

PHB Education

**Government Exam and D. Pharm Exit Exam Preparation
Questions Bank**

Subject: *Pharmaceutics*

Chapter 12 : *Liquid Dosage Forms*

Section A: Solutions (1–20)

1. A solution is a _____ system.
 - a) Homogeneous
 - b) Heterogeneous
 - c) Colloidal
 - d) Coarse→ **a**
2. The solvent used most commonly in pharmaceutical solutions is:
 - a) Water
 - b) Alcohol
 - c) Ether
 - d) Acetone→ **a**
3. The solute in a solution is present in a _____ amount.
 - a) Lesser
 - b) Greater
 - c) Equal
 - d) Variable→ **a**
4. Oral solutions are administered by:
 - a) Mouth
 - b) Injection
 - c) Rectum
 - d) Skin→ **a**
5. Solutions can be classified based on:
 - a) Route of administration
 - b) Solvent used
 - c) Composition
 - d) All of these→ **d**

6. Saturated solution contains:

- a) Maximum solute dissolved
- b) Minimum solute
- c) No solute
- d) None

→ **a**

7. Supersaturated solutions are:

- a) Unstable
- b) Stable
- c) Solid
- d) None

→ **a**

8. The process of dissolving solute in solvent is called:

- a) Solubilization
- b) Diffusion
- c) Precipitation
- d) None

→ **a**

9. Solubility increases with:

- a) Temperature (for solids)
- b) Pressure (for gases)
- c) Stirring
- d) All of these

→ **d**

10. Co-solvents are used to:

- a) Increase solubility
- b) Decrease viscosity
- c) Add color
- d) None

→ **a**

11. Common co-solvents used:

- a) Glycerin, ethanol, propylene glycol
- b) Benzene
- c) Chloroform
- d) None

→ **a**

12. Sweeteners improve:

- a) Palatability
- b) Solubility
- c) Stability
- d) None

→ **a**

13. Preservatives prevent:

- a) Microbial growth
- b) Crystallization
- c) Oxidation
- d) None

→ **a**

14. The main preservative in oral solution is:

- a) Sodium benzoate
- b) Formalin
- c) Phenol
- d) None

→ **a**

15. Filtration of solutions removes:

- a) Undissolved particles
- b) Microbes
- c) Air bubbles
- d) None

→ **a**

16. Clarification refers to:

- a) Removal of suspended impurities
- b) Color enhancement
- c) Flavoring
- d) None

→ **a**

17. Common evaluation test for solutions:

- a) Clarity test
- b) Viscosity test
- c) pH test
- d) All of these

→ **d**

18. Solutions are stored in:

- a) Well-closed containers
- b) Open bottles
- c) Paper packs
- d) None

→ **a**

19. Oral solutions must be:

- a) Palatable and stable
- b) Sterile
- c) Injectable
- d) None

→ **a**

20. Solutions for external use are labeled:

- a) "For External Use Only"
- b) "Sterile"
- c) "For Oral Use"
- d) None

→ **a**

Section B: Elixirs (21–35)

21. Elixirs are:

- a) Sweetened hydro-alcoholic solutions
- b) Pure alcohol
- c) Syrupy liquids
- d) Suspensions

→ **a**

22. The alcohol content of elixirs ranges between:

- a) 4–40%
- b) 60–90%
- c) 1–5%
- d) None

→ **a**

23. Elixirs are more stable than:

- a) Syrups
- b) Emulsions

c) Suspensions

d) None

→ **a**

24. The main solvent in elixirs is:

a) Alcohol-water mixture

b) Ether

c) Oil

d) None

→ **a**

25. Elixirs are prepared by:

a) Simple dissolution

b) Filtration

c) Mixing with flavor and color

d) All of these

→ **d**

26. Alcoholic content improves:

a) Solubility of poorly soluble drugs

b) Color

c) Flavor only

d) None

→ **a**

27. Flavored elixirs are known as:

a) Aromatic elixirs

b) Medicinal elixirs

c) Both a & b

d) None

→ **a**

28. Elixirs are clear due to:

a) Homogeneous nature

b) Suspended solids

c) Emulsion

d) None

→ **a**

29. Elixirs should be stored in:

a) Tightly closed amber-colored bottles

b) Open jars

c) Plastic pouches

d) None

→ **a**

30. Elixirs are evaluated for:

a) Clarity and alcohol content

b) Color only

c) Smell

d) None

→ **a**

31. Alcohol acts as:

a) Solvent and preservative

b) Colorant

c) Flavoring agent

d) None

→ **a**

32. Elixirs are more palatable than:

a) Bitter solutions

b) Suspensions

c) Emulsions

d) None

→ **a**

33. Non-medicated elixirs are used as:

a) Vehicles

b) Drugs

c) Coloring agents

d) None

→ **a**

34. Example of medicated elixir:

a) Phenobarbital elixir

b) Simple syrup

c) Orange oil

d) None

→ **a**

35. The main disadvantage of elixir:

a) Alcohol content unsuitable for children

b) Low solubility

c) Unstable

d) None

→ **a**

Section C: Syrups (36–50)

