Q.1 - 30 Carry One Mark Each. 1. The structural feature common fro propranolol, atenolol, pindolol, metopropol in the side chain is (A) Isopropylamino propan-2-ol (B) Dimethylamino propan-2-ol (C) Diethylamino propan-2-ol (D) Dibutylamino propan-2-ol 2. When N-methyl group of morphine is replaced with an allyl group, the compound formed is (A) Naloxone-morphine antagonist (B) Natrexone-morphine agonist (C) Nalorphine-morphine antagonist (D) Nalbuphine-morphine agonist/antagonist Nitrazepam can be synthesized from 3. (A) 2-Bromo-5-amino benzophenone (B) 2-Nitro-2-chloro acetophenone (C) 2-Amino-5-nitro cyclohexanone (D) 2-Amino-5-nitro benzophenone 4. Clavulanic acid has a beta lactam ring fused to (A) Thienyl system (B) Thiadiazole system (C) Thiazolidine system (D) Oxazolidine system 5. A drug which has antipyretic, anti-inflammatroy and antiplatelet activity is (A) Sulfinpyrazone (B) Aspirin (C) Ticlopidine (D) Acetaminophen 6. Wild cherry bark contains prunasin which is a (A) Phenolic glycoside (B) Isothiocyanate glycoside (C) Coumarin glycoside (D) Cyanogenetic glycoside 7. Ephedra sinica and Ephedra equisetina can be distinguished by type of (A) Branching (B) Stomata (C) Scaly leaves (D) Alkaloids 8. Microprapagation of the plants is carried out through (A) Cross fertilization (B) Seed germination (C) Plant tissue culture (D) Grafting 9. Acontitine belongs to the group of (A) Steroidal alkaloids (B) Terpenoidal alkaloids (C) Indole alkaloids (D) Ouinoline alkaloids 10. Crude fiber value of a drug is a measure of (A) Soft tissue matter (B) Woody matter (D) Organic matter (C) Mineral matter 11. One of the units used for expressing pressure is 'torr' and it is equal to (A) cm of Hg (B) mm of Hg (C) psi (D) gauss 12. Removal of a single electron from a molecule results in the formation of (A) Fragment ion (B) Metastable ion (C) Molecular ion (D) Rearrangement ion 13. Nuclear magnetic moment is NOT shown by

(C) 1H

(C) KCI

A conductance cell is calibrated by using a solution of known conductivity, i.e.,

(B) Detectability

(D) Reproducibility

(A) 13C

(A) NaCl

(A) Molecular weight

usually a solution of

(C) Reversibility

14.

15.

