

## Gastrointestinal Drugs

### Subcommittee:

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### Recommended Curriculum Equivalent: 1.5 h

Acid Reducers and Drugs for the Treatment of Peptic Ulcer Disease	
Proton pump inhibitors	
First generation	Second generation
OMEPRAZOLE	ESOMEPRAZOLE LANSOPRAZOLE PANTOPRAZOLE RABEPRAZOLE
Learning Objectives	
<b>Physiology and pathophysiology</b> Describe the synthesis and mechanism of H <sup>+</sup> secretion by the parietal cells	
<b>Mechanism of action</b> Describe the mechanism of action of proton pump inhibitors and why they are selective for the parietal cell proton pump.	
<b>Actions on organ systems</b> Describe the pharmacological effects of the drugs on gastric function. Are there effects on other organ systems?	
<b>Pharmacokinetics</b> Describe the pharmacokinetics of proton pump inhibitors? Are there significant differences among the different drugs in this class?	
<b>Adverse effects, drug interactions and contraindications</b> Describe the principal adverse effects of proton pump inhibitors. Describe the clinically important drug interactions of proton pump inhibitors. Describe the principal contraindications of proton pump inhibitors.	
<b>Therapeutic uses</b> Describe the current therapeutic uses of proton pump inhibitors.	

**Clinical Pharmacology**

Omeprazole is perceived to be the most potent of this drug class in inhibiting CYP2C19 activity and is proposed to have potential drug interactions with other drugs metabolized by this P450 isoform. Concern has been raised about potential inhibition of clopidogrel activation in patients taking both drugs concurrently.

Current consensus is that in such patients clopidogrel with pantoprazole may be a safer choice to reduce the probability of a drug interaction involving CYP2C19.

Prolonged use of PPI drugs is associated with hypomagnesemia. This possibility is a concern in debilitated patients especially in those having underlying cardiac arrhythmias and/or osteoporosis where associated muscle weakness might increase risk of a fall and subsequent fracture.

**Relevance****USMLE topic**

**Gastrointestinal System, Normal Processes** – organ structure and function including digestion; cell/tissue structure & function including endocrine and neural regulatory functions; gastrointestinal secretory products

**Principles of therapeutics**

Mechanisms of action and use of drugs for treatment of disorders of the gastrointestinal system; treatment and prophylaxis of peptic ulcer disease and gastroesophageal reflux

**AAMC Medical School Objectives**  
**Project Report X Patient Safety-Table 1**

**Topic C**—Drug treatment of common conditions and diseases, using frequently prescribed classes of drugs for the treatment and prevention of disease

<b>Acid Reducers and Drugs for the Treatment of Peptic Ulcer Disease</b>		
Endogenous Substance	H <sub>2</sub> Receptor Antagonists	
	First Generation	Second Generation
histamine	cimetidine	FAMOTIDINE NIZATIDINE RANITIDINE
<b>Learning Objectives</b>		
<b>Physiology and pathophysiology</b> Describe the neurohumoral control of H <sup>+</sup> secretion by gastric parietal cells. Describe the role of histamine in the different phases H <sup>+</sup> secretion. Describe the causes of H <sup>+</sup> hypersecretion.		
<b>Mechanism of action</b> Explain the molecular mechanism of action H <sub>2</sub> receptor antagonists.		
<b>Actions on organ systems</b> Describe the pharmacological effects of the drugs on the stomach. Do these antagonists have effects on other organ systems?		
<b>Pharmacokinetics</b> Describe the pharmacokinetics of the H <sub>2</sub> receptor antagonists.		
<b>Adverse effects, drug interactions and contraindications</b> Describe the principal adverse effects of each H <sub>2</sub> receptor antagonist. Describe the clinically important drug interactions of H <sub>2</sub> receptor antagonists. Describe the principal contraindications of H <sub>2</sub> receptor antagonists.		
<b>Therapeutic uses</b> Identify current therapeutic uses of H <sub>2</sub> receptor antagonists.		
<b>Clinical Pharmacology</b> Tolerance and loss of efficacy occurs with prolonged use of drugs in this class. Due to its ability to inhibit activity of many CYP isoforms, cimetidine is not the preferred drug in this class when administered concurrently with other drugs whose elimination is cytochromes P450-dependent. Use of ranitidine or famotidine is considered preferable in such patients.		
<b>Relevance</b>		
<b>USMLE topic</b> <b>Gastrointestinal System, Normal Processes</b> – organ structure and function including digestion; cell/tissue structure & function including endocrine and neural regulatory functions; gastrointestinal secretory products <b>Abnormal Processes</b> – idiopathic disorders	<b>Principles of therapeutics</b> Mechanisms of action and use of drugs for treatment of disorders of the gastrointestinal system; treatment and prophylaxis of peptic ulcer disease and gastroesophageal reflux	

