

## Autacoids/Nonsteroidal Antiinflammatory/Asthmatic Drugs

### Subcommittee:

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<b>Histamine and Antagonists</b>		
<b>Recommended Curriculum Equivalent: 1.5 hr</b>		
<b>Drug Classes and Drugs to consider</b>		
H <sub>1</sub> Receptor Antagonists		
First Generation	Second Generation	
DIMENHYDRINATE DIPHENHYDRAMINE PROMETHAZINE chlorpheniramine hydroxyzine	FEXOFENADINE LORATADINE cetirizine	
Endogenous Substances	H <sub>2</sub> Receptor Antagonists	Histamine Release Modifiers
HISTAMINE	CIMETIDINE FAMOTIDINE RANITIDINE nizatidine	CROMOLYN omalizumab
<b>Learning Objectives</b>		
<b>Physiology and pathophysiology</b> Describe the synthesis, storage and release of histamine. Describe the metabolism and elimination of histamine. Identify the major classes of histamine receptors - H <sub>1</sub> , H <sub>2</sub> (with mention of H <sub>3</sub> and H <sub>4</sub> ) and discuss their tissue distribution and function.		
<b>Mechanism of action</b> Explain the molecular mechanism of action of each drug in each drug class.		
<b>Actions on organ systems</b> Describe the pharmacological effects of the drugs in each class on various organ systems. Differentiate the histamine receptor subtypes responsible for mediating the effects of histamine in each organ system.		
<b>Pharmacokinetics</b> Describe the pharmacokinetics of the second generation antihistamines.		
<b>Adverse effects, drug interactions and contraindications</b> Describe the principal adverse effects of the drugs in each class. Describe the clinically important drug interactions of the drugs in each class. Describe the principal contraindications of the drugs in each class.		
<b>Therapeutic uses</b> Differentiate the use of the antihistamines in allergies, sedation, and motion sickness.		

**Clinical Pharmacology**

Of the first generation antihistamines, chlorpheniramine is the least sedating and is often preferred because of this characteristic

Dimenhydrinate and diphenhydramine have a relatively narrow therapeutic index and are often used recreationally as over-the-counter hallucinogens.

Cimetidine is no longer recommended as an H<sub>2</sub> receptor antagonist due to its property of being a suicide substrate for CYP 3A4 and the potential for toxicity of concurrently ingested drugs that are metabolized by this CYP isoform.

**Relevance****USMLE topic**

Respiratory System-Abnormal Processes-  
Immunological Disorders and  
Inflammatory Disorders

**Principles of therapeutics**

Decongestants, cough suppressants,  
expectorants, mucolytics & other  
therapeutic modalities

**AAMC Medical School Objectives**

**Project Report X Patient Safety-Table 1**

**Topic C**

Drug treatment of common conditions  
and disease

**Notes**

Objectives for H<sub>2</sub>-receptor antagonists are covered in Gastrointestinal Drugs.

Objectives for Histamine Release Modifiers are covered in Asthma Drugs.

<b>5-Hydroxytryptamine (5-HT, Serotonin): Agonists &amp; Antagonists</b>		
<b>Recommended Curriculum Equivalent: 0.5 hr</b>		
<b>Drug Classes and Drugs to consider</b>		
Drugs used for Migraine		
Abortive Agents		Prophylactic Agents
Triptans	Ergot Alkaloids	
SUMATRIPTAN zolmitriptan	ergotamine	AMITRIPTYLINE PROPRANOLOL TOPIRAMATE
Serotonin Agonists and Antagonists		Antiemetic
SEROTONIN cyproheptadine		ONDANSETRON
<b>Learning Objectives</b>		
<p><b>Physiology and pathophysiology</b>  Describe the synthesis, storage and release of serotonin.  Describe the metabolism and elimination of serotonin.  Identify the major types of serotonin receptors relevant to therapeutic drugs acting in the brain, the vasculature and the g.i. tract.  Describe the roles of serotonin in migraine, carcinoid syndrome, and in the CNS (emesis; mood disorders and other psychiatric conditions, covered with CNS drugs).</p>		
<p><b>Mechanism of action</b>  Explain the molecular mechanism of action of each drug in each drug class.</p>		
<p><b>Actions on organ systems</b>  Describe the pharmacological effects of the drugs in each class on various organ systems.</p>		
<p><b>Pharmacokinetics</b>  Describe the pharmacokinetics of abortive therapy for migraine.</p>		
<p><b>Adverse effects, drug interactions and contraindications</b>  Describe the principal adverse effects of the drugs in each class  Describe the clinically important drug interactions of the drugs in each class.  Describe the principal contraindications of the drugs in each class.</p>		
<p><b>Therapeutic uses</b>  Differentiate the use of these drugs in migraine (prophylaxis vs. abortive therapy) and as antiemetic agents.  Note the use of the 5HT<sub>2A</sub> and H<sub>1</sub> antagonist cyproheptadine in carcinoid syndrome.</p>		
<p><b>Clinical Pharmacology</b>  A combination of aspirin, acetaminophen and caffeine in clinically effective doses should be tried prior to use of triptan drugs for the treatment of migraine headaches. This drug combination is available as an over-the-counter product and has been demonstrated to be effective in the majority of migraine headache attacks.</p>		
<b>Relevance</b>		

