

# **Esophageal and Gastric Motility Disorders: A case based approach**

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## **Conflicts of Interest:**

- **None**

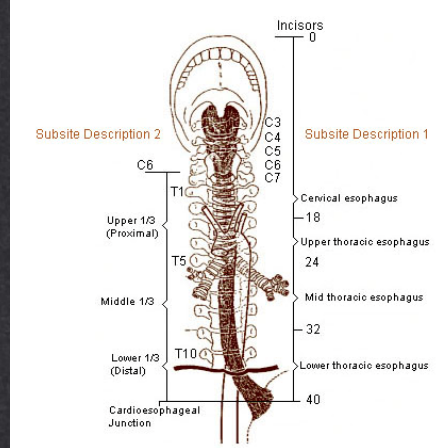
## Overview

- Esophageal anatomy
- Dysphagia-case based approach
- Reflux disease-case based approach
- Gastric physiology
- Gastroparesis-case based approach

## Dysphagia-Case based approach

# Esophagus: Anatomy

- 25 cm muscular tube.
- Extends from upper esophageal sphincter to stomach.
- Proximal 1/3<sup>rd</sup> consist of striated muscles while distal 2/3<sup>rd</sup> is formed by smooth muscles.
- Lined squamous epithelium.



# Terminology

- **Dysphagia:** derived from the Greek word *dys* (difficulty, disordered) and *phagia* (to eat).
- **Odynophagia:** painful swallowing.
- **Globus Sensation:** Sensation of lump in throat between meals.



# History

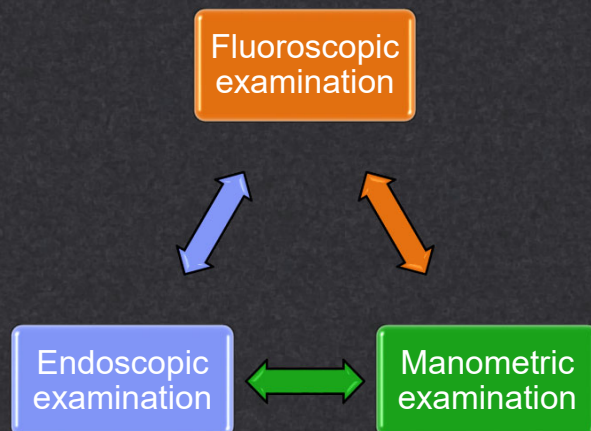
## Oropharyngeal

- Oral:
  - Drooling of saliva
  - Food spillage
  - Sialorrhea
  - Piecemeal swallows
  - Associated dysarthria
- Pharyngeal:
  - Choking/cough during swallow
  - Associated dysphonia

## Esophageal

- Food stuck in suprasternal notch or retrosternal region
- Motility:
  - dysphagia to solids and liquids
  - Associated with heartburn or chest pain.
- Mechanical:
  - progressive dysphagia to solids; may involve liquids at later stages

# Dysphagia Assessment

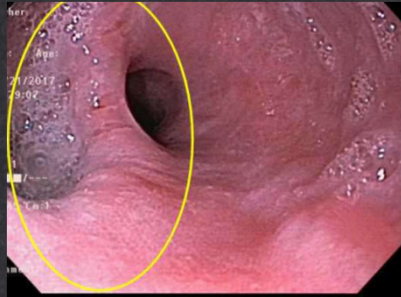




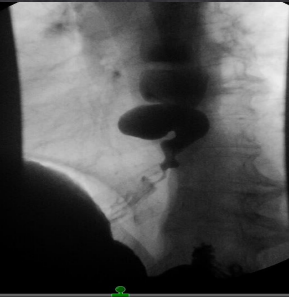
## Case Study 1:

78-year-old female with no significant medical history presenting with:

- Dysphagia to both solids and liquids
- Chest pain
- Denies any heartburn
- 50 lb weight loss



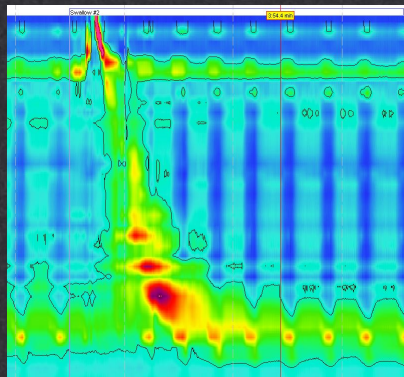
- Epiphrenic diverticulum
- Resistance at GEJ



- Epiphrenic diverticulum
- Beaking at GEJ

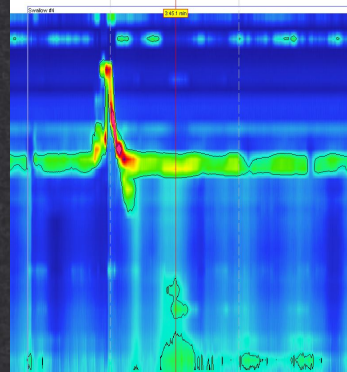
## Case Study 1:

- Mean DCI: 2380
- Mean LES IRP: 32 mm Hg
- Mean DL: 3.8 sec



## Case Study 1:

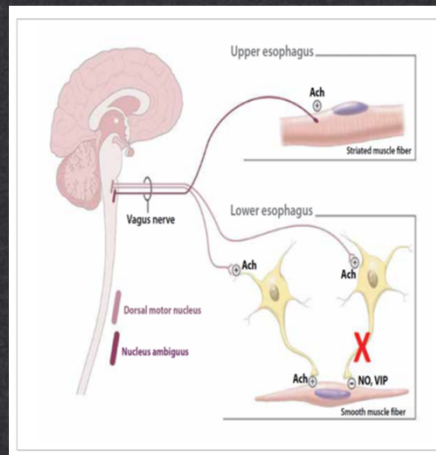
- Post extended myotomy and diverticulectomy
- Fairly doing



