screening methods for anti-parkinsonism activity.
- 21. Explain one preclinical screening method for anti-dyslipidemic activity.
- 22. Enumerate any two preclinical screening methods for anti-asthnatic activity.

Code No: G-13070/PCI

# **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Make-up) Examination, November 2024 Subject: Dietary Supplements and Nutraceuticals (Elective-II)

Time: 3 Hours Max. Marks: 75

PART - A

Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. What are probiotics and give examples.
- 2. Write the difference between natural and synthetic anti oxidants.
- 3. Give the occurrence and medical benefits of Lycopene.
- 4. Explain the importance of nutraceuticals in weight control.
- 5. Write about dietary fibres as functional food ingredients.
- 6. Give the source, chemical nature and uses of Oats and Rice bran.
- 7. Explain about enzymatic antioxidant defence.
- 8. Explain the role of melatonin and glutathione peroxidase.
- 9. Write about AGMARK on food safety.
- 10. What are tocopherols and give examples.

PART - B

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Explain in detail the effect of processing, storage and interactions of various environmental factors on the potential of neutraceuticals.
- 12.(a) Explain the role of Glutathione peroxidase and Superoxide dismutase.
  - (b) Write about public health nutritional benefits in a community.
- 13. Explain the role of Reactive Oxygen Species involvement in the treatment of disorders.

#### PART - C

Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Explain the role of anti-oxidants in the treatment of kidney damage.
- 15. Explain in detail about Carotenoids.
- 16. Explain the free radicals in the treatment of Diabetes milletus.
- 17. Give the pharmacopeial specifications for complex carbohydrates.
- 18. Give the occurrence, chemical nature and uses of Gingko and Ginseng.
- 19. Explain the regulatory aspects of FSSAI on food safety.
- 20. Explain the role of free radicals with lipids.
- 21. Define flavonoids and give the source and medicinal benefits of any two flavonoids.
- 22. Write in detail about adulteration of foods?

Code No: G-13071/PCI

# **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Make-up) Examination, November2024 Subject: Advanced instrumentation techniques (Elective-II)

Time: 3 Hours Max. Marks: 75

PART - A

Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. What is meant by ionisation in MS? List any 3 ionisation techniques used in MS.
- 2. Explain Base peak and molecular ion in MS.
- 3. What is meant by MALDI and FAB?
- 4. Define chemical shift. What is the effect of shielding and deshielding effect on chemical shift?
- 5. List the differences between single crystal and powder X-ray diffraction.
- 6. List the validation parameters as per ICH guidelines.
- 7. Write a note on hyphenated techniques. What are their advantages? Give suitable examples.
- 8. Define calibration and validation. What are the differences between calibration and validation?
- 9. Briefly explain the principle of DSC?
- 10. Give suitable applications of radioimmunoassay.

#### PART - B

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Explain the principle of Mass spectrometry. With a labelled diagram, explain MS instrumentation.
- 12. (a) List the calibration of UV- Visible spectrophotometer and explain any 2 parameters in detail
  - (b) Explain the Principle of Solid phase Extraction.
- 13. Explain the principle and instrumentation of GC-MS/MS.

#### PART - C

Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. With a neat labelled diagram explain the principle, instrumentation and application of Differential Thermal Analysis (DTA).
- 15. What is the role of mass analyser in MS. Explain any two in detail.
- 16. What are the differences between C13 and H1 NMR spectroscopy?
- 17. Describe the calibration of GC.
- 18. Explain the principle and procedure involved in liquid-liquid extraction.
- 19. How X-rays are generated? Derive Bragg's equation.
- 20. Explain any three ionisation methods in MS.
- 21. Explain the coupling constant, shielding and deshielding with suitable examples.
- 22. Briefly explain the process of radioimmunoassay.

Code No: F-7267/PCI

# **FACULTY OF PHARMACY**

# B. Pharmacy VIII - Semester (PCI) (Main & Backlog) Examination, July 2024 Subject: Biostatistics & Research Methodology

Time: 3 Hours Max. Marks: 75

#### **PART-A**

# Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Define the terms Biostatistics & Research.
- 2. The absorbance values of aspirin solution obtained by UV- Visible spectrophotometer as 0.273, 0.275, 0.271, 0.274, 0.275, 0.279, 0.278 and 0.281. Calculate the mean absorbance value.
- 3. Mention applications of regression in Pharmacy.
- 4. Write the properties of normal distribution.
- 5. Give different non-parametric tests.
- 6. What is the need for Research?
- 7. List out MS EXCEL statistical functions.
- 8. Write features of SPSS.
- 9. Mention advantages of factorial design.
- 10. Give different phase of clinical trials.

#### PART-B

# Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Explain about report writing in research methodology.
- 12. Discuss in detail about one way ANOVA with one example.
- 13. Write the details of response surface methodology.

#### **PART-C**

# Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Define sampling. Explain different sampling techniques.
- 15. The relative humidity in tablet production department of a pharmaceutical manufacturing unit is given below. Calculate the standard deviation in percent relative humidity.

DAY	1	2	3	4	5	6
Х	60	62	65	69	75	65

- 16. Explain the theory of probability.
- 17. The following figure shows disease count from a region over a period of 1 year. Represent the data by a pie diagram.

