



**Brigham and Women's Hospital**

Founding Member, Mass General Brigham

# Updates in Upper GI Disorders: GERD, Barrett's Esophagus, and *H. pylori*

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# Disclosures

I have no disclosures to report

I will discuss the use of medications for non-FDA approved indications



# Learning Objectives

## GERD

- Recognize the indications for EGD
- Review current and new treatments

## Barrett's esophagus

- Discuss screening recommendations
- Summarize treatment options

## *H. pylori*

- Explain current treatment recommendations



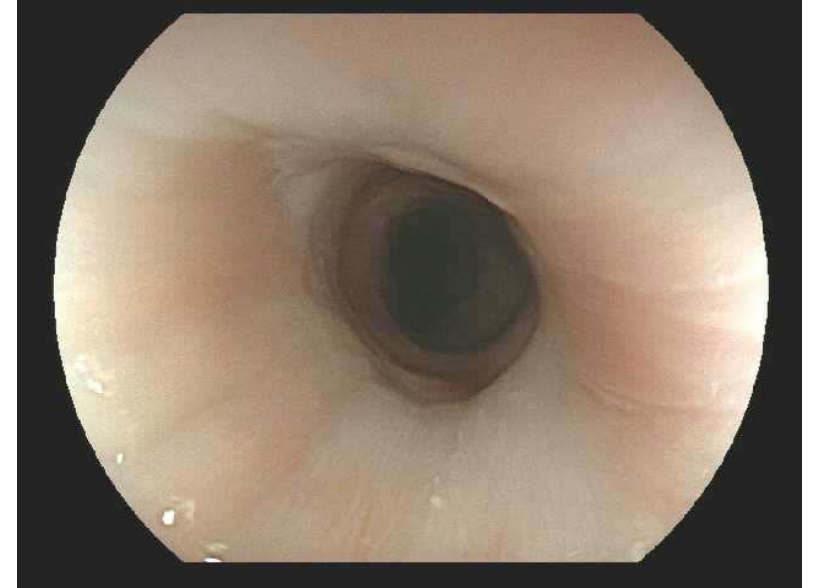
# Gastroesophageal Reflux Disease (GERD)



# Indications for EGD in GERD

Not for diagnosis of GERD

- Most patients will have a normal esophagus



Used to diagnose complications, other causes for symptoms, and evaluate for malignancy

- Erosive esophagitis, Barrett's esophagus, strictures, eosinophilic esophagitis, esophageal cancer



# Indications for EGD in GERD

## 1. Alarm features

- New dyspepsia at age  $\geq 60$  years
  - No age criteria for GERD symptoms
- GI bleeding (overt, occult)
- Iron deficiency anemia
- Dysphagia, odynophagia
- Anorexia, weight loss, persistent vomiting
- Family history of upper GI cancer



# Indications for EGD in GERD

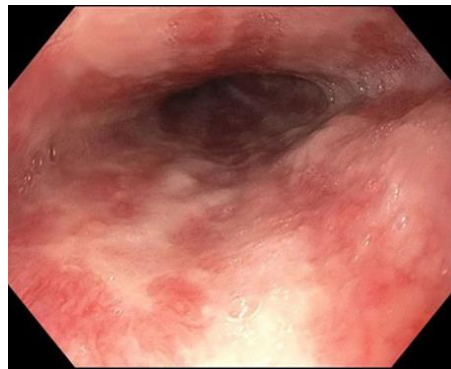
## 1. Alarm features

## 2. GERD symptoms which

- Do not respond to an 8 week trial of standard-dose PPI (ex. omeprazole 20mg daily)
- Recur <3 months after stopping PPI
- [Ideally do the EGD off PPI for 2-4 weeks to see if esophagitis is present]

## 3. Severe esophagitis (LA Grade C and D)

- After PPI x 8-12 weeks to assess healing and r/o Barrett's





# Indications for EGD in GERD

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2. GERD symptoms which

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3. Severe esophagitis (LA Grade C and D)

- After PPI x 2-3 months to assess healing and r/o Barrett's

4. Screen for Barrett's in patients with multiple risk factors

5.  Abnormal upper GI tract imaging

# Prescribing Proton Pump Inhibitors (PPI)

## Optimal dosing

- 30-60 min before eating (breakfast and/or dinner)
- Avoid giving with H2-blockers (can give H2B at bedtime)

