MLT 2nd

Course Title: General pharmacology I

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Note:

- Paper is divided into two questions, Q1 includes 15 MCQs and Q2 includes 15 True/False statements
- Each MCQ or T/F carry one mark with grand total of 30 marks
 Highlight or underline the appropriate option
- Before marking, read every statement carefully to understand the actual sense of question

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- In the process of metabolism introducing the polar functional group into molecule may the pharmacological activity of drug
 - **Increase**
 - Decrease
 - No change
 - All of the above
- Drugs having the properties of ______ may have prolong half life
 - Hydrophobic
 - Pass from enterohepatic circulation
 - Both a. and b.
 - None of the above
- Rapid signal transmission and processing occur through
 - G-protein coupled receptor
 - Ligand-gated receptor
 - Enzyme linked receptor

- Intracellular receptor
- Which of the following statement is not true
 - Mechanistically, every drug can illicit its action via enzymes, ion channels, transporters, receptors
 - Majority of drugs show its effects by interacting with its target biomolecules
 - The targeted biomolecules for initiating drug action are proteins in nature
 - Mechanism of drug action always depends on its selectivity
- The therapeutic index represents an estimate of the______ of a drug, because a very safe drug might be expected to have a very large toxic dose and a much smaller effective dose.
 - Efficacy
 - Safety
 - Toxicity
 - Both a. and b.
- Which of the following statement is incorrect
 - Oral drug administration is easy to self-administer
 - In emergency situation best choice of drug administration is I/M
 - Drugs with poor penetrability is usually administered through I/V
 - Inhalation means administration through mouth in small, atomized droplets form
- Signals to the skeletal muscles are controlled by
 - G-protein receptors
 - Second messenger system only
 - Nicotinic receptors

- None of the above
- Every drug follows same path/phases of metabolism such as
 - Only one phase at a time
 - Phase I followed by Phase II
 - Phase II followed by Phase I
 - Depending upon chemical nature of drug
 - Both a. and d.
- Which of the following statement is the false one
 - Majority of drugs are equally eliminated via urine and feases
 - Nephron is the structural and functional unit of kidney
 - Only appropriately metabolized can be eliminated efficiently by kidneys
 - If kidneys failed to eliminate then workload is eased by intestinal route.
- Degradative enzymes of GI tract cause drugs to denature which in turn lower its
 - Absorption
 - Distribution
 - Bioavailability
 - Both a. and c.
 - All of the above
- Which receptor require preliminary formation of ligand bonding that lead to moveable complex
 - Enzyme linked receptors
 - Ligand-gated ion channels
 - Intracellular receptors

	G-protein coupled receptors				
•	A patient having certain type of infections showed altered				
	Biological half-life				
	Biological effect half-life				
	Plasma half-life				
	All of the above				
•	Drug distribution of any class of drug can be determined in terms of				
	Hydrophilicity				
	Molecular weight				
	Hydrophobicity				
	Plasma proteins binding				
	• All of the above				
•	Which of the following statement is not related to exact principle of drug action				
	• Treatment of low heart rate by giving adrenaline				
	Treatment of acidity with omeprazole				
	• Treatment of diabetes by giving insulin as external hormone				
	Treatment cancer by giving them genetic therapy				
	• All are true				
•	Which of the following directly alter membrane potential				
	• Enzymes				
	• Transporters				
	• Ion-linked channels				
	• All of the above				

Q2. Select and highlight true and false for the given statements

- Clinical pharmacology includes the investigation of drug's efficacy and safety in various form of subjects (True/False)
- Drug X shows its action by releasing cholinergic neurotransmitter, due to large particulate nature its release will occur through endocytosis (True/False)
- Drug metabolism in humans usually results in a product that is more lipid soluble than the original drug (True/False)
- Various doses, dosage form and frequency is adjusted in pharmacotherapeutics (True/False)
- In passive diffusion, aqueous soluble drugs cannot move easily across cell membrane (True/false)
- A patient is administered with drug A started toxic reaction but that drug is slowly
 metabolized by enzymes, he/she should be treated with cimetidine to overcome toxicity
 (True/False)
- In enterohepatic circulation, reactivated metabolites comes from the metabolism via enzymes other than CYP P450 (True/False)
- Drug A is hydrophobic in nature and having high plasma proteins binding are likely to distributed effectively to the target organ (True/False)
- A patient has some allergic reactions to the drug A, for avoiding any adverse reaction he administered Drug B with same desired properties, both of the drugs are therapeutically equivalent (True/False)

- Different dosage forms of same therapeutic class of drug will always effect the time to achieve peak plasma concentration (True/False)
- In case of drug action, preliminary bonding of drug with specified receptor will always lead to alteration of receptor's structure (True/False)
- Mechanistically, every drug can illicit its action via enzymes, ion channels, transporters, receptors (True/False)
- As compared to other receptors, intracellular receptors take long for initiating its effect (True/False)
- Depending upon the nature of ligand, majority of drugs interact with receptors that are present across the cell membrane (True/False)
- Various subunits of second messenger system of G-proteins is always linked other receptors i.e. intracellular receptors, enzymes or ion linked channels (True/False)