#### Section I

### **Pharmaceutics**

- 1. History of the Profession of Pharmacy in India
- 2. Packaging Materials
- 3. Pharmaceutical Aids and Preservatives
- 4. Unit Operations
- 5. Tablets and Capsules
- 6. Liquid Oral Preparations
- 7. Topical Preparations
- 8. Sterile Formulations and Immunological Products
- 9. Quality Control and Quality Assurance
- 10. Novel Drug Delivery System

	Syllabus (As Per ER 2020) Pharmaceutics
Chapter	Topics
1	History of the profession of pharmacy in India in relation to pharmacy education, industry, pharmacy practice, and various professional associations.  Pharmacy as a career
	<b>Pharmacopoeia:</b> Introduction to IP, BP, USP, NF and extra pharmacopoeia. Salient features of Indian pharmacopoeia
2	<b>Packaging materials:</b> Types, selection criteria, advantages and disadvantages of glass, plastic, metal, rubber as packaging materials
3	Pharmaceutical aids: Organoleptic (colouring, flavouring, and sweetening) agents
	Preservatives: Definition, types with examples and uses
4	Unit operations: Definition, objectives/applications, principles, construction, and working of: ■ Size reduction: Hammer mill and ball mill
	■ Size separation: Classification of powders according to IP, cyclone separator, sieves standards of sieves
	Mixing: Double cone blender, Turbine mixer, Triple roller mill and Silverson mixer homogenizer
	Filtration: Theory of filtration, membrane filter and sintered glass filter
	<b>Drying:</b> Working of fluidized bed dryer and process of freeze drying
	Extraction: Definition, Classification, method, and applications
5	Tablets—coated and uncoated, various modified tablets (sustained release, extended-release, fast dissolving, multi-layered, etc.)
	Capsules—hard and soft gelatine capsules
	Liquid oral preparations—solution, syrup, elixir, emulsion, suspension, dry powder for reconstitution
	Topical preparations—ointments, creams, pastes, gels, liniments and lotions, suppositories, and pessaries Nasal preparations, Ear preparations
	Powders and granules—Insufflations, dusting powders, effervescent powders, and effervescent granules Sterile formulations—Injectables, eye drops and eye ointments
	Immunological products: Sera, vaccines, toxoids, and their manufacturing methods.
6	Basic structure, Layout, sections, and activities of pharmaceutical manufacturing plants
~	Quality control and quality assurance: Definition and concepts of quality control and quality assurance,
	current good manufacturing practice (cGMP)
_	Introduction to the concept of calibration and validation
7	<b>Novel drug delivery systems:</b> Introduction, classification with examples, advantages, and challenges

#### History of the Profession of Pharmacy in India in Relation to Pharmacy Education, Industry, Pharmacy Practice and Various Professional Associations; Pharmacy as a Career; Pharmacopoeia

1.	The Greek word <i>Poeio</i> mean	ns:		C. Indian Pharmaceutical Congress Association		
	A. To cure	<b>B.</b> To make		D. Indian Pharmaceutical Association		
	C. To diagnose	<b>D.</b> To treat	8.	Earliest pharmacies were known as		
2.	J	<ul><li>B. William Martindale</li><li>D. ML Schroff</li></ul>		<ul><li>A. Apothecary shops</li><li>B. Pharmacy</li><li>C. Medicine point</li><li>D. Drugstore</li></ul>		
3.	The first British pharmaco	ppoeia was published in	9.	The architect of Indian pharmaceutical industry is		
	<b>A.</b> 1890 <b>C.</b> 1870	<b>B.</b> 1864 <b>D.</b> 1895		A. Acharya SB Manikdas B. Dr RC Subhedar C. Dr APJ Abdul Kalam		
4.	In the year 2018 which edition was published?	of Indian pharmacopoeia	10.	D. Acharya PC Ray  At least is required to practice as a		
	<ul><li>A. 2nd edition</li><li>C. 7th edition</li></ul>	<ul><li>B. 5th edition</li><li>D. 8th edition</li></ul>	100	<ul><li>pharmacist in India.</li><li>A. Degree in Pharmacy</li></ul>		
5.	The British Pharmacopoeia copoeia of the  A. Europe			<ul><li>B. Masters in Pharmacy</li><li>C. Pharm D</li><li>D. Diploma in pharmacy</li></ul>		
	<ul><li>B. United Kingdom</li><li>C. France</li><li>D. South Africa</li></ul>		11.	Pharmacy education in India as a university level programme started in 1937 at  A. Delhi University  B. University of Madras		
6.	DEC stands for  A. Drug Education Committee  B. Drug Ethical Committee			C. Banaras Hindu University D. Patna University		
	<ul><li>B. Drug Ethical Committee</li><li>C. Drug Enquiry Committee</li><li>D. None of the above</li></ul>			12. According to the reference, the commonly worshipped Hindu God of Medicine, "" is the original exponent of the Indian medicine.		
7.	Who organizes the Indian F every year?	<u> </u>		A. Ganesh B. Mahadev		
	<ul><li>A. Indian Pharmaceutical Co</li><li>B. Indian Pharmacy Graduate</li></ul>			<ul><li>C. Dhanvantari</li><li>D. Bramha</li></ul>		

8 **Pharma Success** was enacted as the nation's first 21. \_ were medical doctors as well as minimum standard of educational qualification for pharmacists. pharmacy practice. A. Charaka and Chanakya A. Drugs and Cosmetics Act 1940 (1945) B. Chanakya and Sushruta C. Charaka and Sushruta B. DPCO Act C. Pharmacy Act 1948 D. Aryabhatta and Valmiki **D.** All of the above 22. The Drugs and Cosmetics Act was passed in the 14. Pharmacy practice in India is governed by which of year the following Act? **A.** 1940 **B.** 1945 A. Pharmacy Act 1948 **C.** 1948 **D.** 1955 **B.** Drugs and Cosmetics Act 1940 (1945) 23. Who is known as the "Father of Pharmacy Education C. DPCO Act in India"? **D.** All of the above A. Acharya PC Ray 15. Key strengths of pharmaceutical industry are \_ B. Prof Mahadev Lal Schroff A. Strong manufacturing base C. Sushruta **B.** Cost competitiveness D. Charaka C. Fast growing healthcare industry 24. Pharmacy education in India at the certificate level **D.** All of the above was started in 1842 in Goa by 16. The word 'pharmacy' is derived from the Greek word **B.** Jews A. Britishers **C.** Native businessmen D. Portuguese B. Pharma A. Pharmaces was the first pharmaceutical manufac-C. Pharmacisto D. Pharmakon turing facility establishment in India. 17. The present education regulations 2020 framed replaced A. Indian Pharmaceutical Company Ltd. **B.** Bengal Chemicals and Pharmaceutical Works Ltd. A. Education Regulations 1981 C. Calcutta Chemicals **B.** Education Regulations 1991 **D.** Ranbaxy Pharmaceuticals Ltd. C. Education Regulations 1982 26. The first edition of IP was published in \_\_\_\_\_ D. Education Regulations 2014 **A.** 1948 **B.** 1955 is the federation of five national **C.** 1940 **D.** 1960 pharmaceutical associations as its constituents. 27. Extra pharmacopoeia is also termed A. Indian Pharmaceutical Congress A. Martindale: The Complete Drug Reference B. Indian Pharmaceutical Congress Association **B.** Pharmacopoeia codex C. Indian Pharmacy Graduates Association **C.** National formulary D. Indian Pharmaceutical Association **D.** International pharmacopoeia is the premier professional association of is pharmacy as the minimum required pharmacists in India. qualification to enter into the pharmacy profession A. Indian Pharmaceutical Congress in India. B. Indian Pharmacy Graduates Association B. Postgraduation A. Degree C. Indian Pharmaceutical Association D. Pharm D C. Diploma **D.** Indian Pharmaceutical Congress Association 29. Dioscorides wrote \_ has the highest number of US FDA approved drug manufacturing units (outside USA) A. Merck Index today. **B.** Indian Pharmacopoeia

C. Martindale

D. De Materia Medica

**A.** India

C. China

B. Russia

D. America

30.	which edition of Indian Ph in 2018?	narmacopoeia was published	41.	A. Chopra committee is	s known as
	<b>A.</b> 3rd	<b>B.</b> 7th		<b>B.</b> Bhor committee	
	<b>C.</b> 6th	<b>D.</b> 8th		C. Import of drug bill	
31	Who is the Father of Med	licine?		<b>D.</b> Indian Pharmacopoeia Co	ommittee
01.	A. Ebers	<b>B.</b> Hippocrates	42	The Pharmacy Education	in India is regulated by
	C. Egyptian	<b>D.</b> Pontus	72.	Pharmacy Council of India	
22	57.1			A. The Pharmacy Act 1948	
32.	_	e form are known as		B. Drugs And Cosmetics Act	t 1940
	A. Excipient	B. Source of drug		C. DPCO Act	
	C. Dosage forms	<b>D.</b> API		<b>D.</b> The Industry Act	
33.	-	nbination of drug and diffe-	43.	The first 2 years of profession	onal course "Chemist and
	rent kinds of non-drug con			Druggist Diploma" was star	
		<b>B.</b> Non-additives		<b>A.</b> Banaras Hindu University	
	C. New chemical entity	<b>D.</b> All of the above		B. Bengal Chemical and Pha	rmaceuticals
34.	The 7th edition of IP was	published in .		C. Madras Medical College	
	<b>A.</b> 2007	<b>B.</b> 1996		<b>D.</b> Indian Pharmaceutical As	sociation
	C. 2014	<b>D.</b> 2012	44	The second edition of Ind	ian nharmaconoeia was
35	USP first edition was pub	lished in		published in	
55.	A. Latin	B. English		<b>A.</b> 1985	<b>B.</b> 2007
	C. English and Latin	•		C. 1955	<b>D.</b> 1966
	C .		45	All the following are official l	hooks <i>excent</i>
36.	The meaning of Pharmak	con means	13.	A. Merk Index	воокз ехсері
	A. Powder			B. The Drug Bill	
	<b>B.</b> Medicine or drug			C. National Formulary	
	C. Emulsion			<b>D.</b> Indian Pharmaceutical Co	odex
	<b>D.</b> None of the above		4.0		·
37.	The fourth edition of IP w	as published in	46.	The Indian Government ap Committee' under the Chair	
	<b>A.</b> 1965	<b>B.</b> 1975		·	i manship or
	<b>C.</b> 1996	<b>D.</b> 1985		A. Dr BN Ghosh	B. Dr Mukharji
38.	The content of pharmaco	poeia includes		C. William Martindale	D. Lt Col RN Chopra
	A. Monograph of drug/sub		47	The long form of AIOCD is	
	<b>B.</b> Standard test		47.	A. All India Organization of	
	C. Description, formulae			<b>B.</b> All India Organization of	
	<b>D.</b> All of the above			C. All India Organization of	
20	The first chamist show in I	udia waa ananad hy		<b>D.</b> None of the above	Circinist and Bistiloator
39.	The first chemist shop in In in 1811 in Kolkata.	nuia was openeu by	40		
	A. Scotch Bathgate		48.	The current Education Reg	
	B. Acharya PC Ray			PCI are known as A. ER 1981	• B. ER 1991
	C. Cal RN Chopra			C. ER 2014	<b>D.</b> ER 2020
	<b>D.</b> Haroon Zaffer				
40		n at 11 (DCT)	49.	The 'Import of Drugs B	
40.	The first Pharmacy Council of India (PCI) was constituted by Central Government in			of drugs.	ssembly to control import
	A. 1947	B. 1949		A. 1940	<b>B.</b> 1937
	<b>C.</b> 1948	<b>D.</b> 1950		<b>C.</b> 1935	<b>D.</b> 1931
	C. 1740	<b>U.</b> 1730		C. 1733	<b>ט.</b> 1731