DISEASE	COUNT		
Jaundice	22		
Tuberculosis	18		
Typhoid	32		
Malaria	15		
Dengue	26		

- 18. What is experimental design? Write its principles.
- 19. Write notes on MINITAB.
- 20. Define Plagiarism. Write the types of it.
- 21. Discuss student t-test in brief.
- 22. Explain 2<sup>2</sup> factorial design with an example.

Code No: F-7268/PCI

# **FACULTY OF PHARMACY**

# B. Pharmacy VIII - Semester (PCI) (Main & Backlog) Examination, July 2024 Subject: Social & Preventive Pharmacy

Time: 3 Hours Max.Marks:75

#### PART-A

#### Note: Answer all the questions

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Mention the social causes of disease.
- 2. Define health and hygiene.
- 3. Write a short note on types of respiratory tract infections.
- 4. What is the mode of transmission of the Ebola virus?
- 5. Write the functions of the pulse polio program.
- 6. What are the objectives and national program for the prevention and control of deafness?
- 7. Write about the social health program.
- 8. Write functions of WHO.
- 9. Write down the objectives of improvement in rural sanitation.
- 10. What is the importance of health education?

#### PART-B

#### Note: Answer any two questions

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. (a) Define malnutrition and write about its types and prevention.
  - (b) What are different avoidable habits from the health and hygiene point of view?
- 12. (a) What is SARS. Write its symptoms, prevention, and control.
  - (b) Elaborate community services with health promotion activities in school.
- 13. (a) Write in detail about the Integrated disease surveillance program.
  - (b) Explain the national tobacco control program.

#### PART-C

# Note: Answer any seven questions

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Explain the concept of prevention and control of the disease.
- 15. Explain the prevention and control of diabetes mellitus.
- 16. Write the mode of transmission, prevention, and control of cholera.
- 17. Explain national mental health program objectives, functioning, and outcomes.
- 18. Explain the objectives, functioning and outcomes of the national program for the control of blindness.
- 19. Explain the national malaria prevention program.
- 20. Explain in detail the national family welfare program.
- 21. Write a note on the objectives, functions and staffing pattern of PHC.
- 22. Define community health. Classify and explain the principles of community health services.

Code No: F-7272/PCI

# **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Main & Backlog) Examination, July 2024 Subject: Quality control and standardization of herbals (Elective-I)

Time: 3 Hours Max. Marks: 75

**PART-A** 

Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Define Palisade Ratio.
- 2. Write any four natural pesticide and their uses.
- 3. What are the application of Gas Chromatography?
- 4. What is secondary processing of medicinal plants?
- 5. Write any two biological markers in standardization of herbal products.
- 6. Write any two identification tests for glycosides.
- 7. Lycopodium Spore Method Formula.
- 8. Mention any four examples of herbal drug interactions.
- 9. What are weedicides? Mention two examples.
- 10. Test for teratogenicity.

**PART-B** 

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Describe the guidelines on GACP for medicinal plants.
- 12. Explain the quality control of herbal drugs as per WHO guidelines.
- 13. Enumerate the regulatory requirement of herbal drugs.

**PART-C** 

Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Explain the various herbal pharmacopoeias.
- 15. Write the GMP requirement for herbal medicine.
- 16. Explain the preparation of document for new drug application.
- 17. Discuss the protocol for clinical guidelines in herbal medicine.
- 18. Enumerate various aspects of GLP.
- 19. Write the applications of chromatography technique its standardization of herbal drugs.
- 20. Write briefly about stability studies of herbal medicinal products.
- 21. Describe the basic test for medicinal plant materials.
- 22. Write a note on research guidelines for evaluating safety and efficacy of herbal medicine.

Code No: F-7271/PCI

# **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Main & Backlog) Examination, July 2024 Subject: Pharmacovigilance (Elective - I)

Time: 3 Hours Max. Marks: 75

#### **PART-A**

#### Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. What is a Good Pharmacovigilance Practice?
- 2. What is the significance of Vigimed and Vigiflow?
- 3. Define probabilistic method.
- 4. Explain adverse events following immunization.
- 5. Write a detailed note on ICD
- 6. Describe phase I of clinical trial.
- 7. Explain WHO scale.
- 8. Describe the Types of services provided by CROs.
- 9. Define the terms (a) Case Reports (b) Cohort studies
- 10. Define WHO drug dictionary.

#### PART-B

# Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Explain in detail on communication with regulatory agencies, business partners. Describe health care facilities and media.
- 12. Describe pharmacogenetic variations attributed to CYP450 isoenzymes inhibition and induction.
- 13. Briefly explain International Non-Proprietary Name(INN) for drugs.

## **PART-C**

#### Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Discuss on organization of WHO-DD.
- 15. Describe the classification and significance of adverse events following immunization programme.
- 16. Explain individual case safety reports.
- 17. Discuss on the history of ICD.
- 18. Write about drug safety evaluation in pediatrics.
- 19. Write a short note on types of vaccine failure.
- 20. Write about the Pharmacovigilance program of India.
- 21. Briefly describe safety evaluation at pre-clinical and clinical trial phase.
- 22. What is the importance of CIOMS in Pharmacovigilance?

