

Lecture 2: Gastric Cancer (Gastric CA)

● General Characteristics:

- 9th M/C cancer in males and 8th in females
 - 145 new cases yearly
 - More common in Japan and asia
 - Male > Female = 5:3
 - 7th decade is the peak incidence
 - Less than 5% are under the age of 55
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◆ Etiology:

- There are 2 types of gastric cancers:
 1. Cardiac CA: associated with developed or endemic countries
 2. Non-cardiac CA: associated with developing countries or non-endemic areas like Japan
 - Highly associated with H. pylori
 - H. pylori:
 - Most important factor
 - Found in poor hygiene, crowded, low socio-economic areas
 - Increases the risk (×2) of non-cardiac tumors
 - Risk even higher in CagA-positive patients
 - Chronic atrophic gastritis caused by H. pylori increases the risk (×11) of Cardiac cancer → the only case where H. pylori is associated with cardiac cancer
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◆ Cardiac cancer risk factors:

1. Low fat / protein diet
 2. N-nitroso compounds → inc risk for non cardia cancer in those with h.pylori infx
 3. High salt food (x3)
 4. Processed meat (ham, sausage, bacon)
 5. Fried or grilled food → heterocyclic amines
 6. Smoking (x2)
 7. nitrosamine
 8. Alcohol
 9. Obesity (×2)
 10. Pernicious anemia (2–3%)
 11. post gastric surgery → especially post-antrectomy (after 15–20 years)
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✍ Classification:

1. WHO Morphology: widely used but little in in pt management

- Squamous cell
- Adenosquamous
- Undifferentiated
- Unclassified
- Adenocarcinoma → subclassified into:

- Papillary
 - Signet ring
 - Mucinous
 - Tubular
- all depending on **growth pattern (degree of differentiation)**

2. Bormann's classification:

- Based on **macroscopic appearance**:
 1. Polypoid or fungating lesion
 2. Ulcerated with raised edges
 3. Ulcerated with no edges
 4. Diffuse infiltrating lesion
 5. Doesn't fit any

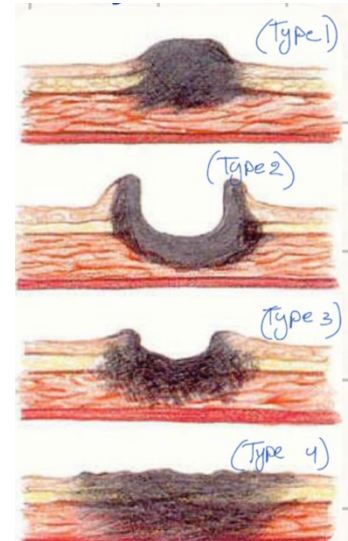
◆ **Not very useful** for diagnosis or prognosis // degree of cellular differentiation

- **Broders classification:**

- Based on **cellular differentiation**:
 - Type 1: well differentiated
 - Type 4: anaplastic

3. Lauren's classification:

Intestinal type	Diffuse type
<ul style="list-style-type: none"> ○ Older men ○ Well-differentiated ○ Spreads through blood ○ Good prognosis ○ Limited (non-cardiac) 	<ul style="list-style-type: none"> ○ Young women ○ Poorly differentiated ○ Spreads? submucosa+ lymphatics ○ Poor prognosis ○ Lacks gland formation ○ Composed of signet-ring cells ○ Associated with blood type A and familial cases



▼ Symptoms:

- **At first:** It's often **asymptomatic**, but you might notice:
 - **Pain** (constant, especially **doesn't relieve by drugs or food**)
 - **Dysphagia**
 - **Weight loss**
 - **Loss of appetite**
 - **Early satiety**
 - **Vomiting**
- On PE: they develop **late**, they are associated with **metastasis signs**:
 1. **Virchow's node** (left supraclavicular node)
 2. **Sister Mary Joseph nodule** (umbilical nodule)
 3. **Palpable epigastric mass**
 4. **Jaundice**
 5. **Ascites**
 6. **Cachexia**

Clinical Evaluation and Staging:

1. flexible upper Endoscopy:

