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DATE:

## TANTA UNIVERSITY FACULTY OF PHARMACY DEPARTMENT OF PHARMACEUTICAL CHEMISTRY



## FINAL EXAM FOR THIRD YEAR PHARMACY STUDENTS

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	03/06/2021	SECOND SEMESTER	TOTAL ASSESSMENT MARKS: 150	TIME ALLOWED: 120 MINUTES

## This exam booklet contains 14 pages and 100 MCQ questions (1.5 marks each). Fill the separate answer sheet using blue pen only for electronic correction machine.

1) Epinephrine is not released from peripheral sympathetic nerve endings. It is synthesized and stored in the adrenal medulla, from which it is released into the circulation

a) True

b) False

2) Isosorbide mononitrate is the only one capable of stimulating the production of coronary collateral circulation and the only one able to prevent experimental myocardial infarction in coronary occlusion

a) True

b) False

3) Ortho-nitrophenyl group of nifedipine is essential for calcium channel blocking activity

a) True

b) False

4) The chemical name of verapamil is 5-[(3,5-dimethoxyphenyl)methylamino]-2-(3,5-dimethoxyphenyl)- 2-isopropyl valeronitrile hydrochloride

a) True

b) False

$$H_3C$$
 $H_3C$ 
 $H_3C$ 
 $H_3C$ 
 $H_3C$ 
 $H_4$ 
 $H_4$ 
 $H_5$ 
 $H_6$ 
 $H_6$ 
 $H_7$ 
 $H_8$ 
 $H_8$ 

5) Compound I is

a) More active than nifedipine

- b) Less active than nifedipine
- c) Equal active as nifedipine
- d) Not active as calcium antagonist

6) Compound I is

a) Chiral molecule

b) Achiral molecule

- 7) II is used as a racemic mixture for treatment of congestive heart failure and hypotension
  - a) True

- 8) Which is not true concerning III?
  - a) It is a selective α<sub>1</sub>-agonist
  - b) It is an orally active, potent vasoconstrictor
  - c) It is not metabolized by MAO or COMT
  - d) It is used in the treatment of severe hypotension
  - e) None

$$H_3C$$
 $H_3C$ 
 $H_3C$ 

- 9) IV is Bepridil HCl
  - a) True

- b) False
- 10) Which is not true concerning methacholine?
  - a) It acts specifically on muscarinic receptors
  - b) It acts selectively on GIT
  - c) It is not easily hydrolyzed by cholinesterase enzyme
  - d) The (S) isomer is 240 times more active than the (R) one
  - e) None
- 11) V is
  - a) Terbutaline

b) Metaproterenol

c) Salbutamol

- d) None
- 12) Which is **not** true concerning indirect-acting sympathomimetics?
  - a) Clinically used are not catechols
  - b) β-hydroxyl group reduces effectiveness of indirect effect
  - c)  $\alpha$ -methyl group reduces effectiveness of direct effect
  - d) Nitrogen substitution reduces indirect effect
  - e) None
- 13) **VI** is
  - a) Digoxin
- b) Digitoxin
- c) None

- 14) Both (R)- and (S)-enantiomers of propafenone exert similar Na<sup>+</sup> channel-blocking effects, the (R)-enantiomer also produces a  $\beta$ -adrenergic blockade because of the correct configuration similar to (-)-(R)-epinephrine
  - a) True

- 15) Which drug is reserved for use in ventricular arrhythmias that are resistant to other therapies
  - a) Lidocaine

b) Procainamide

c) Phenytoin

d) Amiodarone

e) Bretylium tosylate

- 16)  $\alpha$ -Methylnorepinephrine acts on  $\alpha_2$ -adrenergic receptors to inhibit the release of NE, resulting in a decrease of sympathetic outflow from the central nervous system and an activation of parasympathetic outflow
  - a) True

b) False

- 17) Which is **not** true concerning propranolol?
  - a) It is an aryloxypropanolamine derivative
  - b) It is the first one used clinically
  - c) It is a nonselective  $\beta$ -adrenergic antagonist
  - d) It is used for treatment of angina, hypertension arrhythmia and in prophylaxis of migraine headache
  - e) None