## Overprescribed

- Clear indication in only 1/3 of patients

## Similar efficacy

- Differences may relate to drug metabolism polymorphisms – can switch if needed

## Possible adverse effects

- Most data is retrospective (residual confounding) – need more prospective data
- Few concerns are supported by consistent data demonstrating a causal relationship



# Potential PPI Adverse Effects

## GI issues

- *C. diff* and other enteric infections
- SBP in cirrhosis
- Small intestinal bowel overgrowth (SIBO)
- Microscopic colitis
- Possible increased risk of IBD
- ↑ gastrin and atrophy – but no clear increased risk of GI cancer

## Renal

- AIN – idiopathic
- ? CKD – unclear mechanism

## Bone fracture

- ? ↑ osteoclast activity, ↓ Ca absorption (Ca carbonate >> dietary)
- WHI study (2010) – PPI not associated with hip fracture, but was modestly associated with spine, wrist, and total fractures



# Potential PPI Adverse Effects

## Malabsorption

- Magnesium – check prior to long-term use, esp. if diuretics/elderly or hx arrhythmias/prolonged QTc – then check periodically/yearly
- Vitamin B12 – consider periodic/yearly monitoring
- Iron – no clear recommendation to monitor

## Unclear significance

- Dementia
- CV/stroke events
- COVID-19
- Pneumonia

## Drug interactions

- CYP2C19 metabolism
  - Omeprazole & esomeprazole the most
  - Pantoprazole the least
- Clopidogrel – no clear evidence of increased adverse effects
- Protease inhibitors – ↓ absorption



# Prospective PPI Trial (2019)

- Large, prospective, randomized trial
- PPI (pantoprazole 40mg daily) vs placebo
- 17600 older patients (age >65, 78% men, 23% smokers)
  - Pts with stable CAD/PAD receiving rivaroxaban or ASA
- During median f/u of 3 years
  - No difference in pneumonia, hip fracture, CKD, dementia, COPD, gastric atrophy, or cancer
  - Enteric infections other than *C. diff* were more common in pts taking PPI (1.4% vs 1%, p 0.04)
  - *C. diff* 2x more common in pts taking PPI, but not statistically significant



