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1. Glycerin suppositories containing 92% glycerin are solidified by addition of

(a) White wax
(b) Stearic acid
(c) Sodium stearate
(d) PEG 4000





1. Glycerin suppositories containing 92% glycerin are solidified by addition of

(a) White wax
(b) Stearic acid
(c) Sodium stearate
(d) PEG 4000







Soap glycerin Suppository

In glycero-gelatin base, the gelatin is replaced with either curd soap or sodium stearate which makes the base sufficiently hard to prepare good quality of suppositories. Soap also helps in the evacuation action of glycerin.







2. What is the weight of rectal suppository
(a) 3-5 gm
(b) 4 gm
(c) 2 gm
(d) 6 gm





2. What is the weight of rectal suppository
(a) 3-5 gm
(b) 4 gm
(c) 2 gm
(d) 6 gm





Rectal Suppository

These are meant for introduction into the rectum for their systemic effect. These are generally made from theobroma oil and are available in various sizes to meet the needs of infants, children and adults.

Rectal suppositories are usually available in weight about 1-2 g. They are either cone or torpedo shaped.







3. If the suppository base is fatty, what is the range of saponification value
(a) 100 m 120
(b) 200 to 245
(c) 100 to 150
(d) 200 to 300





3. If the suppository base is fatty, what is the range of saponification value
(a) 100 m 120
(b) 200 to 245
(c) 100 to 150
(d) 200 to 300





IDEAL PROPERTIES

- Should be completely non-toxic & non-irritant.
- Should be compatible with a broad variety of drugs.
- Should be non-sensitizing.
- Should have wetting & emulsifying properties.
- Should be stable on the storage i.e. does not change color, odor or drug release pattern.
- Should have acid value below 0.2.
- Should have iodine value less than 7.
- The water no. is high i.e high percentage of water can be incorporated in it.



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4. An estimate of the water absorption capacity of **100 gms of material as water number is a parameter** for evaluation of (a) Hydrogels (b) Absorption bases (c) Gelling agents ARMAC INDIA (d) Wetting agents



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4. An estimate of the water absorption capacity of **100 gms of material as water number is a parameter** for evaluation of (a) Hydrogels (b) Absorption bases (c) Gelling agents ARMAC INDIA (d) Wetting agents



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An estimate of the water absorption capacity of 100 gms of material as water number is a parameter for evaluation of Absorption bases.







5. The synthetic fat bases consisting of a mixture of tri di and mono glycerides saturated fatty acid are known as (a) Witepsol (b) Massa estarinum (c) Massuppol HARMACY INDIA (d) Wecobee



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5. The synthetic fat bases consisting of a mixture of tri di and mono glycerides saturated fatty acid are known as (a) Witepsol (b) Massa estarinum (c) Massuppol HARMACY INDIA (d) Wecobee





Massa estarinum

It is a mixture of mono, di- and triglycerides of saturated fatty acids having the formula $C_{11}H_{23}COOH$ to $C_{17}H_{35}COOH$. This is also known as adeps solidus.







6. Liquid oral preparations of which one or two large doses of about 50ml are given in a single dose container is known as

(a) Tinctures(b) Drops(c) Linctuses(d) Draughts





6. Liquid oral preparations of which one or two large doses of about 50ml are given in a single dose container is known as

(a) Tinctures
(b) Drops
(c) Linctuses
(d) Draughts





Droughts

Droughts are oral liquid preparation meant to take as a single dose. A single dose of mixture is usually known as draught.



Drops are liquid oral preparation of potent drugs or vitamin which are given in to original form without dilution.







Linctuses

They are viscous, sweet, liquid oral preparations that are usually prescribed for the relief of cold. They consist of simple solutions or admixture containing a high amount of syrup and sometimes, glycerine which in addition to give sweet taste to the preparation have a demulcent action on the mucous membranes of the throat.





7. Boric acid is freely soluble in

(a) Water(b) Alcohol(c) Glycerin(d) Chloroform







7. Boric acid is freely soluble in

(a) Water
(b) Alcohol
(c) Glycerin
(d) Chloroform







Boric Acid Solubility

Boric acid is freely soluble in water and glycerol, and sparingly soluble in pyridine. It is also slightly soluble in acetone and ethanol, but practically insoluble in diethyl ether.





8. This preparation is intended to meant for introduction into one of the body cavities

(a) Douches(b) Draught(c) Gargles(d) Throat paints







8. This preparation is intended to meant for introduction into one of the body cavities

(a) Douches
(b) Draught
(c) Gargles
(d) Throat paints







Douches

Douches are aqueous solution directed against a part or into a body cavity for cleansing and antiseptic agent.

