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WINTER - 16 EXAMINATION

Model Answer

Subject Code:

0812

Important Instructions to examiners:

- 1) The answers should be examined by key words and not as word-to-word as given in the model answer scheme.
- 2) The model answer and the answer written by candidate may vary but the examiner may try to assess the understanding level of the candidate.
- 3) The language errors such as grammatical, spelling errors should not be given more Importance (Not applicable for subject English and Communication Skills.
- 4) While assessing figures, examiner may give credit for principal components indicated in the figure. The figures drawn by candidate and model answer may vary. The examiner may give credit for any equivalent figure drawn.
- 5) Credits may be given step wise for numerical problems. In some cases, the assumed constant values may vary and there may be some difference in the candidate's answers and model answer.
- 6) In case of some questions credit may be given by judgement on part of examiner of relevant answer based on candidate's understanding.
- 7) For programming language papers, credit may be given to any other program based on equivalent concept.

Q. No.	Sub Q. N.	Answer	Marking Scheme
Q.1		Attempt any <u>FIVE</u> of the following	5x4=20M
	a)	Write the structure of the following organic groups (Any four)	
		i) Benzoyl	
		Or ——CO-C ₆ H ₅	1M
		ii) Guanidino NH NH C NH ₂	1M

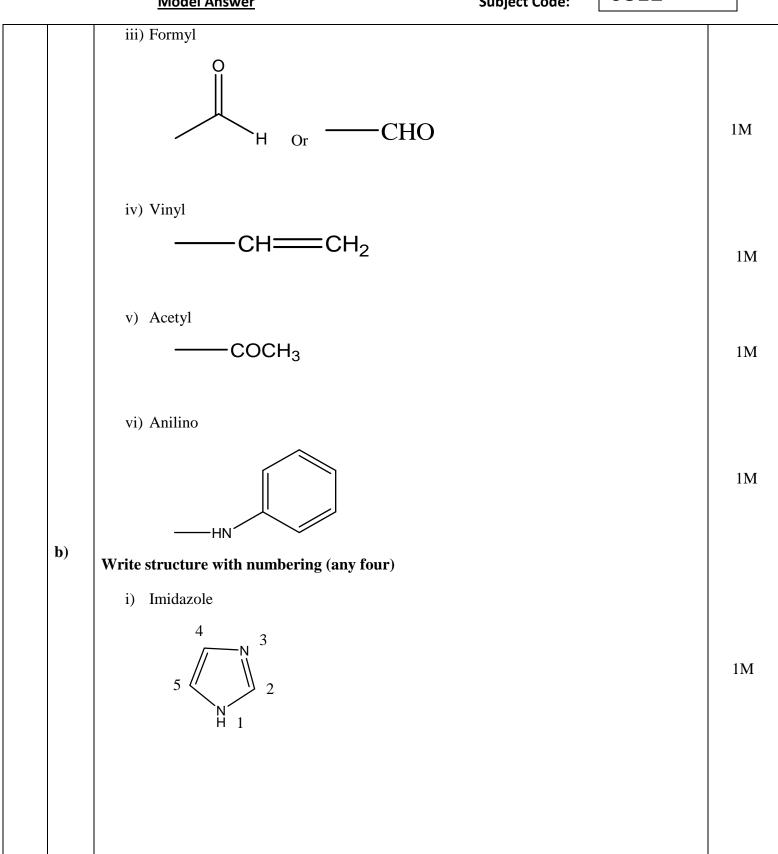


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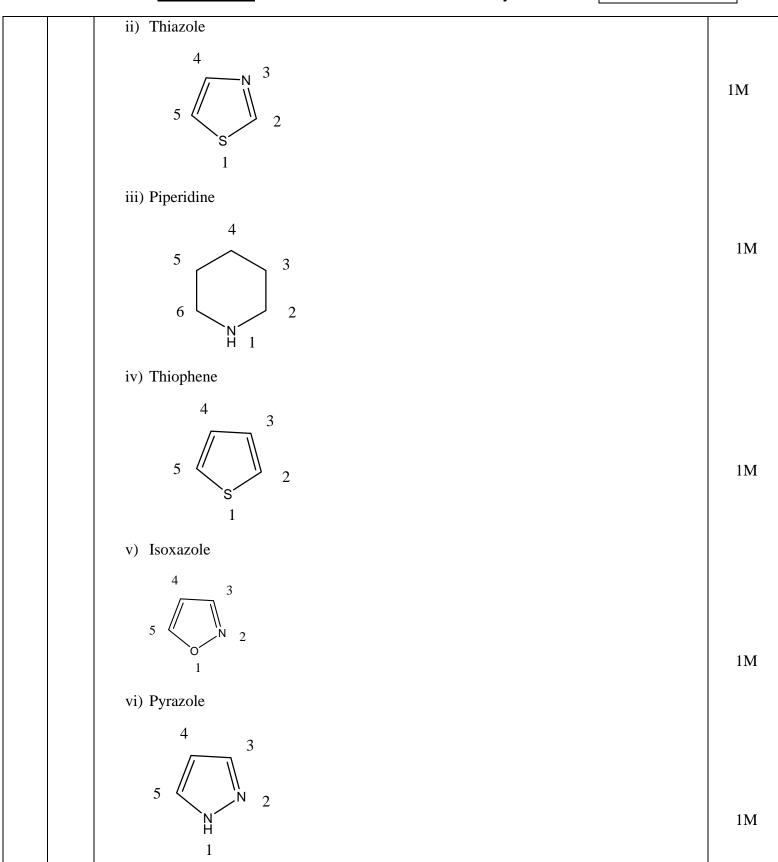