1 B	2 C	3 B	4 D	5 B	6 C	7 C	8 A	9 D	10 D
11 C	12 C	13 C	14 B	15 D	16 D	17 B	18 B	19 C	20 A
21 C	22 A	23 B	24 D	25 B	26 B	27 A	28 C	29 D	30 D
31 B	32 C	33 A	34 C	35 C	36 B	37 C	38 D	39 A	40 B
41 A	42 A	43 C	44 D	45 B	46 D	47 A	48 D	49 B	50 C

56 C

57 A

58 A

59 A

60 B

55 B

51 D

52 C

53 A

54 D

**Answer Key** 

### Packaging Materials

1.	Which one of the following rubber closure?  A. Extractive test C. Permeability test	<ul><li>g is not an official test for</li><li>B. Pyrogen test</li><li>D. Compatibility test</li></ul>	9.	Which of the following pack protect the drug against th A. Plastic containers B. Amber coloured glass co	e lig	ht?		
2.	Following are the materials			C. Both A and B D. None of the above				
	A. Glass	B. Plastic	10.	Type-I glass is also known	as	•		
	C. Rubber	<b>D.</b> Metal		A. Borosilicate glass				
3.	PVC means  A. Polyvinyl chloride C. Polyvinyl carbide			<ul><li>B. Regular soda-lime glass</li><li>C. Treated soda-lime glass</li><li>D. None of the above</li></ul>				
4.		, ,	11.	Which of the following meth	ods a	are used in the produc-		
7.	packaging.	i iai oi choice ioi a nexibie		tion of glass?				
	A. Metal	<b>B.</b> Glass		A. Blowing	В.	Drawing		
	C. Cellulose	<b>D.</b> Cellophane		C. Pressing and casting	D.	All of the above		
5.	is a device by can be closed and opened.	12.	To protect the contents of a sunlight by UV rays, which					
	A. Aerosol	B. Closure		A. Amber coloured glass	_			
	C. Container	<b>D.</b> None of the above				None of the above		
6.	Which of the following mat	erials are used in pharma-	13.	Soda ash is also known as		•		
	ceutical packaging?	F		<b>A.</b> Pure silica	В.	Treated soda-lime		
	A. Glass	B. Plastic		C. Limestone	D.	Sodium carbonate		
	C. Metal	<b>D.</b> All of the above	14.	Plastic containers are gene	rally	made from which of		
7.	Major disadvantages of gla	ss as a nackaging material		the following materials?				
	are			A. Polyethylene	В.	Polypropylene		
	<b>A.</b> Fragility	B. Weight		C. Polystyrene		All of the above		
	C. Both A and B	<b>D.</b> None of the above	15	The neelesse that is divect	ler in	contact with formu		
8.	Composition of glass is		15.	The package that is direct lation is called	-	Contact with formu-		
	A. Sand	B. Soda ash		A. Secondary package		Tertiary package		
	C. Lime stone and cullet	<b>D.</b> All of the above		C. Primary package	D.	All of the above		

16.		vantages of are	f plastic con	tainer	s over	glass	con-	23.	Plasti A. Au			s are s	teriliz	zed usi	ing		•
		y formation							<b>B.</b> Bl			nt					
	-	istance to br							C. Et								
		edom of desi	_						<b>D.</b> Al	•							
		of the above	_											_			
17.		stands for _						24.	The purchase that h		_	-		•		_	avities called
	A. Low	density pol	lyethylene														
	B. Ligh	nt density po	olyethylene						A. St		_						
	C. Low	density po	lypropylene									packa	ge				
	<b>D.</b> Non	e of the abo	ove						<b>C.</b> Bl	-	_						
18.	Herme	tic containe	ers are also l	known	as		<u>_</u> .				•	ckage					
	A. Air-	tight contain	ners					25.	A ligh	t-resis	stant c	contain	ier pro	ovides	prote	ction a	gainst
	<b>B.</b> Mul	ti-dose cont	ainers												_		
	C. Sing	gle dose con	tainers						<b>A.</b> M					B. Li	_		
	<b>D.</b> Non	e of the abo	ove						<b>C.</b> Bo	oth A a	and B			<b>D.</b> No	one of	the ab	ove
19.	Thermo	oplastics ar	·e					26.	Glass	conta	iners	are		ir	ı natu	re.	
	A. Doe	s not softene	ed upon expo	sure to	high t	emper	ature		A. Fr	_				<b>B.</b> Li	~		
	B. Hard	d and brittle							<b>C.</b> Bo	oth A a	and B			<b>D.</b> No	one of	the ab	ove
	C. Softened upon exposure to high temperature						27.	Whic	h is th	ie mos	t resis	tant t	vpe of	glass	?		
	<b>D.</b> None of the above							<b>A.</b> Ty					<b>В.</b> Ту	_			
20.	HDPE	stands for _		_•					C. Ty					<b>D.</b> Ty	-		
	<ul><li>A. High density polyethylene</li><li>B. High density polyethene</li><li>C. High density polypropylene</li></ul>						28.	the m	<b>anufa</b> ctivate	cture d carb		ber c	d as vu losure		zing ag	gent in	
	<b>D.</b> Non	e of the abo	ove						<b>B.</b> Su	_							
21.	Type-ll	glasses are		_•					C. Ta								
	A. Trea	ited soda-lin	ne glass						D. Sto	earic a	acid						
	<b>B.</b> NP §	glass						29.			glass	is a gl	ass of	highe	st pha	rmace	eutical
	C. Bore	osilicate gla	SS						grade	;							
	<b>D.</b> Gen	eral soda-lii	me glass						-	_				<b>B.</b> Ty	pe II		
22.	Rubbei	rs are used	l as materi	al for	const	ructio	on of		<b>C.</b> Ty	pe III				<b>D.</b> Ty	pe IV		
		•						30.	Soda-	lime ş	glass i	s also	know	n as			
	A. Con	tainer	E	3. Clos	sure				<b>A.</b> Ha	ard gla	ass			B. So	ft glas	SS	
	C. Both	n A and B	Ι	). Non	e of th	ne abo	ve		C. Po	tash g	glass			<b>D.</b> Le	ad gla	ISS	
							•	17									
							Answe	er Key									
	1 B	2 C	3 A	4	D	5	В	6	D	7	С	8	D	9	В	10	Α
	1 D	12 C	13 C	14		15		16		17		18		19		20	
	1 A	22 B	23 C	24		25		26		27		28		29		30	

#### Pharmaceutical Aids and Preservatives

1.	Identify the artificial sweet	tener among the following:	8.	Preservatives improve _	•		
	A. Sucrose	<b>B.</b> Dextrose		A. Shelf life of the produ	ct		
	C. Sucralose	D. Mannitol		<b>B.</b> Product stability			
2	To in average viscosity of lie	wid which of the following		C. Appearance of the pro	duct		
2.	To increase viscosity of liq reagent is used?	uid which of the following		<b>D.</b> All of the above			
	A. PVP		9.	Lakes are example for _			
	<b>B.</b> Sodium carboxymethyl	cellulose		A. Colouring agent			
	C. Methyl cellulose			<b>B.</b> Flavouring agent			
	<b>D.</b> All of the above			C. Sweetening agent			
3.	Which of the following is r	not used as a diluent?		<b>D.</b> None of the above			
	<ul><li>A. Lactose</li><li>B. Microcrystalline cellulos</li><li>C. Calcium carbonate</li></ul>	se	10.	sweetening agent?	an example of a non-nutritive		
				A. Sucralose	B. Sucrose		
	<b>D.</b> Polyvinyl Pyrrolidine			C. Mannitol	<b>D.</b> None of the above		
4.	Which among the following natural colors?  A. Annatto C. Saffron	B. Carotene D. Naphthalene blue	11.	<ul><li>EDTA is an example of _</li><li>A. Chelating agent</li><li>B. Antifungal preservative</li><li>C. Anti-oxidant</li></ul>			
5.	Chelating agents act by	mechanism.		<b>D.</b> None of the above			
	<b>A.</b> Complexation	<ul><li>B. Hydration</li><li>D. None of the above</li></ul>	12.	A flavouring agent is use the product.	ed to enhance of		
6.	Diluents are used to	•		A. Smell			
	A. Add cohesiveness to the	powder blend		<b>B.</b> Appearance			
	<b>B.</b> Make up the required bu	lk of the tablet	C. Palatability				
	C. Both A and B			<b>D.</b> None of the above			
	<b>D.</b> None of the above		13	Which of the following a	re widely used and excellent		
7.	Antioxidants act by inhibiti	ngmechanism.	10.	preservatives?	ire widely used and excellent		
	A. Complexation			A. Quaternary ammonium	n compounds		
	<b>B.</b> Hydration			<b>B.</b> Mercurials			
	C. Oxidation			C. Chelating agents			
	<b>D.</b> None of the above			D. Chloroform			