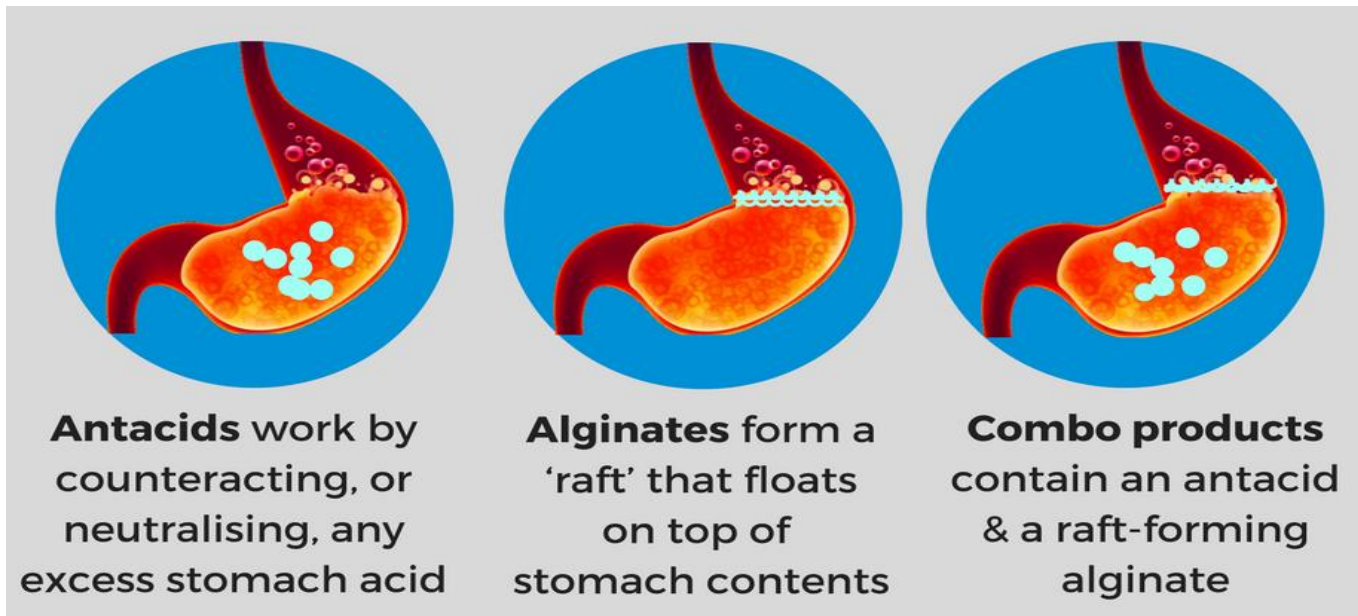
# PPI – General Thoughts

- Overall safe and well-tolerated
- Possible safety issues – need more data
- Balance possible risk of PPI with risk of uncontrolled GERD (symptoms, bleeding, strictures, Barrett's)
- Consider discontinuing PPI w/ frequent symptom re-evaluation
- Use lowest dose needed
- Titrate down dose to avoid rebound acid hypersecretion
- Switch to H2-blockers when possible
- Maintenance PPI recommended if severe esophagitis and Barrett's



# Alginates

- Sodium alginate – polysaccharide derived from seaweed
- Forms a gel that floats above the food/liquid in the stomach
  - Neutralizes the postprandial acid pocket in the proximal stomach
  - Prevent reflux from entering the esophagus



# Alginates

- Used in
  - Mild GERD
  - Refractory acid reflux (on PPI)
  - Non-acid reflux
- Dosing
  - Liquid, tablet
  - After meals and before bed
- Limited efficacy data
  - Mild GERD > refractory GERD or non-acid reflux
- Minimal side effects





# Vonoprazan – New Treatment for Erosive GERD


- Potassium-competitive acid blocker (PCAB)
  - May decrease intragastric pH and maintains it to a greater degree than PPI
- RCT (Laine, et al. Gastro, 2023)
  - Vonoprazan was noninferior and superior to the PPI lansoprazole in healing and maintenance of healing of erosive esophagitis
    - Benefit was seen predominantly in more severe erosive esophagitis
- Adverse events - abdominal pain, dyspepsia, hypertension, UTI
  - Longer terms concerns may be similar to PPI, but less is known at this time
- Approved by FDA in October 2023
- Role in treatment still being determined



# Barrett's Esophagus



# Screening for Barrett's Esophagus

- Expert opinion, not based on RCT data
  - A single screening EGD for patients with chronic GERD symptoms\*\* and 3+ additional risk factors for Barrett's
    - Male sex
    - Age >50 years
    - White race
    - Obesity
    - Tobacco smoking
    - Hiatal hernia?
    - Family hx of Barrett's/esophageal adenocarcinoma in first- degree relative
- \*\* 40% of patients with esophageal adenocarcinoma do not have chronic GERD symptoms
- If initial EGD is negative, repeating EGD for BE screening not recommended
  - If EGD reveals severe esophagitis (LA Classification B/C/D) → repeat EGD after  PPI for 2-3 months

# Novel Screening Modalities for Barrett's

- Swallowable, non-endoscopic capsule device combined with a biomarker
- Swallowed then withdrawn orally – obtaining esophageal cytology samples
- Can be performed in an office setting without sedation
- May make screening for Barrett's easier and more cost-effective
- More research is needed




# PPI for Barrett's Esophagus - 2021 Meta-Analysis

- Meta-analysis of 12 studies (>155,000 patients) with Barrett's esophagus
- PPI use associated with a significantly lower risk of progression to high-grade dysplasia and/or esophageal cancer compared to no PPI (OR 0.47)
- Prospective studies needed



# Management of Barrett's Esophagus

- PPI
  - Data suggests decrease progression to cancer
  - Generally recommend indefinitely at lowest dose to control GERD symptoms (including asymptomatic GERD)
- ASA/NSAIDs
  - May decrease progression to cancer
  - Not recommending solely for Barrett's
- Endoscopic surveillance
  - No dysplasia: 3-5 years
    - Segment  $\geq$  3cm in 3 years; segment  $<$  3cm in 5 years (ACG Guidelines 2022)
  - Low-grade dysplasia (LGD): 6-12 months
  - High-grade dysplasia (HGD): 3 months
-  Radiofrequency ablation – HDG, ? LGD