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	c)	Write structure with numbering (any four)			
		i) Indole			
		5 6 7 1 ii) Isoquinoline			1M
		5 4			
		6 7 8 1			1M
		iii) Acridine			
		$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$			1M
		iv) Phenothiazine			
		6 5 4 7 S 3			1M
		8 2			11

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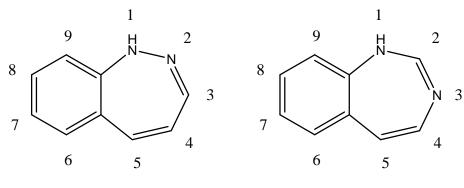
v) Benzimidazole

5

N

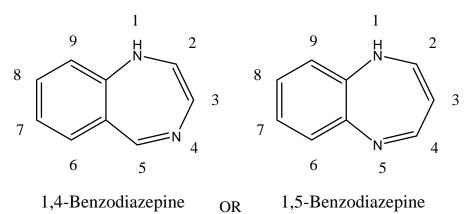
1M

vi) Benzodiazepine



 $1,2\text{-Benzodiazepine} \qquad \text{OR} \qquad \quad 1,3\text{-Benzodiazepine}$

OR



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d) Write the name and structure of drug possessing following moiety (Any two)

i) Barbituric acid

Name of Drug: Phenobarbitone

Name 1M

Struct.

ii) Acridine

Name of drug: Proflavine

₂N NH₂

Name 1M

Struct. 1M

iii) Pyridine

Isoniazid OR

$$C_2H_5$$

Nikethamide

Name 1M Struct

Struct.



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e) Draw structure of drug from given chemical name (any Two)

i) 7-Chloro-1,3-dihydro-1-methyl-5-phenyl-1,4-benzodiazepine-2-one

2M

ii) 1-(Isopropyl amino)-3-(1-naphthyloxy)propan-2-ol

2M

iii) 4-Buty-1,2-diphenyl pyrazolidine-3,5-dione



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- i) Metronidazole: Aristogyl, Flagyl, Metrogyl, Aldezol, Unimezol
- ii) Furosemide: Fru, Lasix, Frusenex
- iii) Diazepam: Calmpose, Valium, Elcion-CR
- iv) Metformin: Dideta SR, Formin, Metchek, Forminal
- v) Propranolol: Ciplar, Inderal, Corbeta, Betacap TR
- vi) Ibuprofen: Ibugesic, Ibuspan SR, Ibuflamar, Brufen

Write structure and chemical name of (any two)

i) Neostigmine

g)

$$H_3C$$
 CH_3
 CH_3
 CH_3
 CH_3

(3-Dimethylcarbamoyloxy-phenyl)-trimethyl-ammonium

ii) Furosemide

$$H_2NO_2S$$
 H_2NO_2S
 H_2NO_2S
 H_2NO_2S

4-Chloro-N-furfuryl-5-sulphamoyl anthranilic acid OR

4-Chloro-2-[(furan-2-ylmethyl)-amino]-5-sulfamoyl-benzoic acid

Struct.

Two brands

1M

1M

Name 1M

Chem.

1**M**

Struct.