- 18) The above equation shows the preparation of isosorbide mononitrate
  - a) True

b) False

- 19) The chemical name of diazoxide is 7-Chloro-2-methyl-2*H*-1,2,4-benzothiadiazine 1,1-oxide
  - a) True

b) False

- 20) Timolol is a nonselective  $\beta$ -blocker used for treatment of glaucoma and hypertension
  - a) True

b) False

21) VII is considered as DHP

a) True

b) False

22) 3-Methoxy-4-hydroxymandelic acid is the principal urinary metabolite of NE

a) True

b) False

$$H_2NO_2S$$
 $SO_2NH_2$ 
 $CI$ 
 $H_2NO_2S$ 
 $SO_2NH_2$ 
 $SO_2NH_2$ 

Chloraminophenamide

Dichlorphenamide

23) Which is **not** true concerning the above structures?

- a) Both chloraminophenamide and dichlorphenamide are active diuretics
- b) Chloraminophenamide is less active than dichlorphenamide as CAI
- c) Dichlorphenamide is more active than chloraminophenamide as CAI
- d) All
- e) None

24) Which ACE inhibitor does not have 2-(S)-aminophenylbutyric acid ethyl ester moiety

a) Lisinopril

b) Fosinopril

c) Enalapril

e) None

d) Ramipril

$$H_3C-N^+$$
  $O-C-NH_2 \cdot C1^ CH_3$ 

(VIII)

(IX)

- 25) VIII is nonselective muscarinic
  - a) Agonist
- b) Antagonist
- c) None
- 26) VIII has longer duration than ACh
  - a) True

- 27) IX acts as
  - a) α<sub>1</sub>-blocker

b) α<sub>2</sub>-blocker

c) \alpha\_1-agonist

d) None

- 28) The product of the above equation is
  - a) 2,4,7-triamino-6-phenylpteridine
  - b) 2,5,7-triamino-6-phenylpteridine
  - c) 3,4,7-triamino-6-phenylpteridine
  - d) 3,5,7-triamino-6-phenylpteridine
- 29) The chemical name of acetazolamide is N-(5-sulfamoyl-1,3,4-thiadiazol-2-yl)acetamide
  - a) True

b) False

- 30) X is
  - a) Indapamide

b) Xipamide

c) Chlorthalidone

- d) None
- 31) XI is an active diuretic similar to ethacrynic acid
  - a) True

- b) False
- 32) Hydralazine is usually used with other antihypertensive agent
  - a) True

b) False

	33) Both nitrovasodilators and poxide	phosphodiesterase inhibit	ors increase the level of nitric
	a) True		b) False
	34) Which drug was the first disable a) α-methyldopa	scovered α <sub>2</sub> -agonist? b) Clonidine	c) Guanabenz
	35) Salmetrol is used in the formal True	m of inhalation only for t	reatment of bronchial asthma b) False
	36) Naphazoline and xylometa: receptors	zoline are partial agonists	s at both $\alpha_1$ - and $\alpha_2$ -adrenergic
	a) True		b) False
37) Guanethidine, unlike reserpine has no CNS effects such as depression, beca the drug is highly polar and does not easily cross the blood-brain barrier a) True b) Faise			
38) α-Methyldopa lowers blood pressure by inhibiting the outflow of sympa vasoconstrictor impulses from the brain			ng the outflow of sympathetic b) False
a) True  39) Nitroglycerin is  a) Calcium channel blocker b) Nitrovasodilator c) Direct acting vasodilator d) Both (a) and (b) e) Both (b) and (c)  40) Which is <b>not</b> true for SAR of high-ceiling diuretics? a) The substituent at the 1-position must be acidic b) A sulfamoyl group in the 4-position is essential c) The "activating group" in the 5-position is essential d) Both (b) and (c) e) None		die dial	

41) Which of the following modifications will decrease the activity of benzodiazepines?