(B) 16O

Derivatisation techniques in HPLC are intended to enhance

(B) Hg,Cl,

(D) 15N

(D) Na,SO₄

PHARMACOPHORE SOLUTIONS

16.	Metoclopramide is generally used for				
	(A) Prophylaxis of vomiting	(B) Preventing motion sickness			
	(C) Treating irritable bowel syndrome				
	(D) Treatment of pancreatic insufficient	су			
17.	DNA amplification by the polymerase ch	ain reaction uses			
	(A) Thermus aquaticus DNA polymerase	e (B) DNA topoisomerase			
18.	(C) RNA polymerase Identify the non-pathogenic organism	(D) DNA helicase			
	(A) Mycobacterium bovis	(B) Mycobacterium smegmatis			
19.	(C) Mycobacterium avium Bioassays are carried out to	(D) Mycobacterium intracellulare			
	(A) Measure the pharmacological activit	y of a drug			
	(B) Avoid clinical trails for new drugs				
	(C) Detect the impurity in a given drug				
20.	(D) Screen fro pharmacognetic influence A direct way of studying idiosyncratic re				
	(A) changing the route of drug administ	ration			
	(B) changing the assay method				
	(C) pharmacogenomics				
21.	(D) structure activity relationship studieAn example of haemopoietic growth fact				
	(A) platelet derived growth factor	(B) epidermal growth factor			
22.	(C) iron dextran Safranin is used as a reagent to detect	(D) erythropoietin			
	(A) Gram-negative bacteria	(B) Gram-positive bacteria			
(C) Acid fast bacteria (D) Myx 23. Sulphonamides do not have adverse drug interaction		(D) Myxozoa g interaction with			
	(A) Oral anticoagulants				
	(B) Sulfonylurea hypoglycemic agents	(C) Hydantoin anticonvulsants			
24.	(D) Dihydrofolate reductase inhibitors Simvastatin belongs to				
	(A) HMG CoA reductase inhibitor type of antilipidemic agents				
	(B) HMG CoA reductase inhibitor type of	- 10 No.			
	(C) Fibrate type of anticoagulant agents				
25.	(D) Fibrate type of antilipidemic agents HIV infection can be clinically controlled to	with			
26.	(A) Cytarabine (B) Acyclovir (C) Zidovudine (D)Amantadine The measure of cohesive strength of the cross linking that occurs between gelatin molecules and is proportional to the molecular weight of gelatin is called				
	(A) Bloom strength	(B) Viscosity			
27.		(D) Partition coefficient ag material in microencapsulation process			
	is:	(B) Cilianna			
	(A) Polyethylene	(B) Silicone			
28.	One of the following is used as a solubi pharmaceutical liquid dosage forms.	(D) Paraffin ilizing agent to solubilize testosterone in			
	(A) Sucrose monoesters	(B) Lanolin esters			
	(C) Lanolin ethers	(D) Tweens			

29.	One of the following is used a	s a pH depen	dent	controlled rele	ase excipient.
	(A) Carnauba wax				
	(B) Hydroxy proply methyl ce	llulose phtha	late		
	(C) Methyl cellulose				
30.	(D) Glyceryl monostearate The Schedule in D & C act tha	t deals with t	he s	tandards for dis	sinfectant fluids is:
	(A) Schedule B (B) S€h			Schedule O	(D) Schedule M
31.	Q.31 – 80 The carboxyl group of aspirin i	Carry Two			onhanal to got
J1.	(A) 3-Acetamidophenly-O-ace		VICH	v acetyr p arriir	iophenor to get
	(B) 4-Acetamidophenly-O-ace				
	(C) O-(2-hydroxy benzoyl) sa				
	(D) 2-acetamidophenyl-O-ace				
32.	IUPAC system of nomenclature	e for diclophe	enac	sodium (BP) is	
	(A) Sodium 2-[(2, 6-Dichlorop	henyl) amin	o] ph	nenyl acetate	
	(B) Sodium 3-[(2, 6-Dichlorop		_		
	(C) Sodium 2-[(2-Chlorophen	7. N			
33.	(D) Sodium 2-[(6-Chlorophen 1-(2-Aminoethyl) perdydroazo rise to an adrenergic neuron b	ocine on trea	tmer		yl isothiourea give
34.	(A) Bethanidine (B) Med Quercetin is	amylamine	(C)	Guanadrel	(D) Gauenthidine
	(A) 5, 7, 3-Trihydroxy flavone		(B)	5, 7, 3, 4-Tetr	adydroxy flavone
	(C) 3, 5, 7, 3, 4-Pentahydroxy	flavonol			
35.	(D) 3, 5, 7, 3, 4-Pentahydroxy Meconic acid is a chemical mar	flavonone ker for the g	enus	5.	
36.	(A) Piper (B) Piloo A novel diterpenoid isolated fro			Prunus xus brevifolia is	(D)Papaver
37.	(A) Demecolcine (B) Pacli The absorption maximum for p	taxel oolar compou	(C) unds	Vinblastin is usually shift	(D)Brevifolicin ed with change in
	polarity of the solvents due to		,		2
	(A) Hydrogen bonding	i.		Chemical reacti	
38.	 (C) Ionization of the compound A titration in which potential constant value and the current is 	applied acros	ss tv		s maintained at a
	(A) Potentiometric titration		(B)	Amperometric t	titration
39.	(C) Displacement titration The parameter in the elution of compound in gas chromatograp		oropo		titration concentration of a
	(A) Number of peaks			Width of the pe	ak
40.	(C) Area under the peak A drug solution has a half life o order kinetics, how long will it potency?	f 21 days. As	(D) s	Shape of the pe	ak ug undergoes first
	(A) 3.2 days (B) 9.6 d	ays	(C)	16 days	(D)6.4 days
41.	An amphoteric surfactant used	in pharmace	utica	l disperse syste	em is:
	(A) Bile salts		(B)	Lecithin	
	(C) Sorbitan monolaurate		(D)	Sorbitan monos	stearate

