

GPAT - 2014

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GPAT QUESTIONS

1.	, system does	not have orifice to release	the	drug.		
	(a) Elementary Osmotic F	Pump	(b)	L-OROS		
	(c) Sandwich Osmotic Pu	ımp Tablet	(d)	Controlled Porosity O	smot	ic Pump Tablet
2.	In which rearrangement	reaction, Isocyanate is forr	ned?			
	(a) Curtious	(b) Lossen	(c)	Both A & B	(d)	None
3.	Chitin gets converted in to	Chitosan upon:				
	(a) Acetylation	(b) Deacetylation	(c)	Oxidation	(d)	Reduction
4.	All of the following are lea	of constants EXCEPT				
	(a) Vein-islet number		(b)	Vein-termination nu	ımbe	r
	(c) Stomatal number		(d)	Leaf fiber		
5.	Zink chloride is added to n	nouth wash because it acts	as	SION		
	(a) Fragrance	(b) Astringent	(c)	Cooling agent	(d) A	Antibacterial
6.	The choice of route of adr	ninistration plays an impo	rtant	role in action of direct	ly ac	ting cholinomimetic
	Adverse effect of choli	ne esters that may be a	avoi	ded by selection of a	an ap	propriate route o
	administration is					
	(a) Bradycardia	(b) Hypotension	(c)	Delirium	(d)	Sweating
7.	Sieve size 80 has opening	g of				
	(a) 0.100 mm	(b) 0.125 mm	(c)	0.150 mm	(d)	0.180 mm
8.	The ideal saponification v	ralue for suppository base	is			
	(a) 50-100	(b) 100-150	(c)	150-200	(c)	200-500
9.	o, m, p- isomers can be di	fferentiated on the basis o	f:			
	(a) Chemical shift		(b)	Coupling constant		
	(c) Extinction coefficient		(d)	Dipole moment		
10.	Which of the following dr	ug comes under Schedule (Z_1			
	(a) Opium	(b) Ergot	(c)	Fish liver oil	(d)	Insulin
11.	Source of amla is					
	(a) Phyllanthus inruri		(b)	Terminiliachebula		
	(c) Terminalia Bacteria		(d)	Embilca officinalis		
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12.	What is the unit of diek	ectric constant		
	(a) Dyne	(b) Debey	(c) Farad/meter	(d) No Unit
13.	Monitoring of plasma d	rug concentration is requ	ired while using:	
	(a) Antihypertensive of	lrugs	(b) Levodopa	
	(c) Lithium carbonate		(d) MAO inhibitors	
14.	Of the following antibi	otics, which one would b	pe acceptable to use who	en treating penicillin resistan
	S. pnumoinae otitis med	dia		
	(a) Azithromycin	(b) Clarithromycin	(c) Cefuroxime	(d) Cefaclor
15.	Addtion of which of the	following to a large volun	ne parntral product is no	t advised
	(a) Active pharmaceut	ical ingredient	(b) Preservatives	
	(c) Buffering agens		(d) Tonicity adjuste	rs
16.	A drug suspension de	composes by zero-order	kinetics with a rate co	onstant of 2 mg mL ⁻¹ month
	if the initial concentrati	on is 100 mg mL ⁻¹ what is	s the shelf life	
	(a) 2 months	(b) 3 months	(c) 4 months	(d) 5 months
17.	Sanguinarine belongs to	o the subgroup of:	PAT	
	(a) Morphinans		— (b) Benzyl isoquino	lines
	(c) Phthalide isoquinol	lines \ DIS	(d) Benzophenanth	renes
18.	Antidote for paracetam	ol overdosing is C E	NTER	
	(a) Atropine	(b) N- Acetly cysteine	(c) Glutathione	(d) Theophylline
19.	Which one of the follow	ving drug combination is o	contraindicated	
	(a) Glyceryl trinitrate a	ınd sildenafil	(b) Amoxicillin	and clavulanic acid
	(c) Losartan and hydro	ochlorothiazide	(d) Pyrimthemaine	and sulfadoxine
20.	Which sugar is suitable	for diabetic patient		
	(a) Fructose	(b) Lactose	(c) Mannitol	(d) Sucralose
21.	Headquarter of Bureau	of Indian standards is situ	iated at	
	(a) New delhi	(b) Mumbai	(c) Kolkata	(d) Chennai
22.	Identity the structure of	barbituric acid		
	(a)	(b) O	e) (d)
			O H	,
	ЙН	ЙН	ŅΗ	

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23.	Ehtics	for 1	oharmacist are	put forth	bv

(a) PCI

- (b) CDSCO
- (c) AICTE
- (d) WHO

24. What is the IUPAC name of the following compound



(a) Bicyclo[2.2.2] octane

(b) Tricyclo[2.2.2] ontane

(c) Bicyclo[2.2.0] ontane

(d) Bicyclo [2.2.1] heptanes

- 25. Morphine does not cause:
 - (a) Constriction of pupil

(b) CNS depression

(c) Respiratory depression

(d) Diarrhoea

- 26. Which of the following is seed gum
 - (P) Gaur gum

(Q) Locust bean gum

(R) Xanthan gum

(S) Gellan gum

(a) P and Q

(b) R and S

(c) Q and R

- (d) P and S
- 27. The cancer that derived form ectoderm of endoderm of epithelial cells is
 - (a) Carcinoma
- (b) Sarcoma
- (c) Leukaemia
- (d) Myloid

