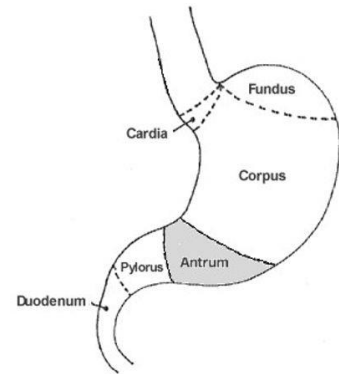


# GASTRIC CANCER

## General Overview

- 5<sup>th</sup> leading cause of cancer worldwide (4<sup>th</sup> leading cause of cancer deaths). Highest rates in Eastern Asia. Global incidence has declined (thanks to refrigerators (less salt-based preservation))
- Risk factors: H. Pylori infection, family history, salt and low vegetables intake, smoking, alcohol
- 2 main histologic variants: “intestinal type” and “diffuse type”
- Diffuse type more frequent in young patients and female, some with CDH1 mutations
- True hereditary (diffuse type) only in 1-3% but familial aggregation in about 10% of cases.
- Association with Lynch, FAP, Peutz-Jeghers, Li Fraumeni and Cowden syndrome.
- Clinical symptoms: weight loss, abdominal pain,...



## Staging (AJCC Version 8) and Prognosis

- CT thorax/abdomen, endoscopy, (endoscopic ultrasound)
- Tumors involving the GEJ with the tumor epicenter no more than 2cm into the proximal stomach are staged as esophageal cancer while GEJ tumors with their epicenter >2cm into the proximal stomach as gastric cancer, as are all cardia cancers not involving the GEJ (even if <2cm of the GEJ)

Primary Tumor (T)	Regional Lymph Nodes (N)	Distant Metastasis (M)
<b>Tx:</b> Primary tumor cannot be assessed	<b>Nx:</b> LN cannot be assessed	<b>M0:</b> no distant M+
<b>T0:</b> No evidence of primary tumor	<b>N0:</b> no regional LN	<b>M1:</b> distant M+
<b>Tis:</b> Ca in situ, high grade dysplasia	<b>N1:</b> M+ in 1 or 2 regional LN	
<b>T1:</b> invasion lamina propria, muscularis mucosae(T1a) or submucosa (T1b)	<b>N2:</b> M+ in 3 to 6 regional LN	
<b>T2:</b> invasion muscularis propria	<b>N3:</b> M+ in 7 or more reg LN	
<b>T3:</b> penetrates the subserosal connective tissue without invasion off the visceral peritoneum or adjacent structures	N3a: 7-15 regional LN	
<b>T4:</b> Tumor invades :	N3b: ≥16 regional LN	
T4a: serosa (visceral peritoneum)		
T4b: adjacent structures/organs		

- Prognosis: 5y survival
  - I: 86%
  - II: 69%
  - III: 21%
  - IV: 4%

- Pathological stages posttreatment (ypTNM)

ypT	ypN	M	Stage
T1-2	N0	0	I
T1	N1	0	I
T3,4	N0	0	II
T2,3	N1	0	II
T1,2	N2	0	II
T1	N3	0	II
T4a	N1	0	III
T3,4	N2	0	III
T2-4	N3	0	III
T4b	N0,1	0	III
Any T	Any N	1	IV

## Treatment

- Among persons with *H. pylori* infection who had a family history of gastric cancer in first-degree relatives, *H. pylori* eradication treatment reduced the risk of gastric cancer (NEJM 2020)
- For patients with T2-4N0 and node-positive disease we recommend peri-operative chemotherapy with FLOT
- For patients with primary surgery adjuvant chemo(RT) is recommended (ex. FOLFOX 6m)
- Follow-up after surgery:
  - Every 3-4 months for the first 2y with imaging (preferably CT), followed by 6 monthly until 5 years.
- Metastatic disease
  - Many trials included both esophageal and gastric cancer regardless of histology and therefore general treatment such as chemotherapy regimens converged.
  - All gastric cancers should be tested for HER2 (IHC + ISH), MSI and PD-L1
  - 1<sup>st</sup> line (1-4):
    - HER2+: pembrolizumab + trastuzumab + 5FU + platinum
    - HER2- / CPS ≥10: chemo + pembrolizumab or nivolumab
    - HER2- / CPS ≥5: chemo + nivolumab
    - HER2- / CPS ≥1: chemo + pembrolizumab
    - Preference for FOLFOX as platinum based chemotherapy
  - 2<sup>nd</sup> line (5-7):
    - MSI-H: pembrolizumab monotherapy
    - HER2+ (confirmed on repeated biopsy): trastuzumab deruxtecan, based on DESTINY Gastric01
    - HER2-: Paclitaxel + ramucirumab or ramucirumab monotherapy
  - Later lines: FOLFIRI, TAS102
- Pembrolizumab reimbursement Belgium:
  - 1<sup>st</sup> line HER2+ AC gastric or GEJ, CPS ≥ 1 in combination with trastuzumab, 5FU and platinum (in theory no reimbursement in combination with capecitabine or oxaliplatin)
  - 1<sup>st</sup> line HER2- AC gastric or GEJ, CPS ≥ 1 in combination with platinum and 5-FU

- 1<sup>st</sup> line HER2 neg esoph / gastric / GEJ, CPS  $\geq$ 10 in combination with platinum and 5FU
- 2<sup>nd</sup> or later lines: MSI-H gastric
- Nivolumab reimbursement Belgium:
  - Adjuvant esophageal / GEJ after neo-adj chemoRT and residual disease (no pCR)
  - 2<sup>nd</sup> line monotherapy in SCC after platinum+5FU
  - 1<sup>st</sup> line SCC in combination with platinum/5FU if TPS $\geq$ 1
  - 1<sup>st</sup> line HER2 negative esoph, gastric or GEJ, CPS $\geq$ 5 in combination with platinum /5FU
- Trastuzumab deruxtecan reimbursement Belgium
  - HER2+ AC gastric or GEJ
  - Previously treated with trastuzumab
  - HER2+ ISH positive

## References

- 1) TOGA trial : Lancet 2010 (Bang YJ et al)
- 2) Janjigian YY et al Lancet Oncol 2020
- 3) Checkmate 649 : Nature 2022 (Shitara K et al)
- 4) Keynote 859 : Lancet oncol 2023 (Rha SY et al)
- 5) DESTINY-Gastric01 : NEJM 2020 (Shitara K et al)
- 6) REGARD trial: Lancet 2014 (Fuchs CS et al)
- 7) RAINBOW trial: Lancet Oncol 2014 (Wilke H et al)

## What's new ?