42. An abrasive used in dentifrices is (A) Dicalcium phosphate (B) Sodium carboxy methyl cellulose (D) Dioctyl sodium sulfosuccinate (C) Sodium lauryl sulfate 43. An electrochemical method that enhances the transport of some solute molecules by creating a potential gradient through the skin tissue with an applied electrical current or voltage is called (A) Electrophoresis (B) Iontophoresis (C) Osmosis A patient with rheumatoid arthritis has been taking acetyl salicylic acid regularly. 44. However, recently she has been experiencing stiffness, swelling and pain due to salicylate resistance. She has occult blood in her faeces. Suggest an appropriate drug suitable for her from those mentioned below: (A) Paracetamol (B) Celecoxib (C) Piroxicam (D) Naproxen 45. The break down of fibrin is catalyzed by (A) Plasmin (D) Ptylin (B) Renin (C) Urokinase Which one of the these best describes a process carried out to render a drug 46. pharmacokinetically more acceptable? (A) Enteric coating if diclofenac. (B) Co-administration of aspirin with antacids (C) Use of colloidal suspensions or liposomes for administering amphotericin-B. (D) Synthesis of an analogue to obtain high receptor specificity. 47. Azithromycin is clinically administered once daily as compared to erythromycin which is administered every 6 hours because, azithromycin (A) Penetrates into most tissues and is released very slowly. (B) Has a methylated nitrogen in its lactone ring which renders it much more potent than erythromycin. (C) Is a very potent antibiotic but not tolerated well in the gastrointestinal tract. (D) Is usually presented in a sustained release dosage form. 48. A patient showing muscle rigidity, bradykinesia, tremors and postural instability was administered levo-dopa. Which of the properties of levo-dopa is not true? (A) Levo-dopa is preferred over dopamine because it can cross the blood brain barrier. (B) Levo-dopa is the levorotatory stereoisomer of 3, 4-dihydroxy phenylalanine. (C) Levo-dopa gets decarboxylated in the brain to dopamine. (D) Levo-dopa is administered because of its strong antagonistic action on dopamine receptors. 49 Autoimmunity refers to (A) an automatic trigger of the immune system directed against a specific (B) failure to distinguish between self and non-self (C) an automatic segregation of T and B cells. (D) failure of B-cells to interact with T-cells. 50. Which of these is true about the discovery of HB antigen in the blood of people infected with Hepatitis-B? (A) It provided a basis for vaccine design. (B) It indicated that specific vaccines cannot be designed for Hepatitis-B.

(A) Amphetamine

51.

(C) It has not been of much significance.

(D) It indicated that Hepatitis-B is a viral disease

Which drug molecule DOES NOT have phenylethyl amine moiety?