Throat paint

Throat Paints are solutions or dispersions of one or more active ingredients intended for application to the mucosa of the throat or mouth. Used for pharyngitis or tonsillitis. Iodine throat paint is designed to kill germs.



Gargles are a type of pharmaceutical liquid dosage form used for local treatment of the throat and oral cavity.





9. What percentage of Benzoic acid is used as preservative in liquid preparations

(a) 0.01% to 0.02%
(b) 0.001% to 100.01%
(c) 0.0001% to 0.001%
(d) 0.1% to 1.01%







9. What percentage of Benzoic acid is used as preservative in liquid preparations

(a) 0.01% to 0.02%
(b) 0.001% to 100.01%
(c) 0.0001% to 0.001%
(d) 0.1% to 1.01%



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Preservatives

Phenyl mercuric nitrate - 0.001% Phenyl mercuric acetate - 0.002% Methyl paraben - 0.01 to 0.18% Propyl paraben - 0.005 to 0.035% Thiomersal - 0.001 to 0.02% Benzyl alcohol - 0.5 to 10.0% Phenol - 0.065 to 0.5% Chlorobutanol - 0.25 to 0.50% Benzoic acid – 0.01-0.02%





10. The label "Shake well before use" is indicated on the mixture containing

(a) Soluble medicament(b) Potent medicament(c) Diffusible medicament(d) Miscible liquid







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(a) Soluble medicament
(b) Potent medicament
(c) Diffusible medicament
(d) Miscible liquid





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STORAGE REQUIREMENTS & LABELLING

- Shake well before use.
- Do not freeze.
- Protect from direct light (for light sensitive drugs).
- In case of dry suspensions powder the specified amount of vehicle to be mixed may indicated clearly on label.



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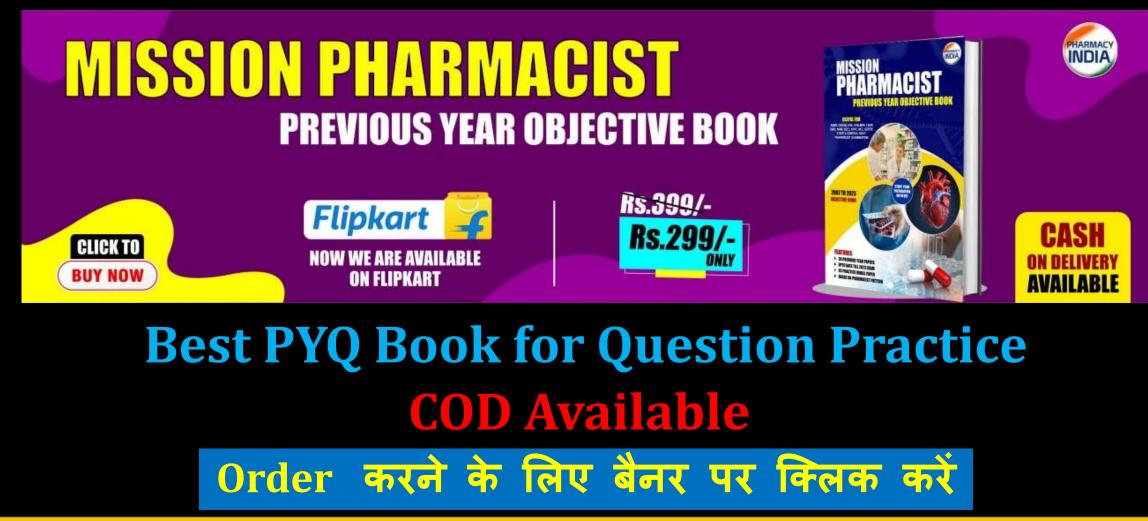


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11. Which one of the following is used as an antimicrobial preservative (a) Propylene glycol (b) Procaine hydrochloride (c) Propyl hydroxyl benzoate (d) Sodium hydrogen carbonate





11. Which one of the following is used as an antimicrobial preservative (a) Propylene glycol (b) Procaine hydrochloride (c) Propyl hydroxyl benzoate (d) Sodium hydrogen carbonate







Antimicrobial preservative

- Propyl hydroxyl benzoate
- Benzyl alcohols
- Chlorobutanol
- Methyl paraben
- Propyl paraben
- Phenol
- Phenyl mercuric acetate
- Phenyl mercuric nitrate







12. On commercial scale, emulsions are prepared by (a) Dialysis (b) Freezing (c) Homogenisation (d) Centrifugation





12. On commercial scale, emulsions are prepared by (a) Dialysis (b) Freezing (c) Homogenization (d) Centrifugation





On commercial scale, emulsions are prepared by

- Mechanical Stirrer
- Homogenizers
- Ultrasonifiers
- Colloidal Mills





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13. To increase the viscosity of liquid, which of the following agents are used (a) PVP (b) Benzalkonium chloride (c) Sodium Carboxy Methyl Cellulose (d) All of these





