

**GATE - 2003** 

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#### PHARMACEUTICAL SCIENCE

Time: 3 hours Maximum Marks: 150

### Read the following instruction carefully.

- 1. This question paper contains 90 objective questions. Q. 1-30 carry 1 mark each and Q. 31-90 carry two marks each.
- 2. Answer all the questions.
- 3. Questions must be answered on special machine gradable Objective Response Sheet (ORS) by darken-ing the appropriate bubble (marked A, B, C, D) using HB pencil against the question number on the left hand side of the ORS. Each question has only one correct answer. In case you wish to change an answer, erase the old answer completely using a good soft eraser.
- 4. There will be NEGATIVE marking. For each wrong answer, 0.25 mark for Q. 1-30 and 0.5 mark for Q. 31-90 will be deducted. More than one answer marked against a question will be deemed as an incorrect response and will be negative marked.
- 5. Write your registration number, name and name of the Centre at the specified locations on the right half of the ORS.
- 6. Using HB pencil, darken the appropriate bubble under each digit of your registration number.
- 7. Using HB pencil, darken the appropriate bubble under the letters corresponding to your paper code.
- 8. No charts or tables are provided in the examination hall.
- 9. Use the blank pages given at the end of the question paper for rough work.
- 10. This question paper contains 20 pages. Please report, if there is any discrepancy.

## (Q. 1 - 30) CARRY ONE MARK EACH

- 1. Colchicine is biogenetically derived from one of the following
  - (a) Tyrosine and Phenylalanine

(b) Tryptophan and phenylalanine

(c) Ornithine and Tryptophan

- (d) Ornithine and phenylalanine
- 2. The diagnostic character for the microscopically identification of Kurchi bark is
  - (a) Fibers with Y-shaped pits

