Small and Large Intestinal pathology, part 1

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Diseases of the intestines

- Intestinal obstruction
 - Vascular disorders
 - Malabsorptive diseases and infections
 - Inflammatory bowel disease.
 - Polyps and neoplastic diseases

Intestinal obstruction

- Mechanical obstruction:
- Intussusception
 - Hernias.
- ► Adhesions.
 - Volvulus



- ► Tumors.
- Diverticulitis —
- Infarction

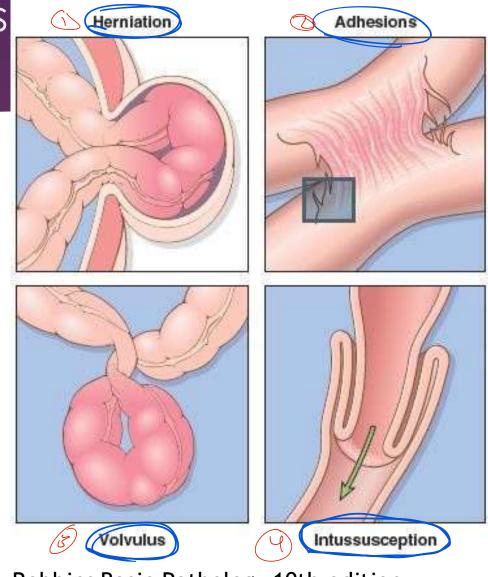
- Non-mechanical obstruction
- Hurschsprung disease
 - Neurological disorders.
 - Drugs....etc



Clinical picture of intestinal obstruction.

- Abdominal pain
- Distention gas raid (uman/ bockerie
- Vomiting
- Constipation.
- Acute or chronic.

80% of mechanical obstructions



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Intussusception

- Segment of the <u>intestine constricted</u> by a wave of <u>peristalsis</u>, <u>telescopes</u> into the immediately distal segment.
- Once trapped, invaginated segment is propelled by peristalsis, and pulls mesentery with it.
- Most common cause of intestinal obstruction in children younger than 2 years of age. pain contribution in children younger than
- Untreated progresses to infarction.



proximal in distal rad

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Causes of intussusception

- < 2years : Idiopathic in most cases.</p>
- Peyer patches hyperplasia (rotavirus vaccine, viral infections)
- Meckles diverticulum (ileum)

10-15 y.

Old children & adults: Intraluminal mass or tumors

Clinical features:

- Abdominal swelling
- Vomiting
- Passing stools mixed with blood and mucus (currant jelly stool)
- ► Pain.

 irrhelle due to

 pain

 iran

 ira

Management



- Contrast enemas in uncomplicated idiopathic cases.
- Surgery if complicated or if masses are the leading point.

Non machanitel.

Hirschsprung Disease

rectom

- Congenital defect in colonic innervations
- Congenital aganglionic megacolon



More common in males



- More severe in females
- Risk increase in siblings.
- Typical presentation:

- Neonatal failure to pass meconium
- Obstructive constipation.

Pathogenesis

- During embryogenesis

 Disrupted migration of neural crest cells from cecum to rectum.
- Lack of Meissner submucosal plexus and the Auerbach myenteric plexus.

 Sun thickness body from rectum not secure,
- ▶ Failure of coordinated peristaltic contractions.
- Mutations in RET: in familial cases and 15% of sporadic
- Other genes and environmental factors play role.

