QP Code: 90284

SET 2 QUESTION PAPER

Subject: Pharmacognosy & Phytochemistry I

Year and Sem: S.Y. B.Pharm (SEM-IV) (Choice Based) (R-2019)

Duration: 3 hours **Total marks: 80**

N.B.: 1. All questions are compulsory

2. Figures to right indicate full marks

Q. I	Choose appropriate option for following multiple choice based	20M
	questions:	
1	Artificial invert sugar, an adulterant for honey is detected by	88.8
a	Selivonanoff's test	
b	Fiehe's test	2 X 12 C
c	Ninhydrine test	
d	Fehling test	X TO
		2
2	Position of Plant in taxonomy and chemical nature of drugs is included in which of the following system of classification	
a	Taxonomical Classification	
b	Chemotaxonomical Classification	
С	Chemical classification	
d	Serotaxonomical classification	
3	Total Ash value in case of crude drug signifies	
a	Organic content of the drug	
b	Cellular content of the drug	
С	Inorganic content of the drug	
d	Phytoconstituents of the drug	
4	Palisade ratio is	
a	Total number of palisade cells beneath each epidermal cell	
b &	Total number of Palisade cells beneath mesophyll	
C M	Average number of Palisade cells beneath each epidermal cell	
d	Average number of Palisade cells beneath four continuous epidermal cells	
525		
55	A change of the DNA sequence within a gene or a chromosome of an organism resulting in the creation of a new character or trait not found in the parental type	
V a	Chemodemes	
b	Hybridization	
o e T	Polyploidy	
of do	Mutation	
7966	12, 2, 4, 4, 6, 6, 6, 8, 8, 8, 8, 8, 8, 8, 8, 8, 8, 8, 8, 8,	
6	The natural plant growth regulator which promote cell division, cell elongation and useful in root formation, phototropism, geotropism and apical dominance	
a	Cytokinins	
b b	Abscisic acid	
c	Auxins	

d	Gibberellins	SO A A
7		
7	The method of collection of gum from the plant	0,400
a	felling	X1. 20 (2. V
b	cutting	60 (X, X-)
d d	tapping	2 2 2 6 6 C
u	digging	
8	Following are the methods of in -situ conservation except	
a	National Park	\$ 5° 5° 5°
b	Botanical Garden	0000
С	Statuary	X 33.00
d	Biosphere reserve	
		X 40
9	Cytokinins are derivatives of	Z.
a	Proteins)
b	Adenine	
c	Glycosides	
d	Indole PROPERTY OF THE PROPERT	
	LE LA LE CALLE CALLES CONTRACTOR	
10	Following are the macronutrient; except	
a	Nitrogen Nitrogen	
b	Sulphur	
c	Copper	
d	Potassium	
11	The unorganized mass of cells which proliferates from the cells of an explant is termed as	
a	Callus culture	
b	Protoplast culture	
C &	Suspension culture	
d	Culture	
25/5/2		
12	The system of medicine which put forth the Laws of Similars which says that like cures like (Similae similibus curentur)	
a	Unani V P P P P P P P	
(b) (c)	Homeopathy	
V CON	Ayurveda	
T'd T	Siddha	
13	Identify the class of compound from the given basic nucleus	
	N H	
$\hat{\beta}$	Indole Alkaloids	
b	Anthraquinone Glycosides	

C	Quinoline Alkaloids	5.22 A X 4
c d	Cardiac Glycosides	10.00 A. V.
u	Cardiac Grycosides	4 8 8 8 0 V
14	Identify the class of compound from the given basic nucleus	
a	Isoflavonoids	DO T
b	Neo Flavonoids	TX.
c	Flavonoids)*
d	Flavan-3-ol	
	\$\$\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\	
15	C-Anthraquinone glycosides are confirmed by which of the following tests	
a	Borntrager's test	
b	Modified Borntrager's test	
С	Keller Killiani test	
d	Shinoda test	
16	Home is all property of Charles to the place.	
16	Hemp is an example of fibre belonging to the class	
a b	Regenerated fibre Mineral fibre	
	Carbohydrate fibre	
c d	Protein fibre	
u	1 Totem Hote	
17	are agents that can cause a birth defect by permanently altering the structure and functions of organs exposed to them	
a	Teratogens	
\$ b 8	Allergens	
50C 4	Hallucinogens	
Sy rd Co	Enzymes	
18	The proteolytic enzyme derived from the bacteria present in the gut of silk	
Z X X E	worm S S S S S S S S S S S S S S S S S S S	
o a	Urokinase	
3, 8, p , 2,	Streptokinase	
2,30°	Serratiopeptidase	
12 d 23	Pepsin	
26, 74 LO.		
19	Acacia and tragacanth are examples of natural gums classified in specific	
	group	
a	Seed gums	
\$\frac{1}{2} \b \frac{1}{2} \cdot \frac{1}{2}	Marine gums	

