# PHARMACOLOGY NOTES NURSING IMPLICATIONS FOR CLINICAL PRACTICE

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Adverse Effects



Teaching

Therapeutic Effects

# **PHARMACOLOGY NOTES**

## **NURSING IMPLICATIONS FOR CLINICAL PRACTICE**

## **Overview**

There are currently nine (9) units comprising this *Pharmacology Notes* resource. Units are broken down by body system and published individually for ease of retrieval:

- Unit A: Autonomic Nervous System (ANS) Pharmacology
- Unit B: Cardiovascular (CV) System Pharmacology
- Unit C: Hematological System Pharmacology
- Unit D: Central Nervous System (CNS) Pharmacology
- Unit E: Skeletal System: Bone and Joint Pharmacology
- Unit F: Immune System Pharmacology
- Unit G: Digestive System Pharmacology
- Unit H: Endocrine System Pharmacology
- Unit I: Respiratory System Pharmacology



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# **UNIT G**

# **DIGESTIVE SYSTEM PHARMACOLOGY**

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(MC) Major Class or Therapeutic Class (SC) Subclass or Pharmacologic Class (SSC) Selective Subclass – more specific action within Subclass

- (MC) Antiulcer Agents
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  (MC) Prokinetics
  (MC) Laxatives
  (MC) Antidiarrheals
  (MC) Anti-IBD
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  (MC) Miscellaneous GI Drugs

  (SC) Antiinflammatory
  (SC) Adsorbents
  (SC) Antiflatulents
- (MC) Nutritional Aids

## **Digestive System Pharmacology**

### I. ANATOMY AND PHYSIOLOGY/PATHOPHYSIOLOGY REVIEW

#### A. Anatomic Divisions: GI Structures and Functions

- 1. Gastrointestinal (GI) Tract or Alimentary Canal:
  - a. Upper GI tract structures and functions:
    - 1) mouth food entry/processing
    - 2) esophagus food passage
    - 3) stomach breaking down food  $\rightarrow$  chemical digestion:
      - a) chief cells pepsinogen  $\rightarrow$  pepsin
      - b) parietal cells hydrochloric acid, intrinsic factor
  - b. Lower GI tract structures and functions:
    - 1) small intestine absorption of nutrients
    - 2) large intestine elimination of wastes
- 2. Accessory Organs of Digestion:
  - a. Salivary glands
  - b. Liver
  - c. Gall bladder
  - d. Pancreas

#### **B.** Factors Affecting Digestion

- 1. Enzymes
- 2. Peristalsis speed
- 3. Enteric Nervous System (part of ANS) regulates digestive tract motility, secretion and blood flow
- 4. Mucosa Layer = inner lining of GI tract:
  - a. Mucous cells secrete *mucous* to protect stomach lining; keeps stomach from digesting itself
  - b. Bicarbonate ion to *neutralize* acid

#### C. Common GI Disorders

- 1. Peptic Ulcer Diseases (PUD):
  - a. Description: erosion of mucosa layer of GI tract associated with acute inflammation (e.g. gastric or duodenal ulcers)
  - b. Causes:
    - 1) helicobacter pylori infection\*
    - 2) excessive acid secretion
    - 3) hyposecretion of adequate mucous protection
  - c. Risk factors:
    - 1) other drugs (e.g. NSAID's, corticosteroids)
    - 2) stress