<b>AAMC Medical School Objectives</b> <b>Project Report X Patient Safety-Table 1</b>	<b>Topic C</b> —Drug treatment of common conditions and diseases, using frequently prescribed classes of drugs for the treatment and prevention of disease
<b>Notes</b>	

<b>Acid Reducers and Drugs for the Treatment of Peptic Ulcer Disease</b>		
<b>Acid Neutralizers (Classic Antacid preparations)</b>		
Endogenous substance	Single agent	Mixed preparations
H <sup>+</sup>	ALUMINUM HYDROXIDE CALCIUM CARBONATE MAGNESIUM HYDROXIDE sodium bicarbonate	ALUMINUM HYDROXIDE MAGNESIUM HYDROXIDE/
<b>Learning Objectives</b>		
<b>Physiology and Pathophysiology</b> Describe the mechanisms of H <sup>+</sup> secretion in the stomach		
<b>Mechanism of action</b> Describe the mechanism of action of antacid medications. Describe the differences in onset and duration of action of each antacid preparation.		
<b>Actions on organ systems</b> Describe the pharmacological effects of the drugs in each class on the stomach.		
<b>Pharmacokinetics</b> Describe the absorption and systemic actions of antacid preparations		
<b>Adverse effects, drug interactions and contraindications</b> Describe the principal adverse effects of each antacid preparation. Describe the clinically important drug interactions with antacids. Describe the principal precautions and contraindications in the use of antacids.		
<b>Therapeutic uses</b> Describe the primary indication of antacid use.		
<b>Clinical Pharmacology</b> The main concern with these antacids is chelation of co-administered drugs that may be interpreted as disease progression due to reduction of systemic bioavailability of these other agents. Some older patients ingest calcium carbonate tablets as a cheap source of calcium supplementation as a means to preserve bone mass. There is no good evidence to support the claim that long-term use of antacids prevents to occurrence of peptic ulcers.		
<b>Relevance</b>		

<b>USMLE topic</b> <b>Gastrointestinal System, Normal Processes</b> – organ structure and function including digestion; cell/tissue structure & function including endocrine and neural regulatory functions; gastrointestinal secretory products <b>Abnormal Processes</b> – idiopathic disorders	<b>Principles of therapeutics</b> Mechanisms of action and use of drugs for treatment of disorders of the gastrointestinal system; treatment and prophylaxis of peptic ulcer disease and gastroesophageal reflux
<b>AAMC Medical School Objectives Project Report X Patient Safety-Table 1</b>	<b>Topic C</b> Drug treatment of less common but severe conditions
<b>Notes</b>	

<b>Drugs for the Treatment of Peptic Ulcer Disease</b>		
Cytoprotectant agents		
Endogenous substance	Analog	Surface protectant
PGE <sub>2</sub>	MISOPROSTOL	SUCRALFATE
<b>Learning Objectives</b>		
<b>Physiology and Pathophysiology</b> Describe the mechanisms for production of the gastric cytoprotective barrier. Describe causes for disruption of the cytoprotective barrier.		
<b>Mechanism of action</b> Explain the mechanism of action of each drug.		
<b>Actions on organ systems</b> Describe the pharmacological effect of the each drug on the cytoprotective barrier.		
<b>Pharmacokinetics</b> Describe the absorption, distribution metabolism and excretion of each drug.		
<b>Adverse effects, drug interactions and contraindications</b> Describe the principal adverse effects of each drug. Describe clinically important drug interactions of the drugs in each class.		
<b>Therapeutic uses</b> Describe the primary indications for use of each drug.		
<b>Clinical Pharmacology</b> As a prostaglandin analog, misoprostol is contraindicated in pregnancy. Its dose-related GI toxicity has relegated its use to second-line strategy. In patients with renal impairment, hypophosphatemia and aluminum intoxication become a concern with prolonged use of sucralfate.		
<b>Relevance</b>		