- 28. Which of the following is/are marine anticancer
 - (a) Trabectadine

(b) Eribulin

(c) Cytarabine

- (d) All of the above
- 29. Identity the compound which is derived form typtophan
 - (a) Pilocarpine
- (b) Ephedrine
- (c) Muscarine
- (d) Quinoline

- 30. Opium, cocoa, poppy straw are given in
 - (a) Schedule H
 - (b) Schedule X
 - (c) Narcotic drugs and Psychotropic substances act 1998
 - (d) Schedule C
- 31. Which of the following will be inert in NMR spectrometry
 - (a) 13C

(b) 31P

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(c) 2H

(d) 1H

- 32. What is the relationship between keto and enol tautomers
 - (a) Resonance forms

- (b) Steriosomers
- (c) Constituonal isomers
- (d) Different conformations of the same compound

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33.	Which of the following is following is true for natura	l kill	er cells		
	(a) They may phagocytose tumor cells				
	(b) Killing of cells is enhanced by interleukin-2				
	(c) They recognize and kill some virus-infected cells	S			
	(d) Killing of cells is stimulated by prostaglandin E2				
34.	Evaluation of colour is tablets is done by				
	(a) Reflectance spectrophotometer	(b)	Tristimulus colorimet	er	
	(c) Microreflectance photometer	(d)	All of the above		
35.	The disintegration time of the effervescent tablets is				
	(a) 2 minutes (b) 2.4 minutes	(c)	3.5 minutes	(d)	5 minutes
36.	Identify the false statements about magmas:				
	(P) The addition of suspending agents to magmas is	alw	ays necessary		
	(Q) Magmas differ from gels in that their suspended	l par	ticles are larger		
	(R) Magmas are two- phase systems				
	(S) Magmas basically are gets				
	(a) P and Q (b) Q and R	(c)	Only P	(d)	Only S
37.	All of the following ACE inhibitors are prodrugs EXC	EPT	SION		
	(a) Ramipril (b) Lisinopril C E 1	(c)	Enalapril	(d)	Perindopril
38.	All of the following is resistant to both true and pseu	ıdo c	holinesterase enzyme	S	
	(a) Carbachol	(b)	Acetylcholine		
	(c) Methacholine	(d)	Pilocarpine		
39.	Globule size of parenteral emulsion should be				
	(a) $0.1 \text{ to } 0.5 \mu\text{m}$ (b) $0.5 \text{-} 5 \mu\text{m}$	(c)	5-10 μm	(d)	Any of the above
40.	The objective of audit is to				
	(a) Improve the product quality	. ,	Find out the fault		_
	(c) Improve the product value	(d)	Find and process fau	lt an	d to improve
41.	is an alkaloid derived form aliphatic amino a				
	(a) Reserpine (b) Nicotinic acid	(c)	Anabasine	(d)	Vinblastine
42.	The drug sulphan blue is obtained fromsou				
	(a) Plant (b) Animal	(c)	Synthetic	(d)	Mineral
43.	In mammals, The major fat in adipose tissue is:				
	(a) Triglyceride	. ,	Cholesterol		
	(c) Sphingophospholipids	(d)	Phospholipids		



