

Pharmacology Examination Topics

1st semester

1. Pharmacodynamic principles. Receptors and subtypes
 - General description of parasympathetic nervous system from pharmacological point of view (neurotransmitters and receptors)
 - Antihypertensive mode of action of thiazide diuretics and the side effects, osmotic diuretics
2. Dose-response relationships. Efficacy and potency
 - Directly acting parasympathomimetics
 - Calcium channel blockers
3. Graded and quantal dose-response relationships. Therapeutic index, therapeutic window
 - Parasympatholytics
 - Centrally acting sympathoplegic drugs
4. Agonists and antagonists. Antagonism on the receptor level
 - Sympathomimetics
 - Pharmacology of renin/angiotensin system
5. Antagonism. Non-receptorial antagonism
 - Non-selective α -adrenoceptor blockers
 - General description of antiarrhythmic drugs. Vaughan Williams classification
6. Control of receptor expression. Receptor diseases and receptors and disease
 - β -adrenoceptor blockers
 - Treatment of myocardial ischemia especially the treatment of angina pectoris
7. Desensitization, tachyphylaxis and tolerance
 - Indirectly acting parasympathomimetics
 - Drugs used in the treatment of hyperlipidemias
8. The movement of drugs through biological membranes
 - Structure-activity relationships demonstrated among sympathomimetics
 - Drugs used for the treatment of congestive heart failure
9. Distribution of drugs in the body: the apparent volume of distribution (V_d)
 - General description of sympathetic nervous system from pharmacological point of view (neurotransmitters and receptors)
 - Characterization of quinidine, lidocaine, and amiodarone
10. Elimination of drugs: the half-life ($T_{1/2}$)
 - Pharmacological tools to influence the sympathetic neurotransmission
 - Expectorants and antitussives
11. The clearance
 - Selective α -adrenoceptor blockers
 - Pharmacology of the liver and the gall bladder
12. Plasma concentrations after repeated administration: loading dose and maintenance dose
 - Metabolism of catecholamines and pharmacological modulation
 - Pharmacological treatment of bronchial asthma
13. Absorption of drugs and ion trap
 - Comparison of elimination of acetylcholine (ACh) and norepinephrine/noradrenaline from the synaptic cleft and the possibilities of pharmacological modulation
 - Therapeutic importance of diuretics, mode of action and classification. Antialdosterone compounds and other potassium-sparing diuretics

14. Bioavailability. AUC
 - Compare the effects of norepinephrine/noradrenaline, epinephrine/adrenaline and isoprenaline
 - Inhibitors of carboanhydrase enzyme, thiazides and other sulfonamide type diuretics, high-ceiling diuretics (loop diuretics) and antidiuretics
15. First pass effect
 - Synthesis, storage, release and elimination of acetylcholine (Ach). Demonstration of Dale's experiment
 - Agents used in anemias
16. Drug elimination: I. Biotransformation
 - Non-adrenergic, non-cholinergic (NANC) transmission
 - Drugs used in coagulation disorders
17. Factors influencing the drug elimination
 - Uptake mechanisms, substrates and inhibitors
 - Drugs used in acid-peptic disease
18. Drug elimination: II. Excretion
 - α_2 sympathomimetics and the concept of "false transmitter"
 - Laxatives, antidiarrheal drugs. Drugs in the treatment of chronic inflammatory bowel disease, antiobesity drugs
19. Factors influencing the drug effect. Preclinical phase of drug development
 - Pharmacology of cardiac glycosides
 - Drugs promoting gastrointestinal motility. Emetics and antiemetic drugs
20. Drug interactions. Biologicals (biological therapy), special considerations with respect to their development.
 - Positive inotropic substances except cardiac glycosides
 - Pharmacotherapeutic approach to exocrine pancreatic diseases
21. Clinical phase of drug development.
 - Adrenergic neuron blockers and reserpine. Antihypertensive mode of action of β -blockers
 - Botanical/herbal remedies