## Achalasia

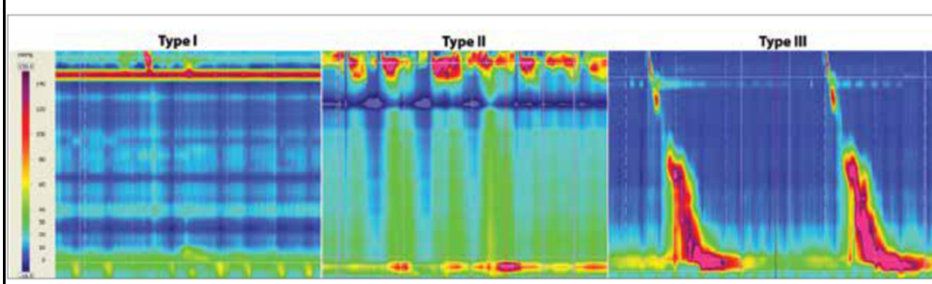
- Rare esophageal motility disorder
- Esophageal aperistalsis
- Impaired LES relaxation

*Loss of inhibitory neurons secreting VIP and NO leads to unopposed excitatory activity and failure of LES relaxation*



DA Patel. An Overview of Achalasia and Its Subtypes.  
Gastroenterology & Hepatology. Volume 13, Issue 7 July 2017

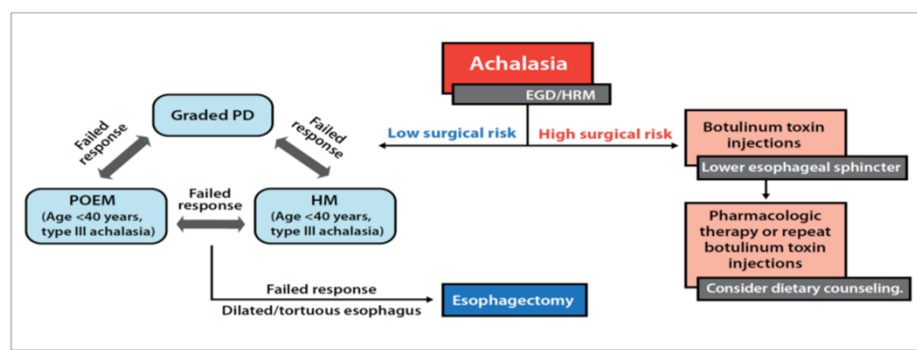
## Achalasia: Subtypes



Type I is characterized by a quiescent esophageal body, type II has pan-esophageal pressurization, and type III is characterized by simultaneous contractions.

DA Patel. An Overview of Achalasia and Its Subtypes.  
Gastroenterology & Hepatology. Volume 13, Issue 7 July 2017

## Achalasia: Treatment Algorithm



DA Patel. An Overview of Achalasia and Its Subtypes.  
Gastroenterology & Hepatology. Volume 13, Issue 7 July 2017



## Achalasia: Treatment Options

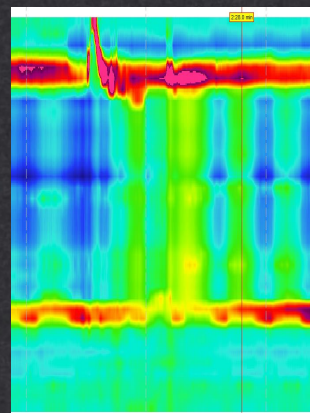
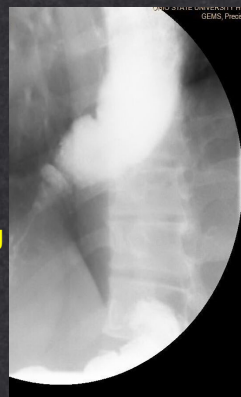
Treatment Options	Pros	Cons
Medications(CaCB/Nitrates)	<ul style="list-style-type: none"> <li>On Demand</li> <li>Minimal risk</li> <li>For non-operative candidates</li> </ul>	<ul style="list-style-type: none"> <li>Least effective</li> <li>Not durable</li> </ul>
Botulinum toxin injection	<ul style="list-style-type: none"> <li>Good option for nonoperative candidates</li> <li>Short procedure time</li> </ul>	<ul style="list-style-type: none"> <li>Durability of 6–12 months</li> </ul>
Pneumatic dilation	<ul style="list-style-type: none"> <li>Most effective nonsurgical option</li> <li>Short recovery time</li> <li>Durability 2–5 years</li> <li>Procedure time &lt;30 minutes</li> </ul>	<ul style="list-style-type: none"> <li>Perforation (1%–5%)</li> </ul>
Surgical myotomy	<ul style="list-style-type: none"> <li>Durability 5–7 years</li> <li>Procedure time ~90 minutes</li> </ul>	<ul style="list-style-type: none"> <li>General anesthesia required</li> <li>Hospital stay of 1–2 days</li> </ul>
Esophagectomy	<ul style="list-style-type: none"> <li>For end-stage disease</li> </ul>	<ul style="list-style-type: none"> <li>High morbidity and mortality</li> </ul>

## Case Study 2:

24-year-old female presented with dysphagia to solids and liquids.

- Mean DCI: NA
- Mean LES IRP: 24 mm Hg
- Mean DL: NA

Diagnosis??

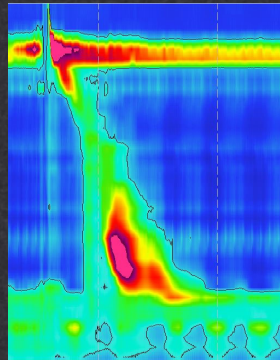


Type 2 Achalasia. Patient sent for myotomy

## Case Study 3:

64-year-old female with CAD, chronic backache on morphine is presenting dysphagia and spasmodic pain in the neck and chest.

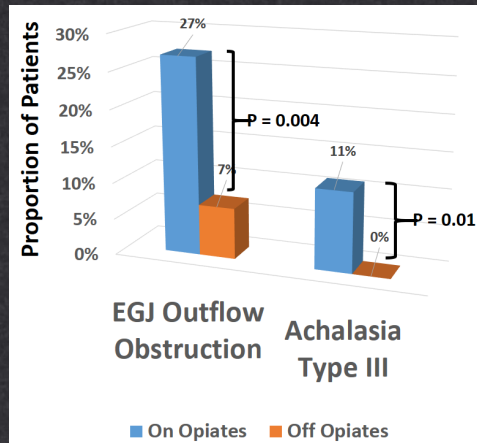
- Mean DCI: 2765
- Mean LES IRP: 18 mm Hg
- Mean DL: 3.8s



Diagnosis?? **Opioid induced esophageal dysfunction**

## Opioid-induced esophageal dysfunction

Opioid-induced esophageal dysfunction is often characterized by EGJ outflow obstruction and type III achalasia pattern.