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44.	Dovers powder used as a diaphoretic contains:	
	(a) Ipecac & Opium	(b) Ipecac, Senna & Cinchona
	(c) Opium, Ipecac & Cinchona	(d) All
45.	Biological active form of Vit D in man is:	
	(a) Cholecalciferol	(b) Calcifediol
	(c) Calciferol	(d) Calcitriol
46.	Actions and clinical uses of muscarinic cholinocepto	or agonists include which on of the following
	(a) Bronchodilation (asthama)	
	(b) Improved aqueous humor drainage (glaucoma	
	(c) Decreased gastrointestinal motility (diarrhea)	
	(d) Decreased neuromuscular transmission and rela	xation of skeletal muscle (During surgical anesthesia)
47.	Regarding the role of surfactants in pharmaceutic	al suspensions for oral administration which of the
	following statements is false	
	(a) Surfactants decrease the water contact angle of	dispersed drug particle
	(b) Surfactants promote flocculation	DAT
	(c) Surfactants with high HLB stabilize oral suspen	sions
	(d) Surfactants increase the viscosity of the continu	ious phase of pharmaceutical suspensions
48.	Which of the following drug is NOT used in treatme	nt of H. Pylori infection
	(a) Ampicillin	(b) Clarithromycin
	(c) Mosapride	(d) Bismuth subgallate
49.	The most suitable disinfectant for decontamination	of HIV contaminated endoscope is
	(a) 1% Sodium hypochlorite	(b) 2% Glutaraldehyde
	(c) 5% phenol	(d) 70% ethanol
50.	Which rule does provide the most accurate method	to calculate the dose for child based on adult dose
	(a) Age is months	(b) Age in years
	(c) Weight in pounds	(d) Body surface area
51.	Chemokine co-receptor 5 (CCR 5) inhibitor is	
	(a) Enfuvirtide	(b) Maraviroc
	(c) Raltegravir	(d) Atazanavir
52.	The Franz diffusion cell which is used for the evaluation	tion of transdermal drug delivery systems consists of
	(a) 1 chamber (b) 2 chamber	(c) 3 chamber (d) None
53.	Whick of the following plastic is transparent and fle	xible
	(a) Silicon rubber (b) PVP	(c) HDPED (d) PE
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predetermined point (a) A-B-C method (b) Maximum and minimum method (c) Open-to-buy method (d) Economic order quantity 55. Choose the option with two reducing sugars (a) Lactose and maltose (b) Trehalose and surcrose (c) Maltose and tredhalose (d) Economic order quantity 56. The Local anesthetic with highest cardiotoxicity is (a) Lingocaine (b) Bupivacaine (c) Levo- bupivacaine (d) Procaine 57. Homatropine is (a) Tropine ester of amino acetic acid (b) Tropine ester of mendelic acid (c) Tropine methyl bromide ester of mendelic acid (d) Tropine ester of amino formic acid 58. Tranexamic acid is (a) Antithrombotic (b) Antifibrinolytic (c) Fibrinolytic (d) Styptic 59. Which of the antihistaminic compound has antiadrogenic effect (a) Famotidine (c) Nizatidine (c) Nizatidine (d) Cimetidine 60. Which of the following drug is used prefenntially as preanesthetic mediation (a) Midazolam (b) Oxazepam (c) Alprazolam (d) Alirozepam 61. Proton pump inhibitors are most effective when given (a) Half hour before meals (b) With meal (c) After prolonged fasting (d) Along with H2 blockers 62. Match compounds is Group I with inhibitory activities in Group II Group I (P) Vancomycin (1) Folate metabolism (2) DNA synthesis (3) Procin synthesis (5) Ciprofloxacin (4) RNA synthesis (5) Cell wall systhesis (6) P-4, Q-3, R-1, S-2 (6) P-4, Q-1, R-3, S-2 (7) P-4, Q-1, R-3, S-2 (8) P-5, Q-3, R-2, S-4	54.	In which method an order of a fixed number of ite	ms is placed every time an inventory level falls to a
(c) Open-to-buy method (d) Economic order quantity 55. Choose the option with two reducing sugars (a) Lactose and maltose (b) Trehalose and surcrose (c) Maltose and tredhalose (d) Economic order quantity 56. The Local anesthetic with highest cardiotoxicity is (a) Lingocaine (b) Bupivacaine (c) Levo-bupivacaine (d) Procaine 57. Homatropine is (a) Tropine ester of amino acetic acid (d) Tropine ester of mendelic acid (c) Tropine methyl bromide ester of mendelic acid (d) Tropine ester of amino formic acid 58. Tranexamic acid is (a) Antithrombotic (b) Antifibrinolytic (c) Fibrinolytic (d) Styptic 59. Which of the antihistaminic compound has antiadrogenic effect (a) Famotidine (c) Nizatidine (d) Cimetidine 60. Which of the following drug is used prefemitally as preanesthetic mediation (a) Midazolam (b) Oxazepam (c) Alprazolam (d) Nitrozepam 61. Proton pump inhibitors are most effective when given (a) Half hour before meals (b) With meal (c) After prolonged fasting (d) Along with H2 blockers 62. Match compounds is Group I with inhibitory activities in Group II Group I Group II (P) Vancomycin (1) Folate metabolism (Q) Rifampin (2) DNA synthesis (R) Puromycin (3) Protein synthesis (R) Puromycin (4) RNA synthesis (S) Ciprofloxacin (4) RNA synthesis (S) Cill wall systhesis (a) P-5, Q-4, R-3, S-2		predetermined point	
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(c) Tropine methyl bromide ester of mendelic acid (d) Tropine ester of amino formic acid 58. Tranexamic acid is (a) Antithrombotic (b) Antifibrinolytic (c) Fibrinolytic (d) Styptic 59. Which of the antihistaminic compound has antiadrogenic effect (a) Famotidine (b) Ranitidine (c) Nizatidine (c) Nizatidine (d) Cimetidine 60. Which of the following drug is used prefenntially as preanesthetic mediation (a) Midazolam (b) Oxazepam (c) Alprazolam (d) Nitrozepam 61. Proton pump inhibitors are most effective when given (a) Half hour before meals (b) With meal (c) After prolonged fasting (d) Along with H2 blockers 62. Match compounds is Group I with inhibitory activities in Group II Group I Group II (P) Vancomycin (1) Folate metabolism (Q) Rifampin (2) DNA synthesis (R) Puromycin (3) Protein synthesis (S) Ciprofloxacin (4) RNA synthesis (5) Cell wall systhesis (a) P-5, Q-4, R-3, S-2 (b) P-4, Q-3, R-1, S-2	57.	Homatropine is	
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(a) Antithrombotic (b) Antifibrinolytic (c) Fibrinolytic (d) Styptic 59. Which of the antihistaminic compound has antiadrogenic effect (a) Famotidine (c) Nizatidine (c) Nizatidine (d) Cimetidine 60. Which of the following drug is used prefenntially as preanesthetic mediation (a) Midazolam (b) Oxazepam (c) Alprazolam (d) Nitrozepam 61. Proton pump inhibitors are most effective when given (a) Half hour before meals (b) With meal (c) After prolonged fasting (d) Along with H2 blockers 62. Match compounds is Group I with inhibitory activities in Group II Group I Group I Group II (P) Vancomycin (1) Folate metabolism (Q) Rifampin (2) DNA synthesis (R) Puromycin (3) Protein synthesis (5) Ciprofloxacin (4) RNA synthesis (5) Cell wall systhesis (a) P-5, Q-4, R-3, S-2		(c) Tropine methyl bromide ester of mendelic acid	(d) Tropine ester of amino formic acid
59. Which of the antihistaminic compound has antiadrogenic effect (a) Famotidine (b) Ranitidine (c) Nizatidine 60. Which of the following drug is used preferntially as preanesthetic mediation (a) Midazolam (b) Oxazepam (c) Alprazolam (d) Nitrozepam 61. Proton pump inhibitors are most effective when given (a) Half hour before meals (b) With meal (c) After prolonged fasting (d) Along with H2 blockers 62. Match compounds is Group I with inhibitory activities in Group II Group I (P) Vancomycin (1) Folate metabolism (Q) Rifampin (Q) Rifampin (2) DNA synthesis (R) Puromycin (3) Protein synthesis (5) Ciprofloxacin (4) RNA synthesis (5) Cell wall systhesis (a) P-5, Q-4, R-3, S-2 (b) P-4, Q-3, R-1, S-2	58.	Tranexamic acid is	
(a) Famotidine (c) Nizatidine CEN(d) Cimetidine 60. Which of the following drug is used prefenntially as preanesthetic mediation (a) Midazolam (b) Oxazepam (c) Alprazolam (d) Nitrozepam 61. Proton pump inhibitors are most effective when given (a) Half hour before meals (b) With meal (c) After prolonged fasting (d) Along with H2 blockers 62. Match compounds is Group I with inhibitory activities in Group II Group I (P) Vancomycin (1) Folate metabolism (Q) Rifampin (Q) Rifampin (2) DNA synthesis (R) Puromycin (3) Protein synthesis (5) Ciprofloxacin (4) RNA synthesis (5) Cell wall systhesis (a) P-5, Q-4, R-3, S-2 (b) P-4, Q-3, R-1, S-2		(a) Antithrombotic (b) Antifibrinolytic	(c) Fibrinolytic (d) Styptic
(c) Nizatidine C E N(d) Cimetidine 60. Which of the following drug is used prefenntially as preanesthetic mediation (a) Midazolam (b) Oxazepam (c) Alprazolam (d) Nitrozepam 61. Proton pump inhibitors are most effective when given (a) Half hour before meals (b) With meal (c) After prolonged fasting (d) Along with H2 blockers 62. Match compounds is Group I with inhibitory activities in Group II Group I (P) Vancomycin (1) Folate metabolism (Q) Rifampin (Q) Rifampin (2) DNA synthesis (R) Puromycin (3) Protein synthesis (5) Ciprofloxacin (4) RNA synthesis (5) Cell wall systhesis (a) P-5, Q-4, R-3, S-2 (b) P-4, Q-3, R-1, S-2	59.	Which of the antihistaminic compound has antiadro	genic effect
60. Which of the following drug is used prefenntially as preanesthetic mediation (a) Midazolam (b) Oxazepam (c) Alprazolam (d) Nitrozepam 61. Proton pump inhibitors are most effective when given (a) Half hour before meals (b) With meal (c) After prolonged fasting (d) Along with H2 blockers 62. Match compounds is Group I with inhibitory activities in Group II Group I (P) Vancomycin (1) Folate metabolism (Q) Rifampin (2) DNA synthesis (R) Puromycin (3) Protein synthesis (S) Ciprofloxacin (4) RNA synthesis (5) Cell wall systhesis (a) P-5, Q-4, R-3, S-2 (b) P-4, Q-3, R-1, S-2		(a) Famotidine DISCI	(b) Ranitidine
(a) Midazolam (b) Oxazepam (c) Alprazolam (d) Nitrozepam 61. Proton pump inhibitors are most effective when given (a) Half hour before meals (b) With meal (c) After prolonged fasting (d) Along with H2 blockers 62. Match compounds is Group I with inhibitory activities in Group II Group I (P) Vancomycin (1) Folate metabolism (Q) Rifampin (Q) Rifampin (2) DNA synthesis (R) Puromycin (3) Protein synthesis (5) Ciprofloxacin (4) RNA synthesis (5) Cell wall systhesis (a) P-5, Q-4, R-3, S-2 (b) P-4, Q-3, R-1, S-2		(c) Nizatidine CEN	(d) Cimetidine
(c) Alprazolam (d) Nitrozepam 61. Proton pump inhibitors are most effective when given (a) Half hour before meals (b) With meal (c) After prolonged fasting (d) Along with H2 blockers 62. Match compounds is Group I with inhibitory activities in Group II Group I (P) Vancomycin (1) Folate metabolism (Q) Rifampin (Q) Rifampin (2) DNA synthesis (R) Puromycin (3) Protein synthesis (5) Ciprofloxacin (4) RNA synthesis (5) Cell wall systhesis (6) P-4, Q-3, R-1, S-2	60.	Which of the following drug is used prefenntially as	preanesthetic mediation
61. Proton pump inhibitors are most effective when given (a) Half hour before meals (b) With meal (c) After prolonged fasting (d) Along with H2 blockers 62. Match compounds is Group I with inhibitory activities in Group II Group I (P) Vancomycin (1) Folate metabolism (Q) Rifampin (2) DNA synthesis (R) Puromycin (3) Protein synthesis (5) Ciprofloxacin (4) RNA synthesis (5) Cell wall systhesis (a) P-5, Q-4, R-3, S-2 (b) P-4, Q-3, R-1, S-2		(a) Midazolam	(b) Oxazepam
(a) Half hour before meals (b) With meal (c) After prolonged fasting (d) Along with H2 blockers 62. Match compounds is Group I with inhibitory activities in Group II Group I (P) Vancomycin (1) Folate metabolism (Q) Rifampin (Q) Rifampin (2) DNA synthesis (R) Puromycin (3) Protein synthesis (5) Ciprofloxacin (4) RNA synthesis (5) Cell wall systhesis (a) P-5, Q-4, R-3, S-2 (b) P-4, Q-3, R-1, S-2		(c) Alprazolam	(d) Nitrozepam
(c) After prolonged fasting (d) Along with H2 blockers 62. Match compounds is Group I with inhibitory activities in Group II Group I (P) Vancomycin (Q) Rifampin (Q) Rifampin (Q) Puromycin (3) Protein synthesis (5) Ciprofloxacin (4) RNA synthesis (5) Cell wall systhesis (a) P-5, Q-4, R-3, S-2 (b) P-4, Q-3, R-1, S-2	61.	Proton pump inhibitors are most effective when giv	en
62. Match compounds is Group I with inhibitory activities in Group II Group I (P) Vancomycin (Q) Rifampin (Q) Rifampin (Q) Puromycin (S) Ciprofloxacin (A) RNA synthesis (B) Puromycin (C) DNA synthesis (C) Ciprofloxacin (D) Protein synthesis (E) Cell wall systhesis (E) Cell wall systhesis (E) P-4, Q-3, R-1, S-2		(a) Half hour before meals	(b) With meal
Group I (P) Vancomycin (Q) Rifampin (Q) Rifampin (Q) Puromycin (S) Ciprofloxacin (P) Vancomycin (C) DNA synthesis (D) Protein synthesis (E) Cell wall systhesis (E) Cell wall systhesis (E) P-4, Q-3, R-1, S-2		(c) After prolonged fasting	(d) Along with H2 blockers
(P) Vancomycin (1) Folate metabolism (Q) Rifampin (2) DNA synthesis (3) Protein synthesis (5) Ciprofloxacin (4) RNA synthesis (5) Cell wall systhesis (6) P-4, Q-3, R-1, S-2 (7) P-4, Q-3, R-1, S-2	62.	Match compounds is Group I with inhibitory activities	es in Group II
(Q) Rifampin (2) DNA synthesis (R) Puromycin (3) Protein synthesis (5) Ciprofloxacin (4) RNA synthesis (5) Cell wall systhesis (6) P-4, Q-3, R-1, S-2		Group I	Group II
(R) Puromycin (S) Ciprofloxacin (4) RNA synthesis (5) Cell wall systhesis (a) P-5, Q-4, R-3, S-2 (b) P-4, Q-3, R-1, S-2		(P) Vancomycin	(1) Folate metabolism
(S) Ciprofloxacin (4) RNA synthesis (5) Cell wall systhesis (a) P-5, Q-4, R-3, S-2 (b) P-4, Q-3, R-1, S-2		(Q) Rifampin	(2) DNA synthesis
(5) Cell wall systhesis (a) P-5, Q-4, R-3, S-2 (b) P-4, Q-3, R-1, S-2		(R) Puromycin	(3) Protein synthesis
(a) P-5, Q-4, R-3, S-2 (b) P-4, Q-3, R-1, S-2		(S) Ciprofloxacin	(4) RNA synthesis
			(5) Cell wall systhesis
(c) P-4, Q-1, R-3, S-2 (d) P-5, Q-3, R-2, S-4		(a) P-5, Q-4, R-3, S-2	(b) P-4, Q-3, R-1, S-2
		(c) P-4, Q-1, R-3, S-2	(d) P-5, Q-3, R-2, S-4