<b>USMLE topic</b> Central and Peripheral Nervous System- Normal processes: cell/tissue structure and function-synthesis, storage, release, reuptake, and degradation of neurotransmitters and neuromodulators	<b>Principles of therapeutics</b>  Mechanisms of action and use of drugs for treatment of disorders of the nervous system-antimigraine agents
<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 disease

<b>Nitric Oxide and Drugs for Erectile Dysfunction</b>			
<b>Recommended Curriculum Equivalent: 0.5 hr</b>			
<b>Drug Classes and Drugs to consider</b>			
Agonist	PDE inhibitor	NO Donors	PGE <sub>1</sub> Analog
NITRIC OXIDE	SILDENAFIL	SODIUM NITROPRUSSIDE nitroglycerin	alprostadil
<b>Learning Objectives</b>			
<b>Physiological roles of NO and cGMP</b> Describe the mechanisms and cellular site of endogenous synthesis of NO and its interactions with guanylate cyclase to regulate cellular levels of cGMP. Explain the roles of NO and cGMP in local control of blood flow, erectile dysfunction and relaxation of the pulmonary vasculature .			
<b>Mechanism of action</b> Explain the molecular mechanism of action of NO, guanyl cyclase and each drug in each class.			
<b>Actions on organ systems</b> Describe the pharmacological effects of the drugs in each class on various organ systems.			
<b>Pharmacokinetics</b> Describe the synthesis (constitutive vs. inducible) and release of nitric oxide.			
<b>Adverse effects, drug interactions and contraindications</b> Describe the principal adverse effects of the drugs in each class. Describe the clinically important drug interactions of the drugs in each class. Describe the principal contraindications of the drugs in each class.			
<b>Therapeutic uses</b> Differentiate the use of these drugs in erectile dysfunction. Describe the use of NO gas to induce pulmonary vasodilation in persistent pulmonary hypertension in newborn and in adult respiratory distress syndrome.			
<b>Clinical Pharmacology</b>			
<b>Relevance</b>			
<b>USMLE topic</b> Cardiovascular System Reproductive System Pharmacodynamic and Pharmacokinetic Processes-general principles		<b>Principles of therapeutics</b> Mechanisms of action, use, and adverse effects of drugs for treatment of disorders of the cardiovascular system antihypertensive drugs, coronary and peripheral vasodilators Restoration of potency Signal transduction	
<b>AAMC Medical School Objectives</b> Project Report X Patient Safety-Table 1		<b>Topic C</b> Drug treatment of common conditions and disease	

**Notes**

Objectives for NO donors are covered under Cardiac Drugs.

Objectives for drugs related to prostaglandins in ED are covered under Eicosanoids.

