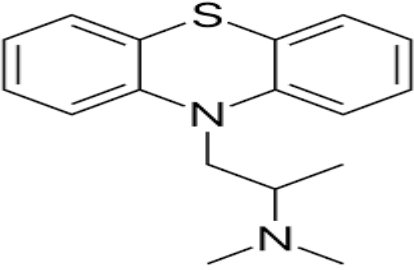
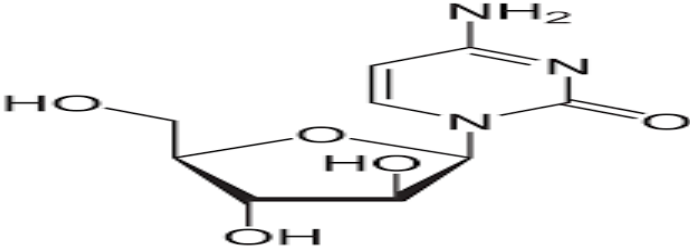
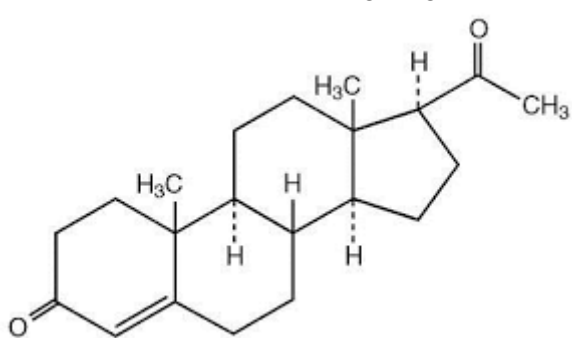


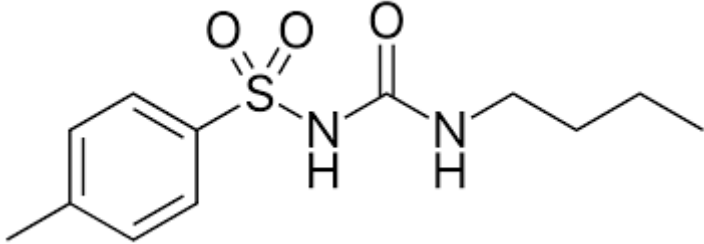
Q. 1	Choose appropriate option for following multiple choice based questions.	20 mks
1	H3 receptors are mostly found high density in	
a	brain	
b	respiratory system	
C	GIT system	
d	nerve cells	
2	H2 antihistaminic have following structural feature as	
a	indole	
b	imidazole	
C	ethylenediamine	
d	ethanolamine	
3	identify the following drug. <div style="text-align: center; margin: 10px 0;">  </div>	
a	promethazine	
b	triprolidine	
C	azatidine	
d	buclizine	
4.	what is mode of action of ranitidine?	
a	H⁺/K⁺ ATPase Proton pump inhibitor	
b	H1 antagonist	

C	H2 antagonist	
d	topical antihistaminics	
5.	<p>What is this drug's mode of action?</p> 	
a	DNA methylating agent	
b	Intercalater	
C	Chain terminator	
d	antimetabolites	
6.	Identify structural feature in dactinomycine from the following	
a	oxazole	
b	ketone	
C	thiazole	
d	phenoxazine	

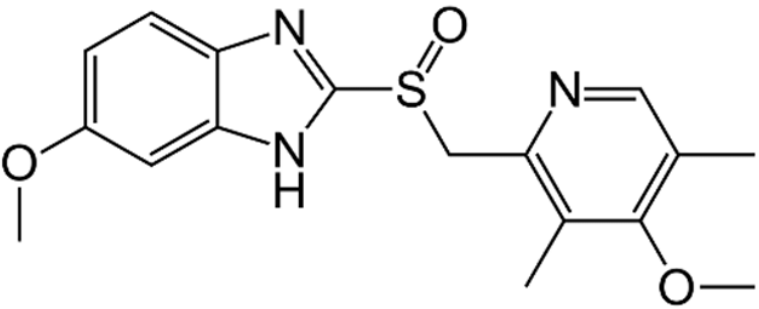
7.	Identify the following structure	
a	Mechlorethamine	
b	busulfan	
C	cyclophosphamide	
d	melphalan	
8.	zinc binding group in ACE inhibitor is	
a	alkyl group	
b	sulphydryl group	
C	ester group	
d	amide group	
9.	Mode of action of dipyridamole is	
a	Ca channel blocker	
b	loop diuretics	
C	vasodilation	
d	potassium sparing diuretic	
10	furosemide is	
a	salicylic acid derivative	
b	anthranilic acid derivative	
C	benzoic acid derivative	