С	Exudate gums	30°77
d	Microbial Gums	6 6 6
		720
20	Gelatin is the example of source of drug of natural origin which is obtained	TO OF
	from the following	O YL
a	Vegetable source	1. 65° 65° (
b	Animal source	× 60 × 60
c	Mineral source	100 m
d	Marine Source	
u	Triume bource	
Q. II A	Answer any one of the following.	12M
1a	Draw the nucleus, example, use and chemical test of the following-	6M
1 a	Draw the nucleus, example, use and enemical test of the following-	301
	i. Tropane alkaloid	
	ii. Cardiac glycosides	KIKE
	iii. Triterpenoidal pentacyclic saponin	5 ⁷
	in. Therpenoidal pentacyche saponin	
1 ե	Define Ash value and availing Index 2 White Days as an army attention of	61/
1b	Define Ash value and swelling Index? Write a note on wax obtained from	6M
	animal source.	
		(3.7
2a	Draw the nucleus, example, use and chemical test of the following	6M
	i. Anthraquinone Glycosides	
	ii. Steroidal saponin	
	iii. Isoquinoline alkaloids	
2b	Define and write the significance of moisture content as physical parameter	6M
	in the evaluation of Drugs of Natural origin.	
	Give complete pharmacognostic account of any one oil having cathartic	
	property	
Q.II B	Answer any four out of the five	48 M
1. 29	The little of the same of the same of	
1a	Explain the role of Pharmacognosy in Allopathy and Ayurveda system of	6 M
- 10 No.	medicine with examples.	
1b	Write in detail about any three factors influencing cultivation of medicinal	6 M
	plants with one example each.	
2000	5 5 7 7 5 5 6 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8	
2a	Discuss transgenic plants and their use in the production of edible vaccines.	6 M
100 m	Give the advantages of edible vaccine.	
2b	Classify following with examples:	6 M
ENTER!	i. Volatile oil	
O A AN	ii. Tannins	
SO TO ST		
-3a	With the help of suitable examples differentiate between organized and	6 M
0,230	unorganized drugs. Write a note on Agar.	
3b	Write the source, preparation, constituents, chemical tests and uses of	6 M
COOK! AT	'Acacia'.	0 141
97 75 79 V	A A A A A A A A A A A A A A A A A A A	
W 97 4 9	Define Polymoidy and Hybridization	2 1/
4a	Define Polyploidy and Hybridization.	3 M
4b	Differentiate between Callus and Suspension culture.	3M

4c	Compare and contrast Absorbent and Non-absorbent cotton.	
4d	Write about Sero-taxonomical classification of Drugs of natural origin with	3M
	suitable examples.	2 2 20 6
		X 20 7 20
5a	Write a note on	6 M
	i. Conservation of medicinal plants.	1 1 2 0 0 V
	ii. Novel medicinal agents from marine source.	
5b	Explain any two proteolytic enzymes obtained from plant source.	6 M

Q.P.Cods:90422

Subject: Pharmacology I (Theory) Year and Sem: S. Y. B. Pharm. Sem IV Rev. 2019 **Duration: 3 Hrs** Total marks: 80 N.B.: 1. All questions are compulsory 2. Figures to right indicate full marks Q. 1. Choose appropriate option for following multiple choice based 20 marks questions. 1. The phenomenon in which the action of one drug is abolished by the other is known as a. Antagonism **b.** Synergism c. Dose-response relationship **d.** Desensitization 2. The theoretical volume of plasma from which the drug is completely removed in unit time signifies _____ of a drug. **a.** Absorption **b.** Metabolism c. Volume of distribution **d.** Clearance 3. Which of the following effect can be seen in competitive antagonism in a drug-response curve? a. Non-parallel left shift **b.** Non-parallel right shift c. Parallel right shift d. Parallel left shift 4. Idiosyncrasy is a. Type A ADRs **b.** Type B ADRs c. Type C ADRs d. Type D ADRs **5.** Latanoprost is used in the treatment of a. Myasthenia gravis **b.** Glaucoma c. Alzheimer's disease **d.** Epilepsy **6.** An example of surface anaesthetic is ______. a. Prilocaine **b.** Bupivacaine c. Benzocaine