<b>Eicosanoids: Agonists &amp; Antagonists</b>			
<b>Recommended Curriculum Equivalent: 0.5 hr</b>			
<b>Drug Classes and Drugs to consider</b>			
Prostanoids		Leukotrienes	
Endogenous	Analog	Endogenous	Leukotriene Modifiers
PGE <sub>2</sub> PGF <sub>2A</sub> PROSTACYCLIN THROMBOXANE A <sub>2</sub>	ALPROSTADIL MISOPROSTOL Latanoprost	LTB <sub>4</sub> LTC <sub>4</sub> ,D <sub>4</sub> ,E <sub>4</sub>	ZAFIRLUKAST Montelukast Zileuton
<b>Learning Objectives</b>			
<p><b>Physiology and Pathophysiology</b> Describe the synthesis of prostaglandins, thromboxanes, leukotrienes from arachidonic acid. Explain physiologic and pathophysiologic roles of eicosanoids in regulation of local blood flow, airway resistance, inflammation and nociception.</p>			
<p><b>Mechanism of action</b> Explain the molecular mechanism of action of each drug in each drug class.</p>			
<p><b>Actions on organ systems</b> Describe the pharmacological effects of the drugs in each class on various organ systems. Differentiate drugs inhibiting leukotriene synthesis (zileuton) from leukotriene action at CysLT1 receptors (zafirlukast).</p>			
<p><b>Pharmacokinetics</b> Describe the metabolism and elimination of eicosanoids</p>			
<p><b>Adverse effects, drug interactions and contraindications</b> Describe the principal adverse effects of the drugs in each class. Describe the clinically important drug interactions of the drugs in each class. Describe the principal contraindications of the drugs in each class. Describe the shunting of arachidonic acid metabolism to the production of leukotrienes by inhibition of COX enzymes, leading to bronchoconstriction.</p>			
<p><b>Therapeutic uses</b> Describe the clinical utility of prostaglandin analogs.</p>			
<p><b>Clinical Pharmacology</b> The leukotriene antagonists remain as secondary choices for the management of patients with asthma. Prostaglandin analogs are contraindicated in women of child-bearing age who are or are likely to become pregnant. Misoprostol is most often used together with an NSAID (most often diclofenac) to reduce the risk of a GI bleed in patients requiring chronic drug therapy with an anti-inflammatory analgesic.</p>			
<b>Relevance</b>			
<p><b>USMLE topic</b> Musculoskeletal System</p>		<p><b>Principles of therapeutics</b> Mechanisms of action and use of drugs for treatment of disorders of the musculoskeletal system</p>	

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

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

**Notes**

Objectives for the Leukotriene Modifiers are covered under Asthma Drugs.



Bioactive Peptides		
<b>Recommended Curriculum Equivalent: 0.25 hr</b>		
Drug Classes and Drugs to consider		
Kinins	Neuropeptides	
	Endogenous	Antagonist
BRADYKININ	CGRP substance P VIP	aprepitant
<p><b>Physiology and Pathophysiology:</b>  Describe the synthesis and metabolism of kinins, and the pathological factors that can trigger kinin formation.  Describe the roles of substance P, neurokinins and CGRP in pain perception and local inflammation, and the probable role of substance P in emesis.  Describe briefly the receptors activated by bradykinin, and substance P and other neurokinins.</p>		
<p><b>Therapeutic uses:</b>  Describe the use of the neurokinin antagonist, aprepitant, as an anti-emetic.</p>		
<p><b>Drug interactions</b>  Describe the effects of ACE inhibitors on the metabolism of bradykinin and the production of cough related to ACE inhibitor therapy.</p>		
<p><b>Clinical Pharmacology</b>  Where cough is induced by treatment with an ACE inhibitor, this side effect may be eliminated by substitution with an angiotensin receptor antagonist. There is no good clinical evidence for the concurrent use of an ACE inhibitor and an angiotensin receptor antagonist.</p>		
<b>Relevance</b>		
<p><b>USMLE topic</b>  Pharmacodynamic and Pharmacokinetic  Processes-general principles  Gastrointestinal System</p>	<p><b>Principles of therapeutics</b>  Signal transduction  Mechanisms of action and use of drugs  for treatment of disorders of the  gastrointestinal system</p>	
<p><b>AAMC Medical School Objectives</b>  Project Report X Patient Safety-Table 1</p>	<p><b>Topic C</b>  Drug treatment of common conditions  and disease</p>	