Ratuapli S, et al. Opioid-Induced Esophageal Dysfunction (OIED) in Patients on Chronic Opioids. *Am J Gastroenterol* 2015; 110:979-984;

## Achalasia syndromes beyond the CC v3.0

CC v3.0 diagnosis	IRP > ULN?	Oesophageal contractility	Notes
Oesophagogastric junction outflow obstruction	Yes	Sufficient peristalsis to exclude type I, II or III achalasia	<ul style="list-style-type: none"> <li>Heterogeneous group</li> <li>Early or incomplete achalasia</li> <li>Can resolve spontaneously</li> <li>Recording artefacts</li> </ul>
Absent contractility	No	Absent contractility	<ul style="list-style-type: none"> <li>Can be achalasia</li> <li>Abnormal FLIP distensibility index supports achalasia</li> <li>Oesophageal pressurization with swallows or MRS supports achalasia</li> </ul>
Distal oesophageal spasm	Yes or no	≥20% premature contractions (DL <4.5s)	Might be spastic achalasia
Jackhammer	Yes or no	≥20% of swallows with DCI >8,000 mmHg·s·cm	Might be spastic achalasia if DL <4.5 s with ≥20% swallows
Opioid effect (not in CC)	Yes	Normal, hypercontractile or premature	Can mimic EGJOO, type III achalasia, DES or jackhammer
Mechanical obstruction (not in CC)	Yes	Absent, normal or hypercontractile	EUS, CT or MRI of the EGJ might clarify the aetiology

Kahrilas, P. J. *et al.* (2017) Advances in the management of oesophageal motility disorders. in the era of high-resolution manometry: a focus on achalasia syndromes. *Nat. Rev. Gastroenterol. Hepatol.* doi:10.1038/nrgastro.2017.132

## GERD-Case based approach



## Gastroesophageal Reflux Disease Definition

**GERD is a condition that develops when the reflux of gastric content causes troublesome symptoms or complications.**

- Mild symptoms once in > 2 days/week
- Moderate/Severe once in >1 day/week

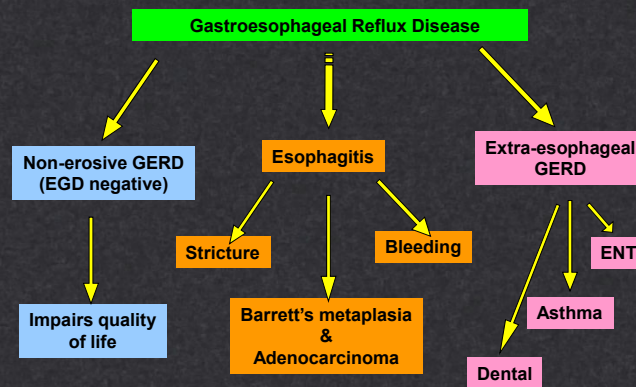
Vakil N, van Zanten SV, Kahrilas P, et al. Global Consensus Group. The Montreal definition and classification of gastroesophageal reflux disease: a global evidence-based consensus. Am J Gastroenterol. 2006;101:1900–1920.

## Risk factors:

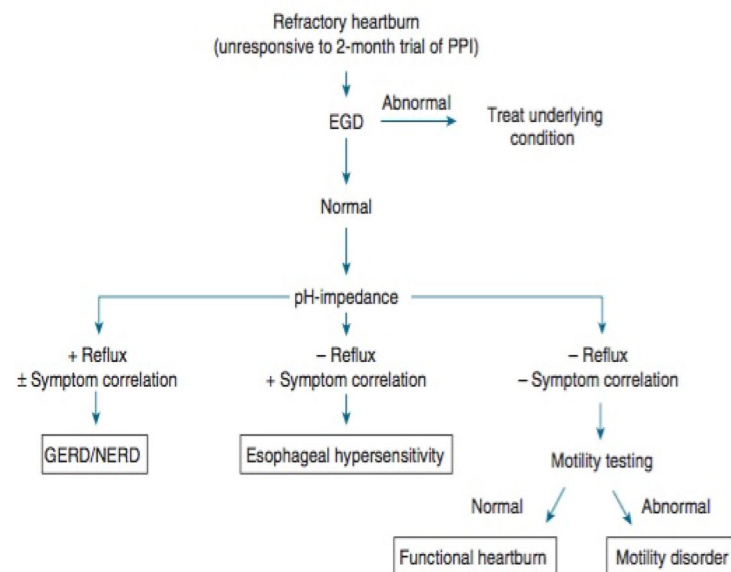
- Obesity
- Family history for GERD
- Tobacco smoking
- Alcohol consumption
- Associated psychosomatic complaints

Locke GR, et al. The American Journal of Medicine. 1999;106(6):642-649  
Hampel H. Ann Intern Med. 2005;143(3):199-211.

# Impact of Gastroesophageal Reflux Disease



Irvine EJ, Hunt RH. *Evidence-Based Gastroenterol*. BC Decker Inc. Hamilton and London. 2001.



## **Goals for Treatment of GERD**

- **Eliminate symptoms**
- **Heal erosive esophagitis**
- **Prevent the relapse of erosive esophagitis and complications from GERD**

## **Life-Style Modifications include:**

- **Elevate the head of the bed on 4" to 6" blocks.**
- **Advise weight loss for obese patients.**
- **Avoid recumbency for 3 hours after meals.**
- **Avoid bedtime snacks.**
- **Avoid fatty foods, chocolate, peppermint, onions, and garlic.**
- **Avoid cigarettes and alcohol.**
- **Avoid drugs that decrease LES pressure and delay gastric emptying.**