Chem. Name



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iii) Atropine

3-Hydroxy-2-phenyl-propionic acid 9-methyl-9-aza-bicyclo[3.3.1]non-3-yl ester

OR

(RS)-(8-Methyl-8-azabicyclo[3.2.1]oct-3-yl) 3-hydroxy-2-phenylpropanoate

Give uses of (any two) h)

- **Iopanoic acid:**
- 1) It is used in Cholecystography (X ray examination of Gall bladder)
- 2) Treatment of Thyrotoxicosis
- ii) Nystatin
- 1) It has wide range of activity against fungi and yeast
- 2) Treatment of candida local infection of mucous membrane, skin, nails
- 3) Pessaries are used in the treatment of vaginal candidiasis
- 4) Treatment of GIT candidiasis
- iii) Fluorescein sodium
- 1) Fluorescein sodium is a diagnostic agent.
- 2) It is used to detect diseased or damaged areas of cornea.
- 3) It is used to detect foreign bodies in the eye.

Struct.

1M

Chem. Name

1M

2M

2M

2M

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3x4 = 12MQ. 2 Attempt any **THREE** of the following Define 'Cardiovascular agents'. Classify them based on their therapeutic applications a) with examples. Def. 1M Cardiovascular agents:- Cardiovascular agents represents a group of drugs which have direct action on heart or other parts of vascular system so that they modify the total output to the heart or the distribution of blood to certain parts of circulatory system. These drugs are used in the treatment of various cardiac diseases like hypertension, angina pectoris, arrhythmia, CHF, myocardial infarction etc. **Classification:** Class. 1) Cardiotonic drugs: 3M Cardiac glycosides obtained from Digitalis, stropanthus like Digoxin, Digitoxin, Gitoxin. 2) Antiarrhythmic agents: a) Membrane stabilizing agent: - Quinidine, Procainamide, Diisopyramide, Phenytoin b) Beta blockers: Propranolol c) Drugs that prolong the duration of action potential:-Amiodarone d) Calcium channel blocker: Verapamil, Amlodipine, Diltiazem 3) Antianginal agents: - a) Organic nitrates – Amyl nitrate, Glyceryl trinitrate, Isosorbid nitrate b) Calcium channel blocker:-Verapamil c) Beta blockers:-Propranolol 4) Antihypertensive drugs: a) Centrally acting drugs:- alpha-methyl Dopa, Clonidine b) Ganglionic blockers - Pentolinium, Mecamylamine



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	Model Answer Subject Code:	
	c) Adrenergic neuron blockers -Reserpine, Guanethidine	
	d) Beta blockers: - Propranolol, Atenolol	
	e) Alpha blockers: - Prazocin, Tolazoline	
	f) Direct acting vasodilators -Hydralazine ,Minoxidil	
	g) Calcium channel blocker:-Verapamil, Diltiazem, Nifedipine	
	h) Angiotensin converting enzyme inhibitors:-Captopril, Enalapril, Lisinopril	
	5) Antihyperlipidemic drugs:-Clofibrate, Simvastatin, Atorvastatin	
	6) Anticoagulants:-Heparin	
	7) Antiplatelet agents:-Aspirin	
	8) Diuretics:-Frusemide, Thiazides	
b)	Write the composition, mechanism of antibacterial action and official preparations of Cotrimoxazole.	
	Cotrimoxazole is the combination of two drugs i.e. Sulphamethoxazole and Trimethoprim.	1M
	It is a mixture of 5 parts of Sulphamethoxazole and 1 part of Trimethoprim. Sulphonamides	
	block the biosynthesis of folic acid from p-amino benzoic acid. Trimethoprim inhibits the	2M
	enzyme folate reductase and blocks the conversion of folic acid to tetrahydofolic acid (THF).	
	THF is the form required for coenzyme synthesis. Combination of Sulphamethoxazole and	
	Trimethoprim by synergism produces bactericidal effect.	
	Official preparations:	
	1. Cotrimoxazole Tablets I.P., B.P.C.	13.4
	2. Cotrimoxazole Mixture B.P.C.	1M
	3. Cotrimoxazole Injection B.P.C.	
	4. Cotrimoxazole Dispersible tablets B.P.C.	
	 Pediatric Cotrimoxazole mixture B.P.C. 	
	2. I calatile Confinement in the Diff. C.	