- **First choice** once you suspect **gastric cancer**
- You should take **biopsies** from:
 - **The edge** and not the crater
 - *(Since it may show necrotic debris only)*
- You need **at least 7 biopsies** to assess:
 - **Size**
 - **Site**
 - **Morphology**




2. Blood tests:

- Should appear **normal** unless there's a **sign of mets** :
 - Anemia (check **full blood count**)
 - **LFTs** for liver mets and advanced disease
 - **Coagulation panel** → abnl in advanced disease


◆ 3) Double contrast barium swallow

- We don't use it anymore since it gives **false +ve** or **-ve**
- It's **not helpful** for differentiating benign and malignant tumors
- cost-effective
- accuracy (~90%)

◆ 4) EUS (Endoscopic Ultrasound)

-  Used for **staging**
- **Extent of wall invasion and nodal status**
- **Good for evaluating LN / added advantage of FNA**
- Perfect for **T1 and T3 (superior to CT)**
-  Not useful for **T2**, since it **can't reach the muscularis propria**
-  Not suitable for therapy response since it :
 - **Cannot differentiate fibrosis from cancer**
 - Gives ~80% accuracy

◆ 5) CT scan

- Useful for **T4, nodal enlargement, liver mets**
-  Not suitable for **T1/T2 tumors (early)** <5 mm mets to liver and peritoneum
- If nodal involvement(size) is seen in the **chest**, it's a **poor prognosis**
- **Chest , abdomn , pelvis**
- **PET-CT** → detect distant mets , used to **followup** (اللي عندهم احتمال لتطور المرض)
- Accuracy: **80–85%**

◆ 6) Diagnostic laproscopy

- Used for **locoregional disease**
- Detects **liver and peritoneal mets** <5 mm
- Can **detect mets** in 30% that were thought resectable in CT or EUS
- ● Once you see tumor cells in **peritoneal fluid** → (presence of free intraperi. Gastric cells) T4

◆ L.N. classification: -regional lymph node (N)-

- Nx ; nodes can't be assessed
- **N0**: No lymph node mets
- **N1**: 1–6 positive nodes
- **N2**: 7–15 positive nodes
- **N3**: >15 positive nodes
- **M1**: Distant metastasis

■ Note:

- **T1 and T2** are **early-stage** gastric cancer even if there's nodal involvement
- T3/N2 = **advanced stage cancer**

● Treatment:

- **Stage IV disease (M1)**: **palliation therapy**. ^{to chemo}
- All other stages if medically fit consider for diagnostic laparoscopy for further staging.

After laparoscopy

- Stage M1 palliation only.
- Stage M0, but medically unfit either palliation only or radiotherapy and 5-FU radiosensitization.
- Stage M0 and medically fit, T1 or less. **For surgery**.
- Stage M0 and medically fit, **T2 or higher**. **Neoadjuvant chemotherapy with ECF (MAGIC trial protocol) followed by surgery**.

▲ Surgical Treatment:

- First tx of gastric cancer
- <50% at presentation are resectable, extent of resection depends on R0
- **6 cm from edge** is required to **decrease risk of local recurrence**

Types:

1. **Proximal tumor (cardiac)** → Total gastrectomy
2. **Distal tumor** → Subtotal gastrectomy

? Do you need to remove lymph nodes?

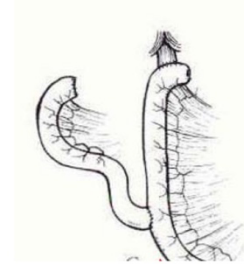
- Controversial
- You can only **remove group 1 LN** (found on **lesser & greater** curvature)
- **✗** Group 2 (found on common hepatic, left gastric, coeliac, splenic) **can't** be removed
- Group 3 (para – aortic)

! D1 vs D2 surgery:

- D2 = higher morbidity & mortality
- BUT no difference in **overall survival**

Type of surgery :

- **Roux-en-Y** reconstruction → connects **esophagus to jejunum** after total gastrectomy