Drugs used for treating Asthma and COPD			
Recommended Curriculum Equivalent: 1 hr			
Drug Classes and Drugs to consider			
Antiinflammatory Drugs		Leukotriene Modifiers	
Steroids	Modulators of mast cell degranulation	Leukotriene receptor antagonists	5-LO inhibitor
BECLOMETHASONE FLUTICASONE	CROMOLYN omalizumab	ZAFIRLUKAST montelukast	zileuton
Bronchodilators			
$\beta_2$ Agonists	Methylxanthines	Muscarinic receptor antagonists	
ALBUTEROL PIRBUTEROL SALMETEROL terbutaline	THEOPHYLLINE aminophylline	IPRATROPIUM tiotropium - now preferred due to once daily dosage	
Learning Objectives			
<p><b>Physiology and Pathophysiology</b> Describe the disease processes of asthma and COPD including airway inflammation, bronchial smooth muscle constriction, and mast cell degranulation. Describe the role of various mediators (histamine, acetylcholine, proteases, leukotrienes C4, D4; prostaglandins; cytokines) in asthma and COPD.</p>			
<p><b>Mechanisms of action</b> Explain the molecular mechanism of action of each drug in each drug class.</p>			
<p><b>Actions on organ systems</b> Differentiate the effects on the lung of the quick relief drugs and the drugs used for long-term control.</p>			
<p><b>Pharmacokinetics</b> Describe the routes of administration of each drug. List the main drugs and clinical situations that can alter the pharmacokinetics of theophylline.</p>			
<p><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.</p>			
<p><b>Therapeutic uses</b> Differentiate the use of these drugs in asthma (short term relief and long term control) and their use in COPD.</p>			
<p><b>Clinical Pharmacology</b> Long-acting beta-2 receptor agonists are not to be used alone, due to down-regulation of the beta-2 receptor. These drugs are to be used only concurrently with inhaled steroids. Many clinicians prefer beclomethasone as an inhaled steroid therapy because of its high first pass metabolism for that portion of the dose that is swallowed during the inhalational ingestion. There is no good evidence for superiority of any of the short-acting beta-2 receptor agonist congeners. Theophylline remains an effective bronchodilator, but the recommended dose has been lowered in recent times. Tiotropium has supplanted ipratropium as an antimuscarinic bronchodilator strategy due to its more convenient once daily dosage recommendation. Anticholinergic and antileukotriene drug therapy remain secondary drug management strategies.</p>			

<b>Relevance</b>	
<b>USMLE topic</b> Respiratory System	<b>Principles of therapeutics</b> Mechanisms of action and use of drugs for treatment of disorders of the respiratory system-bronchodilator drugs
<b>AAMC Medical School Objectives Project Report X Patient Safety-Table 1</b>	<b>Topic C</b> Drug treatment of common conditions and disease
<b>Notes</b> Objectives for steroids are covered under Adrenal Cortex	

## Hypersensitivity and Immunopharmacology

Recommended Curriculum Equivalent: 1 hr

### Drug Classes and Drugs to consider

#### Immunosuppressants

Cytotoxic drugs	Lymphotoxic drugs	Drugs acting on Immunocompetent cells	Drugs acting on cytokines or on cytokine receptors	Other
AZATHIOPRINE	PREDNISONE Antithymocyte immunoglobulin	CYCLOSPORINE TACROLIMUS MYCOPHENOLATE MOFETIL Muromonab Sirolimus	DACLIZUMAB INFLIXIMAB Lenalidomide Etanercept Thalidomide	Rh <sub>0</sub> (D) immune globulin

#### Immunostimulants and colony stimulating factors

EPOETIN ALFA  
FILGRASTIM (G-CSF)  
INTERFERONS  
SARGRAMOSTIM (GM-CSF)  
Idesleukin  
BCG vaccine

### Learning Objectives

#### Physiology and Pathophysiology

Describe the role of immunoglobulins (IgE, IgG, IgM) and cytokines in the immune response.  
Differentiate different types of allergic reactions (Type I-IV) and factors (e.g. cytokines, MHC) involved  
Describe the release of allergic mediators and processes leading to hypersensitivity.

#### Mechanisms of action

Explain the molecular mechanism of action of each drug in each drug class.

#### Actions on organ systems

Relate the main effects of each drug to its molecular mechanism of action.

#### Pharmacokinetics

Describe the route of administration and the relevant pharmacokinetic features of each drug in each drug class.

#### Adverse effects, drug interactions and contraindications

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.

#### Therapeutic uses

Outline the main therapeutic uses of the drugs of each class.