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	6. Cotrimoxazole Oral suspension I.P.	
c)	Define 'General anesthetics'. Classify based on their route of administration with examples.	13.5
	Definition: General anesthetics are the central nervous system depressant drugs which bring	1M
	about loss of all modalities of sensations along with a reversible loss of consciousness.	
	Classification:	
	1) Inhalation anaesthetics:-,which include the liquids of volatile nature and gaseous	2) (
	substances used by inhalation to produce anaesthesia. These may be sub-classified as	3M
	follows:	
	i) Volatile liquids:	
	(a) Halogenated hydrocarbons: e.g. Chloroform, Halothane, Trichloroethylene,	
	Ethylchloride	
	(b) Ethers : e.g. Diethyl ether, Vinyl ether	
	ii) Gases: e.g. Cyclopropane, Nitrous oxide	
	2) Intraveneous anaesthetics:-	
	i) Barbiturates: Ultra short acting barbiturates such as Methohexitone, Thiopentone	
	sodium	
	ii) Non-barbituates:	
	a) Eugenol derivatives. e.g. Propanidid	
	b) Phencyclidine derivatives. e.g Ketamine	
	c) Steroids. e.g. Althesin	
	d) Miscellaneous. such as Etomidate, Propofol	



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Define and classify Antimalarials. Give structure of Chloroquine.

1**M**

Definition: Antimalarial drugs are used in the treatment of malaria caused due to Plasmodium Vivax, P. falciparum, P. Ovale, P. malariae, P. Knowlesi.

Classification:

1. Cinchona alkaloids: Quinine

2M

- 2. 4-amino quinolines: Chloroquine, Amodiaquine, Santaquin
- 3. 8- Amino quinolines: Primaquine, Pentaquine, Isopentaquine
- 4. 9-Amino acridines: Quinacrine
- 5. Pyrimidines: Pyrimethamine, Trimethoprime
- 6. Biguanides: Proguanil, Cycloguanil
- 7. Sufonamides and sulfone: Sulfadoxin, Dapsone
- 8. Miscellaneous: Tetracycline, Artesunate, Artemether

$$C_2H_5$$
 C_2H_5

1M

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Model Answer Subject Code:	0812
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e)	Define Diabetes Mellitus and Hypoglycemic agents. Enumerate different Insulin	
	preparations.	
	Diabetes Mellitus: - Diabetes Mellitus is a condition characterized by hyperglycemia	1M
	(excessive sugar in blood, than the threshold value) & glycosuria (presence of sugar in	
	urine). The disease is caused by deficiency of insulin, a protein hormone secreted by beta	
	cells of islets of Langerhans, responsible for proper carbohydrate metabolism.	
	Hypoglycemic agents are the drugs that decrease the level of glucose in the blood and are	1M
	used in the treatment of diabetes mellitus characterized by hyperglycemia, glycosuria,	11/1
	polyuria, polydypsia etc.	
	Insulin Preparations:	
	1) Insulin injection	23.4
	2) Insulin injection biphasic	2M
	3) Neutral insulin injection	
	4) Globin zinc insulin injection	
	5) Isophane insulin injection	
	6) Protamine zinc insulin injection	



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Q. No.	Sub Q. N.	Answer	Marking Scheme				
Q. 3		Attempt any THREE of the following	3x4=12M				
	a)	Classify antihistaminic agents giving suitable examples. Write structure of Diphenhydramine					
		Classification:					
		1. H ₁ blockers or H ₁ antagonist:					
		a) Aminoalkylethers e.g. Diphenhydramine	3M				
		b) Ethylenediamine e.g.Mepyramine, Tripelennamine					
		c) Alkylamines e.g. Pheniramine, Chlorpheniramine, Triprolidine					
		d)Phenothiazine derivatives e.g. Promethazine, Trimeprazine					
		e) Piperazine derivatives. E.g Meclizine, Cyclizine, Chlorcyclizine					
		f) Miscellaneous e.g. Cyproheptadine, Phenindamine tartrate					
		2. H ₂ Blockers or H ₂ receptor antagonist					
		e.g. Ranitidine, Cimetidine, Famotidine					
		3. An inhibitor of histamine release					
		e.g. Sodium Cromoglycate					
		Structure: Diphenhydramine					
			1M				
	b)	Define local anesthetics. Write structure, chemical name, physical properties and brand name of Lignocaine hydrochloride.	age 15 of 31				



