Lecture 3: Neoplasm of the Stomach

General Note:

- Gastric adenocarcinoma >90% of malignant lesions of stomach
- The symptoms range from asymptomatic to bleeding, obstruction, or perforation.
- To identify these lesions, you would need:
 - Endoscopy
 - **Histopathological examination** of the lesion and the surrounding mucosa.



Gastric Polyps:

- Found in 6% of asymptomatic patients who undergo endoscopy (found incidentally).
- Types:
 - 1. **Fundic** (most common)
 - 2. Hyperplastic (~17%)
 - 3. Adenomas (~0.69%)
 - 4. Inflammatory (~0.1%)
- 2% are diagnosed as "malignant polyps".
- Note: Hyperregenerative epithelium can result in development of hyperplastic polyps.
 - We need to examine the <u>whole</u> abdomen

1. Fundic polyps: FGP

- Most common type.
- Associated with:
 - Long-term **PPI** therapy (4x risk)
 - Gastrinoma († gastrin levels)
 - **FAP** (familial adenomatous polyposis) and polyposis syndromes (MUTH-associated polyposis / gastric adenocarcinoma
 - proximal polyposis of stomach
- Genetic factors :
 - sporadic FGP and those linked to <u>PPI</u> are at low risk
 - Who assosciated with <u>polyposis syndrome</u> \rightarrow >70% have <u>APC mutations</u> that are linked to <u>malignancy</u>
- Screening → colonoscopy is recommended in pt who have 1->20FGP 2-<40 onset 3- duodenal adenomas 4-polyps outside the gastric corpus →suspect FAP.

2. Hyperplastic polyps:

- Caused by chronic inflammatory stimuli (e.g., H. pylori) / Adjacent to gastric ulcers or gastroesophageal or gastrointestinal anastomosis.
- Diagnosis:
 - 1. Endoscopy
 - 2. Biopsy is required
 - 3. Larger polyps (>1cm) and pedunculated ones have a higher risk of malignancy
 - 4. Tx \rightarrow >0.5 cm should be resected via endoscopic or histopathological examination
 - 5. H.pylori Should be eradicated

Gastric Adenoma:

- Most frequent **neoplastic lesion** of the stomach.
- Most commonly found in the **distal stomach** (antrum) \rightarrow unlike **FGP which is proximal**.
- Risk of malignancy = 8-59% are assodciated with gastric ca so the whole abdomen should be examined
- Management \rightarrow Endoscopic resection should be done ASAP with regular follow-up.

1	Hyperplastic polyps	Adenomas	Fundic gland polyps	
Macroscopy	most frequently in the antrum, often multiple, usually smooth, dome-shaped, 0.5 - 1.5 cm in diameter; large hyperplastic polyps often become lobulated and pedunculated, and the surface epithelium is typically eroded a reter is may beck	Velvety lobulated appearance, usually <u>solitary</u> : located more often in the antrum	multiple, small (<1 cm), appear smooth, glassy, and sessile constr in proceed domain with tog lik 4 PPC inc., solor pathecuated	



<u>Fig. 1</u> a Fundic gland polyps. **b** Fundic gland polyps and adenoma.

GIST (Gastrointestinal Stromal Tumor)

General Characteristics:



- GIST = mesenchymal tumor (not epithelial)
- Typically present \rightarrow Most common **subepithelial neoplasm** of the stomach.
- GIST is the most prevelent entity
- Other mesenchymal tumors (less frequent + significant) = leiomyoma, leiomyosarcoma, schwannoma

+ epidemiology:

- 2.2% of Malignant cancer tumors with non gender prefrences
- Peak age: 60s–80s
- Distribution:
 - Stomach (60–70%)
 - o midgut (20–30%)
 - Esophagus and lower GIT (10%)

Histological & Genetic Features:

- Histology: Spindle cell, epithelioid, or rarely pleomorphic
- They arise from cells of cajal(pacemaker)
- Genetics:
 - KIT mutations
 - PDGFRA receptor tyrosine kinase
 - o Less commonly: SDH-mutations, BRAF, NF1

📟 Diagnosis:

- Based on: (together)
 - Conventional pathology
 - Immunohistochemistry stain (IHC)
- IHC markers:
 - \mathbf{V} CD117 (c-KIT) → 95%
 - $\boxed{\bigcirc}$ DOG-1 $\xrightarrow{\rightarrow}$ useful in dx
- A Useful in diagnosing when KIT-negative GIST

***** Clinical Considerations:

- GIST spreads locally and does not involve lymph nodes
- Chance of recurrence or resection decision depends on:
 - o Tumor size / site
 - Mitotic number
- If ≤ 1 cm and asymptomatic \rightarrow can leave it and observe

A Complications:

- 1. Obstruction
- 2. Bleeding
- 3. Perforation

NET: Neuroendocrine Tumors

1. Incidence:

• NETs are on the rise, with gastric NETs accounting for 6–23% of all GI NETs.