<p><b>USMLE topic</b>  <b>Gastrointestinal System, Normal Processes</b> – organ structure and function including digestion; cell/tissue structure &amp; function including endocrine and neural regulatory functions; gastrointestinal secretory products; gastrointestinal defense mechanisms  <b>Abnormal Processes</b> – drug-induced adverse effects on the gastrointestinal system</p>	<p><b>Principles of therapeutics</b>  Mechanisms of action and use of drugs for treatment of disorders of the gastrointestinal system; treatment of peptic ulcer disease</p>
<p><b>AAMC Medical School Objectives</b>  <b>Project Report X Patient Safety-Table 1</b></p>	<p><b>Topic C</b>  Drug treatment of less common but severe conditions</p>
<p><b>Notes</b></p>	

Drugs for the Treatment of Peptic Ulcer Disease				
Helicobacter pylori eradication				
Triple therapy		Quadruple therapy		
Antibiotics	Acid suppression	Antibiotics	Acid suppression	other
AMOXICILLIN CLARITHROMYCIN METRONIDAZOLE TETRACYCLINES	PPI	CLARITHROMYCIN TETRACYCLINES	H <sub>2</sub> - BLOCKERS PPI	BISMUTH SUBSALICYLATE
<b>Physiology and Pathophysiology:</b> Describe the role of H. pylori in peptic ulcer disease. Describe tests for evaluating H. pylori infection.				
<b>Therapeutic uses:</b> Describe the use of triple and quadruple therapy regimens used for H. pylori eradication. Describe factors to be used in selecting the best therapeutic options for a given patient.				
<b>Mechanism of action:</b> Describe the contribution of each agent in triple or quadruple therapy regimens in H. pylori eradication.				
<b>Drug interactions</b> Describe potential drug interactions. Describe potential for antibiotic resistant strains of H. pylori.				
<b>Clinical Pharmacology</b> Amoxicillin, clarithromycin and a PPI is the most commonly recommended triple therapy regimen. With initial treatment failure, bismuth subsalicylate is often added to the second attempt at eradication and the approach is then referred to as quadruple therapy. Bismuth subsalicylate ingestion has been associated with salicylate intoxication, especially in children, and in adults ingesting other salicylate-containing drug preparations concurrently.				
Relevance				
<b>USMLE topic</b> <b>Gastrointestinal System, Normal Processes</b> – organ structure and function including digestion; cell/tissue structure & function including endocrine and neural regulatory functions; gastrointestinal secretory products <b>Abnormal Processes</b> – infectious disorders		<b>Principles of therapeutics</b> Mechanisms of action and use of drugs for treatment of disorders of the gastrointestinal system; treatment and prophylaxis of peptic ulcer disease		
<b>AAMC Medical School Objectives</b> <b>Project Report X Patient Safety-Table 1</b>		<b>Topic C</b> Drug treatment of common conditions and diseases, using frequently prescribed classes of drugs for the treatment and prevention of disease		

