



Pharmacology



Drug Fundamentals, Plus the Most Frequently Prescribed Drug Classifications-Including Indications, Reactions, Examples & More

definitions

- phermacodynamics Study of the mechanisms of action of drugs within the body and how drugs produce their effects in the body
- pharmacogenetics Study of drug reactions in the body that are manticipated or unusual, and may have a hereditary basis for the response pharmacokinetics Study of drug actions as they move through the
 - body, the way the body absorbs, distributes, metabolizes and excretes drugs; mathematical study of drugs based on time and dose
- pharmacology Study of biologically active compounds, how they react n the body and how the body reacts to them
- pharmacetherapeutics Study of drugs used to prevent, treat or diagnose disease
- m pharmacy Preparation and dispensing of drugs.
- texicology Study of harmful or poisonous effects of drugs

erue mannes CHEST Chemical Name: Scientific Generale Drogs Progress Palament. name, also other the provi no, networked by the he structures reducedant form and these advantages to provide pharmadeutical company that make the drug

pharmacodynamics

- . Protein molecules with one or more binding sites, located on cell
- Receive a signal from the body's chemicals: neurotransmitters, hormones, enzymes
- Signal will cause a molecular event on the inside of the cell to occur
- Drugs Enhance (agonist), diminish (partial agonist) or block (antagonist) the generation, transmission or receiving of the signal
- Affinity Attraction between a drug and a receptor
- High affinity Drug will bind easily to the receptor
- . Low affinity Requires a higher concentration of the drug to get a therapeutic response

Driving Plotterscy

Amount of drug required to produce a therapeutic response

Dose Response Curve

- Effective Dose (ED) Amount of drug that produces a therapeutic response in 50% of the people taking it
- Toxic Dose (TD) Amount of drug that produces adverse effects in 50th of the people taking it.
- Therapeutic Index (Ti) Margin of safety; ratio between the TD and the ED
- The higher the TI, the safer the drug is considered to be; in general. nonprescription drugs have much higher TIs than prescription drugs

pharmacokinetics

Routes drugs take to get into the body

- - Enters the body through the GI tract.
 - Taken by mouth, through the rectum, under the tongue or held in the cheeks.
 - Enters the body through a different means (i.e., other than the GI tract) Can be injected into the veins, arteries, muscles, spinal cond, or under the skin; inhalled through the lungs; transdermally through the skin via

Absorption

- Biographisty Percent absorbed into systemic circulation after administration · Bioavailability depends on route of administration as well as the drug's ability to cross membranes and reach its target
- First Pass Effect:
 - . Drugs absorbed through the stomach and small intestine must pass through the liver before circulating systemically
- · Liver can inactivate the drug, making less of the drug available to reach the target organ
- Absorption at cellular level occurs through passive transport, active transport, pinocytosis and facilitated diffusion

Distribution

- · Influenced by several factors:
 - . Tissue permeability: Ability of drug to pass through the membranes rapidly affects the extent to which the drug moves around in the body.
- a Blood flow: Once in the blood stream, will get to the organs and
- tissues that are highly perfused.

 Plasma proteins binding: Drug can bind to a protein that will render. the drug inactive; only an unbound drug can attach to the receptors
- # Binding to subcellular components
- e blood pH

Drug Storage Sites

- Adipose tissue Primary site; lipid-soluble; drugs tend to remain for long. periods of time due to low metabolic rates of drugs and poor blood perfusion of tissue
- Bone Toxic agents like heavy metals.
- Muscle Binding can cause muscle to store drugs
- Oceans Liver and kidneys

Metabolism

- Biotramiformation Chemical changes that occur in the drug following
- Metabolite Altered version of the chemical compound
- Can have a higher or lower rate of activity than the original drug, if higher, drug is given as an inactive or prodrug form
- Prodrug Requires metabolism or activation of drug in order for it to act within the body

Exponention.

- First order Rate of removal of drug from the body is proportional to the concentration of the drug in the plasma
- Half-life Time required to decrease the blood levels of a drug by one-half.
 A one-time drug dosage will be eliminated almost completely by 5.
- Population of the Parket St.
- A drug given on a continuous dosage schedule will reach steady state concentration after 5 half-lives
- Steady state Rate of drug administration is equal to the rate of drug excretion
- Organs that excrete drugs Kidneys, lungs, sweat glands, mammary glands, salivary glands, skin and Gl tract

chedule dans



Schedule Class Schedule 1: C-1 Schedule 2: C-III

- Characteristics.
- High abuse potential, not legal, no acceptable medical use; no prescriptions available. High abuse potential and severe dependence liability; current, accepted medical use; prescription drug signed, not stamped prescription; 30-day supply, no refills
- Schedule 3: C-III . Less abuse potential; low-moderate physical dependence; high psychological dependence; by prescription only, expires within 6 months; max. 5 refills on one
- Schedule 4: C-IV . Less abuse potential than C-III drugs; accepted medical use; limited physical and psychological dependence, written or verbal prescription, expires in 6 months.
- Schedule St C-V .
- max. 5 refills on one script
- Limited abuse potential, accepted medical use; small amounts of narcotics used as antitussives (cough medicine) or antidianheals; may not need a prescription but must be recorded as a transaction.

Examples (C-I to C-IV)

- . Heroin, LSD, cocaine, marguana, methaqualone
- Opium, morphine, coca. methadone
 - · Amphetamines, codeine, burbituratos, Valium, Xanox, amaboolic steroids.
- · Chloral hydrate, meptrobarmate. paraldehyde, phonobarbital

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