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63.	Formation of Okazaki occurs in		
	(a) Transcription	(b) Replication	
	(c) Translation	(d) Reverse Transcription	
64.	Drug used in ventricular arrhythmia is		
	(a) Flecainide	(b) Verapamil	
	(c) Esmolol	(d) Diltazem	
65.	The lipoprotein with the fastest electrophoretic mob	oility and the lowest TG content	is
	(a) VLDL	(b) HDL	
	(c) LDL	(d) Chylomicrons	
66.	As per schedule 'Y' of the drugs and cosmetics act, th	e animal toxicity study requiren	nents for marketing of
	a drug depends upon tentative route and duration of	administration in humans. In Th	nis context, which one
	of the following statements is incorrect		
	(a) Single dose human use-animal toxicity for 2 week	eks in 2 species	
	(b) Oral use for 2 weeks in humans- animal toxicity	for 4 week in 2 species	
	(c) Aerosol use by repeated use in humans- anima	toxicity for 24 weeks in 2 spec	cies
	(d) Multiple daily ocular application for short durat	ion-iregation test in 1 species fo	or 3 weeks
67.	For determining the efficacy of sterilization in an au	oclave, the spores of the followi	ing organism are used
	as test organisms \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \	NTER	
	(a) Bacillus cereus	(b) Clostridium pefringens	
	(c) Bacillus stearothemophilus	(d) Clostridium histolyticum	
68.	Which of the following pairs is mismatched		
	(a) Aerobic, helical bacteria- gram negative	(b) Entrics- gram negative	
	(c) Myconbacteria – acid fase	(d) Pseudomonas –gram posi	itive
69.	List of drugs whose import, manufacture and sale, labe	eling and packaging are governed	d by special provisions
	are included in schedule:		
	(a) X (b) K	(c) H	(d) G
70.	Sigma minus method is used in assessment of		
	(a) Bioavailability	(b) Absorption	
	(c) Metabolism	(d) Tissue distribuation	
71.	Which of the plant family contains volatile oil in their	trichome	
	(a) Rutaceae	(b) Papaveracease	
	(c) Umbelliferare	(d) Laminaceae	