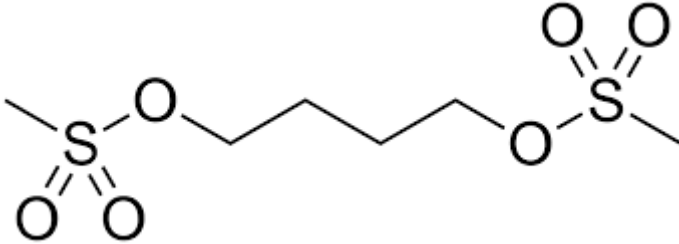
d	Thiazide devative	
11	Mechanism of action of Spironolactone is	
a	interfere with Sodium and chloride reabsorption	
b	Block Calcium channel	
C	Block Potassium channel	
d	ACE inhibitors	
12	mexiletine is specifically used for	
a	supraventricular arrythmia	
b	prevetion of ventricular arrythmia	
C	hypertension	
d	diuretic treatment	
13	phenytoin acts by	
a	shorten phase 3 repolarization	
b	K channel blockade	
C	prolongs phase 3 repolarization	
d	Na channel blockade	
14	clofibrate decrease significantly	
a	cholesterol level	
b	LDL level	
C	blood glucose level	
d	plasma triglycerides	

15	estrogen to 4-methylestrogen metabolism is carried out by	
a	16 α hydroxylase	
b	17 α hydroxylase	
C	COMT and estrogen 2,4,hydroxylase	
d	5 α reductese	
16	Write the name of the following drug.	
		
a	Cortisone	
b	Hydrocortisone	
C	progesterone	
d	estrogen	
17	propylthiouracil is used as	
a	thyroid replacement therapy	
b	natural thyroid	
C	hyperthyroidism	
d	thyrotoxicosis	
18	following statement is wrong about meglitinides	
a	benzoic acid derivatives of glibenclamide	
b	shut down ATP sesnsitive K channel	
C	insulin secretogouges	

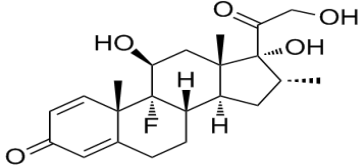
d	α glucosidase inhibitor	
19	What is the name of the structure given below? 	
a	Glipizide	
b	glibenclamide	
C	Tolbutamide	
d	Acrabose	
20	lignocaine is	
a	benzoic acid derivative	
b	aminobenzoic acid derivative	
C	anilide derivative	
D	nitrobenzoic acid derivative	

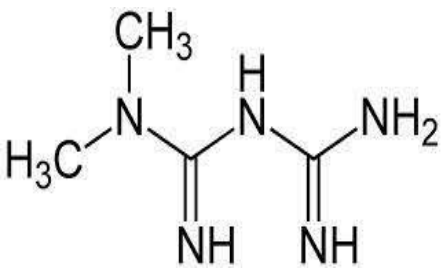
21	H1 Anti-histamines used for motion sickness and nausea include	
a	Diphenhydramine and meclizine	
b	Tornadol and meclizine	
C	Ephedrine and Luratidine	
d	Luratidine	
22	Which of the following statements about histamines is correct?	

a	Histamine is stored in peripheral nerve endings	
b	Histamine is released from mast cells following and allergic challenge	
c	Histamine is a vasoconstrictor	
d	Histamine is an essential amino acid.	
23	Name the following drug.	
		
a	Omeprazole	
b	Lancoprazole	
c	Rabeprazole	
d	Panteprazole	
24.	Which one of the following drugs bind with tubulin and arrest the cell cycle in metaphase.	
a	Vinca	
b	Nitrogen mustards	
c	Antimetabolites	
d	Alkylating agents	