(b) Horse shoe shaped stone cells

(c) Steroids containing calcium oxalate crystals

(d) Stratified cork

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3.	It is possible to initiate the development of complete p	lants from callus cellCultures by suitable manipulatior		
	of the medium with respect to			
	(a) Minerals	(b) Vitamins		
	(c) Carbohydrates	(d) Hormones		
4.	Polyploidy is defined as			
	(a) Addition of one chromosome	(b) Multification of entire chromosome set		
	(c) Submicroscopic change in DNA material	(d) Gross structural change		
5.	The starting material for the synthesis of ALPRAZOL	AM is		
	(a) 3-amino-5-bromoactophenone	(b) 2-amino-5-chloroactophenone		
	(c) 2-amino-5-chlorobenzophenone	(d) 3-amino-5-chlorobenzophenone		
6.	Simplification of Morphinan system gave one BENZO	OMORPHAN derivative		
	(a) Pentazocin	(b) Pethidine		
	(c) Levorphanol	(d) Buprenorphine		
7.	A metabolite of SPIRONOLACTONE is			
	(a) Aldosterone	(b) Canrenone		
	(c) Corticosterone	(d) Pregnenolone		
8.	The IUPAC name for NAPROXEN is DISCU	JSSION		
	(a) (S)-2-(6-ethoxy-2-naphthyl)-acetic acid $\mathbb{C} \ \mathbb{E} \ \mathbb{N}$	(b) (S)-2-(6-methoxy-2-naphthyl)-aceticacid		
	(c) (S)-2-(6-ethoxy-2-naphthyl)-propionic acid	(d) (S)-2-(6-methoxy-2-naphthyl)-propionic acid		
9.	The metabolic function of Riboflavin involves the following	lowing		
	(a) FMN and FAD	(b) NADP and NADPH		
	(c) AMP and ATP	(d) Retin and Retinine		
10.	X-ray spectral lines Ká doublet arises from transition	of electrons from		
	(a) M shell to K shell	(b) L shell to K shell		
	(c) L shell to M shell	(d) M shell to K shell		
11.	The method of expressing magnetic field strength			
	(a) Cycles/sec (b) Pulses/sec	(c) Debye units (d) Gauss		
12.	A solvent used in NMR			
	(a) Chloroform	(b) Acetone		
	(c) Carbon tetrachloride	(d) Methanol		
13.	A widely accepted detector electrode for pH measure $% \left( \mathbf{p}\right) =\left( \mathbf{p}\right) $	ment is		
	(a) Platinum wire	(b) Glass electrode		
	(c) Ag-AgCl electrode	(d) Lanthanum fluoride		
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14.	Commercial production of citric acid is carried	out by the microbial culture of			
	(a) Fusarium moniliformi	(b) Rhizopus nigrican			
	(c) Aspergillus Niger	(d) Candida utilis			
15.	For thermophilic micro-organisms, the minimu	ım growth temperature required is			
	(a) 20°C (b) 37°C	(c) 45°C (d) 65°C			
16.	Obligatory anaerobes				
	(a) Can tolerate oxygen and grow better in its	presence			
	(b) Do not tolerate oxygen and die in its presen	nce			
	(c) Can grow in oxygen levels below normal				
	(d) Can grow in presence of atmospheric oxyg	gen			
17.	Plasmid is a				
	(a) Macromolecule involved in the protein synthesis				
	(b) Circular piece of duplex DNA				
	(c) A hybrid DNA that is formed by joining pieces of DNA				
	(d) Endogenous substances ecreted by one type	e of cell			
18.	Lactose intolerance is because of the lack of				
	(a) Acid phosphates DISCU(b) Lactate dehydrogenase				
	(c) Galactose-1-phosphate-uridyl transferase	E (d) Amylase			
19.	Synthesis of UREA takes place exclusive in				
	(a) Kidney	(b) Liver			
	(c) Gall bladder	(d) Urinary bladder			
20.	A term which describes a cofactor that is finally	bound to an enzyme			
	(a) Holoenzyme	(b) Prosthetic			
	(c) Coenzyme	(d) Transferase			
21.	How many parts of 10 % ointment be mixed with	ith 2 parts of 15 % ointment to get 12% ointment			
	(a) 2 (b) 3	(c) 5 (d) 6			
22.	The correct non-ionic surfactant used as a pene	etration enhancer in the preparation of mucoadhasives			
	(a) Oleic acid	(b) Tween-80			
	(c) Glycerol	(d) Propylene glycol			
23.	One of the ex-officio member of the Pharmacy (	Council of India is			
	(a) Director General of Health Services	(b) Government Analyst			
	(c) Registrar of theState Pharmacy Council	(d) Director General of veterinary Research Institut			

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24.	The Schedule in Drugs and Cosmetics Act that deals with the requirements and guidelines for clinical trials				
	import and manufacture of new drugs is				
	(a) Schedule '0'	(b) Schedule 'M'	(c)	Schedule 'F'	(d) Schedule 'Y'
25.	A retardant material that	forms a hydrophilic matri	x in t	he formulation of matr	rix tablets is
	(a) H.P.M.C	(b) C.A.P	(c)	Polyethylene	(d) Carnauba wax
26.	A drug which causes pink	to brownish skin pigmen	tation	within a weeks of the	initiation of the therapy is
	(a) Itraconazole	(b) Clofazimine	(c)	Lomefloxacin	(d) Neomycin
27.	The risk of Digitalis toxici	ty is significantly increase	d by o	concomitant administr	ation of
	(a) Triamterene		(b)	Lidocaine	
	(c) Captopril		(d)	Hydrochlorothiazide	
28.	An agent used in Prinzme	etal angina has spasmolyti	c acti	on which increasesco	conary blood supply is
	(a) Nitroglycerine		(b)	Nifedipine	
	(c) Timolol		(d)	Isosorbide mononitr	ate
29.	An organism which has b	een implicated as a possil	ole ca	use of chronic gastritis	andpeptic ulcer is
	(a) Campylobacter Jejuni		(b)	Escherichia Coli	
	(c) Helicobacter pylori		(d)	Giardia lambia	
30.	A 5HT <sub>1D</sub> receptor agonist	useful in migraine is S	US	SION	
	(a) Sumatriptan	(b) Ketanserin C E	<b>N</b> (c)	Ergotamine	(d) Methysergide
		(Q.31-90) CARRY T	'WO	MARK EACH	
31.	At present, different spec	ies of Papaver such as <i>P.</i> (	Orient	<i>ale</i> are being cultivate	d instead of <i>P. somniferum</i>
	because they contain	•		G	
	(a) More of morphine	(b) Less of morphine	(c)	Only codeine	(d) Only thebaine
32.	Guggulipid, a resin is	•		•	
	(a) A hypolipidemic agen	t obtained from cotton plai	nts co	ntaining multifunction	al compound (±) Gossypol
	(b) A lipid obtained from	Arctium lappa, Asteraces	e trac	litionally used for the	treatment of dermatoses
		obtained from Ipmoea ori		-	
	(d) A hypolipidemic ager	rt obtained from Commip	hora	mukul consisting of m	nixture of sterols including
	Z-pregna-(20)-diene-	_		_	
33.	In nitrofuantion synthesis		cetate	is treated with one of	the following intermediate
	in presence of CH <sub>2</sub> COOH+	•			G
	(a) Hydantoin	Z 4 Z Z	(b)	1-5-diamino hydanto	in
	(c) 1-3-diamino hydanto	in	• •	1-amino-hydantoin	
	- J		(-)	<b>y</b> 2 22 <del>22</del>	

