Pharmacology - Core Concept Cheat Sheet

01: Introduction to Pharmacology

History Of Pharmacology	Types Of Drug Interactions
 Materia medica: the science of drug preparation and medical use of drugs began to develop around the 17th century. Pharmacological studies developed from this. Magendie and Bernard: Laid the foundations for animal physiology and pharmacology in the 18th and 19th centuries. Pharmacogenomics: the study of how an individual's genetic make up affects his or her response to a drug. 	 Drug interactions can be divided into: Pharmacodynamic: the actions of the drug on the body Agonist Antagonist Pharmacokinetic: the actions of the body on the drug Agonists: initiate changes in cell function, producing effects of various types. Their potency
Introduction To Pharmacology Terminology	depends upon their:
 Pharmacology: study of the way that substances interact with the systems of the body to activate or inhibit its processes. Toxicology: the term used to describe the undesirable effects of drugs. Drug: any substance that brings about a change in biological function through its chemical actions. Prodrug: a substance that is administered in its inactive form, but once absorbed, is converted into an active drug molecule. Xenobiotic: a drug molecule that is a chemical not synthesized by the body. Poison: drugs that cause harmful and undesirable effects. Toxins: poisons that are of biological origin. 	 Affinity: the tendency to bind to receptors Efficacy: the ability to initiate changes once bound Full Agonists: Produce maximal effects Have high efficacy Partial Agonist: Produce a lower response at full receptor occupancy than full agonists Have intermediate efficacy Antagonists: prevent agonists from activating receptors Reversible competitive antagonism: Progressively inhibit the agonist response; at maximal concentrations, they completely prevent the response.
Physical Nature Of Drugs	Non-competitive antagonism:
 There are four characteristics of drugs that affect how they interact with the body. Drug size Drug reactivity and drug-receptor bonds Drug shape Degree of ionization Drug Size: affects binding to receptors as well as the permeability through membranes. Drug Reactivity and Drug-Receptor Bonds: Drugs may bind with receptor molecules through covalent, electrostatic and hydrophobic bonds Covalent bonds: tend to be stronger and hence result in irreversible interactions Hydrophobic bonds: on the other hand are relatively weak 	 Block the chain of events that leads to the production of a response by the agonist. Non-Receptor Antagonists: Chemical antagonism: interaction of two substances in solution so that the effect of the active drug is lost Pharmacokinetic antagonism: one drug affecting the absorption, metabolism or excretion of another drug Physiological antagonism: two agents producing opposing physiological effects Pharmacodynamic effect: the duration of drug action depends upon how long the drug occupies the receptor. The Dose-Response Relationship: for a given dose of a drug, there will be a given biological response that is directly proportional to the given dose.
 Drug Shape: shape of the drug molecule is important factor in determining how well it will fit into its receptor Ionization State of a Drug: Drug molecules exist in ionized and unionized forms Henderson-Hasselbalch Equation: Calculates the percentage of ionized & unionized molecules in solution. 	

How to Use This Cheat Sheet: These are the keys related this topic. Try to read through it carefully twice then recite it out on a blank sheet of paper.