2. Causes of increased incidence:

- Unclear whether it's due to:
 - True increase in cases
 - \circ Or \uparrow (esophagogastroduodenoscopy)

3. Clinical course:

- Most are **asymptomatic and benign**
- Some can become aggressive and resemble gastric cancer

4. Malignant potential?

- All have malignant potential
- In the stomach \rightarrow 3 distinct type of neuroendocrine neoplasm Differentiated by stage and histology





Features of ga	astric NETs acco	ording to [20]
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Feature	Type 1	Type 2	Type 3 Poorly differentiate
Proportion among gastric neuroendocrine neoplasms	70 - 80%	5 - 6%	14 – 25%
Macroscopic aspect	often small (<1 – 2 cm), 78% polypoid, 65% multiple	often small (<1 – 2 cm), polypoid and multiple	unique, large (>2 cm), polypoid, ulcerated
Pathological differentiation	often NET G1	NET G1 -G2	NEC G3
Proportion of metastases	2 - 5%	10 - 30%	50 - 100%
Associated conditions	chronic atrophic gastritis	gastrinoma/MEN – 1	none

◆ 5.secondary neoplasia of the stomach :

- Prevalance \rightarrow gastric **mets** = rare (1.3-1.7%) of cancer pt
- Common origins → breast cancer / melanoma / lung cancer

Lymphoma:

1. Definition:

• Can arise from the **gastric wall** or **adjacent lymph nodes**

2. Prevalence:

- GI lymphoma = 4–18% in the western world
- ~25% in the Middle East
- Stomach is the most common in the West
- Midgut is most common in the Middle East

3. Age of onset:

- Typically in the **5th decade**
- No gender predominance

4. Risk Factors:

- **H. pylori** (Leading cause) \rightarrow 90% in MALT lymphoma
- Immunosuppression (e.g., HIV) $\rightarrow \uparrow risk (\sim 23\%)$

5. Immunohistochemistry:

- Positive for B-cell markers (CD19 / CD20 / CD22)
- Negative for CD5 / 10 / 38 / IgD

+ Types:

- **High-grade** = majority **>high grade b cells**
- Low-grade = B-cell MALT (the only low-grade one)

MALT Lymphoma:

- Symptoms:
 - Abdominal pain
 - Nausea / vomiting
 - Signs of bleeding
 - Palpable mass if large
- Treatment:
 - $\circ \quad \text{Early stage} \rightarrow \text{Regresses after } \underline{\text{H. pylori therapy}}$
 - $\circ \quad \text{Advanced stage} \rightarrow \underline{\text{Radiotherapy}}$
- High-grade lymphomas (Mantle cell / Diffuse B-cell / Burkitt lymphoma) → All treated with chemotherapy

Genetic risk:

- t(11;18), t(q21;q21)
 - \rightarrow Associated with treatment failure
 - \rightarrow And transformation into diffuse large B-cell lymphoma
- **Surgery** only for complications:
 - Bleeding
 - Perforation

Tonic	High-Vield Surgical Evam Points	
<u>Costria Adapaganainama</u>	• >000/ of molionant agating lagions	
Gastric Adenocarcinoma	• -90% of mangnant gastric resions	
	• Can present with bleeding , obstruction , or perforation – important complications	
	surgeons focus on	
Gastric Polyps	• Fundic Gland Polyps \rightarrow PPI use, FAP \rightarrow screen for FAP if:	
	1) >20 polyps 2) <40 age 3) duodenal adenoma 4) polyps outside fundus	
Gastric Adenoma	• Associated with malignancy $(8-59\%) \rightarrow$ full abdominal exam required	
	• Must be resected ASAP with follow-up	
GIST	Most common subepithelial neoplasm	
	• From Cajal cells (pacemaker cells)	
	• Spreads locally, no lymph node involvement (unique among gastric tumors!)	
	 Important complications: bleeding, obstruction, perforation 	
	Diagnosis: CD117 (KIT), DOG-1	
NET (Neuroendocrine	All have malignant potential	
Tumors)	Can be asymptomatic or aggressive like gastric ca	
Secondary Neoplasms	• Rare (1.3–1.7%)	
	• MC origins: breast, melanoma, lung	
Lymphoma (especially	• MC in stomach (West), midgut in Middle East	
MALT)	• H. pylori \rightarrow 90% of MALT	
	• Treated with antibiotics (early stage)	
	Surgery only for: bleeding or perforation	



MALT lymphoma