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72.	Ferritin is:		
	(a) Coenzyme	(b) The stored form of Ir	on
	(c) Non-protein moiety	(d) Isoenzyme	
73.	Which oil is solute is alcohol		
	(a) Arachis oil	(b) Sesame oil	
	(c) Castor oil	(d) Corn oil	
74.	One of the first step of the citric acid cycle is isomeri	zation of citric acid to isoctiric	acid this step is necessary
	because		
	(a) Oxidation of secondary alcohols is very difficu	ılt	
	(b) Reduction of secondary alcohol is very impos	sible	
	(c) Reduction of tertiary alcohols would require a	a very powerful oxidizing age	nt
	(d) Oxidation of tertiary alcohols would require of	xidizing agents	
75.	Which of the following alkyl halides would underg	o SN2 reaction most rapidly	
	(a) CH ₃ CH ₂ -BR	(b) CH ₃ CH ₂ -CL	
	(c) CH ₃ CH ₂ -I	(d) CH ₃ CH ₂ -F	
76.	Mechanism of action of Ketoconazole is:		
	(a) Inhibits Ergosterol synthesis DISC	(b) Inhibits DNA gyrase	
	(c) Inhibits dihydropteroate synthetase $\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \$	(d) Induces translation m	isreading
77.	All are adrenal gland over activity disorders EXCE	PT	
	(a) Addison's disease	(b) Conn's syndrome	
	(c) Cushing's syndrome	(d) Cushing's disease	
78.	The oil used in a parenteral product cannot conta		
	(a) WFI (b) Parffin oil	(c) Peanut oil	(d) Glycerine
79.	Identity the non-absorbable sature		
	(a) Catgut suture	(b) Chromic catgut suture	
	(c) Silk suture	(d) Polydioxanone suture	
80.	The relative lowering of vapour pressure is given		
0.4	(a) Raoult's law (b) Henry's law	(c) Boyle's law	(d) Charles law
81.	Identity the functional group present in meprobar		
02	(a) Amide (b) Ester	(c) Carbamic	(d) Lactam
ŏΖ.	Match the following	(4) WAY(00) 40W 0	
	(P) Gypsum salt	(1) KAI(SO4)2 · 12H2O	
	(Q) Epson salt	(2) FeSO ₄ ·7H ₂ O	