- 3) smoking
- 4) caffeine
- 5) blood type O
- 6) family hx
- d. Consequences/Complications:
  - 1) acute GI bleed/hemorrhage
  - 2) anemia
- 2. Gastroesophageal Reflux Disease (GERD):
  - a. Description: acid contents of stomach move upward into esophagus
  - b. Causes: weakening of lower esophageal sphincter (no longer closes tightly)
  - c. Risk factors:
    - 1) obesity
    - 2) eating large meals or fatty or acid foods
    - 3) tobacco or alcohol use
    - 4) age >40 and infants
  - d. Consequences/Complications:
    - 1) esophagitis
    - 2) esophageal ulcers or strictures
- 3. Inflammatory Bowel Disease (IBD):
  - a. Cause: chronic inflammation of small and/or large intestines; idiopathic with possible autoimmune origin
  - b. Types:
    - 1) Crohn's Disease
    - 2) Ulcerative colitis
  - c. Risk factors:
    - 1) genetic predisposition
    - 2) possible dietary influences (promoting microbial overgrowth)
    - 3) stress
  - d. Consequences/Complications:
    - 1) GI ulcers/bleeding
    - 2) ↑ risk of colorectal cancer
    - 3) formation of GI fistulas, strictures and abscesses
    - 4) malnutrition
- 4. Irritable Bowel Syndrome (IBS):
  - a. Cause: unknown/unclear
  - b. Risk factors:
    - 1) intestinal infections
    - 2) stress
    - 3) dietary factors
  - c. Consequences/Complications:
    - 1) altered bowel elimination (alternating constipation or diarrhea)
    - 2) nutritional imbalances

- 5. Constipation:
  - a. Description: ↓ frequency or number and/or amount of stool/bowel movements (BMs)
  - b. Causes:
    - 1) lack of exercise/activity
    - 2) insufficient food intake
    - 3) ↓ fluid intake
    - 4) adverse effects of medications (e.g. opioids, anticholinergics, antihistamines, antacids, iron)
  - c. Risk factors:
    - 1) age
    - 2) chronic illnesses
    - 3) lifestyle
  - d. Consequences/Complications:
    - 1) overall discomfort
    - 2) fecal impaction  $\rightarrow$  bowel obstruction
    - 3) straining with defecation  $\rightarrow$  Valsalva maneuver with vagal response
- 6. Diarrhea:
  - a. Description: ↑ frequency and fluidity of bowel movement (BMs)
  - b. Cause:  $\downarrow$  reabsorption of water from fecal matter in large intestine
  - c. Risk factors:
    - 1) adverse effects of medications (e.g. antibiotics)
    - 2) bowel infections
    - chronic inflammatory bowel conditions (i.e. Crohn's Disease, ulcerative colitis)
    - 4) irritable bowel syndrome
  - d. Consequences/Complications:
    - 1) fluid loss
    - 2) electrolyte imbalances
    - 3) acid-base imbalances
- 7. Nausea:
  - a. Description: unpleasant subjective sensation accompanied by weakness, diaphoresis, hyper-production of saliva
  - b. Cause: stimulation of vomiting center:
    - 1) digestive tract via vagus nerve  $\rightarrow$  involve **serotonin** and **dopamine**
    - inner ear (vestibular apparatus) → involve *histamine*, and *muscarinic* receptors
    - chemoreceptor trigger zone (CTZ) in cerebral cortex → involve serotonin, dopamine, and muscarinic
  - c. Risk factors:
    - 1) GI infections/GI trauma
    - 2) food poisoning
    - 3) stress
    - 4) motion sickness

- 5) extreme pain
- 6) pregnancy
- 7) other medications (adverse effects)
- d. Consequences/Complications = vomiting/emesis:
  - 1) fluid loss
  - 2) electrolyte imbalances
  - 3) acid-base imbalances

#### **II. PHARMACOLOGY**

#### A. Pharmacological Connections for GI Drugs

- 1. When managing GI Disorders, often agents are given to treat/prevent.
  - a. **Cause**: e.g. PUD  $\rightarrow$  infection (h. pylori)
  - b. *Effect* = s/sx of GI upset or distress:

e.g.