(B) Glyburide

(C) Pheniramine (D) Mescaline

	ons are correct. Ch	oose the correctico	mbination amond	ptions. Two of these J A, B, C and D. on of insulin may be		
	(P) Binding with re	esins.				
		of amio acid residues.				
		nplex of insulin with p				
	(S) Modification of		orotem.			
53.	(A) Q, R The attributes of co	(B) R, S	(C) P, S	(D)P, R		
55.	(P) No tautomeris					
			oric analic form			
	(Q) Exists in equilibrium with its tautomeric enolic form.					
	# # # # # # # # # # # # # # # # # # #	(R) Stable in alkaline solution, destroyed rapidly at neutral or acidic pH.(S) Stable in neutral solution, destroyed in alkaline pH.				
		200	nares es es establica	(D) D D		
54.	(A) R, S Compared to ben biological propertie		(C) Q, R cillin has the fol	(D)P, R lowing advantages in		
	(P) The amino g degradation.	roup renders the	antibiotic resistar	nt to acid catalysed		
	(Q) The spectrum (of acidity is broadene	d.			
	(R) The amino grou	up of renders penicilli	nase resistance to t	the compound.		
55.	(S) The phenolic group renders penicillinase resistance to the compound. (A) P, Q (B) P, R (C) P, S (D)Q, R The identification of propellants in pharamaceutical aerosols is carried out by					
	(P) Gas-chromatog	ıraphy				
	(Q) Tag-open cup a	apparatus				
	(R) Pyknometer					
	(S) IR Spectrophot	ometer				
56.	(A) P, Q Schedule 'H' and S following.	(B) P, S chedule 'S' as per th	(C) Q, R e Drugs and Cosm	(D)R, S etics Act deal with the		
	(P) Prescription drug of R.M.P.	ugs which are requi re	d to be sold by ret	ail only on prescription		
	(Q) Standards for o	cosmetics				
	(R) Biological and	special products				
57.	(S) List of coal tar colours permitted to be used in cosmetics and soaps (A) P, Q (B) P, R (C) Q, S (D)R, S Myristica fragrans Houtt has two of the following characteristics.					
٠,٠	(P) An indeciduous tree, which produces drupaceous, pale yellow fruits.					
	(Q) Each fruit has several round seeds with smooth surface and lignaceous tegument, and the orange red fleshy aril – the mace, is present inside the seed.					
	(R) A deciduous ta	ll tree, which produce	s lignaceous capsu	les.		
		a unique avoid seed niate fleshly aril – the		ument, surrounded by		
58.	(A) Q, R In size exclusion ch	(B) P, R nromatography the st	(C) P, S ationary phases use	(D)Q, S ed are:		
	(P) Alumina (A) P, S	(Q) Dextran (B) Q, R	(R) Agarose (C) Q, S	(S) Styrene (D) P, R		

59.	Group I	Group II	
	Synthetic Drug	Intermediates from which Group I drugs are synthesized	
	(P) Buclizin	(1) Aziridin and thiophosphoryl chloride	
	(Q) Chlorophenesin	(2) 4-Chlorophenol	
	(R) Thiotepa	(3) 4-Chlorobenzhydryl chloride	
Code	(S) Alprazolam	(4) 2-Amino-5-Chloro benzophenone	
(C)	P - 3 Q - 2 R - P - 2 Q - 4 R -	10 mm	
0.	Group I Cardiac Agents	Group II Mechanism of action	
	(P) Digitoxin	(1) Produces negative inotropic effect by blo calcium channels	
	(Q) Dobutamine	(2) Depresses adrenergically enhanced calcium influ through beta receptor blockade	
	(R) Sotalol	(3) Causes elevation of cAMP levels by stimulation of adenylate cyclase.	
	(S) Nicardipine	(4) Inhibits membrane bound sodium potassium ATPas pump.	

61.

(A) P - 4 Q - 3 R - 2 S - 1 (B) P - 3 Q - 4 R - 1 S - 2 (C) P - 4 Q - 2 R - 3 S - 1 (D) P - 4 Q - 3 R - 1 S - 2

	Group I Technique employed	Group II Source of Radiation	
	(P) Visible spectrophotometry	(1) R _f Source transmitter	
	(Q) IR spectrophotometry	(2) Xenon lamp (3) Tungsten lamp	
	(R) NMR spectrophotometry		
Codes:	(S) Fluorescence spectrophotometry	(4) Nernst glower	

(5) Fluorescence	spectrophotometry (4) Nernst glower
- 2 Q - 4 R - 3	S - 1 (B) P - 3 Q - 2 R - 1 S - 4
- 3 Q - 4 R - 1	S - 2 (D) P - 4 Q - 1 R - 3 S - 2
Group I	Group II
Amino acids	Common degradative products that are citric acid cycle intermediates or their precursors.
(P) Aspartic acid	(1) Succinyl CoA
(Q) Arginine	(2) Alpha-Ketoglutarate
(R) Serine	(3) Fumarate
(S) Methionine	(4) Pyruvate
- 3 Q - 2 R - 4	S-1 (B) P-3 Q-1 R-4 S-2
- 1 0 - 2 R - 3	S-4 (D) P-4 Q-2 R-3 S-1
	- 2 Q - 4 R - 3 - 3 Q - 4 R - 1 Group I Amino acids (P) Aspartic acid (Q) Arginine (R) Serine (S) Methionine

63.