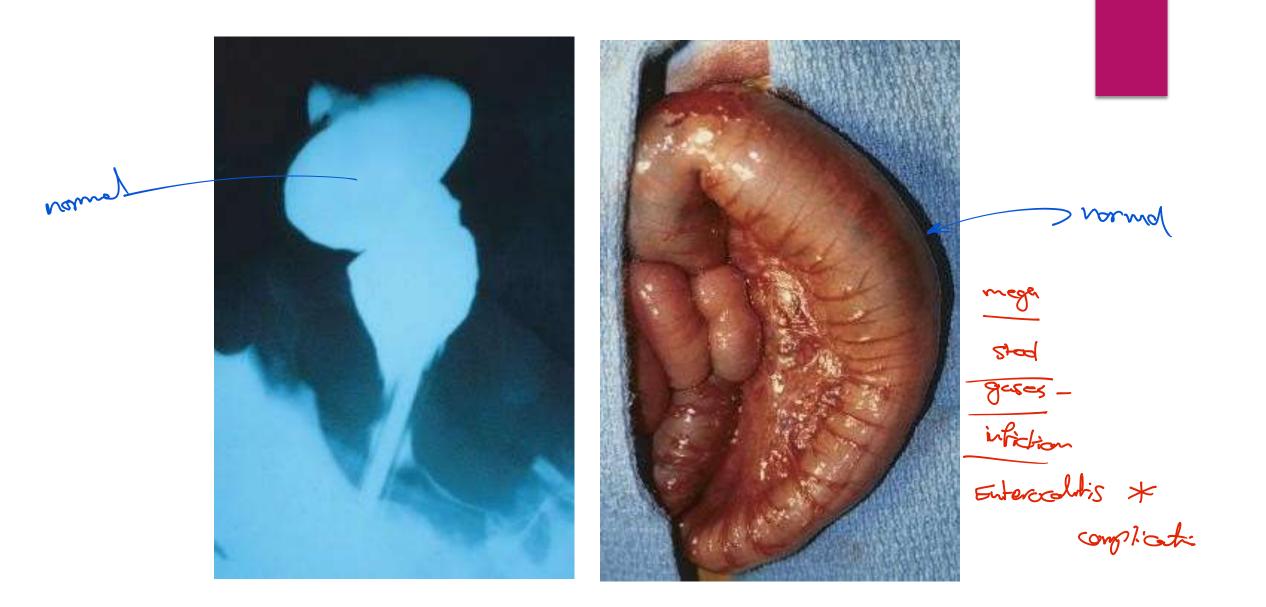
Morphology

- Rectum always involved.
- Extent is variable.
- Most cases in rectosigmoid.

contraction .

- Macroscopic distal.
- Aganglionic region normal or contracted
- Proximal normal segment progressively dilated.
- Diagnostic workup: barium enema and BIOPSY.

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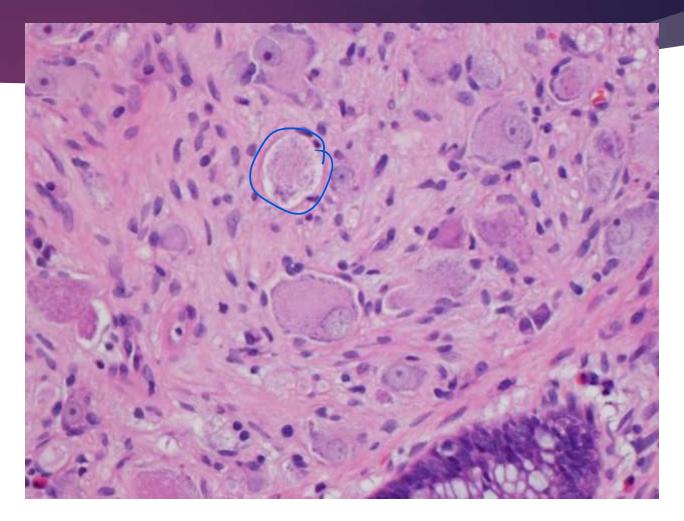
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ganglion cells

LW= G,97 Submices a + Myutric

السواة مرافع

normal



Complications

inflormation of both 5.+ colon.

- Enterocolitis
- Fluid and electrolyte disturbances Johnshien.
- <u>Perforation</u>
- Peritonitis
- Treatment:
- Surgical resection of aganglionic segment and anastomosis of normal segments.