d. Chloroprocaine

7.	The most unwanted stage of anaesthesia which can be escaped with
	newer anaesthetic agents.
a.	Stage of analgesia
b.	Stage of delirium
С.	Medullary paralysis
d.	Surgical anaesthesia
8.	The drug useful in alcohol abstinence is
a.	Disulfiram
b.	Propranolol
c.	Atropine
d.	Tubocurarine
9.	Nootropic drugs are
a.	CNS depressants
b.	Anxiolytics
c.	Cognition enhancers
d.	Antiepileptic drugs
10.	The morphine exerts effect
a.	Miosis
b.	Increased motility
С.	Respiratory stimulation
d.	Algesia
11.	Low volume of distribution indicates that the drug is distributed in the:
a.	Vascular compartment
b.	Extracellular fluid
c.	Intracellular fluid
d.	Tissues
12.	Which of the following is a G protein coupled receptor?
	Muscarinic cholinergic receptor
a. b.	Nicotinic cholinergic receptor
Č.	Glucocorticoid receptor
d.	Insulin receptor
	VIRSUIH ECEPTOR
13.	Which of the following statement is correct for nasal decongestants?
) a.	Safer in hypertensives
b.	Do not produce any systemic effects
) C.C.	They are alpha antagonists
d.	Cause impairment of mucosal cilary function

14.	The treatment involved in barbiturate poisoning include
a.	Flumazenil
b.	Urine alkalization
c. d.	Pralidoxime Atropine
u.	Attopine
15.	Monoamine oxidase B subtype cause oxidation of
a.	Histamine
b.	Hydroxytryptamine
c.	Adrenaline
d.	Phenylethylamine
16.	Atypical antipsychotics are preferred over typical antipsychotics mainly because
a.	Atypical antipsychotics are potent dopamine blockers
b.	Atypical antipsychotics are specific dopamine receptor blocking
c.	Excreted unchanged in the urine
d.	Minimal extrapyramidal side effect
17.	Glutathione conjugation detoxifies which of the following drug?
a.	Proguanil
b.	Acetazolamide
c.	Paracetamol 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5
d.	Dopamine
18.	A partial agonist can antagonize the effects of a full agonist because it has
a.	High affinity but low intrinsic activity
b.	Low affinity but high intrinsic activity
c.	No affinity and low intrinsic activity
d.	High affinity but no intrinsic activity
19.	Dry, flushed and hot skin, dilated pupil, photophobia, dry mouth, excitement, convulsions and coma are the manifestations of
a.	Organophosphate poisoning
b.	Morphine poisoning
	Belladonna poisoning
d.	Heavy metal poisoning
20.	The drawback of nitrous oxide as anesthetic agent is
 a.	It may lead to diffusion hypoxia
b.	It has hangover effect
€.	It is highly explosive
d.	Incompatibility with other anesthetic agents

Q. 2 A Answer ANY ONE question.

12 marks

- **a** Define metabolism. Enlist various Phase I and Phase II reactions. Add a note on enzyme induction and inhibition.
- **b** Classify anti-epileptics. Give the mechanism of action and adverse effects of Phenytoin and Valproic acid.

Q. 2 B Answer ANY FOUR questions.

48 marks

- **a. i.** Define absorption. Add a note on factors affecting absorption.
- **a. ii** Give the advantages and disadvantages of the oral route.
- **b. i.** Classify the receptors along with the examples. Explain in brief ion channel receptors.
- **b. ii.** Define clinical trials, enlist their various phases and write a note on preclinical studies.
 - **c** What are sympatholytics? Classify them and add a note on the treatment of Glaucoma.
 - **d** Give mechanism of action and anyone therapeutic use of the following drugs: Thiopental, Disulfiram, Ketamine, and Baclofen.
- **e. i.** Write a detailed note on Psychostimulants.
- e. ii. Explain the pharmacology of Levodopa.

Subject: Medicinal Chemistry-I Year and Sem: S.Y. B.Pharm.(SEM-IV)

Duration: 3 hours Total marks: 80 M

N.B.: 1. All questions are compulsory

2. Figures to right indicate full marks

Q. 1	Choose the appropriate option for following multiple choice-based questions.	M
	Each question carries one mark.	Z.Y
1	The type of metabolic reaction which occurs in the following biotransformation is	37
	H ₃ C ₋	/
	N N N N N N N N N N N N N N N N N N N	
	[a]Oxidation at benzylic carbon [b]Oxidation of Aromatic ring	
	[c]Oxidation of C -S system [d] S-demethylation	
	Which of the fallowing statement is incorrect the standard of the set of the set	
2	Which of the following statement is incorrect about metabolism of drugs [a] Metabolism is also called a detoxification process	
	[b] Phase I and Phase II reactions are metabolism pathways	
	[c] Phase II reactions are also called as functionalization reactions	
	[d] Cytochrome enzymes play an important role in the metabolism of drugs	
	[u] eyecemente enzymes play an importante reclassionary of drugs	
3	Which of the following is a selective α -1 receptor agonist?	
	$\begin{array}{ c c c c c c c c c c c c c c c c c c c$	
4	Which drug contains a 4-amino- 6,7-dimethoxyquinazoline ring system attached to an acyl piperazine moei	ty?
	[a] Tolazoline [b] Phentolamine [c]Phenoxy-benzamine [d] Prazosin	
5	What is the name of this cholinergic drug? $H_2N \longrightarrow 0 \longrightarrow N \longrightarrow CI^{\Theta}$	
30,00	[a]Bethanechol chloride [b]Carbachol chloride [c]Methacholine chloride [d] Acetylcholine chloride	

6 Which drug is synthesised using phenyl acetonitrile and 1,5-dibromopentane as precursors?

[a] Cyclopentolate [b]Tacrine [c] Neostigmine [d] Dicyclomine

Select the INCORRECT statement with respect to the SAR of adrenergic agonists with specific reference to 3',5'-dihydroxy ring substitution pattern.