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Local anesthetics: Local anesthetics are drugs which produce insensitivity in a limited area	
around the site of application or injection of the drug by preventing generation and	1M
conduction of impulses along nerve fibers and nerve ending and the effects are reversible.	1111
Structure	
CH ₃	
∠C₂H₅	
NHCOCH ₂ —N . HCI	1M
C ₂ H ₅	114
Сн ₃	
Chemical Name: 2-(diethylamino)-N-(2,6-dimethylphenyl)acetamide hydrochloride	1 N
Physical Properties:Hydrochloride salt occurs as white crystalline powder	0.5
Odourless	0.5
Slightly bitter numbing taste	
Very soluble in water	
Freely soluble in alcohol and soluble in chloroform	
Brand names: Anestacon, Dalcaine, Ultacaine, Xylocain, Xylocard, Lignox, (Any one)	0.5
Define and classify Sympathomimetics. Write structure of Isoprenaline.	
Sympathomimetics: Drugs that mimic the actions obtained as a result of stimulation of the	
sympathetic or adrenergic nerves are called Sympathomimetics.	
sympathetic of adrenergic herves are canca sympathonimicaes.	1N
OR	
The drugs that produce pharmacological effects like adrenaline or nor adrenaline or drugs	
which bring about stimulation of adrenergic nerves are called Sympathomimetics.	
wanta camp account canada ca autonorgeo and camp a camp and camp account canada	
Classification:	2N
Catecholamines e.g : Adrenaline, Nor-adrenaline, Isoprenaline, Dopamine	
Non-Catecholamines eg. Phenylephrine, Salbutamol, Terbutaline, Ephedrine,	
Pseudoephedrine	
1 seudoepheurine	



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• Imidazoline derivatives eg. Naphazoline, Xylometazoline

OR

- 1. Vasoconstrictors († B. P.): Noradrenaline (Norepinephrine), Dopamine, Ephedrine
- 2. Cardiac stimulants: Dopamine, Adrenaline (Epinephrine), Isoprenaline
- 3. **CNS stimulants:** Amphetamine
- 4. Smooth muscle relaxants: Adrenaline, Isoprenaline, Salbutamol, Terbutaline
- 5. **Drugs used in allergic reactions:** Ephedrine
- 6. **Local vasoconstrictor/ nasal decongestants:** Phenylephrine, Pseudoephedrine, Naphazoline
- 7. **Anorectics** (↓ **Appetite**): Amphetamine, Phentermine.

Structure:

HO CH-CH₂-NH-CH

Isoprenaline

d) Define vitamins. Classify giving suitable examples.

Vitamins: Vitamins are the constituents of the diet other than carbohydrates, fats, proteins and inorganic salts and are necessary for the normal metabolic functions of the body.

Classification:

3M

1**M**

- I) Fat soluble vitamins: Vitamin A (Retinol), Vitamin D (Calciferol), Vitamin E (Tocopherol), Vitamin K (Menadione)
- II) Water Soluble vitamins: Water Soluble vitamins (includes Vitamin B-Complex and Vitamin C): Vitamin B_1 (Thiamine), Vitamin B_2 (Riboflavin), Vitamin B_3 (Niacin), Vit. B5 (Pantothenic acid), Vitamin B_6 (Pyridoxine), Vitamin B_7 (Biotin), Folic acid, Vitamin B_{12} (Cyanocobalamine) Vitamin C (Ascorbic acid).



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	9)	Define diuretics, Write structure, chemical name and mechanism of action of urea as a diuretic.			
	e)	Diuretics: The drugs which increase the rate of formation & excretion of urine through	13.5		
		kidneys primarily by inhibiting tubular reabsorption of sodium and its osmotic equivalent	1M		
		amount of water.			
		Structure NH ₂ -CO-NH ₂			
			1M		
		Chemical Name: Carbonyl diamide, Carbamide			
		Mechanism of action: Urea is an osmotic diuretic. Osmotic diuretics work by expanding			
		extracellular fluid and plasma volume, therefore increasing blood flow to the kidney. This			
		washes out the cortical medullary gradient in the kidney. This stops the loop of Henle from	1M		
		concentrating urine, which usually uses the high osmotic and solute gradient to transport			
		solutes and water.			
Q. 4		Attempt any <u>THREE</u> of the following.	3x4=12M		
	a)	Name one drug each used in (any four)			
	,	i) Gout- Diclofenac, Ibuprofen, Naproxen, Celecoxib, Allopurinol, Colchicine	1M each		
		ii) Myasthenia gravis – Neostigmine, Physostigmine, Pyridostigmine			
		iii) Parkinsonism-Atropine, Levodopa, Amantadine, Biperiden			
		iv) Raynaud's disease- Calcium channel blockers- e.g. Amlodipine, Felodipine			
		v) Vasodilators- Nitroglycerine, Losartan, Alpha blockers- Prazosin, Doxazosin			
		vi) Fungal infection- Griseofulvin, Amphotericin, Nystatin, Miconazole, Ketoconazole,			
		Econazole, Fluconazole			
	b)				
	,	i) Define coagulants. Write structure and chemical name of Menadione.			
	Coagulants: These are the agents used in the treatment of severe hemorrhage, causing		2M		
		coagulation of blood.			



