

<p>d. Dose-response relationship</p>	<p>Draw the concentration-effect curves for the relationship of the effect against,</p> <ul style="list-style-type: none"> • full agonist concentration • logarithm of full agonist concentration • log partial agonist concentration • log full agonist concentration in the presence of a fixed/increasing amounts of competitive antagonist • log full agonist concentration in the presence of a non-competitive antagonist • log full agonist concentration in the presence of a partial agonist • log inverse agonist 		
<p>2016-SBM/MED2214/03</p>			
<p>Pharmacokinetics</p> <p>a. Transport across cell membrane:</p> <p>b. Absorption</p> <p>c. Routes of administration</p> <p>d. Distribution in tissues, body compartments and across barriers</p> <p>e. Metabolism (Biotransformation)</p> <p>f. Elimination</p>	<p>Describe the mechanisms of transport of drug molecules across the cell membrane and the factors that influence such mechanisms</p> <p>1. explain how drugs are absorbed into blood after administration</p> <p>2. list the factors that influence the absorption of drugs</p> <p>1. list different routes of administration of drugs</p> <p>2. list the different types of dosage forms/special drug delivery systems</p> <p>3. explain the advantages and disadvantages of different routes of administration</p> <p>1. list the different compartments of the body into which drugs are distributed</p> <p>2. describe the factors which influence the distribution of drugs into different compartments</p> <p>3. explain the concept of redistribution of drugs</p> <p>4. explain the concept of barriers across tissues for transport of drugs</p> <p>1. explain the basic mechanisms by which drugs undergo biotransformation in the body</p> <p>2. list the common drugs which induce/inhibit the cytochrome P 450 enzyme system</p> <p>1. define elimination of drugs</p> <p>2. list the physiological processes of different organ-systems that are involved in drug elimination</p> <p>3. describe the mechanisms by which drugs are eliminated from the body</p>	<p>6</p> <p>2</p>	<p>Lectures</p> <p>SGD</p>



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<p>g. Pharmacokinetic parameters</p> <p>h. Drug concentration vs time curve in different dosing regimes</p> <p>i. First-order & Zero-order kinetics</p>	<p>1. define the following</p> <ul style="list-style-type: none"> • bioavailability • bioequivalence • first pass effect • area under the Concentrate-time curve (AUC) • (apparent) volume of distribution • clearance • half life • steady state concentration • loading dose • maintenance dose • dosage regimen <p>2. explain the principles of calculating the bioavailability, volume of distribution, clearance, loading dose & maintenance dose</p> <p>draw the concentration-time curves for</p> <ul style="list-style-type: none"> • single intravenous bolus injection • intermittent intravenous bolus injection • continuous intravenous infusion • single intramuscular injection • single subcutaneous injection • single-dose oral administration • intermittent oral administration • modified-release formulations <p>explain first order kinetics and zero order kinetics</p>		
<p>j. Clinical application of pharmacokinetic parameters</p>	<p>explain the clinical significance of pharmacokinetic principles</p>		
<p>2016-SBM/MED2214/04</p>			
<p>Adverse and Toxic effects</p>			
<p>a. Basis of adverse/toxic effects</p>	<p>1. define adverse effects and toxic effects of drugs</p> <p>2. describe the mechanisms of adverse effects of drugs</p> <p>3. classify adverse effects based on their mechanisms briefly explain teratogenicity, mutagenicity and carcinogenicity</p> <p>5. explain how these reactions could be minimized/prevented.</p> <p>6. define therapeutic index</p> <p>7. explain the clinical significance of therapeutic index</p>	<p>4</p> <p>2</p>	<p>Lecture</p> <p>SGD</p>
<p>b. Drug interactions</p>	<p>1. classify drug interactions (eg. Drug-drug, drug-food and drug-herb)</p> <p>2. describe mechanisms of drug interactions</p> <p>3. explain the clinical significance of drug interactions</p>		



c. Pharmacogenetics	describe the influence of genetic variation on response to drug therapy		
d. Drug therapy in special populations	describe the principles underlying the precautions that should be taken during drug therapy in special situations such as pregnancy, breast feeding, renal and hepatic dysfunction, extremes of age		
e. Measurement & monitoring of drug effect	<ol style="list-style-type: none"> 1. describe the methods by which the effects of drug therapy could be measured 2. describe how the measurement of plasma drug concentrations helps in monitoring drug therapy 		
2016-SBM/MED2214/05			
Autonomic Nervous System	<ol style="list-style-type: none"> 1. recall the anatomical and functional organization of autonomic nervous system 2. recall the anatomy and the physiology of the cholinergic and the noradrenergic 'junctions 3. recall the types of autonomic receptors with examples of typical sites 4. describe the mechanisms of action and clinical uses of drugs acting on autonomic nervous system 	2	Lecture
		1	SGD
2016-SBM/MED2214/06			
Pain Control			
a. Physiology of pain	<ol style="list-style-type: none"> 1. recall the definition of pain and briefly explain theories of pain 2. list the types of pain 3. recall physiology of pain perception (stimuli, receptors, pathways and central connection) 4. list methods of pain relief 5. classify pharmacological agents used in pain relief (with main indications) 6. explain the basis of neuropathic pain 		
b. Opioid Analgesics	<ol style="list-style-type: none"> 1. classify the agents acting on opioid receptors 2. describe the mechanisms of action of opioid analgesics. 3. describe the pharmacokinetics of the drugs acting on opioid receptors. 4. describe the adverse effects of opioid analgesics. 5. list the clinical uses of opioid receptor antagonists 	2	Lecture
c. Non-steroidal anti-inflammatory drugs (NSAIDs)	<ol style="list-style-type: none"> 1. describe the physiological/pathological roles of cyclo-oxygenase-1 (COX-1) and COX-2 enzymes. 2. describe the pharmacokinetics, clinical uses, important adverse effects and drug interactions of NSAIDs (including COX-2 inhibitors). 3. list the commonly used NSAIDs 	2	SGD



2016-SBM/MED2214/07			
Drug therapy in neoplastic disease	<ol style="list-style-type: none"> 1. state how neoplastic cells/tissues differ from normal cells/tissues with respect to potential targets for drug therapy in neoplastic disease 2. explain the basis of <ul style="list-style-type: none"> • combination chemotherapy • resistance to chemotherapy • adverse effects of chemotherapy 3. classify antineoplastic drugs based on the mechanism of action 	1	Lecture
2016-SBM/MED2214/08			
Drug Information			
a. Sources, Reliability and Interpretation	<ol style="list-style-type: none"> 1. identify different sources of drug information 2. differentiate unbiased information from promotional material. 3. critically analyse the information in a given source of drug information. 4. carry out a literature search on drug information 	2	SGD
b. Drug Discovery and Development	<ol style="list-style-type: none"> 1. state the history of drug discovery 2. list the sources from which new drugs are developed 3. describe the different stages of the development of a new drug 	1	Lecture
2016-SBM/MED2214/09			
Antimicrobial agents	<ol style="list-style-type: none"> 1. define an “antimicrobial agent” 2. classify antimicrobial agents based on their chemical structure/mechanism of action with examples 3. describe the mechanism of action, pharmacokinetics, clinical uses, adverse effects, interactions and limitations for the use of commonly used antimicrobial drugs 4. explain the basis of chemoprophylaxis in infections 5. explain the principles underlying the selection of appropriate antimicrobial agents in infectious diseases 	9 4	Lecture SGD

