

BACHELOR OF PHARMACY (PRACTICE)
F.Y. B. Pharm. (Practice) :SUMMER- 2022
SUBJECT : APPLIED PHARMACEUTICS

Day : Friday
Date : 3/6/2022

S-20430-2022

Time : 10:00 AM-01:00 PM
Max. Marks : 60

N.B.

- 1) **Q.No. 1 and Q.No. 5** are **COMPULSORY**. Out of remaining attempt **ANY TWO** questions from Section – I and **ANY TWO** questions from Section – II.
- 2) Figures to the **RIGHT** indicate **FULL** marks.
- 3) Answer to both the sections should be written in **SEPARATE** answerbook.

SECTION – I

- Q.1 A)** Attempt **any seven** objective questions (MCQs / fill in the blanks) of the following. **(07)**
- i) A drug has dose of 15 mg per Kg body weight t.i.d. for 3 days. For a kid of 20 Kg body weight which of the following composition and pack is most suitable.
 - a) 150 mg / 5 ml, 30 ml
 - b) 125 mg / 5 ml, 60 ml
 - c) 300 mg / 5 ml, 45 ml
 - d) 300 mg / 5 ml, 30 ml
 - ii) An ideal requirement of a drug with usual dose of one tablet three times a day to formulate its sustained release tablet are _____.
 - a) Short biological half life
 - b) Conventional dose should be 500 mg or more
 - c) Narrow therapeutic window
 - d) a & c
 - iii) One of the following is used as disintegrating agent in tablets -
 - a) Talc
 - b) Maize starch
 - c) Gum acacia
 - d) Magnesium stearate
 - iv) Topically applied Keratolytic drug should reach at following site for drug action.
 - a) Skin surface
 - b) Within skin layer
 - c) Systemic circulation
 - d) Any of these
 - v) A liquid dosage form called _____ is hydroalcoholic solution of a drug for oral administration.
 - vi) Topically applied soft paraffin minimizes dryness of skin due to its _____ effect.
 - vii) Sodium chloride solution of _____ % w/v is isotonic with body fluids.
 - viii) Disintegrating time for film coated tablet in 0.1 N HCl may be _____ minutes.
- Q.1 B)** Discuss approaches to select flavours in Pharmaceutical formulation. **(03)**
- Q.2** Discuss various quality control tests for tablets. **(10)**
- Q.3 a)** Write reasons to formulate suppositories. **(05)**
b) Write method to inject insulin and related precautions. **(05)**
- Q.4** Answer **ANY FIVE** of the following. **(10)**
- i) Write ideal properties of capsule shell.
 - ii) Write difference between water for injection and sterile water for injection.
 - iii) Write any one approach to remove pyrogens from water.
 - iv) Write two merits of DPI over MDI.
 - v) Write two reasons to formulate transdermal patch.
 - vi) Write difference between QA and QC.

P.T.O.

SECTION – II

- Q.5 A)** Attempt **any seven** objective questions (MCQs / fill in the blanks) of the following. **(07)**
- i)** Which of the following routes of administration will show 100% bioavailability _____.
 - a) Oral
 - b) Subcutaneous
 - c) Intravenous
 - d) Rectal
 - ii)** Which drug undergo endocytosis absorption mechanism _____.
 - a) Sugar
 - b) All ionic drugs
 - c) Oral polio vaccine
 - d) thiouracil
 - iii)** A drug having Low solubility and High Permeability is classified as _____.
 - a) BCS Class I
 - b) BCS Class II
 - c) BCS Class III
 - d) BCS Class IV
 - iv)** Following is the primary drug excretion fluid _____.
 - a) Saliva
 - b) Urine
 - c) Sweat
 - d) Fecal matter
 - v)** Unionized form of drug shows _____ absorption and ionized drug shows _____ absorption.
 - vi)** _____ molecular size of drug pass from extracellular fluid to intracellular fluid.
 - vii)** Name of enzyme involved in the liver metabolism _____.
 - viii)** The pH of Blood plasma, extra cellular fluid and CSF is _____.
- Q.5 B)** Describe plasma drug concentration-time profile **(03)**
- Q.6** Define Bioavailability and explain in detail factors affecting it. **(10)**
- Q.7 a)** Explain Phase I and Phase II metabolism of drugs. **(05)**
- b)** Explain in details about various drug absorption mechanisms. **(05)**
- Q.8** Answer **ANY FIVE** of the following. **(10)**
- i)** What is mean by pre-systemic metabolism?
 - ii)** What are physicochemical properties affects the drug absorption?
 - iii)** Why metabolic acidosis patients should not take basic drugs. Give example.
 - iv)** What will happen if a lipophilic drug that is absorbed into systemic circulation is not Metabolised?
 - v)** Define drug distribution in body and what are the steps in distribution of drugs?
 - vi)** Differentiate between generic and branded medicine.

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