#### PHARMACOLOGY & THE NURSING PROCESS

- Gather data necessary to evaluate therapoutic effects.
- Cathor data recessary to evaluate adverse reactions
  Judge if patient is capable of self-administration of treatment.
- Assess any potential risk factors Analysis/Diagnosis

- Determine if prescribed therapeutic plan is appropriate.
- Identify possible drug interactions, edverse reactions, and health problems

Compose a care plan, including nursing interventions and expected outcomes.

#### **Implementation**

- Administer the drug.
- . Educate the patient about the drug-
- . Enact measures that will enhance drug effectiveness and enintenize adverse reactions.

#### Wanted Street

- . Determine the presence or absence of the desired therapeutic response or adverse reactions
- · Identify if patient followed the regimen prescribed and is satisfied with treatment

#### DRUG NAMES

#### Chemical Name

Scientific name, describes the atomic and molecular structure of a drug

#### Generic Name

Nonproprietary name, abbreviation of the chemical name

#### Brand Name

Trade name, selected by the pharmacoutical company that made the drug

#### **EDUCATING PATIENTS**

When instructing patients on a prescribed drug, share the following:

- Drug name/category
- Dosage size/schedule
- What to do in the event of skipped dose
- How to administer. Expected response
- How the patient can enhance response. Duration of treatment.
- \*How to store the drug
- · Major and minor side effects.
  - Important possible interactions.
  - How to react to an emergency.

#### DRUG INTERACTION

Additive: Two drugs with similar therapeutic effect - combined potency

Antagonistic: Two drugs taken together = decreased effectiveness of both Potentiation: One drug's potency is enhanced when combined with another

Absorption: One drug affects absorption rate of another

Metabolism: One drug affects metabolism of another, if inhibited, can cause **BORRETTY** 

#### Adverse Reactions

Caused by Patient Sensitivity

Altergic/anaphylactic reaction lidiosymcratic response

Caused by Dose

Secondary/side effects

Hypersusceptibility futrogenic effects

Overdone

#### ROUTES OF ADMINISTRATION

Orak Capsule, tablet, liquid; absorbed in Gil tract

Intravenous DVs Injection into bloodstream via vein

Intramuscular DMs: Injection into muscle: can use large doors: fast systemic action.

Intrathecal: Injection into spinal canal; affects spinal fluid

Subcutaneous (sub-Q): Injection into tissue below dermis

Sublingual: Absorbed under the tongue

Rectal/vaginal: Suppositories or preams; usually for local distribution.

Inhalation: Absorbed in lungs; gaseous form; rapid absorption

#### PHARMACOLOGY BASICS

Pharmacokinetics: Study of drugs' actions as they move through the body

Absorption: Affected by administration route, formulation, patient stress level or stomach contents, blood flow

Distribution: Affected by solubility, protein activity, blood flow

Metabolism: Affected by diseases, environment, age, genetics Exception: Through kidneys, skin, intestines, lungs, exocrine glands

Pharmacodynamics: Study of the mechanism of action of drugs within the body and how drugs produce their effects in the body

Pharmacetherapeutics: Study of drugs used to prevent, treat, or diagnose disease

#### CONSIDERATIONS ACROSS THE LIFESPAN.

#### Pregnant/Breastfeeding Women

- c metabolism are accelerated w consider higher dosages. · intestinal excetion rate decreased - longer absorption time - consider reduced dosages
- Assume all drugs can enter the fetus via placenta.
- Consult FDA risk categories for every drug before considering administration
  Weeks 3-8 present greatest risk of drug-induced malformation of fetus

- After pregnancy, doses should be taken directly after breestfeeding to ensure minimal drug concentration in breast milk for next feeding.

#### Pediatric Patients

- Assume increased drug sensitivity due to immeture organ system.
- Infants have imegular gastric patterns absorption rates may vary
- Infants/young children have thin skin = rapid topical drug absorption.
- \*Infants' blood-brain barrier not fully developed increased sensitivity to CNS drugs, risk
- Neonates absorb IM drugs slower than adults: infants absorb IM drugs faster
- Infants have reduced protein-binding ability = high free concentrations of drugs
- User and kidneys not fully developed until after 1 year of life; assume reduced ability for hepotic and renal metabolism in infants
- In children over 1 year of age, drug metabolism rate is higher than adults.
- Children may have unique side effects to certain drugs, including suppressed growth

General rule for dooing adjustment:
 Body surface area of shild x Ashift dots: = approx. dose for child

Patient education should include instructions for how to account for spitting or spilling of medication, avoiding multiple desing

#### Geriatric Patients

- Assume increased drug sensitivity due to deteriorated organ system
  Reactions vary greatly based on individual patient condition

- Rate of absorption generally slowed = delayed therapoutic response
  Hepatic metabolism rate likely slowed = longer therapoutic response
  Hepatic metabolism as accumulation of drug, increased risk of authorizer afflication
- Determine creatinine dearance prior to drug administration to assess. nemal functions
- Intentional failure to follow prescribed regimen a common problem;
- longer or more extensive patient education may be required Anticipate, assess, and/or manage drug interactions carefully



Schedule Class	Characteristics	
Schedule 1: C-II	High abuse potential; not legal; no acceptable medical use; no prescriptions available	
Schedule 2: C-8	High abuse potential and severe dependence liability, current, accepted medical use; prescription drug-signed, not stamped prescription; 30-day supply, no refills.	1
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 script.	3
Schedule 4: C-IV	Less abuse potential than C-81 drugs; accepted medical use; limited physical and psychological dependence; written or verbal prescription, expires within 6 months; max. 5 refills on one script	0
Schedule Sr C-V	Limited abuse potential; accepted medical use: small amounts of narcotics used as antitusives icough medicine) or antidianheals; may not need a prescription but must be recorded as a transaction	1

Heroin, LSO, cocaine, marijuana, methaqualone Opium, morphine, coca, methadone	
Amphetamines, codeine, barbiturates, Valium, Kanax, anabolis, steroids	

Chloral hydrate, meptrobamate, paraldehyde, phenobarbital. Promethatine with codeline (cough medicine) oligiberronylate (arctidiarcheal)

# **Study Guide For Nursing Pharmacology**

Norma L. Pinnell

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