VASCULAR DISORDERS OF BOWEL

Ischemic Bowel Disease why / less vosolety (hopelic Plexer)
Hemorrhoids

with other disore.

Hemorrhoids

Dilated anal and perianal collateral vessels that connect the portal and caval venous systems.

portosystanic shoul

Predisposing factors:

- Constipation and <u>straining</u>
- Venous stasis of pregnancy,
- Portal hypertension.
- External and internal hemorrhoids

above donetal line

Thin -walled, dilated, submucosal yessels beneath anal or rectal mucosa.

Symptoms: Bleeding, pain, thrombosis and inflammation

not mixed with

conflicated

Surgery in comblicati.

DIARRHEAL DISEASE

- Diarrhea: increase in stool mass, frequency or fluidity.
- Dysentery: painful, bloody, small volume diarrhea.
- Malabsorptive Diarrhea

Pancreatic insuffciency.

Celiac disease

Crohn disease

Cystic Fibrosis

Lactase (Disaccharidase) Deficiency

<u>Abetalipoproteinemia</u>

- Infectious Enterocolitis
- Inflammatory bowel diseases.....

Malabsorptive Diarrhea

- Chronic.
- Defective absorption of fats, fat- and water-soluble vitamins, proteins, carbohydrates, electrolytes, minerals and water
- Hallmark is: steatorrhea.

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abrormed quality of fat in shoot.

+ Vitamines

yellow - chem
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Malabsorptive diarrhea Defect in one of the following:

- Intraluminal digestion.
- Terminal digestion.
- ► Transepithelial transport. When G
- Lymphatic transport.

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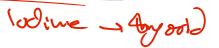
Manifestations:

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Weight loss, anorexia,

Flatus, abdominal distention,

- Borborygmi, Muscle wasting
- Anemia and mucositis (iron, pyridoxine (VB6), folate, or vitamin B12 deficiency)
- Bleeding (vitamin K deficiency)
- Osteopenia and tetany (calcium, magnesium, or vitamin D deficiency)
- ► Neuropathy (vitamin A or B12 deficiency)
- ► Skin and endocrine disorders.



Cystic Fibrosis

- Mutations in cystic fibrosis transmembrane conductance regulator (CFTR)
- Defects in ion transport across intestinal and pancreatic epithelium.
- Thick viscous secretions. Was water in ilven dicorhe
- Mucus plugs in pancreatic ducts >>> pancreatic insufficiency (80% of patients).
- Defect in intraluminal digestion.

Celiac Disease

تحسی العر

- Gluten sensitive enteropathy
- Immune mediated enteropathy
- Wheat, rye or barley.
- Genetically predisposition, HLA-DQ2 or HLA-DQ8.
- Treatment: gluten free diet.

Ab > = interior 2000 = chiltren.

(Dudenon)



