

10. An ideal enteric polymer should dissolve or become permeable near and above pH.....
- | | |
|------|------|
| a. 4 | c. 7 |
| b. 5 | d. 8 |
11. is NOT an example of enteric film former.
- | | |
|--------------------------------|-----------------------------------|
| a. Shellac | c. Cellulose acetate trimellitate |
| b. Cellulose acetate phthalate | d. Polyvinyl phthalate |
12.solid dissolves and saturates a thin film of water on its surface.
- | | |
|---------------------|------------------|
| a. Hygroscopic | c. Deliquescent |
| b. Very hygroscopic | d. Efflorescence |
13. Surfactant blends with HLB value of is preferred for making O/W emulsions
- | | |
|-----------|------------|
| a. 1 to 3 | c. 7 to 9 |
| b. 4 to 6 | d. 8 to 16 |
14. Emulsifier withpH is not suitable for topical formulation
- | | |
|-------------|-----------|
| a. Alkaline | c. Neural |
| b. Acidic | d. Any |
15. If a drug shows polymorphism, then for suspension we require polymorphism.
- | | |
|---------------|-------------|
| a. Stable | c. Unstable |
| b. Metastable | d. Any |
16. Preparation of emulsion using hard water might cause
- | | |
|--------------------|------------------|
| a. Phase inversion | c. Sedimentation |
| b. Creaming | d. Flocculation |
17. Tag Open Cup Apparatus is used to determine.....
- | | |
|---------------------|-------------------|
| a. Flame projection | c. Vapor pressure |
| b. Flash point | d. Spray pattern |
18. In cold filling process of pharmaceutical aerosol
- | |
|---|
| a. It is applicable to aqueous product only |
| b. Active ingredients should withstand temperature as low as -40°C |
| c. Propellant and concentrate are mixed and chilled to -40°F. |
| d. The chilled product are added into a previously valve crimped container. |
19. Term 'leaching' is used to describe migration of from to drug product.
- | | |
|-----------------------|--------------------|
| a. gas, atmosphere | c. dyes, container |
| b. dyes, drug product | d. gas, container |
20. provides very good resistance to acids.
- | | |
|--------------------------|-----------------------|
| a. Acrylic multi polymer | c. Polyethylene |
| b. Nitrile polymer | d. Polyvinyl chloride |

KATHMANDU UNIVERSITY
End Semester Examination
March/April 2017

APR 07 2017

Level : B.Pharm.
Year : IV
Time : 2 hrs. 30 mins

Course : PHAR 404
Semester: I
F. M. : 55

Note: Check (✓) the number of each question of Section B, C and D you have answered in the front page of main answer book. Unnecessary writing will deduct your marks.

SECTION "B"

[5Q.×3=15 marks]

- II. Answer *ANY FIVE* questions:
1. What is anhydrous lactose? Mention its pharmaceutical application.
 2. Write a short note on "Mouth-dissolving tablet".
 3. Draw a well-labeled diagram of any one type of perforated pan based coating equipment.
 4. Write about the materials that are used for preparation of non-gelatin capsules.
 5. Write about the tests that are performed for identification of emulsion type.
 6. Briefly describe the manufacturing of pharmaceutical aerosol by cold filling method.
 7. What are feed additives? Mention its advantages and limitations.

SECTION "C"

[5Q. × 5=25 marks]

- III. Answer *ANY FIVE* questions:
8. Why polymorphism study of a drug is important during preformulation study. List out the tools that are used for polymorphism study
 9. Describe how you will perform disintegration test of uncoated conventional tablets. Discuss on possible results and acceptance or rejection criteria.
 10. List the film defects that you might face in film coating. Write about the source and suitable approaches for remedy of the defects.
 11. Write about the various physical, chemical and biological tests that are performed for evaluation of pressurized packages.
 12. Write about the various factors that affects penetration rate of drug through human skin.
 13. Why vortex formation is not desired during mixing of liquids? Give your approach for preventing it?
 14. Suggest a suitable plastic resin that can be used for packaging of 'Dry Syrup'. Mention strength and weakness of the resin.

SECTION "D"

[2 Q. × 7.5=15 marks]

- IV. Answer *ANY TWO* questions:
15. What pharmaceutical benefits do granulation of powder gives? Write about different methods and related granulators used for preparation of granules. Discuss on various aspects that you should consider while selecting granulation methods.
 16. Briefly describe the operation steps of tablet compression machine. Write about the process variables related to the machine and formulation that must be properly controlled in order to achieve minimum weight variation of compressed tablets.
 17. Write about the physical properties that you seek in a pharmaceutical suspension. Give your reasons. Write down the formulation ingredients that are required for preparation of pharmaceutical suspension. Discuss on the function of each ingredients.