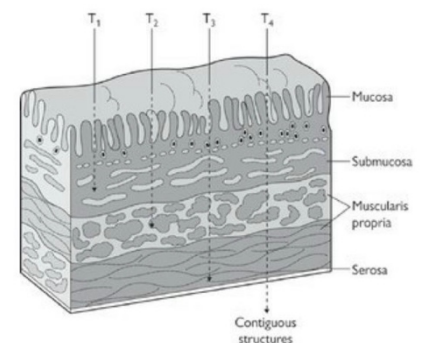


! Complications of Gastric Surgery:

- **Early:**
 - Anastomotic leak
 - bleeding
 - infx
- **general:**
 - DVT
 - PE
 - Cardiorespiratory complications
- **Late complications:**
 - **Early dumping** → 20-30min after ingestion of a meal (Acute destruction of the bowel) // Autonomic response // serotonin, bradykinin-like substances, neurotensin, and enteroglucagon
 - **GI symptoms** → nausea and vomiting, sense of fullness, belching, abdominal cramps, and explosive diarrhoea.
 - **Cardiac symptoms** → palpitations, tachycardia, sweating, fainting, dizziness, flushing, and visual disturbance // Symptoms usually subside with time
 - **Late dumping** → 2-3h after ingestion of a meal // large amount of carbohydrates to the proximal small intestine (quickly absorbed, sudden hyperglycaemia, large amount of insulin, profound hypoglycaemia, catecholamines from the adrenal gland, tachycardia, sweating, confusion, and dizziness)
 - Symptoms similar to hypoglycaemic shock
 - Also reduces with time

Stage 0	Tis, N0, M0
Stage 1A	T1, N0, M0
Stage 1B	T1, N1, M0 T2a/b, N0, M0
Stage II	T1, N2, M0 T2a/b, N1, M0 T3, N0, M0
Stage IIIA	T2a/b, N2, M0 T3, N1, M0 T4, N0, M0
Stage IIIB	T3, N2, M0
Stage IV	T4, N1-3, M0 T1-3, N3, M0 Any T, any N, M1

Staging systems



Adjuvant Therapy:

M1	Palliative therapy
R0 resection and T1, NO	Observe
R0 and <u>T2</u> and higher	ECF chemotherapy
R1 resection	Radiotherapy plus concurrent 5-FU sensitization followed up by ECF chemotherapy if T2 and higher <i>Radio then chemo</i>
R2 resection	5-FU based radiosensitization or ECF chemotherapy or best supportive care if unfit <i>Radio and chemo</i>
Primary palliative chemotherapy	Reassess and if good response consider surgery

Follow-up:

Follow ups?

Surveillance

- First 3 years follow-up should be intensive, since recurrence is most common at this stage
- Follow-up should be 4-6 months for first 3 years, thereafter annually
- History, physical examination, and routine blood profile should be conducted at each followup visit
- CT scan should be performed yearly for first 3 years or sooner if suspicious
- Yearly gastroscopy on patients who have undergone subtotal gastrectomy

In General:

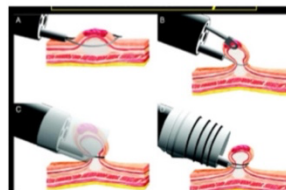
- 1) Total → use CT
- 2) Subtotal → use endoscopy

Endoscopic mucosal resection (EMR)

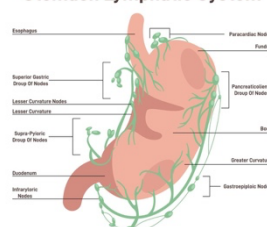
Most experience with EMR is in Japan where there is higher incidence of early gastric cancer and an active screening program.

Indications for EMR

- well or moderately differentiated
- tumour size less than 30mm
- absence of ulceration
- no evidence of invasive findings



Stomach Lymphatic System



Anatomy?

Gross anatomy

- Relations of the stomach:
- anteriorly – left lobe of the liver;
- superiorly – the diaphragm;
- medially – the liver;
- laterally – spleen;
- inferiorly – transverse mesocolon, caudate lobe of liver, crura of the diaphragm, and retroperitoneal nerves and vessels.