Association with: type 1 diabetes, thyroiditis, and Sjogren syndrome

Autoimme diseases

Pathogenesis

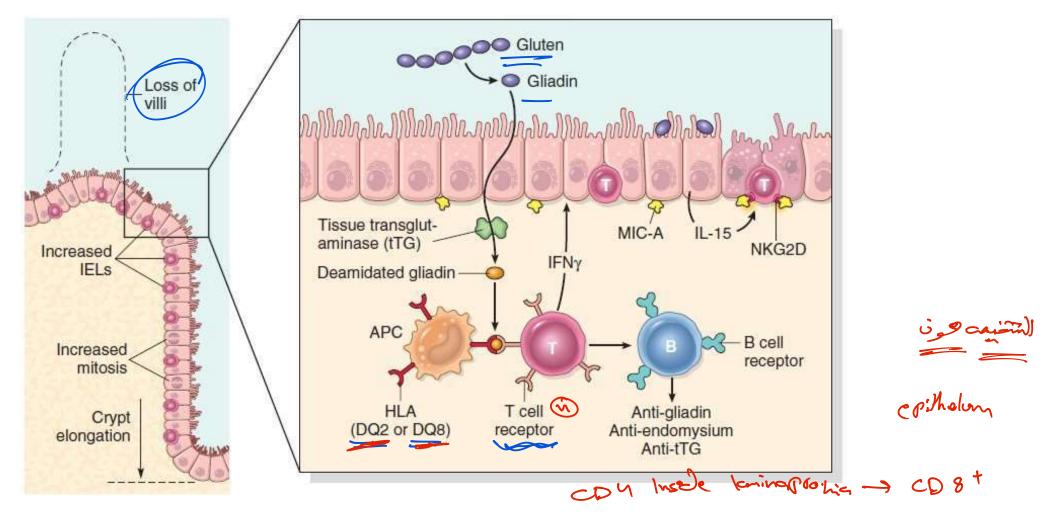
Gluten >>> gliadin >>> react with HLA-DQ2 or HLA-DQ8 on antigen-presenting cells >>> CD4+ T cells activation >>> cytokines >>> tissue damage.

Serology:

- Anti- tissue transglutaminase antibodies 🖈 0
- Anti-gliadin antibodies
- Anti -endomysial antibodies (3)

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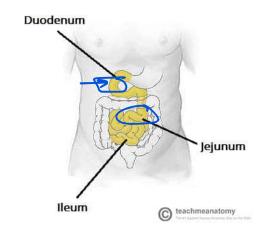


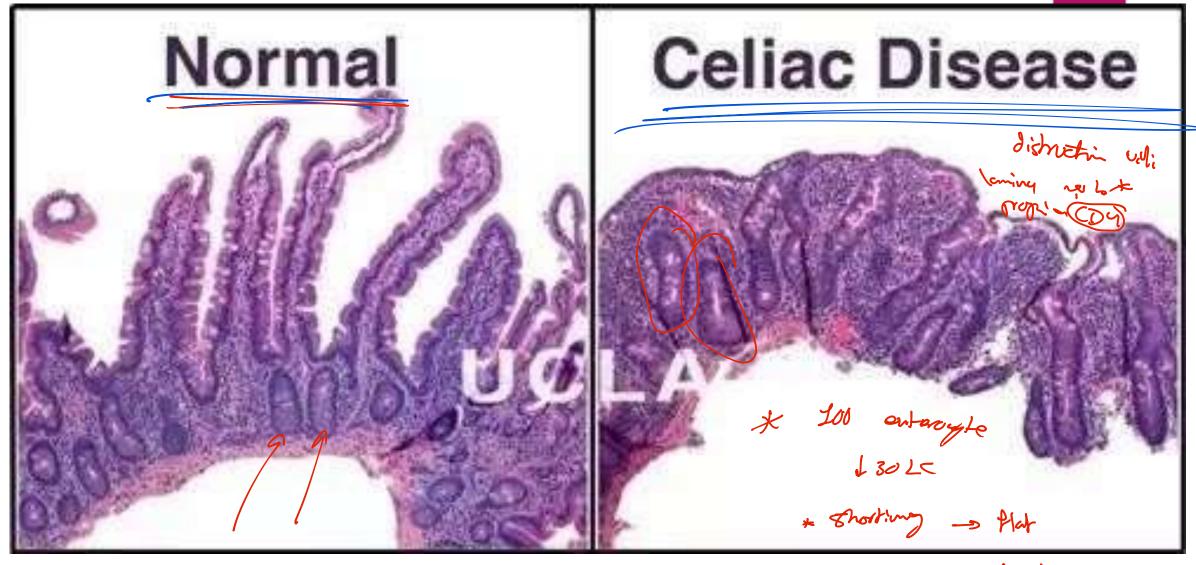
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MORPHOLOGY

ocid stormoch in the

- Second portion of the duodenum or proximal jejunum.
- Triad: intraepithelial lymphocytosis (CD8+ T cells), crypt hyperplasia, and villous atrophy.
- Lamina propria lymphocytes, plasma cells, eosinophils......
 - IEL & villous atrophy are not pathognomonic, seen in viral enteritis.
- Diagnosis: Clinical, histologic and serologic correlation.