<b>Prokinetic Drugs and Laxatives</b>		
<b>Recommended Curriculum Equivalent: 1 hr</b>		
<b>Drug Classes and Drugs to consider</b>		
Drugs used to treat upper GI motility disorders (Gastroparesis, dyspepsia, GERD)	Drugs used to treat lower GI motility disorders (Constipation)	
<b>METOCLOPRAMIDE</b> cisapride domperidone erythromycin tegaserod	Receptor mediated	Laxatives
	<b>ALVIMOPAN</b> <b>LUBIPROSTONE</b> <b>METHYLNALTREXONE</b> bethanechol neostigmine renzapride tegaserod	<b>DOCUSATE</b> <b>LACTULOSE</b> <b>METHYLCELLULOSE</b> <b>POLYETHYLENE GLYCOL</b> <b>SODIUM PHOSPATE</b> <b>SODIUM CITRATE</b> <b>PSYLIUM</b> castor oil, bisacodyl, senna, cascara, mineral oil
<b>Learning Objectives</b>		
<b>Physiology and Pathophysiology</b> Describe the neural and hormonal mechanisms controlling stomach and intestinal motility Describe the changes in neural and hormonal control of stomach and intestinal motility that lead to delayed gastric emptying or accommodation.		
<b>Mechanisms of action</b> Explain the molecular mechanism of action of each drug.		
<b>Actions on organ systems</b> Describe why some drugs are selective for upper GI motility disorders and why others are selective for lower GI motility disorders.		
<b>Pharmacokinetics</b> Describe the relevant pharmacokinetic features of each drug		
<b>Adverse effects, drug interactions and contraindications</b> Describe the principal adverse effects of the drugs of each class. Describe the clinically important drug interactions of the drugs of each class Describe the principal contraindications of the drugs of each class		
<b>Therapeutic uses</b> Outline the main therapeutic uses of the drugs of each class.		

## Clinical Pharmacology

Cisapride is no longer favored because of its potent inhibition of CYP3A, the cytochrome P450 isoform responsible for metabolism of a large proportion of drugs ingested by patients, and the fact that it is associated with prolongation of the QTc interval.

Drugs to treat constipation are frequently abused by the general population. Use should be progressive from stool softeners (docusate) to fibre supplementation (methylcellulose) and finally to propulsive agents. Recently polyelectrolyte lavage solution has been found to be acceptable with less cramping than seen with the use of castor oil.

## Relevance

### USMLE topic

**Laxatives: Gastrointestinal System, Normal Processes** – organ structure and function including motility, digestion and absorption; cell/tissue structure & function including endocrine and neural regulatory functions; gastrointestinal defense mechanisms and normal flora

**Abnormal Processes** – infection, inflammatory and immunologic disorders; systemic disorders affecting the gastrointestinal system; drug-induced adverse effects on the gastrointestinal system; idiopathic disorders

**Prokinetics: Gastrointestinal System, Normal Processes** – organ structure and function including motility; cell/tissue structure & function including endocrine and neural regulatory functions **Abnormal Processes** – systemic disorders affecting the gastrointestinal system; idiopathic disorders

### Principles of therapeutics

**Laxatives:** mechanisms of action and use of drugs for treatment of disorders of the gastrointestinal system; drugs to alter gastrointestinal motility

**Prokinetics:** mechanisms of action and use of drugs for treatment of disorders of the gastrointestinal system; drugs to alter gastrointestinal motility; treatment and prophylaxis of gastroesophageal reflux

**AAMC Medical School Objectives  
Project Report X Patient Safety-Table 1**

**Topic C**  
Drug treatment of common conditions and diseases, using frequently prescribed classes of drugs for the treatment and prevention of disease

### Notes

Cisapride is available in the U.S. only in special cases.

Tegaserod has been removed from the U.S. market and is now available only in emergency cases requiring special FDA approval.

Domperidone is available in Europe and Canada but not the U.S. However, patients in the U.S. have access to this drug via the internet or by travel to Europe and Canada.