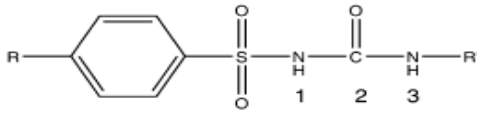
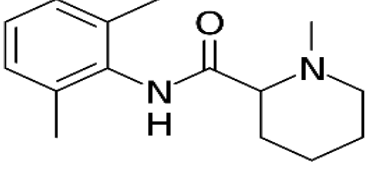
25.	What is this drug's mode of action? 	
a	Metallating agents	
b	Intercalater	
c	Chain terminator	
d	Alkylating agents	
26.	Which of the following type of angina is caused by Coronary Spasm	
a	Stable Angina	
b	Unstable Angina	
c	Crescendo Angina	
d	Prinzmetal Angina	
27.	The following drug acts at the proximal convulated tubule	
a	Hydrochlorothiazide	
b	Furosemide	
c	Spironolactone	
d	Acetazolamide	
28.	4-chloro-2-[(furan-2-ylmethyl) amino] 5-sulphamoylbenzoic acid is the IUPAC name of	
a	Hydrochlorothiazide	
b	Furosemide	
c	Spironolactone	

d	Acetazolamide	
29.	Which one is not PDE inhibitor	
a	Sildenafil	
b	Tadalafil	
C	Testosterone	
d	Acetildenafil	
30	Initial medication orders for the treatment of hypertension are likely to include any of the following except	
a	ACE Inhibitors	
b	Beta blockers	
C	Calcium channel blockers	
d	Thiazide type diuretics	
31	Mechanism of action of Mexiletine hydrochloride is	
a	Block Sodium channel	
b	Block Calcium channel	
C	Block Potassium channel	
d	Beta blockers	
32	The anti-arrhythmic drug with local anaesthetic action is	
a	Verapamil	
b	Procainamide	
C	Encainide	
d	Digoxin	

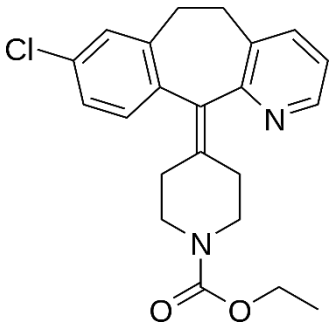
33	Which one of the following drug act by sequestering the bile acids in the GIT?	
a	Colestipol	
b	Lovastatin	
c	Probucol	
d	Niacin	
34	Mechanism of action of Digitoxin is	
a	Decrease in calcium uptake	
b	Increase in ATP synthesis	
c	Modification of of actin molecules	
d	Increase in intracellular calcium level	
35	Pregn-4-en-3,20 Dione is the IUPAC name for	
a	Estradiol	
b	Prednisone	
c	Progesterone	
d	Sildenafil	
36	<div style="text-align: center;">  <p>The image shows the chemical structure of Dexamethasone, a corticosteroid. It features a four-ring steroid nucleus with a ketone group at C3, a double bond between C4 and C5, a fluorine atom at C6, a hydroxyl group at C11, and a diene side chain at C17 consisting of a double bond, a ketone group, and a hydroxyl group.</p> </div> <p>Write the name of the following drug.</p>	
a	Cortisone	
b	Hydrocortisone	
c	Dexamethasone	

d	Betamethasone	
37	Following is a synthetic derivative of thyroxine	
a	Propylthiouracil	
b	Methimazole	
C	Levothyroxine	
d	Levothyronine	
38	Chlorpropamide is an example of	
a	Byguanides	
b	Thiazolidinediones	
C	Meglitinides	
d	Sulphonylureas	
39	What is the name of the structure given below? 	
a	Glipizide	
b	Tolbutamide	
C	Metformin	
d	Acrabose	
40	Which one of the following is a natural local anesthetic?	

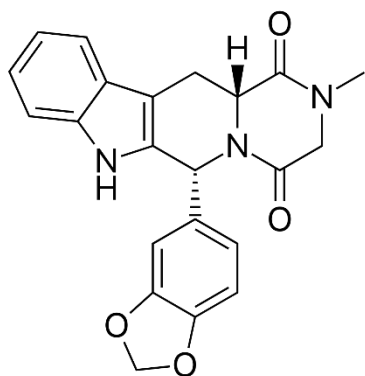
a	Lignocaine	
b	Benzocaine	
C	Cocaine	
D	Procaine	
41	The gauche conformer of histamine has a preferred affinity towards _____receptor/s	
a	H1	
b	H2	
C	H3	
d	H1 and H2	
42	A nitrate vasodilator causes	
a	Inhibition of soluble guanylate cyclase	
b	Activation of soluble guanylate cyclase	
C	Activation of myosin light chain kinase	
d	Inhibition of cGMP-dependent protein kinases	
43	With respect to chemical structure, Methotrexate differs from folic acid by	
a	Presence of -NH ₂ at position 4 and absence of -CH ₃ at N10	
b	Presence of -CH ₃ at position 4 and -NH ₂ at N10	
C	Presence of -NH₂ at position 4 and -CH₃ at N10	
d	Presence of -CH ₃ at position 4 and absence of -NH ₂ at N10	
44.	Which of the following statement is false related to structural features of cardiac glycoside?	
a	The aglycone portion of the cardiac glycosides is a steroid nucleus	