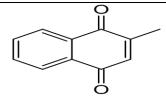
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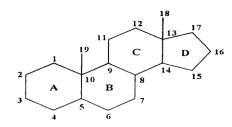
0812



Chemical Name: 2-Methyl-1,4-naphthoquinone

ii) What are steroids? Draw the basic steroidal nucleus with numbering. Give properties and uses of testosterone.

Steroids are polycyclic organic compounds containing 1,2 cyclopentanoperhydrophenanthrene skeleton. i.e. it contains four fused rings A, B, C, D and thus are polycyclic hydrocarbons. The ring A, B and C are six membered and ring D is five membered.



Properties: White crystalline powder, tasteless, odorless, practically insoluble in water, soluble in fixed in oils. It is incompatible with oxidizing agent.

Uses:

- As a substitute in the male for replacement therapy in hypogonadism
- To correct penile size in childhood
- To treat male infertility
- For muscle development
- In the palliative treatment of disseminated breast cancer in females and management of some menopausal disorders.
- Labor pain relief



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Model Answer Subject Code: What does the abbreviation NSAIDs stand for? Why are they so named? Give c) structure and chemical name of Indomethacin. **Ans.** Abbreviation NSAIDs stand for Non steroidal anti-inflammatory drugs. 1M They are named so because they do not contain the steroidal nucleus & mechanism of action of these agents differ from those of anti-inflammatory steroids and they are devoid of 1M undesirable effects of steroid therapy. Structure: 1M Chemical name: 1-(4'-chlorobenzoyl)-5-methoxy-2- methylindo-3-ylacetic acid 1M **Define and classify Antineoplastic with examples.** d) **Definition:** Antineoplastic agents, also known as cytotoxic agents are used in the treatment of malignant diseases when surgery or radiotherapy is not possible or has proved ineffective, 1M in other words, antineoplastic agents are used in the treatment of cancer. Classification: 3M 1. Alkylating Agents. a) Nitrogen mustard drugs: Mustine, Chormabucil, cyclophosphamide b) Aziridines: Thiotepa c) Alkyl sulphonate: Busulphan d) Nitrosourea group compound: Lomustine 2) Antimetabolites: Methotrexate, Mercaptopurine, Azathioprine, Fluorouracil 3) Antibiotics: Actinomycin, Daunorubicin, Doxorubicin

4) Plant Products: Sulphates of vinblastin and vincristine.

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	<u>Mo</u>	<u>del Answer</u>	Subject Code:	0012	
	5) Hormone	s and related drugs: Glucocortico	ids, Tamoxifen		
	6) Miscellan	neous agents: Hydroxyurea, cispla	ntin		
e)	Define and	classify Antitubercular agents v	with example.		
	Definition :	The agents used in treatment of to	aberculosis, a disease caused by	Mycobacterium	1M
	species (M.	Tuberculosis, M. bovis or M. afri	canum) characterized by formati	on of nodular	1111
bodies or tubercles.					
	Classification	on of Antitubercular drugs:			
	i) p-am	nino salicylic acid derivative – e.g	g. PAS		
	ii) Pyrio	dine derivatives – e.g. Isoniazid, I	Ethionamide		3M
	iii) Pyra	zine derivatives- e.g. Pyrazinamio	de		
	iv) Ethy	lene diamine derivatives – e.g. Et	thambutol		
	v) Antil	biotics – e.g. Cycloserine, Strepto	omycin, Rifampicin		
			OR		
	i) First l	ine drugs			
	e.g.Iso	oniazid, Rifampin, Ethambutol, P	yrazinamide, Streptomycin, Thio	pacetazone etc.	
	ii) Second	d line drugs			
	e.g.Et	hionamide, Kanamycin, capreom	ycin, Cycloserin, Para amino sal	icylic acid etc.	
	ĺ	line drugs			
	e.g.Cla	arithromycin, Thioacetazone			
			OR		
	1. Synthe	tic anti-tubercular drugs:			
	Para An	nino Salicylic acid (PAS), Isoniaz	zide, Ethambutol, Pyrazinamide	, Ethionamide	
	2. Antibio	otics:			
	Streptor	mycin, Cycloserine, Rifampin, Cl	arithromycin		
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0.5		Attended one Tipper of the following	2-:4 1214
Q.5		Attempt any <u>THREE</u> of the following.	3x4=12M
	a)	Define 'Analeptics'. Name an analeptic each, which is	
		(i) a Xanthine derivative	
		(ii) a Pyridine derivative	
		Also write structure with chemical name of the two.	13.6
		Definition: Analeptics increases activity in certain areas or the whole of the brain. These	1M
		drugs are used to stimulate central nervous system, so it reduces narcosis brought about by	
		excess of depressant drugs.	
		Name an analeptic:	1M
		(i) a Xanthine derivative :- Caffeine	1 1V1
		(ii) a Pridine derivative :- Coramine (Nikethamide)	
		Structure of caffeine	
		H ₃ C N N N N CH ₃	1M
		Chemical name of Caffeine: 1,3,7-Trimethyl xanthine	
		Structure of Coramine	
			1M
		Chemical name of Coramine: N, N-diethyl, pyridine-3-carboxamide.	



