

GASTROINTESTINAL PATHOLOGY

Non-infectious causes of malabsorption:

1) Celiac sprue :- Aka gluten sensitive enteropathy. (Present in wheat, rye, barley, old rice)

Antibodies produced :- Anti gliadin, Anti endomysial Ab, Anti tissue transglutaminase Ab

2) Ileostigam of choia - Biopsy - flattening or loss of microvilli

Skin involvement :- Dermatitis herpetiformis *

Rx :- Avoid gluten containing foods. They can eat MAIZE

POC for dermatitis herp... => DAPSONE => Infliximab

2) INFLAMMATORY BOWEL DS :-

1) CROHN'S DS :-

Any part of GIT can be affected. Muc sit - Pleum

* Usually - rectum spared.

* Ulcers - longitudinal serpiginous ulcers

* Skip lesions non-caseating microscopy - granulomatous infection

- Transmural inflammation
- Mucosa gives cobblestone appearance
- Ulcer -> heal by fibrosis.

pulling up of muscularis into bowel (creeping fold)
Excessive fibrosis -> stricture (narrowing)

Histology :- granulomatous inflammation

Radiology - String of Kantor *

Lab's - Ab - ASCA +ve
↓
saccharomyces cerevisiae

Hallmark :- fistulas features 

ULCERATIVE COLITIS

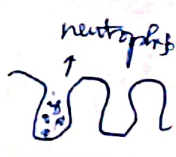
- Colon affected -> continuous retrograde manner (PANCOLITIS)
- Superficial inflammation - only mucosa & submucosa

Histology - Non-granulomatous inflammation

Formation of pseudopolyps.

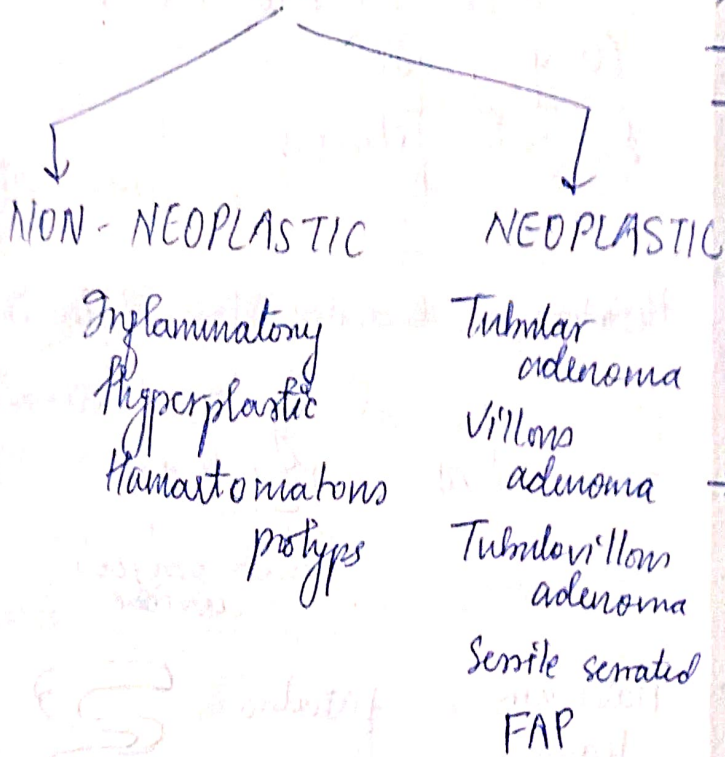
Associated PSC - Primary sclerosing cholangitis

LABS - P-ANCA +ve

Histology - crypt abscess
Radiological - Lead pipe appearance 

Rx:- 5-aminosalicylic acid
Sulphasalazine

POLYPS



* Hyperplastic Polyp :-

- Upper crypts are serrated.
- 60 years; m/c - rectum
- Biopsy - m/s
 - ↳ serrated upper crypts
 - ↳ narrow uniform basal crypts

* ~~Hamartoma~~ Sessile serrated adenoma
+ basal crypts are also serrated.
+ ^{m/s} t shape or boot shaped crypts

* Hamartomatous Polyps :-

A mass of mature but disorganised tissue native to the organ

1. Peutz Jegher Syndrome
 2. Juvenile polyposis
 3. Cowden DS
- } Hamartoma polyp

Juvenile Polyposis - < 5 years -

- Rectal polyp (m/c)
- hamartomatous
- non-cancerous

CF - Hirschsprung (froh)
Iron def. anemia

Mutation:- SMAD4

BMPRIA

① Hereditary hemorrhagic Telangiectasia
Complications - ② Malformation

- Colonic / enterocolonic adenoma
- GIT cancers

* Histology - mucin filled crypts

Peutz Jegher Syndrome

- Mucocutaneous pigmentation
- GIT hamartomatous polyp

⇒ STK11 gene mutation

- m/c site - jejunum
- multiple hamartomatous polyps
- family h/o - +ve
- hists - arborizing network of lamina propria