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	(R) Alum		(3)	CaSO ₄ ·2H ₂ O	
	(S) Green vitriol		(4)	$MgSO_4 \cdot 7H_2O$	
	(a) P-1, Q-2, R-4, S-3		(b)	P-3, Q-4, R-1, S-2	
	(c) P-4, Q-3, R-1, S-2		(d)	P-2, Q-4, R-1, S-3	
83.	Tinea capitis is ringworm in	fection of			
	(a) Feet (b	o) Groin	(c)	Head	(d) Nails
84.	Rank the following compound	ds is order of increasing re	activ	ity in electrophilic arom	natic Substitution reactions
	(P) C_6H_6 (0	Q) C ₆ H ₅ CH ₃	(R)	$C_6H_5NO_2$	
	(a) Q <p<r (b)<="" th=""><th>o) R<p<q< th=""><th>(c)</th><th>Q<r<p< th=""><th>(d) P<q<r< th=""></q<r<></th></r<p<></th></p<q<></th></p<r>	o) R <p<q< th=""><th>(c)</th><th>Q<r<p< th=""><th>(d) P<q<r< th=""></q<r<></th></r<p<></th></p<q<>	(c)	Q <r<p< th=""><th>(d) P<q<r< th=""></q<r<></th></r<p<>	(d) P <q<r< th=""></q<r<>
85.	Dose dumping may be a gen	neral problem in the form	nulat	ion of:	
	(a) Soft gelatin capsules		(b)	Suppositories	
	(c) Modified release drug p	roducts	(d)	None	
86.	Codeine differ in structure fr	rom morphine by:			
	(a) N-methyl group		(b)	Acetyl group at C1 an	d C6
	(c) -OC2H5 group	Y/ GF	(d)	-OCH3 group	
87.	Isotopes differ in:				
	(a) The number of protons	\		The valency number	
	(c) The chemical activity			The number of neutr	
88.	Arrange the given acids in in		ie nu	imber of carbons pres	ent in them
		Q) Caprylic		Caproic	(D) Lauric
	(a) P <q<r<s< th=""><th></th><th>. ,</th><th>R<q<p<s< th=""><th></th></q<p<s<></th></q<r<s<>		. ,	R <q<p<s< th=""><th></th></q<p<s<>	
	(c) R <q<p<s< th=""><th></th><th>. ,</th><th>Q<p<r<s< th=""><th></th></p<r<s<></th></q<p<s<>		. ,	Q <p<r<s< th=""><th></th></p<r<s<>	
89.	E1cb (Elimination reaction v	via conjugate base) which			ll form?
	(a) Carbocation			Carbanion	
	(c) Free radical		(d)	All	
90.	What are the three basic ste			_	
	(a) Denature, anneal, & Stra	•	. ,	Denature, anneal & e	
	(c) Strand displacement, syn	nthesis & release	(d)	Reverse-Transcription	n, anneal & extend
91.	Rabies bodies are		<i>a</i> >		
	(a) Negri bodies			Cowdry type B inclusion	ion bodies
0.0	(c) Cowdry type A inclusion		. ,	Bollinge bodies	
92.	When glucose reacts with br	-			
	(a) Glucaric acid (b	o) Glucoronic acid	(c)	Sorbitols	(d) Gluconic acid