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34.	4-hydroxy-3-hydroxymet	thyl benzaldehyde is treated	with acetic anhydride and	l then kept with other solvent,
	t-butyl cyanide and acetic	acid for ten days. Resulting	g compound is reduced w	ith LiAIH <sub>4</sub> in tetra hydrofuran.
	The final product is			
	(a) Isoprenaline	(b) Dobutamine	(c) Salbutamol	(d) Orciprenaline
35.	2-iminothiazolidine is tr	eated with phenyl oxirane	to get a drug used in rou	indworm infection
	(a) Piperazine	(b) Tetramisole	(c) Thiabendazole	(d) Levamisole
36.	Thiamine hydrochloride	on treatment with alkaline	potassium ferricyanide g	gives
	(a) Thymochrome with	fluorescence	(b) Oxythiamine with	golden yellow color
	(c) Neopyrithiamine wi	th orange yellow color	(d) Thiochrome with	blue fluorescence
37.	A new drug delivery syst	em which is composed of p	phospholipids that spont	aneously form a multiamellar,
	concentric bilayer vesicle	es with layers of aqueous m	nedia separating the lipid	l layers is
	(a) Prodrugs	(b) Liposomes	(c) Osmotic pumps	(d) Nanoparticles
38.	Unless otherwise stated	in the individual monograp	ph of the pharmacopeia,	in the disintegration test for
	enteric coated tablets, fir	st the dissolution is carried	d out in	
	(a) 0.1 MHCI	(b) Phosphate buffer	(c) Water	(d) $0.1  MH_2 SO_4$
39.	What us the proportion	of NaCl required to render	r a 1.5% solution of drug	g isotonic with blood plasma?
	The freezing point of 1%	$6~\mathrm{w/v}$ solution of drug is $0$	,122 <sup>0</sup> C and that of NaCl i	is -0.576 <sup>0</sup> C
	(a) 0.65%	(b) 0.585% C E	$N(c)\Gamma 0.9\% R$	(d) 0.5%
40.	IR Spectra appear as dip	s in the curve rather than	maxima as in UV-Visible	spectra because it is a plot of
	(a) % Absorbance again	nst wave no.	(b) % Transmittance a	against concentration
	(c) % Absorbance again	nst Concentration	(d) % Transmittance a	against wave no
41.	ESR is applied to only the	ose substances showing par	ra magnetism which is du	e to the magnetic moments of
	(a) Neutrons	(b) Protons	(c) Paired electrons	(d) Unpaired electron
42.	Rotation of electrons about	ut the proton generates a se	econdary magnetic field w	which may oppose the applied
	magnetic field. The proto	on is then said to be		
	(a) Shielded	(b) Shifted	(c) Hydrogen	(d) Deshielded
43.	The analyte is used in th	e form of a solution flame	photometry because it sh	nould undergo
	(a) Evaporation	(b) Condensation	(c) Nebulization	(d) Precipitation
44.	The mechanism of antip	arasitic action of Mebenda	zole and thiabendazole in	nvolves
	(a) Stimulation of acetyl	choline receptors at neuro	muscular junctions	
	(b) Inhibition of dihydro	opolate reductase		
	(c) Interference with m	icrotubule synthesis and as	ssembly	
	(d) Block thiamine trans	sport		
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#### 45. Isoniazid is a primary anti-tubercular agent that

