

Overview of Gastrointestinal Bleeding

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Overview

- Epidemiology
- Definitions of gastrointestinal (GI) bleeding
- Differential diagnosis
- Clinical history and examination- key points
- Initial evaluation and management
- Diagnostic evaluation options

Epidemiology

- Upper GI bleed is approximately 67/100,000 people
- Lower GI bleed is approximately 36/100,000 people
- Morbidity and mortality with over \$1 billion in direct medical costs annually
- Hospitalization rate of upper GI bleed in the USA decreased by 21% from 2002 to 2012
 - Increase use of treatments, improved hemostatic techniques.

Definitions of GI Bleed

- Hematemesis
 - Vomiting of fresh blood
- Coffee ground emesis
 - Slowed or stopped
 - Within red blood cells, iron oxidizes following exposure to gastric acid

Definitions of GI Bleed

- Melena
 - Black tarry stool
 - NOT typically dark, formed stool
 - Only needs 50-100cc of blood to become melena
 - Upper GI bleed vs lower GI bleed
 - ~5-10% can be from small bowel or proximal colon
- Hematochezia
 - Passage of bright red blood per rectum (BRBPR), maroon colored, or clots

Definitions of GI Bleed

- Overt vs Occult
- Overt:
 - Visible blood
 - Bright red, altered blood (melena)
 - Occult:
 - No visible blood identified
 - Presents as iron deficiency anemia, positive stool test for occult blood
- Obscure:
 - No bleeding source identified
 - May be overt or occult

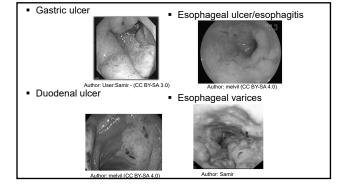
Upper vs lower GI Bleed

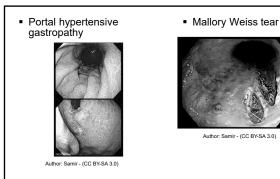
- Factors that increase the likelihood of upper GI bleed:
 - Patient history of melena (LR 5.1-5.9)
 - Melena on examination (LR 25)
 - Nasogastric lavage with blood or coffee ground contents (LR 3.6)
 - BUN/Cr >30 (LR 7.5)

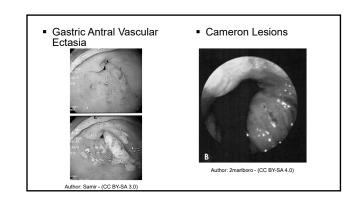
Differential Diagnosis- Upper

- Gastric/ duodenal ulcers*
- Esophagitis/ gastritis
- Esophageal or gastric varices
- Portal hypertensive gastropathy
- Arteriovenous malformations (AVM)
- Mallory-Weiss tear
- Erosions
- Dieulafoy lesion

- Gastric antral vascular ectasia (GAVE)
- Mass lesions
- Hemobilia
- Hemosuccus pancreaticus
- Aortoenteric fistula
- Cameron lesions
- latrogenic







Differential Diagnosis-Lower

- Diverticulosis
- Angiodysplasia
- Hemorrhoids
- Ischemic
- Post biopsy or polypectomy
- Anal fissures
- Radiation-induced telangiectasia
- Infectious
- Inflammatory bowel disease
- Ulcers
- Polyp
- Carcinomas

Diverticulosis



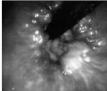
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Angiodysplasia



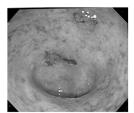
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Hemorrhoids



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Ulcerative colitis



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History

- Past medical history
 - Prior episodes of bleeding
 - Liver disease, cardiac disease (including aortic aneurysms), kidney disease, hematologic disorders
 - History of peptic ulcer disease (PUD) or H pylori
 - Malignancy
 - History of alcohol abuse
 - Recent procedures: colonoscopy, AAA repair, radiation
 - History of gastroenteric anastomosis

History

- Medications review
 - Non steroidal anti inflammatory drugs (NSAIDs)
 - Aspirin
 - Medications associated with pill esophagitis
 - Antiplatelet and anticoagulants
 - Other less obvious medications have been associated with GI bleeding
 - Psychiatric medications, blood pressures medications
 - Bismuth, iron can turn the stool black

Physical

- Evaluate for signs of hemodynamic instability
 - Vitals/orthostatic
- Abdominal exam
- Rectal exam- evaluate for fissures, hemorrhoids, mass, stool exam

Initial Evaluation and Management

- Assessment of hemodynamic status
- Placement of 2 large bore IV lines or central line
- Secure airway if needed
- Labs: Complete blood count, PT/INR, lactate, liver function tests, type and cross
- Transfuse for hemoglobin <7 (or <8 if cardiac), platelet >50
- Resuscitate!

