

Pharmacology Concepts For Emergency Medical Technicians

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Drug Names

- Chemical Name
 - Precise description of the drug's chemical composition and molecular structure
 - 7-chloro-1, 3-dihydro-1methyl-5-phenyl-2H-1, 4benzodiazepin-2-one
- Generic Name (Non-proprietary Name)
 - Official name approved by the FDA
 - Usually suggested by the first manufacturer
 - diazepam



Drug Names

- Official Name
 - The name assigned by the USP
 - diazepam, USP
- Trade Name (Proprietary Name)
 - The brand name registered to a specific manufacturer or owner
 - Valium ®

Sources of Medications

- Plants
 - morphine sulfate, atropine
- Animals and/or Humans
 - insulin, ACTH
- Minerals
 - sodium bicarb, calcium
- Synthetic (Chemical Substances)
 - lidocaine, diazepam



Drug Classification

- By Body System
 - Sympathetic Agonist, Anticholinergic
- Class of Agent
 - Antidysrhythmic, Analgesic
- Mechanism of Action
 - Calcium Channel Blocker, Diuretic



Sources of Drug Information

- Physicians Desk Reference (PDR)
- Hospital Formulary (HF)
- Drug Inserts
- Other texts/sources
 - Brady
 - Internet
 - Mosby
 - Jones and Bartlett

Components of a Drug Profile

- Brand name
- Generic name
- Classification
- Mechanism of action
- Indication
- Contraindication
- Route of administration
- Side effects/adverse reaction
- Adult & Pediatric dose
- Onset & duration



Scope of Management

- You are held responsible for safe and therapeutically effective drug administration
- Personally responsible for each drug you administer
 - Legally
 - Morally
 - Ethically

The "Six Rights" of Medication Administration

- Right Medication
- Right Dose
- Right Time
- Right Route
- Right Patient
- Right Documentation

General Properties of Drugs

- Drugs <u>do not</u> confer any new functions on a tissue or organ, they only <u>modify</u> existing functions
- Drugs in general exert <u>multiple</u> effects rather than a <u>single</u> effect
- Drug action results from a physiochemical interaction between the drug and a functionally important molecule in the body



Pharmacology

- Pharmacokinetics
- The study of how a drug enters the body, reaches its site of action, and is eliminated
- Principles of pharmacokinetics (M/A)
 To exert its biomechanical effects on the body a drug must reach its targeted tissues in suitable form and significant concentration
- Principle of pharmacodynamics
 - Interactions at receptor sites



Pharmokinetics

- Mechanisms that affect pharmokinetics
 - Absorption
 - Distribution
 - Biotransformation
 - Excretion



Basic Drug Processes

- To produce a desired effect must be present in an appropriate concentration
 - At various sites of action
- Each drug undergoes four basic processes
- These processes influence the concentration at the site of action



The 4 Basic Drug Processes

Absorption into the circulatory system

• Distribution throughout the body

Biotransformation into its active form

• Elimination from the body



Absorption

- Diffusion
- Tendency of molecules to move from an area of higher concentration to lower

The molecules may be in the form of a gas, solid, or liquid

• Diffusion occurs through a semipermeable membrane



Absorption

Osmosis

• Diffusion of fluid through a semipermeable membrane

• The principle flow being from the less dense to a more dense solution



Drug Concentration

 Concentration of a drug also affects rate of absorption

 Drugs administered in high concentrations are absorbed more rapidly than those administered in low concentrations

Rate of Drug Absorption

- Areas of high vascular supply have more rapid rate of absorption
- Areas of lower vascular supply (subcutaneous tissues) have slower rate
- A general rule is muscle has a higher vascular supply than sub-q tissues
 - Thus rate of absorption is faster in muscle than in sub-q tissues



Distribution

 The process whereby a drug is transported from the site of absorption to the site of action

 Once in the circulatory system a drug will be distributed throughout various body tissues



Distribution

- Organs with the greatest blood supply will receive drugs the fastest
 - Heart, brain, lungs, kidney
- Organs with a smaller blood supply will receive drugs more slowly
 - Skin, GI system, fat