- 1) abdominal pain
- 2) indigestion/heartburn
- 3) nausea/vomiting
- 4) anorexia/weight loss
- 5) bowel irregularity (constipation/diarrhea)
- c. **Complications**:
  - e.g.
  - 1) GI inflammation or bleeding ulcers
  - 2) fluid and electrolyte imbalances
  - 3) nutritional deficiencies/malnutrition
  - 4) fecal impaction  $\rightarrow$  bowel obstruction
- 2. *Goals* of Drug Therapy:
  - a. Eliminate cause(s) if possible
  - b. Reduce or eliminate GI upset or distress
  - c. Prevent consequences/complications:
    - 1) maintain weight
    - 2) maintain or restore normal fluid and electrolyte balance
    - 3) prevent acute GI bleeding

## Nursing Implications: GI Pharmacology: Antiulcer Agents

Subclass	MOA	Prototype – generic	Prototype – trade	A – Admin	T – ✓ Therapeutic Effects – General (MC)	A – ✓ Adverse Effects – Specific (SC)	T – Teaching – General (MC)	T – Teaching – Specific (SC)
histamine <sub>2</sub> receptor	Blocks histamine <sub>2</sub> - receptors in	rantidine	Zantac	Timing: pc/ HS	↓ s/sx PUD/GERD:	<ul> <li>GI: D/C</li> <li>CNS: drowsiness,</li> </ul>	<ul> <li>Take as directed</li> </ul>	<ul> <li>✓ CBC, liver &amp; renal fn tests</li> </ul>
antagonist/ blockers	stomach leading to ↓ acid production	famotidine	Pepcid		<ul> <li>↓ abdominal pain</li> <li>↓ heartburn -</li> </ul>	dizziness; confusion, restlessness,	<ul> <li>Diet: Ø foods that</li> </ul>	<ul><li>Ø antacids</li><li>Report: fever,</li></ul>
(H2RB)	(suppresses volume & acidity of parietal	cimetidine	Tagamet		<ul> <li>Ø c/o N/V</li> </ul>	hallucinations, depression	↑gastric acid	excessive bruising
	cell secretions)				Ø s/sx GI bleed:	• Heme: ↓ B <sub>12</sub>	secretion	
proton pump	Blocks (H+, K+-	omeprazole	Prilosec	Timing: ac	<ul> <li>(-) stool guaiac</li> <li>Ø blood in emesis</li> </ul>	<ul> <li>absorption</li> <li>GI: N/V/D/ abd. pain</li> </ul>	<ul> <li>Ø smoking or drinking</li> </ul>	Short-term use
inhibitor (PPI)	ATPase) enzyme			, and go are	or stool	• Other: rash, HA	<ul> <li>HOB ↑ for</li> </ul>	<ul> <li>✓ liver fn tests if</li> </ul>
	responsible for secreting	pantoprazole	Protonix		<ul> <li>improved blood counts:</li> </ul>	If used long-term: risk of gastric cancer	reflux • Report s/sx	<ul><li>taken long-term</li><li>Contraception</li></ul>
	hydrochloric acid leading to ↓ acid				o ↑ RBCs o ↑ Hgb		GI bleed	
	production (enzyme acts as 'pump' to				o ↑ Hct			
	release acid)							
antacids	Alkaline, inorganic substance that	aluminum hydroxide	AlternaGel, Amphojel	Timing: Do <u>not</u> give		<ul> <li>GI: N/V/C/D, abd. cramping, distention,</li> </ul>		<ul> <li>✓ renal function tests; electrolytes</li> </ul>
	neutralizes stomach	calcium	r -7-	within 2 hrs		belching		<ul> <li>Do <u>not</u> take w/ other</li> </ul>
	acid by raising pH (Does not promote	carbonate	Tums	meds; ac		<ul> <li>Electrolyte imbalances</li> </ul>		meds – space by 2 hrs
	healing of ulcer – nor eradicate	sodium				depending on compound used (i.e.		
	h. pylori)	bicarbonate				aluminum, Mg⁺⁺, `		
mucosal	Produces thick, gel-	sucralfate	Carafate	Timing: ac		Ca <sup>++</sup> , Na+) • GI: constipation, dry		Do not take 1-2 hrs of
protectants	like substance that coats ulcer,			& HS; space 1-2		mouth		other meds
	protecting from			hr of other				
	further erosion/ promoting healing;			meds				
	(does not â gastric							
	acid secretion nor eradicate h. pylori)							