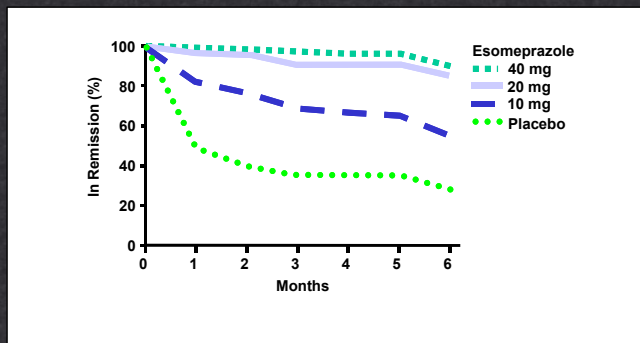


## Medical treatment options:

### Proton Pump Inhibitors:

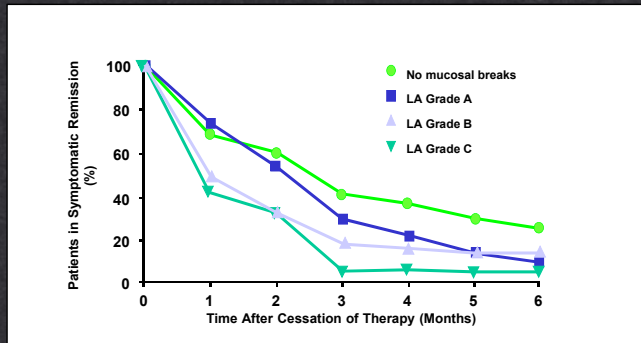
- Higher healing rates in mild to moderately severe reflux esophagitis(80% to 100%).
- Improves dysphagia.
- Decreases the need for esophageal dilation in patients who have peptic esophageal strictures.
- About 70% may have nocturnal acid breakthrough that requires H2RA.

## Maintenance of Healing Erosive Esophagitis



Pooled from Johnson DA, et al., *Am J Gastroenterol*, 2001;96:27-34 and Vakil NB, et al., *Aliment Pharmacol Ther*, 2001;15:927-935.

## GERD Is a Chronic Condition Likely to Relapse



Lundell LR, et al. *Gut*. 1999;45:172-180.

## Appropriateness of PPI use

	Reason for use
Long-term PPI therapy appropriate	<ul style="list-style-type: none"> <li>Barrett's esophagus</li> <li>Healing and maintenance of healed Los Angeles grade C or D erosive esophagitis<sup>a</sup></li> <li>PPI-responsive esophageal eosinophilia</li> <li>Idiopathic (<i>H. pylori</i> and NSAID/aspirin negative) peptic ulcer disease</li> <li>Zollinger-Ellison disease<sup>a,b</sup></li> <li>PPI-responsive GERD/non-erosive reflux disease<sup>a,c</sup></li> <li>Long-term non-selective NSAID users at high-risk for upper GI complications or long-term cox-2 inhibitor users with a prior episode of GI bleeding<sup>a</sup></li> <li>Anti-platelet therapy in patients at high-risk for upper GI complications (age &gt; 65 years or concomitant use of corticosteroids or anticoagulants or history of peptic ulcer disease)</li> <li>Steatorrhea refractory to enzyme replacement therapy in chronic pancreatitis</li> </ul>
Short-term PPI therapy appropriate (4- to 12-week course)	<ul style="list-style-type: none"> <li>Healing of Los Angeles grade A or B erosive esophagitis<sup>a</sup></li> <li>Eosinophilic esophagitis</li> <li><i>H. pylori</i> eradication (in combination with antibiotics)<sup>a,d</sup></li> <li>Stress ulcer prophylaxis in high-risk patients (i.e., critically ill patients with respiratory failure or coagulopathy)</li> <li>Functional dyspepsia</li> <li>Treatment and maintenance of peptic ulcer disease<sup>a</sup></li> <li>Prior to endoscopy for acute upper GI bleeding</li> <li>Following endoscopic treatment of a high-risk ulcer GI bleed</li> </ul>
PPI use not appropriate	<ul style="list-style-type: none"> <li>Corticosteroid users without concomitant NSAID therapy</li> <li>To prevent bleeding from hypertensive gastropathy in cirrhotic patients</li> <li>Acute pancreatitis</li> <li>Stress ulcer prophylaxis in non-critically ill hospitalized patients that are not at high-risk for ulcer formation and GI bleeding</li> </ul>
PPI use of uncertain benefit	<ul style="list-style-type: none"> <li>PPI non-responsive GERD</li> <li>Extra-digestive GERD</li> </ul>

Yadlapati and Kahrilas *BMC Medicine* (2017) 15:36

**Table 2** Quality of evidence and risks of adverse effects associated with long-term proton pump inhibitors (PPIs)

	Potential adverse effect	Nature of evidence	Risk estimate
Causality established, idiosyncratic, rare	Acute interstitial nephritis	Observational, case-control	OR 5.16 (2.21–12.05)
Causality proven but of minimal significance	Fundic gland polyps	Observational	OR 2.2 (1.3–3.8) [6]
	B12 deficiency	Observational, case-control	OR 1.65 (1.58–1.73) [7]
Weak association, causality probable	Small intestinal bacterial overgrowth	Meta-analysis	OR 2.28 (1.23–4.21) [8]
	Spontaneous bacterial peritonitis in cirrhotic patients	Systematic review/meta-analysis	OR 2.17 (1.46–3.23) [9]
Weak association, causality			ent response, (1.78–5.10) [10]
			3.50)
			2.64) [11]
			2.93) <sup>a</sup> [12]
			3.90)
			1.96) [13]
			1.52)
			1.24) <sup>b</sup>
			1.92) <sup>b</sup>

**Decisions to start, properly dose, continue, or discontinue PPI therapy should be personalized based on indication, effectiveness, patient preferences, and risk assessment.**

Yadlapati and Kahrilas BMC Medicine (2017) 15:36

## Medical treatment options:

- **Antacids and Alginic Acid:**
  - Temporarily relieve episodic heartburn
  - Useful add on therapy
- **Histamine H2-Receptor Blocking Agents:**
  - Safe and effective in mild esophagitis
  - Not useful in severe esophagitis
  - Useful for breakthrough symptoms
  - Concern for tachyphylaxis
- **Prokinetic Agents:**
  - Limited efficacy and side effects in up to 30%
- **TLESR Inhibitors:**
  - As addon for non-acid reflux/post prandial reflux



## Indications for anti-reflux surgery

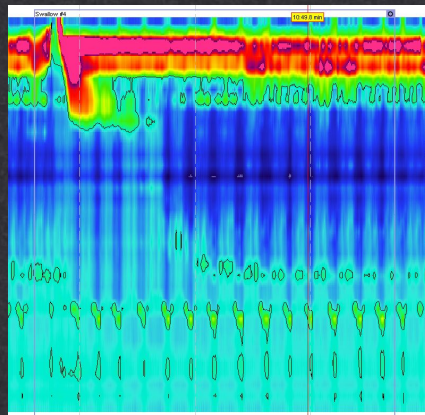
- Unwillingness to remain on medical therapy
- Intolerance of medical therapy
- Medically refractory symptoms with objective evidence of GERD
- GERD in the setting of a large hiatal hernia

Badillo R, Francis D. Diagnosis and treatment of gastroesophageal reflux disease. *World J Gastrointest Pharmacol Ther.* 2014;5(3):105-12.

