## Medicinal alemostry - III (REVISED COURSE).

Sem VIII (P)

RV-4664 (2 Hours) [Total Marks: 40 N.B.: (1) Question No. 1 is compulsory. Attempt four more questions from the remaining six. Write the structure, generic name and major therapeutic use for the following (any three):-(i) Ethyl 2 -(4 - chlorophenoxy) - 2- Methylpropanoate (ii) 1, 2, 3, 4 - tetrahydro - 9 - aminoacridine (iii) 4' - [1- hydroxy - 2 - (isopropylamino) ethyl] Methylsulfonamide (iv) N, 2, 3, 3 - tetramethyl - 2 - norbornanamine hydrochloride Write the structure, generic name and major therapeutic use of the drugs with 2 the following description (any two) :--(i) A direct acting vasodilator (ii) An adrenergic neuron blocking agent containing guanidine. A depolarising neuromuscular blocking agent of the bis quaternary ammonium compound class. Explain the following statements. Support your answer with relevant structures :-(a) Pralidoxime is ineffective if administered 36 hours after the exposure to insecticide. (b) Succinylcholine chloride produces short duration of neuromuscular blockade. (c) Enalapril, a prodrug, has a better bioavailability as compared to Enalaprilat. (d) Replacement of catechol ring by resorcinol imparts oral bioavailability to  $\beta$  – agonists. Give schematic synthesis of the following, specifying the names of the reactants 6 and reaction conditions (any two) :--(i) Captopril (ii) Propranolol (iii) Nifedipine. Write the structure of one active metabolite of each of the following:---Lovastatin (ii) Losartan. Give schematic metabolism of the following drugs and label the metabolites as 4 active / inactive (any two) :--(i) Captopril (ii) Verapamil (iii) Gemfibrozil. Classify antiarrhythmic drugs on the basis of their mechanisms of action. 4 Give examples of drugs belonging to each subclass. Explain the structure Activity Relationship of Angiotensin - II Receptor antagonists. Support your answer with relevant structures. Explain the development of  $\beta$  – blockers from dichloroisoproterenol. Discuss the similarities between aryloxypropanolamines and arylethanolamines. 6. (a) Discuss the Mechanisms of action of — (i) HMG-CoA Reductase Inhibitors (ii) Organic nitrates. Explain why acetylcholine is a poor therapeutic agent. Outline the structural 4 modifications in acetylcholine which resulted in therapeutically useful drugs with agonistic activity at the muscarinic receptor. Write short notes on any two of the following:-(a) Alpha adrenergic antagonists

Angiotensin Converting Enzyme Inhibitors

(c) Alzheimer's Disease and its Treatment.