Subclass	МОА	Prototype – generic	Prototype – trade	A – Admin	T – ✓ Therapeutic Effects – General (MC)	A – ✓ Adverse Effects – Specific (SC)	T – Teaching – General (MC)	T – Teaching – Specific (SC)
prostaglandin E analogs	Inhibits gastric acid secretions & stimulates production of protective mucous	misoprostol	Cytotec	With meals & HS		<ul> <li>GI: diarrhea, abd. cramping</li> </ul>		Contraception
Other meds: antiinfec- tives	<b>#1 drug to treat</b> <b>PUD</b> – Irradiates h. pylori organism	metronidazole clarithromycin amoxicillin	Flagyl Biaxin Amoxil	Refer to Ant	iinfective lecture			

## Nursing Implications: GI Pharmacology: Antiemetics/Prokinetics

		Prototype –	Prototype –		T – ✓ Therapeutic Effects – General	A –✓ Adverse Effects	T – Teaching –	T – Teaching –
Subclass	MOA	generic	trade	A – Admin	(MC)	– Specific (SC)	General (MC)	Specific (SC)
anticholinergics	Blocks action of acetylcholine → blocks impulses from vestibular apparatus of inner ear to vomiting center	scopolamine	Transderm Scop	Can use > 1 antiemetic – selection based on <i>emetogenic</i> <i>risk</i> Route: • dependent	<ul> <li>↓s/sx N/V</li> <li>Ø complications 2° N/V: dehydration, electrolyte imbalances</li> </ul>	ALL Antiemetics: CNS depression • Dry mouth, drowsiness, urinary retention, constipation	<ul> <li>Safety, ↓ activity</li> <li>Timing of meds – take before N/V</li> <li>Diet: NPO during acute vomiting;</li> </ul>	Related to managing adverse effects • Good oral care • Give fluids & fiber
antihistamines	Blocks <i>histamine</i> <sup>1</sup> & muscarinic receptors between inner ear & vomiting center	dimenhydrinate meclizine hydroxyzine	Dramimine Antivert Atarax, Vistaril	on nausea vs. vomiting PO route discouraged w/ active vomiting		<ul> <li>Dry mouth, drowsiness, urinary retention, constipation</li> </ul>	advance as tolerated • Avoid other CNS depressants; • ✓ labs 2° N/V	<ul> <li>Good oral care</li> <li>Give fluids &amp; fiber</li> </ul>
phenothiazines/ dopamine antagonists	Blocks <i>dopamine</i> receptors in brain → block signals to vomiting center	prochlorperazine metoclopramide	Compazine Reglan			<ul> <li>CV: hypotension (orthostatic), tachycardia</li> <li>EPS (dyskinesia, dystonia akathisia, Parkinsonism)</li> </ul>		• ✓ EPS • ✓ BP
serotonin anatagonists/ SSRIs	Block <i>serotonin</i> receptors in CTZ; & vagal neurons traveling from UGI tract to CTZ	ondansetron	Zofran			<ul> <li>GI: diarrhea/ constipation</li> <li>Other: dizziness, HA, fatigue</li> </ul>		<ul> <li>Monitor bowel function</li> </ul>
cannabinoids	Same active ingredient as marijuana; produces both antiemetic & relaxation effects – mechanism unknown	dronabinol nabilone	Marinol Cesamet			<ul> <li>Altered mood, cognition, perception, conjunctivitis; potential for abuse</li> </ul>		<ul> <li>Watch for habit forming</li> </ul>
Substance P/ neurokinin 1 antagonist (SPA)	Inhibits substance P in brain (vomiting center in medulla contains high conc. of substance P; their activation stimulates vomiting reflex)	aprepitant	Emend			<ul> <li>Fatigue, diarrhea, dizziness, hepatotoxicity</li> </ul>		<ul> <li>✓ liver enzymes</li> </ul>