Group I	Group II
Tablet defects	Explanation
(P) Picking	(1) A term used to describe the surface material from a tablet that is sticking to and being removed from the tablet's surface by a punch.
(Q) Sticking (2) Term refers to tablet material adhering to the	
(R) Mottling (3) Term refers to an unequal distribution of colou tablet.	
(S) Lamination	(4) Term refers to separation of a tablet into two or more distinct layers.

(A) P - 1 Q - 2 R - 3 S - 4 (B) P - 1 Q - 3 R - 4 S - 2 (C) P - 2 Q - 4 R - 3 S - 1 (D) P - 3 Q - 1 R - 2 S - 4

(0)	Ρ.	-570	4
65.			

Group I	Group II	
Specific chemical test	Phytoconstituents	
(P) Thalleioquin Test	(1) Hyoscyamine	
(Q) Murexide Test	(2) Barbaloin	
(R) Vitali-Morin Test	(3) Quinine	
(S) Modified Borntrager's Test	(4) Theobromine	

Data for Q.66 - 90 are based on the statement/problem. Choose the correct answer for each question from among the options A, B, C and D. Data for questions 66 to 68:

In a formulation development laboratory a tablet is to be formulated. The care tablet has a bad taste and requires physical and chemical protection of the drug from moisture. The tablet should also deliver the drug for local action in the intestine.

- Suggest a suitable method.
- (C) Enteric coating (D) Sub coating
- (A) Sugar coating (B) Film coating (C) Choose the correct coating material to be used. 67.

(B) Acacia

(C) Ethyl cellulose

- (D) Cellulose acetate phthalate
- 68. Choose the correct solvent for the coating material.
- (A) Acetone
- (C) Propylene glycol (D) Glycerine

Data for questions 69 and 70:

Compound A with the formula C_2H_2N shows the following important bands in the IR spectra; (a) 3423 cm⁻¹ (b) 3236 cm⁻¹

- Assign these bands to the important group in the compound A. 69.
- (A) ${\text{-CH}_3}$ (B) ${\text{-NH}_3}$ (C) ${\text{-CN}}$ (D)=C=N_ On treatment with nitrous acid the compound A is converted to B, which shows a 70. strong band at 3430 cm⁻¹. Assign the absorption band for the group formed in the product.
- (A) -OH (B) = $C=N_{-}$ Data for questions 71 to 73:
- (C) -COOH
- (D)-N=N

In the assay of sulfamethoxazole I.P. $(C_{10}H_{11}N_3O_3S)$, 0.2 g of the sample was dissolved in 50 ml of 2M HCl. To this was added 3 g of KBr and the titration was carried out.