[a] Increases the drug distribution

[b] Increases resistance to metabolism by COMT

[c] Provides selectivity for β2-receptors

[d] Gives orally active bronchodilator

8	Following are structural requirements essential for sympathomimetic activity of arylethanolamines EXCEPT?
	[a] (1S)-OH [b] Catechol ring [c] β-phenylethylamine [d] (1R)-OH
	\$\times\$\times\text{\$\times\times\text{\$\times\times\text{\$\times\times\text{\$\times\times\text{\$\times\times\times\text{\$\times\times\text{\$\times\times\times\text{\$\times\times\times\times\times\text{\$\times\time
9	Identify the triazole ring fused benzodiazepine from the following.
	[a] Chlordiazepoxide [b] Diazepam [c] Oxazepam [d] Alprazolam
10	The benzodiazepine analog which has the least sedative activity
	[a] ortho-substituted 5-aryl benzodiazepine [b] di-ortho-substituted 5-aryl benzodiazepine
	[c]para-substituted 5-aryl benzodiazepine [d]unsubstituted 5-aryl benzodiazepine
11	Droperidol is a member of class of antipsychotic agents.
	[a]Phenothiazine [b] Butyrophenone [c] Benzazepine [d] Benzoisoxazole
12	
12	The spacer group present between the ring nitrogen and the side chain amino nitrogen in phenothiazines
	for optimum antipsychotic activity is
	[a]Butyl [b] Methyl [c] Ethyl [d] Propyl
13	Identify the name of ring present in phonytain from the following
13	Identify the name of ring present in phenytoin from the following [a]Succinimide [b] Oxazolidinedione [c]Hydantoin [d]Iminostilbene
	[a]succinimide [b] Oxazolidinedione [c]Hydantoin [d]iminostiibene
14	Which of the following phenothiazine derivatives contains piperidine side chain.
	[a] Thioridazine [b] Prochlorperazine [c] Triflupromazine [d] Chlorpromazine
	[a] Thoritazine [b] Frochio perazine [c] minapionazine (a) enorpromazine
15	Which of the following is structural isomer of Enflurane
	[a] Isoflurane [b] Sevoflurane [c] Methoxyflurane [d]Desflurane
16	Which of the following is not an example of Inhalation anaesthetics
	[a] Halothane [b] Enflurane [c] Ketamine [d] Sevoflurane
17	Which of the following is INCORRECT statement about Methadone
	[a] Methadone is a synthetic opioid
	[b] R-enantiomer is more potent than S enantiomer
	[c] Methadone is opioid antagonist
	[d] N-demethylation is major metabolic pathway for Methadone
18	Which of the following is not a structural feature of Opioid Antagonist
	[a] Presence of allyl/cyclopropyl methyl group at 17th position
200	[b] Replacement of 6-OH with keto group
	[c] Presence of 7-8 double bond
	[d]Substitution of 14 OH
37 50 C	
19	The isosteric replacement of the indole ring with the Indene ring system resulted in which of the following
	anti-inflammatory drug
37.65	[a] Sulindac [b] Diclofenac [c] Tolmetin [d]Naproxen
47 9 S	Identify the given apticipatement on agent
20	Identify the given anti-inflammatory agent
STATE OF THE PARTY	
162	
3000	OH OH
6,60	[a] Piroxicam [b] Tolmetin [c] Phenacetin [d] Mefenamic acid