#### Clinical Pharmacology

These drugs are almost always used in combinations. Emphasize the increased risk of activation of latent infection and increased susceptibility to tuberculosis. Some of these drugs suspected to increase the risk of cancer.

### Relevance

<b>USMLE topic</b> Immune System	<b>Principles of therapeutics</b> Mechanisms of action and use of drugs that specifically affect immune function-immunomodulating drugs
<b>AAMC Medical School Objectives Project Report X Patient Safety-Table 1</b>	<b>Topic C</b> Drug treatment of common conditions and disease
<b>Notes</b> Objectives for Corticosteroids are covered under Adrenal Cortex.	

<b>Analgesic, Antipyretic, Antiinflammatory</b>		
<b>Recommended Curriculum Equivalent: 1 hr</b>		
<b>Drug Classes and Drugs to consider</b>		
Nonsteroidal Antiinflammatory Drugs (NSAIDS)		
Salicylic acid derivatives	Nonselective COX inhibitors	Selective COX-2 inhibitors
ACETYLSALICYLIC ACID mesalamine sodium salicylate	IBUPROFEN INDOMETHACIN - NO LONGER A FIRST CHOICE NSAID DUE TO ITS GI TOXICITY NAPROXEN diclofenac ketorolac piroxicam sulindac	CELECOXIB
Analgesic, Antipyretic Drugs		Antidote for acetaminophen
ACETAMINOPHEN		acetylcysteine
<p><b>Physiology and Pathophysiology of pain, inflammation &amp; hyperthermia</b> Outline the physiological basis of temperature control and peripheral sensory pain fibers. Describe the role of eicosanoids and bradykinin in causing local pain, edema and fever. Outline the pathophysiology of acute and chronic inflammation.</p>		
<p><b>Mechanisms of action</b> Explain the molecular mechanism of action of each drug in each drug class. Differentiate the mechanisms of action of acetylsalicylic acid, acetaminophen, and NSAIDS.</p>		
<p><b>Actions on organ systems</b> Differentiate the effects on pain, fever, and inflammation of the drugs in each class.</p>		
<p><b>Pharmacokinetics</b> Describe the metabolism of and mechanism of toxicity of acetaminophen. - Why? This is not an NSAID and is not included in the list above. No acceptable evidence for clinically important anti-inflammatory activity - can be used together with an NSAID to treat acute pain before turning to opioids - See above for migraine headache therapy strategies. Describe the factors that affect the renal elimination of acetylsalicylic acid. Emphasize the dose-dependent disposition within the therapeutic dose range and the ability to dose less than QID if administered as a 900 mg dose.</p>		
<p><b>Adverse effects, drug interactions and contraindications</b> Describe the principal adverse effects of the drugs of each class. Describe consequences of protein binding, zero order metabolism, and irreversible inhibition related to acetylsalicylic acid. Describe the clinically important drug interactions of the drugs of each class. Describe the principal contraindications of the drugs of each class.</p>		
<p><b>Therapeutic uses</b> Differentiate the use of these drugs in treatment of pain, fever, and inflammation. Describe the principles of treatment for acetaminophen toxicity. Describe the principles of treatment for salicylate toxicity.</p>		

**Clinical Pharmacology**

Never use two NSAIDs concurrently. Naproxen is indicated in patients resistant to other propionic acid congeners because it is the only congener that is present in the pure active isomer. Resistance may include a component of inability to convert the inactive isomer to its active form, since this metabolic pathway is highly variable among patients.

Sulindac should be included, since it is the only member of this class where inactivation of the active sulfide metabolite occurs in the kidney of the majority of patients, and may allow use of an NSAID in a patient with impaired renal function.

Selective COX-2 inhibitor drugs have a slower onset of analgesia and should not be used for management of acute pain. Remember that COX-2 is the primary isoenzyme in the kidney and the brain. One should emphasize the danger when two highly-bound drugs are used together – e.g. most NSAIDs and sulfonyleureas. Important also is the risk of bleeds with warfarin + an NSAID due to different mechanisms of anticoagulant effect.

**Relevance****USMLE topic**

Musculoskeletal System

**Principles of therapeutics**

Mechanisms of action and use of drugs for treatment of disorders of the musculoskeletal system-non-steroidal anti-inflammatory drugs and analgesics

**AAMC Medical School Objectives**

Project Report X Patient Safety-Table I

**Topic D**

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

**Notes:**

Objectives for glucocorticoids are covered under Adrenal Cortex.