<b>Anti-diarrheal drugs</b>			
<b>Recommended Curriculum Equivalent: 0.5 hr</b>			
<b>Drug Classes and Drugs to consider</b>			
Opioid- compounds	Serotonergic- compounds	Adrenergic compounds	Probiotics
LOPERAMIDE diphenoxylate	alosetron cilansetron	clonidine	bifidobacterium infantis
somatostatin analog	muscarinic antagonists		other
octreotide	ATROPINE DICYCLOMINE hyoscyamine scopolamine	BISMUTH SUBSALICYLATE bismuth citrate	
<b>Learning Objectives</b>			
<b>Physiology and Pathophysiology:</b>			
Describe the neural and hormonal mechanisms controlling colonic motility and water and electrolyte absorption and secretion.			
Describe the conditions under which neural mechanisms controlling colonic motility and water and electrolyte absorption and secretion are impaired.			
Describe the neural mechanisms of visceral sensation and visceral pain.			
Describe the importance of maintaining normal gut flora and how disruption can lead to altered motility and absorption and secretion in the colon.			
<b>Mechanisms of action:</b>			
Explain the molecular mechanism of action of each drug in each drug class.			
<b>Actions on organ systems:</b>			
Describe the effects of each drug on the colon and also on other organ systems.			
<b>Pharmacokinetics:</b>			
Describe the absorption distribution metabolism and secretion of each drug.			
<b>Adverse effects, drug interactions and contraindications:</b>			
Describe the principal adverse effects of the drugs of each class.			
Describe the clinically important drug interactions of the drugs of each class			
Describe the principal contraindications of the drugs of each class			
<b>Therapeutic uses</b>			
Identify the specific therapeutic applications of each class of drug.			

**Clinical Pharmacology**

Adequate replacement of fluid and electrolytes remains the primary approach to management so that clearance of the offending intestinal stimulant is removed.

Loperamide represents the primary opioid receptor agonist drug intervention that has minimal effects on the central nervous system. It is often useful in patients with chronic diarrhea secondary to treatment with chemotherapeutic drugs.

Use of bismuth subsalicylate is discouraged because of its potential to cause salicylate intoxication if it is used chronically. Other drug interventions are considered second line and rarely used because of associated adverse reactions.

**Relevance****USMLE topic****Gastrointestinal System, Normal**

**Processes** – organ structure and function including motility, digestion and absorption; cell/tissue structure & function including endocrine and neural regulatory functions; gastrointestinal defense mechanisms and normal flora

**Abnormal Processes** – infection, inflammatory and immunologic disorders; systemic disorders affecting the gastrointestinal system; drug-induced adverse effects on the gastrointestinal system; idiopathic disorders

**Principles of therapeutics**

Mechanisms of action and use of drugs for treatment of disorders of the gastrointestinal system; drugs to alter gastrointestinal motility

**AAMC Medical School Objectives****Project Report X Patient Safety-Table 1****Topic C**

Drug treatment of common conditions and diseases, using frequently prescribed classes of drugs for the treatment and prevention of disease

**Notes:**

Drugs used for the Treatment of Inflammatory Bowel Disease			
Recommended Curriculum Equivalent: 0.5 hr			
Drug Classes and Drugs to consider			
Salicylates	Steroids	Anti-mitotic agents	Monoclonal antibodies
5-AMINO SALICYLIC ACID SULFAPYRIDINE SULFASALAZINE balsalazide	HYDROCORTISONE METHYLPREDNISOLONE PREDNISONE budesonide	6-mercaptopurine azathioprine cyclosporine methotrexate	INFLIXIMAB
Antibiotics		Probiotics	
ciprofloxacin clarithromycin metronidazole		VSL #2 lactobacillus	
<b>Learning Objectives</b>			
<p><b>Pathophysiology:</b> Describe the differences between ulcerative colitis and Crohn's disease. Describe the mechanisms responsible of intestinal and extraintestinal symptoms of inflammatory bowel disease. Describe the contribution of intestinal bacteria to the pathophysiology of inflammatory bowel disease.</p>			
<p><b>Mechanism of action</b> Describe the mechanism of action of each of the major classes of drugs.</p>			
<p><b>Pharmacokinetics</b> List the routes of administration of drugs in each class. Describe the absorption and distribution of each class of drug and how this impacts on the choice of the route of administration. Describe the mechanisms for bioactivation of the salicylates and how this impacts on their use for the treatment of inflammatory bowel disease.</p>			
<p><b>Adverse effects, drug interactions and contraindications</b> Describe the main adverse effects of the drugs of each class. Describe the clinically important drug interactions of the drugs of each class. Describe the principal contraindications or precautions of the drugs of each class.</p>			
<p><b>Therapeutic uses</b> Know the selective use of each class of drug for the treatment of ulcerative colitis vs. Crohn's disease.</p>			