Code No: F-7270/PCI

# **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Main & Backlog) Examination, July 2024 Subject: Pharmaceutical Regulatory Science (Elective - I)

Time: 3 Hours Max. Marks: 75

#### **PART-A**

# Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Discuss briefly about Purple Book
- 2. Define the terms (a) WHO (b) CDSCO (c) EMA
- 3. What is a generic product?
- 4. What is meant by 'double blind trial'?
- 5. Give a note on Investigational new drug.
- 6. What is Phase 3 clinical trial?
- 7. Define ASEAN common technical documents (ACTD).
- 8. Mention the different types of DMFs.
- 9. What are the functions of CDSCO?
- 10. What is good clinical practice

#### PART-B

# Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Write a detailed on the following.
  - (a) Timeline and types of IND.
  - (b) Institutional review board.
- 12. How to manage and monitor clinical trials.
- 13. Write about common Technical Document.

#### PART-C

# Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Compare the documentation requirements of ANDA and NDA submissions.
- 15. Discuss about various Stages of drug discovery
- 16. Describe general check list for 21CFR part 11.
- 17. What is the general procedure for export of pharmaceutical product?
- 18. Explain the GCP obligations of investigators and sponsers
- 19. Describe in detail new drug approval process along with its documentation requirements as per USFDA.
- 20. How innovator drug are different from generics drugs?
- 21. What is the constitution and purpose of Ethics Committee?
- 22. Explain the Orange Book features.

Code No: F-7269/PCI

# **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Main & Backlog) Examination, July 2024 Subject: Pharmaceutical Marketing Management (Elective-I)

Time: 3 Hours Max. Marks: 75

PART - A

Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Define Marketing and selling.
- 2. List factors influencing the choice of physician.
- 3. List the stages of product life cycle.
- 4. Classify pharmaceutical market.
- 5. What is consumer profiling?
- 6. What is retailing and mention its advantages.
- 7. Write the roles and responsibilities of distributors?
- 8. List the duties of professional sales representatives.
- 9. What is vertical marketing?
- 10. Define pricing and mention its objectives.

PART-B

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Explain various approaches of market research for analysing market.
- 12. Describe the different channels in distribution management and mention the conflicts to be considered while selecting them.
- 13. Explain different pricing methods and strategies.

**PART-C** 

Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Explain the factors affecting industrial buying behaviour.
- 15. What is product portfolio analysis and its role in product positioning?
- 16. Write the roles of advertising and public relations in promotion of pharmaceutical products.
- 17. Explain the steps involved in personal selling.
- 18. Describe the role of journals and medical exhibition in product promotion.
- 19. Write the evaluation criteria and compensation planning for professional sales representatives.
- 20. Write the regulatory norms applicable to customer calls.
- 21. Describe the determinants for fixation of prices.
- 22. Write the functions and role of NPPA.

Code No: F-7273/PCI

# **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Main & Backlog) Examination, July 2024 Subject: Computer-Aided Drug Design (Elective-I)

Time: 3 Hours Max. Marks: 75

PART-A

Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Write a note on serendipitous discovery of drugs.
- 2. Define the term lead optimization.
- 3. Differentiate between SAR and QSAR.
- 4. Write about Free-Wilson analysis.
- 5. What is virtual screening?
- 6. What is rigid docking?
- 7. Define the terms: Bioactive conformer & Force field.
- 8. What is global conformational minima?
- 9. Enlist the applications of bioinformatics tools in drug design.
- 10. Give examples for pharmaceutical databases.

PART-B

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. What is 3D QSAR? Write about CoMFA and CoMSIA methods.
- 12. Describe the concept of pharmacophore-based virtual screening in drug design.
- 13. Define and classify bioisosteres. Explain its significance in drug design with suitable examples.

**PART-C** 

Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Explain about Hansch analysis with a model QSAR equation.
- 15. Highlight the significance of partition coefficient in QSAR analysis and write its determination.
- 16. Describe metabolism-based lead discovery with specific examples.
- 17. Explain the steps involved in molecular docking.
- 18. Explain drug-likeness screening along with the tools used for its determination.
- 19. List out various protein databases. Explain the significance of these databases in drug design with a specific example.
- 20. Discuss the significance of in silico ADME databases in drug design.
- 21. Discuss briefly about conformational analysis.
- 22. Explain various stages involved in molecular mechanics.

# **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Main & Backlog) Examination, July 2024
Paper: Cell and Molecular Biology (Elective-II)

Time: 3 Hours Max. Marks: 75

PART-A

Note: Answer all questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Write a brief note on functions of m-RNA?
- 2. Write the functions of cell membrane.
- Differentiate between t-RNA and m-RNA
- 4. Write the significance of Lac operon pathway.
- 5. Define Transgenics.
- 6. Define genetic code.
- 7. Enlist the functions of Okasaki fragments.
- 8. What are SSB proteins?
- 9. Enlist the properties of the cells.
- 10. What are spindle fibres?

PART-B

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Describe in detail about the enzymes involved in DNA replication.
- 12. Write in detail about cell signaling pathways and its misregulation.
- 13. What are chromosomes? Write a detailed account on discovery, structure, number and significance of chromosomes in prokaryotic and eukaryotic cells. Draw labelled diagrams wherever necessary.