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13. To increase the viscosity of liquid, which of the following agents are used (a) PVP (b) Benzalkonium chloride (c) Sodium Carboxy Methyl Cellulose (d) All of these





Viscosity Enhancers

- Viscosity enhancers are avoided for the preparation of parenterals, owing to their high viscosity.
- Viscosity enhancers are prepared by hydrocolloids such as methyl cellulose, Sodium CMC, HPMC, or bentonite etc. produces high viscosity at low concentration.







14. Which one of these paraffin wax is used in eye ointment

(a) White soft paraffin(b) Yellow soft paraffin(c) Hard paraffin(d) None of these







14. Which one of these paraffin wax is used in eye ointment

(a) White soft paraffin
(b) Yellow soft paraffin
(c) Hard paraffin
(d) None of these







- Eye ointments containing yellow soft paraffins are used to relieve eye dryness and irritation.
- They moisten, soothe and lubricate the surface of your eye, making it feel more comfortable.







15. Which is an example of absorption base (a) Macrogol (b) Coconut oil (c) PEG (d) Wool fat







15. Which is an example of absorption base (a) Macrogol (b) Coconut oil (c) PEG (d) Wool fat



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Ointment bases

Ointment Bases	Properties	Examples	
Oleagenous bases	Occlusive, hydrophobic,	White petrolatum (Soft	
	greasy, non-washable.	paraffin), Hard paraffin,	
		Liquid paraffin	
Absorption bases	Occlusive, Water absorbent,	Wool fat, Hydrous wool	
	greasy, anhydrous.	fat (lanolin), Wool	
		alcohol, Bees wax	
Emulsion bases			
W/O type emulsion	Occlusive, hydrous,	Lanolin, cold cream	
bases	hydrophilic, greasy, non-		
	washable.		
O/W type emulsion	Non-occlusive, can be diluted	Hydrophilic ointment	
bases	with water, non-greasy,		
	washable.		
Water soluble bases	Water-soluble, washable,	Polyethylene glycol	
	non-greasy, non-occlusive,	(Macrogals, Carbowax)	
	lipid free.		





16. W/O emulsifiers have HLB scale value range (a) 1-3 (b) 3-6 (c) 7-9 (d) 13-16





16. W/O emulsifiers have HLB scale value range (a) 1-3 (b) 3-6 (c) 7-9 (d) 13-16





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Explanation -

HLB RANGE	CATEGORY	EXAMPLE	
1-3	Antifoaming agents	Simethicone	
3-6	w/o emulsifying agents	Span 80,	
		Span 60	
7-9	Wetting agents	Span 20,	
		Span 40	
8-16	o/w emulsifying agents	Tween 20, 40, 60, 80	
13-15	Detergents	Tween 20, 40	
15-18	Solubilizing agents	SLS	





17. "Zephiran" is a
(a) Anionic
(b) Cationic
(c) Non-ionic
(d) Ampholytic





17. "Zephiran" is a
(a) Anionic
(b) Cationic
(c) Non-ionic
(d) Ampholytic





Zephiran

- Zephiran, also known as benzalkonium chloride, is a cationic detergent and ammonium compound that's used as a disinfectant and antiseptic in the medical field.
- It's also a germicide and topical antiseptic.
- Zephiran is chemically stable, has low surface tension, and is an efficient wetting agent.







18. The most important influence of temperature on emulsion stability
(a) Creaming
(b) Caking
(c) Cracking
(d) Phase inversion





18. The most important influence of temperature on emulsion stability
(a) Creaming
(b) Caking
(c) Cracking
(d) Phase inversion





Phase inversion:

- Inversion in which O/W emulsion inverts to become a W/O emulsion and vice versa.
- The reason for the phase inversion is using of wrong type of emulsifying agents.

Causes

- Addition of an electrolyte may lead to phase inversion.
- May arise due to inappropriate emulsifier.





19. Example for a non-ionic emulsifying agent
(a) Sodium lauryl sulphate
(b) Polyoxyethylene sorbitan monooleate
(c) Cetyltrimethyl ammonium bromide
(d) Dodecyl pyridinium chloride







19. Example for a non-ionic emulsifying agent
(a) Sodium lauryl sulphate
(b) Polyoxyethylene sorbitan monooleate
(c) Cetyltrimethyl ammonium bromide
(d) Dodecyl pyridinium chloride





Tween 80

- Polyoxyethylene sorbitan monooleate.
- Polysorbate 80 is a nonionic surfactant and emulsifier often used in pharmaceuticals, foods, and cosmetics. This synthetic compound is a viscous, water-soluble yellow liquid.