<p><b>Clinical Pharmacology</b>  Subsequent to the finding of efficacy of sulfasalazine, it was determined that efficacy was mediated mostly by the 5-aminosalicylic acid that was released by sulfasalazine hydrolysis in the lower GI tract. Thus patients intolerant to sulfonamide drugs can be given 5-aminosalicylate directly to eliminate exposure to the sulfonamide moiety.  From a steroid perspective, prednisone is most cost effective with change to methylprednisolone in patients with hepatic impairment.  Use of infliximab remains a second line intervention for those patients having failed more established drug therapies.  Side effects from TNF-alpha inhibitions are significant, especially related to the increased susceptibility to infection.</p>	
<b>Relevance</b>	
<p><b>USMLE topic</b>  <b>Gastrointestinal System, Normal Processes</b> – organ structure and function including motility, digestion and absorption; cell/tissue structure &amp; function including endocrine and neural regulatory functions; gastrointestinal defense mechanisms and normal flora  <b>Abnormal Processes</b> – inflammatory and immunologic disorders; systemic disorders affecting the gastrointestinal system; idiopathic disorders</p>	<p><b>Principles of therapeutics</b>  Mechanisms of action and use of drugs for treatment of disorders of the gastrointestinal system; anti-inflammatory, immunosuppressive drugs</p>
<p><b>AAMC Medical School Objectives</b>  <b>Project Report X Patient Safety-Table 1</b></p>	<p><b>Topic C</b>  Drug treatment of common conditions and diseases, using frequently prescribed classes of drugs for the treatment and prevention of disease</p>
<p><b>Notes</b>  Objectives for salicylates and steroids are covered under Analgesic, Antipyretic, Antiinflammatory Drugs  Objectives for mitotic inhibitors are covered under Chemotherapy Drugs</p>	

Drugs used to Treat Nausea and Vomiting	
<b>Recommended Curriculum Equivalent: 1 hr</b>	
<b>Drug Classes and Drugs to Consider</b>	
<b>Emetic drugs</b>	
Dopamine receptor agonist	Non-selective emetics
Apomorphine	Syrup of Ipecac

<b>Anti-Emetic Drugs</b>			
Dopamine receptor antagonists	5-HT <sub>3</sub> receptor antagonists	Cannabinoid receptor antagonists	Histamine receptor antagonists
METOCLOPRAMIDE PROCHLORPERAZINE haloperidol	DOLASETRON GRANISETRON ONDANSETRON PALONOSETRON ramosetron	DRONABINOL nabilone	DIMENHYDRINATE DIPHENHYDRAMINE cyclizine hydroxyzine meclizine promethazine
Neurokinin receptor antagonists	Corticosteroids	Benzodiazepines	Muscarinic receptor antagonists
APREPITANT	METHYL - PREDNISOLONE PREDNISON dexamethasone	LORAZEPAM ALPRAZOLAM DIAZEPAM	SCOPOLAMINE
<b>Learning Objectives</b>			
<b>Physiology and Pathophysiology</b> Describe the central and peripheral nervous system mechanisms responsible for nausea and vomiting.			
<b>Mechanisms of action</b> Describe the mechanism of action of emetic drugs. Explain the cellular and molecular mechanisms of action of each drug class. Describe the use of multi-drug treatment of nausea and vomiting.			
<b>Actions on organ systems</b> Describe the pharmacological effects of each drug in each class.			
<b>Pharmacokinetics</b> Know the absorption, distribution, metabolism and excretion of each drug class.			
<b>Adverse effects, drug interactions and contraindications</b> Describe the principal adverse effects of the drugs of each class. Describe the clinically important drug interactions of the drugs of each class Describe the principal contraindications of the drugs of each class			
<b>Therapeutic uses</b> Know the appropriate uses of anti-emetic drugs for the treatment of specific conditions such as chemotherapy-induced nausea and vomiting, postoperative nausea and vomiting, motion sickness, vertigo, and nausea and vomiting during pregnancy.			