b	The steroid nucleus also carries two angular methyl groups at C-10 and C-13	
c	The lactone ring is present at C17 in aglycone part	
d	Hydroxy group located at C14 in a steroid nucleus is a point of sugar attachment	
45.	<p>Which of the following statement is false about sulfonylureas?</p> 	
a	These are urea derivatives with an arylsulfonyl group in the 1-position and an aliphatic group at the 3-position	
b	The aliphatic group, R', confers hydrophilic properties to the molecule.	
c	Maximal activity results when R' consists of three to six carbon atoms	
d	Aryl groups at R' generally give toxic compounds	
46.	Hydralazine shows the presence of	
a	Phthalazine	
b	1,2,4-benzothiazine	
c	N-oxide of a piperidinopyrimidine	
d	Indole nucleus	
47.	<p>Identify the following local anaesthetic agent.</p> 	
a	Dibucaine	
b	Mepivacaine	
c	Prilocaine	
d	Etidocaine	

48.	The 9 α -fluoro group in betamethasone	
a	increases the glucocorticoid activity and decreases the mineralocorticoid activity	
b	decreases the glucocorticoid activity and increases the mineralocorticoid activity	
C	decreases the glucocorticoid activity and decreases the mineralocorticoid activity	
d	increases the glucocorticoid activity and increases the mineralocorticoid activity	
49.	Which of the following local anesthetic is an ester of phenyl carbamic acid	
a	Dibucaine	
b	Diperodon	
C	Phanacaine	
d	Etidocaine	
50	Which of the following structural features are found in quinidine	
a	Quinoline ring, quinuclidine ring and hydroxymethylene bridge	
b	Isoquinoline ring, decaline ring and methylene bridge	
C	Indole ring, quinuclidine ring and methylene bridge	
d	Benzofurane, decaline ring and hydroxymethylene bridge	
51	The piperazine nucleus is present in all of the following drugs except	
a	Cetirizine	
b	Chlorocyclizine	
C	Meclizine	
d	Phenindamine	

52	The potential nucleophilic site on DNA, which are susceptible to electrophilic attack by an alkylating agent is	
a	N-7 position of guanine	
b	N-7 position of adenine	
c	N-3 position of cytosine	
d	N-1 position of guanine	
53	Choose the correct non-dihydropyridine calcium channel blocker	
a	Nifedipine	
b	Amlodipine	
c	Verapamil	
d	Nicardipine	
54	Identify the class to which the following drug belongs	
		

a	First generation H1 antihistaminics	
b	Second generation H1 antihistaminics	
C	H2 receptor antagonists	
d	Proton pump inhibitors	
55	Identify the active form of 5-Fluorouracil	
a	5-FdUMP	
b	5-FUMP	
C	5-FUTP	
d	5-FdUTP	
56	Name functional group present in methimazole	
a	Thioamide	
b	Carboxamide	
C	ketone	
d	Primary amine	
57	Select the correct name and use of the drug.	

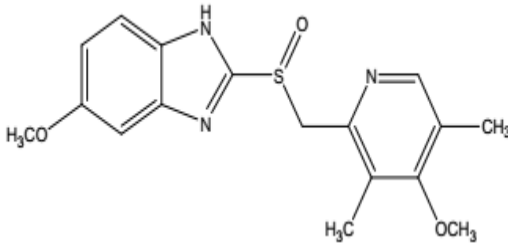


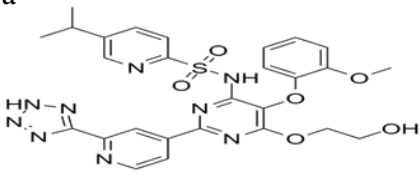
	a	Testosterone, androgen
	b	Mifipristone, oral contraceptive
	C	Tadalafil, used for erectile dysfunction
	d	Estradiol, estrogen
58		Which of the following antihyperlipidemic agent is a fibric acid derivative?
	a	Clofibrate
	b	Lovastatin
	C	Cholesteramine
	d	Cholestipol
59		To which Chemical Class does Warfarin belongs?
	a	Indanedione derivative
	b	Coumarin derivative
	C	Thienopyridine-class
	d	Naphthoquinone
60		Which form of thyroxine is biologically synthesized and is active
	a	levo