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PREVIOUS YEAR PAPER



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93. Increase in melting temperature of DNA is due to high content of

(a) A+T

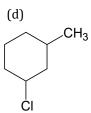
(b) G+C

(c) A+G

(d) T+G

94. What will be the primary product of the following reaction

- (a) CH₃
- (b) CH₃
- (c)



- 95. The mass spectrum of a compound with an approximate MW 137 shows tow equally intense. peaks at m/z 136 and 138. What does the suggest
 - (a) The compound is alkyl iodide

(b) The compound is alkyl bromide

(c) The compound is alkyl chloride

(d) The compound is aryl fluoride

- 96. Capping in tablets mainly due to:
 - (a) Less upper punch pressure
 - (c) Proper formulation design

- (b) Poor flowability of granules
- DISCU(d) Entrapment of air in tablet during compression
- 97. How can we detect the rhizome from the root of the Rauwolfia?
 - (a) By the presence of small central pith
- (b) By the absence of small central pith
- (c) By the presence of vascular bundle
- (d) None
- 98. Drug of choice to treat H1N1 influenza is
 - (a) Adefovir

(b) Cidofovir

(c) Oseltamivir

(d) Tenofovir

- 99. Identify the correct statement
 - (P) Condensed tannins are polymers flavans
 - (Q) Condensed tannins do not contain sugar redidues
 - (R) Hydrolyzabletannis are polymers of gallic acid or ellagic acids
 - (S) Gallic acid and catechin are psedotannins
 - (a) Only Q

(b) P and Q

(c) P, Q and R

- (d) P, Q, R and S
- 100. Quick breaking aerosols are applicable:
 - (a) Orally

(b) Parenterally

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- (c) Topically
- (d) Ophthalmically



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101.	In t	the Reimer-Tiemann reaction	Re	acts wi	th	phenol to give the ortho-formylated produc
	(a)	Carbene		((b)	Carbocation
	(c)	Carbanion		((d)	Free redical
102.	Whi	ich of the following in not added	l to chewing	g tablet		
	(a)	Gildant			(b)	Disintegrant
	(c)	Lubricant			(d)	Anitadhesive
103.	Ran	ge of C=O stretching in enol is				
	(a)	1800 cm ⁻¹			(b)	1710 cm ⁻¹
	(c)	1685 cm ⁻¹		((d)	1655 cm ⁻¹
104.	Mat	ch the following phytochemical	s with their	source	and	d use
	(P)	Shatavrin	(1) Buckw	heat ar	nd c	itrus fruits, strengthens capillary walls
	(Q)	Resvertatrol	(2) Brocco	oli and	cab	bage, protects against bladder cancer
	(R)	Glucosinolates	(3) Purple	grape,	ant	i inflammatory, anticancer
	(S)	Rutin	(4) Aspara	agus, ga	alact	togogue
	(a)	P-4, Q-3, R-2, S-1		TD	(b)	P-4, Q-2, R-3, S-1 P-2, Q-3, R-4, S-1
	(c) l	P-3, Q-1, R-4, S-2		JI	(d)	P-2, Q-3, R-4, S-1
105.	Whi	ich RNA polymerase is the only	\			
	(a)	RNA polymerase I	V C	EN	(b)	RNA polymerase II
	(c)	RNA polymerase III			(d)	RNA pimase
106.	Mat	ch the scientist awarded with N	obel prize	with the	eir c	contributions
	(P)	Alexander Fleming			(1)	GPCR
	(Q)	Kobilka			(2)	β-blocker
	(R)	Banting			(3)	Penicillin
	(S)	Black			(4)	Insulin
	(a)	P-4, Q-3, R-2, S-1			(b)	P-4, Q-2, R-3, S-1
	(c)	P-3, Q-1, R-4, S-2			(d)	P-2, Q-3, R-4, S-1
107.	Mea	an arterial pressure is				
	(a)	Systolic pressure – Diastolic P	ressure		(b)	(Diastolic pressure + Diastolic Prssure)/2
	(c)	Diastotic alcohol + $(1/3) \times pu$	lse pressure	9	(d)	Stroke valume X heart rate
108.	Eug	enol is				
	(a)	Monoterpene alcohol			(b)	Sesquiterpene alcohol
	(c)	Aliphatic alcohol			(d)	Phenylpropene



