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Total Number of Pages: 02

Course: B.Pharm
Sub_Code: BP701T

7th Semester Regular/Back Examination: 2024-25

SUBJECT: Instrumental Methods of Analysis

BRANCH(S): B.Pharma

Time: 3 Hours

Max Marks: 75

Q.Code: R616

Answer Question No.1 (Part-1) which is compulsory, any eight from Part-II and any two from Part-III.

The figures in the right hand margin indicate marks.

Part-I

Q1 Answer the following questions : (2 x 10)

- What are chromophores and auxochromes?
- Define Fluorescence and Phosphorescence.
- Differentiate between normal phase & reverse phase chromatography.
- What are the different electronic transitions that occur due to the absorption of UV radiation?
- What is finger print region? Give its significance.
- What is the difference between isocratic and gradient elution in chromatography?
- What is the principle involved affinity chromatography?
- What is molar extinction co-efficient?
- Write the principle of ion exchange chromatography.
- Name two detecting reagents used in thin layer and paper chromatography.

Part-II

Q2 Focused-Short Answer Type Questions- (Answer Any Seven) (5 x 7)

- Write the statement and derive the equation for Beer's – Lambert's law.
- Explain the instrumentation and working of flame emission spectrometry.
- What are Nephelometry and turbidometry? Write principle involved for the same.
- Discuss the instrumentation and application of HPLC.
- Discuss briefly rate and plate theory.
- Explain the principle and theory of gel chromatography.
- Describe the principle, working, and instrumentation of Atomic Absorption Spectroscopy.
- What is Quenching? Enumerate the various factors which influence quenching effect.
- Define electrophoresis and explain the various factors affecting electrophoresis.

Part-III

Long Answer Type Questions (Answer Any Two)

- Q3** Describe the principle, instrumentation, and applications of UV-Visible Spectrophotometer. (10)
- Q4** Write a note on theory and applications of IR spectrophotometry. Explain different sampling techniques employed in IR spectroscopy. (10)
- Q5** Describe in brief the principle, instrumentation, and applications of gas chromatography. (10)
- Q6** Describe principle and different techniques of paper electrophoresis. Add a note on applications of paper electrophoresis. (10)

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Total Number of Pages: 02

Course: B.Pharm
Sub_Code: BP702T

7th Semester Regular/Back Examination: 2024-25

SUBJECT: Industrial Pharmacy II

BRANCH(S): B.Pharm

Time: 3 Hours

Max Marks: 75

Q.Code: R619

Answer Question No.1 (Part-1) which is compulsory, any seven from Part-II, and any two from Part-III.

The figures in the right hand margin indicate marks.

Part-I

Q1 Answer the following questions : (2 x 10)

- Write the role of CDL.
- Mention important data documents for ANDA.
- What is Quality risk management?
- What are the components of non-clinical drug development?
- Mention the importance of NABL accreditation.
- Differentiate between GMP and cGMP.
- Mention various TT agencies in India.
- Write the use of Quality by Design (QbD).
- Mention the objectives of plant scale up techniques.
- Differentiate between qualification and validation.

Part-II

Q2 Focused-Short Answer Type Questions- (Answer Any Seven) (5 x 7)

- Write a note on platform technology and mention the needs of regulatory affairs.
- Discuss the various aspects of TQM
- Give a brief account on SUPAC guidelines.
- What are the quality management systems, discuss.
- Discuss all the responsibilities of state licensing authority.
- Write the importance of Six Sigma concept in detail.
- Explain certificate of pharmaceutical product.
- Write a short note on COPP.
- Explain factors considered in scale-up process of dry granulation.