Subclass	МОА	Prototype – generic	Prototype – trade	A – Admin	T – ✓ Therapeutic Effects – General (MC)	A –✔ Adverse Effects – Specific (SC)	T – Teaching – General (MC)	T – Teaching – Specific (SC)
Others (FYI): benzodiazepine corticosteroids		lorazepam	Ativan	Refer to Neuro a	nd Endocrine Pharma	acology		
MAJOR CLASS: PROKINETIC/ GI stimulants	<ul> <li>TWO (2) MOAs:</li> <li>1. blocks <i>dopamine</i> &amp; serotonin receptors in CTZ;</li> <li>2. parasympathomimetic – mimics actions of <i>Ach</i> → <i>cholinergic</i></li> </ul>	metoclopramide	Reglan	(See previous page)	<ul> <li>↓ N/V &amp; other complications</li> <li>↓s/sx of GERD</li> <li>Improved GI function – +BS, +BM</li> </ul>	<ul> <li>GI: diarrhea</li> <li>+ EPS; CNS depression/ sedation</li> </ul>	<ul> <li>Similar w/ other antiemetics – watch use of other ANS drugs</li> </ul>	<ul> <li>✓ bowel fn</li> <li>✓ EPS</li> </ul>

## Nursing Implications: GI Pharmacology: Laxatives

Subclass Bulk-forming/ mechanical stimulants (can also be used for diarrhea)	MOA Absorbs water → add size to fecal mass → ↑ defecation reflex r/t bowel wall stretching	Prototype – generic psyllium muciloid	Prototype – trade Metamucil	A – Admin Route: PO Mix w/ 8 oz of water – give <u>immediately</u>	<ul> <li>T - ✓ Therapeutic Effects - General (MC)</li> <li>↓ s/sx constipation:</li> <li>+ BM</li> <li>soft-formed stools</li> <li>no abd pain or fullness or distention</li> </ul>	<ul> <li>A - ✓ Adverse Effects - General (MC)</li> <li>GI: diarrhea, abd cramping, nausea</li> </ul>	<ul> <li>A – ✓ Adverse Effects – Specific (SC)</li> <li>Aspiration, esophageal obstruction due to thickened drink</li> </ul>	<ul> <li>T – Teaching – General (MC)</li> <li>Take as directed – no overuse</li> <li>Hold for diarrhea</li> <li>Diet: high fiber, fluids</li> <li>Activity as</li> </ul>	<ul> <li>T – Teaching – Specific (SC)</li> <li>Mix well – drink promptly</li> <li>Aspiration precaution as indicated</li> </ul>
stool softener/ surfactant/ lubricants	↑absorption of water & fat in stools → aids in easier stool passage by serving as a 'wetting agent' (lowers surface tension)	docusate sodium	Colace	Route: PO Freq: Take daily	Ø complications assoc. w/ constipation: • no straining (Valsalva maneuver)			tolerated • NOTE: laxatives may interfere w/ absorp- tion of other meds	
stimulant (chemical)	Irritates bowel to increase peristalsis by directing & stimulating nerve plexus in intestinal wall	bisacodyl senna tabs (herbal agent)	Dulcolax Senakot, Ex-Lax	Route: PO – Don't chew; if given pr – insert finger length			Laxative     dependency		<ul> <li>Contraception</li> <li>Do <u>not</u> take w/ milk or antacids</li> </ul>
osmotics	Not absorbed in intestine; pulls water into fecal mass to create more watery stool	polyethylene glycol magnesium hydroxide lactulose	Milk of Magnesia Miralax	Route: PO Timing: Give on empty stomach			<ul> <li>Electrolyte imbalances, esp. Mg++</li> <li>Dehydration or fluid retention</li> <li>Used as an ammonium detoxicant</li> </ul>		<ul> <li>Monitor labs as indicated, esp. w/ h/o renal &amp; liver dz</li> <li>Increase fluids</li> <li>Safety during bowel cleansing prep</li> </ul>