Initial Evaluation and Management

- Risk Stratification Scores
 - Glasgow Blatchford Score
 - Stratifies upper GI bleeding patients who are "low risk" and candidates for outpatient treatment
 - Score 0 is low risk
 - Evaluates: hemoglobin, systolic blood pressure, pulse, BUN, "no melena or syncope", no past or present liver disease or heart failure

Medication Management

- Proton pump inhibitorInhibit gastric acid secretion
 - Heal ulcers, improve platelet aggregation and clot development by raising gastric pH
 - Has been shown to reduce risk of rebleeding (high risk stigmata) and the need for endoscopic intervention
 - High dose PPI- comparable outcomes in dosing (bolus + drip vs bolus + 40mg IV BID)
- If concerned for variceal bleeding:
 - IV somatostatin like octreotide
 - IV antibiotic to empirically cover for spontaneous
- bacterial peritonitis

 Consider holding patient's home blood thinners (risk vs benefits)

Medication Management

- Pro-motility agent
 - Helps to clear the stomach for improved visualization and decreases the need for repeat endoscopy
 - Erythromycin 250 mg IV over 20-30 minutes about 30-120 minutes before EGD
 - Metoclopramide 10mg over 1-2 minutes

Diagnostic Evaluation

- Nasogastric tube
 - Used less often
 - Negative (clear) nasogastric tube aspirate does not rule out an upper GI
 - Bile can help confirm tube in duodenum however may see in stomach due to reflux
 - Can be helpful for gastric emptying however inferior to pro-motility agents



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Diagnostic Evaluation

- Stool guaiac/ hemoccult
 - Great tool for colon cancer screening
 - NOT a test for acute GI bleed
 - False positives:
 - Medications (ASA, NSAIDs)
 - Extra-intestinal blood loss (epistaxis, hemoptysis)
 - Trauma
 - Exogenous peroxidase activity: red meat, fruits, uncooked vegetables

Diagnostic Evaluation

- Endoscopy
 EGD/ upper endoscopy
 Evaluates up to duodenum

 - Push enteroscopy
 Evaluates small bowel
 - Capsule endoscopy
 Evaluates entire GI tract
 - Single balloon enteroscopy (upper and
 - Evaluates small bowel- much further

 - than push enteroscopy

 Colonoscopy
 Evaluates terminal ileum and colon
- Timing:

 Within 24-48 hours after presentation





Diagnostic Evaluation

- Imaging
 - CT angiography
 - Diagnostic and therapeutic
 - Bleeding rate at least 0.3-0.5 to 1.0cc/min
 - Tagged RBC scan
 - Not therapeutic
 - Bleeding rate at least 0.1-0.5cc/min

Therapeutic Management

- Some bleeds typically resolve on their own!
- Endoscopic therapy
 - Epinephrine injection
 - Coagulation
 - Hemoclip
 - Band ligation
- Interventional Radiology
- Surgery



Key Takeaway Points

- Although upper GI bleed typically refers to melena and lower GI bleed to hematochezia, this is not absolute
- There is no utility in hemoccult in active signs of
- Placement of nasogastric tube for the evaluation of GI bleeding is less frequently used
- Resuscitate!