Objectives for opiates are covered under CNS.

<b>Antirheumatic Drugs</b>		
<b>Recommended Curriculum Equivalent: 0.5 hr</b>		
<b>Drug Classes and Drugs to consider</b>		
COX Inhibitors	DMARDS (Disease Modifying Antirheumatic Drugs)	
	Biologics	Traditional
ASPIRIN IBUPROFEN NAPROXEN celecoxib	ANAKINRA ETANERCEPT INFLIXIMAB abatacept adalimumab rituximab	METHOTREXATE hydroxychloroquine leflunomide sulfasalazine
<b>Learning Objectives</b>		
<p><b>Mechanism of action</b> Explain the molecular mechanism of action common to all nonsteroidal antiinflammatory drugs (NSAIDs). Describe the likely mechanisms of antirheumatic action of the DMARDS.</p>		
<p><b>Pharmacokinetics</b> List the routes of administration of drugs in each class. Recognize the time required before the onset of action of the DMARDS.</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> Outline the use of the NSAIDs and DMARDS in arthritic disorders.</p>		
<b>Clinical Pharmacology</b>		
<b>Relevance</b>		
<p><b>USMLE topic</b> Musculoskeletal System</p>	<p><b>Principles of therapeutics</b> Mechanisms of action and use of drugs for treatment of disorders of the musculoskeletal system- antigout therapy and 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 COX inhibitors are covered under Analgesic, Antipyretic, Antiinflammatory Drugs. Rasburicase is used in pediatric patients receiving cancer chemotherapy.</p>		



<b>Gout</b>		
<b>Recommended Curriculum Equivalent: 0.5 hr</b>		
<b>Drug Classes and Drugs to consider</b>		
Drugs for the gouty attack	Decrease urate formation	Increase urate excretion
INDOMETHACIN Colchicine	ALLOPURINOL Febuxostat Rasburicase	PROBENECID Sulfapyryrazone
<b>Learning Objectives</b>		
<b>Physiology and Pathophysiology</b> Describe the causes and pathophysiology of acute gouty arthritis and chronic tophaceous gout.		
<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 pharmacological effects of each drug in each class. Differentiate the effects of the drugs in the treatment of gout.		
<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. List the drugs that interfere with the renal excretion of uric acid. Describe the mechanism of gouty flare-up associated with the treatment of chronic tophaceous gout.		
<b>Therapeutic uses</b> Differentiate the use of these drugs in the treatment of acute gout attacks and as prophylactic therapies.		
<b>Clinical Pharmacology</b> Probably Indomethacin is no longer the preferred NSAID due to its GI toxicity. Almost all NSAIDs will manage the acute symptoms of gout. Importance of colchicine is noted as an alternate drug therapy for chronic gout in patients allergic to allopurinol. In the elderly, probenecid is probably no longer indicated due to decreased renal function.		
<b>Relevance</b>		
<b>USMLE topic</b> Musculoskeletal System	<b>Principles of therapeutics</b> Mechanisms of action and use of drugs for treatment of disorders of the musculoskeletal system- antigout therapy and immunosuppressive drugs	
<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	

**Notes**

Drugs for specific clinical entities					
Drugs for headaches					
Drug Classes and Drugs to consider					
Acute or Abortive Treatment			Prophylaxis		
Analgesics	Triptans	Ergot alkaloids	β-Blockers	Antiepileptics	Others
acetaminophen aspirin ibuprofen (NSAIDS) (N-acetylcysteine)	sumatriptan	ergotamine	propranolol	valproate	amitriptyline verapamil
<b>Therapeutic uses</b> Outline the use of these drugs in the acute and prophylactic treatment of headaches including migraine, tension and cluster headaches. Outline the management of treatment of overdose to acetaminophen.					
Drugs Used for Treating Alopecia					
Drug Classes and Drugs to consider					
Alopecia Areata			Androgenetic Alopecia		
cyclosporine glucocorticoids			finasteride minoxidil		
<b>Therapeutic uses</b> Outline the use of these drugs in the treatment of alopecia areata and androgenic alopecia.					
Drugs Used for Treating HPV & Molluscum, Actinic Keratoses, BCC and SCC, and Psoriasis					
Drug Classes and Drugs to consider					
HPV & Molluscum, Actinic Keratoses, BCC and SCC	Psoriasis				
	Topical		Systemic		
5-fluorouracil imiquimod podofilox	calcipotriene glucocorticoids tazarotene		acitretin adalimumab alefacept cyclosporine etanercept infliximab methotrexate		
<b>Therapeutic uses</b> Outline the use of these drugs in the treatment of HPV & molluscum, actinic keratoses, BCC and SCC, and psoriasis.					
Drugs Used for Treating Inflammatory Bowel Disease					
Drug Classes and Drugs to consider					
Mesalamine-based therapy	Steroids	Immuno-suppressives	Biologicals	Drugs altering balance of enteric bacteria	