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71.	Titration was carri	ed out using		
	(A) NaNO₂ to esti	mate the amino grou	ıp.	
	(B) NaNO ₂ to estimate	nate the sulphonam	ido group.	
	(C) NaOH to estin	nate the amino grou	p.	
	(D) NaOH to estin	nate the sulphonami	do group.	
72.	The end point in the	ne assay was determ	nined by	
	(A) Conductometr		(B) Using an	
72	(C) Potentiometric		(D) Photomet	
73.	sample.	.1 M titrant consume	ed was 7.8 mi, caict	ılate the % purity of the
	(A) 99.70%	(B) 9.97%	(C) 8.87%	(D)98.79%
	for questions 74 a	nd 75:		20.000
				s degraded in presence
74.		formulated in the formulated in the formulated in the formulated in the formula i		
	(A) Preservative		(B) Chelating	-
	(C) Buffer		(D) Tonicity c	
75.		iate filling and packi	. 	
		nber colored ampou with nitrogen and se		of antioxidant, replacing
				d sealing the ampoule.
	S 12	nber colored ampou		
		npoule, sealing and		
	for questions 76	and 77:		
The us 76.	sual adulterants for Clove stalks can b	clove buds are clove e identified by the pr	e stalks and anthoph resence of	ıylli.
	(A) Starch grains			
	(B) Cystoliths			
	(C) Lignified sclere			
77.	(D) Acicular crysta Anthrophylli can b	als of calcium oxalate e identified by the p	e resence of	
	(A) Lignified scler	eids		
	(B) Acicular crysta	als of calcium oxalate	e	
	(C) Cystoliths			
Data	(D) Starch grains for questions 78	to 90.		
Plant	tissue culture of ca	rrot is beina develor	ed in the laborator	y on a semisolid White's
mediu	ım.	W W		 In the state of the broad restrict of the region of the state of the contract of the both of the state of the
78.		essential in the med		
70	(A) NaCl	(B) CoCl ₂	(C) KCl	(D) CaCl ₂
79.	The pH of the med		(C) F C	(D) F 0
	(A) 6.6	(B) 6.0	(C) 5.6	(D)5.0
80.	The tissue growth			
		ed cells suspended i		
		ed cells in clusters d	istributed in the me	dium.
	(C) Differentiated			
	(D) Surface growt	h of undifferentiated	I mace of colle	

	or questions 81 and 82:			
In glue	cose metabolism, name the enzymes catal Conversion of glucose to glucose-6-phosp			
01.	(A) Hexokinase	mate		
	(B) Glucose-6-phosphate dehydrogenase	25		
	(C) Glycogen phosphorylase			
	(D) Glycogen synthase			
82.	Conversion of 2-phosphoglycerate to pho	sphoenol pyruvate.		
	(A) Pyruvate kinase	(B) Phosphoglycerate mutase		
	(C) Phosphoglycerate kinase	(D) Enolase		
	or questions 83 and 84:			
	trexate, Trimethoprim and Pyrimethami ofolate reductase. Yet, they are classified			
83.		methotrexate in its therapeutic category		
	(A) Trimethoprim binds to bacterial DH compared to the host DHFR.	FR about 50,000 times more strongly as		
	(B) Trimethoprim can be administered o	rally.		
	(C) Trimethoprim exhibits no significant	adverse effects.		
84.	(D) Trimethoprim has additional anti-infl Methotrexate is thought to exert its action			
	(A) Interfering with purine synthese.			
	(B) Intracellular formation of an amine a	adducts.		
	(C) Forming a conjugate with nucleic acids.			
	(D) Inhibiting the synthesis of folic acid.			
	for questions 85 to 87:	proceure gostrie edidity and dishetes is		
	ribed famotidine, enalapril and tolbutamid	pressure, gastric acidity and diabetes is e.		
85.		drugs, predict which will be ionized in the		
	(A) Famotidine	(B) Enalapril		
86.	(C) Tolbutamide The patient cannot tolerate enalapril. W	(D) Enalapril and tolbutamide hich of the following can be substituted?		
	(A) Omeprazole (B) Losartan	(C) Rosiglitazone (D)Clofibrate		
87.	Famotidine acts as			
	(A) H ₁ – histamine antagonist	(B) H ₂ – histamine antagonist		
2-Met acety meth Ester	le-6-methoxy naphthalene. This is conv oxy-2-naphthyl acetic acid, which is ester	rified with methanol to the methyl ester. hthoxy-2-naphthyl)-propionic acid methyl nd).		
:		90 The final compound 7 is:		

(A) Morpholine/Sulphur followed by H2SO4/H2O

(A) Naphazoline (B) Carprofen (C) Pranoprofen (D) Naproxen

(B) Morphine/Sulphur followed by HCl/H2O

(C) Pranoprofen

(C) Formic acid/Cu followed by acetic acid

(D) Hydroiodic acid followed by $\rm H_2SO_4/H_2O$ Identify the reagents – Y. 89.

(A) NaOH/CH₃OH

(B) NaH/CH₃I

(C) Hydrazine/CH3I

(D) LiAlH₄/CH₃OH

Best of Luck www.gpatindia.com