20. Example for a cationic emulsifying agent (a) Dioctyl sodium sulphosuccinate (b) Polyoxyethylene sorbitan mono-oleate (c) Cetyl trimethyl ammonium bromide (d) Sodium carboxymethyl cellulose







20. Example for a cationic emulsifying agent (a) Dioctyl sodium sulphosuccinate (b) Polyoxyethylene sorbitan mono-oleate (c) Cetyl trimethyl ammonium bromide (d) Sodium carboxymethyl cellulose





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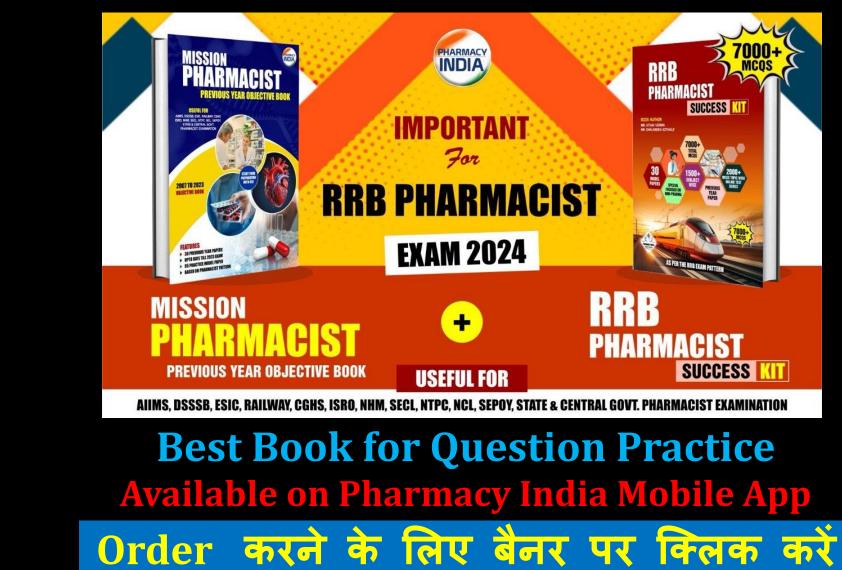
Types	Example
Anionic	Sodium lauryl sulphate(SLS), Triaton-X200,
	Sodium acetyl sulphate, Docusate
Cationic	Benzalkonium chloride, Cetyl trimethyl
	ammonium, Cetrimide
Non-ionic	Tween, Span, Myrj, Brij, Diethanolamine, PEG,
	Tween 80
Amphoteric	N- alkylamino acid, Lecithin, Betaines



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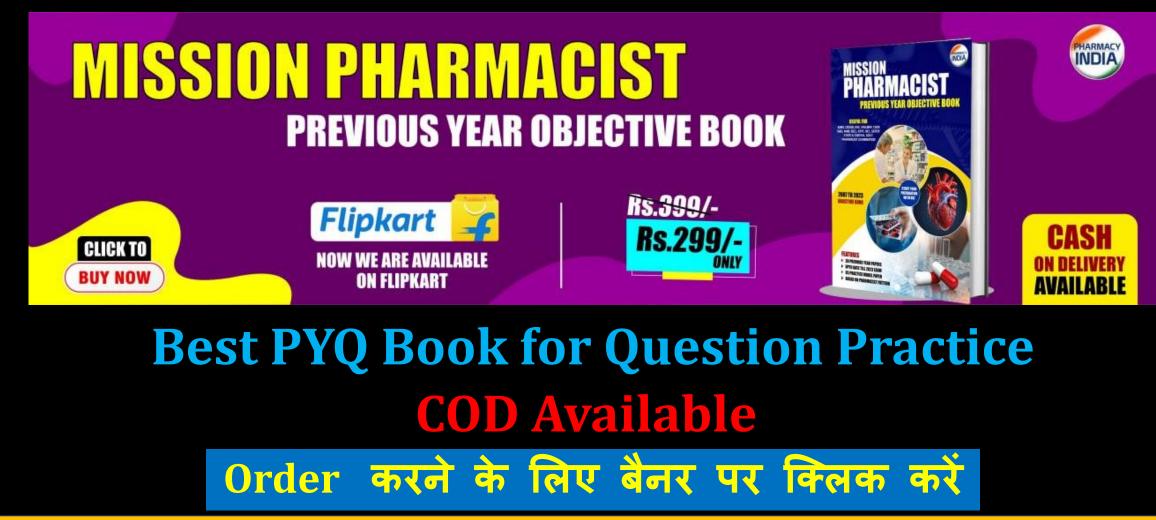




