

POLYPS OF COLON:

- A polyp is a growth arising out of the mucosa.
Polyps may be: (A mass that protrude into the lumen of the gut)

Pedunculated: with a well developed stalk

Sessile: without polyp stalk

Histopathology:

I $\Rightarrow$ Non-Neoplastic - Inflammatory, Hamartomatous, Hyperplastic.

II $\Rightarrow$ Neoplastic / Adenomas - Tubular, Villous, Tubovillous, secreted Adenomas. NO

Inflammatory Polyps:

$\Rightarrow$ Raised inflamed / regenerating epithelium.

$\Rightarrow$ Associated with solitary rectal ulcer syndrome, IBD, ischemic colitis, infections.

$\Rightarrow$ Impaired Relaxation of Anorectal sphincter $\rightarrow$ sharp angle that damage of ulcerate Rectal Mucosa $\rightarrow$ Repeated cycles of Ulceration & healing $\rightarrow$ polyp formation.

$\Rightarrow$ P/w Rectal bleed & Mucous discharge, inflammatory polyp. (mixed)

Hyperplastic Polyp

Most common left type - Sessile Rubrum (left colon) $\rightarrow$ MC

Usually occurs in 6th to 7th decade

piling up of epithelium (cell turnover & delayed shedding)

No malignant potential

Single / Multiple

Sessile < 5 mm

Elongated crypts with serrated fray tooth appearance

Appearance $\rightarrow$ smooth nodular protrusion of the mucosa.

Microscopy $\rightarrow$ expanded population of mature goblet & Absorptive cell.

Juvenile Peutz-Jeghers poly

Hamman-Rich polyp: - (Rac tumor like growth)

- Hamartoma - A benign mass of disorganized tissue native to a particular anatomical site.
- Hamartomatous polyps can occur as part of a syndrome
- neoplastic potential
- They may be associated with intestinal and extraintestinal manifestations.

Juvenile Polyp (most common type)

- Usually seen in children < 5 years
- Can be sporadic or syndrome (juvenile polyposis)

AD inheritance with mutation in TGFβ signaling

multiple polyps are associated with adenocarcinoma of colon & other sites.

clinical location, Rectum, presents with Rectal bleed.
Gross - pedunculated, spherical, smooth-surface Red to color.

Microscopy

- edematous lamina propria with cryptically dilated gland lined by cuboidal or columnar epithelium with mixed inflammatory infiltrates.

Peutz-Jeghers Syndrome

Rare AD disease with multiple GI hamartomas

mucocutaneous hyperpigmentation of Risk of Malignancies

⇒ Genetic Loss of function mutation in STK11

- ↑ Risk of colon of cancer, lung, pancreas, ovary, testis
- HC location of poly, small intestine
- Median Age - 11 years

Gross → large, pedunculated polyp with lobulated surface
→ present dozen seen in ST

Histology

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- Destructive Papillary villous Architecture with tree-like - Arborescence and Arborescence Intermixed with loose propria
- hyperplastic mucosal epithelium divided by broad bands of mucosa smooth muscle.

Neoplastic Polyps: Adenoma

- Precursor to colorectal adenocarcinoma
- 1) → Growth pattern - flat/sessile/pedunculated
- 2) → Architecture - Tubular / Tubulovillous / villous.

Size is the most important risk factor for carcinoma

Polyp > 2cm, Multiple, Villous Architecture, Dysplasia, Development of cancer

% Adenomas

→ Tubular

→ villous

→ Tubulovillous

Sessile Serrated Adenoma

→ Right colon

→ Serrated polyp (hyperplastic polyp)

- Malignant polyp doesn't show dysplasia

Familial Adenomatous Polyposis (FAP)

AD, numerous colorectal Adenomas by teenage

History of polyposis in the

clinical FAP has at least 100 polyps

Prophylactic colectomy is done

Morphology of polyps is the same as that of sporadic Adenomas

50% of cases → develop CA colon

APC - Adenomatous polyposis coli gene

Variants

FAP (Familial Adenomatous Polyposis) Syndrome

- Association of FAP
- epidermal cysts
- Desmoid
- Thyroid tumors
- Dental abnormalities, including unerupted & supernumerary teeth

Turcot syndrome

- colonic Adenomas
- tumor of the central nervous system
- 2/3rd APC gene mutation & develop medulloblastoma
- 1/3rd mutation in DNA Repair gene develop glioblastoma

Lynch syndrome (Hereditary Non Polyposis colorectal cancer)
HNPCC

- Associated defect in DNA mismatch Repair gene (MMR)
- Majority of APC/LCC cases involve either HSB12 or MLH1
- Microsatellite Instability
- AD transmission, ↑ Risk of cancer of colorectum.
- Younger Age
- Few Adenomas & are usually more serrated Adenomas
- Sporadic colon cancer is usually left

Condon syndrome

mutation → PEN gene

Pancreatic silicosis → APC

MORPHOLOGY OF NEOPLASTIC POLYPS

- ⇒ General features: site almost 50% all adenomatous polyps of colon located in Rectum sigmoid Region.
- ⇒ Macroscopy: All colorectal Adenomas are low grade dysplasia + low in

TUBULAR ADENOMAS

- constitute 2/3 of the Adenomas of LT.
- Appear as small, smooth, surfaced, pedunculated polyps. less than 2 cm.

Micro → closely packed small rounded or tubular glands embedded in stroma. They may show variable degree of epithelial dysplasia.

VILLOUS ADENOMAS (Villous Papilloma)

- predominantly found in the Rectum sigmoid Region.
- Gross: Large, broad-based, sessile (cauliflower like surface)
- Micro → thin long finger like projections, superficially resemble the villi of the small intestine.
- They contain foci of carcinoma more commonly than tubular Adenoma.

TUBULOVILLOUS ADENOMAS → mixture of both. 15% will be villous type

humanes histology of colorectal polyps: first factor: genetic factors (mostly in familial polyposis)

second factor: environmental factors (diet, lifestyle, etc.)

third factor: unknown factors

fourth factor: unknown factors (possibly related to inflammation, infection, etc.)

Adenoma is premalignant polypoid