Subclass	MOA	Prototype – generic	Prototype – trade	A – Admin	T – ✓ Therapeutic Effects – General (MC)	A – ✓ Adverse Effects – General (MC)	A – ✓ Adverse Effects – Specific (SC)	T – Teaching – General (MC)	T – Teaching – Specific (SC)
Miscellaneous laxatives	Lubricates stool & colon mucosa	mineral oil		Route: PO Do <u>not</u> give w/ other meds or vitamins	(See previous page)	(See previous page)	<ul> <li>Interferes w/ absorption of fat-soluble vitamins</li> </ul>	(See previous page)	<ul> <li>Discourage use of mineral oil for treating constipation</li> <li>Do <u>not</u> take w/ other meds or vitamins</li> </ul>

## Nursing Implications: GI Pharmacology: Antidiarrheals & Miscellaneous GI Drugs

Major Class/ Subclass	MOA	Prototype – generic	Prototype – trade	A – Admin	T – ✓ Therapeutic Effects – General (MC)	A – ✓ Adverse Effects – Specific (SC)	T – Teaching – General (MC)
Antidiarrheal: Opioid	Acts directly on intestine to slow peristalsis →↑ absorption of fluids & electrolytes	diphenoxylate w/ atropine loperamide	Lomotil Imodium	Route: PO Schedule V	<ul> <li>↓# of BMs</li> <li>↓abd. pain, cramping</li> <li>↑formed stools</li> </ul>	<ul> <li>GI: constipation</li> <li>CNS: dizziness, drowsiness</li> <li>Other: anticholinergic effects</li> </ul>	<ul> <li>Take as directed – short term use</li> <li>Encourage fluids</li> <li>Report worsening GI sx</li> <li>Safety</li> <li>Avoid CNS depressants</li> <li>Labs: lytes, renal, liver function tests</li> </ul>
Anti-IBD: NSAIDs – Step 1	↓ inflammation by inhibiting prostaglandin synthesis	sulfasalazine	Azulfidine	Route: PO • Consider contraindications of NSAIDs	<ul> <li>↓inflammation →</li> <li>↓# of BMs &amp;</li> <li>↓abd pain, cramping</li> </ul>	<ul> <li>GI: N/V/D, dyspepsia, abd pain</li> <li>Blood: dycrasias</li> <li>Other: hepatotoxicity</li> </ul>	<ul> <li>Take as directed</li> <li>Report worsening GI sx</li> <li>Labs: CBC, liver enzymes</li> <li>DDIs</li> </ul>
Anti-IBD: Corticosteroids – Step 2	↓ inflammation by increasing cortisol	prednisone budesonide	Deltasone Entocort – EC, Uceris	Refer to Endocrine F	Pharmacology	· · · · ·	
Anti-IBD: Immunosuppressant – Step 3	Suppresses immune response	azathioprine methotrexate	Imuran MTX, Rheumatrex, Trexall	Refer to Immune Ph	armacology		
Anti-IBS: Diarrhea form	Blocks serotonin receptors in colon → relaxes colon to slow movement → ↓ urgency/freq of BMs	a <b>los</b> etron ("loose stools")	Lotronex (name similar to Lomotil → antidiarrheal)	Route: PO	<ul> <li>↓# of BMs</li> <li>↓ urgency/freq of stools</li> </ul>	<ul> <li>GI: constipation</li> <li>GI toxicity: ischemic colitis, bowel obstruction, impaction or perforation</li> </ul>	<ul> <li>Take as directed</li> <li>Report worsening GI sx</li> <li>Diet: limit/eliminate caffeine, alcohol, milk products, high sugar, fatty food, gas-producing, artificial sweeteners</li> <li>Stress mgmt.</li> </ul>

