

Harvard-MIT Division of Health Sciences and Technology
HST.121: Gastroenterology, Fall 2005
Instructors: Dr. Jonathan Glickman

Vascular and Inflammatory Diseases of the Intestines

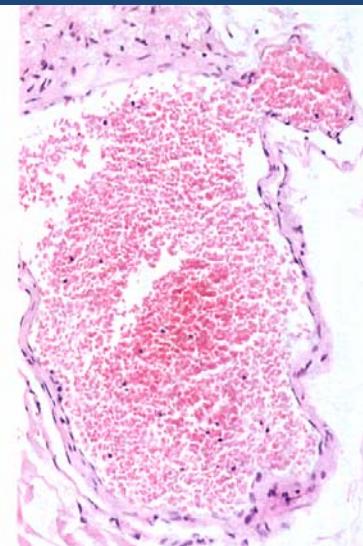
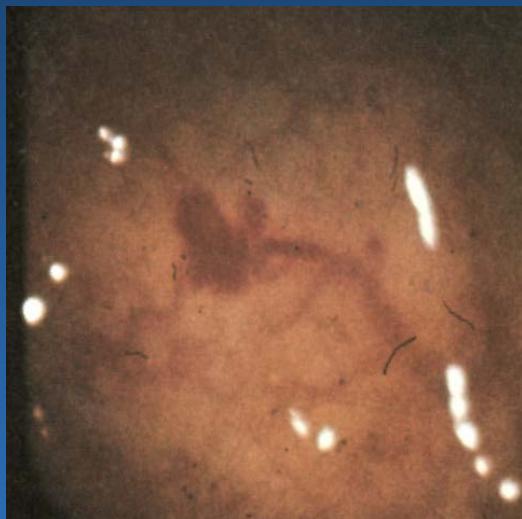
Overview

- **Vascular disorders**
 - **Vascular “malformations”**
 - **Vasculitis**
 - **Ischemic disease**
- **Inflammatory disorders of specific etiology**
 - **Infetious enterocolitis**
 - **“Immune-mediated” enteropathy**
 - **Diverticular disease**
- **Idiopathic inflammatory bowel disease**
 - **Crohn’s disease**
 - **Ulcerative colitis**

Sporadic Vascular Ectasia (Telangiectasia)

- Clusters of tortuous thin-walled *small* vessels lacking muscle or adventitia located in the mucosa and the submucosa
- The most common type occurs in cecum or ascending colon of individuals over the age of 50 and is commonly known as “**angiodysplasia**”
- Angiodysplasias account for 40% of all colonic vascular lesions and are the most common cause of lower GI bleeding in individuals over the age of 60

Angiodysplasia



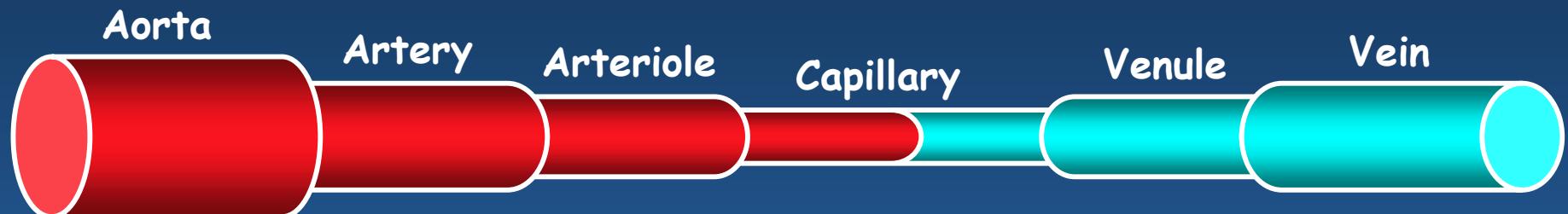
Hereditary Vascular Ectasia

- Hereditary Hemorrhagic Telangiectasia (HHT) or Osler-Webber-Rendu disease
- Systematic disease primarily involving skin and mucous membranes, and often the GI tract
- Autosomal dominant disease with positive family history in 80% of cases
- After epistaxis which occurs in 80% of individuals, GI bleed is the most frequent presentation and occurs in 10-40% of cases

Arteriovenous Malformations (AVM's)

