

Total No. of printed pages = 4

**BP 404T**

Roll No. of candidate

W16/24

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Hatkhowapara, Azara, Ghy-17

2024

**B.Pharm. 4<sup>th</sup> Semester End-Term Examination**

**PHARMACOLOGY – I**

Full Marks – 75

Time – Three hours

The figures in the margin indicate full marks for the questions.

**Section A**

**1. Multiple Choice Questions :**

(20 × 1 = 20)

- (i) Bioavailability is highest in following routes of drug administration
  - (a) Subcutaneous
  - (b) Intravenous
  - (c) Intramuscular
  - (d) Intradermal
- (ii) Following factors influence absorption and bioavailability
  - (a) Particle size
  - (b) Degree of Ionization
  - (c) Food
  - (d) All the above
- (iii) A constant fraction (Percentage) of drug is eliminated in constant interval of time in following elimination kinetics
  - (a) First order
  - (b) Zero order
  - (c) Mixed order
  - (d) None of above
- (iv) Following is a second messenger for GPCR
  - (a) cAMP
  - (b) ATP
  - (c) Ion channel
  - (d) Tyrosine kinase
- (v) Therapeutic index should be
  - (a) More than 1
  - (b) Equal to 1
  - (c) Less than 1
  - (d) None of above

**[Turn over**

- (vi) Following is the example of microsomal enzyme
- Monoamine oxidase
  - Amidase
  - Transferase
  - Cytochrome p450
- (vii) Following is the compartment/reservoir for distribution of drugs in our body
- Cell
  - Fat
  - Plasma protein
  - All the above
- (viii) In voltage operated ion channel
- Ion conductance is regulated by a drug
  - Ion conductance is regulated by alteration in voltage gradient across plasma membrane
  - Agonist open the channel
  - Antagonist prevent the channel from opening
- (ix) Neurotransmitter at the sympathetic ganglia is
- Noradrenaline
  - Acetylcholine
  - Histamine
  - Serotonin
- (x)  $\alpha_2$  - agonist
- Increase release of Nor adrenaline
  - Decrease release of Nor adrenaline
  - Increase release of Dopamine
  - None of above
- (xi) The anxiolytic which is a potent enzyme inducer and exhibit more drug interactions
- Diazepam
  - Phenobarbital
  - Zolpidem
  - Buspirone
- (xii) Majority of antipsychotic drugs act by
- Blocking dopamine receptors
  - Activating dopamine receptors
  - Blocking muscarine receptors
  - Activating nicotine receptor
- (xiii) Following is the excitatory neurotransmitters
- NMDA
  - GABA
  - Glycine
  - None of above

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- (xiv) Dissociative anesthesia is produced by
- |              |                |
|--------------|----------------|
| (a) Ketamine | (b) Propofol   |
| (c) Fentanyl | (d) Thiopental |
- (xv) Following is a respiratory stimulant
- |                |                |
|----------------|----------------|
| (a) Picrotoxin | (b) Doxapram   |
| (c) Strychnine | (d) Bicuculine |
- (xvi) Following are the pharmacological actions of alcohol
- |                                       |
|---------------------------------------|
| (a) Decrease myocardial contractility |
| (b) Diuresis                          |
| (c) Cutaneous vasodilation            |
| (d) All the above                     |
- (xvii) Following is a drug for glaucoma that blocks acetylcholine esterase enzyme
- |                   |                   |
|-------------------|-------------------|
| (a) Physostigmine | (b) Timolol       |
| (c) Pilocarpine   | (d) Acetazolamide |
- (xviii) "Side effects" are coming under following type of ADR
- |       |       |
|-------|-------|
| (a) A | (b) B |
| (c) C | (d) D |
- (xix) Atracurium works by
- |                                  |
|----------------------------------|
| (a) Blocking GABA receptor       |
| (b) Blocking $N_M$ receptor      |
| (c) Blocking $\alpha_2$ receptor |
| (d) Blocking $M_1$ receptor      |
- (xx) Morphine causes
- |                            |                          |
|----------------------------|--------------------------|
| (a) Miosis                 | (b) Suppression of cough |
| (c) Respiratory depression | (d) All the above        |

#### Section B

2. Short Type Questions (Answer any Seven): (5 × 7 = 35)
- |  |
|--|
| (a) What are the mechanisms of bio transportation? Write in brief about carrier mediated transport system. |
| (b) Explain the synthesis, release and degradation of acetylcholine.                                       |
| (c) Define Competitive and Noncompetitive antagonism.  |
| (d) Classify antipsychotics and explain their MOA.   |

- (e) Define sedatives and hypnotics. Explain the mechanism of Barbiturates with ADR.
- (f) Explain in brief about Drug-Drug Interactions.
- (g) Classify antiadrenergic drugs and mention a few therapeutic applications.
- (h) Write a short note on drug dependence and drug abuse.
- (i) What are the different types of seizures? Write in brief about Phenytoin including MOA and ADR.

### Section C

#### 3. Long Type Questions (Answer Any Two)

- (a) Explain in details about G-protein coupled receptor and intracellular receptor. (2 × 10 = 20)
- (b) Define local anaesthetics. Classify local anaesthetics with mechanism of action and ADR. (7+3 = 10)
- (c) Explain the classic symptoms of Parkinsons Disease. Classify antiparkinson drugs with Mechanism of action and ADR. (2+8 = 10)