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MPC 103T

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2021

M.Pharm. 1st Semester End-Term Examination

ADVANCED MEDICINAL CHEMISTRY – I

(New Regulation)

Full Marks – 75

Time – Three hours

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

Answer question No. 1 and any ~~seven~~ from the following.

1. (A) Multiple choice questions : (5 × 1 = 5)

- (i) Which of the following is a H3 receptor blocker
- (a) Pitolisant
 - (b) Ranitidine
 - (c) Cimatidine
 - (d) All of the above
- (ii) Which one of the following is H1 receptor antagonist Ethylene diamine derivatives?
- (a) Pyrilamine
 - (b) Diphenhydramine
 - (c) Pheneramine
 - (d) Promethazine

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(iii) Which of the following NSAIDs has greater inhibitory selectivity for COX-1 than COX-2?

- (a) Flurbiprofen
- (b) Indomethacin
- (c) Diclofenac
- (d) Celecoxib

(iv) Which is a property of eicosanoids?

- (a) All eicosanoids contain three conjugated double bonds
- (b) All eicosanoids contain arachidonic acid and sphingosine
- (c) Prostaglandins and leukotrienes both contain a ring structure
- (d) Thromboxanes and prostaglandins both contain a carboxyl group

(v) Which of the following is GABA analogue?

- (a) Progabide
- (b) Diazepam
- (c) Clonazepam
- (d) Phenobarbital

2. Describe the stages of drug discovery in details. (10)
3. Classify receptors and discuss elaborately about G-Protein coupled receptor and Enzyme linked receptors. (2 + 4 + 4 = 10)
4. Write a note on Enzyme mimics and bio precursors. (5 + 5 = 10)
5. Give schematic illustration of the prodrug concept. Explain the application of Prodrug. (4 + 6 = 10)
6. Explain the role of chirality in selective and specific therapeutic agents. (10)
7. Discuss the chemistry of prostaglandins and thromboxanes. (5 + 5 = 10)
8. Define Analogue design of drug. Differentiate classical and non-classical Bioisosteres. Briefly discuss about Bioisosteric replacement. (1 + 3 + 6 = 10)

9. Write down the synthesis of following drugs : (any five) (5 × 2 = 10)
- (a) Promethazine
 - (b) Prazosin
 - (c) Clonidine
 - (d) Hydralazine
 - (e) Diazepam
 - (f) 5-fluoro uracil
10. Explain the structure activity relationship of Hydantoins, Nitrogen mustards. Write a note on COX-I and COX-II inhibitors citing example with structure. (5 + 5 = 10)
11. Discuss the Peptidomimetic drug design principle. Write down the recent advances in peptidomimetics. (6 + 4 = 10)
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