atrophy. Erght hyporphesis

Clinical Features

1. Joi d'126

- ► Children 6-24 months: classical or non classical symptoms
- Classical: Irritability, abdominal distention, anorexia, diarrhea, failure to
- thrive, weight loss, or muscle wasting
- Non-classical: abdominal pain, nausea, vomiting, bloating, or constipation.

shoot human

Blistering skin lesion, dermatitis herpetiformis, in 10% of Pnts.

Dermatitis herpetiformis.



- Adults (30-60 years)
- Anemia: iron deficiency
- B12 and folate deficiency: less common.
- Diarrhea , bloating, and fatigue.
- Missed diagnosis: Silent celiac or latent celiac.

No symbon + serdogy but biopsy name.

Increased risk of enteropathy associated T cell lymphoma & Small intestinal adenocarcinoma

de 200 60-109

edubouties

Diagnosis:

- 1. Non invasive serologic tests:
- Most sensitive:

Anti tissue transglutaminase antibody, IgA *

Anti deamidated gliadin antibodies, IgA & IgG 💥

Most specific, but less sensitive

Antiendomysial antibody.

2. Invasive tests: small bowel biopsy.

normal ofter

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Lactase (Disaccharidase) Deficiency

- Osmotic diarrhea No Jigistan
- Lactose remains in the gut lumen.
- Lactase found at apical brush border membrane
- Normal biopsy findings.
- Two types:

Congenital: <u>AR</u>, <u>genetic mutation</u>, <u>rare</u>, explosive diarrhea, watery, frothy stools & abdominal distention, after milk ingestion

Acquired: follow viral or bacterial enteritis, after childhood.

well

Stop with and it's devalues.

Abetalipoproteinemia

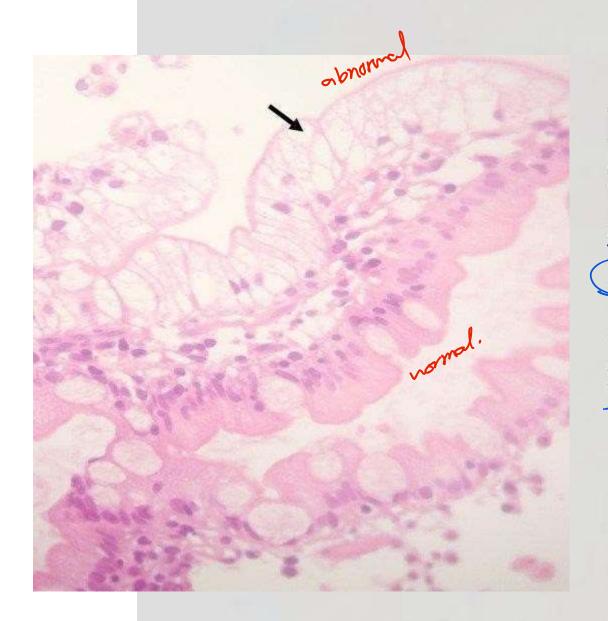
- Autosomal recessive, rare.
- Infants w/ failure to thrive, diarrhea, and steatorrhea
- Lack of absorption of fat and fat soluble vitamins



- nability to synthesize triglyceride-rich lipoproteins.
- Transepithelial transport defect of TG and FAs.
- Monoglycerides and triglycerides accumulate in epithelial cells.







Micrograph showing enterocytes with a clear cytoplasm (due to lipid accumulation) characteristic of abetalipoproteinemia.



Micrograph showing enterocytes with a clear cytoplasm (due to lipid accumulation) characteristic of abetalipoproteinemia.