- Irregular meshwork of structurally abnormal medium to large ectatic vessels
- Unlike small vessel ectasias, AVM's can be distributed in all layers of the bowel wall
- AVM's may present anywhere at any age, although some are thought to be congenital

Vasculitis



Goodpasture

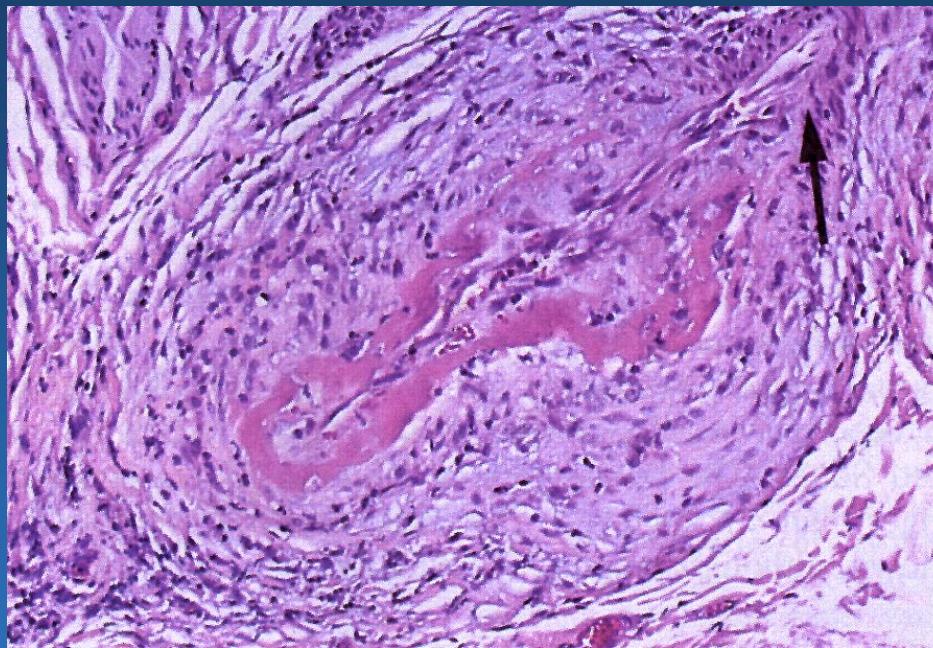
Giant-Cell Arteritis,
Takayasu Arteritis

Microscopic Polyangiitis, Wegener's,
Churg-Strauss Syndrome, SLE

Polyarteritis Nodosa,
Kawasaki Disease

Henoch-Schonlein Purpura

Vasculitis (PAN)