## Case Study 4:

42-year-old female with prior history of scleroderma is presenting with persistent reflux in spite of twice daily PPI, referred for fundoplication.

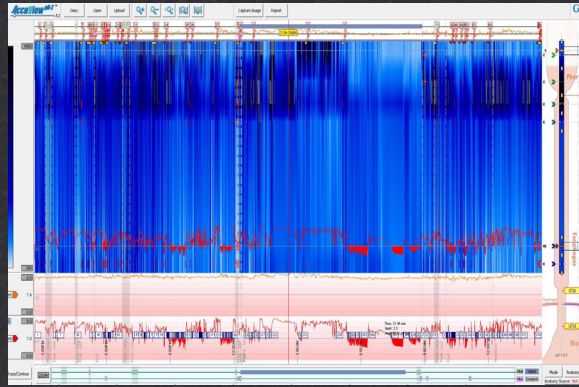
- Mean DCI: NA
- Mean LES IRP: 2mm Hg
- Mean DL: NA



## Case Study 4:

- Acid exposure:
  - Total AET:14.5%
  - Reflux events:112
- Reflux symptom analysis:
  - SI:54
  - SAP: 98

What would be the next step?

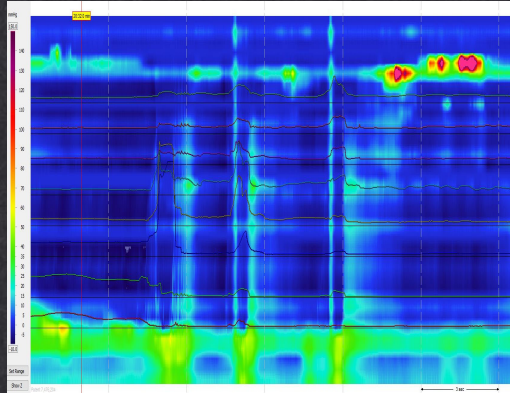


## Case Study 4:

- Educated on lifestyle measures.
- Added H2B at bedtime.
- Was doing much better.

## Case Study 5:

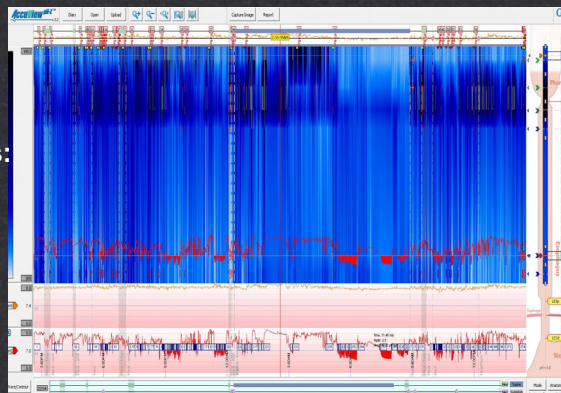
- 28 yr old female with anxiety presenting with persistent heartburn inspite of PPI twice daily
- EGD: normal esophagus with biopsy



## Case Study 5:

- Acid exposure:
  - Total AET:10.5%
  - Reflux events:119
- Reflux symptom analysis:
  - SI:50
  - SAP: 96

What would be the next step?

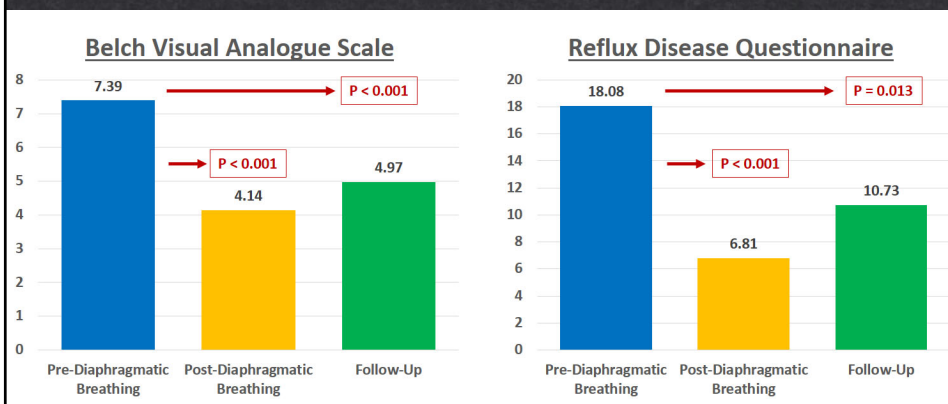




## DDx to PPI-Refractory GERD

- **Refractory reflux symptoms with esophagitis**
  - Eosinophilic esophagitis
  - Pill induced esophagitis
  - Skin disorders like Lichen planus
  - Hypersecretory condition like ZES
  - Genotypic differences in CYP450 2C19
- **Refractory reflux symptoms with normal esophagus**
  - Eosinophilic esophagitis
  - Achalasia
  - Gastroparesis
  - Aerophagia and Belching disorder
  - Rumination syndrome
  - Functional heartburn

## Effect of DBT on belching and GERD



Ong et al. Diaphragmatic Breathing Reduces Belching and Proton Pump Inhibitor Refractory Gastroesophageal Reflux Symptoms. Clinical Gastroenterology and Hepatology 2018;16:407–416