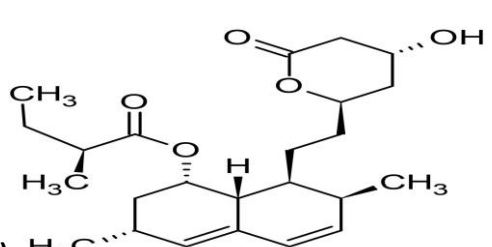
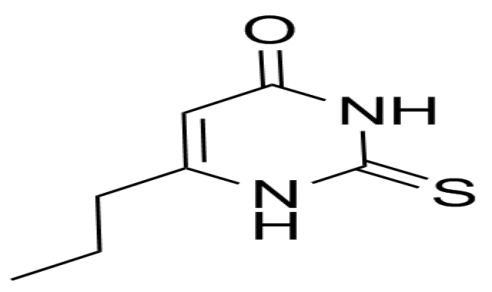
b	dextro	
C	meso	
D	racemic	

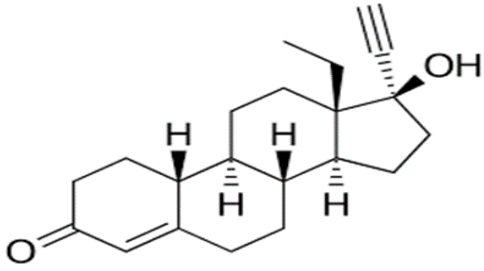
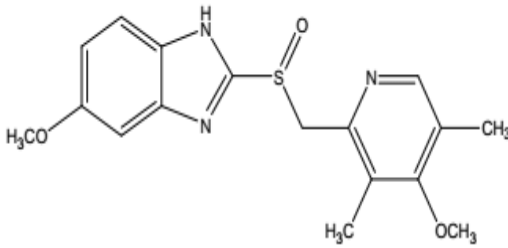
1	Give two examples (structures required) and the therapeutic indication of H1 antagonists using suitable examples and explain the mechanism of action of these compounds	4mks
2	Illustrate the mechanism of action of antimetabolites by giving suitable examples.	4mks
3	Outline synthesis and clinical use of Furosemide.	4mks
4	What are diuretics. Classify with examples. Discuss the SAR of Thiazide Diuretics.	4mks
5	Classify antiarrhythmic agents with examples and discuss Class III antiarrhythmic drugs.	4mks
6	Write a note on HMG-CoA reductase Inhibitor.	4mks
7	Draw the structures. Write the mechanism of action, uses and metabolism of Benzocaine, Procaine, Lidocaine and Prilocaine.	4 mks
8	Discuss SAR of Sulfonylureas by citing suitable examples.	4mks
9	Draw the structure of Testosterone and give its chemical name. What is the effect of the following structural changes on the activity i) Addition of methyl group at C-17 α ii) Removal of C-19 carbon	4mks

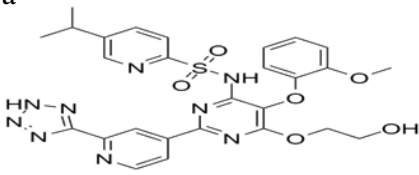
10	Answer the following with respect to given structure.	4mks

	a) Name this structure.	
	b) Comment on its mechanism of action.	
	c) Write its therapeutic use.	
	d) List the adverse reactions.	
11	Answer the following with respect to given structure.	4mks
		
	i) Name this drug.	
	ii) Comment on its mechanism of action.	
	iii) Write its therapeutic use.	
	iv) Comment on its metabolism.	
12	Outline chemical synthesis with reagents and reaction conditions for any one of the following.	4mks
	i) Meclorothamine	
	ii) Mercaptopurine	
13	Match the following	4mks