**PART-C** 

Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Explain the DNA replication mechanism in eukaryotes.
- 15. Explain in detail significance of protein synthesis.
- 16. Write an account on the types of RNA. Discuss their functions.
- 17. Describe the stages of mitosis.
- 18. What is Bacterial Transduction? Explain the process of Transduction in Bacteria.
- 19. What are the structural and regulatory genes? Explain genetic control of protein synthesis.
- 20. Discuss the role of the enzyme DNA ligase plays during DNA replication.
- 21. Describe the stages of prophase -1 of meiosis.
- 22. Construct a complete transcription unit with promnoter and terminator on the basis ATGCATGCATAC

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Code No: F-7275/PCI

### **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Main & Backlog) Examination, July 2024 Subject: Cosmetic Science (Elective-II)

Time: 3 Hours Max. Marks: 75

#### **PART-A**

Note: Answer all questions.  $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Define Cosmetic and Cosmeceuticals.
- 2. What are the preservatives used in cosmetic products?
- 3. Write a note on face wash.
- 4. Write a note on mouth washes.
- 5. What are the uses of clove in oral care?
- 6. Write a note on Tewameter (TEWL).
- 7. Differentiate between soaps and syndet bars.
- 8. Write a note on hair combing properties.
- 9. Write a note on comedogenic and dermatitis.
- 10. What is the reasons for bad body odour.

#### **PART-B**

#### Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Draw basic structure of skin and write functions of skin. Explain formulation of Moisturizing cream and Cold Cream as skin care products.
- 12. Write the role of herbs in cosmetics. Write the role of aloe, turmeric and neem in cosmetic formulation.
- 13. Write about blemishes, wrinkles, acne and prickly heat in skin problems.

#### PART-C

# Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Write the classification of cosmetics and cosmeceuticals with examples.
- 15. Write the classification and applications of i) Surfactants ii) Rheology modifiers
- 16. Write formulation and mechanism of action of Antiperspirants & deodorants.
- 17. Write the formulation of toothpaste for bleeding gums and sensitive teeth.
- 18. Classify sunscreen formulations and explain SPF.
- 19. Write a note on evaluation of Shampoo and skin cream as per BSI.
- 20. Write the principles and applications of Sebumeter and Corneometer
- 21. Write a note on oily and dry skin. Write causes for dry skin.
- 22. Write the hair fall causes and dandruff.

Code No: F-7278/PCI

# **FACULTY OF PHARMACY**

B. Pharmacy VIII – Semester (PCI) (Main & Backlog) Examination, July 2024
Subject: Experimental Pharmacology
(Pharmacological Screening Methods)(Elective-II)

Time: 3 Hours Max. Marks: 75

PART-A

Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. What are common laboratory animals?
- 2. Differentiate between sedative and hypnotic agents.
- 3. Write the composition of IAEC.
- 4. What is one way ANOVA?
- 5. Enlist various techniques of blood collection in common laboratory animals
- 6. Write about Mutant animals.
- 7. What are nootropics?
- 8. What is the importance of sham negative and positive control groups?
- 9. List out screening methods for drugs acting on eye.
- 10. Write about Euthanasia?

#### PART-B

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Explain various *in vivo*, *in vitro* methods to evaluate a compound for antidiabetic activity.
- 12. Describe in detail about regulations for laboratory animal care as per CPCSEA Guidelines.
- 13. Write in detail about the methods of screening for antihypertensives and anti arrhythmics.

#### **PART-C**

Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Write a note on *in vivo* methods for screening antiepileptic drugs.
- 15. Explain the screening methods for skeletal muscle relaxants.
- 16. Describe various routes of drug administration in animals with advantages and disadvantages.
- 17. Write a brief note on screening methods of analgesic drugs.
- 18. Explain the applications of transgenic animals in pharmacological research.
- 19. Write any two preclinical screening methods for antidepressant activity.
- 20. Enumerate any two preclinical screening methods for local anaesthetics.
- 21. Explain the criteria of dose selection and calculations of dose for animals.
- 22. Explain the interpretation of results using Students-t test?

Code No: F-7276/PCI

# **FACULTY OF PHARMACY**

B. Pharmacy VIII – Semester (PCI) (Main & Backlog) Examination, July 2024 Subject: Dietary Supplements and Nutraceuticals (Elective-II)

Time: 3 Hours Max. Marks: 75

## **PART-A**

Note: Answer all questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Write short notes on Reactive Oxygen Species.
- 2. Define flavonoids and polyphenolics.
- 3. Give the source, chemical nature and uses of Tea and coffee.
- 4. Explain the importance of synthetic anti-oxidants.
- 5. Write about GMPs on food safety.
- 6. Write about complex carbohydrates as functional food ingredients.
- 7. Give the occurrence and medical benefits of Lignans and Rutin.
- 8. What are phyto estrogens?
- 9. Write about production of free radicals in cells.
- 10. Write about storage potential of neutraceuticals.

#### **PART-B**

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. What are various endogeneous antixodants give its enzymatic and non enzymatic defence mechanism of action.
- 12. Explain various mechanisms of free radicals involved in Diabetes milletus and renal failure.
- 13. Write the various components of dietary supplements and their applications. Add a note on deficiency of dietary supplements.

#### **PART-C**

Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Explain in detail about Carotenoids.
- 15. Give the occurrence, chemical nature and uses of Gingko and Ginseng.
- 16. Explain the regulatory aspects of FSSAI on food safety.
- 17. Explain the role of free radicals with lipids.
- 18. Write in detail about adulteration of food.
- 19. Explain in detail about sulphides and xanthophylls.
- 20. Give the importance of proteins and vitamins as functional food.
- 21. Explain the free radicals theory of ageing.
- 22. Write about various pharmacopoeial specification of neutraceuticals.