- (a) Requires pyridoxine supplementation
- (b) Causes ocular complication that are reversible if the drug is discontinued
- (c) Is ototoxic and nephrotoxic
- (d) Should never be used due to its hepatotoxicity potential

#### 46. Decreased risk of Atherosclerosis is associated with increase in

(a) Very-low-density lipoproteins

(b) Low-density lipoproteins

(c) Cholesterol

(d) High-density lipoproteins

#### 47. The mechanism of action of Paclitaxel is

- (a) Bing to DNA through intercalation between specific bases and block the synthesis of new RNA or DNA, cause DNA strand scission
- (b) Mitotic spindle poison through the enhancement of tubulin polymerization
- (c) Competitive partial agonist-inhibitor of estrogen and binds to estrogen receptors
- (d) S-Phase specific antimetabolite that is converted by deoxy kinase to the 5'-mononucleotide

### 48. Lycopodium spore method can be used to find out percentage purity of crude drug which contain

- (a) Multi-layered tissues or cells
- (b) Well defined particles which can be counted CUSSION
- (c) Oil globules
- (d) Characteristic particles of irregular thickness the length of which can measured

### 49. The microscopical character flower buds of Eugenia caryophyllus is

(a) Collenchymatous parenchyma containing in its outer part numerous ellipsoidal schizolysigenous oil glands

CENTER

- (b) Small translucent endosperm containing aleurone grains
- (c) Wide parenchymatous starchy cortex, the endosperm containing volatile oil
- (d) Outer surface consisting of external perisperm, rough, dark brown with reticulate furrows

### 50. In protein blosynthesis, each amino acid

- (a) Recognises its own codon by a direct interaction with the m-RNA template
- (b) Is added in its proper place to a growing peptide chain through "adaptor" function of t-RNA
- (c) Is first attached to an anti codon specific for the amino acid
- (d) Undergoes fidelity translation which is assured by the presence of traces of DNA on the ribosome

#### 51. Rabies Antiserum I. P. is

- (a) A freeze dried preparation containing antitoxic
- (b) A preparation containing specific globulin or its derivatives obtained by purification of hyper immune

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serum or plasma of healthy horses

- (c) A sterile preparation containing antitoxic globulin
- (d) A sterile preparation containing antioxic globulin obtained by purification of hyper immune serum of horses

Q. 52-58 are multiple selection items. P, Q, R, S are the options. Two of these options are correct. Choose

the	corr	ect combination fron	n among the alt	ternatives A, B,	C and D.		
52.	2. Total ash value in case of crude drug signifies						
	(P)	Organic content of th	e drug				
(Q) Mineral matter in the drug							
	(R)	Addition of extraneou	ıs matter such a	s stand stone e	tc		
	(S)	Woody matters prese	ent in the drug				
	(a)	R, S	(b) Q, R	(c)	P, Q	(d)	P, S
53.	The	compounds listed be	low contain $\alpha$ , $\beta$	and η electrons	S		
	(P)	Acetaldehyde	N	(Q)	Butadiene		
	(R)	Formaldehyde			Benzene		
	(a)	R,S	(b) Q,R	DISCI (c)	P,R	(d)	P,S
54.	A 60	0 year old patient pres	sents with glauce	oma. Therapy s	hould include		
	(P)	Topical atropine	4	(Q)	Topical pilocarp	oine	
	(R)	Oral acetazolamide		(S)	Oral pilocarpine	e	
	(a)	P,Q	(b) Q,R	(c)	R,S	(d)	P,S
55.	Mea	asurement of particle s	size in pharmace	eutical Aerosols	is by		
	(P)	Cascade impactor		(Q)	Light scatter de	cay	
	(R)	Karl-Fischer method		(S)	IR spectrophoto	ometry	
	(a)	P,Q	(b) Q,R	(c)	R,S	(d)	P,S
56.	The	common attributes of	f ascorbic acid, a	n antiscorbutio	vitamin, are		
	(P)	Exit in nature in both	reduced and ox	xidized form an	d in a state of rev	versible eq	uilibrium
	(Q)	Has keto-enol system	in the molecule				
	(R)	Has an aldehyde grou	p since it gives j	positive Shiff's	reaction		
	<b>(S)</b>	Salt forming properti	ies are due to th	e presence of t	he free carboxyl	group	
	(a)	P,R	(b) Q,R	(c)	R,S		(d) Q,S