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21. In a suspension, the particle size is
(a) 5-10 micron
(b) 10-15 micron
(c) 0.5-5 micron
(d) 8-12 micron





21. In a suspension, the particle size is
(a) 5-10 micron
(b) 10-15 micron
(c) 0.5-5 micron
(d) 8-12 micron





Suspension

- Pharmaceutical suspension may be defined as a dispersion in which insoluble solids (drugs) are suspended in a liquid medium.
- Pharmaceutical suspension is a heterogeneous system consisting of two phases in which internal phase is dispersed uniformly throughout the external phase.
- ✓ Particle size ranges from 0.5 to 5 micron.







22. A suspensoid in suspension formulation is (a) An active drug (b) A solubilizing vehicle (c) A wetting agen (d) A preservative





22. A suspensoid in suspension formulation is (a) An active drug (b) A solubilizing vehicle (c) A wetting agen (d) A preservative





Suspensoid

➤A suspensoid is a substance that is dispersed throughout another substance, also known as a suspended phase.





23. Which of the following statements is **TRUE for flocculated suspensions** (a) Hard Cake is formed (b) Rate of sedimentation is high (c) They are difficult to re-disperse (d) Suspension is pleasing in appear







23. Which of the following statements is **TRUE for flocculated suspensions** (a) Hard Cake is formed (b) Rate of sedimentation is high (c) They are difficult to re-disperse (d) Suspension is pleasing in appearance





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Difference Between Flocculated and Deflocculated Suspension		IN
DEFLOCCULATED SUSPENSION	FLOCCULATED SUSPENSION	
Pleasant appearance , because of uniform dispersion of particles.	Slightly sediment and clear supernatant layer.	a
Supernatant remains cloudy	Supernatant is clear	
Particles experience repulsive force	Particles feel attractive forces	
Particles exist as separate entities	Particles forms loose aggregates	
Rate of sedimentation is slow as the size of the particles are small	Rate of sedimentation is high , as flocs are the smaller particles (higher size)	
The sediment is closely packed and form hard cake.	Sediment is loosely packed network and hard cake cannot form.	PHARN
Can not be redispersed	Easy to redisperse	CIVE - MINEL - CANAGE
In the potential energy curves , it represents the primary minimum.	In the potential energy curve , it represents the secondry minimum	
Bioavailability is relatively high	Bioavailability us comparatively less.	App from p



24. Which among the following is a thickening agent for suspension EXCEPT (a) Acacia (b) Tragacanth (c) Starch (d) Lactose





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24. Which among the following is a thickening agent for suspension **EXCEPT** (a) Acacia (b) Tragacanth (c) Starch (d) Lactose





Thickening Agents

They act as structured vehicle.
 Xanthan Gum
 Tragacanth
 Acacia
 Starch





25. The particles which are deeply bonded, settle rapidly, easily redispersible are known as

(a) Coagules
(b) Defloccules
(c) Floccules
(d) Sediments





25. The particles which are deeply bonded, settle rapidly, easily redispersible are known as

(a) Coagules
(b) Defloccules
(c) Floccules
(d) Sediments





- The particles which are deeply bonded, settle rapidly, easily redispersible are known as floccules.
- Flocs are loosely bound clusters having the open type of structures.
- Aggregates are strongly bound particles and more difficult to redisperse.







26. Indian pharmacopoeia of India has adopted type of classification system

(a) Morphological(b) Pharmacological(c) Chemical(d) Alphabetical







26. Indian pharmacopoeia of India has adopted type of classification system

(a) Morphological
(b) Pharmacological
(c) Chemical
(d) Alphabetical







Indian Pharmacopoeia

 The alphabetical system of classification was adopted in Indian Pharmacopoeia e.g., A → Aspirin, B → Bumetanide, C→ Cetirizine etc.





27. Which of the following is NOT an unofficial pharmacopoeia

(a) British Pharmaceutical Codex
(b) Remington's Pharmaceutical Sciences
(c) Merck Index
(d) The United States Dispensary







27. Which of the following is NOT an unofficial pharmacopoeia

(a) British Pharmaceutical Codex
(b) Remington's Pharmaceutical Sciences
(c) Merck Index
(d) The United States Dispensary





History of British Pharmacopoeia

- The first list of approved drugs with information on how they should be prepared was the London Pharmacopoeia published in 1618.
- In 1907 the British Pharmacopoeia was supplemented by the **British Pharmaceutical Codex**, which gave information on drugs and other pharmaceutical substances not included in the BP, and provided standards for these.