a) The chloride present at position (7)

b) The presence of (4,5) double bond

c) The presence of phenyl group at 5th position

d) The presence ethyl group at 3rd position

42) Which of the following benzodiazepines does not contain a carbonyl group at position 2?

a) Diazepam

b) Flurazepam

c) Chlordiazepoxide

d) Oxazepam

43) Which of the following is the active form of Clorazepate Dipotassium?

a) Nordazepam

b) Lorazepam

c) Oxazepam

d) Alprazolam

44) Which of the following is non-benzodiazepine hypnotics and has no affinity for the GABA receptor complex?

a) Flumazenil

b) Eszopiclone

c) Ramelteon

d) None

45) Which of the following is essential for sedative and hypnotic activity of barbiturates?

a) Alkylation at 1 or 3 position

b) Two alkyl groups at 5 position

c) Sulfur atom at 2 position

d) Hydrogen atom at 5 position

46) Which of the following is correct regarding the barbiturate hereunder?

a) It has a short duration of action

b) It has a long duration of action

c) It has an intermediate duration of action

d) It is inactive

47) Which of the following is a barbiturate with a short duration of action?

a) Thiopental sodium

b) Cyclobarbitone

c) Pentobarbital-Sodium

d) Mephobarbital

48) Which of the following drugs has the chemical name: 3-ethyl-3-phenylpiperidine-2,6-dione?

a) Glutethimide

b) Meprobamate

c) Methaqualone

d) Zaleplon

49) Which of the following inhalation anesthetics have chloride, bromide and fluoride atoms?

a) Chloral hydrate

b) Nitrous oxide

c) Methoxyflurane

d) Halothane

50) Which of the following drugs is an arylcyclohexylamine intravenous anesthetic?

a) Thiamylal sodium

b) Thiopental sodium

c) Ketamine HCl

d) Midazolam

51) Potency of phenothiazines is maximum when there are (5) carbons between two 'N' atoms?

a) True

b) false

52) Which of the following modifications will increase the activity of phenothiazines?

a) Introduction of methyl group at C-1 of alkyl side chain.

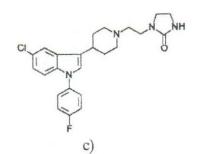
b) Oxidation of sulphur at fifth position.

c) Replacement of dimethylamino group by morpholine.

d) Introduction of oxygen into C-1 of alkyl side chain.

53) Which of the following drugs is atypical antipsychotic agent with partial agonist on specific dopamine and serotonin receptors while antagonizing other serotonin receptors?

b)



54) Which of the following is the mechanism of anticonvulsant activity of phensuximide?

a) Blocking sodium channels.

a)

b) Inhibiting gamma amino butyric acid transaminase enzyme.

c) Blocking T-type calcium channels.

d) GABA agonist.

55) Which of the following drugs belongs to oxazolidinedione anticonvulsants?

- 56) Lamotrigine, a 1,2,4-triazine derivative, is potent and long acting anticonvulsant, used as an add-on therapy for treatment of generalized seizures and acts as sodium channel blocker.
- a) True

- b) False
- 57) Which of the following antidepressant drugs is the product of the reaction hereunder?

a) Phenelzine

b) Meclobemide

c) Selegiline

- d) Tranylcypromine
- 58) Which of the following groups is not essential for activity of tricyclic antidepressant?
- a) A protonable nitrogen

b) Two aromatic rings

- c) The C10-C11 bridge
- d) Approximately a 3-carbon distance between the protonable nitrogen and aromatic ring
- 59) Which of the following antidepressant drugs is selective serotonin reuptake inhibitor and has anxiolytic activity?
- a) Maprotiline Hydrochloride
- b) Fluoxetine

c) Nisoxetine

- d) Venlafaxine
- 60) COX-2 is responsible for the production of prostaglandins at the inflammation site?
  - a) True

- b) False
- 61) Which of the following modifications could abolish activity of salicylates as antiinflammatory drug?
- a) Salt of salicylic acid with choline or magnesium.
- b) Introduction of (F) at 5 position.
- c) The presence of phenolic hydroxyl group at ortho position to the carboxyl group.
- d) Placement of the phenolic hydroxyl group at meta or para to the carboxyl group.