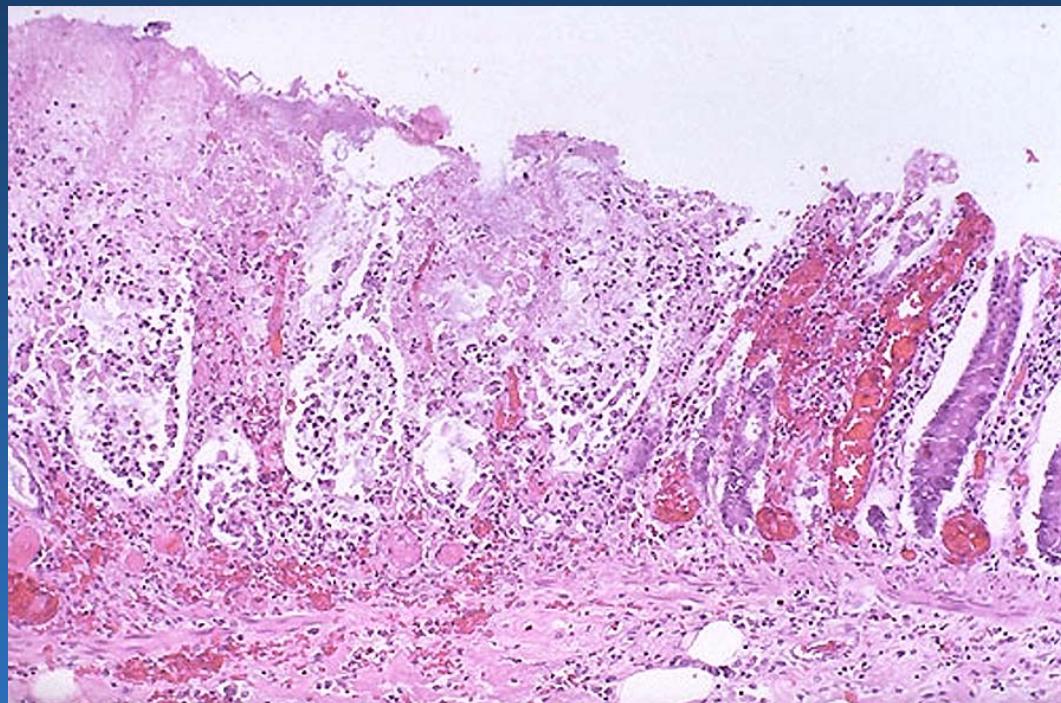
Vascular Insufficiency

- **Vaso-occlusive Diseases**
 - Mesenteric arterial occlusion (embolism/thrombosis)
 - Mesenteric vein thrombosis
 - Bowel strangulation (volvulus, hernia)
- **Non-Occlusive Vascular Insufficiency**
 - Systemic hemodynamic disturbances
 - Local hemodynamic disturbances

Ischemic colitis



Ischemic Colitis



Infectious Enteritis

- The most common GI problem worldwide
- Most symptomatic infections produce diarrhea and some produce malabsorption
- Diagnosis is most often by stool culture or O&P
- Organisms **rarely** produce a pathognomonic pattern of injury

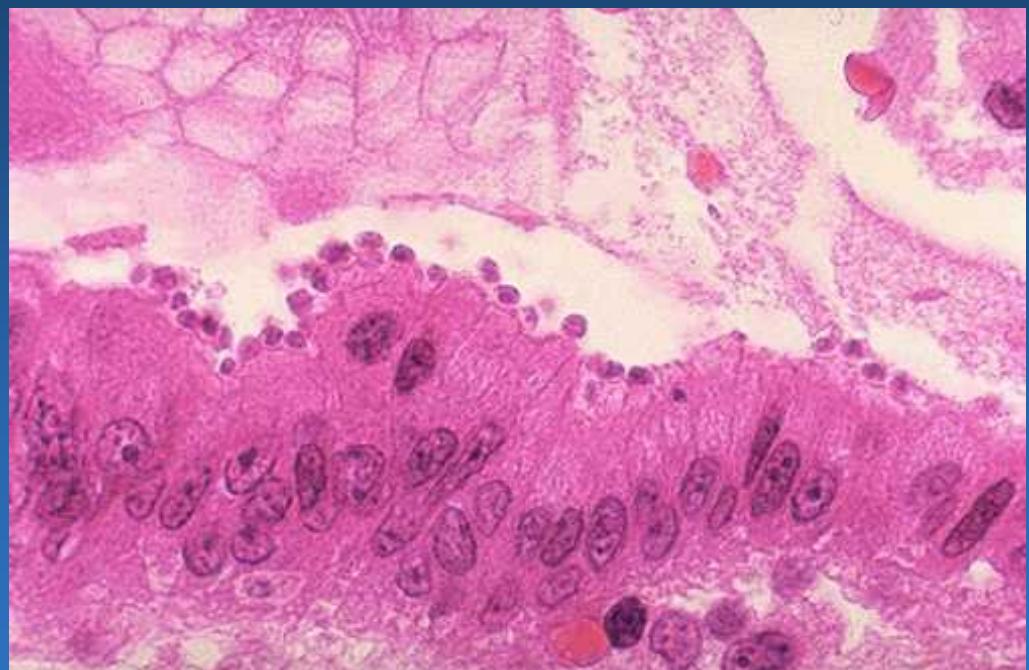
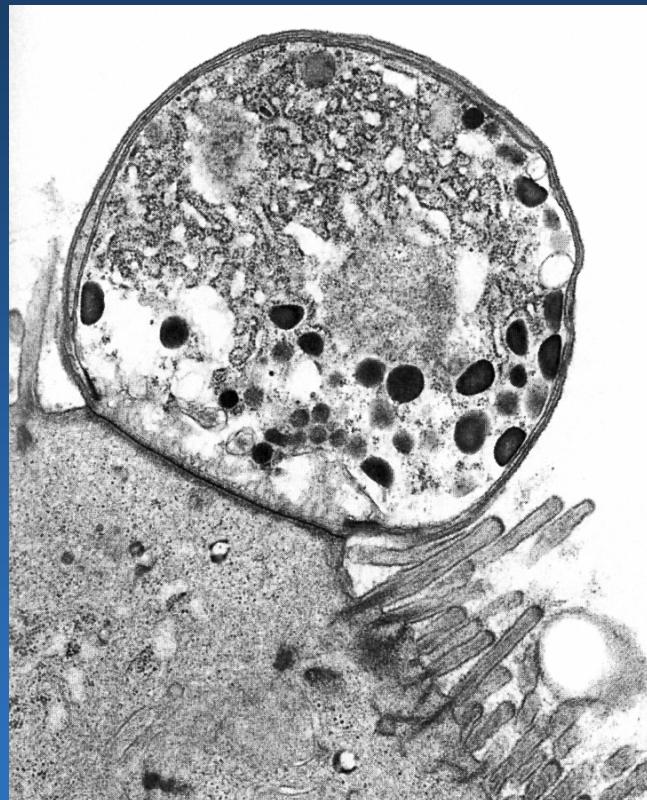
Mechanism of Injury: Toxin Production