28. Which of the following substances liberate water of crystallization

(a) Hygroscopic
(b) Efflorescent
(c) Deliquescent
(d) Eutectic mixtures





28. Which of the following substances liberate water of crystallization

(a) Hygroscopic
(b) Efflorescent
(c) Deliquescent
(d) Eutectic mixtures





- Efflorescence Substances can lose water of crystallization through a process called efflorescence, which occurs when the crystals are exposed to dry air, even for a short time.
- Deliquescent If a hydroscopic substance absorbs so much moisture that an aqueous solution is formed, the substance becomes deliquescent.







29. The solvates can exist in different crystalline forms called as

(a) Enantiotropic
(b) Monotropic
(c) Pseudopolymorphs
(d) Amorphous





29. The solvates can exist in different crystalline forms called as

(a) Enantiotropic
(b) Monotropic
(c) Pseudopolymorphs
(d) Amorphous





Pseudopolymorphism

Solvates, also known as pseudopolymorphs, can exist in different crystalline forms. Solvates are crystalline solids that form during crystallization with the help of a solvent, and incorporate solvent molecules into their crystal lattice.







30. Which of the following is a type of Oral dosage form

(a) Aerosol
(b) Nebulizer
(c) Subcutaneous administration
(d) Tablet





30. Which of the following is a type of Oral dosage form

(a) Aerosol
(b) Nebulizer
(c) Subcutaneous administration
(d) Tablet



Tablet



According to the Indian Pharmacopoeia :-Pharmaceutical tablets are solid, flat or biconvex dishes, unit dosage form, prepared by compressing a drugs or a mixture of drugs, with or without diluents meant for oral administration.



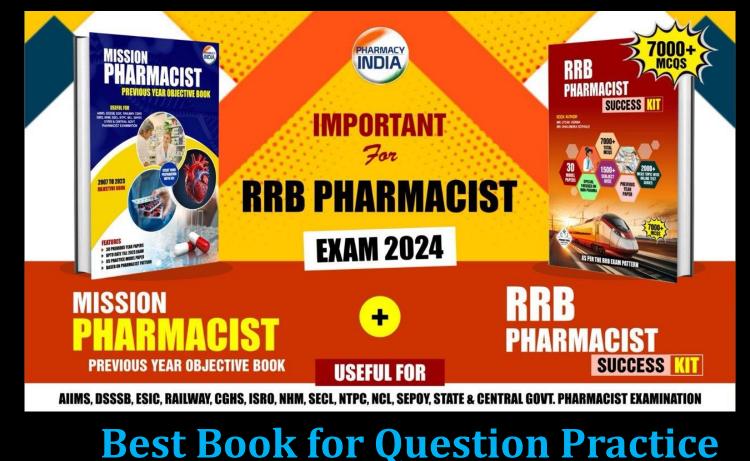


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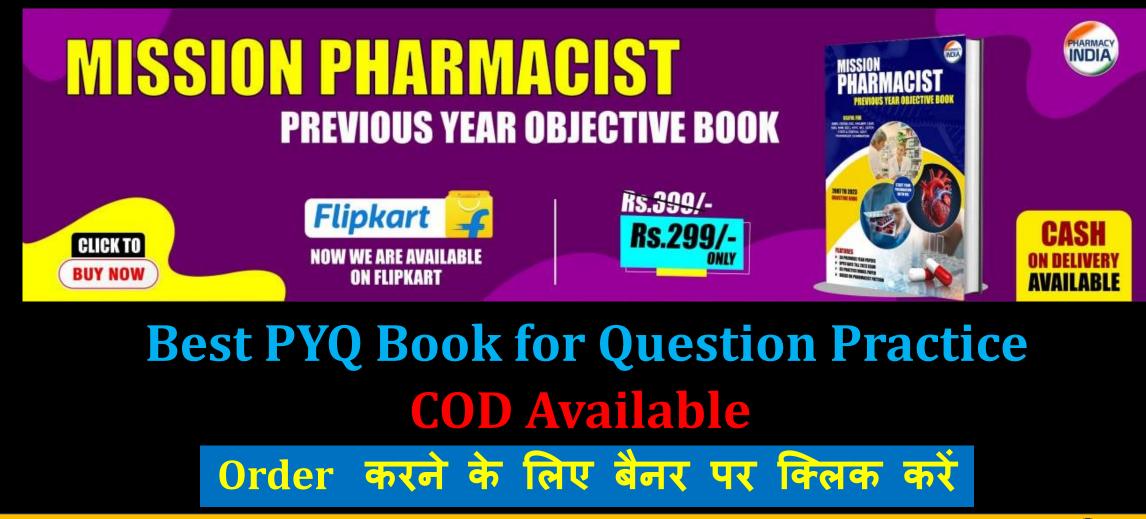