Part-III

Long Answer Type Questions (Answer Any Two)

- Q3** Explain in detail about central drug standard control organization (CDSCO). (10)
- Q4** What are the clinical research protocols? Briefly discuss on clinical research and BE studies. (10)
- Q5** What are the different types of drug applications that can be submitted to FDA? Discuss in detail about ISO 9000 series of quality systems standards. (10)
- Q6** What is scale up process? Explain in detail about pilot plant scale up considerations for solids. (10)

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Total Number of Pages: 01

Course: B.Pharm
Sub_Code: BP703T

7th Semester Regular/Back Examination: 2024-25

SUBJECT: Pharmacy Practice

BRANCH(S): Pharmacy

Time: 3 Hours

Max Marks: 75

Q.Code: R626

Answer Question No.1 (Part-I) which is compulsory, any seven from Part-II, and any two from Part-III.

The figures in the right hand margin indicate marks.

Part-I

- Q1 Answer the following questions :** (2 x 10)
- a) Define Idiosyncrasy.
 - b) Define and classify hospitals based on number of beds.
 - c) Write code of ethics for community pharmacy.
 - d) Write about beneficial interactions with examples.
 - e) What is patient compliance?
 - f) Define hospital formulary.
 - g) What is drug addiction?
 - h) What genetically determined toxicity.
 - i) What is drug dependence?
 - j) Classify the types of allergic drug reactions.

Part-II

- Q2 Focused-Short Answer Type Questions- (Answer Any Seven)** (5 x 7)
- a) Define and classify ADRs.
 - b) Write notes on PTC.
 - c) What is the importance of communication skills for a Pharmacist?
 - d) Discuss rational use of over the counter medication.
 - e) Brief the functions and responsibilities of clinical pharmacist.
 - f) Explain Therapeutic drug monitoring.
 - g) How to prepare and implement budget in hospital.
 - h) Discuss the purchase procedure of Hospital.
 - i) Write the sources of drug information services.

Part-III

Long Answer Type Questions (Answer Any Two)

- Q3** Define Hospital. Write its organizational structure and explain services in details. (10)
- Q4** Discuss in details Hematological parameters along with their significance. (10)
- Q5** Give details on drug distribution system in hospital with special emphasis on in-patient and out-patient department. (10)
- Q6** Explain various drug interactions in details with suitable examples. (10)

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Total Number of Pages: 01

Course: B.Pharm
Sub_Code: BP704T

7th Semester Regular/Back Examination: 2024-25

SUBJECT: Novel Drug Delivery System

BRANCH(S): B.Pharm

Time: 3 Hours

Max Marks: 75

Q.Code: R634

Answer Question No.1 (Part-1) which is compulsory, any seven from Part-II, and any two from Part-III.

The figures in the right hand margin indicate marks.

Part-I

- Q1 Answer the following questions:** (2 x 10)
- Define NDDS and mention few advantages.
 - What are the limitations of TDDS?
 - Write the major drug release mechanisms from CR dosage forms.
 - What is Mucosal Drug Delivery system? Mention its usefulness.
 - Mention the importance of partition co-efficient in drug designing.
 - Differentiate between CR and SR.
 - Name the chemical cross linking agent mostly used for nano-preparation.
 - Mention the importance of niosomes.
 - What are the general characteristics of Ocular preparations?
 - Define implants.

Part-II

- Q2 Focused-Short Answer Type Questions - (Answer Any Seven)** (5 x 7)
- Write mucosal permeation enhancers with examples
 - Write about Biological factors affecting controlled drug designing.
 - Explain in detail about solid lipid nanoparticles.
 - Discuss about Floating drug delivery system.
 - What is Ion exchange principle for drug release?
 - Differentiate in between liposomes and niosomes.
 - Mention the applications of monoclonal antibodies.
 - What are various approaches of transdermal drug delivery?
 - Define buccal drug delivery and write in detail about its components.

Part-III

Long Answer Type Questions (Answer Any Two)

- Q3** Define liposomes and classify them. Discuss various passive loading methods for its preparation. (10)
- Q4** Define bio-adhesive drug delivery system. What are the various theories involved in bio-adhesion? Describe in detail. (10)
- Q5** Define microencapsulation. Write the applications of microencapsulation. Explain phase separation-Coacervation technique. (10)
- Q6** What is polymer? Classify it and write in detail polymer application in pharmaceutical Formulations. (10)