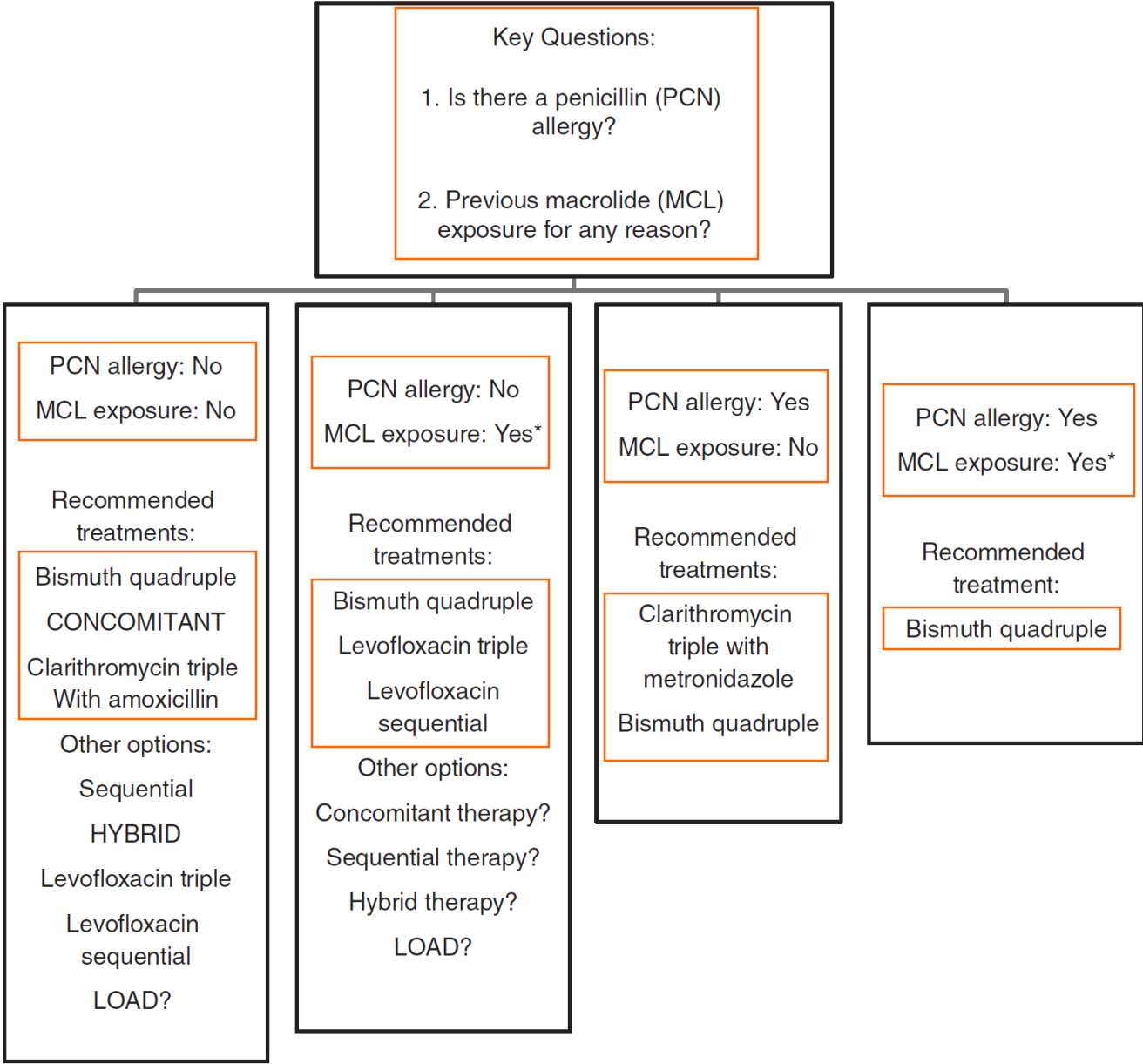
# *H. Pylori* Treatment



# H. pylori Treatment Guidelines

American College of Gastroenterology (ACG)

Concomitant – PPI, amoxicillin, clarithromycin, metronidazole



\*In regions where clarithromycin resistance is known to be >15% utilize recommendations for patients with a history of macrolide exposure

For drugs, doses, and durations of specific first-line regimens, see Table 2.