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31. The sweetening agent commonly used in chewable tablet is

(a) Sucrose
(b) Honey
(c) Mannitol
(d) Saccharin







31. The sweetening agent commonly used in chewable tablet is

(a) Sucrose
(b) Honey
(c) Mannitol
(d) Saccharin





Sweetening Agent

- ✓ For chewable tablets:
- ✓ Mannitol \rightarrow 72% as sweet as sucrose
- ✓ Saccharine (artificial): 500 time's sweeter than sucrose
- Disadvantage: Bitter after taste and carcinogenic
- ✓ Aspartame (artificial)
- Disadvantage: Lack of stability in presence of moisture.







32. Which of the following is the example of "Invert Sugar"
(a) Sucralose
(b) Lactulose
(c) Lactose
(d) Sucrose





32. Which of the following is the example of "Invert Sugar"
(a) Sucralose
(b) Lactulose
(c) Lactose
(d) Sucrose





Invert sugar

Sucrose is called an invert sugar because there is a change in the sign of rotation from dextro before hydrolysis to leavo after hydrolysis. Sucrose on hydrolysis gives equimolar mixture of D-(+) glucose and D-(-) fructose.





33. Which of the following is used as a binder (a) Talc (b) Starch (c) Cellulose (d) Kaolin





33. Which of the following is used as a binder (a) Talc (b) Starch (c) Cellulose (d) Kaolin





Binders and Adhesives

✓ These materials are added either dry or in wet form to form granules or to form cohesive compacts for directly compressed tablet.

Example

Starch, Carboxymethylcellulose sodium, Cellulose, Microcrystalline, Ethyl cellulose, HPMC, Magnesium aluminium silicate, Methylcellulose, Poly dextrose







34. Lactose is used as
(a) Diluent
(b) Glidant
(c) Lubricant
(d) Disintegrant





34. Lactose is used as
(a) Diluent
(b) Glidant
(c) Lubricant
(d) Disintegrant





Diluents

Microcrystalline cellulose	Avicel Aricel Emocel	
Starch	Sta-Rx-1500	
Sucrose (sucrose dextran ppt)	Di-Pac, Sugar tab. Nu- tab.	
Anhyd. Lactose	DCL-30	
Spray dried lactose	Fast flow Zeparox TM	
Hydrolysed starch Dextrates	Celutab, Emdex	
CaHP04	Emcompaces	
Microfine cellulose	Elcema	







35. Why sub coating is provided on the tablets (a) To avoid deterioration to microbial attack (b) To prevent the solubility in acidic media (c) To avoid stickiness (d) To increase the bulk size of the tablets





35. Why sub coating is provided on the tablets (a) To avoid deterioration to microbial attack (b) To prevent the solubility in acidic media (c) To avoid stickiness (d) To increase the bulk size of the tablets







SUB COATING

Sub coating is applied :

- To form uniform edges
- To build up the tablet size
- Sub coating increases the tablet weight from 50 to 100 percent
- Examples Gelatin, sugarcane powder, corn syrup, syrup, distilled water, Gum acacia.







36. According to Indian Pharmacopoeia the statement of sparingly soluble at 20° to 30° means approximate volume of solvent in ml per gm of solute is

(a) From 1 to 10
(b) From 30 to 100
(c) From 10 to 30
(d) From 100 to 1000





36. According to Indian Pharmacopoeia the statement of sparingly soluble at 20° to 30° means approximate volume of solvent in ml per gm of solute is

(a) From 1 to 10
(b) From 30 to 100
(c) From 10 to 30
(d) From 100 to 1000





Solubility Studies

Seneral terms used for expressing solubility:

Terms	Parts of solvent required to	
	dissolve 1 part pf solute	
Very soluble	Less than 1 part	
Freely soluble	1 to 10 parts	
Soluble	10 to 30 parts	
Sparingly soluble	30 to 100 parts	PHARMACY
Slightly soluble	100 to 1000 parts	
Very slightly soluble	1000 to 10000 parts	
Practically insoluble	More than 10000 parts	Download PHARMACY INDIA App from play store





37. Simple syrup USP contains _____ w/v sugar

(a) 85%
(b) 68%
(c) 75%
(d) 50%





37. Simple syrup USP contains _____ w/v sugar

(a) 85%
(b) 68%
(c) 75%
(d) 50%





Syrups

- Syrups are concentration aqueous preparation of sugar or sugar substances with or without flavoring agent and medical substances.
- The syrups containing medicinal substances are called "medicated syrup" and those containing aromatic or flavoured substances are known as "flavoured syrups".
- Simple syrup I.P contains 66.7%w/w sucrose in purified water (100 ml).
- Simple syrup USP contains 85%w/v sucrose in purified water (100 ml).







