#### Final Year B.Pharm.(Sem VIII) 2020-21

## BPH\_E\_811\_T-NovelDrug Delivery Systems

### **Practice Question Bank**

### 1. A non-ionic surfactant based multilamellar or unilamellar vesicular structure

- A. Microspheres
- B. Liposome
- C. Niosome
- D. Nanoparticle

#### 2. This particulate system is also known as "bodies of water".

- A. Aquasome
- B. Liposome
- C. Niosome
- D. Dendrimer

#### 3. Which of the following is a non- erodible insert?

- A. Ocusert
- B. Collagen shield
- C. NODS
- D. SODI

## 4. An ocular device that has the shape of a flag

- A. Ocusert
- B. Lacrisert
- C. NODS
- D. SODI

## 5. Which of the following does not constitute an appendageal route?

A. Sweat glands

- B. Hair follicle
- C. Sebaceous gland
- D. Stratum corneum

#### 6. A Polymer used for colonic systems is

- A. carboxymethyl cellulose
- B. cellulose acetate phthalate
- C. gelatin
- D. acacia

## 7. Drug release from osmotic drug delivery systems depends on

- A. osmotic pressure
- B. ionic strength
- C. osmotic pressure & ionic strength
- D. osmotic pressure & environment in git

## 8. \_\_\_\_\_ is a dispersed matrix system

- A. nanospheres
- B. nanoparticles
- C. nanocapsules
- D. nanopolymers

#### 9. Chitosan is a \_\_\_\_\_ mucoadhesive polymer

- A. cationic
- B. anionic
- C. synthetic
- D. non-ionic

# 10. \_\_\_\_\_ is a drug unsuitable for GRDDS

- A. ciprofloxacin
- B. diazepam
- C. furosemide
- D. aspirin

#### 11. A microcapsule has\_\_\_\_\_

- A. Drug dispersed in matrix
- B. Dug core surrounded by distinct wall
- C. Drug adsorbed on the surface
- D. Drug distributed in polymeric matrix

#### 12. 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

#### 13. Stealth liposomes

- A. have short half-life
- B. are taken up by macrophages
- C. have very large size
- D. are sterically stabilized

#### 14. An example of a polymer incorporated into dendrimers is

- A. propylene glycol
- B. polyethyleneimine
- C. polyurethane
- D. styrene copolymers

#### 15. Hydrogen bonds in mucoadhesion are formed by

- A. dipole moment
- B. non polar groups
- C. dispersion forces
- D. electronegative atoms

# **16.** Ideal glass transition temperature for a pressure sensitive adhesive used in transdermal system should be

- A. 20° C to 40° C
- B. 2° C to 4° C
- C. 20° C to 40° C
- D. 2° C to 4° C

#### 17. Ocusert is an example of

- A. Feedback regulated system
- B. Activation modulated system

#### C. Bio -responsive system

D. Membrane permeation system

### 18. \_\_\_\_\_ is an advanced method of determining size of nano particles

- A. Atomic force microscopy
- B. Ultrasound scattering
- C. Compound microscopy
- D. Molecular microscopy

#### 19. Chimeric peptides have

- A. chylomicrons
- B. polymeric micelles
- C. peptidomimetic antibodies
- D. polymeric nanoparticles

#### 20. \_\_\_\_\_ is an example of a synthetic biodegradable polymer

- A. acrolein
- B. polyethylene glycol
- C. LDPE
- D. polystyrene

## 21. Carbopols are:

- a. Synthetic vinyl polymers with ionizable carbonyl group
- b. Polyoxythylene ethers with carboxy groups
- c. Mineral waxes with hydrocarbon content ranging from C35 to C55
- d. Polyoxyethylene derivatives of plyoxypropylene

## 22. Which amongst the following are the smallest liposomes?

- A Large unilamellar vesicles
- **B** Oligolamellar vesicles
- C Multilamellar vesicles
- D Multivesicular vesicles

# 23. Which of the following is used as chemical cross-linking agent in preparation of nanoparticles?

- A Glutaraldehyde
- B 2,2, di-methyl propane
- C Lactides and glycolides
- d Poly (acryl) starch

