

**HNN215: Quality Use of Medicines**

**Week 1- Introduction to QUM (Part A) & Medicines used to treat pain (Part B):**

**Introduction to QUM:**

**Demonstrate an understanding of the principles of quality use of medicines:**

Quality use of medicines is using medicines safely and effectively, choosing suitable medicines that are relevant to the situations and managing options wisely

**Identify the causes and effects of drug errors and explain relevant nursing actions to minimise risks**

- Medication errors increase the risk of adverse drug reactions
- Causes: distraction, poor communication, lack of documentation, wrong medication calculation, wrong patient, wrong time, wrong dose, wrong medication, wrong route, tiredness, stress, medications that sound/ look alike
- Effects: illness, death, allergic reaction
- To minimise risks:
  - right patient, right medication, right time, right dose, right route
  - patient participation, listen to patients and include them in medication management
  - be confident in medication calculations
  - documentation
  - communicate effectively with patient, family, nurses, pharmacists, doctors etc
  - review medications frequently
  - check that medication prescription is legal and medication order is legible
  - assess the patient and situation before and after administration

**Apply clinical decision making to ensure safe medication practices**

**Calculate medication doses**

**Interpret medication prescriptions accurately**

**Demonstrate evidence-based practices in medication documentation**

**Explain and apply relevant pharmacodynamic and pharmacokinetic concepts (eg bioavailability, half-life, steady state concentration, first pass effect, metabolism and excretion)**

- Bioavailability: percentage of agent administered that reaches the systemic circulation; absorption efficiency of a medication
- Half life: time interval required for elimination processes to reduce the concentration of a medication in the body by one half
- Steady state concentration: the amount of drug in the plasma has built up to a concentration level that is therapeutically effective and as long as regular doses are administered to balance the amount of drug being cleared, the drug will be active
- First pass effect: medications administered orally enter the liver first and are metabolised before they enter the systemic circulation
- Metabolism: process of energy and material transformation in all living cells; the sum of all physical and chemical changes that take place within a microorganism
- Excretion: process by which a drug is eliminated from an organism either in an unaltered form (unbound molecules) or modified as a metabolite

**Identify the factors that may result in altered medication effects in older persons and paediatrics**

Older persons:

- Absorption of some nutrients from the GIT diminishes with advancing age

## Intended Learning Outcomes

- Some medications may pass through the digestive tract unchanged
- Slowed rate of passage of medications through the lower GIT
- Reduction in blood flow due to age results in a decrease in the absorptive surface in the intestine
- Faster topical absorption due to thinner skin surface
- IM absorption is difficult to anticipate because peripheral circulation is affected by environmental changes
- Total body water is decreased resulting in diminished volume of distribution of some water soluble medications
- Total body fat content is increased, altering the distribution of fat soluble medications
- Age related muscle tone loss alters distribution of some medications
- Decrease in protein binding ability
- Enzyme levels decrease because of decline in liver function
- Liver blood flow reduces leading to a decline in the body's ability to transform active medications into inactive metabolites
- More likely to experience medication toxicity because of the accumulation of medications

### Paediatrics:

- Reduced gastric acidity due to immature gastric acid producing cells- medication may pass through the digestive tract unchanged
- Slower gastric emptying
- Faster topical absorption because of thinner skin surface
- IM absorption is difficult to anticipate because peripheral circulation is affected by environmental changes
- Greater total body water content, require higher doses of water soluble medications
- Lower total body fat content, fat soluble medications must be varied
- Decreased protein binding
- Blood brain barrier is immature, leading to more medications being able to enter the brain
- Decreased enzyme levels
- May require higher levels of medication due to higher metabolic rates
- Decreased tubular secretion and reabsorption due to immature kidneys
- Decreased perfusion to the kidneys
- Lower urine pH, medications circulate longer and have the potential of reaching toxic levels
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**Utilise a drug diary to consolidate your knowledge of broad drug groups**

### Medicines used to treat pain:

**Identify the major classes of analgesic agents and examples of medications found in each:**

**Non-narcotics:** paracetamol, Panadol rapid, herron

**NSAIDS:** ibuprofen, diclofenac, celecoxib (COX2)

**Narcotics:** oxycodone, morphine, fentanyl, targine

**Adjuvants:** diazepam, gabapentin

**Briefly describe the actions of each major class (one to two sentences):**

**Non-narcotics:** given for mild to moderate pain. Provides an analgesic effect by inhibiting prostaglandin synthesis in the CNS and activating descending pathways- also has antipyretic effect (reduces temperature)

## Intended Learning Outcomes

**NSAIDs:** given for pain and inflammation for injury, analgesic/ anti- inflammatory and anti- pyretic actions. It inhibits the production of prostaglandins by blocking COX reducing inflammation and producing analgesia

**Narcotics:** indicated for acute or chronic mild, moderate to severe pain. Act on opioid receptors in CNS and GIT, producing analgesia

**Adjuvants:** not pain medications but can work with analgesic to enhance pain relief

### **Identify significant adverse drug reactions (ADRs) for each drug class:**

**Non-narcotics:** significant risk of hepatotoxicity or liver toxicity with elevated doses

**NSAIDs:** GIT bleeding, NSAID induced renal impairment, increased risk of cardiovascular events

**Narcotics:** respiratory depression, constipation, bradycardia and hypotension

**Adjuvants:** injection site pain, skin ulceration

### **Explain relevant nursing actions to manage risk when administering these medications:**

**Non-narcotics:** monitor liver function

**NSAIDs:** use lowest effective dose for the shortest period of time, monitor renal function

**Narcotics:** assessment of sedation and associated side effects, give laxative with analgesics, be aware of legal obligations of administering narcotic analgesics

**Adjuvants:** monitor injection site

### **Synthesise knowledge to inform clinical decision making for patient assessment, education and evaluation**

**Demonstrate evidence-based practice in medication documentation**