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b) Define 'Antiseptics and Disinfectants'. Classify with examples and write the structure of chlorocresol.

Antiseptics: Antiseptics are the agents that are used on living tissues & act as antimicrobial but don't kill them necessarily.

1M

Disinfectants:- Disinfectants are agents which are applied on inanimate objects & kill the microbes outright.

Classification:-

- 1) Phenols & related compounds: Phenol, Chlorocresol. Chloroxylenol, Hexachlorophene
- 2) Alcohols & aldehydes: Alcohol, Formaldehyde
- 3) Halogen compounds: Chloramine t, Chorhexidine acetate, Dibromopropamidine
- 4) Organic mercurials: Merbromin (mercurochrome), Thiomersal

2M

- 5) Dyes: Aminacrine hydrochloride, Brilliant green, Proflavine hemisulfate, Crystal Violet (gentian violet), Acriflavine.
- Cationic surface-active agents. e.g. Cetylpyridinium chloride, Benzalkonium chloride, Cetrimide
- 7) Miscellaneous agents. e.g. Dequalinium chloride, Nitrofurazone

Structure: Chlorocresol

1**M**

Define the term 'Cardiotonics'. Write about their hydrolysis products.

1**M**

Cardiotonics:

c)

These are the drugs which have stimulating action on the cardiac muscles. They increase the force of muscle contraction without increasing oxygen consumption capacity of heart.

Cardiac glycosides on hydrolysis yield corresponding sugar and aglycones.

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d)

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Cr. No.	Glycoside	Products of Hydrolysis	
Sr. No.		Sugar	Aglycone
1	Digitoxin	3moleculesof digitoxose	Digitoxigenin
2	Digoxin	3 molecules of digitoxose	Digoxigenin
3	Lanatoside	2 molecules of digitoxose; 1 molecule of acetyl digitoxose and 1 molecule of glucose	Digoxigenin

Define antidepressants. Give mechanism of action of MAO inhibitors. Write structure of Imipramine.

Antidepressants: Antidepressants are drugs which counteract or overcome mental depression. These drugs are therapeutically useful in a variety of cases pertaining to mentally ill patients. Mental depression is a phenomenon which may arise in normal individuals or in mentally ill persons.

Mechanism of mono amino oxidase Inhibitor

These drugs block oxidative deamination of naturally occurring amines. MAO enzyme is present intracellularly in most of the tissues (highest conc. in liver). Enzyme oxidises active biogenic amines like 5HT, noradrenaline, & dopamine to inactive compounds. These amines are normally stored in granules in the neurons & get liberated by nervous stimuli. MAO inhibitors inhibit the enzyme & result in accumulation of these amines in the brain. Ultimately excitement, enhanced motor activity is observed. So MAO inhibitors are used as antidepressants.

3M

1M

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Imipramine:

1M

1M

Penicillin is the first antibiotic to be discovered. The various penicillins are obtained by fermentation using various strains of mold penicillium. The penicillins are strong monobasic acids. They readily form salts and esters. The penicillins are derivatives of 6- amino penicillanic acid. Penicillin contains β -lactum ring, a 4-membered cyclic amide, which is fused with thiazolidine ring. This bicyclic heterocyclic system is called penam. –COOH functional group is present at C_3 and amide group at C_6 .

 $R - H_2C - C - HN - S - CH_3$ $R - H_2C - C - HN - S - CH_3$ $R - H_2C - C - CH_3$ $R - H_2C - C - CH_3$ $R - H_2C - C$

1**M**

Different penicillin:

- 1. Benzyl penicillin (Penicillin G)
- 2. Phenoxy methyl penicillin (Penicillin V)
- 3. Ampicillin
- 4. Amoxicillin
- 5. Cloxacillin
- 6. Carbenicillin



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Q.6

a)

Attempt any <u>THREE</u> of the following.

What are Beta adrenergic blockers? Give two examples and structure of any one of

them and therapeutic uses.