<p><b>Clinical Pharmacology</b>  Syrup of ipecac is no longer recommended as an antiemetic, because of its lack of efficacy relative to the time of poison ingestion. It also has a potent sedative effect that could increase risk of aspiration pneumonia after induction of emesis.  Response to antiemetic drug treatment is highly variable among patients. These drugs are used most often in conjunction with chemotherapy, and are usually specified by protocol. Administration of dimenhydrinate or diphenhydramine should be done with caution because of their risk of abuse, and the narrow therapeutic index between efficacy and substantial toxicity.  The scopolamine patch is an effective antinauseant intervention and is successfully used for space travelers.</p>	
<b>Relevance</b>	
<p><b>USMLE topic</b>  <b>Gastrointestinal System, Normal Processes</b> – organ structure and function including motility; <b>Central and Peripheral Nervous Systems</b> – brain stem; blood-brain barrier  <b>Abnormal Processes</b> – idiopathic disorders; systemic disorders affecting the gastrointestinal system; drug-induced adverse effects on the gastrointestinal system</p>	<p><b>Principles of therapeutics</b>  Mechanisms of action and use of drugs for treatment of disorders of the gastrointestinal system; drugs to alter gastrointestinal motility; drugs affecting the autonomic nervous system</p>
<p><b>AAMC Medical School Objectives</b>  <b>Project Report X Patient Safety-Table 1</b></p>	<p><b>Topic C</b>  Drug treatment of common conditions and diseases, using frequently prescribed classes of drugs for the treatment and prevention of disease</p>
<b>Notes</b>	

Miscellaneous Gastrointestinal Drugs		
Antigas/Antiflatulent	Digestive Enzyme	Drug for Gallstones
SIMETHICONE activated charcoal	α-GALACTOSIDASE LACTASE PANCRELIPASE	ursodiol
Learning Objectives		
Explain mechanism of action, effects, side effects, toxicities, pharmacokinetics and therapeutic uses of each agent.		
<p><b>Clinical Pharmacology</b>  Activated charcoal should be used with caution because of the risk of impaction in the GI tract.</p>		
<b>Relevance</b>		

<p><b>USMLE topic</b> Gastrointestinal System-Principles of therapeutics</p>	<p><b>Principles of therapeutics</b> Mechanisms of action and use of drugs for treatment of disorders of the gastrointestinal system Pancreatic replacement therapy and treatment of pancreatitis Other therapeutic modalities</p>
<p><b>AAMC Medical School Objectives</b> <b>Project Report X Patient Safety-Table 1</b></p>	<p><b>Topic C</b> Drug treatment of common conditions and diseases, using frequently prescribed classes of drugs for the treatment and prevention of disease</p>
<p><b>Notes</b></p>	

<b>Use of Drugs in Pharmacotherapy of Common Gastrointestinal Problems</b>	
<p>The course of instruction should include the pharmacotherapy of common gastrointestinal disorders including: “heart burn,” gastroesophageal reflux disorders, gastritis, bloating, distension, constipation (functional, acute, chronic, idiopathic, iatrogenic), diarrhea (functional, pathogenic, iatrogenic), peptic ulcer disease (duodenal, esophageal, gastric and inflammatory bowel diseases (Crohn’s disease and ulcerative colitis), cramping and irritable bowel syndrome.</p>	
<p><b>Clinical Pharmacology</b>            Already addressed above in the specific section. When using opioid analgesics to manage pain associated with IBD, sufficient drug needs to be administered so as to counter that smooth muscle spasm induced by low doses of this class of drug. Dose should be mg/kg tailored to each patient.</p>	
<b>Relevance</b>	
<p><b>USMLE topic</b>  <b>Gastrointestinal System, Normal Processes</b> – organ structure and function including digestion; cell/tissue structure &amp; function including endocrine and neural regulatory functions; gastrointestinal secretory products  <b>Abnormal Processes</b> – idiopathic disorders</p>	<p><b>Principles of therapeutics</b>            Mechanisms of action and use of drugs for treatment of disorders of the gastrointestinal system</p>
<p><b>AAMC Medical School Objectives</b>  <b>Project Report X Patient Safety-Table 1</b></p>	<p><b>Topic C</b>            Drug treatment of common conditions and diseases, using frequently prescribed classes of drugs for the treatment and prevention of disease</p>
<p><b>Notes</b></p>	