Barriers to Distribution

 Drug distribution is also affected by physiological barriers

 These barriers may inhibit the movement of certain substances, but permit the passage of others



Barriers to Distribution

- Drug reservoirs
 - Plasma protein binding (molecules)
 - Tissue binding (adipose or fat)
- Barriers
 - Blood-brain barrier
 - Placental barrier



Barriers to Distribution

• Examples of these barriers are:

Blood-brain barrier

Blood-testes barrier

Blood-placental barrier



- In pharmacology the term *metabolism* often refers to making a drug more polar or water soluble
- Although this often results in drug inactivation and excretion, it's incorrect to assume that a metabolite will be less active or more easily excreted than the parent drug



 Drugs, chemicals, and toxins are foreign to the body

 The body will attempt to rid itself of these substances, regardless of the therapeutic value of that substance



 Most drugs must be biotransformed, or metabolized, before they can be excreted

 This occurs through a complex cascade of chemical reactions until they become chemically inactive



Metabolic reactions can transform.....

 An active drug into less active or inactive forms

• A prodrug (inactive or less active drug) into a more active drug



 All inactive chemicals, chemical byproducts, and waste(often referred to as metabolites) are eventually excreted

• This is the process of elimination



The 4 Basic Drug Processes

- Elimination
- K= kidneys
- I= intestines
- L= liver
- L= lungs
- S= skin



 The rate of elimination varies with the drug and state of the body

 If the liver or kidneys do not function properly, drugs may not be properly metabolized or eliminated



- This may cause higher doses of medication to circulate for a longer period of time
- This produces symptoms of toxicity and over dosage
- A standard method of expressing how long it takes to metabolize and eliminate a drug is called the *half-life*



 The half-life is the time it takes to eliminate 50% of the drug from the body

 Because these rates are usually the same for most people, half-life helps determine dosage and frequency required for the administration of different drugs





- Liquid
- Solid
- Gas

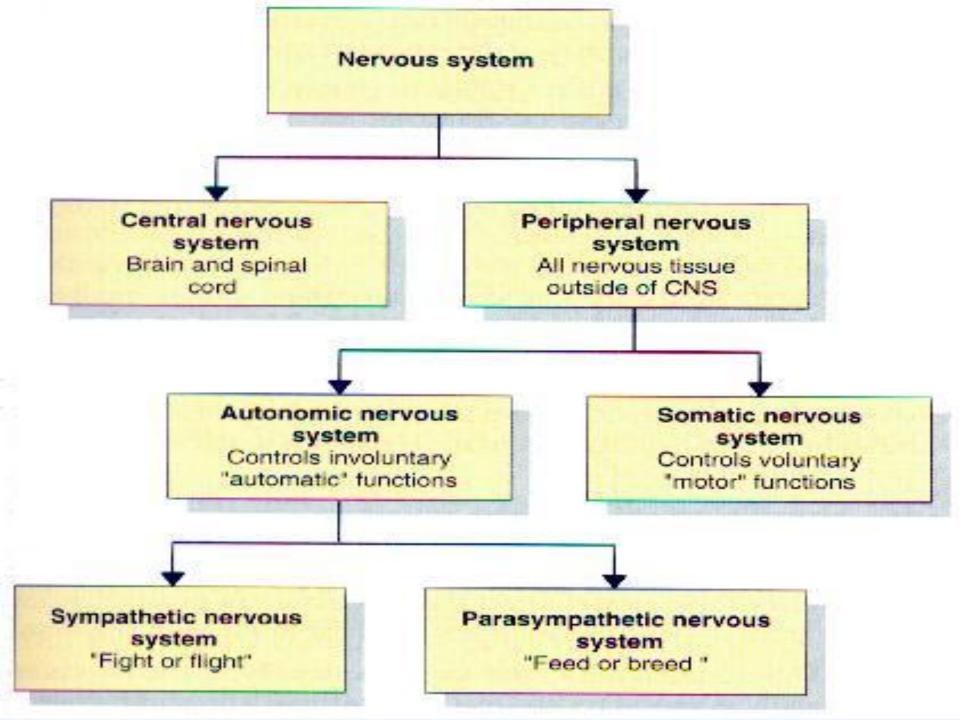


Routes of Drug Administration

 Effects the rate at which the onset of action occurs and may effect the therapeutic response that results