14.	<b>A.</b> Acid <b>B.</b> Neu <b>C.</b> Mer	·	22.	<ul> <li>Which of the following flavour is not responsible sour taste?</li> <li>A. Citrus flavour</li> <li>B. Liquorice</li> <li>C. Raspberry</li> <li>D. Mint spice</li> </ul>							ole for					
	<b>D.</b> Qua	ternary amn	nonium com	pounds			23.			ne follo	owing	agent	s are	e used a	s flavo	uring
15.	<ul><li>5. Which of the following is an example of synthetic colouring agent?</li><li>A. Amaranth</li><li>B. Curcumin</li></ul>					hetic								Chloroform None of the above		ove
	C. Coc			None of t		ve	24.				owing			ar colou		
16.	Which	of the follow	wing sugar l	ıas bitter ta	aste?			A. Ar			· · · · · · · · · · · · · · · ·			Caramel		
	A. Glu	cose	В	3. Sucrose				C. Tit	aniun	n dioxi	de		<b>D.</b> (	Cochine	al	
	<b>C.</b> Saccharine <b>D.</b> None of the above					ve	25.	The c	olour	ing ag	gents	are pe	ermi	tted un	der tl	ne Act
17.	7. Which of the following is a synthetic sweetener?					•					1049		р г	NAD A a	+ 1054	
	A. Glu			Sucrose				<b>A.</b> Ph <b>C.</b> NI		-				OMR Ac D and C		
	C. Sorbitol D. Aspartame						26	C. NDPS Act 1985 D. D and C Act 1946. Which of the following is used as a colouring ag								
18.	8. At which concentration, phenol acts as preservative?						20.	A. Le		ic ion	ywing			a colou Sorbitol	i ing a	gent.
	<b>A.</b> 0.2- <b>C.</b> 0.05			3. 0.5–0.8 3. None				C. Clove				<b>D.</b> Caramel				
10			lavouring a		Lindus	try is	27.	. Pharmaceutical aids are also l						wn as _		
1).	A. Mer	-	iavouring a	Sent in 1000	illuus	ti y 15		<b>A.</b> Ac	•					Adjuncts		
	B. Chloroform							C. Ex	•					All of th		e
	<ul><li>C. Monosodium glutamate</li><li>D. None of the above</li><li>Ingredients used to enhance the elegance of pharma-</li></ul>					28.										
												Colourin				
20.						C. Sw						All of the		e		
			ons are calle	d	<b>-•</b>		29.	A. Buffers  B. Antioxidants								
		minatives rmaceutical	aide					<b>A.</b> Bu <b>C.</b> Di						Antıoxıd /ehicles		
	C. Astı		aius				•				_					_
		e of the abo	ve				30.	<b>A.</b> Be					r sp	rays an	d aero	sols.
21.	Cochin	eal is a	8	gent.				<b>B.</b> Tri								
	A. Flav			G. Colouring	g			C. Ca								
		etening		. Thickening	_			D. Pro								
			_			Answe	er Key	<u>'</u>								
	1 A	2 D	3 D	4 D	5	Α	6	В	7	С	8	D		9 A	10	Α
	1 A	12 C	13 A	14 D	15			С	17	-	18		-	9 C	20	
2	1 B	22 D	23 C	24 A	25	I)	26	D	27	l D	28	В	」っつり	9 B	30	в

# Unit Operations Size Reduction, Size Separation, Mixing, Filtration, Drying, Extraction

1.		t for extraction of alkaloids,	8.	The process of separa		-
	glycosides, volatile oils an			suspension or slurry is		
	A. Water	<b>B.</b> Alcohol		<b>A.</b> Filtration	B	. Sieving
	C. Kerosene	<b>D.</b> Light petroleum		C. Distillation	D.	. Drying
2.	•	esses of extraction except	9.	Which one of the follow	ing does r	ot influence filtration?
	·			A. Temperature	В.	Density
		B. Digestion		C. Viscosity	D.	. pH
	C. Drying	<b>D.</b> Maceration	10	Matafiltania alsa knav	WM 00	
3.	is used for o	drying of biological product	10.	Metafilter is also know		
	like plasma, serum, vacci			A. Filter press		
	-	<b>B.</b> Spray dryer		C. Edge filter	D.	. Sintered filter
	C. Rotary dryer		11.	Filter aids are added t	o the liqu	ıid
				<b>A.</b> To increase the porc	osity	
4.	Thermolabile material	can be sterilised by using		<b>B.</b> To increase cake per		<b>√</b>
	•			C. Both A and B		
	<b>A.</b> Filter leaf	<b>B.</b> Sintered glass filter		<b>D.</b> None of the above		
	C. Filter press	<b>D.</b> Filter paper				
_	The efficiency of ball mill is maximum at			Size reduction is also	called	•
5.	<del>-</del>	is maximum at		<b>A.</b> Comminution		
	A. Low speed			<b>B.</b> Continuation		
	<b>B.</b> High speed			C. Communication		
	C. Very high speed			<b>D.</b> Elutriation		
	<b>D.</b> 2/3 speed (critical speed	d)	12	*****	6 4	
6	Most simple and most free	quently used method for size	13.	Which of the followin		is not anecting on the
υ.	separation is			process of size reducti		Cut 1:
	A. Sieve Shaker	<b>-</b> •		A. Hardness		Stickiness
	B. Cyclone separator			C. Viscosity	D.	. Abrasiveness
	C. Air separator		14.	In the ball mill maxis	mum siz	e reduction occurs at
	<b>D.</b> Elutriation			speed.		
	D. Elutriation			A. Low		
7.	Which type of mixtures is	s easily formed?		B. High		
	A. Positive	<b>B.</b> Negative		C. Optimum/critical		
	C. Neutral	<b>D.</b> Ampholytic		<b>D.</b> None of the above		
	- : - : - : - : - : - : - : - : - : - :	·				

ieves to  Nibrate  Oscillate	B. None D. Spin	26.	<ul><li>A. Density in shape</li><li>B. Shape and surface are</li></ul>	
A. Convection B. Diffusion		27.	<b>D.</b> Surface texture and si	is the type of size separation?
Which of the following is N eduction? A. Cutting C. Burning	OT a method used for size  B. Impact D. Shear	28.	A powder of which the	entire particles pass through more than 40.0% through a
Impact     Impact and attrition	<ul><li><b>B.</b> Attrition</li><li><b>D.</b> None of the above</li></ul>		<ul><li>A. Coarse powder</li><li>B. Moderately coarse po</li><li>C. Fine powder</li><li>D. Very fine powder</li></ul>	owder
Attrition is a mechanism   A. Ball mill  C. Roller mill	<ul><li>B. Plate mill</li><li>D. All of the above</li></ul>	29.	The filtration which in intermicellar liquid fr membrane under reduc	avolves the separation of the com solid by semipermeable ced pressure is
<ul><li>Increase size</li><li>Decrease absorption</li><li>Decrease solubility</li></ul>		30.	C. Cross flow	f asbestos is  B. Carbon
For ease in reduction of size,  A. Hard, brittle  C. Hard tough	material must be  B. Soft tough D. Soft brittle	31.	Surface type cartridges candles made up of por	s are usually in the shape of celain, and  B. Cotton
Principle of hammer mill is  A. Cutting C. Attrition	B. Crushing D. Impact	32.	·	on the principle of
A. Compression  B. Impact	the ball mill is		<ul><li>B. Freezing</li><li>C. Sublimation</li><li>D. Melting</li></ul>	шоп
O. Attrition and impact  of the space of the balls in ball mill.		33.		is obtained after the complerocess is called  B. Marc D. Extract
C. 80%	<b>D.</b> 20%	34.	Continuous hot percola A. Soxhlation	tion is called  B. Maceration
A. Mass B. Volume in litres per unit C. Weight		35.	<ul><li>C. Infusion</li><li>Size classification is also</li><li>A. Size separation</li></ul>	<b>D.</b> Decoction
	ieves to A. Vibrate C. Oscillate Che mechanisms of mixing A. Convection B. Diffusion C. Convection, shear, diffus D. Both A and B  Which of the following is Needuction? A. Cutting C. Burning A ball mill works on the process. A. Impact C. Impact and attrition Attrition is a mechanism A. Ball mill C. Roller mill C. Roller mill C. Roller mill C. Decrease absorption C. Decrease size and increase Cor ease in reduction of size, A. Hard, brittle C. Hard tough Crinciple of hammer mill is C. Cutting C. Attrition Che mechanism involved in C. Compression C. Attrition Che mechanism involved in C. Attrition Che mechanism involved in C. Attrition and impact  Of the space of the space of the balls in ball mill. C. 100% C. 80% Che rate of filtration is exp C. Mass C. Volume in litres per unit	A. Vibrate B. None C. Oscillate D. Spin  The mechanisms of mixing are A. Convection B. Diffusion C. Convection, shear, diffusion D. Both A and B  Which of the following is NOT a method used for size eduction? A. Cutting B. Impact C. Burning D. Shear A ball mill works on the principle of A. Impact B. Attrition C. Impact and attrition D. None of the above attrition is a mechanism of size reduction found in A. Ball mill B. Plate mill C. Roller mill D. All of the above dize reduction helps in A. Increase size B. Decrease absorption C. Decrease solubility D. Decrease size and increase stability  For ease in reduction of size, material must be A. Hard, brittle B. Soft tough C. Hard tough D. Soft brittle  Principle of hammer mill is A. Cutting B. Crushing C. Attrition D. Impact  The mechanism involved in the ball mill is A. Compression B. Impact C. Attrition D. Attrition and impact of the space of the cylinder is occupied by the balls in ball mill. A. 100% B. 30 to 50% C. 80% D. 20%  The rate of filtration is expressed as A. Mass B. Volume in litres per unit time C. Weight	The week to  A. Vibrate B. None C. Oscillate D. Spin  The mechanisms of mixing are  A. Convection B. Diffusion C. Convection, shear, diffusion D. Both A and B  Which of the following is NOT a method used for size eduction? A. Cutting B. Impact C. Burning D. Shear B. Attrition C. Impact B. Attrition D. None of the above Attrition is a mechanism of size reduction found D. None of the above Attrition is a mechanism of size reduction found D. All of the above D. All of the above D. Soft tough D. Decrease size and increase stability  For ease in reduction of size, material must be  A. Hard, brittle B. Soft tough D. Hard tough D. Soft brittle  Principle of hammer mill is  A. Cutting B. Crushing J. Attrition D. Impact  The mechanism involved in the ball mill is  A. Compression B. Impact D. Attrition and impact  of the space of the cylinder is occupied by the balls in ball mill.  A. 100% B. 30 to 50% D. 20%  The rate of filtration is expressed as  A. Mass B. Volume in litres per unit time D. Weight	A. Density in shape A. Density in shape B. None C. Oscillate D. Spin C. Oscillate D. Spin C. Convection C. Convection C. Convection C. Convection, shear, diffusion D. Both A and B D. Both A and B Cassifying and scree C. Sieving D. All of the following C. Convection? C. Cutting D. Shear C. Sieving D. All of the above C. Fine powder D. Very fine