- **V. cholera**
- **E. coli**
- “Food poisoning”
 - **Staphylococcus**
 - **Clostridium**

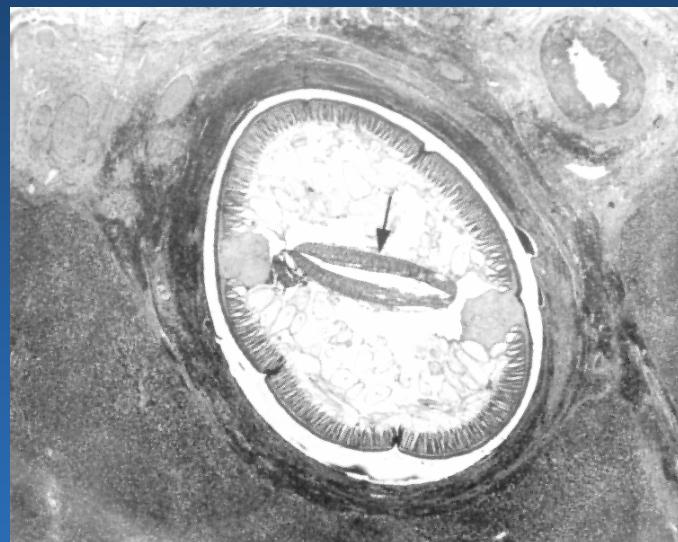
Mechanism of Injury: Invasion

- **Bacteria**
 - **Salmonella, shigella, campylobacter, E. coli, yersinia, mycobacteria**
- **Protozoa**
 - **Cryptosporidia, isospora, microsporidia**
- **Viruses**
 - **Rotaviruses, adenovirus, CMV, HSV**
- **Fungi**
 - **Histoplasma, candida**