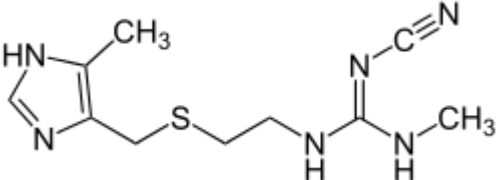
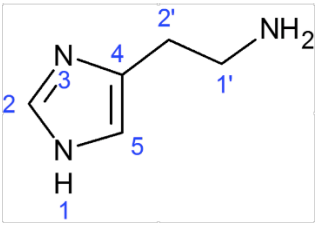
	<p>i. Digoxin</p> <p>ii . Nesiritide</p> <p>iii. Bosentan</p> <p>iv. Tezosentan</p>	<p>a</p>  <p>b. Endothelin-1 antagonists</p> <p>c. Cardiotonic glycosides obtained from Digitalis lanata</p> <p>d. Recombinant human B-type natriuretic peptide</p>	
14	Outline the synthesis of Nitroglycerin OR Isosorbide dinitrate with reaction conditions and necessary reagents and write its use.	4mks	
15	Answer the following.	4mks	
	i) Enlist the zinc-binding groups of ACE inhibitors.		
	ii) What happens when Esterification of carboxylate group of ACE inhibitors takes place.		
	iii) Write down the mechanism of action of ACE inhibitors.		
	What happens after introduction of large hydrophobic heterocyclic rings on ACE inhibitors.		
16	Write a note on stereochemistry and metabolism of steroids	4mks	
17	Answer the following.	4mks	
	What will happen		
	i) After introduction of unsaturation in 'B' ring of estrangement.		
	ii) Epimerisation of 17 β-OH group		
	iii) Removal of 3-OH group		
	iv) Removal of ring 'A'		
18	Match the following	4mks	

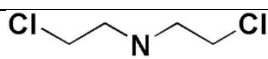
	i. Byguanides a. Tolbutamide	
	ii. Meglitinides b. Pioglitazone	
	iii. Thiazolidinediones c. Repaglinide	
	iv. Sulfonyl ureas d. Metformin	
19	<p>Identify the given structures and indicate their therapeutic use</p> <p>i)</p>  <p>ii)</p> 	4mks
20	Classify histamine H1 antagonist and comment on its structural requirements.	4mks
21	Classify anti-neoplastic agents with suitable examples .	4mks
22	Answer the following with respect to given structure.	4mks

		
a)	Name this structure.	
b)	Comment on its mechanism of action.	
c)	Write its therapeutic use.	
d)	List the adverse reactions.	
23	Answer the following with respect to given structure.	4mks
		
i)	Name this drug.	
ii)	Comment on its mechanism of action.	
iii)	Write its therapeutic use.	
iv)	Comment on its metabolism.	
24	Outline chemical synthesis with reagents and reaction conditions for any one of the following.	4mks
i)	Meclorothamine	
ii)	Mercaptopurine	
25	Match the following.	4mks

	<p>i. Digoxin</p> <p>ii . Nesiritide</p> <p>iii. Bosentan</p> <p>iv. Tezosentan</p>	<p>a</p>  <p>b. Endothelin-1 antagonists</p> <p>c. Cardiotonic glycosides obtained from Digitalis lanata</p> <p>d. Recombinant human B-type natriuretic peptide</p>	
26	Outline the synthesis of Nitroglycerin OR Isosorbide dinitrate with reaction conditions and necessary reagents and write its use.	4mks	
27	Answer the following.	4mks	
	i) Enlist the zinc-binding groups of ACE inhibitors.		
	ii) What happens when Esterification of carboxylate group of ACE inhibitors takes place.		
	iii) Write down the mechanism of action of ACE inhibitors.		
	iv) What happens after introduction of large hydrophobic heterocyclic rings on ACE inhibitors.		
28	Write a note on stereochemistry and metabolism of steroids	4mks	
29	Answer the following.	4mks	
	What will happen		
	i) After introduction of unsaturation in 'B' ring of estrangement.		
	ii) Epimerisation of 17 β-OH group		
	iii) Removal of 3-OH group		
	iv) Removal of ring 'A'		

30	Match the following	4mks
	i. Biguanides a. Tolbutamide	
	ii. Meglitinides b. Pioglitazone	
	iii. Thiazolidinediones c. Repaglinide	
	iv. Sulfonyl ureas d. Metformin	
31	Classify anti-neoplastic agents with suitable examples (Structures needed).	4mks
32	A. Illustrate the activation pathway for Pantoprazole. Identify its target enzyme and the type of interaction	4mks
33	B. State whether following statements are true or false giving a suitable explanation. i. Fibrates are chemically propanoic acid derivatives. ii. Amiodarone is class-I antiarrhythmic agent.	4mks
34	Outline synthesis and clonical use of Furosemide.	4mks