36. Syrups are:

a) Concentrated sugar solutions

b) Alcoholic

c) Suspensions

d) None

→ **a**

37. Sucrose concentration in syrup is:

a) 60–70% w/v

b) 20%

c) 90%

d) None

→ **a**

38. Syrups are used as:

a) Vehicles

b) Sweetening agents

c) Both a & b

d) None

→ **c**

39. Syrups prevent microbial growth due to:

a) High osmotic pressure

b) Preservatives

c) Alcohol

d) None

→ **a**

40. Simple syrup contains:

a) Only sugar and water

b) Drug

c) Alcohol

d) None

→ **a**

41. Medicated syrup contains:

- a) Drug + sweet vehicle
- b) Alcohol
- c) Only color
- d) None

→ **a**

42. Syrups are prepared by:

- a) Solution with heat or agitation
- b) Emulsification
- c) Suspension
- d) None

→ **a**

43. Excessive heating of syrup may cause:

- a) Inversion of sugar
- b) Crystallization
- c) Separation
- d) None

→ **a**

44. Flavored syrups mask:

- a) Bitter taste of drug
- b) Solubility
- c) Stability
- d) None

→ **a**

45. Syrups are stored in:

- a) Tightly closed glass bottles
- b) Plastic jars
- c) Open vessels
- d) None

→ **a**

46. Syrup spoilage may occur due to:

- a) Fermentation
- b) Evaporation
- c) Sublimation
- d) None

→ **a**

47. Syrups are evaluated for:
- a) Viscosity and microbial stability
 - b) Color
 - c) Weight
 - d) None
- **a**

48. Artificial sweeteners used:
- a) Saccharin, aspartame
 - b) Alcohol
 - c) Citric acid
 - d) None
- **a**

49. Preservatives used in syrup:
- a) Benzoic acid
 - b) Glycerin
 - c) Propylene glycol
 - d) All of these
- **d**

50. Syrup clarity test is done to ensure:
- a) No suspended particles
 - b) No color
 - c) Odorless
 - d) None
- **a**

Section D: Suspensions (51–70)

51. A suspension is a _____ system.
- a) Heterogeneous
 - b) Homogeneous
 - c) Clear
 - d) None
- **a**

52. Dispersed phase in suspension is:
- a) Insoluble solid particles
 - b) Solvent

c) Liquid

d) None

→ **a**

53. Dispersion medium is usually:

a) Water

b) Alcohol

c) Ether

d) None

→ **a**

54. Suspensions are used when:

a) Drug is insoluble

b) Drug is soluble

c) Volatile

d) None

→ **a**

55. Suspending agents prevent:

a) Sedimentation

b) Crystallization

c) Evaporation

d) None

→ **a**

56. Example of suspending agent:

a) CMC, tragacanth, xanthan gum

b) Alcohol

c) Ether

d) None

→ **a**

57. Wetting agents reduce:

a) Surface tension

b) Density

c) Viscosity

d) None

→ **a**

58. Deflocculated suspensions are:

a) Uniform and slow to settle

b) Quickly settling

c) Clear

d) None

→ **a**

59. Flocculated suspensions are easy to:

a) Redisperse

b) Separate

c) Filter

d) None

→ **a**

60. Sedimentation volume measures:

a) Stability

b) Clarity

c) Density

d) None

→ **a**

61. Suspensions are prepared by:

a) Wet gum or dry gum method

b) Melting

c) Heating

d) None

→ **a**

62. Suspensions are stored in:

a) Wide-mouth bottles

b) Metal tins

c) Plastic bags

d) None

→ **a**

63. Label instruction for suspension:

a) "Shake well before use"

b) "Keep away from sunlight"

c) "For external use only"

d) None

→ **a**

64. Evaluation test:

a) Redispersibility

b) Sedimentation

c) Viscosity

d) All of these

→ **d**

65. Flocculation improves:

a) Physical stability

b) Color

c) Odor

d) None

→ **a**

66. Viscosity influences:

a) Sedimentation rate

b) Solubility

c) Evaporation

d) None

→ **a**

67. Antimicrobial agents are added to prevent:

a) Microbial contamination

b) Sedimentation

c) Crystallization

d) None

→ **a**

68. Suspensions must have:

a) Uniform distribution of solids

b) Layer separation

c) High density

d) None

→ **a**

69. Example of oral suspension:

a) Paracetamol suspension

b) Glycerin

c) Syrup

d) None

→ **a**

70. Parenteral suspensions must be:

a) Sterile

b) Non-sterile

c) Viscous

d) None

→ **a**

Section E: Emulsions (71–85)