Major Class/ Subclass	MOA	Prototype – generic	Prototype – trade	A – Admin	T – ✓ Therapeutic Effects – General (MC)	A – ✓ Adverse Effects – Specific (SC)	T – Teaching – General (MC)
Anti-IBS: Constipation form	↑ fluid secretion in intestine to promote motility	<b>lub</b> iprostone ("lubricate" bowel)	Amitiza	Route: PO w/ food	<ul> <li>Relief of constipation</li> <li>Return of normal bowel function</li> </ul>	• GI: N/diarrhea; abd. pain	<ul> <li>Take as directed w/ food</li> <li>Report worsening GI sx</li> <li>Diet: add fiber, water, exercise</li> <li>Stress mgmt.</li> <li>Support groups</li> </ul>
Antiinflammatory/ Antacid	↓ inflammation & acts as antacid	bismuth subsalicylate	Pepto-Bismol	Route: PO – liquid, tablets	<ul> <li>↓ inflammation →</li> <li>↓ # of BMs &amp;</li> <li>↓ abd pain, cramping</li> </ul>	<ul> <li>GI: blacken stools &amp; tongue</li> <li>Salicylism: long- term use</li> </ul>	<ul> <li>Take as directed</li> <li>Report worsening GI sx</li> <li>Do <u>not</u> give to children</li> </ul>
Adsorbents	Preferred method for removing ingested poisons from the GI tract by adhering to drug molecules :→ prevents their absorption into the blood → eliminates in stools	charcoal tablets		Route: PO (fine powder consistency) Adults: 60-100 gm Children: 15-30 gm Contraindications: bowel perforation or obstruction	<ul> <li>↓ Poison absorption &amp; toxicity</li> </ul>	● GI: ↑ BMs	<ul> <li>Mix w/ water for oral administration</li> <li>Administer within 30 minutes after poison ingestion (best results)</li> <li>Do <u>not</u> give antidote immediately before, with, or shortly after the charcoal</li> </ul>
Antiflatulents	↓ gas bubbles → allows easier passing of flatulence (antifoaming agent)	simethicone	Mylicon, Gas-X	Route: PO	<ul> <li>Relief of bloating &amp; abdominal discomfort</li> </ul>	<ul> <li>GI: minimal as it is not absorbed in body; mild N/V/D</li> </ul>	<ul> <li>Take as directed w/ food</li> <li>Report worsening GI sx</li> </ul>

## Nursing Implications: GI Pharmacology: Nutritional Aids

Major Class	MOA	Prototype – generic	Prototype – trade	A – Admin	T – ✓ Therapeutic Effects – General (MC)	A – ✓ Adverse Effects – Specific (SC)	T – Teaching – GENERAL (MC)
Digestive enzymes	Pancreatic enzymes	pancrelipase [lipase, alpha- galactosidase]	Creon, Viokase, Pancrease	Route: PO – capsule (delayed release), powder, tablet Should be taken w/ each meal & snack	Improvement in nutritional markers – esp. fat digestion • ↓fatty stools	<ul> <li>GI: N/V/D</li> <li>F&amp; E: hyperuricemia (high doses)</li> </ul>	<ul> <li>DDI: iron, antacids</li> <li>✓ uric acid levels</li> <li>Use w/ caution in DM:</li> <li>✓ BG</li> </ul>
Probiotics	Nonherbal dietary – maintains intestinal health (lactobacilli – normal components of gut flora)	lactobacillus acidophilus		Route: PO – daily	<ul> <li>Improvement in food metabolism; promote nutrient absorption</li> <li>Reduce diarrhea caused by clostridium difficile (c. diff colitis)</li> </ul>	<ul> <li>GI: flatulence, bloating (generally well-tolerated) if taken in recommended doses</li> </ul>	Maintain balanced diet