# 24. What type of protein binding characteristics of a drug are desirable to be formulated into an ocular system?

a. Low

B Medium

C High

D It has no bearing

25. A positive temperature-sensitive hydrogel has ------ critical solution temperature

- a. Upper
- b. Lower
- c. Hybrid
- d. Mixed

26. The stratum corneum consists of -----layers of keratinized cells

A 10 to 25 B 0 to 10 C 25 to 50 D Above 50

27. Peel adhesion is tested by measuring the force required to pull a single coated tape, applied to a substrate at a ......° angle

a.180 b.360 c 45 d 90

28. Which of the following is the Noyes – Whitney equation?

a. 
$$\frac{dC}{dt} = -k(c_r - c)$$
  
b. 
$$\frac{dC}{dt} = \frac{DAk_{o_{fw}}(c_s - c_b)}{Vh}$$
  
c. 
$$\frac{M_0^{1/3} - M^{1/3} = Kt}{M_0} = k\sqrt{t}$$
  
d. 
$$\frac{M_t}{M_0} = k\sqrt{t}$$

**29.** Which among the following can be used as a hydrophobic matrix to formulate SRDDS?

- a. Ethyl cellulose
- b. Hydroxyporpyl methylcellulose
- c. Hydroxypropylcellulose
- d. Sodium carboxymethylcellulose

**30.** Which amongst this is a physicochemical factor of the drug that should be considered while formulating a controlled drug delivery system?

- a. Diffusivity
- b. Half life
- c. Side effects
- d. Absorption

# **31.** Based on their half-lives, which drug would you select to make a sustained release tablet?

a.Metformin (6 hr)
b.Heroin (2 – 6 min)
c.Cocaine (50 mins)
d Amlodipine (20 hrs)

# 32. Which of the following is an effective barrier for drug?

- a. Tight junctions
- b. Pinocytes
- c. Glucose transporters
- d. Protein carriers

# **33.** To prevent the loss of drug that has migrated into the adhesive layer during storage, this is used

- a. Release liner
- b. Rate controlling membrane
- c. Adhesive layer
- d. Backing membrane

#### 34. These noninvasive techniques have been used for drug delivery to brain

- a. Nanogels
- b. Bradykinin administration
- c. Onmaya reservoir
- d. Microgel

# 35. OROSCT Approach is used in

- a. Colon targeting
- b. Lymphatic targeting
- c. Brain targeting
- d. Mucoadheisve delivery

# 36. The dissolution study of colon targeted drugs is carried by

- a. Bio Dis III apparatus
- b. Beaker Method
- c. Flow through cell
- d. USP Type I AND II Apparatus

# 37. Super critical fluid technology is used to prepare:

- a. Nanoparticle
- b. Neosome
- c. Aquasomes
- d. Liposomes

**38.** These are a unique class of synthetic macromolecules having highly branched, three dimensional, nanoscale architecture with very low polydispersity index and high functionality

- a. Dendrimers
- b. Neosomes
- c. Auasomes
- d. Nanoparticles

# 39. \_\_\_\_\_ is carrier for Haemoglobin

- a. Neosome
- b. Nanoparticle
- c. Aquasomes
- d. Phytosomes

## 40. Following is the example of invasive brain targeting

- a. Osmogens
- b. Colloidal carriers
- c. Amino acid transporters
- d. Neosomes

## 41. The force required to remove an adhesion coating from test substrate is determined by

- a. Peel adhesion test
- b. Shear adhesion test

- c. Rolling ball tack test
- d. Probe tack test

# FINAL YEAR NDDS - SET 2

- 1. Hydrogen bonding capacity is related to which type of factor affecting mucoadhesion
- A) physiological
- B) polymer
- C) environment
- D) physicochemical
- 2. What type of process does the liposomes undergoes?
- A) Oxidation
- B) Acetylation
- C) Reduction
- D) Isomerization
- 3. Find out the odd type of ocular inserts
- A) Lacrisert
- B) Occusert
- C) SODI
- D) Minidisc
- 4. What is extrusion?
- A) pushing the heated material through an orifice
- B) producing a hole by a punch

- C) making cup shaped parts from the sheet
- D) process of mixing the ingredient
- 5. Which from the following factor does not affect Osmotic systems
- A) Osmotic pressure gradient
- B) Delivery orifice
- C) Membrane permeability, Surface area, thickness
- D) Change in pH of environment
- 6. Which of the following drugs cannot be given as transdermal DDS
- A) Drugs with very short half life
- B) Drugs with narrow therapeutic indices
- C) Easy removal & termination
- D) Drugs against peptic ulcer
- 7. Which of the following is the example of Physical theory of mucoadhesion
- A) Wetting
- B) Electronic
- C) Adsorption
- D) Adhesion
- 8. Niosomes are prepared from which of the following
- A) Phospholipids
- B) Lecithin
- C) Spingolipid
- D) Surfactants