36. Which mechanism help in size separation by sieve			48.	48. Membrane filter works on the principle of					
	shaker?			A. Physical separation	<b>B.</b> Chemical separation				
	$\mathcal{E}$	<b>B.</b> Sedimentation		C. Shearing force	<b>D.</b> Gravitational force				
37	<ul><li>C. Brushing</li><li>Movement of particle can</li></ul>	D. Shearing forces  be enhanced during size	49.	'Kozeny-Carman equation' the theories of					
57.	separation by which one of			A. Extraction					
		<b>B.</b> Attrition		C. Filtration	•				
	C. Gravitation		<b>=</b> 0						
38.	The solvent used for extrac	_	50.	Final removal of liquid from of heat is called					
	A. Menstruum	B. Marc		A. Extraction					
	C. Expression	D. Galenical		C. Separation	<b>D.</b> Mixing				
39.	For mixing eac	ch particle of one material	51.	is also known	as freezer drying.				
	lies as nearly adjacent as po	ssible to a particle of other		A. Homogenization					
	material.			C. Clarification	<b>D.</b> Filtration				
	<ul><li>A. Positive</li><li>C. Perfect</li></ul>	_	52.	In cyclone separator the power	der is separated depending				
40.	The Pharmacopoeia has p	rescribed only the unner		on its  A. Particle size	<b>B.</b> Density				
	limit for the fir			C. Particle size and density	•				
	<b>A.</b> 2	<b>B.</b> 3		•					
	C. 5	<b>D.</b> 4	53.	Which of the following equipmixing of powder?	ipment can be used for				
41.	The preparations which are			<b>A.</b> Turbine mixer					
	method are called			C. Sigma blade mixer	<b>D.</b> All of the above				
	<ul><li>A. Biologicals</li><li>C. Galenicals</li></ul>		54.	A clear liquid passing thr	ough the filter is called				
42.	is not used for	mixing, in dispensing.		A. Filtrate	B. Slurry				
		<b>B.</b> Spatulation		C. Filter media	•				
	C. Trituration								
43.	Which type of the following		55.	The separation process in whiliquid is not more than 1%	w/v is called				
	blender?	D. Tymshlan miyan		A. Clarification					
	<ul><li>A. Double cone blender</li><li>C. Sigma blade mixer</li></ul>	<ul><li>B. Tumbler mixer</li><li>D. Paddle mixer</li></ul>		C. Centrifugation	<b>D.</b> Evaporation				
			56.	Leaching means	_•				
44.	Degree of mixing is also kn			<b>A.</b> Liquid–liquid extraction	<b>B.</b> Solid–liquid extraction				
	A. Degree of homogeneity			C. Solid-phase extraction	<b>D.</b> All of the above				
	C. Random mixing	<b>D.</b> None of the above	57.	is the process	of removal of an active				
45.	Sieve through which all the of moderately fine particles		0	constituent from crude drug in which it is soluble.					
	<b>A.</b> 44	<b>B.</b> 85		A. Filtration	<b>B.</b> Size reduction				
	C. 22	<b>D.</b> 10		C. Drying	<b>D.</b> Extraction				
46.	The principle behind cyclon	e separator is	58.	Continuous hot percolation i	s also known as				
	A. Agitation	B. Centrifugal force		<b>A.</b> Lyophilization	<b>B.</b> Soxhalation				
	C. Sieves	<b>D.</b> None of the above		C. Homogenization	<b>D.</b> None of the above				
47.	Sintered glass filter is made	e from	59.	Which of the following is sta	andard for sieve?				
	A. Limestone	<b>B.</b> Soda-lime glass		A. Sieve number	<b>B.</b> Nominal aperture size				
	C. Sulphur glass	<b>D.</b> Borosilicate glass		C. Nominal diameter of wire	<b>D.</b> All of the above				

60.	Which materials are not dryer?	used in drying in a freeze	71.	Which of the following the filtration?	eory not d	lescribe rate of		
	A. Seafood	<b>B.</b> Fruits		A. Darcy Law				
	C. Pharmaceuticals	<b>D.</b> Dyes		<b>B.</b> Poiseulle's equation				
61.	In drying process, the fina	al product is in the form of		<b>C.</b> Kozeny-Carman equation <b>D.</b> Noyes-Whitney equation				
	A. Slurry	<b>B.</b> Solution	72.	The slurry is				
	C. Solid	<b>D.</b> Solvent concentrate		A. A suspension to be filtered	d			
62.	·	ritical in drying process is	<ul><li>B. A porous membrane used to retain the solids</li><li>C. The solids which are present on the filter</li><li>D. A clear liquid passing through the filter</li></ul>					
		<b>B.</b> Pressure		D. A clear fiquid passing thi	ough the h	itei		
	C. Temperature	<b>D.</b> Volume	73.	Which of the following drye	r is knowr	ı as lyophilizer?		
63.	Advantage of fluidized be	d dryer is .		A. Fluidized bed dryer	B. Spray	y dryer		
	<b>A.</b> No attrition			C. Freeze dryer	<b>D.</b> Vacu	um dryer		
	C. Fluppy mass is formed	nously exposed to heat source	74.	means extraction		-		
	<b>D.</b> Humidity can be increased	sea		<b>A.</b> Evaporation	_			
64.	In sieving, sieves are arrai	nged in orders.		C. Galenicals	<b>D.</b> Filtra	ition		
	A. Ascending	B. Random	75.	Proper drying can prevent_		_ of the product.		
	C. Descending	<b>D.</b> Horizontal		A. Loss	B. Weig	ht		
65.	Mechanism of mixing in sil	verson mixer is .		C. Deterioration	D. Solul	oility		
	A. Convective	B. Laminar	76.	Drying involves	transfe	r operation.		
	C. Random	D. Turbulent		A. Mass	<b>B.</b> Heat			
"	A double cone mirror is us	ad fan miving		C. Mass and heat		er mass or heat		
00.	A double cone mixer is use A. Solids	B. Semi-solids						
	C. Liquids		77.	For extraction of thermolab process should be used?	ile drugs v	which extraction		
	C. Liquius	D. Suspensions			B. Mace	anati an		
<b>67.</b>	Triple roller mill works on			<ul><li>A. Percolation</li><li>C. Decoction</li></ul>	<b>D.</b> Both			
	<b>A.</b> Diffusion mixing			C. Decoction	<b>D.</b> Dom	A and D		
	<b>C.</b> Dry mixing	<b>D.</b> Shear mixing	<b>78.</b>	The solvent used for extrac	tion is kno	own as		
68.	The rate of flow of the filt	rate through the filter cake		<b>A.</b> Distillate	<b>B.</b> Extra			
	is to the thick	_		C. Marc	<b>D.</b> Mens	struum		
	<ul><li>A. Inversely proportional</li><li>B. Directly proportional</li></ul>		79.	The inert insoluble material t	nat remain	s after extraction		
	C. Remain constant			A. Distillate	<b>B.</b> Extra	ict		
	<b>D.</b> None of the above			C. Marc	<b>D.</b> Mens	struum		
69.	Which of the following is	not a filter aid?	80.	Which of the following proc	ess is used	for extraction?		
	A. Diatomaceous earth	B. Perlite		A. Infusion	B. Deco			
	C. Cellulose	<b>D.</b> Cotton		C. Digestion	<b>D.</b> All o	of the above		
70	Filter aids are added to th	e liauid	01	_	ge and the	on allowing it to		
, 0.	Filter aids are added to the liquid  A. To increase the porosity			Pouring water over the drukeep in contact with water	_	_		
	<b>B.</b> To increase the porosity			called	asumiy 10	10 111111 13		
	C. Both A and B			A. Infusion	B. Deco	oction		
	<b>D.</b> None of the above			C. Percolation	<b>D.</b> Dige			

82.	Extraction does not involve ponents	one of the following com-	92.		ation in which two or more her in such a way that each
	<b>A.</b> Active constituents			particle of material looks	similar to particles of other
	B. Solvent			material.	
	C. Vapour			<b>A.</b> Filtration	<b>B.</b> Extraction
	<b>D.</b> Crude solids			C. Mixing	<b>D.</b> Drying
83.	Which of the following is	not a mechanism of size	93.	Following is the mechanis	m of mixing
	reduction?			A. Shear mixing  B. Convective mixing	
	A. Compression			<b>B.</b> Convective mixing	
	C. Attrition	<b>D.</b> Elutriation		<ul><li>C. Diffusion mixing</li><li>D. All of the above</li></ul>	
84.	Following are the processes	s used for extraction except			
		•	94.	_	e mixer are used for mixing
	A. Maceration	B. Sublimation		of	
	C. Digestion	<b>D.</b> Percolation		A. Solids	<b>B.</b> Liquids
0.5	D	1		C. Semi-solids	<b>D.</b> Powders
<b>ช</b> 5.	Preparations like infusions are commonly known as _		95.	Double cone blender wo	rks on the mechanism of
	A. Pharmaceuticals			A. Convective mixing	
	<b>B.</b> Galenicals			<b>B.</b> Diffusion mixing	
	C. Nutraceuticals			C. Shearing mixing	
	<b>D.</b> Cosmeceuticals			<b>D.</b> All of the above	
86.	Solvent used for extraction	ı is	06		_
	A. Water	<b>B.</b> Alcohol	90.	Silverson mixer is used fo	
	C. Ether	<b>D.</b> All of the above		<ul><li>A. Homogenization</li><li>C. Extraction</li></ul>	
87	What is the disadvantage	of alcohol as solvent for		C. Extraction	<b>D.</b> Separation
07.	extraction?	of alcohol as solvent for	97.	Clarification is used for th	e removal of small amount
	A. Toxicity	<b>B.</b> Stability		of suspended solids	•
	C. Cost	<b>D.</b> Selectivity		<b>A.</b> Not more than 2%	
00	D 11 11 1 1 1 1 1	•		<b>B.</b> Not more than 0.15%	
88.	Ball mill is based on princi	ple of		C. Not more than 1.5%	
	A. Cutting			<b>D.</b> Not more than 2.5%	
	B. Compression		98.	The deposited layer of so	lid on the filter medium is
	<ul><li>C. Impact and attrition</li><li>D. Attrition</li></ul>		, , ,	called .	
	D. Authon			A. Filter cake	<b>B.</b> Filter medium
89.	Hammer mill is based on t	he principle of		C. Filtrate	<b>D.</b> Slurry
	A. Impact	<b>B.</b> Attrition	00	Which of the fellowing	•
	C. Cutting	<b>D.</b> Compression	99.	filtration?	filters are used for sterile
90.	Elutriation is a method of_	•		A. Cotton wool and filter p	aper
	<b>A.</b> Size reduction	<b>B.</b> Extraction		<b>B.</b> Cotton wool and glass v	
	C. Size separation	<b>D.</b> Filtration		C. Membrane filter and sin	_
91.	In cyclone separator	force is used to		<b>D.</b> Muslin cloth and asbest	os
	separate solid from the flui		100.	are the examp	les of filter aids.
	A. Fractional force			A. Diatomic and Kieselgul	
	B. Centrifugal force			<b>B.</b> Asbestos and cotton wo	
	C. Electromagnetic force			C. Glass wool and cotton v	vool
	<b>D.</b> Nuclear force			<b>D.</b> Muslin cloth and asbest	os

#### **Answer Key**

1 B	2 C	3 D	4 B	5 D	6 A	7 A	8 A	9 D	10 C
11 C	12 A	13 C	14 C	15 D	16 C	17 C	18 C	19 D	20 D
21 A	22 D	23 D	24 B	25 B	26 C	27 D	28 A	29 B	30 A
31 D	32 A	33 B	34 A	35 A	36 D	37 A	38 A	39 C	40 A
41 C	42 A	43 A	44 A	45 A	46 B	47 D	48 A	49 C	50 B
51 B	52 C	53 B	54 A	55 A	56 B	57 D	58 B	59 D	60 D
61 C	62 A	63 B	64 C	65 D	66 A	67 D	68 A	69 D	70 C
71 D	72 A	73 C	74 B	75 C	76 C	77 C	78 C	79 C	80 D
81 A	82 C	83 D	84 B	85 B	86 D	87 C	88 C	89 A	90 C
91 B	92 C	93 D	94 B	95 B	96 A	97 B	98 A	99 C	100 A