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#### 57. Two properties of Radiopharmaceuticals are

- (P) Slow localization in target issue
- (Q) Very long half-life to minimize radiation exposure yet long enough to get imaging information
- (R) Short half-life to minimize radiation exposure yet long enough to get imaging information
- (S) Rapid localization in target tissue and quick clearance from non-target organs
- (a) P,Q

(b) Q,R

(c) R,S

(d) P,S

#### 58. Two correct statements concerning vitamin D are

- (P) The active molecule 1,25-dihydroxy cholecalciferol binds to intracellular receptor proteins
- (Q) Cholecalciferol is found in vegetables
- (R) 1,25-dihydroxy-D<sub>3</sub> is the potent vitamin D metabolite
- (S) It is required in the diet of individuals exposed to sunlight
- (a) P,S

(b) P,R

(c) R,S

(d) P,Q

## Q. 59-65 are "Matching" exercises. Match Group I with Group II. Choose the correct combination from

among the alternatives A,B,C and D,

### 59. Group I (Tablet Additives)

- (P) Binder
- (Q) Insoluble lubricant
- (R) Film coating material
- (S) Direct compression diluents
- (a) 2-P, Q-1, 3-R, 4-S
- (c) 4-P, 3-Q, 2-R, 1-S

### 60. Group I (IR Detectors)

- (P) Themocouple
- (Q) Pyroelectric Detector
- (R) Golay cells
- (S) Thermistor
- (a) P-4, Q-2, R-3, S-1
- (c) P-1, Q-3, R-2, S-4

### 61. Group I (Alkaloid)

- (P) Coniine
- (Q) Papaverine
- (R) Anabasine

## Group II (Examples)

#### DISCUIS SAcacia

## C E N2. TLight mineral oil

- 3. Hydroxy ethyl cellulose
- 4. Microcrystalline cellulose
- (b) 3-P, 2-Q, 1-R, 4-S
- (d) 1-P, 4-Q, 3-R, 2-S

### Group II (Composition)

- 1. Oxides of Mn, Co and Ni
- 2. Bi-Sb
- 3. Xenon
- 4. Triglycine sulphate
- (b) P-3, Q-1, R-4, S-2
- (d) P-2, Q-4, R-3, S-1

## Group II (Ring system)

- 1. Isoquinoline
- 2. Pyridine-Piperdine
- 3. Yohimbane



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- (S) Reserpine
- (a) P-2, Q-3, R-1, S-4
- (c) P-4, Q-1, R-2, S-3
- 62. Group I (Immunoglobulins[Ig])
  - (P) IgG
  - (Q) IgA
  - (R) IgM
  - (S) IgE
  - (a) P-4, Q-3, R-2, S-1
  - (c) P-2, Q-3, R-4, S-1
- 63. Group I (Antibiotics)
  - (P) Streptomycin
  - (Q) Erythromycin
  - (R) Gentamycin
  - (S) Tetracycline
  - (a) P-4, Q-3, R-1, S-2
  - (c) P-3, Q-2, R-3, S-4
- 64. Group I (Synthetic estrogenic drug)
  - (P) Ethinyl Estradiol
  - (Q) Dienoestrol
  - (R) Chlorotrainisine
  - (S) Stilboestrol
  - (a) P-4, Q-3, R-1, S-2
  - (c) P-1, Q-4, R-2, S-3