Code No: F-7277/PCI

# **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Main & Backlog) Examination, July 2024 Subject: Advanced instrumentation techniques (Elective-II)

Time: 3 Hours Max. Marks: 75

## **PART-A**

# Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. What are the Molecular ions in MS?
- 2. List the different ionisation techniques in MS.
- 3. Explain Sheilding and Desheliding in NMR.
- 4. Define chemical shift. List the factors affecting chemical shift
- 5. What is the internal standard in HPLC? Justify its selection.
- 6. What is the principle of Thermogravimetric Analysis (TGA)?
- 7. List the important steps in solid phase extraction.
- 8. Give suitable applications of radioimmunoassay.
- 9. List the parameters for the calibration of UV Visible spectrophotometer.
- 10. Define validation? List out various validation parameters.

#### **PART-B**

# Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Explain the principle of Mass spectrometry. With a labelled diagram, explain MS instrumentation.
- 12. Explain the Principle and instrumentation of RP-HPLC?
- 13. Explain the principle and applications of LC-MS/MS.

#### **PART-C**

#### Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. With a neat labelled diagram explain the instrumentation and application of DTA.
- 15. Explain Fragmentation techniques in MS. Explain any two methods in detail.
- 16. Explain Time of flight and quadrupole mass analysers in MS.
- 17. What are the differences between C13 and H1 NMR spectroscopy?
- 18. Explain the origin of X-rays. Derive Bragg's equation
- 19. Explain the phenomena of spin spin coupling with a suitable examples.
- 20. Explain the principle and procedure involved in liquid-liquid extraction.
- 21. What is the difference between calibration and validation? List the validation parameters as per ICH guidelines and explain any two.
- 22. Explain the principle, advantages and applications of RIA.

Code No: F-7195/PCI

# **FACULTY OF PHARMACY**

# B. Pharmacy VIII - Semester (PCI) (Backlog) Examination, March 2024 Subject: Biostatistics & Research Methodology

Time: 3 Hours Max. Marks: 75

#### PART-A

# Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Find the median of the disintegration times of tablets for a data set 1, 4, 8, 3, 5, 7, 2, 10 and 6.
- 2. Calculate the range for individual series-X: 120,170,240,100, 105, 205, 300, 160, 150, 180.
- 3. Define the term Probability.
- 4. What is meant by population?
- 5. Give different non-parametric tests.
- 6. What is the need for design of experiments?
- 7. Mention different types of graphs.
- 8. Write statistical features of EXCEL.
- 9. Mention advantages of factorial design.
- 10. Give different phase of clinical trials.

#### PART-B

# Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Explain about report writing in research methodology.
- 12. Write informative notes on (a) Plagiarism (b) Types of sampling.
- 13. What is SPSS? Write about its models.

#### **PART-C**

# Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Discuss about Karl Pearson's coefficient of correlation.
- 15. Find the standard deviation of incubation period of small pox in 9 patients where it was found to be 15,12,10,15,11,7,9,17 and 14.
- 16. What is normal distribution? Explain the properties with a suitable example.
- 17. Write short notes on ANOVA.
- 18. A quality control analyst finds that on the average the sample passes the test 4 times out of 5. If the sample is tested 4 times, what is the probability of
  - (a) Sample passing more than 2 times
- (b) At least 3 failures
- 19. Write notes on any one non-parametric test.
- 20. Discuss about statistical features of MINITAB.
- 21. Explain designing of a clinical trial.
- 22. Explain 2<sup>2</sup> factorial design with an example

Code No: F-7205/PCI

# **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Backlog) Examination, March 2024 Subject: Advanced instrumentation techniques (Elective-II)

Time: 3 Hours Max. Marks: 75

#### **PART-A**

#### Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. What are the important steps in MS?
- 2. List the different ionisation techniques in MS.
- 3. Explain Base peak and molecular ion in MS.
- 4. Define chemical shift. List the factors affecting chemical shift
- 5. What is the internal standard in NMR spectroscopy? Justify its selection.
- 6. What is the principle of TGA?
- 7. List the important steps in solid phase extraction.
- 8. Give suitable applications of radioimmunoassay.
- 9. List the parameters for the calibration of UV Visible spectrophotometer.
- 10. What is the difference between calibration and validation?

#### **PART-B**

# Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Explain the principle of Mass spectrometry. With a labelled diagram, explain MS instrumentation.
- 12. Explain HPLC calibration process.
- 13. Explain the principle of LC/MS/MS.

#### PART-C

#### Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. With a neat labelled diagram explain the instrumentation and application of DTA.
- 15. Classify the ionisation techniques in MS. Explain any two methods in detail.
- 16. Explain Time of flight and quadrupole mass analysers in MS.
- 17. What are the difference s between C<sup>13</sup> and H<sup>1</sup> NMR spectroscopy?
- 18. Explain the origin of X-rays. Derive Bragg's equation
- 19. Explain the phenomena of spin spin coupling with a suitable example.
- 20. Explain the principle and procedure involved in liquid-liquid extraction.
- 21. Write a note on hyphenated techniques. Give suitable examples. What are their advantages? Add a note on interfaces.
- 22. Explain the principle advantages and applications of RIA.