9. Select the physical mechanism by which in situ gelling system is formed

A) Change in pH

- B) Change in glucose level
- C) Change in electric field
- D) Change in ion concentration
- 10. What are the characteristics of matrix diffusion controlled release system?
- A) Release the drug along the entire length of GIT
- B) Drug disperse in an insoluble matrix of rigid hydrophobic material
- C) Employ waxes to control the rate dissolution
- D) Release only at specific site
- 11. Which of the following is not the advantage of Transmucosal DDS?
- A) Drugs sensitive to pH change can be administered via this route
- B) Drug having poor bioavailability through oral route can be administered via this route
- C) Various hormone, steroids, enzymes can be administered by this route
- D) Ease of administration
- 12. Ocular iontophoresis is a process which does not involve
- A) Electrical potential driving charged ions into eyes
- B) Delivers high concentration to specific sight
- C) Good bioavailability
- D) Disadvantage of epithelial on conjunctival edema
- 13. Which of the following is not a component of dendrimer?
- A) Central core
- B) Stem
- C) Interior dendritic structure

## D) Exterior surface

- 14. Which of the following is incorrect about Transdermal DDS?
- A) A stable and controlled blood level can be attained
- B) All potent drugs can be administered as TDDS
- C) Drugs with narrow therapeutic window can be administered as TDDS
- D) Self- medication is possible
- 15. Which of the following is not a disadvantage of conventional dosage form?
- A) Poor patient compliance
- B) Change in concentration may lead to under or over medication
- C) Attainment of steady state condition difficult.
- D) have high cost
- 16. Which polymers occur naturally?
- A) Starch and Nylon
- B) Starch and Cellulose
- C) Proteins and Nylon
- D) Proteins and PVC
- 17. Which of the following is a thermosetting polymer?
- A) polystyrene
- B) polyolefins
- C) nylons
- D) phenolic resins
- 18. Which of the following characteristics is suitable for transdermal drug?
- A) Large drug dose
- B) Large molecular size

- C) Drug with narrow therapeutic indices
- D) Drugs which are metabolized in the skin
- 19. Which among the following polymers have lowest solubillty?
- A) polyethylene
- B) polystyrene
- C) nylon 6
- D) epoxy resin
- 20. Which of the following is not a component of buccal patch?
- A) Polymer
- B) Active substance
- C) Flavouring agent
- D) Counter irritant
- 21. Example of hydrophobic polymer used in nanoparticles is
- A) Gelatin
- B) Alginate
- C) Acrylate
- D) Lectin
- 22. Which of the method is not used for preparation of nanoparticle?
- A) Imersion polymerization
- B) Dispersion polymerization
- C) Interfacial polymerization
- D) Emulsion polymerization

23. What are the characteristics of continuous release systems?

- A) Release the drug along the entire length of GIT
- B) Prolonged their residence in the GIT and release
- C) Release only at a specific drug
- D) Release as soon as comes in contact to the saliva

24. What is the characteristic of dissolution controlled release systems?

- A) Release the drug along the entire length of GIT
- B) Prolonged their residence in the GIT and release
- C) Release only at a specific drug

D) Very slow dissolution rate

25. The absorption of the ophthalmic drug does not depend on which of the following?

- A) Physicochemical properties of the permeating molecule
- B) Drainage of tears
- C) Output of tears

D) Size of the eyeball

26. Which of the following is not a property of Bio-adhesive microspheres?

A) Achieved by making use of adhesive properties of water soluble polymers

B) Adhesion of drug delivery device to the mucosal membrane such as buccal, ocular, rectal, nasal.

C) Exhibit a prolonged residence time at the site of application and causes intimate contact with the absorption site and produces better therapeutic action.

D) They contain radioisotope i.e. either  $\alpha$ ,  $\beta$  or  $\gamma$  emitters.

27. What are the characteristics of the reservoir or membrane devices?

- A) The drug has a large therapeutic index
- B) Drug permeation rate is high
- C) Control drug release by partitioning the drug from the oil
- D) Administration of emulsions