# First Line Treatment Summary


- Limited information on *H. pylori* resistance in the US → assume clarithromycin resistance is > 15% unless local resistance is known
- Macrolide use is common and patients don't remember if they have taken it in their lives



- Generally avoid “triple therapy”
- Bismuth “quad therapy” increasingly used as 1<sup>st</sup> line



# Regimens Beyond Triple & Quad Therapy

- In general:
  - Clarithromycin, metronidazole, and fluoroquinolones should not be used again due to concern for resistance
  - Amoxicillin and tetracycline can often be reused as resistance is rare
- Levofloxacin – resistance a concern (consider prior exposure)
  - Levofloxacin, amoxicillin, and PPI BID x 14 days
- Rifabutin (anti-TB drug) – H. pylori resistance rare
  - Rifabutin, amoxicillin, and PPI BID x 14 days
  - Recent study supports use as salvage therapy (Chen J, et al. J Infect Dis 2023)
  - Side effects
    - Cytopenias (less likely with short course), uveitis/myelotoxicity (rare)
    - Drug-drug interactions (similar to rifampin)
    -  – May increase mycobacteria resistance

# Consider Penicillin Allergy Testing

- Most patients with a history of penicillin (PCN) allergy do not have true PCN hypersensitivity
  - 5-10% of US population report PCN allergy →
  - 90% have negative skin testing and can tolerate PCN
- Consider referral for allergy testing after failure of a 1<sup>st</sup> line treatment to see if an amoxicillin-containing regimen can be given



# Prevent Treatment Failures

- Think about resistance
  - Especially to clarithromycin, metronidazole, and levofloxacin
- Explain regimen & reinforce compliance
- Consider sensitivity testing after 2 failed regimens
  - Culture – requires EGD and special processing, increasingly available via commercial labs
  - Molecular testing – the future, but not widely available and insurance may not cover




# Confirmation of *H. pylori* Eradication

- Who?
  - Anyone that has been treated, but especially PUD, gastric cancer/MALT
- Which tests?
  - Breath test or stool antigen; not serology
  - EGD if doing for another reasons (e.g. gastric ulcer follow-up, persistent dyspepsia)
- When?
  - Avoid false negatives due to bacterial suppression
    - 4-8 weeks after *H. pylori* treatment
    - Off antibiotics/bismuth for 4 weeks and PPI for 2 weeks



# Vonoprazan – New Treatment for *H. pylori*

- Potassium-competitive acid blocker (PCAB)
  - May decrease intragastric pH and maintains it to a greater degree than PPI
- Use of this instead of a PPI may improve *H. pylori* eradication rates
- Previous data limited to East Asian countries
- RCT (Chey, et al. Gastro 2022) – first clinical trial from the US and Europe
  - Vonoprazan triple (with amoxicillin and clarithromycin) and dual (with amoxicillin) were superior to PPI-based triple therapy (80.8% and 77.2% vs 68.5%), especially in clarithromycin-resistant strains (65.8% and 69.6% vs 31.9%)
- Notes:
  - Cure rate for PPI-based triple therapy was 68% - we should be avoiding clarithromycin-based regimens
  - Overall efficacy of vonoprazan-based triple therapy improved to 81%, still not at recommended target of >90% for first line treatment
  - Efficacy of vonoprazan-based therapy of clarithromycin-resistant strains still low
- Approved by FDA in May 2022
-  Role in treatment still being determined

# Summary

## GERD

- EGD should be performed to diagnose complications, other causes for symptoms, and evaluate for malignancy – and should be repeated after an EGD shows severe esophagitis
- Be mindful about PPI therapy
- Consider sodium alginates
- Vonoprazan a new possible treatment (instead of PPI), role in treatment TBD

## Barrett's esophagus

- Screening can be considered in patients with multiple risk factors
- PPI therapy is generally continued at lowest dose to control GERD symptoms

## *H. pylori* treatment

- Review prior antibiotic exposure, more commonly using quadruple therapy
- Consider penicillin allergy testing to allow use of amoxicillin
- Consider sensitivity testing after 2 failed regimens



Vonoprazan a new possible treatment (instead of PPI), role in treatment TBD

# Selected References

## GERD

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