# **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Backlog) Examination, March 2024
Paper: Cell and Molecular Biology (Elective–II)

Time: 3 Hours Max. Marks: 75

#### **PART-A**

Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Differentiate Prokaryotic cell versus Eukaryotic cell.
- 2. Write a note on power house of the cell.
- 3. Differentiate between DNA and RNA.
- 4. Write the components of Lac-operon.
- 5. Define chromatin.
- 6. What is osmosis and diffusion?
- 7. Differentiate SER and RER.
- 8. Discuss the role of DNA ligase during DNA replication.
- 9. Mention different sub-stages of prophase -2 of meiotic cell division.
- 10. Differentiate microtubules and microfilaments.

#### PART-B

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. What is Bacterial Transduction? Explain the process of Transduction in Bacteria.
- 12. What are the structural and regulatory genes? Explain genetic control of protein synthesis.
- 13. Explain about giant chromosomes with their structure, functions of nucleus and its components.

#### **PART-C**

Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Write an account on the types of RNA. Discuss their functions.
- 15. Write in detail about cell signaling pathways and its misregulation.
- 16. Explain the role of DNA -dependent RNA polymerse in transcription.
- 17. Distinguish between mitosis and meiosis with appropriate diagrams.
- 18. Write a short note on classification of cell types.
- 19. Describe the Watson and Crick model of DNA structure with labelled diagram.
- 20. Explain in detail functioning of protein kinases.
- 21. Write in detail about definition, theory, basics and applications of cell and molecular Biology.
- 22. Explain Chargaff's law.

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Code No: F-7203/PCI

# **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Backlog) Examination, March 2024 Subject: Cosmetic Science (Elective-II)

Time: 3 Hours Max. Marks: 75

#### **PART-A**

# Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Write a note on preservatives used in cosmetics
- 2. Explain cosmetics as Quasi and OTC drugs.
- 3. What are emollients?
- 4. Write a note on moisturizing cream.
- 5. Explain sun protection formulations.
- 6. What are mouthwashes?
- 7. Write the role of neem in oral care.
- 8. What is the difference between soap and syndet bar.
- 9. What are the reasons and prevention of dry skin.
- 10. Write a note on reasons and prevention of body odor.

#### PART-B

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Write a brief note on following excipients with examples a) Surfactantsb) Humectantsc) Rheology modifiersd) Emollientse) Preservatives
- 12. Write the causes and prevention of blemishes, wrinkles, acne and hair fall.
- 13. Explain Sebumeter, Corneometer, Tewameter (TEWL) in cosmetic evaluation.

#### **PART-C**

#### Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Classify cosmetics and cosmeceuticals with examples.
- 15. Write a note on basic structure of hair and hair growth cycle.
- 16. Write the principle involved in formulation of cold cream and vanishing cream.
- 17. Write formulation and mechanism of action of Antiperspirants & deodorants.
- 18. Write a note on conditioning shampoo, antidandruff shampoo in hair care.
- 19. Write the formulation of toothpaste for bleeding gums and sensitive teeth.
- 20. Write a note on henna and amla in hair care.
- 21. Write the causes and prevention of blemishes, wrinkles and acne.
- 22. Discuss the role of importance of Aloe and turmeric in Herbal cosmetics.

Code No: F-7206/PCI

### **FACULTY OF PHARMACY**

# B. Pharmacy VIII – Semester (PCI) (Backlog) Examination, March 2024 Subject: Experimental Pharmacology (Pharmacological Screening Methods)(Elective-II)

Time: 3 Hours Max. Marks: 75

PART-A

Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. List a few laboratory animals and their use in research.
- 2. What are transgenic animals?
- 3. List the common routes of drug administration in animals.
- 4. What are coagulants and anticoagulants?
- 5. List out the drugs acting on the eye. Name the models.
- 6. What is Euthanasia and list the techniques of euthanasia.
- 7. List various agents which cause inflammation.
- 8. How is dose selected in preclinical screening methods?
- 9. What is Students-t test and where is it used?
- 10. What is preclinical data analysis?

PART-B

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Describe the screening models for evaluation of a compound for Antihypertensive activity.
- 12. Discuss the *in vitro* and *in vivo* techniques for screening of anticancer agents.
- 13. Describe in detail about regulations for laboratory animal care as per CPCSEA guidelines.

**PART-C** 

Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Write a brief note on screening methods of antinflammatory drugs.
- 15. What is Research? Mention the significance of selection of research topic.
- 16. Explain the screening methods for diuretics.
- 17. Describe the techniques for collection of blood in the animals?
- 18. Write about One-way ANOVA and its importance in preclinical studies.
- 19. Write a note on methods involved in the screening of nootropics.
- 20. Enumerate any two preclinical screening methods for local anaesthetics.
- 21. What are antiasthamatic agents? Discuss the methods involved in their screening.
- 22. Write the preclinical screening methods of sympathomimetics.

Code No: F-7204/PCI

# **FACULTY OF PHARMACY**

B. Pharmacy VIII – Semester (PCI) (Backlog) Examination, March 2024 Subject: Dietary Supplements and Nutraceuticals (Elective-II)

Time: 3 Hours Max. Marks: 75

**PART-A** 

Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Write the difference between dietary supplements and neturaceuticals.
- 2. Write about polyphenols and tocopherols.
- 3. Give the occurrence and medical benefits of Lycopene.
- 4. Write about dietary fibres as functional food ingredients.
- 5. Give the source, chemical nature and uses of Oats and Rice bran.
- 6. Explain about enzymatic antioxidant defence.
- 7. What are phytosterols give its uses?
- 8. Write about AGMARK on food safety.
- 9. Write the benefits of Public health nutrition.
- 10. Name the marker compounds of spirulina and ginko.