- 62) Which of the following is essential for the activity of the 3,5-pyrazolidinedione?
- a) Replacement of one of the nitrogen atom in the pyrazolidinediones with an oxygen atom.
- b) The dicarbonyl function at the 3rd and 5th positions.
- c) m-Substitution of aryl rings with a methyl or chloride.
- d) 4, 4-dialkyl derivatives.
- 63) Which of the following drugs is an anthranilic acid derivative?
- a) Flufenamic Acid

b) Phenylbutazone

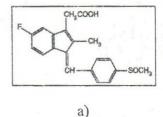
c) Paracetamol

d) Phenacetin

- 64. The presence of indole ring nitrogen is essential for activity of indole acetic acid derivatives.
- a) True

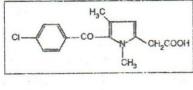
b) false

65. Which of the following is zomepirac?



CH<sub>3</sub> CH<sub>2</sub>COÖNÉ

b)



c)

- 66) Which of the following is the most active isomer of ibuprofen?
- a) The (s)-(-) isomer

b) the (s)-(+) isomer

c) The (R)-(+) isomer

d) the (R)-(-) isomer

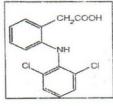
- 67) Which of the following drugs is a cyclizd hetroarylpropionic acid derivative with the alpha methyl group being fused to the pyrrole ring and is accepted as alternative to narcotic analgesics?
- a) Diclofenac

b) Naproxen

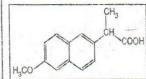
c) Ketoprofen

d) Ketorolac

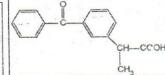
68) Which of the following drugs possesses structural characteristic of both arylalkanoic acid and the anthranilic acid classes?

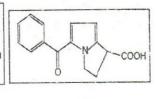


a)



b)



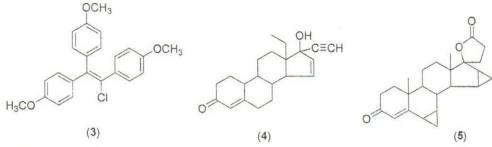


c)

d)

69) The introduction of heterocyclic ring in the amide chain significantly increases the anti- inflammatory activity of oxicams.					
a) True	b) false				
70) Which of the following drugs is a prodrug, second-generation COX-2 inhibitor developed for injectable use and is rapidly converted by hepatic enzymes to the active form?					
a) Valdecoxib	b) Celecoxib				
c) Parecoxib	d) Nimesulide				
71) Which of the following is correct regarding the drug hereunder?					
NH:	SO <sub>2</sub> CH <sub>3</sub>				
<ul><li>a) It contains a sulphonamide moiety as</li><li>b) It exhibits significant selectivity towa</li><li>c) It is a prodrug.</li><li>d) It is nimesulide</li></ul>	basic group, ards COX-1.				
72) Which of the following classes contain the endorphins?					
a) Natural opiates	b) Semi-synthetic opiates				
c) Endogenous opioid peptides	d) Fully- synthetic opioids				
73) Which of the following is essential for activity of morphine?					
a) Alcoholic OH	b) The phenolic OH				
c) Double bond between C7 & C8	d) The ether bridge				
74) Which of the following makes morphina	as more potent agonists and antagonists?				
a) Removal of ring D	b) The phenolic OH				
c) The introduction of a hydroxyl group at position 14					
d) Removal of ring E					
75) Which of the following opiates does not have rings C and D?					
a) Heroine	b) Nalorphine				
c) Oxymorphine	d) Benzomorphans				
76) Which of the following is true about structure of methadone?					
a) Does not have rings B, C and D	b) Does not have rings B, C and E				
c) Does not have rings B and D	d) Does not have rings B, D and E				