#### Tablets and Capsules

1.	The disintegration time li is	mit for film coated tablet	7.	Disintegration time for is	enteric coated tablet as per IP
	A. 15 minutes	<b>B.</b> 30 minutes		A. 1 hour	<b>B.</b> 2 hours
	C. 45 minutes	<b>D.</b> 60 minutes		C. 5 hours	<b>D.</b> 3 hours
2.	Which of the following is formulation?	used as glidant in tablet	8.	_	et is measured by
	A. Cellulose acetate phthala	ate		A. Thickness	
	B. Talc	ato		<ul><li>B. Weight variation</li><li>C. Friability</li></ul>	
	C. Polyvinyl chloride			<b>D.</b> None of the above	
	<b>D.</b> Carbowax			<b>D.</b> None of the above	
3.	Which type of coating is do	ne to disintegrate the tablet	9.	How many tablets are retest?	equired for content uniformity
	in intestine?			<b>A.</b> 10	<b>B.</b> 20
	A. Sugar coating			<b>C.</b> 30	<b>D.</b> 40
	<b>B.</b> Film coating		10	Th	
	C. Enteric coating		10.	at by the constant temp	ssolution media is maintained
	<b>D.</b> None of the above			A. $37 \pm 0.5$ °C	
4.	Which of the following is a	method of preparation of		<b>C.</b> $37 \pm 0.5$ °C	
	granules?	1 1		C. 37 ± 1.3 C	<b>D.</b> 3/ ± 2 C
	<ul><li><b>A.</b> Slugging</li><li><b>B.</b> Dry granulation</li></ul>		11.	Weight variation for 80–250 mg is	tablet weight ranging from
	C. Wet granulation			<b>A.</b> 10%	<b>B.</b> 7.5%
	<b>D.</b> All of the above			<b>C.</b> 5%	<b>D.</b> None of the above
5.	The partial or complete	removal of top or bottom	12.	Lactose is used as	·
	portion of tablet is known	_		A. Diluent	B. Glidant
	A. Picking	B. Capping		C. Lubricant	<b>D.</b> Disintegrant
	C. Sticking		13.	To increase bulk of the	tablet, which of the following
6.	Shellac is the material use	d in of tablets			ed to tablet formulation?
	A. Film coating			A. Glidants	
	<b>B.</b> Enteric coating			B. Lubricants	
	C. Sugar coating			C. Diluents	
	<b>D.</b> Control release shooting			<b>D.</b> Disintegrants	
				-	

14.	Which of the following ebinder in granulation?	xcipients can be used as a	24.	should be stored at temperature no exceeding 30°C.			
	A. Magnesium stearate	B. Starch mucilage		A. Tablet			
	C. Kaolin	<b>D.</b> Fuller's earth		<b>B.</b> Capsule			
15.	Unequal distribution of c	olour in tablet is known as		<ul><li>C. Sustained release ta</li><li>D. Sublingual tablet</li></ul>	ıblet		
	A. Capping	B. Picking	25.	The disintegration ti	me for soft gelat	in capsule is	
	C. Mottling	<b>D.</b> Lamination		<b>A.</b> 45 minutes	<b>B.</b> 60 min	utec	
16.	is performed to determine the ability of tablet withstand wear and tear during packing,			C. 20 minutes	<b>D.</b> 10 min		
	handling and transportat		26.	Gelatin is a hydrolyse	d product of	·	
	A. Disintegration test			A. Pectin	B. Tannin		
	<b>B.</b> Friability test			C. Alkaloid	<b>D.</b> Collage	en	
	C. Dissolution test		27.	The elasticity of the so	ft gelatin capsule s	shell is impro-	
	<b>D.</b> All of the above			ved with the addition		•	
17.	means for	mation of film surface like		A. Lubricant			
	orange peel due to rapid			<b>B.</b> Increasing water co	ontent		
	A. Wrinkling			C. Plasticizer			
	C. Sweating	D. Orange peel		<b>D.</b> None of the above			
18.	Disintegration time of effe	rvescent tablet is	28.	Rotary die process is capsules.	used for	filling into	
	<b>A.</b> 3 minutes	<b>B.</b> 5 minutes		A. Powders	<b>R</b> Liquids	•	
	C. 15 minutes	<b>D.</b> 30 minutes		C. Pellets	• .		
19.	Separation of tablet in tw	o or more distinct layers is	29.	The moisture conter	nt of a soft gelat	in capsule is	
	A. Picking	B. Capping		A. < 10%	<b>B.</b> 9–13%		
	C. Lamination	<b>D.</b> Broken		<b>C.</b> >16%	<b>D.</b> 20–26%	<b>6</b>	
20.	Appearance of uneven sp tablet is known as	oot on a film surface of the	30. Capsules normally fall into two main categories a				
	A. Splitting			A. Hard gelatin capsul	es and soft gelatin	capsules	
	C. Blistering	<b>D.</b> Flaking		B. Hard gelatin capsul	es and layered cap	sules	
21.	Disintegration time for	hard gelatin capsule is		<ul><li>C. Soft gelatin capsule</li><li>D. Compressed and lay</li></ul>		capsules	
	A. 30 minutes		31.	The largest size of caps	ule is denoted by w	hich number?	
	<b>B.</b> 50 minutes			<b>A.</b> 000	<b>B.</b> 1		
	C. 60 minutes			<b>C.</b> 2	<b>D.</b> 5		
	<b>D.</b> 70 minutes		32.	Which of the following	g steps NOT invol	ved in manu-	
22.	are solid do	sage form in which drugs or		facturing of "hard gel			
		osed in small water-soluble		A. Dipping	B. Spinnir	~	
	A. Tablet	<b>B.</b> Pills		C. Trimming	<b>D.</b> Roughi	ng	
	C. Capsule	<b>D.</b> Granules	33.	The plasticiser used in		of soft gelatin	
22	•			capsule is	<b>-•</b>		
23.	capsule.	ient of opaque hard gelatin		<ul><li>A. Sorbitol</li><li>B. Povidone</li></ul>			
	A. Glycerine	<b>B.</b> Titanium dioxide		<b>C.</b> Polyethylene glyco	1		
	C. Polyhydric alcohol			<b>D.</b> Hydroxypropyl met			

		lablets and	a Cap	osuies					23
34.	Which step is involved in h production? A. Dipping C. Trimming	<ul><li>ard gelatin capsule shells</li><li>B. Spinning</li><li>D. All of the above</li></ul>	38.	<b>D.</b> All	s dosage for	ve	hell can be	made	from
35.	The moisture content of is A. <10% C. 12-16%	a hard gelatin capsule  B. 10–13% D. >16%	39.	Empty	rragenan y <b>capsule</b> l	has moistu	B. HPMC D. All of the		
	Which capsule size has the A. 0 C. 3 Capsules are	smallest capacity?  B. 1  D. 5	40.	A. 609 C. 50-	–70% ipsule size i	number for drug is	B. 12–15% D. 30–64% the capacity	1	milli-
57.	A. Unit solid dosage forms  B. Liquid dosage form			<b>A.</b> 00 <b>C.</b> 2		ur ug 13	<b>B.</b> 1 <b>D.</b> 3		
		Answe	er Ke	у		_			
	1 B 2 B 3 C	4 D 5 B	6	В	7 D	8 C	9 Д	10	Δ

1	В	2 B	3 C	4 D	5 B	6 B	7 D	8 C	9 A	10 A
11	С	12 A	13 C	14 B	15 C	16 B	17 D	18 B	19 C	20 C
21	Α	22 C	23 B	24 B	25 B	26 D	27 C	28 B	29 B	30 A
31	Α	32 D	33 A	34 D	35 C	36 D	37 A	38 A	39 B	40 B

#### Liquid Oral Preparations Solutions, syrups, Elixirs, Emulsion, Suspensions

1.	is a formulation layer of emulsion from the o		7.	For an ideal suspension, the should be	the sedimentation volume
	<ul><li>A. Cracking</li><li>C. Creaming</li></ul>	<ul><li>B. Flocculation</li><li>D. Disintegration</li></ul>		<ul><li>A. Equal to one</li><li>B. Less than one</li></ul>	
2.		stability in the emulsion?  B. Cracking  D. All of the above	8.	<ul><li>C. More than one</li><li>D. Zero.</li><li>The size of dispersed part ranges from</li></ul>	ticles in coarse dispersion
3.	HLB means A. Hydrolytic lipolytic balan. B. Hydrophilic lipophilic bal C. Hydrogen lipid balance D. High lipid balance			<b>A.</b> 1 μm to 100 μm <b>B.</b> 1 nm to 100 nm <b>C.</b> 1 mm to 100 cm <b>D.</b> Less than 1 μm	
			9.	The primary emulsion form	
4.	HLB plays an important rolemulsion			<b>A.</b> 4:2:1 <b>C.</b> 2:2:1	
	<ul><li>A. Dissolution and disintegra</li><li>B. Formation and solubility</li><li>C. Formulation and stability</li><li>D. None of the above</li></ul>		10.	In suspensions insoluble drug in the vehice A. Suspending agents C. Surfactants	le.
5.	A. Griffin B. Darcy C. Poiseulle	ered HLB method.	11.	In suspensions, the stability upon  A. Sedimentation volume C. Particle size	B. Zeta potential
6.	A wetting agent is included suspension, particularly whe		12.	Simple syrup contains sucrose. A. 50% w/v C. 66.7% w/v	B. 60% w/v
	<ul><li>A. Are hydrophobic</li><li>B. Are denser than the vehicl</li><li>C. Are water soluble</li><li>D. Have lesser interfacial ten</li></ul>		13.	Elixir contains A. 10–50% C. 4–40%	amount of alcohol. B. 5–15% D. 100%