71. An emulsion is a:

- a) Heterogeneous system of two immiscible liquids
- b) Solid mixture
- c) Gas mixture
- d) None

→ **a**

72. The dispersed phase is:

- a) Internal phase
- b) External phase
- c) Solvent
- d) None

→ **a**

73. Emulsions are stabilized by:

- a) Emulsifying agents
- b) Preservatives
- c) Colorants
- d) None

→ **a**

74. Example of emulsifying agents:

- a) Tween, Span, acacia
- b) Alcohol
- c) Oil
- d) None

→ **a**

75. Types of emulsions:

- a) O/W and W/O
- b) Solid/Solid
- c) Gas/Liquid
- d) None

→ **a**

76. O/W emulsion means:

- a) Oil dispersed in water
- b) Water in oil
- c) Both
- d) None

→ **a**

77. Primary emulsion is prepared by:

- a) Dry gum or wet gum method
- b) Fusion
- c) Crystallization
- d) None

→ **a**

78. Dry gum method ratio (oil:water:gum) is:

- a) 4:2:1
- b) 1:2:3
- c) 2:1:1
- d) None

→ **a**

79. Emulsions are evaluated for:

- a) Type, stability, droplet size
- b) Color
- c) Odor
- d) None

→ **a**

80. Phase separation in emulsions is known as:

- a) Cracking
- b) Creaming
- c) Sedimentation
- d) None

→ **a**

81. Creaming is:

- a) Reversible
- b) Irreversible
- c) Permanent
- d) None

→ **a**

82. Coalescence leads to:

- a) Breaking of emulsion
- b) Stability
- c) Thickening
- d) None

→ **a**

83. Emulsions should be stored in:

- a) Well-closed, cool containers
- b) Open jars
- c) Metal tins
- d) None

→ **a**

84. Label instruction for emulsion:

- a) "Shake well before use"
- b) "Refrigerate"
- c) "Keep away from light"
- d) None

→ **a**

85. Example:

- a) Cod liver oil emulsion
- b) Paracetamol
- c) Glycerin
- d) None

→ **a**

Section F: Dry Powder for Reconstitution (86–100)

86. Dry powder for reconstitution is:

- a) Sterile powder to be mixed with solvent before use
- b) Tablet
- c) Suspension
- d) None

→ **a**

87. This dosage form is useful for:

- a) Unstable drugs in solution form
- b) Stable drugs

c) Creams

d) None

→ **a**

88. Common example:

a) Amoxicillin dry syrup

b) Paracetamol tablet

c) Glycerin

d) None

→ **a**

89. Reconstitution involves adding:

a) Specified volume of water

b) Alcohol

c) Oil

d) None

→ **a**

90. After reconstitution, the product is usually:

a) Suspension

b) Solution

c) Cream

d) None

→ **a**

91. Shelf life after reconstitution:

a) 7–14 days (commonly)

b) 1 month

c) 1 day

d) None

→ **a**

92. Dry powders are stored in:

a) Airtight glass bottles

b) Open jars

c) Plastic pouches

d) None

→ **a**

93. Storage temperature:

a) Cool and dry place

b) Freezer

c) Hot shelf

d) None

→ **a**

94. Evaluation includes:

a) Reconstitution time

b) Appearance after mixing

c) Sedimentation and uniformity

d) All of these

→ **d**

95. Preservatives in dry powder:

a) Methylparaben, propylparaben

b) Alcohol

c) Oil

d) None

→ **a**

96. Buffers maintain:

a) pH stability

b) Color

c) Odor

d) None

→ **a**

97. Label instruction:

a) "Add water to mark and shake well"

b) "For external use only"

c) "Keep frozen"

d) None

→ **a**

98. Flavors are added to improve:

a) Palatability

b) Solubility

c) Viscosity

d) None

→ **a**

99. Before use, the dry powder must be:

a) Reconstituted with prescribed volume of water

b) Swallowed dry

c) Mixed with milk

d) None

→ **a**

100. The main advantage of dry powder formulation

a) Better stability during storage

b) Easier manufacturing

c) Lower viscosity

d) None

→ **a**



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