- 77) Estrogens reduce bone resorption and increase bone formation a) True b) False 78) Males also possess estrogen receptors and estrogen to some extent and levels in the male blood are higher than post-menopausal women. a) True b) False 79) To prevent oxidation of hydroxyl group at C17 in estradiol to be taken orally, we should: a) Replace the hydrogen atom with ethinyl (alkyne) group. b) Convert the cyclopentyl ring to phenyl ring. c) Block OH by esterification. d) All of the above. (1) 80) Compound 1 is a) Mestranol b) Quinestrol c) Estradiol 3-benzoate d) Estradiol Cypionate 81) Compound 2 is a) Mestranol b) Quinestrol c) Estradiol 3-benzoate d) Estradiol Cypionate 82) To improve oral absorption of Ethinyl Estradiol, make a) Etherification at OH group at position 3 b) Esterification at OH group at position 3 c) Oxidation CH group at position 3 d) Alkylation at OH group at position 3 83) To slower rate of estradiol release for depot IM preparation, make the esterification at OH group at position 3 or 17. a) True b) False 84) Letrozole inhibits the conversion of estradiol to androstenedione and
- 84) Letrozole inhibits the conversion of estradiol to androstenedione and testosterone
  - a) True b) False
- 85) Clomiphene can be synthesized from acetophenone derivative
  a) True
  b) False



- 86) Compound 3 acts as
  - a) Antiestrogenic agent
  - c) Progesterone antagonist
- b) Non-steroidal Estrogenic agent
- d) Aromatase Inhibitors

- 87) Compound 4 is
  - a) Mestranol
- b) Quinestrol
- c) Estradiol 3-benzoate
- d) Gestodene
- 88) Compound 5 is
  - a) Mestranol
- b) Quinestrol
- c) Drospirenone
- d) Gestodene
- 89) Tamoxifen acts as antiestrogenic agent by:
  - a) Block Estrogen receptor
  - b) Inhibit Aromatase enzyme
  - c) Block Progesterone receptor
  - d) Kills sperm or stops it from moving
- 90) Physiological function(s) of progesterone is (are)
  - a) increase thickness of endometrium of uterus.
  - b) desensitization of uterus to oxytocin
    - c) decrease the sperm motility
    - d) all of the above
- 91) Norethistrone can be characterized by
  - a) Has moderate to high potency and has androgenic action
  - b) It is progesterone derivatives
  - c) Has high potency with anti-mineralocorticoid action
  - d) It is Spironolactone derivatives
- 92) If progesterone is taken orally, it is metabolized by reduction to inactive metabolite that is called:
  - a) Dydrogestrone

- b) Prognanediol
- c) Medroxy-progesterone
- d) Norgestimate
- 93) Medroxy-progesterone can be taken orally by:
  - a) Remove two hydrogen atoms at C6 & C7 of progesterone

b)	Add methyl gp at C6 and ester gp at C17 of progesterone			
c)	c) Add methoxy gp at C6 and ester gp at C17 of progesterone			
d)	Replace hydrogen atoms at C6 & C7 of prog	gesterone with halogens		
94) Mifepr	ristone:			
a)	Used as abortifacient			
b)	Decrease endometrium thickness of uteru motility.	as and increase the uterus		
c)	Block progesterone receptor			
d)	All of the above			
	tin component of the Yasmin® is			
	Drospirenone			
b)	Levonorgestrol			
c)	Dydrogestrone			
d)	Lynestrenol			
96) Nonox	xynol-9 act as:			
a)	Male hormonal contraceptive			
b)	Spermicides			
c)	Mono-phasic oral contraceptive pills			
d)	Bi-phasic oral contraceptive pills			
	phasic oral contraceptive pills contain a constin in each active pill throughout entire cycle			
	a) True	b) False		
	ubstitution of position 16 with hydroxyl kable decrease in activity as in	group of estradiol lead to		
a)	Estriol	b) Estrone		
, c)	Ethinyl Estradiol	d) Estradiol 3-benzoate		
99) Dydro	gestrone can be characterized by			
a) Has moderate to high potency and has androgenic action				
b)	It is progesterone derivatives			
c)	Has high potency with anti-mineralocortico	oid action		

d) It is Spironolactone derivatives

a) True

100) Phenolic moiety at ring A in estradiol is an important in activity.