#### PART-B

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Explain in detail the effect of processing, storage and interactions of various environmental factors on the potential of dietary supplements.
- 12. (a) Explain the role of antioxidants in the treatment of Cancer.
  - (b) Write about various nutritional benefits in a community.
- 13. (a) Classify various nutraceuticals with examples.
  - (a) Explain the role of Reactive Oxygen Species involvement in the treatment of disorders.

#### **PART-C**

Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Explain the role of anti-oxidants in the treatment of kidney damage.
- 15. Give the pharmacopeial specification for complex carbohydrates.
- 16. Explain the regulatory aspects of FSSAI on food safety.
- 17. Explain the role of melatonin, Vitamin E and Catalase.
- 18. Define flavonoids and give the source and medicinal benefits of any two flavonoids.
- 19. Write in detail about adulteration of foods.
- 20. Explain the role of various endogenous anti-oxidants.
- 21. Give the importance of proteins and vitamins as functional foods.
- 22. Give the occurrence, chemical nature and uses of Garlic and Flax seeds.

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Code No: F-7197/PCI

# **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Backlog) Examination, March 2024 Subject: Pharmaceutical Marketing Management (Elective-I)

Time: 3 Hours Max. Marks: 75

#### **PART-A**

# Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Differentiate between Marketing and selling.
- 2. Classify pharmaceutical products.
- 3. What is product branding?
- 4. What are the free samples and norms applicable to them,
- 5. What is significance of direct mail in product promotion?
- 6. Write the quantitative and qualitative aspects of pharmaceutical market.
- 7. Name different channels of distribution.
- 8. List factors influencing the choice of retail pharmacist.
- 9. What is product line and give example.
- 10. Write the merits and demerits of wholesale distribution channel.

#### PART-B

### Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Explain stages in product life cycle and mention its role in product positioning and new product decisions.
- 12. Describe the selection, training and future prospects of professional sales representatives.
- 13. Compare and contrast between rural, industrial and global marketing.

#### PART-C

#### Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Explain the factors affecting consumer buying behaviour.
- 15. Describe salient features applicable for product packaging and labelling.
- 16. Explain the demographic descriptions and socio-psychological characteristics of consumers.
- 17. Write the motivational factors and prescribing habits of physician.
- 18. Write the promotional mix factors to be considered for fixing the promotional budget.
- 19. Explain online promotional techniques relevant to OTC products.
- 20. Differentiate between vertical and horizontal marketing.
- 21. Write the challenges of price management in pharmaceutical marketing.
- 22. Write an overview on DPCO and its functions.

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# **FACULTY OF PHARMACY**

# B. Pharmacy VIII - Semester (PCI) (Backlog) Examination, March 2024 Subject: Pharmaceutical Regulatory Science (Elective-I)

Time: 3 Hours Max. Marks: 75

#### PART-A

#### Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. What are the functions of US regulatory authority?
- 2. What are non-clinical studies?
- 3. Write the difference between brand and generic products.
- 4. Describe the modules in ACTD.
- 5. Mention the general list of CFR.
- 6. Write a note on purple book.
- 7. Define a. eCTD b. CFR.
- 8. List out the items in module III in ANDA.
- 9. Explain the exclusion criteria for clinical trials.
- 10. Define ICF.

#### PART-B

# Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Discuss different stages of preclinical studies.
- 12. Write the process for export of pharmaceutical products.
- 13. Explain in detail on DMF system in India.

#### **PART-C**

# Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Differentiate CTD and eCTD.
- 15. Write a note on submission of DMF.
- 16. Explain the protocol of clinical trials.
- 17. What are the stages of drug discovery process?
- 18. Write the salient features of pharmacovigilance.
- 19. Discuss code of federal regulation.
- 20. Explain the objectives of regulatory affairs department in pharma industry.
- 21. Explain the different modules of CTD in detail.
- 22. Write the steps involved in changing an approved NDA /ANDA.

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# **FACULTY OF PHARMACY**

# B. Pharmacy VIII - Semester (PCI) (Backlog) Examination, March 2024 Subject: Pharmacovigilance (Elective - I)

Time: 3 Hours Max. Marks: 75

PART-A

Note: Answer all the questions.  $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Explain Development Safety Update Report(DSUR)
- 2. What are the concepts of DDD?
- 3. Define Data mining
- 4. Give the purpose of MedDRA.
- 5. Enlist any two source of ADR reporting.
- 6. Define pharmacogenetics and pharmacogenomics.
- 7. What is post approval phase?
- 8. What is Vaccine pharmacovigilance.
- 9. What are responsibilities of CROs?
- 10. Describe CIOMS working group.

PART-B

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Explain individual reporting and spontaneous 'epo' ling.
  Enlist the steps recommended by WHO for establishing a PV centre.
- 12. Explain about Good clinical practices in pharmacovigilance.
- 13. Write a detailed note on ICD.

**PART-C** 

Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Describe in detail the organisational structure & functions of CDSCO in India.
- 15. Write the differences in Indian and global pharmacovigilace requirements.
- 16. Describe drug safety in pregnancy and lactation.
- 17. Explain in detail the organization and objectives of ICH
- 18. Write about effective communication in drug safety crisis management
- 19. Explain the following a. case control study b. cohort study.
- 20. List out the adverse events following immunization.
- 21. Discuss the establishment of national pharmacovigilance program.
- 22. Write about anatomical and therapeutic classification of drugs.

