



# BRAINWARE UNIVERSITY

Term End Examination 2022

Programme – B.Pharm-2019

Course Name – Novel Drug Delivery System

Course Code - BP704T

( Semester VII )

Full Marks : 75

Time : 3:0 Hours

[The figure in the margin indicates full marks. Candidates are required to give their answers in their own words as far as practicable.]

## Group-A

(Multiple Choice Type Question)

1 x 20=20

1. Choose the correct alternative from the following :

- (i) Biodegradable polymer is:
- a) PLGA  
b) Ethyl Cellulose  
c) Polydimethyl siloxane  
d) All
- (ii) Which of the following is used to increase the density
- a) Potassium permanganate.  
b) Boric Acid.  
c) Iron Powder.  
d) Glucose Powder.
- (iii) Conventional hydrogels usually have porous size of  $10^4\mu\text{m}$  hence require much time to reach equilibrium while their modified version Super porous Hydrogels' ' reach equilibrium faster has an average pore size of
- a)  $>100^4\mu\text{m}$   
b)  $> 100\text{nm}$   
c)  $10^4\mu\text{m}$   
d)  $10\text{nm}$
- (iv) Identify the approach not useful to increase gastric retention time for GRDDS.
- a) High density Systems  
b) Floating Systems.  
c) Swelling Systems.  
d) Compressing Systems.
- (v) Tensile strength of ideal polymer is
- a) High  
b) Low  
c) Medium  
d) None
- (vi) Protein stability in the formulation is a major issue with Pulmonary Drug Delivery System, it can be addressed by adding Surfactants which act by,
- a) Reducing protein aggregation.  
b) Forms protective complexes  
c) Decreases their absorption  
d) Prevent coagulation
- (vii) The curve of Controlled drug delivery remains at:
- a) Toxic level  
b) Therapeutic level  
c) Subtherapeutic level  
d) All levels
- (viii) An advantage of Novel Drug Delivery Systems is
- a) It causes fluctuation of blood levels  
b) It cannot be target specific  
c) It increases toxicity of the drug  
d) It reduces side effects of the drug
- (ix) Bile salts like Sodium deoxycholate, sodium glycocholate are used in Nasal drug

delivery systems as a

- a) Propellant
  - b) Bioadhesives
  - c) Permeation enhancers
  - d) Antiallergics
- (x) Solubility & pKa include in the following factor:
- a) Patient related factor
  - b) Biological Factor
  - c) Physico-Chemical factor
  - d) None
- (xi) Which of the following equation follows Zero Order Release:
- a)  $dM_t/dt = k(M_0 - M_t)$
  - b)  $dM_t/dt = k$
  - c)  $dM_t/dt = k t^{1/2}$
  - d) All
- (xii) Which of the following characteristics is suitable for selection of a candidate for TDDS?
- a) Large Dose.
  - b) Larger molecular Size.
  - c) Higher first pass effect.
  - d) Metabolism in Skin.
- (xiii)  $t_{1/2} = 0.693$  is the value of half life of the following order of drug release:
- a) Zero order
  - b) Pseudo zero order
  - c) First order
  - d) None
- (xiv) Which of the following drugs cannot be given as transdermal administration?
- a) Drugs with very short half-lives
  - b) Drugs with narrow therapeutic index
  - c) Easy removal and termination
  - d) Drugs against peptic ulcer
- (xv) Which of the following characteristics is suitable for transdermal drug?
- a) Large drug dose
  - b) Large molecular size
  - c) Drugs with narrow therapeutic indices
  - d) Drugs which are metabolized in the skin
- (xvi) From which of the following mechanisms most of the drugs get absorbed via skin.
- a) Active transport
  - b) Passive Transport
  - c) Facilitated transport
  - d) Osmosis
- (xvii) In Noyes-Whitney equation  $[dC/dt = KD A(CS - C)]$ , CS is the:
- a) The concentration of drug in bulk of the solution
  - b) Saturation solubility of drug
  - c) The concentration of drug in the blood
  - d) None
- (xviii) From which of the following mechanisms most of the drugs get absorbed via skin.
- a) Active transport
  - b) Passive Transport
  - c) Facilitated transport
  - d) Osmosis
- (xix) Polymer is defined as:
- a) Very small molecule
  - b) Very large molecule
  - c) Medium molecule
  - d) None of these
- (xx) Nylon is:
- a) Natural polymer
  - b) Synthetic polymer
  - c) Semisynthetic polymer
  - d) None

### Group-B

(Short Answer Type Questions)

5 x 7=35

2. Write the difference between dissolution and diffusion (5)
- OR**
- Write the applications of liposomes (5)
3. Write the advantages and disadvantages of liposomal drug delivery systems (5)
- OR**
- Write the advantages and disadvantages of niosomal drug delivery systems (5)
4. Describe the ion exchange resins methods used in controlled drug delivery formulations (5)
- OR**
- Write the properties of ideal targeted drug delivery systems (5)
5. Write the mechanism of respiratory deposition (5)
- OR**
- Write about the formulation of pressurized metered dose inhalers (5)
6. Write the disadvantages of drug targeting (5)

**OR**

- Write the applications of nanoparticulate drug delivery systems (5)  
7. What are the advantages and disadvantages of controlled drug delivery systems (5)

**OR**

- Explain the need for gastro-retention for therapeutic agents (5)  
8. Discuss the advantages and disadvantages of transdermal drug delivery system (5)

**OR**

- Explain the applications of gastroretentive drug delivery system (5)

**Group-C**

(Long Answer Type Questions)

10 x 2=20

9. Explain the principle involved in the design of controlled drug delivery systems (10)

**OR**

- Write about the properties influencing drug targeting (10)  
10. Write about the evaluation parameters of gastroretentive drug delivery system (10)

**OR**

- Explain the concept and approaches for the Controlled release formulations (10)

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