14.	<ul> <li>A is a mixture in which one substance of microscopically dispersed insoluble particles is suspended through another substance.</li> <li>A. Suspension B. Emulsion</li> <li>C. Colloid D. None of the above</li> </ul>						les is	23.	<ul> <li>A. Compound iodine throat paint</li> <li>B. Aqueous iodine solution</li> <li>C. Strong iodine solution</li> <li>D. Lugol's solution</li> </ul>								
15.	The pr  A. 1:2:			mula B. 2		latile	oil is	24.	requi	reluents			sser co				sucrose
	<b>C.</b> 3:2:	1		<b>D.</b> 1	:1:2				C. Bi	nders				<b>D.</b> A	Adjuva	ants	
16.				<b>B.</b> E			ve	25.	A. Di	n and soluents				<b>B.</b> \			
17.	Monophasic liquid dosage f				include			26.	Whic	h metl	od is	used	l for e	valu	ation	or stal	bility of
	A. Aro C. Solu	matic waters itions	S		Γinctures All of the				suspe A. Se	nsion?	tation	meth	od	В. І	Electro	kinetic	method
18.	Simple	syrup is a s	aturated	soluti	ion of		<u></u> .		C. Rh	neologi	cal m	ethod	[	<b>D.</b> A	All of	the abo	ove
	A. Suc							27.	Linim	nents m	ust no	ot be	applie	d on t	he		skin.
	C. Dex	trose		<b>D.</b> N	None of the	he abo	ve			velled					Brokei		
19.		alue of SLS	is						C. Pa	inful				<b>D.</b> 1	Vorma	ıl	
	<b>A.</b> 10 <b>C.</b> 40			<b>B.</b> 1 <b>D.</b> N		he abo	ve	28.	Immi	scibilit	ty of	oil aı 	nd wa	ter c	an be	overc	ome by
20.		mponent pr vn as vent		<b>B.</b> S	Solution	all qua	ntity		<ul><li>A. Formulating an emulsion</li><li>B. Formulating suspension</li><li>C. Formulating an insufflation</li><li>D. Formulating an elixir</li></ul>								
	C. Solu	ite		<b>D.</b> I	Liquid				<b>D.</b> Fo	rmulat	ing ar	elixi	ir				
21.	Which form?  A. Solu	of the follow	Ü		-	quid d	osage	29.	A. Concentrated C. Diluted					B. Warmed			
22	C. Susj				Enemas			30.	All aı		tests	used					mulsion
22,		lro-alcoholic			Aqueous Semi-soli	ds			A. Fl	uoresce ectric c	ence to	est	y test			ng test est	
Answe																	
	1 C	2 D	3 B		4 C	+	A		A	7			D	+	A 6	_	A C
1	1 D	12 C	13 C		14 A	15	В	16	Α	17	ם	18	Α	19	9 C	20	) C

21 C

22 A

23 A

24 B

25 C

26 D

27 B

28 A

29 C

30 D

Topical Preparations
Ointments, Creams, Pastes, Topical Gels, Liniments, Lotions, Suppositories, Pessaries, Nasal Preparations, Ear Preparations, Powders, Granules

1.	disintegrate o	r melt at body temperature.	9.	Suppositories are generally e	evaluated by
	A. Powders	<b>B.</b> Suppository		A. Melting range test	<b>B.</b> Breaking test
	C. Tablet	<b>D.</b> Capsule		C. Liquefaction	<b>D.</b> All of the above
2.	Suppositories intended	for the vagina are called	10.	The intracellular absorption in which the drugs diffuse	through the
	A. Cone	<b>B.</b> Suppositories		present in the stratum corn	eum.
	C. Bougies	<b>D.</b> Pessaries		<b>A.</b> Lipids	<b>B.</b> Proteins
3.	Weight of rectal supposito	ry is generally grams.		C. Minerals	<b>D.</b> None of the above
	<b>A.</b> 5–6	<b>B.</b> 1–2	11.	Toothpastes are	_ type of pastes.
	<b>C.</b> 3–4	<b>D.</b> 6–7		<b>A.</b> Fatty	<b>B.</b> Aqueous
4	Cumpositorios intended fo	with a mass and weathers are		C. Non-aqueous	<b>D.</b> Hydrocolloids
4.	called	or the nose and urethra are	12.	Agents which prevent produ	cts from drying are called
	A. Bougies	<b>B.</b> Pessaries		•	
	C. Suppositories	<b>D.</b> None of the above		A. Astringent	B. Antiseptic
5	Theobroma oil is also kno	awn as		C. Humectants	<b>D.</b> Antimicrobial
٥.	A. Bees wax	B. Lanolin	13.	Bentonite is an example of	•
		<b>D.</b> Macrogols		A. Hydrogel	B. Organogel
				C. Xerogel	D. Non-aqueous gel
6.	Witepsol and Massupol ar to type	e suppository bases belongs	14.	Gels are prepared by	method.
	A. Fat bases			A. Fusion	<b>B.</b> Dispersion
	<b>B.</b> Water soluble bases			C. Cold	<b>D.</b> All of the above
	C. Emulsifying bases		15.	Liniments are applied with	•
	<b>D.</b> None of the above			<b>A.</b> Without friction	
7.	of suppositors	mould is necessary to avoid		C. Without massage	<b>D.</b> None of the above
٠.	sticking of suppository to		16.	Calamine lotion is used as	•
	A. Lubrication	<b>B.</b> Purification		A. Topical protectant	
	C. Calibration	<b>D.</b> Washing		C. Antidote	<b>D.</b> Antipyretic
8.	Weight of pessaries varies	s from gm.	17.	Macrogols are	
	<b>A.</b> 1–2	<b>B.</b> 2–3		A. Hydroalcohols	
	<b>C.</b> 4–8	<b>D.</b> 8–10		C. Ethylene dioxides	<b>D.</b> Methylsalicylate

18.		weight of medicament that	30.	are water-sol	uble ointment basis.			
	displaces one part by weight	ght of base is		A. Cetrimide	B. Macrogols			
	A. Density value	<b>B.</b> Saponification value		C. Wool fat	D. YSP			
	C. Iodine value	<b>D.</b> Displacement value	31	YSP stands for				
19	Petrolatum are example of	of	51.	A. Yellow soft polymer	- <b>'</b>			
1).	<b>A.</b> Water-soluble bases	B. Water removable		<b>B.</b> Yellow soft polycarbon				
				C. Yellow soft paraffin				
	C. Emuision bases	<b>D.</b> Hydrocarbon bases		<b>D.</b> Yellow soft polyhalide				
20.	Sulphur ointment is prepare	ared by		1 2				
	<b>A.</b> Emulsification method		32.	Wool fat is also called				
	<b>B.</b> Fusion method			<b>A.</b> Anhydrous lanolin				
	C. Levigation method			C. Petrolatum	<b>D.</b> Vaseline			
	D. Chemical reaction meth	od	33.	Paste usually contains				
21	Pastes consists of	% of solids		A. Beewax	B. Glycerin			
21.	<b>A.</b> 0–10	B. 20–50		C. Wool alcohol	•			
	<b>C.</b> 100	<b>D.</b> 0	2.4					
	<b>C.</b> 100	<b>D.</b> 0	34.	Liniments usually contain co	ounter irritants like			
22.		solids to allow		A. Almond oil				
	perspiration.			B. Beewax				
	<b>A.</b> Porous	<b>B.</b> Larger		C. Lanolin				
	C. Smaller	D. Fine		<b>D.</b> Methyl salicylate				
23.	The length of urethral bo	ugies is	35.	Gum Arabic is a	·			
	<b>A.</b> 100–150 mm			<b>A.</b> Anionic polysaccharide				
	<b>C.</b> 10–20 mm	<b>D.</b> 10–50 mm		<b>B.</b> Cationic polysaccharide				
				C. Neutral polysaccharide				
24.	Which of the following mechanical equipment can be			<b>D.</b> None of the above				
	used for emulsification?		36.	Which of the following is no	ot a semisolid dosage form?			
	A. Homogenizers			A. Paste	<b>B.</b> Creams			
	C. Ultrasonifiers	<b>D.</b> All of the above		C. Ointments				
25.	Which of the following is	not used as an emulsifying	25		-			
	agent?		3/.	Generally, pastes contain				
	A. Surfactant	<b>B.</b> Hydrophilic colloids		A. High percentage of insol				
	C. Electrolytes	• •		B. Low percentage of insolu	uble solids			
	•	•		C. Both A and B				
26.		earbon in semisolid dosage		<b>D.</b> None of the above				
	forms is		38.	Lotion is fluid preparation	applied to skin			
	A. Petrolatum	B. Mineral oil		A. Without friction				
	C. Both A and B	D. None of the above		<b>B.</b> With friction				
27.	A suppository is generally	intended for use in		C. With massage				
	A. Rectum	B. Vagina		<b>D.</b> None of the above				
	C. Urethra	<b>D.</b> All of the above	39.	are used for soothing, cooling and				
				softening effect on the skin	•			
28.	Vaginal suppositories also			A. Liniment	B. Paste			
	A. Pessaries	<b>B.</b> Simple suppositories		C. Lotion	D. Gargles			
	C. Bougies	<b>D.</b> None of the above	40		•			
29	Which of the following is	not a vegetable oil?	40.	The most common vehicle	t tot masat preparation is			
•	A. Peanut oil	B. Almond oil		A. Water	<b>B.</b> Glycerine			
	C. Olive oil	<b>D.</b> Petrolatum		C. Ethyl alcohol	<b>D.</b> Propylene glycol			
	C. Onvoon	D. I Cabiatain		C. Laryr alconor	D. Tropyrene grycor			

41.		with nasal fluid.	50.		ective ingredients for relief			
	<b>A.</b> Hypertonic	<b>B.</b> Hypotonic		from itching and hard wa				
	C. Isotonic	<b>D.</b> Paratonic		<b>A.</b> Sodium bicarbonate	•			
42.	are also call	ed otic or aural preparations.		C. Both A and B	<b>D.</b> None of the above			
	A. Nasal drop	<b>B.</b> Ear drop	51.	One of the common uses	of nasal solution is for the			
	C. Eye drop	<b>D.</b> None of the above		relief of				
				A. Cough	<b>B.</b> Congestion			
43.		erfere with the		C. Throat infection	<b>D.</b> Body pain			
	action of epithelial cilia o	of nasal mucosa.						
	<b>A.</b> Smoothening action		52.		os should be			
	<b>B.</b> Cleansing action				<b>B.</b> Non-toxic			
	<b>C.</b> Relieving action			C. Non-irritant	<b>D.</b> All of the above			
	<b>D.</b> All of the above		53	The wrenning of nowde	r in two papers (packing)			
44	Ear drop solution is generally prepared by using			covering is called				
77.	Lat utop solution is ge	nerally prepared by using		A. Flexible packing	•			
	A. Water			B. Blister packing				
	<b>B.</b> Glycerine and propyler	ne alveol		C. Double wrapping				
	C. Dil alcohol	ne grycor		D. Strip packing				
	<b>D.</b> All of the above			D. Strip packing				
	D. All of the above		54.	Double wrapping is essen	tial for drugs			
45.		to instilling the nasal drops		A. Hydrophilic				
	in nasal cavity.			<b>B.</b> Hygroscopic				
	A. Soft containers			C. Lipophilic				
	<b>B.</b> Sifters			<b>D.</b> None of the above				
	C. Dropper							
	<b>D.</b> None of the above		55.		powders are applied on			
16	Nosal duan should be isa	tania with		umbilical cord of infants?				
40.	Nasal drop should be iso A. 0.3% Sodium chloride			A. Dusting powders				
				B. Surgical dusting powders				
	<b>B.</b> 0.4% Sodium chloride			C. Cosmetic dusting powd	lers			
	C. 0.5% Sodium chloride			<b>D.</b> All of the above				
	<b>D.</b> 0.9 % Sodium chloride		56.	Effervescent granules usu	ially contain .			
47.	Ear drops are generally	used for		A. Sodium bicarbonate, ci	-			
	A. Cleaning			B. Sodium sulphate, citric				
	<b>B.</b> Softening the wax			C. Calcium carbonate, citr				
	C. Treating the mild infec	etions		<b>D.</b> Sodium chloride, citric				
	<b>D.</b> All of the above							
			57.		e medicated powders which			
48.	Most preferred vehicles f	for ear drop is			or nasal decongestant action			
	A. Glycerin			<b>A.</b> Dusting powders				
	<b>B.</b> Propylene glycol			<b>B.</b> Tooth powders				
	C. Both A and B			C. Snuffs				
	<b>D.</b> None of the above			<b>D.</b> Medicated dusting pow	ders			
49.	The ear drop label should	d be stated as	58.	must be ste	rile.			
	<b>A.</b> For external use only			A. Face powder				
	<b>B.</b> For internal use only		<b>B.</b> Surgical powder					
	C. Dilute before use			C. Both A and B				
	<b>D.</b> None of the above			<b>D.</b> None of the above				