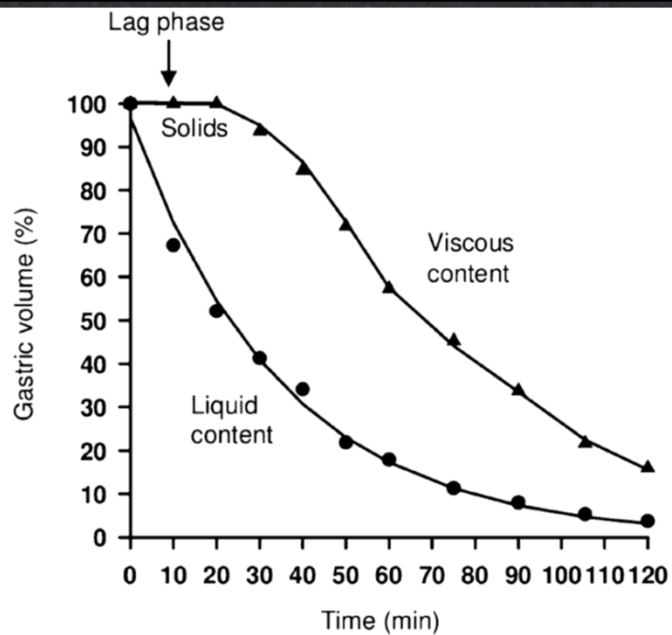
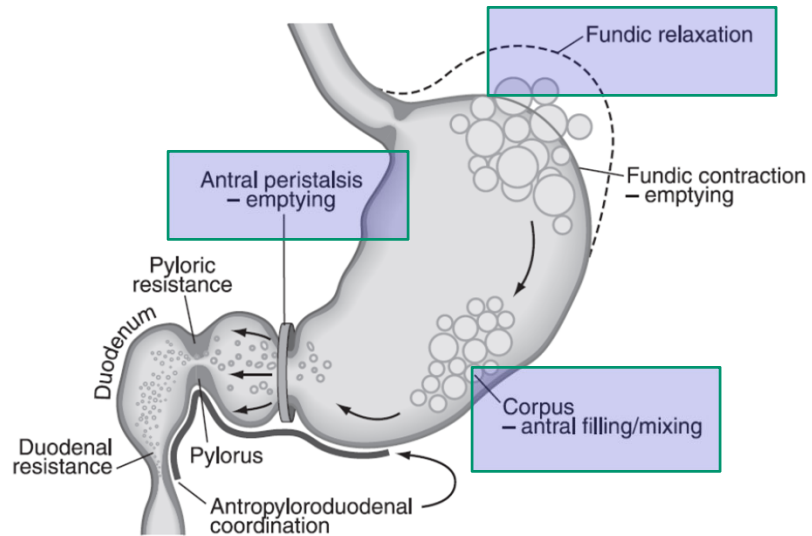


## Case Study 5:

- *Continued PPI,*
- *Started on behavioral therapy and anti-anxiety medication,*
- *Educated on DBT*

## Gastroparesis-Case based approach

# Physiology of stomach



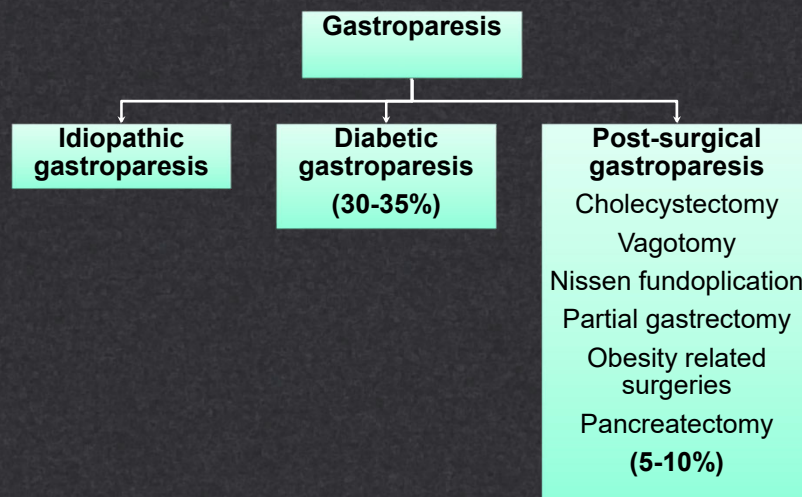
Normal Velocities of emptying of solid and liquid chyme.

## Definition:

***Gastroparesis is defined as a delay in the emptying of ingested food in the absence of mechanical obstruction of the stomach or duodenum.***

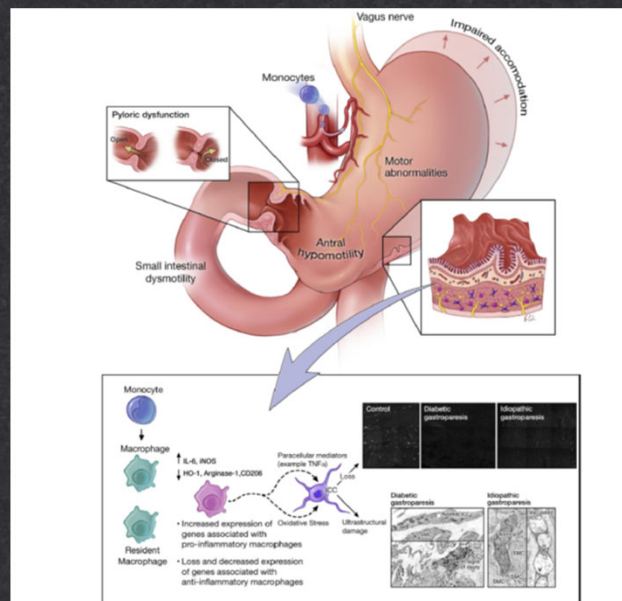
Camilleri M, Parkman H, Shafi M, et al. Clinical guideline: management of gastroparesis. Am J Gastroenterol 2013;108:18–37.

## Etiology of Gastroparesis





## Pathophysiology



Grover, M et al. Gut. 2019 Dec;68(12):2238-2250.