38. A solution of local anaesthetic contains 1:100,000 adrenaline. How much adrenaline have been added to make the solution

(a) 0.01%
(b) 0.1%
(c) 100 mcg/ml
(d) 10 mcg/ml





38. A solution of local anaesthetic contains 1:100,000 adrenaline. How much adrenaline have been added to make the solution

(a) 0.01%
(b) 0.1%
(c) 100 mcg/ml
(d) 10 mcg/ml





Ratio Interpretation:

The ratio 1:100,000 means there is 1 part adrenaline to 100,000 parts of the solution.

Amount of Adrenaline Added:

To find out how much adrenaline is added, we need to know the total volume of the solution. However, if we assume a standard context (e.g., 1 liter), then the calculation would proceed as follows:

Amount of adrenaline = $\frac{1}{100,000} \times \text{total volume of solution}$ For example, if the total volume of the solution is 1 liter (1000 milliliters) Amount of adrenaline = $\frac{1}{100,000} \times 1000 \text{ ml}$ = 0.01 mL = 10 mcg/ mL



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39. Use of Potassium Iodide in Iodine solution

(a) Increase the solubility of Iodine(b) Decrease the solubility of Iodine(c) Reduce the toxicity of Iodine(d) Improve the appearance of Iodine solution





39. Use of Potassium Iodide in Iodine solution

(a) Increase the solubility of Iodine
(b) Decrease the solubility of Iodine
(c) Reduce the toxicity of Iodine
(d) Improve the appearance of Iodine solution







Iodine Solution

Iodine is wider antiseptic agent which is effective against bacteria, fungi, yeast, protozoa and viruses. It is slightly soluble in water. Its solubility in water can be increased by addition of complex forming agent like potassium iodide.





40. The dosage form in which fluid containing the drug is retained in the rectum for the drug to act either locally or systemically

(a) Retention enema(b) Evacuant enema(c) Suppositories(d) Lotion







40. The dosage form in which fluid containing the drug is retained in the rectum for the drug to act either locally or systemically

(a) Retention enema
(b) Evacuant enema
(c) Suppositories
(d) Lotion







Enema

Retention enemas require you to "hold it" or retain the fluid for a bit so the enema has time to work inside your colon.
 Enemas are rectal injections of fluid intended to cleanse or stimulate the emptying of your bowel.



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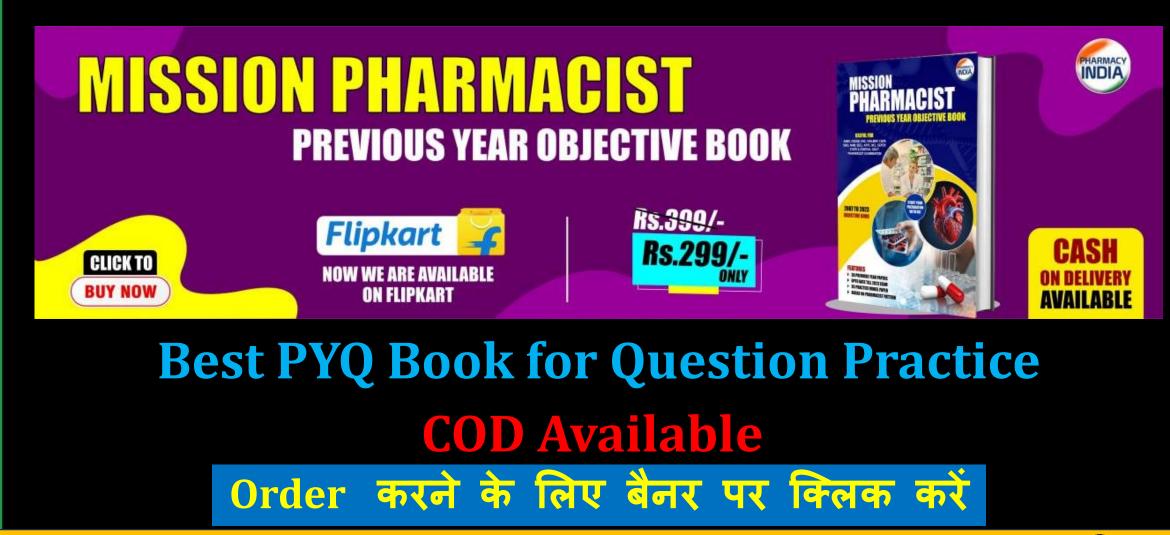


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