| | Utech |
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| Name: | |
| Roll No.: | A Grand (V Knowledge Stad Excellent) |
| Invigilator's Signature : | |

PHARMACEUTICAL CHEMISTRY (MEDICINAL CHEMISTRY)

Time Allotted: 3 Hours Full Marks: 70

The figures in the margin indicate full marks.

Candidates are required to give their answers in their own words as far as practicable.

GROUP - A

(Multiple Choice Type Questions)

- 1. Choose the correct alternatives for any ten of the following: $10 \times 1 = 10$
 - i) Which one of the following G-protein increases adenylyl cyclase activity and increases Ca-21 currents?
 - a) Gs

b) Gi

c) Gq

- d) Go.
- ii) Which of the following ganglionic blockers contains quaternary ammonium groups?
 - a) Pempidine
- b) Nicotine
- c) Mecamylamine
- d) Pentolinium.

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- iii) Which one of the following is the prodrug of Epinephrine, which is particularly useful for open angle glaucoma?
 - a) Phenylephrine
 - b) Norepinephrine
 - c) Dipivefrin
 - d) Oxidative metabolite of epinephrine.
- iv) Which of the following pairs is the starting materials for the synthesis of Propranolol ?
 - a) Benzene with cyclohydrine
 - b) 1-naphthol with chlorohydrine
 - c) 1-naphthol with NaOH
 - d) 1-naphthol with epichlorohydrin.
- v) To synthesize diazonium salt the starting material required is
 - a) primary amine
 - b) secondary amine
 - c) tertiary amine
 - d) any type of amine.

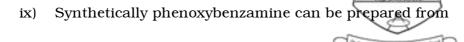
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- vi) In the case of nomenclature of heterocyclic compounds the preference is made on which of the following basis?
 - a) P > O > S > N > Si
- b) O > N > S > P > S
- c) O > S > N > P > Si
- d) O > Si > P > N > S.
- vii) Intracellular storage house of calcium ion is
 - a) endoplasmic reticulum
 - b) sarcoplasmic reticulum
 - c) golgi apparatus
 - d) none of these.
- viii) g-protein coupled receptor is known as
 - a) 7-transmembrane receptor
 - b) metabotrophic receptor
 - c) both (a) and (b)
 - d) nuclear receptor.

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- a) Phenol and propylene oxide
- b) Phenol and *p*-chlorobenzyl chloride
- c) Phenol and trichloropropane
- d) Phenol and 2-chloro-1-methyl propane.
- x) Which one of the following drugs is used for the organophosphate toxicity?
 - a) 2-pyridine aldoxime methyl chloride
 - b) 2-pyridine aldoxamine methyl chloride
 - c) 2-piperidine aldoxime methyl chloride
 - d) 2-pyrimidine aldoxime methyl chloride.
- xi) Thiazole ring containing H $_{2}$ blocker is
 - a) Ranitidine
- b) Nizatidine
- c) Cimetidine
- d) Burimamide.

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- xii) The starting material in the biosynthesis of Histamine is
 - a) Histidine
- b) 5-Hydroxy tryptamine
- c) Homocystaine
- d) Homovaline.
- xiii) Ergot alkaloid contains which of the following heterocyles?
 - a) Indene

b) Indoline

c) Indole

d) none of these.

GROUP - B

(Short Answer Type Questions)

Answer any *three* of the following.

 $3 \times 5 = 15$

- 2. a) What is the mechanism of action of \mathbf{H}_1 receptor antagonists?
 - b) What are the major side effects and use of \mathbf{H}_1 receptor antagonists ?
 - c) Give the synthesis of any one H_1 receptor antagonist.

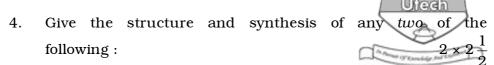
2 + 1 + 2

3. What are irreversible acetyl cholinesterase inhibitors? What is the significance of 2-PAM?

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- a) Prazocin
- b) Indomethacin
- c) Atenolol
- d) Diphenhydranin
- e) Propanolol.
- 5. Explain the cycloxygenase pathway.
- 6. Write a note on β -adrenergic blockers.

GROUP - C

(Long Answer Type Questions)

Answer any *three* of the following. $3 \times 15 = 45$

- 7. a) Classify sympatholytic agents.
 - b) Write down the SAR for adrenergic receptor agonist.
 - c) Sketch the synthesis procedure on any *two* of the following drugs:
 - (i) Clonidine
 - (ii) Salbutamol
 - (iii) Guanethidine
 - (iv) Isoprenaline. $4 + 5 + (2 \times 3)$

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- 8. a) Describe Hansch analysis in brief.
 - b) Write down the methodology in molecular modelling for drug molecules.
 - c) Mention and discuss steps involved in QSAR based drug design. 5 + 5 + 5
- 9. a) Write down the chemical classification of NSAIDs.
 - b) Write down the SAR of anthranilic acid derivatives.
 - c) Write down the synthesis of any *two* of the following :
 - (i) Ibuprofen
 - (ii) Diclofenac sodium
 - (iii) Phenyl butazone.

3 + 6 + 6

- 10. a) Define drug potency.
 - b) With the help of dose-response curve explain the dose-response relationship with example.
 - c) Derive the Hill-Langmuir equation to establish the drugreceptor relationship. 2 + 5 + 8
- 11. a) Write down the metabolic pathway of catecholamines and identify the major metabolite.
 - b) Explain the depolarizing blocking mechanism of (+) tubocurarine with its structure.
 - c) Briefly explain the transducer mechanism for IP3 DAG pathway of GPCR. 5+5+5

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