59.	is a finely divided powder meant for	67.	is supplied in a sterile form.			
	introduction into the body cavities		<b>A.</b> Tooth powder			
	A. Inhalation		B. Surgical powder			
	<b>B.</b> Dusting powders		C. Both A and B			
	C. Medical powder		<b>D.</b> None of the above			
	<b>D.</b> Insufflation	60	The method used when notent substances are to be			
60.	powder contains more than one	00.	The method used when potent substances are to be mixed with a large amount of diluents is			
ov.	ingredient.		A. Solubility			
	A. Simple powder		B. Salvation			
	B. Compound powders		C. Geometric dilution			
	C. Both A and B		D. Complexation			
	<b>D.</b> None of the above		•			
	B. None of the above	69.	Abrasive agents used in dentifrices are			
61.	Method of granule formulation is		A. Calcium sulphate			
	<b>A.</b> Spatulation <b>B.</b> Fusion method		<b>B.</b> Magnesium carbonate			
	<b>C.</b> Wet method <b>D.</b> Both B and C		C. Sodium carbonate			
62	Dusting powders are dispensed in		<b>D.</b> All of the above			
02.	containers.	70.	Advantage of powders is			
	A. Plastic	, 0.	A. Easy to adjust the dose			
	B. Sifter top container		B. Rapid onset of action			
	C. Cardboard box		C. No difficulty in swallowing			
	<b>D.</b> None of the above		<b>D.</b> All of the above			
	B. None of the above					
	Medicated dusting powders are mainly used for  A. Superficial skin conditions B. In body cavities C. During surgery D. In major wounds	71.	particles which are supplied in cylindrical or flat metal boxes with hinge lid meant for inhaling through the nostrils for decongestion.  A. Snuffs B. Dentifrices C. Eutectic powder			
64.	Liquefaction of some solid substances when mixed		<b>D.</b> Explosive powder			
	in a dry form due to adsorption of moisture from environment is due to	72.	are intended for cleaning the teeth or			
	A. Immiscibility		other parts of oral cavity using a toothbrush.			
	B. Insolubility		A. Snuffs			
	C. Hygroscopicity		<b>B.</b> Dentifrices			
	<b>D.</b> None of the above		C. Eutectic powder			
	b. None of the above		<b>D.</b> Explosive powder			
65.	Containers are labelled with direction, immerse in water for a few seconds and then swallow with draught of water.	73.	Powders are meant for cleansing or irrigation of body cavity with a suitable liquid especially for cleansing vaginal cavity are			
	A. Dusting powder B. Cachets		<b>A.</b> Douches <b>B.</b> Snuffs			
	C. Insufflations D. Snuffs		C. Insufflations D. Eutectic			
66.	Powders consisting of more than one ingredient are called  A. Compound powders  B. Simple powders  C. Both A and B  D. None of the above	74.	Powder containing mixture of powdered ingredients which produces explosion on trituration is  A. Snuffs B. Dentifrices C. Eutectic powder D. Explosive powder			

1 E	В	2 D	3 B	4 A	5 C	6 C	7 A	8 C	9 D	10 A
11 [	D	12 C	13 A	14 D	15 B	16 A	17 B	18 D	19 C	20 C
21 E	В	22 A	23 A	24 D	25 C	26 C	27 A	28 A	29 D	30 B
31 (	C	32 A	33 B	34 D	35 C	36 D	37 A	38 A	39 C	40 A
41 (	С	42 B	43 B	44 D	45 C	46 D	47 D	48 C	49 A	50 C
51 E	В	52 D	53 C	54 B	55 B	56 A	57 C	58 B	59 D	60 B
61 [	D	62 B	63 A	64 C	65 B	66 A	67 B	68 C	69 D	70 D
71 /	Α	72 B	73 A	74 D	75 A	76 C	77 A	78 D	79 C	80 A

## Sterile Formulations and Immunological Products

1.	TEN means		0.	LAL stanus for	<u> </u>	
	<b>A.</b> Total pure nutrition			<b>A.</b> Limulus acyte lysate		
	<b>B.</b> Total polymer nutrition			<b>B.</b> Limulus amoebocyte ly	ysate	
	C. Total parenteral nutrition			C. Left auricular lymphoc	yte	
	<b>D.</b> Total paratonic nutrition			<b>D.</b> Lipo acrylic lysate		
2.	SVPs means		9.	pH of tear fluid is		_•
	<b>A.</b> Slow volume parenterals			<b>A.</b> 2.1	В.	4.5
	<b>B.</b> Small value parenterals			C. 9.2	D.	7.4
	C. Small volume parenterals		4.0		***	
	<b>D.</b> Small viscous parentals		10.	Which of the following is	_	
3	The volume of small volum	ne narenteral is less than		A. EDTA		Lanolin
٥.	•	ie parenterar is less than		C. Paraben	D.	Sodium alginate
	<b>A.</b> 100 ml	<b>B.</b> 200 ml	11.	Injections are	pre	parations.
	<b>C.</b> 300 ml	<b>D.</b> 500 ml		A. Conventional	B.	Non-sterile
1	SVPs are manufactured as			C. Sterile	D.	Traditional
٦.	A. Multidose containers	•	12	Drugs are injected into su	uhara	phnoid space is called
	B. Single dose containers		14.	route.	uparac	chinoid space is caned
	C. Both A and B			A. Intravenous	R	Intrathecal
	<b>D.</b> None of the above			C. Intra-articular		
5.	is used to det matter in parental products		13.	Viscosity enhancer in o	phtha	ilmic preparation is
	A. Pyrogen test			A. Poly vinyl alcohol	В.	Povidone
	C. Sterility test			C. Dextran	D.	Macrogol
6.	Bacterial endotoxin test is a	lso known as	14.	Ophthalmic solution is st	erilize	d by
	A. Clarity test	<b>B.</b> LAL test		A. Autoclave	В.	Hot air oven
	C. Leaker test	<b>D.</b> Sterility test		C. Membrane filter	D.	Bacterial filters
7.	Drugs are injected between nerve is called	first and second cervical route.	15.	Drugs are injected into	bone i	s called
		<b>B.</b> Intrathecal		A. Intravenous	В.	Intrathecal
	C. Intra-articular	<b>D.</b> Peridural		C. Intra-articular		Peridural

16.	Which of the following is:	antibacterial agent?	26.	LVPs mean	
	A. Methylcellulose			A. Less volume parenterals	
	<b>B.</b> Paraffins			<b>B.</b> Large volume parenteral	S
	C. Benzalkonium chloride			C. Large vacuum parentera	ls
	<b>D.</b> Tween			<b>D.</b> Large volume polymers	
17.	Which one of the following	is used to adjust the isotoni-	27.	LVPs are packed in large	•
	city?			A. Multi dose containers	
	A. Dextrose	B. Boric acid		<b>B.</b> Single dose containers	
	C. NaCl	<b>D.</b> All of the above		C. Both A and B	
10	To increase contact time of	f preparation to eye which		<b>D.</b> None of the above	
10.	of the following additives		28.	is performed	to test degree of immunity
	A. Carboxy methyl cellulos			of an individual against di	
	<b>B.</b> Polyethylene glycol			<b>A.</b> Killer test	<b>B.</b> Tuberculin test
	C. Methyl cellulose			C. Schick test	
	<b>D.</b> All of the above		20	The smallpox vaccines as	
10	Delvasubate 00 is used in a	uhthalmia nuonauatiana aa	29.	between	ie storeu at temperature
19.	·	ophthalmic preparations as		<b>A.</b> 8° and 10°C	<b>B.</b> 2° and 8°C
	<b>a A.</b> Preservative			C. 5° and 7°C	<b>D.</b> 10° and 12°C
	<b>B.</b> Antioxidants				
	C. Wetting agents		30.	The following are endotox	
	<b>D.</b> All of the above			-	<b>B.</b> Opsonins
				C. Agglutinins	<b>D.</b> Virulence
20.	Thermolabile drug contai sterilized by	ning ophthalmic solution is	31.	The immunity developed at from a disease is called	
	A. Oven	<b>B.</b> Autoclave		A. Passive immunity	
	C. Membrane filtration	<b>D.</b> All of the above		C. Both A and B	<b>D.</b> None of the above
21.	Which of the following w	vax is used to prepare eye	32.	Vaccines are prepared from	n immune
	ointment?			A. Vitamins	B. Blood
	A. Bees wax	<b>B.</b> White soft wax		C. Serum	<b>D.</b> Plasma
	C. Carnauba wax	<b>D.</b> None of the above	22		
2.2	Ontimum viscosity of or	ohthalmic solution ranges	33.	BCG vaccine is used for _	
	between	minumic solution ranges		A. Tuberculosis	
	<b>A.</b> 15 and 25 cps	<b>B.</b> 10 and 15 cps		C. Both A and B	<b>D.</b> None of the above
	C. 25 and 30 cps	<b>D.</b> 30 and 40 cps	34.	Which of the following provi against pathogens?	des the long-term immunity
23.	% bioavailab	ility is given by IV route.		A. Naturally acquired passi	ve immunity
	<b>A.</b> 50	<b>B.</b> 80		<b>B.</b> Artificially acquired pass	•
	<b>C.</b> 100	<b>D.</b> 70		C. Naturally acquired active	•
24.	LVPs are administered in	volume of ml		<b>D.</b> All of the above	•
	per day by slow IV drip.		35.	The branch of biology which	h involves immune systems
	<b>A.</b> 1–10	<b>B.</b> 20–50		in all organisms is known	as
	<b>C.</b> 50–100	<b>D.</b> 250–1000		A. Zoology	<b>B.</b> Microbiology
25.	is a non-an	ueous oily vehicle used in		C. Immunology	<b>D.</b> Biotechnology
<b>4</b> J.	parenterals.	acous ony venicie useu III	36.	The first vaccine was devel	loped by
	A. BHT	<b>B.</b> BHA	•	A. Louis Pasteur	B. Edward Jenner
	C. WFI	<b>D.</b> Arachis oil		C. Carl Landsteiner	

