BIRLA INSTITUTE OF TECHNOLOGY, MESRA, RANCHI (END-SEMESTER EXAMINATION)

CLASS: PHARMACY

BRANCH: PHARMACY

SEMESTER: IV

SESSION: SP/2019

SUBJECT: BP402T MEDICIANAL CHEMISTRY I

TIME: 3.00 Hours FULL MARK: 75 INSTRUCTIONS:

1. The missing data, if any, may be assumed suitably.

- 2. Before attempting the question paper, be sure that you have got the correct question paper.
- 3. Tables/Data hand book/Graph paper etc. to be supplied to the candidates in the examination hall.
- 4. This question paper consists of (03) three parts. Read the part wise instructions before attempting the questions.

PART-I

Objective types questions (Instruction: Answer all questions)

Q1. $(4 \times 5 = 20 \text{ Marks})$

A. Fill in the blanks

- (i) Root of used since ancient times for the treatment of dysentery in Brazil
- (ii) Type of adrenergic receptors found in cardiac muscle is
- (iii) Torsion angle between quaternary nitrogen and esteric oxygen in anticlinal conformation of acetylcholine is
- (iv) is a benzodiazepine with anticonvulsant activity.
- (v) Inhalational anaesthetic has good skeletal muscle relaxant property

B. State True of False

- (i) First pharmacopoeia has the monograph of Ephedrine
- (ii) Nor-epinephrine is biosynthesised from tryptophan
- (iii) Carbachol is slowly hydrolysed by acetyl cholinesterase
- (iv) Sodium valproate acts as anticonvulsant by inhibiting Na-channel
- (v) Aspirin is an ester of acetic acid.

C. <u>Match</u> the following (Single answer)

- (i) Diethyl stilbosterol
- (ii) Selective Beta-1 blocker with shortest half-life
- (iii) Pralidoxime
- (iv) Ultra-short acting barbiturate
- (v) Dissociative anaesthesia

- (a) Cholinesterase reactivator
- (b) Ketamine (v)
- (c) Thiopental sodium
- (d) Esmolol
- (e) Geometrical isomer

D. Identify the following

- (i) Draw the structure of Methotrexate and identify the bioisosteric replacement in it with reference to substrate
- (ii) Identify the drug from its IUPAC nomenclature and draw its chemical structure: 1-(naphthalen-1-yloxy)-3-[(propan-2-yl)amino]propan-2-ol
- (iii) Identify the drug from its structure and write its IUPAC nomenclature:

- (iv) Draw the structure of Chlorprothixene and identify the basic nucleus in it.
- (v) Identify the drug and draw its structure: It is an NMDA inhibitor administered parenterally for producing anaesthesia

PART-II Short Answers

(Instruction: Answer seven out of nine questions)

 $(7 \times 5 = 35 \text{ Marks})$

- Q2. <u>List</u> the factors influencing the aqueous solubility of the drug molecule. <u>Define</u> Polymorphism. <u>Name</u> two drugs that exists in different polymorphic forms. <u>Summarize</u> the effect of polymorphism on drug solubility with one example.
- Q3. <u>Identify</u> the structural components responsible for activity, selectivity and metabolic stability With the general structure for Adrenergic agonists.
- Q4. <u>Design</u> a general synthetic pathway for the synthesis of phenylethylamine class of adrenergic agonists/given below

- Q5. <u>Analyse</u> the structural features required for designing a anticholinesterase inhibitors by considering the structure of Carbidopa and Neostigmine.
- Q6. Construct the synthetic route for anticholinesterase inhibitor starting from 3-amino phenol.
- Q7. Give an account of Structure activity <u>relationship</u> of benzodiazepine class of sedatives and hypnotics
- Q8. <u>Elaborate</u> the <u>synthesis</u> of drug that falls under dibenzazepine class and used in the treatment of trigeminal neuralgia
- Q9. What is meant by pre-anaesthetic medication? Why it is required? <u>List</u> the class of drugs used in pre-anaesthetic medication with suitable example.
- Q10. <u>Elaborate</u> the synthesis of Halothane, write its IUPAC nomenclature and Adverse effects if any.

PART-III Long Answers (Instruction: Answer two out of three questions)

 $(2 \times 10 = 20 \text{ marks})$

- Q11. Present a short <u>survey</u> on optical isomerism in drugs and their influence on pharmacokinetics and pharmacodynamics of drug molecules.
- Q12. <u>Discuss</u> in detail the mechanism of hydrolysis of acetylcholine by acetylcholinesterase and chemistry of organophosphorus poisoning.
- Q13. Elaborate the synthesis of the following: (i) Fentanyl and (ii) Mefenamic acid

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