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

#### Group II (Actions)

- 1. Agglutination and cytolysis
- 2. Antiallergic
- 3. Neutralises toxins
- 4. Antimicrobial
- (b) P-3, Q-4, R-1, S-2
- (d) P-2, Q-1, R-4, S-3

#### Group II (Microrganism used in the I.P. assay)

- 1. Bacillus cereus
- 2. Staphylococcus
- 3. Klebsiella pneumoniac
- 4. Micrococcus luteus
- (b) P-3, Q-4, R-2, S-1
- DISCI (d) P-3, Q-4, R-1, S-2
- C E Group H (Methods of synthesis)
  - 4'4, Dimethoxy of benxophenone is treated with 4 - methoxy benzoly chloride + Mg, resulting product is treated with PTS followed by Cl<sub>2</sub>+CCl<sub>4</sub>
  - Deoxy anisoin is alkylated and product subjected to Grignard reaction, the resulting tertiary alcohol is dehydrated and demethylated with alcoholic KOH
  - reduction 3. By pinacol of p-hydroxy propiophenone and subsequent removal of water
  - 4. From Estrone by the action of Potassium acetylide
  - (b) P-4, Q-1, R-3, S-2
  - (d) P-3, Q-1, R-4, S-2

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#### 65. Group I (Immunosuppressants)

- (P) Azathioprine
- (Q) Tacrolimus
- (R) Glucocorticoids
- (S) Cyclophosphamide
- (a) P-3, Q-2, R-1, S-4
- (c) P-2, Q-1, R-3, S-4

#### Group II (Mechanism of action)

- 1. Destroys proliferating lymphoid cells
- 2. Prodrug transformed to mercaptopurine which on further conversion inhibits purine metabolism
- 3. Inhibits the cytoplasmic phosphatase Calcineurin
- 4. Interferes with the cell cycle of activated lymphoid cells
- (b) P-2, Q-3, R-4, S-1
- (d) P-4, Q-3, R-2, S-1

Data for Q. 66-90 are based on the statement/problem. Choose the correct answer for each question from the option A,B,C,D.

## Data for (Q.66 - 68)

Leaves of Digitalis Purpurea were subjected to morphological, microscopical and chemical screening

- 66. Morphological character with respect to the leaf is
  - (a) Ovate lanceolate with entire margin
- (b) Ovate lanceotlate with crenate margin
- (c) Linear laceolate with serrate margin (d) Linear laceolate with sinuate margin
- 67. Morphological character with respect to the leaf is
  - (a) Ovate lanceolate with entire margin
- (b) Ovate lanceolate with crenate margin
- (c) Linear lanceolate with serrate margin
- (d) Linear lanceolate with sinuate margin

- 68. The drug gives positive
  - (a) Borntrager's test

(b) Murexide test

(c) Legal's test

(d) Thaleoquin test

### Data for (Q.69-70)

In a synthetic procedure -chloro-2,4 diamino sulfomyl aniline is treated with P to obtain 7-amino sulfomyl 6-chloro-3-chloro-methyl-2H-1,2,4-benzothiadiazin-1:1 dioxide. Subsequently it is refluxed with  $C_6H_5$ -CH $_2$ -SH+NaOH+DMF to yield Y

- 69. Select the reagent **P** 
  - (a) Chloroacetyldehyde

(b) Formaldehyde

(c) Formic acid

(d) Acetaldehyde

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#### 70. The final product **Y** is

- (a) 3-benzyl methyl-6-chloro-2H-1, 2, 4-benzothiadiazine-7-sulphonamide-1, 1-dioxide
- (b) 3-benzyl thiomethyl-6-chloro-2H-1, 2, 4-benzothiadiazine-7-sulphonamide1, 1-dioxide
- (c) 3-benzyl thiomethyl-5-chloro-2H-1, 2, 3-benzothiadiazine-7-sulphonamide1, 1-dioxide
- (d) 3-benzyl thiomethyl-6-chloro-2H-1, 2, 4-benzothiadiazine-7-sulphonamide1, 1-dioxide

## Data for (Q.71-73)

Proguanil is synthesized by diazotization of p-chloroaniline and treating with dicynamide to yield p-chlorophenyl dicyandiamide which is converted to proguanil by reaction with an aliphatic amine. Proguanil is metabolized to a triazine derivative which is an active metabolite.