Q.2 A	nswer <u>any one</u> of the following two questions.	20°25	12M
Α	(I) State whether following statements are true or false in relation to the compounds (structure	6M	82,00
	drawn below) active as antimuscarinic agents. If false, correct the statement and justify.		
	Support your answer with relevant structures.	\$ 50 G	
	$ \begin{array}{c c} R_1 \\ R_2 & X - (CH_2)n - N \\ R_3 & R_6 \end{array} $		
	 Substitution of R₂ and R₃ by naphthalene ring increases the anticholinergic activity. Introduction of hydroxyl group at R₁ increases the anticholinergic activity. Compound belong to the amino alcohol ether class if X = -COO- and R₁ = -OH. 		
	(II) Outline the synthesis of Salbutamol along with reaction conditions and necessary reagents and give its mechanism of action.	4M	
	(III) Phenoxybenzamine and Prazosin are two α -adrenergic antagonists. Is their mechanism of action the same? Explain.	2M	
В	(I) Answer the following questions	6M	
	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$		
	a b c d		
	 Indicate the chemical classes of 'a' and 'b' Predict the effect of attaching a methyl group on both the ring nitrogens of 'a' Write the mechanism of action of 'c' Predict the effect of replacing the ring methyl group of 'b' by H Name the enzymes involved in the metabolism of 'd' 		
	(II) Explain the basis of GI side effects, generally caused by the non-selective class of NSAIDs.	2M	
	(III) (1) Give two examples of Narcotic antagonists with structure.(2) Give two examples of flexible opioid agonists with structure.	4M	
N. SO. S.			

	Answer <u>any four</u> of the following five questions. (I) Match the following						121	
		Drugs		Column A	Т	Column B	6M	200
	1	Clonidine	a	Metabolized to α-methyl NE	i	2-Arylimidazoline	X 2000	
	2	Naphazoline	b	Contains resorcinol nucleus	ii	Non-catecholamine β2- selective agonist		266
	3	Methyldopa	С	Indirect acting adrenergic agonist	iii	Phenylethylamine		200
	4	Terbutaline	d	Contains naphthalene ring	iv	2-aminoimidazoline		
	5	Isoproterenol	е	Presence of o-chlorine groups and NH bridge	v	Phenyl propanolamine	8 0 V	
	6	Pseudoephedrine	f	Non-selective β-agonist	vi	Catecholamine with isopropyl N-substituent		
	(2	1) Resistance to CON	ΛT	tions in sympathomimetics besto (2) Resistance to MAO (3) h respect to the structures given	Ora	lactivity	3M	
	но	OH H	\	но М	но	\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\		
	но	(a)	97,907	HO (b)	HO	(c)		
3	1. 2. 3.	Which of the above Predict the effect of Arrange the above by MAO.	f iso _l		ctur	e C. pect to rate of metabolism	6M	121
	1. 2. 3. (I) De	Which of the above Predict the effect of Arrange the above of by MAO.	f iso struc stor	(b) Ictures is a selective β2-agonist? propyl group on selectivity in structures in the increasing order with	ctur 1 res	e C. pect to rate of metabolism	6M 4M	121
	1. 2. 3. (I) De	Which of the above Predict the effect of Arrange the above shy MAO. Escribe biosynthesis, explain the effect of fecture drawn below).	stor	(b) Inctures is a selective β2-agonist? Propyl group on selectivity in structures in the increasing order with rage, release and metabolism of a wing structural changes on the activity.	cety cety H ₃	e C. pect to rate of metabolism		121
	1. 2. 3. (II) Ex (structure) 1. 2. 3.	Which of the above Predict the effect of Arrange the above by MAO. Escribe biosynthesis, eplain the effect of fecture drawn below). Replacement of ace Replacement of all Replacement of ace	storud storud storud storud etyl g	(b) actures is a selective β2-agonist? propyl group on selectivity in structures in the increasing order with rage, release and metabolism of a wing structural changes on the acture of β	ictur i res icety tivity	e C. pect to rate of metabolism Icholine. of muscarinic agonist CH ₃ CH ₃ gen with -C ₂ H ₅		121

С	(I) Answer the following questions. Support your answer with relevant structures wherever required	5M	12M
	 Protein binding can prolong the duration of action. Explain 'Geometrical isomerism influences biological activity'. Explain with suitable examples. Enlist Phase I reductive metabolic reactions Explain the concept of bioisosterism with suitable examples Give an example of 'hydrolysis' as biotransformation pathway. 	3M	
	[II] Elaborate on factors affecting drug metabolism		
	[III] Write the structure of any two Phase I metabolites of following	4M	
	$\begin{array}{cccccccccccccccccccccccccccccccccccc$		
D	(I) Classify antipsychotic drugs based on their chemical structures with at least one example from each class. (Structures needed)	4M	12M
	(II) Outline the synthetic scheme of chlorpromazine indicating the reagents and reaction conditions used.(III) Compare the antipsychotic activity and side effect profile of chlorpromazine with	4M	
	prochlorperazine.	4M	
E	(I) Explain why morphine has poor oral bioavailability. Discuss the structure activity relationship of morphine analogues with suitable examples.	6M	12M
	(II) Classify the following drugs into various subclasses of NSAIDS and give their structures and mechanism of action Indomethacin, Diclofenac, Aspirin, Acetaminophen, Antipyrine, Ketorolac	6M	

Q.P. Code:90507

Subject: Physical Pharmaceutics- II Class: S. Y. B. Pharm. (Sem.-IV) R-2019 Duration: 3 Hrs. Maximum Marks: 80