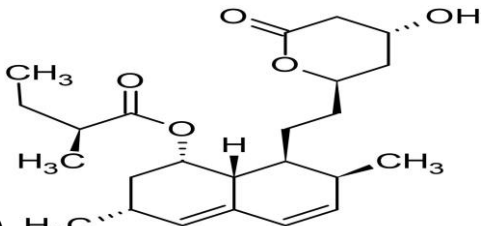
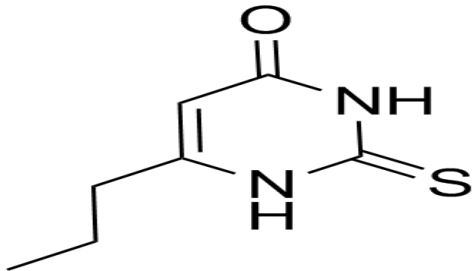
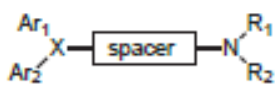
35	<p>Identify the given scaffold. Discuss the strategy that led to its rational development.</p> <div style="text-align: center;">  </div>	4mks
36	<p>Write a detailed classification of antiarrhythmic agents on the basis of mechanism of action giving one example (structures required) from each class. Describe the pH dependent effects of Class I antiarrhythmic agents</p>	4mks
37	<p>Answer following with respect to given structure.</p> <div style="text-align: center;">  </div> <p>a) Identify this structure.</p> <p>b) Comment on its activity at the H₁ and H₂ receptor.</p> <p>c) Predict the effect of the following structural changes on its activity:</p> <ol style="list-style-type: none"> i. Introduction of a CH₃ group at position 5. ii. Replacement of α-amino group by a guanidino group. 	4mks
38	<p>Discuss any two examples and uses of proton pump inhibitors (structures required). Explain the mechanism of action of these compounds.</p>	4 mks
39	<p>Based on the structure given below answer the following questions.</p>	4mks

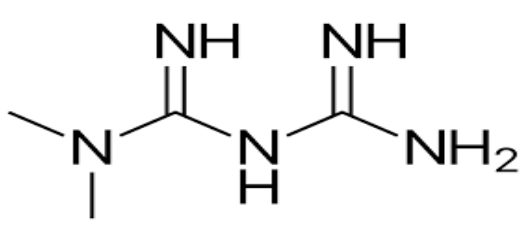


- i) Identify the drug if R=CH₃, draw its structure & give its uses. Also mention its mechanism of action.
- ii) Indicate the advantage of having an aromatic ring on nitrogen instead of methyl group in N-mustards.

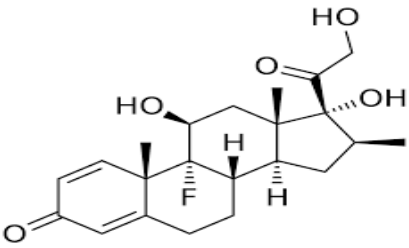
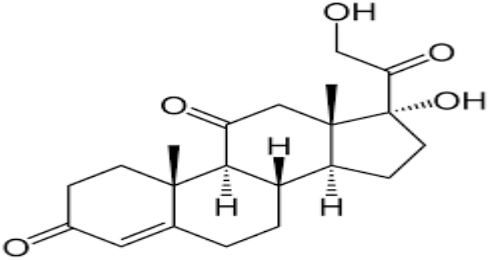
40	What is angina pectoris? Write chemical classification of antianginal class of drugs. Support your answer with relevant structure	4mks
41	What are diuretics? Classify diuretic drugs based upon their site of action. Give one example with structures from each class.	4mks
42	Answer the following	4mks
i)	Enlist any two binding interactions of ACE inhibitors with a target molecule	
ii)	ii. Why o,o dichloro substitution is necessary in clonidine? Justify.	
43	What is erectile dysfunction? Give an example of a pyrazolopyrimidinone containing drug which is used for treatment of erectile dysfunction and describe its mechanism of action.	4mks
44	Discuss the structural activity features required for local anaesthetics activity giving examples.	4mks
45	Outline synthesis of furosemide or acetazolamide with reaction conditions and necessary reagents write its use.	4mks
46	Outline the chemical synthesis of triprolidine with reagents and reaction conditions.	4mks
47	Write a note on stereochemistry and nomenclate the following	4mks