#### 71. What is the reagent used for diazotization

(a) NaNO<sub>2</sub> + dilute HCl

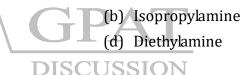
(b)  $KNO_3 + dilute H_2SO_4$ 

(c)  $Zn + dilute H_2SO_4$ 

(d)  $Tin + H_2SO_4$ 

#### 72. Name the aliphatic amine used

- (a) Dimethylamine
- (c) Isobutylamine



#### 73. Name the metabolite

- (a) Thioguanil
- (c) Cycloguanil

# C E N(b) Diguanil

(d) P-chlorphenyl biguanide

### Data for (0.74-76)

Calculate the  $\lambda$  max for the following compounds. Base value for Benzaldehydein ethanol is 250nm.

- 74.  $\lambda$  max of p-bromobenzaldehyde in nm is
  - (a) 265

(b) 255

(c) 275

(d) 260

- 75.  $\lambda$  max of p-hydroxy benzaldehyde in nm is
  - (a) 253

(b) 275

(c) 261

(d) 270

- 76. λ max of o-chlorobenzaldehyde in nm is
  - (a) 275

(b) 265

(c) 255

(d) 250

### Data for (Q.77-78)

In the assay of Folic acid I.P., a weighed quantity is dissolved in 0.1 M NaOH solution and subsequently treated with Zn and HCl. The resulting product is mixed with ammonium sulphate, kept for 2 minutes and a reagent is added to get final colored product whose absorbance is measured.

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77.	Select the product obta	ined when folic acid is heat	ed witl	h Zn + HCl		
	(a) Benzoic acid	(b) P-aminobenzoic	(c)	Glutamic	(d) Succini	c acid
78.	Select the reagent used	d for the development of col	or			
	(a) N-1-napthyl ethyle	ene diamine dihydrochloride	(b)	Ninhydrin reagent		
	(c) P-dimethyl amino	benzaldehyde	(d)	Phloroglucinol		
		Data for (	( <b>Q.7</b> 9	-80)		
		non neurological moven and tremors at rest. Both L		_		y of skeletai
79.	Carbidopa is used beca	ause				
	(a) It crosses blood by	rain barrier				
	(b) It inhibits aromatic	c L-amino acid decarboxylas	se			
	(c) It inhibits MAO typ	ie A				
	(d) It inhibits MAO typ	ре В				
80.	Select the specific unw	anted effect of L-DOPA				
	(a) Dementia	(b) Hypertension	(c)	Dyskinesia	(d) Excito	toxicity
		Data for (	Q.81-	82) N		
m)		CF	NI T	FD		mi
	•	ug in aqueous acid solution	-	-		_
		to be 0.056 M. The concer		on after a period of	12 hours wo	is 4.10×10 <sup>-2</sup>
moi	es/liter. The reaction r	ate constant is 0.02599 hr	-1 <b>,</b>			
81.	What is the quantity of	f drug remaining undecomp	osed a	fter 8 hours.		
	(a) 0.455 moles/liter		(b)	0.25 moles/liter		
	(c) 0.0455 moles/liter	•	(d)	0.10 moles/liter		
82.	What is the amount of	drug deteriorated during th	ıe peri	od of 24 hours.		
	(a) 0.026 moles/liter		(b)	0.0026 moles/liter		
	(c) 0.03 moles/liter		(d)	0.053 moles/liter		
		Data for (	Q.83-	85)		
	formulation developn itamin A and water.	nent laboratory, you have	to for	mulate an oral dosc	age form con	ntaining olive
83.	Suggest a suitable dosa	ge form				
	(a) Solution	(b) Suspension	(c)	Emulsion	(d) Capsul	le
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84.	Suggest a substance to be incorporated into the formulation			
	(a) Glycerin	(b) Acacia	(c) Cetrimide	(d) Alcohol
85.	Select one of the appr	ropriate labeling directio	ns	
	(a) Keep in the refri	gerator	(b) No-preserv	atives
	(c) Schedule 'G'		(d) Shake well	before use
		Data f	or (Q.86-87)	