Beta adrenergic blockers: These drugs inhibit adrenergic responses mediated through the β -receptors. Beta blockers are competitive antagonists that block the receptor sites for the endogenous catecholamines epinephrine (adrenaline) and norepinephrine (noradrenaline) on adrenergic beta receptors, of the sympathetic nervous system. Some block activation of all types of β -adrenergic receptors and others are selective.

β-adrenergic blockers e.g. Propranolol, Atenolol, Metoprolol, Betoxolol. Nodolol, Timolol, Acebutolol, Esmolol etc.

Structure of Propranolol:

Therapeutic uses of Propranolol:

- 1. It is a typical beta adrenergic receptor blocker used in the treatment of cardiac diseases like
 - Angina pectoris
 - Cardiac arrhythmia
 - Hypertension
 - Congestive heart failure
 - Coronary atherosclerosis
- 2. Treatment of Pheochromocytoma

1M

3x4=12M

1**M**

1**M**

1**M**

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b) Define 'Sedatives and Hypnotics'. Give structure, chemical name and uses of a hypnotic having piperdine nucleus in its structure.

Hypnotics: Hypnotics are drugs which induce sleep by depression of central nervous system function.

Sedatives: Sedatives are the agents which reduce excitement & motor activity & produce a calming effect without inducing sleep.

Hypnotic having piperidine nucleus in its structure is Glutethimide.

Structure of Glutethimide:

Uses of Glutethimide:

c)

Chemical name -3-Ethyl-3-phenyl-piperidine-2,6-dione OR 3-Ethyl-3-phenyl-glutarimide

- 1. It is used as Hypnotic and sedative in insomnia.
- 2. It is used in treatment of anxiety and tension

Amoebiasis: Amoebiasis is a parasitic infection of the intestines caused by the protozoan *Entamoeba histolytica*. The symptoms of amoebiasis include abdominal pain, passage of soft stools with mucus & occasional blood, fatigue, excessive gas, rectal pain, unintentional weight loss etc.

What is Amoebiasis? Write structure, chemical name and uses of Metronidazole.

1**M**

1M

1M

1M

1M

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Structure of Metronidazole:

Chemical name: 2-(2'-Methyl-5'-nitroimidazolyl)ethanol OR

1-(2'-hydroxyethyl)-2- methyl- 5-nitro imidazole.

Uses:

- 1. It has antiprotozoal and antibacterial action
- 2. It is used in the treatment of severe intestinal amoebiasis
- 3. It is active against anaerobic bacteria like streptococci and H-Pylori
- 4. It is a primary drug in the treatment of hepatic amoebiasis
- 5. Treatment of *Trichomonous vaginalis*, infection due to *entamoeba histolytica*, *giardia lamblia* etc.

d)

Name one tranquilizer with its structure and chemical name belonging to :-

I) Phenothiazine class: Chlorpromazine

1M

1M

1**M**

1**M**

Structure

1**M**

Chemical name: 2-Chloro-10-(3'-dimethylamino propyl) phenothiazine

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II) Butyrophenone class: Haloperidol.

1**M**

Structure

1M

Chemical name: 4-[4'-(4"-chlorophenyl),4-hydroxy piperidine] 4-fluorobutyrophenone.

e)

i) Write structure and chemical name (Any two):

1) Ampicillin

Structure:

0.5 M

0.5M

 $\textbf{Chemical name:} \ 6\hbox{-}(2\hbox{-amino-}2\hbox{-phenylace} tamido)\hbox{-}3,3\hbox{-dimethyl penam-}2\hbox{-carboxylic acid}.$

OR (6R)-6-(α-phenyl-D-glycylamino)penicillanic acid

0.5M

2) Paracetamol

Structure:

0.5M



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Chemical Name: N-acetyl-p-aminophenol; OR 4-hydroxyacetanilide 0.5M

3) Pethidine

Structure:

0.5M

Chemical name: Ethyl-1-methyl-4-phenyl-piperdine-4-carboxylate.

ii) Write stability storage conditions for (Any two):

1. Diethyl ether:

1M

- It is oxidized by atmospheric oxygen and is affected by light. Hence it is stored in tightly closed, light resistant containers in a cool place.
- If cork is used as a closer than it should be protected with metal foil. An antioxidant like hydroquinone or propyl gallate in suitable proportion should be added.

2. Adrenaline:

1M

- It contains catechol nucleus which can be oxidized readily with air or oxygen to get pink or red colored complex.
- It darkens on exposure to light and air, hence it should be stored in well closed, tight container and protected from light.

3. Insulin:

- As insulin is affected by heat & light, all insulin preparations must be stored at low temperatures between 2-8°C in a dark place.
- It should not be allowed to freeze.



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