	<p>i) <chem>C[C@H]1CC[C@@H]2[C@@H](O)C(=O)C=C[C@@H]2[C@H]1F</chem></p> <p>ii) <chem>C[C@H]1CC[C@@H]2[C@@H](O)C(=O)C=C[C@@H]2[C@H]1CO</chem></p>	
48	Answer the following.	4mks
	What will happen	
i)	After introduction of unsaturation in 'B' ring of estrangement.	
ii)	Epimerisation of 17 β -OH group	
iii)	Removal of 3-OH group	
iv)	Removal of ring 'A'	
49	Match the following	4mks
	i. Biguanides a. chlorpropamide	
	ii. Meglitinides b. Pioglitazone	
	iii. Thiazolidinediones c. Nateglinide	
	iv. Sulfonyl ureas d. Metformin	
50		
51	Identify the given structures and indicate their therapeutic use	4mks

	<p>i)</p>  <p>ii)</p> 	
52	Classify histamine H1 antagonist and comment on its structural requirements.	4mks
53	Classify anti-neoplastic agents with suitable examples and add a note on mechanism of action of antimetabolites (Structures needed).	4mks
54	<p>Identify scaffold below and answer the questions</p>  <p>i) name and draw structure of any drug by substituting X with N</p> <p>ii) name of the prototype drug after replacing X with CHO</p> <p>iii) name the drug and give structure when X is replaced by CH</p>	4mks
55	Illustrate the mechanism of action of nitrogen mustard as an alkylating agent by giving suitable reactions pathway .	4mks
56	Outline synthesis and clonical use of Acetazolamide .	4mks
57	Classify diuretics . Outlie synthesis of chlorothiazide.	4mks

58	Classify antiarrhythmic agents with examples and discuss Class IB antiarrhythmic drugs for mechanism of action . Draw structure of any one.	4mks
59	Describe Mechanism of action of HMG CoA reductase inhibitors and describe SAR .	4mks
60	Classify local anaesthetic chemically with one structure of each class. Describe the mechanism of action of Benzocaine and lidocaine	4 mks
61	Discuss SAR of thiazolidinediones by citing suitable examples.	4mks
62	Draw the structure of estradiol and give its chemical name. What is the effect of the following	4mks
	i) Addition of hydroxyl group at 6,7 and 11	
	ii) substitution of 17 α with ethynyl group	
63	Describe Mechanism of action of Cardiac glycosides and cholestipol	4mks
64	Answer the following with respect to given structure. 	4mks
	i) Name this drug.	
	ii) Comment on its mechanism of action.	
	iii) Write its therapeutic use.	
	iv) Comment on its metabolism.	
65	Outline chemical synthesis with reagents and reaction conditions for any one of the following.	4mks
	i) Meclorothamine	

ii)	Mercaptopurine	
66	Match the following.	4mks
	i. Chlorpheniramine a. antimetabolite ii. Doxylamine b. Nitrogen mustard iii. Melphalan c. ethanolamine ether iv. Azathioprine d. Alkylamine group	
67	Outline the synthesis of Diisopyramide OR Warfarin with reaction conditions and necessary reagents and write its use.	4mks
68	Answer the following.	4mks
i)	Answer the following.	
ii)	Enlist the binding interaction of ACE inhibitors.	
iii)	state condition for optimum activity condition in respect to stereochemistry aspect .	
iv)	Write the mechanism of action of ACE inhibitors.	
69	Write a note on stereochemistry and metabolism of steroids	4mks
70	Answer the following.	4mks
	What will happen in progestins if	
i)	Removal of CH ₃ group at 19 position	
ii)	Unsaturation of B or C ring	
iii)	Introduction of C6 α with halogen	
iv)	Unsaturation at C6 and C7	

<p>71</p>	<p>Identify the given structures and comment on metabolism</p> <p>i)</p>  <p>ii)</p> 	<p>4mks</p>
<p>72</p>	<p>Describe SAR of local anaesthetic and comment on mechanism of action .</p>	<p>4mks</p>
<p>73</p>	<p>Classify anti-neoplastic agents with suitable examples (Structures needed).</p>	<p>4mks</p>