

12. Which of the following is a new drug delivery process which is composed of phospholipids that spontaneously form multilamellar concentric vesicles with layers of aqueous media separating the lipid layer?
 a) Prodrugs b) Liposomes c) Osmotic pumps d) Nanoparticles
13. The measure of cohesive strength of the cross linking that occurs between gelatin molecules and is proportional to the molecular weight of gelatin is called
 a) Bloom strength b) Viscosity c) Surface tension d) Partition coefficient
14. If the Carr's index of a powder is 10% then the type of powder flow is
 a) Poor b) Excellent c) Very poor d) Good
15. Inadequate spreading of the coating solution during tablet coating before drying causes
 a) Orange peel effect b) Sticking c) Blistering d) Picking
16. The ratio of void volume to the bulk volume of the packing of the powder is called as
 a) Porosity b) True density c) Granular density d) Bulk density
17. Which of the following technique is used to formulate depot formulations?
 a) Solubilization b) Parenteral suspension c) Microemulsions d) Encapsulation
18. The following incompatibility occurs in a mixture of menthol and thymol
 a) Chemical incompatibility b) Therapeutic incompatibility
 c) Physical incompatibility d) Tolerance incompatibility
19. Bloom strength is used to check the quality of
 a) Lactose b) Ampoules c) Hardness of tablets d) Gelatin
20. Mixing of semisolids is carried out using
 a) Double cone mixer b) Rotating cube mixer
 c) Planetary mixer d) Fluidized bed mixer
21. In the Drugs & Cosmetics Act and rules, the Schedule relating to clinical trials of new drug is
 a) Schedule M b) Schedule X c) Schedule J d) Schedule Y
22. A co-solvent used in the preparation of parenteral products is:
 a) Benzyl alcohol b) Methyl alcohol c) Dimethyl acetamide d) Phenol

11. Note on Osmotic pressure controlled delivery system.
12. Classify permeability enhancers.
13. Note on Niosomes.
14. List out basic components of TDDS.
15. Classify polymer.

SEC- C

Note: Answer any 05 questions. Each question carries FOUR marks

5x4=20

1. Explain about preformulation stability studies of powder.
2. Explain effect of particle size on strength of tablets.
3. Write a note on chewable tablets.
4. Write a note on applications optimization techniques.
5. Pharmaceutical factors affecting drug absorption.
6. Approaches to improve dissolution of poorly soluble drug.
7. Write a concept of loading dose.
8. Explain combination of dissolution controlled drug delivery systems.
9. Evaluation of Transdermal patches.
10. Formulation aspects of liposome.