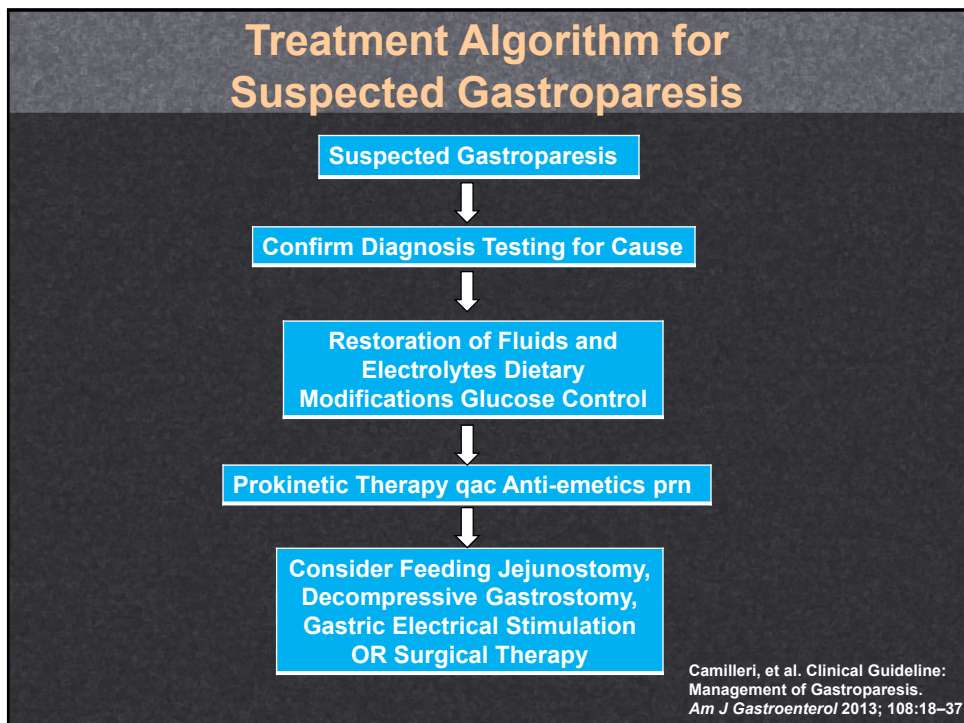
## Clinical Presentation:

- Nausea
- Vomiting
- Early satiety
- Bloating
- Postprandial fullness
- Abdominal pain
- Weight loss/weight gain
- Constipation and/or diarrhea
- Wide glyceemic fluctuations

Characteristic	IG (n 254) N (% or mean) <sup>b</sup>	T1DM (n 78) N (% or mean) <sup>b</sup>	T2DM (n 59) N (% or mean) <sup>b</sup>	Pair-wise P value <sup>a</sup>		
				IG vs all DM	IG vs T1DM	IG vs T2DM
<b>Symptoms prompting evaluation for gastroparesis</b>						
Nausea	214 (84.3)	66 (84.6)	56 (94.9)	.19	.94	.03
Vomiting	152 (59.8)	69 (88.5)	54 (91.5)	<.001	<.001	<.001
Bloating	146 (57.5)	44 (56.4)	37 (62.7)	.75	.87	.46
Early satiety	146 (57.5)	37 (47.4)	44 (74.6)	.75	.12	.02
Postprandial fullness	136 (53.5)	44 (56.4)	39 (66.1)	.18	.66	.08
Abdominal pain	193 (76.0)	47 (60.3)	41 (69.5)	.01	.007	.30
Diarrhea	98 (35.6)	35 (44.9)	30 (50.9)	.09	.32	.08
Constipation	112 (44.1)	32 (41.0)	34 (57.6)	.44	.63	.06
Anorexia	32 (12.6)	12 (15.4)	17 (28.8)	.03	.53	.02
Weight loss	118 (46.5)	41 (52.6)	31 (52.5)	.25	.35	.40
Weight gain	45 (17.7)	14 (18.0)	14 (23.7)	.57	.96	.24
Gastroesophageal reflux	137 (53.9)	43 (55.1)	35 (59.3)	.57	.85	.45
Problems with diabetes control	0 (0.0)	39 (50.0)	27 (45.8)	<.001	<.001	<.001

*In 416 patients from the NIH Gastroparesis Registry, symptoms prompting evaluation more often included vomiting for diabetic gastroparesis and abdominal pain for idiopathic gastroparesis.*

Source: Parkman HP, Yates K, Hasler WL, et al. Similarities and differences between diabetic and idiopathic gastroparesis. *Clin Gastroenterol Hepatol*. 2011;9(12):1056–e134.



## Diagnostic Testing for Gastroparesis:

TABLE 2. Diagnostic Testing for Gastroparesis

Modality	Advantages	Disadvantages
Gastric scintigraphy		
4-hour solid phase	Widely available	Radiation exposure
	Considered the “gold standard” for diagnosis	False positives with liquid phase only studies
Wireless motility capsule		
Smart Pill, given imaging	Avoids radiation exposure	Less validated than scintigraphy
	FDA approved for diagnosis	Cannot be used in those with pacemaker or defibrillator
Radiolabeled carbon breath test		
<sup>13</sup> C-labeled octanoic acid or Spirulina platensis	Low cost	Lack of standardization
		Has primarily been used as a research tool

Gastroparesis: A Review of Current Diagnosis and Treatment Options. Stein, Benjamin; Everhart, Kelly; Lacy, Brian. Journal of Clinical Gastroenterology. 49(7):550-558, August 2015.

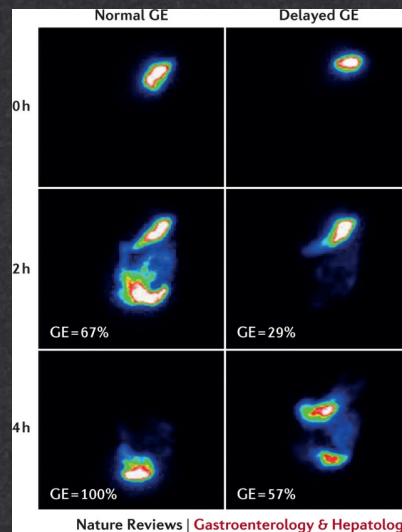
## Radionuclide Gastric Emptying Scintigraphy

- Best current test for measuring gastric emptying because it is sensitive, quantitative, and physiological.
- <sup>99m</sup>Tc sulfur colloid-labeled low-fat egg white meal as a test meal.
- Imaging is performed in the anterior and posterior projections at least at four time points (0, 1, 2, and 4 h).
- The 1 h image is used to help detect rapid gastric emptying.
- The 2 and 4 h images are used to evaluate for delayed gastric emptying.
- **Hyperglycemia (glucose level > 270 mg/dL) delays gastric emptying in diabetic patients.**