N.B.: 1. All questions are compulsory

2. Figures to right indicate full marks

Q. I	Choose the appropriate option for the following multiple choice	20M
1.	based questions. Dilatant flow is characterized as a reverse phenomenon of:	
1.		
	b. Plastic flow	
	c. Pseudoplastic flow	
	d. Rheopexy	
2.	A plot of shear rate, as a function of shear stress is called	
	a Rheogram	
	b Standard Plot	
	c Humidity Chart	
	d Histogram	
3.	Brook-field viscometer is an example ofviscometer.	
	a. Cone and plate	
	b. Extrusion	
	c. Rotating sphere	
	d. Rotating spindle	
4.	During elastic deformation, the stress–strain relationship for a specimen is described by	
	a. Hooke's law	
	b. Boyle's law	
	c. Beer Lambert's law	
	d. Charle's law	
5. 35	A deformation that recover after the release of stress is known as	
	a plastic deformation	
957.45	b elastic deformation	
(2) (2)	c pseudoplastic deformation	
3 6 7 5	d creep	
62	The ratio of void volume to bulk volume is known as	
	a. Porosity	
	b. Tapped density	
	c. Granule volume	
	d. Bulk Density	
7.088	Helium pycnometer is used to determine	
	a. Size	
	b. True density	
	c. Sedimentation rate	
	d. Surface area	
8.	The powder having low bulk density or large bulk volume is known as	
	a. Bulk powder b. Heavy powder	
	e. Light powder	
2.2.14.0	of Eight pondor	

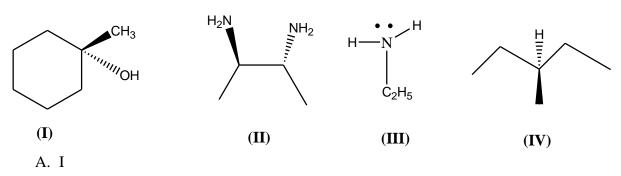
9.	Which of the following is the half-life of First order reaction?
	a. t1/2 =2k
	b. $t1/2 = A0/2k$
	c. $t1/2 = 0.693/2k$
10	d. t1/2 =0.693/k
10.	Climate zone III is
	a. Hot/dry climate
	b. Subtropical and Mediterranean climate
	c. Hot/humid climate
11	d. Moderate climate
11.	The dielectric constant is used to measure
	a. Spreadability of the solventb. Polarity of the solvent
	c. Viscosity of the solvent
	d. Temperature of the solvent
	d. Temperature of the solvent
12.	is the reaction of compounds and molecular oxygen
	a. Photolysis
	b. Hydrolysis
	c. Auto-Oxidation
	d. Thermolysis
13.	The type of emulsion can be easily identified using the following test except
	test. 3 3 4 5 5 5 6 5 5 7 5 5 5 6 6 6 6 6 6 6 6 6 6
	a. Dye solubility
	b. Creaming
	c. Dilution
	d. Redispersibility
14.	As the viscosity of the emulsion is the flocculation of globules will
	be reduced.
	a. Increased
6,65	b. Decreased
\$ 5.74 \$ 5.74	c. Maintained zero
12, 60° E	d. Lowered
15.	In an emulsion, the relative volume of water and oil is expressed as
	a. Phase ratio
	b. Phase volume ratio
	c. Phase inversion
	d. Viscosity
16.	is an example of hydrophilic colloid used in preparation of an
	emulsion
	\$\text{\$\ext{\$\text{\$\exitinx{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\}}}}\text{\$\}\$\text{\$\text{\$\text{\$\text{\$\text{\$\text{\$\tex{

	b. Spans	
	c. Bentonite	
	d. Veegum	
17.	surfactants do not impart charges on interfacial films.	20 20 70 20 75 76
17.		
	a. Ionic	
	b. Non ionic	
	c. Cationic	
18.	d. Anionic Donnan membrane effect means:	
	a. Driving the drug ion of similar charge to the opposite side of the semipermeable membrane	
	b. Driving the drug ion of opposite charge to the opposite side of the semipermeable membrane	6
	c. Driving the drug ion of neutral charge to the opposite side of the semipermeable membrane	
	d. Stopping the transfer of drug ion of similar charge to the opposite side of the semipermeable membrane.	
19.	Which of the following is an example of lyophilic colloid? a. Gold	
	b. Silver	
	c. Sulphur	
20.	d. Albumin Lyophobic colloids are:	
20.	a. Easy to prepare and thermodynamically stable	
	b. Easy to prepare but thermodynamically unstable	
	c. Difficult to prepare but thermodynamically stable	
20,20	d. Difficult to prepare and thermodynamically unstable	
Q. II A)	Answer any one question. Explain the optical properties of colloids in detail	12M
a. b.	Classify viscometers. Describe the principle, construction and working of cup	
	and bob viscometer.	
Q. II B)	Answer any four questions.	48M
l. a.	Describe types of particle deformation.	6M
b.	Describe the mechanical behaviour of solids in terms of elastic modulus.	6M
2. a.	What do you understand by particles packaging arrangements in powders? How is powder porosity evaluated?	6M
b .	What are the methods used for determining particle size? Explain in detail any two.	6M
3. a.		6M
10 C C C C C C C C C C C C C C C C C C C	State Stoke's law and its significance in sedimentation of suspension	6M