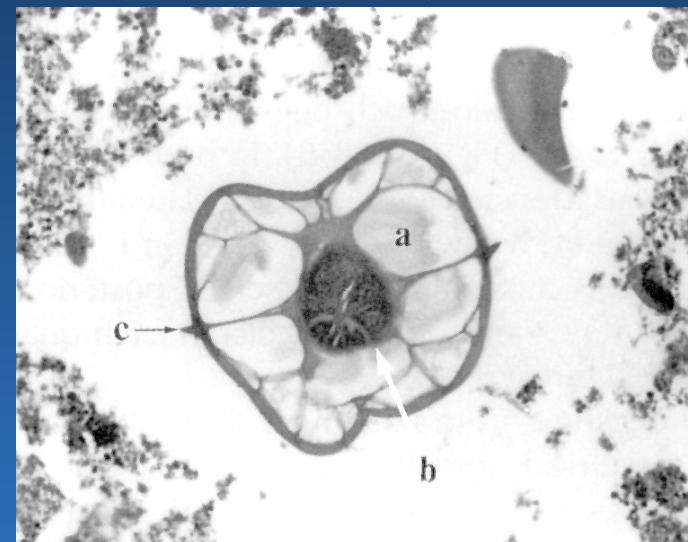
Cryptosporidiosis



The Lumen Dwellers



Ascaris lumbricoides

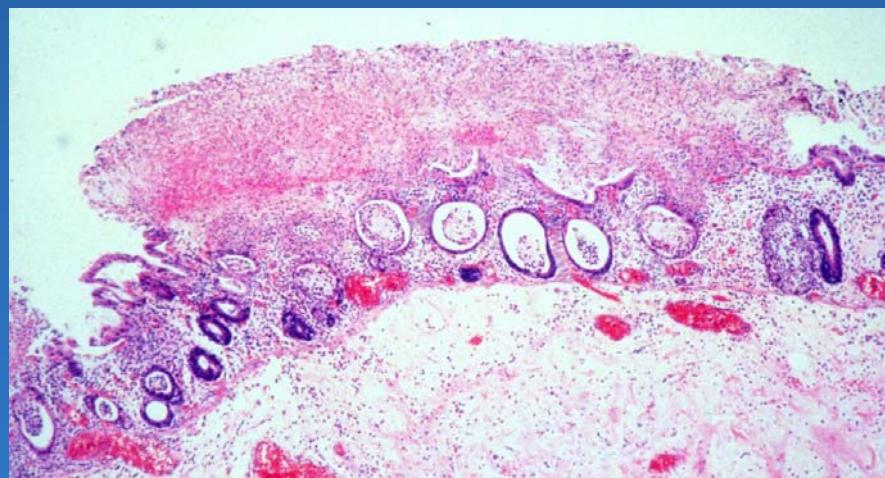


Enterobius vermicularis

Antibiotic-Associated Colitis

- **Antibiotic-associated pseudomembranous colitis (PMC) is an acute colitis characterized by the formation of an inflammatory exudate**
- **PMC is a toxin-mediated colitis usually caused by C. difficile or less commonly by E. Coli**

PMC



“Immune” Enteropathy: Celiac sprue

- **Gluten-sensitive enteropathy, Celiac disease, Non-tropical sprue**
- **Chronic inflammatory disease of the proximal small intestine with generalized malabsorption**
- **Most common in the Irish, British, and other northern European populations**
- **Immune mediated injury to enterocytes accompanied by serum antibodies to gliadin, a component of gluten**

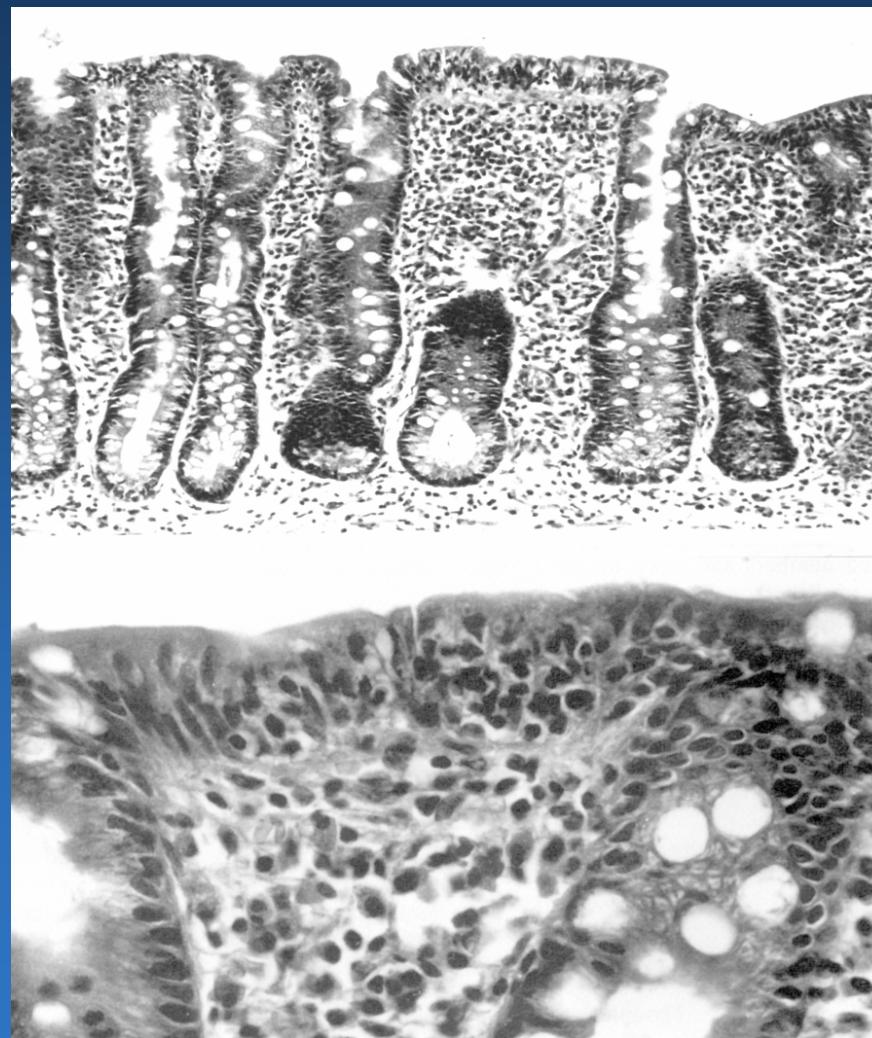
Celiac disease- Immunologic mechanism of damage