Autonomic Pharmacology

- Central Nervous System (CNS)
- Peripheral Nervous System
 - Somatic Nervous System
 - Autonomic Nervous System (ANS)
 - Sympathetic Branch
 - Parasympathetic Branch



Drug Receptor Interaction

- Affinity
 - Drug's propensity to bind or attach itself to a given receptor site
- Efficacy
 - Drug's ability to initiate biological activity as a result of binding to a receptor site

Drug Receptor Interaction

- Agonists
 - Drug that binds to a receptor site and causes a physiological response
- Antagonists
 - Drug that binds to a receptor site and prevents a physiological response or prevents another drug from binding to a receptor site



Type of Receptors

- Beta 1
- Beta 2
- Alpha 1
- Alpha 2



Beta Receptors

- Beta 1 Receptors
 - Located primarily in the heart
 - Cause increase in force of contraction and heart rate
- Beta 2 Receptors
 - Located primarily in the lungs
 - Dilate bronchioles & blood vessels
 - Relax smooth muscle



Alpha Receptors

- Alpha 1 Receptors
 - Located primarily in the vascular bed
 - Cause increase in contraction of arterial vessels



Predictable Responses

- Desired Action
 - Action or effect is seen that is consistent with why the drug was given
- Side Effects
 - Undesirable and often unavoidable effects of a drug
 - Action or effect other than those for which the drug was given



Unpredictable Adverse Reactions

- Allergic Reaction
 - Activates the Immune System
- Anaphylactic Reaction
 - Severe allergic reaction
- Idiosyncracy
 - Drug effect unique to individual
 - Different than expected



Unpredictable Adverse Reactions

- Tolerance
 - Physiologic response that requires a drug dosage to be increased to produce the same effect
- Cross Tolerance
 - Tolerance after administration of a different drug
 - Morphine and other opiod agents

Drugs by Classification

- Analgesics and antagonists
- Anesthetics
- Antianxiety, sedative, and hypnotic drugs
- Anticonvulsants

- Central nervous system stimulants
- Psychotherapeutic drugs
- Antidepressant therapy



Beta Effects

- Cardiac acceleration and increased contractility
- Vasodilation of arterioles supplying the skeletal muscle
- Bronchial relaxation
- Uterine relaxation



Indirect Acting & Dual Acting Drugs

- Act indirectly on receptors by triggering the release of the catecholamines norepinephrine and epinephrine which then activates the alpha and beta receptors
- Example
 - Ephedrine Sulfate



Drugs that Affect the Blood

Anti-Platelett Agrregant



Anticoagulants

- Drugs that interfere with platelet aggregation
- Also known as anitplatelet or antithrombic drugs
 - Aspirin



Bronchodilators

- Administered via inhalation via a nebulizer or a pressurized cartridge
- Sympathomimetics categorized by receptor action
 - Selective beta2 receptor drugs



Bronchodilators

- Albuterol
- Atrovent
- Combi-Vent

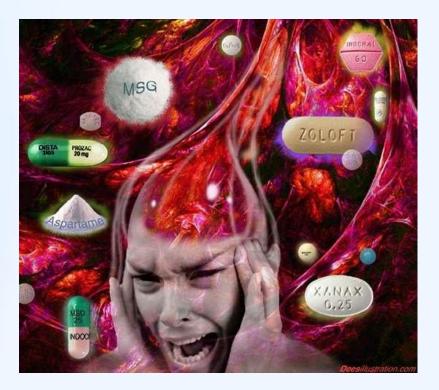


Epinephrine

• EPI-Pen Adult

EPI-Pen Peds





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