4. a.	Discuss the various factors influencing particle settling in suspension	6M
b.	Discuss the various identification tests used to differentiate the type of emulsion	6M
5. a.	What are the limitations of Arrhenius equation for determination of accelerated stability studies?	6M
b.	The half-life of drug which decomposes according first order kinetics is 75 days. Calculate shelf life and k.	6M

S. Y. B. PHARM. SEM-IV-POC-III-MCQ-FH2022

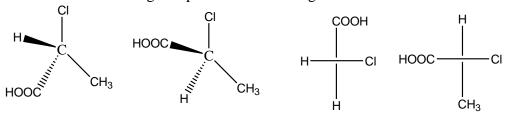
- 1. Identify the relationship between the molecule X and Y.
 - $\begin{array}{c|c}
 CONH_2 & CONH_2 \\
 CI & H & CI & Br \\
 Br & CH_3 & CH_3
 \end{array}$ $\begin{array}{c}
 CONH_2 & CONH_2 &$
 - A. Diastereomers
 - B. Enantiomers
 - C. Identical
 - D. Structural isomers
- 2. Which of the following molecules is chiral?



- B. II
- C. III
- D. IV
- 3. Nomenclate the given molecule:

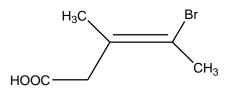
- A. L-Glucose
- B. 2R,3S,4R,5R-Glucose
- C. 2R, 3R, 4S, 5R- Glucose
- D. D-Glucose
- 4. A mixture of equal quantities of two enantiomers is called as.......
 - A. Diastereomeric mixture

- B. Mesomer
- C. Racemic mixture
- D. Optical isomers
- 5. Which of the following compound has R-Configuration?



- Compound-I
- Compound-II
- Compound-III
- Compound-IV

- A. Compound-I
- B. Compound-II
- C. Compound-III
- D. Compound-IV
- 6. Identify the correct name with configuration of the following compound



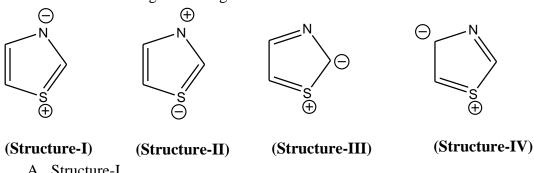
- A. (Z)-4-Bromo-3-methyl-pent-3-enoic acid
- B. (Z)-2-Bromo-3-methyl-pent-2-enoic acid
- C. (E)- 4-Bromo-3-methyl-pent-3-enoic acid
- D. (E)-2-Bromo-3-methyl-pent-2-enoic acid
- 7. Which of the following decrease in order of stability of cyclohexane conformation is correct?
 - A. Chair > twist boat > boat > half chair
 - B. Chair > boat > twist boat > half chair
 - C. Half chair > twist boat > boat > chair
 - D. Chair > boat > half chair > twist boat
- 8.are stereoisomers resulting from hindered rotation about single bonds where the steric strain barrier to rotation is high enough to allow for the isolation of the conformers.
 - A. Enantiomers
 - B. Diastereomers
 - C. Mesomers
 - D. Atropisomers
- 9. Identify the wrong statement about geometrical isomers
 - A. It must contain a carbon-carbon double bond in the molecule
 - B. Two different atoms or groups must be linked to each doubly bonded carbon atoms.

- C. Cis and trans are geometric isomers
- D. It occurs due to the rotation of carbon-carbon single bond
- 10. Nomenclate the following structure

- A. 1H-Pyrazolo[3,4-d]isoxazole
- B. 1H-Pyrazolo[4,3-d]oxazole
- C. 1H-Pyrazolo[2,3-c]oxazole
- D. Oxazolo[4,3-d]1H-pyrazole
- 11. Identify the structure of 3-Acetyl-4-chloro-6-ethyl-1H-indole.

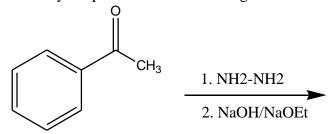
$$C_2H_5$$
 C_2H_5
 C

- A. Compound-I
- B. Compound-II
- C. Compound-III
- D. Compound-IV
- 12. Which of the following resonating structures of Thiazole is incorrect?