Code No: F-7200/PCI

# **FACULTY OF PHARMACY**

B. Pharmacy VIII - Semester (PCI) (Backlog) Examination, March 2024 Subject: Quality control and standardization of herbals (Elective-I)

Time: 3 Hours Max. Marks: 75

#### PART-A

#### Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- Define Stomata.
- 2. What is the difference between TLC & HPTLC?
- 3. Write the names of four markers.
- 4. What is Extractive value and its significance?
- 5. Define SOP.
- 6. What is AYUSH?
- 7. Write four examples of herbal drug interactions.
- 8. What is the significance of ICH?
- 9. Define term herbal medicine & crude drug.
- 10. What is Quantitative microscopy?

#### PART-B

# Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. Describe WHO guidelines for quality control of herbal drugs.
- 12. Explain the infra structural requirements under GMP for herbal industry.
- 13. Describe the preparation of documents for new drug application and export registration.

#### **PART-C**

#### Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. What is GAP? Explain the various parameter of GAP.
- 15. Explain the importance of HPTLC method in the standardization of herbal drugs.
- 16. Write a note on regulatory requirement for herbal drugs.
- 17. Describe the guidelines on safety and efficacy of herbal medicine.
- 18. Explain WHO guidelines on current good manufacturing practices for herbal medicine.
- 19. Discuss about the assessment of Genotoxicity of herbal preparations.
- 20. Define and classify markers with examples.
- 21. Describe the basic test for medicinal plant material.
- 22. Explain ICH guidelines for the quality control of herbal drugs.

Code No: F-7201/PCI

# **FACULTY OF PHARMACY**

# B. Pharmacy VIII - Semester (PCI) (Backlog) Examination, March 2024 Subject: Computer Aided Drug Design (Elective-I)

Time: 3 Hours Max. Marks: 75

#### PART-A

### Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. What is random screening? Give an example.
- 2. What are the advantages of virtual screening over conventional techniques?
- 3. Explain the significance of the partition coefficient.
- 4. Define the terms: Molecular mechanics and quantum mechanics.
- 5. What is 3D QSAR?
- 6. Define and differentiate the following terms: Lead and Drug.
- 7. Write the significance of PDB in drug design.
- 8. Describe De novo drug design.
- 9. What are pharmaceutical databases? Give examples.
- 10. What is global conformational minima?

#### **PART-B**

Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. What is QSAR analysis? List out various physicochemical parameters and explain about electronic parameters. Provide a model QSAR equation.
- 12. Explain various stages involved in drug discovery and development.
- 13. Describe the significance of various bioinformatics tools used in drug design with suitable examples.

#### **PART-C**

# Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Explain Hansch analysis with its advantages & disadvantages.
- 15. What is molecular docking? Explain its significance in drug design.
- 16. Write about various energy minimization methods.
- 17. Explain chemical databases with suitable examples. Give their importance in drug design.
- 18. Write a note on serendipitous discovery of drugs.
- 19. Discuss the methodology involved in CoMSIA.
- 20. Explain drug-likeness screening. Write various tools used for the same.
- 21. Define and differentiate various types of bioisosteres with suitable examples.
- 22. Explain the role of quantum mechanics in drug design.

Code No: F-7196/PCI

# **FACULTY OF PHARMACY**

# B. Pharmacy VIII - Semester (PCI) (Backlog) Examination, March 2024 Subject: Social and Preventive Pharmacy

Time: 3 Hours Max.Marks:75

#### PART - A

#### Note: Answer all the questions.

 $(10 \times 2 = 20 \text{ Marks})$ 

- 1. Write a note on symptoms of vitamin deficiencies?
- 2. Define balanced diet.
- 3. What is the difference between drug abuse and drug addiction?
- 4. Explain the social causes of the disease?
- 5. Explain different types of diabetes mellitus?
- 6. Write the various objectives of HIV and AIDS control programme?
- 7. Write a note on Malaria control strategies?
- 8. What are the objectives of integration with National urban Health Mission (NUHM)?
- 9. Write a note on school health promotion program?
- 10. Write a note on the public health care system in India?

#### PART - B

# Note: Answer any two questions.

 $(2 \times 10 = 20 \text{ Marks})$ 

- 11. (a) Explain about Malnutrition and various methods of its prevention.
  - (b) Explain general principles of prevention and control of Malaria.
- 12. (a) Explain the objectives, functioning and outcomes of national mental health programme.
  - (b) Write a note on role of WHO in Indian National programmes.
- 13. (a) Discuss in detail about National family welfare programme.
  - (b) Write a note on functions of PHC.

#### PART - C

# Note: Answer any seven questions.

 $(7 \times 5 = 35 \text{ Marks})$ 

- 14. Write the various risk factors and diagnosis of Cancer.
- 15. Explain signs, symptoms, transmission and treatment of SARS.
- 16. Define health and explain different dimensions of good health.
- 17. Write a note on objectives and strategies for Leprosy elimination in India.
- 18. Write a note on pulse polio program.
- 19. Write a note on national programme for health care of elderly.
- 20. What are the aims and achievements of National Tobacco Program?
- 21. Explain about the different levels of Evaluation of public health.
- 22. How can you improve sanitization in rural area, explain different schemes and programs.