balsalazide mesalamine olsalazine sulfasalazine	budesonide hydrocortisone prednisone	azathioprine mercaptopurine methotrexate	infliximab natalizumab	ciprofloxacin metronidazole <i>lactobacillus</i> spp. <i>saccharomyces</i> <i>boulardii</i>
<b>Therapeutic uses</b> Outline the use of these drugs in treating ulcerative colitis and Crohn's disease.				
<b>Drugs used for Nausea and Vomiting</b>				
<b>Drug Classes and Drugs to consider</b>				
5-HT Antagonists	Substance P/NK <sub>1</sub> Antagonist	Corticosteroids	Others	
granisetron ondansetron	aprepitant	dexamethasone	metoclopramide nabilone olanzapine prochlorperazine scopolamine	
<b>Therapeutic uses</b> Outline the use of these drugs in treating nausea and vomiting associated with chemotherapy, radiation, and postoperative.				
<b>Drugs used for treating Peptic Ulcer Disease and GERD</b>				
<b>Drug Classes and Drugs to consider</b>				
Inhibitors of Acid Secretion				
H <sub>2</sub> RAs	PPIs	Mucosal Protectants	Drugs for eradicating <i>H.</i> <i>pylori</i>	Antacids
cimetidine famotidine nizatidine ranitidine	omeprazole	bismuth salts misoprostol sucralfate	amoxicillin bismuth clarithromycin metronidazole tetracycline inhibitors of acid secretion	CaCO <sub>3</sub> Al(OH) <sub>3</sub> Mg(OH) <sub>2</sub> NaHCO <sub>3</sub>
<b>Therapeutic uses</b> Differentiate between the use of these drugs in peptic ulcer disease and GERD.				
<b>Drug Treatment for Erectile Dysfunction</b>				
<b>Drug Classes and Drugs to consider</b>				
PDE5 Inhibitors	Prostaglandin Analog	Testosterone Replacement	Others	
sildenafil tadalafil vardenafil	alprostadil	methyltestosterone testosterone topical testosterone enanthate	papaverine phentolamine	
<b>Therapeutic uses</b> Outline the use of these drugs in treating erectile dysfunction.				
<b>Drugs for Allergic Reactions</b>				

<b>Drug Classes and Drugs to consider</b>				
Glucocorticoids	Decongestants	Anticholinergics	Anaphylaxis	Autacoid Antagonists
fluticasone prednisone	phenylephrine pseudoephedrine	ipratropium	epinephrine	antihistamines modulators of histamine release H1 receptor antagonists
<b>Therapeutic uses</b> Outline the use of the drugs in each class in treating allergic disorders.				
<b>Pharmacology of Tocolytics, Antenatal Drugs and Abortives</b>				
<b>Drug Classes and Drugs to consider</b>				
Tocolytics			Abortives	Antenatal/Neonatal Therapy
Ca <sup>++</sup> Blockers	β-Agonists	COX inhibitors		
magnesium sulfate nifedipine	terbutaline	indomethacin	methotrexate mifepristone misoprostol	betamethasone indomethacin
<b>Therapeutic uses</b> Outline the clinical uses of the drugs used in preventing preterm labor and in abortion. Outline the uses of betamethasone and indomethacin in antenatal and neonatal therapy, respectively.				
<b>Clinical Pharmacology</b> Evidence for efficacy of tocolytics in humans is less than Impressive. ACE inhibitors also contraindicated In pregnancy.				