A. Structure-I

- B. Structure-III
- C. Structure-IV
- D. Structure-II
- 13. Which of the following five membered heterocyclic rings is most basic?
 - A. Imidazole
 - B. Pyrazole
 - C. Thiazole
 - D. Oxazole
- 14. Which heterocycle is synthesized from malonic ester and urea?
 - A. Pyridine
 - B. Imidazole
 - C. Pyrimidine
 - D. Indole
- 15. Paal-Knorr synthesis of furan is.....
 - A. Cyclization of 1,4-diketone under acidic condition
 - B. Cyclization of 1,2-diketone under acidic condition
 - C. Cyclization of 1,3-diketone under acidic condition
 - D. Cyclization of 1,4-diketone under basic condition
- 16. What is Chichibabin reaction?
 - A. Conversion of pyridine to 3-bromopyridine
 - B. Conversion of quinoline to 2-hydroxyquinoline
 - C. Conversion of pyridine to 2-aminopyridine
 - D. Conversion of quinoline to 8-bromoquinoline
- 17. Bromination of thiophene gives....
 - A. 2,3-dibromothiophene
 - B. 3-bromothiophene
 - C. 3,4-dibromothiophene
 - D. 2-bromothiophene
- 18. Identify the product of the following reaction



- A. Toluene
- B. Ethylbenzene
- C. Phenol
- D. 1-phenylethanol
- 19. Which of the following drug is used as anti-lipidemic?
 - A. Ranitidine
 - B. Celecoxib
 - C. Atorvastatin

D. Zidovudine

20. Identify the heterocycle in the given molecule

- A. Oxazole
- B. Pyrrole
- C. Isoxazole
- D. Furan

S. Y. B. PHARM. SEM-IV-POC-III-DTQ-FH2022

Q.I Attempt ANY ONE of the following

(12M)

- **Q.1** i) Draw all possible resonating structures of pyrazole. Compare the reactivity of pyrrole, furan and thiophene. Which position of the imidazole ring is susceptible to electrophilic aromatic substitution? Justify your answer. (6M)
- ii) Define the terms 'stereospecific' and 'stereoselective'. Discuss, in details, the mechanism of halogenation of cis-2-butene and trans-2-butene. Comment on whether halogenation is stereoselective &/or stereospecific. (6M)
- Q.2 i) What is resolution of racemic modification? Describe in detail, the methods of resolution of racemic modification by giving examples. (6M)
 - ii) Give mechanism with reagents and reaction conditions for the following synthesis:

(6M)

- a) Van-Leusen oxazole synthesis
- b) Hantzsch Pyridine synthesis

Q.II Attempt ANY FOUR of the following

(48M)

- **Q.1** i) Draw conformations of n-butane with energy profile diagram. Comment on the stability of every conformer. (4M)
- ii) Draw all possible projections of 2-Chloro-2-methyl-3-nitro-3-phenylpropan-1-ol. (4M)
- iii) What will be the product of Birch reduction of benzoic acid? Write its mechanism. (4M)
- **Q.2** i) What is asymmetric synthesis? Enlist methods of asymmetric synthesis; explain any one in detail by giving examples. (4M)
- ii) Identify the products A, B, C and D of the following reactions. (4M)

iii) Explain the two necessary conditions for biphenyl compounds to exhibit optical activity with suitable examples. (4M)

Q.3 i) Give mechanism for the following

(6M)

(5M)

a) Radziszewski Imidazole synthesis

i. Friedlander synthesi s of quinoline

ii) Nomenclate the following structures

SN

III)

I)

 H_2N

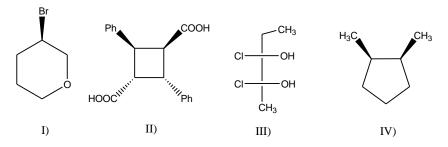
NO₂ IV)

II)

iii) Assign R/S configuration to the given molecule

(1M)

- Q.4 i) Write medicinal uses of Clonidine, Tacrine, Zidovudine and Pyrantel pamoate. (4M)
- ii) Identify whether the following molecules are chiral/achiral. Justify the same. (4M)



- iii) Draw structures of the following
 - a) (Z) 2-Bromo-2-butenoic acid
 - b) (E) 3-Hydroxy-2-butenal
 - c) Cis and trans isomers of 1,2-dimethylcyclohexane
- Q.5 Give the products of following reactions (write structures). (7M)

I) Indole
$$\frac{\text{HCHO}}{\text{NH(CH}_3)_2}$$

II) Pyridine
$$\frac{\text{Br}_2, \text{Oleum}}{130^{\circ}\text{C}}$$

III) Thiophene
$$\frac{\text{HNO}_3/\text{Ac}_2\text{O}}{}$$

ii) Using various oxidising and reducing agents, discuss oxidation and reduction of isoquinoline. (5M)

(4M)