37.	Which a lifetir	of the follo	wing in	nmun	ity is	obtaiı	ned dı	ıring				y acqu Ily acc	-			-			
38.	A. Acq C. Pass for the	uired immusive immuni are t prevention	ty the prepof dise	D parat ases s	B. Active immunity D. None of the above arations which are meant ses such as vaccines or for as antitoxin and antiserum					C. Naturally acquired passive immunity									
39.	<ul><li><b>A.</b> Imn</li><li><b>B.</b> Imn</li><li><b>C.</b> Loti</li><li><b>D.</b> Non</li></ul>	nunological	product ve						45.	<ul><li>sible f</li><li>A. Ac</li><li>B. Pa</li><li>C. In</li></ul>	for pretive it is ssive that in the interior i		ng us f ity nity ty					espon- oecies?	
	B. Wea	antigenic proakened pathore attenuated of the above	ogen pathoge	en					46.	<b>A.</b> Ba <b>B.</b> Ba	acteria acillus	l Colo Cox C	ny Gu Calmet	te					
40.	disease A. Dig	of the follo- causing pa estive system retory system	<b>thogen</b> n	s? B	. Res	piratoı		em	<ul> <li>D. Bacterial Centric Guerin</li> <li>47. Whooping cough is caused by</li> <li>A. Mycobacterium tuberculosis</li> <li>B. Vibrio cholerae</li> </ul>										
41.	A. Rec	tis is an exa ombinant va oids vaccine	accine	В	. Sub		accine cine		48.	<b>D.</b> Bo	ordete	nonas lla per in	tussis						
42.		of the follov form phago	_		the in	ımune	syste	m do		A. Li	ve ant	igen ies				ttenuat		_	
43.	A. Eos. C. Neu Newbo	-	ir antih	D		ropha	_	milk.	49.	A. Ki	lled R	ccine is Licketts ekettsia	sia		<b>B.</b> K	illed ba			
	This is A. Nati	an example urally acquir ficially acqu	of	ve im	· munit	y			50.	Mant	oux to	e <b>st is</b> _ lin test		•	<b>B.</b> So	chick to	est		
								Answ	er Key	,									
-	1 C	2 C	3			Α	<del>                                     </del>	D	+	В		Α		В	-	С	+	D	
	1 C 1 B	12 B 22 A	13 23		14 24		15 25		16 26			D B	18 28		-	C D	30	C D	

31 B

41 A

32 C

42 B

33 A

43 C

34 C

44 B

35 C

45 C

36 B

46 C

37 A

47 D

38 B

48 C

39 D

49 A

40 D

50 A

#### Quality Control and Quality Assurance

1.	SOP means			<b>C.</b> Equipment validation	
	A. Sample operating procedu	ires		<b>D.</b> All of the above	
	B. Standard operating proced	lures	0		
	C. Some operating procedure		8.	of an in	strument of the process of
	<b>D.</b> Standard ongoing procedu			determining its accuracy.	<b>D</b> 0 111
_	0 01			<b>A.</b> Validation	<b>B.</b> Calibration
2.	cGMP means			C. Documentation	<b>D.</b> Quality control
	A. Current Goal Manufacturi	•	9	LAFH is .	
	B. Current Good Manufactur	0	7.	A. Light air and fluid homo	ganizar
	C. Current Good Maintenand	e Practices		<b>B.</b> Left air flow heat	gemzer
	D. Current Good Managing I	Practices		C. Laminar attributes of air	humidita
3	explains the	CMP and requirement		<b>D.</b> Laminar air flow hood	numany
٥.	of premises, plants, equipm			<b>D.</b> Lammar an now nood	
	products.	ients for pharmaceuticar	10.	BMR is	
	•	<b>B.</b> Schedule M		A. Batch manufacturing rec	cord
		D. Schedule Y		<b>B.</b> Bilayer manufacture reg	
	C. Benedule W	D. Schedule 1		C. Batch multilayer record	
4.	cGMP regulations are estab	lished by		<b>D.</b> Bilayer multilayer regist	er
	A. FDA	B. PCI	4.4	, , ,	
	C. IDMA	<b>D.</b> IPA	11.	The dispensing area has the	
_				A. 80 Pascal	
5.	is the docume			C. 20 Pascal	<b>D.</b> 10 Pascal
	any procedure process, equi	_	12.	is the docume	ent evidence for the equin-
	or system actually leads to t		12,	ment tested at the manufac	
	A. Quality control	~ •		by end-user.	pour era prince una aperinca
	C. Validation	D. GMP		A. Site acceptance test (SA)	T)
6.	The soft gelatine capsule invo	olves the process of encap-		<b>B.</b> Factory acceptance test (SA	
	sulation of medicaments with			C. Both A and B	(1711)
	<b>A.</b> 0.1–30	<b>B.</b> 40–50		<b>D.</b> None of the above	
	<b>C.</b> 60–70	<b>D.</b> 80–90			
			13.	Lower the class number, _	the cleansing of
7.	Which of the following is a	type of validation?		air.	
	<b>A.</b> Process validation			<b>A.</b> Higher	<b>B.</b> Lower
	<b>B.</b> Cleaning validation			C. Neutralizes	<b>D.</b> Stabilizes

								,			,									
14. 15.	Quality A. Des B. Mar C. Cor D. Cor the equ and ha facility	tructive nageria rective structi ipmer ve un	re tool al tool e tool ive too  at that dergo	ol is it is no	the o t dan	locum naged (	durin ng at	g tran end v	sport iser's		The resmall use is A. Ac. Proce A. Proce B. Re C. Co	ccuracy ecision ss vali	dation	dication  includation  idation	udes _	B. D.	<b>Abilit</b> Robu Linea	y dun Istne arity	ring n	
	<ul><li>user.</li><li>A. Site</li><li>B. Face</li><li>C. Both</li><li>D. Nor</li></ul>	accep tory ac h A an	tance cceptar d B	test (SA	AT)					19.	The ir of an a A. Ra C. Ac	nterva analyt inge	l betwee is	een l	·	В.	h <b>ighe</b> Linea Preci	arity	ncent	ration
16.	Validat A. Pros B. Rett C. Con D. All	spectiv cospec currer	ve vali tive va it valid	dation alidatio dation			•			20.	FDA s A. Fo B. Fo C. Fo D. Fo	od and od and od and	l Dose l Distr l Drug	Adm ibutio Adm	ninistra on Adr ninistra	ninis ition	stratio	on		
									Answe	er Key	,									
	1 B	2	В	3	В	4	A	5	С	6	Α	7	D	8	В		9 D		10	Α

17 B

18 D

19 A

20 C

11 C

12 B

13 A

14 B

15 A

16 A

#### Novel Drug Delivery System

1.	A. Hydrophilic molecule	· of		<ul><li>C. The membrane is permeable to water</li><li>D. The membrane must swell</li></ul>			
	<ul><li>B. Hydrophobic molecule</li><li>C. Both A and B</li></ul>		8.	is the disadvantage of the NDDS.			
	<b>D.</b> None of the above			A. Higher cost			
2.	Liposomes have	half life.		B. Dose dumping			
_,	A. Longer C. Intermediate	B. Shorter D. Both A and B		<ul><li>C. Poor <i>in-vitro–in-vivo</i> correlation</li><li>D. All of the above</li></ul>			
3.	is a method of pre A. Pan coating C. Solubilization	<ul><li>paration of nanoparticle.</li><li>B. Filtration</li><li>D. Precipitation</li></ul>	9.	<ul> <li>Hydration activated drug delivery systems depend on</li> <li>A. Hydration of polymer</li> <li>B. Ionic strength within the reservoir</li> </ul>			
4.	Which among the following with conventional drug deliver.  A. Lower effectiveness			<ul><li>C. Osmotic pressure and ionic strength</li><li>D. Osmotic pressure and environment in GIT</li></ul>			
	<ul><li>B. Easy of manufacturing</li><li>C. Decrease side effect</li><li>D. Septal and temporal control</li></ul>	ol	10.	Which statement is correct with respect to Novel Dru Delivery Systems?  A. It causes fluctuation of blood levels			
5.	Drug release from the feedback systems depends uponA. Concentration of biochem	·		<ul><li>B. It cannot be target specific</li><li>C. It increases toxicity of the drug</li><li>D. It reduces side effects of the drug</li></ul>			
	<ul><li>B. Physicochemical propertie</li><li>C. Thickness of the polymer</li><li>D. None of the above.</li></ul>	_	11.	Drug release from osmotic drug delivery system depends on  A. Ionic strength			
6.	NDDS improves  A. Solubility properties of dr B. Pharmacokinetics			<ul><li>B. Osmotic pressure</li><li>C. Osmotic pressure and ionic strength</li><li>D. Osmotic pressure and environment in GIT</li></ul>			
	<ul><li>C. Pharmacodynamics</li><li>D. All of the above</li></ul>		12.	NDDS means A. Non-Drug Delivery System			
7.	Osmotic drug delivery systematics. A membrane that is soluble B. The membrane is impermental.	e at intestinal pH		<ul><li>B. Novel Drug Discovery System</li><li>C. Novel Drug Delivery System</li><li>D. Novel Dose Delivery System</li></ul>			

	Novel Drug	Delivery System	37
13.	Nanoparticles have a particle size range from  A. 50–100 nm B. 200–500 nm C. 300–700 nm D. 100–200 nm	. 17. CDDM means  A. Controlled Drug Delivery Molecules  B. Control Drug Dissolution Modules  C. Controlled Drug Delivery Modules  D. Constant Drug Delivery Modules	
14.	Drug release from the activation modulated drudelivery systems depends upon  A. External/internal stimuli  B. Physicochemical properties of the drug  C. Thickness of the polymer coating  D. None of the above	18. Which of the following are 'films and strips' a Drug Delivery system?  A. Spray bandages B. Buccal strips C. Zero order release film D. All of the above	as a Novel
15.	Bio-responsive activated drug delivery systems relearate depends on  A. Hydration of polymer  B. Concentration of biochemical substance  C. Osmotic pressure and ionic strength  D. Osmotic pressure and environment in GIT	are small sterile hypodermic table are placed under skin by minor surgery.  A. Liposomes B. Implants C. Nanoparticles D. Microspheres	lets which
	The size of microsphere ranges from μ A. 1–1000 B. 1000–1500 C. 1500–2000 D. None of the above	20. Liposomes are spherical structures usually in a diameter nm  A. 80–100 B. 60–100 C. 55–1000 D. 15–1000	between
	An	swer Key	

1 C	2 B	3 D	4 A	5 A	6 D	7 C	8 D	9 A	10 D
11 B	12 C	13 B	14 A	15 A	16 A	17 C	18 D	19 B	20 D