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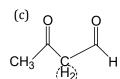
109. The Vitamin required for carboxylation of pyruvate to form oxaloacetate is

- (a) Thiamine
- (b) Biotin
- (c) Pyridoxine
- (d) Niacin

110. Which of the following circled hydrogen is most acidic







111. The drug formulated as suspension follows order reaction

(a) Zero

(b) Pseudo Zero

(c) First

(d) Pseudo first

112. Which diuretic causes decrease in release of insulin

(a) Chlorothiazide

(b) Ethacynic zero

(c) Triamterene

(d) Acetazolamide

113. Match the following drugs with their mode of action

(P) Methotrexate

(1) Mitotic inhibitor

(Q) Cyclophosphamide

(2) Antimetabolite

(R) Vincristine

(3) Alkylating agent

(S) Dactinomycin

DISCU(4) Intercalating agent

(a) P-4, Q-3, R-2, S-1

E N(b) P-2, Q-4, R-1, S-3

(c) P-3, Q-1, R-4, S-2

(d) P-2, Q-3, R-1, S-4

114. Which compound would be expected to show intense IR absorption at 3300 cm⁻¹

(a) $CH_3CH_2CH_2CH_3$

(b) CH₃CH₂C=CH

(c) $CH_3C=CCH_3$

(d) CH₂CHCH₂CH₃

115. In the carbon NMR, in what region of the spectrum does one typically observe carbons which are part of the aromatic ring

(a) -10-0 ppm

(b) 40-60 ppm

(c) 80-100 ppm

(d) 120-150 ppm

116. Meclofenamate belongs to which class of drug

(a) Slicylates

(b) Oxicams

(c) Aryl antaranillic acid

(d) p-Amino phenols

117. Match the following crude with their chemical constituents

(P) Aloe

(1) Hesperidine

(Q) Ginger

(2) Palmitin

(R) Lemon peel

(3) Barbaloin

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			_	
	(S)	Olive oil	(4)	Allin
	(a)	P-4, Q-3, R-2, S-1	(b)	P-3, Q-4, R-2, S-1
	(c)	P-3, Q-4, R-1, S-2	(d)	P-3, Q-1, R-1, S-4
118.	Dop	pamine agonists with tetralene function		
	(a)	Ropinorole	(b)	Pirebidil
	(c)	Pramipixole	(d)	Rotigotine
119.	The	IUPAC name of the compound-(CH ₃) ₂ CHCH ₂ Cl:		
	(a)	2-methyl-3-chloropropane	(b)	1-chloro-3-mehtyl butane
	(c)	1-chloropentane	(d)	2-mehtyl-4-chlorobutane
120.	A po	owerful inhibitor of stomatal opening is		
	(a)	Auxin	(b)	Bytokinin
	(c)	Gibberellin	(d)	Abscisic acid
121.	Wha	at is the renal clearance of a substance, if its conce	entra	ation in plasma is 10mg, concentration in urine is
	100	mg and urine flow is 2 ml/min		
	(a)	0.02 ml/min	(b)	0.2 ml/min
	(c)	2ml/min	(d)	20 ml/min
122.	Aim			SION
	(a)	To monitor drug toxicity $V \subset E $	(b)	To monitor unauthorized drug manufacture
	(c)	To monitor rational use of drugs	(d)	To check and control drug costs
123.	Pha	se zero studies is a/an		
	(P)	Exploratory investigational new drug study		
	(Q)	Human microdosingstudies		
		Step to speed up drug discovery/ development	proc	cess
	(S)	Mandatory by FDA		
	. ,	P and Q (b) Q and R	(c)	P, Q and R (d) P, Q, R and S
124.		enteral product must be:		
	. ,	Packed in bottle		Sterilized
	(c)	Free from viable/living organism	(d)	Pyrogenic

End of paper

125. Quinine present in highest amount in:

(a) C. calisaya

(c) C. ledgeriana

(d) C. succirubra

(b) C. officinalis



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ANSWER KEY GPAT 2014

1-d	2-с	3-b	4-d	5-b	6-b	7-d	8-c	9-b	10-с
11-d	12-d	13-с	14-с	15-b	16-d	17-b	18-b	19-a	20-d
21-a	22-b	23-a	24-a	25-d	26-a	27-a	28-d	29-b	30-b
31-с	32-c	33-с	34-d	35-d	36-с	37-b	38-a	39-a	40-d
41-c	42-c	43-a	44-a	45-d	46-b	47-d	48-с	49-b	50-d
51-b	52-b	53-d	54-d	55-a	56-b	57-b	58-b	59-d	60-a
61-a	62-a	63-b	64-a	65-b	66-d	67-c	68-d	69-a	70-a
71-d	72-b	73-с	74-d	75-c	76-a	77-b	78-b	79-с	80-a
81-с	82-b	83-с	84-a	85-c	86-d	87-d	88-b	89-b	90-b
91-a	92-d	93-d	94-с	95-b	96-d	97-a	98-с	99-d	100-с
101-a	102-b	103-d	104-a	105-b	106-с	107-с	108-d	109-b	110-с
111-a	112-b	113-d	114-b	115-d	116-с	117-с	118-d	119-a	120-d
121-d	122-a	123-с	124-b	125-с					

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