Successive solvent extraction of a crude drug with petroleum ether, benzene, chloroform, ethyl alcohol and water performed. Qualitative chemical testing of petroleum ether extract gave positive keller-killani and salkowski's reaction. Ethyl alcohol and aqueous extract gave positive FeCl, reaction and aqueous axtract gave foamy solution.

	*	•
	(a) Plant sterols	(b) Tropane slkaloids
	(c) Sesquiterpenoids	(d) Purines
87.	What constituents are present in the ethyl a	cohol and aqueous extracts?
	(a) Plant lipids	(b) Anthraquinone glycosides
	(c) Alkaloids	(d) Plant phenols and saponins

86. What constituents are present in the petroleum ether/benzene extract?

A business executive while playing tennis complained of chest pain and was brought to emergency room. He has history of mild hypertension and elevates bood cholesterol. ECG changes confirmed the diagnosis of myocardial infarction. The decision is made to open his occluded artery by using thrombolytic agent and also use aspirin later.

Data for (Q.88-90)

88. The thrombolytic agent used is (a) Heparin (b) Warfarin (c) Anistreptase (d) Vit. K 89. Mechanism of action of aspirin is (a) Inhibit vitamin K absorption (b) Antithrombin activity (c) Inhabit metabolism of heparin (d) Inhibit platelet aggregation 90. Mechanism of action of antithrombic agent is (a) Conversion of plasminogen to plasmin (b) Activation of clotting factors (d) Agonist of vitamin K (c) Inhibit platelet aggregation

End of paper

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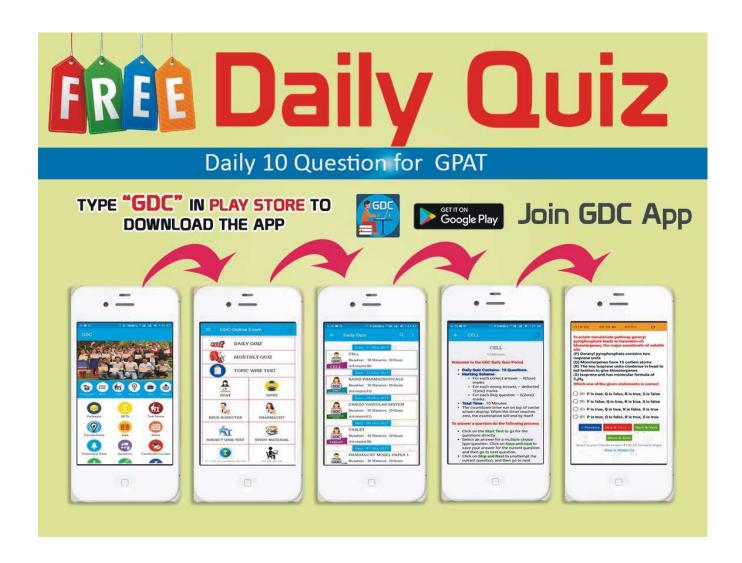
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1- a	2 – b	3 – d	4 – b	5 – b	6 – a
7 – b	8 – d	9 – a	10 – b	11 – d	12 – c
13 – с	14 – с	15 – d	16 – b	17 – c	18 – b
19 – b	20 – c	21 – b	22 – d	23 – a	24 – d
25 – a	26 – b	27 – d	28 – d	29 – c	30 – a
31 – d	32 – d	33 – d	34 – c	35 – a	36 – a
37 – b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

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