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Celiac Sprue

- Symptoms:
 - Steatorrhea, abdominal distention, flatulence, fatigue, and weight loss
- Complications:
 - Iron and vitamin deficiency
 - Risk of lymphoma (T-cell type)
- Extraintestinal manifestation:
 - Dermatitis herpetiformis (a pruritic papulovesicular rash with IgA deposits at the dermoepidermal junction)

CD



Malabsorption- other causes

- Immune conditions
- Hypersensitivity/allergy/eosinophilic gastroenteritis
- Infection- Whipple's dis., tropical sprue, bacterial overgrowth
- Nutritional deficiencies-
- Inherited- Microvillous inclusion dis.,lymphangiectasia
- Infiltrative disorders- amyloidosis, lymphoma
- Systemic disorders- lipid storage
- Other- short bowel

Whipple's disease

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Rosai, Juan, and Lauren Ackerman. *Rosai and Ackerman's Surgical Pathology*. 9th ed. New York, NY: Mosby, 2004. ISBN: 0323013422.

Collagenous, lymphocytic colitis

- Collectively, “microscopic colitis”
- Middle aged to elderly adults
- Chronic watery diarrhea
- Endoscopically normal mucosa

Diverticulosis Coli

- Acquired colonic diverticula are present in nearly half of the population over the age of 50
- Diverticula are associated with low-fiber, low-residue diets
- Etiology is most likely high intraluminal pressure required for propulsion of hard, small stools
- Complications include hemorrhage, acute diverticulitis, perforation, fistula formation

Idiopathic Inflammatory Bowel Disease (IBD)

- Chronic, relapsing, **idiopathic** inflammatory disease of the GI tract
- Crohn's Disease
 - Transmural granulomatous disease affecting any portion of the GI tract
- Ulcerative Colitis
 - Superficial, non-granulomatous inflammatory disease restricted to the colon

Ulcerative Colitis

- Bloody mucoid diarrhea, rarely toxic megacolon
- Can begin at any age, peaks at 20-25 years
- Annual incidence of ~10 per 100,000 in US
- Negligible risk of cancer in the first 10 years, but 1% per year risk of cancer thereafter
- Good response to total colectomy if medical therapy fails

Ulcerative colitis- pseudopolyps

Crohn's Disease

- Variable and elusive clinical presentation with diarrhea, pain, weight loss, anorexia, fever
- Can begin at any age, peaks at 15-25 years
- Annual incidence of ~3 per 100,000 in US
- Many GI complications and extracolonic manifestations
- Risk of cancer less than in UC
- Poor response to surgery

Crohn's disease- gross appearance

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Aphthous ulcer

“Cobblestoning”

Wall thickening

Crohn's disease- stricture

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Crohn's disease- microscopic

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Transmural inflammation

Granuloma formation

Macroscopic Features of CD vs UC

Feature	Crohn's	UC
Region	SI, colon	Colon
Skip areas	Yes	No
Strictures	Yes	No
Wall	Thick	Thin
Creeping fat	Yes	No
Dilatation	No	Yes

Microscopic Features of CD vs UC

Feature	Crohn's	UC
Granulomas	Yes	No
Ulcers	Fissuring	Superficial
Inflammation	Transmural	Mucosal
Sinuses	Yes	No
Fistulas	Yes	No
Fibrosis	Marked	Minimal