Parkman H.P. (2018) Gastric Emptying Studies. In: Bardan E., Shaker R. (eds) Gastrointestinal Motility Disorders. Springer, Cham



## Radionuclide Gastric Emptying Scintigraphy



Keller, J. et al. (2018) Advances in the diagnosis and classification of gastric and intestinal motility disorders. *Nat. Rev. Gastroenterol. Hepatol.*

### Therapeutic options guided by symptom severity

Lifestyle Modifications	Increased meal frequency, decreased meal volume. Aim for low particle size, fat, and fiber diet.	In DM, optimize glycemic control. Balance diet with GP symptoms. Target plasma glucose $\leq 200$ mg/dL	Trial withdrawal of medications known to impair gastric emptying (e.g., GLP-1 analogs, opioids).	→
	Mild	Nutritional assessment with correction of dietary deficiencies.	Consider enteral nutrition if the patient cannot tolerate oral diet.	→
Pharmacotherapy	<b>Prokinetic Agents</b>	Metoclopramide (recommended duration < 12 wk due to risk of TD).	Domperidone (baseline QTC <470 ms in men / <450 ms in women).	→
	<b>Symptom Control</b>	Anti-emetic agents	Erythromycin (ambulatory use limited by tachyphylaxis). Anti-nociceptive agents	→
Interventional Therapy	Gastric Electrical Stimulator May benefit patients with diabetic GP and intractable nausea/vomiting.			→
	Severe	Surgical Options Pyloroplasty and subtotal/total gastrectomy, may benefit patients with post-surgical GP. consider gastric bypass in obese patients		→

Gastroparesis: A Review of Current Diagnosis and Treatment Options. Stein, Benjamin; Everhart, Kelly; Lacy, Brian. *Journal of Clinical Gastroenterology*. 49(7):550-558, August 2015.

## Pro-kinetics:

Medications	Mechanism	Pros	Con
Metaclopramide	D2 Antagonist	Improves gastric emptying. Lowest possible dose (5 mg TID before meals). No long term study available. Efficacy:29-53%. Comparable to Domperidone	Black box warning:>12 weeks use of tardive dyskinesia Acute dystonias Parkinsonism type movements Associated with QTc interval
Domperidone	D2 Antagonist	Improvement in symptoms (54% to 79%). Drug interaction.	Less CNS effects Associated with QTc interval. Increases Prolactin levels. Requires IND for approval.
Erythromycin	Motilin agonist	Useful during acute exacerbation. IV better than PO.	Tachyphylaxis. Associated with QTc prolongation.
Cisapride	5-HT4 agonist	Significant improvement in symptoms.	cardiac arrhythmias and death Requires IND
Prucalopride	5-HT4 agonist	Improves gastric emptying and colon transit times. FDA approved for chronic constipation.	Diarrhea and suicidal ideations. Avoidance in ESRD. No cardiac toxicity document.

## Anti-emetics:

Medications	MOA	Pros	Cons
Diphenhydramine	Antihistamines	Useful in mild nausea/vomiting.	<ul style="list-style-type: none"> <li>Sedative effect.</li> <li>Anticholinergic S/E.</li> </ul>
Hyoscine	Anti-cholinergics	Cheap and widely available. Useful in mild cases.	<ul style="list-style-type: none"> <li>Anti-cholinergic side effects(dry mouth, glaucoma,etc).</li> </ul>
Phenothiazines/ prochlorperazine	D1/D2 Antagonist	Useful in severe nausea and vomiting.	<ul style="list-style-type: none"> <li>EKG changes</li> <li>Psychomotor issues in elderly</li> <li>Dystonia/Parkinsonism</li> </ul>
Ondansetron	5HT3 antagonists	Widely available. Useful in mild vomiting.	<ul style="list-style-type: none"> <li>QT prolongation.</li> <li>Serotonin syndrome.</li> <li>Constipation.</li> </ul>
Transdermal granisetron	5HT3 antagonists	Not widely available/cost. Useful in those who cannot tolerate oral meds.	<ul style="list-style-type: none"> <li>QT prolongation.</li> <li>Serotonin syndrome.</li> <li>Constipation.</li> </ul>
Aprepitant	NK1 receptor antagonists	Not widely available/cost. Useful in reducing N/V.	<ul style="list-style-type: none"> <li>Fatigue.</li> <li>Neutropenia.</li> </ul>
Dronabinol	Agonist of CB <sub>1</sub> and CB <sub>2</sub>	Helpful for N/V when other therapies have failed.	<ul style="list-style-type: none"> <li>Delays gastric emptying.</li> </ul>

## Neuromodulators:

Medications	MOA	Pros	Con
Nortriptyline/ Amitriptyline	TCA	Modest improvement in N/V and abdominal pain	Worsens gastric emptying. Anti-cholinergic side effects. Constipation.
Mirtazapine/ Buspirone	SNRI/SSRI	Improves appetite. Improves fundic accommodation.	Suicidal thoughts. EKG changes. Serotonin syndrome.

## Gastric electric stimulation

- Patient Selection: **Diabetic gastroparesis with refractory N/V even after 1 year of pro-kinetics.**
- Response to therapy:
  - Diabetics.
  - Not on narcotics.
  - Predominant nausea/vomiting.
- Response was modest with 43% over a period of a year and half.

Heckert, J., Sankineni, A., Hughes, W. B., Harbison, S., & Parkman, H. (2016). Gastric Electric Stimulation for Refractory Gastroparesis: A Prospective Analysis of 151 Patients at a Single Center. *Digestive Diseases and Sciences*, 61(1), 168-175.

## Final Case Study

- 42-year-old gentleman with type 2 diabetes(HgbA1c:9) on exenatide presenting with recurrent vomiting and nausea for the last 6 months?

What would be the next step?

Normal upper endoscopy with moderate food retention in the stomach. Bx: negative for H. pylori.

4-hour GES: 43%. What do we do next?

Switch exenatide to insulin+CGM.  
Nutrition consult for gastroparesis.