Management of Acute Gastro-esophageal Variceal Bleed

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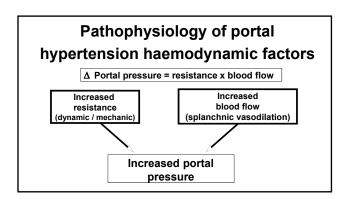
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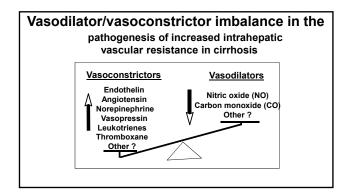
Objective

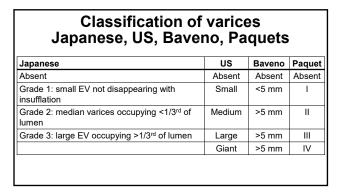
- Introduction
- Various Classifications of EV
- Predictors of bleeding including HVPG
- Varices management
 - Pre-primary prophylaxis
 - Primary prophylaxis
 - Active variceal bleed
 - Secondary prophylaxis

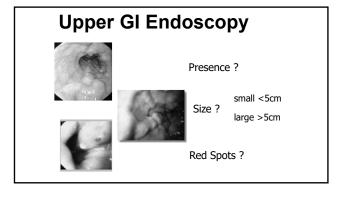
Introduction

- At diagnosis of cirrhosis, varices are present in:
 - 20-40% of compensated patients.
 - 40-60% of patients with ascites. (Schepis F, et al. Hepatology 2001)
 - 5% develop new varices per year.
 - Once developed, varices increase from small to large at 10 – 15% per year.
- Once developed, 25% of varices bleed at 2 years. (deFranchis R, Primigagni M. Clin Liv Dis 2001)
- Mortality due to variceal bleed ranges: 5-15 %







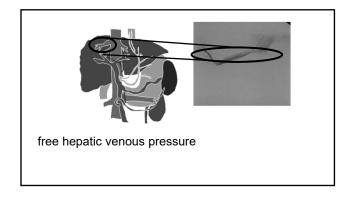


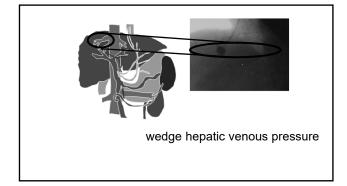
Predictors of Bleeding

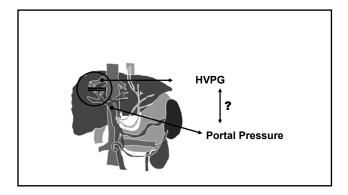
- 1. Variceal size:
 - Small varix 10% at 2 years.
 - Large varix 30% at 2 years.
- 2. Presence of red signs.
- 3. Severity of underlying liver disease:
 - Child A 17%.
 - Child B 31%.
 - Child C 39%. (NIEC New Engl J Med 1988)
- 4. MELD score
- 5. Hepatic Venous Pressure Gradient (HVPG)

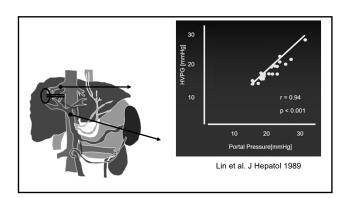
Hepatic Venous Pressure Gradient

- Most commonly used for measurement of portal pressure
- HVPG—gradient between the wedged and free hepatic venous pressure (normal gradient, <5 mm Hg).
 - Polio J, et al. Hemodynamic factors involved in the development and rupture of esophageal varices. Semin Liver Dis 1986;6:318-331

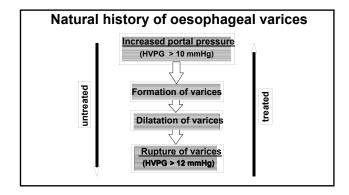








Prognostic value of HVPG in patients with chronic liver disease | Measurement | Significance | |-5 mm Hg | Normal | |-6-10 mm Hg | Preclincal sinusoidal portal HTN | |≥10 mm Hg | Clinically significant portal HTN (CSPH) | |≥ 12 mm Hg | Increased risk for rupture of varices | |≥ 16 mm Hg | Increased risk of morality | |≥ 20mm Hg | Treatment failure and mortality in AVB



Case: Acute Variceal Bleed

- 55 M with HCV and ETOH cirrhosis with moderate ascites. Presenting with UGI bleed. BP: 95/55; HR:110.
- Actively drinking for >10 yrs. H/o IVDU in 1990s.
- Blood work: Hb: 6.5; Plat: 65; LFTs: bill: 3.5, AP: 151; INR: 1.8:
- US shows features of cirrhosis, ++ ascites, no HCC.
- EGD: large >5 mm EV with red wale signs and cherry red spots.

Management of Acute Variceal Hemorrhage

- Prompt resuscitation, hemodynamic support, and correction of hemostatic dysfunction.
- Empirical vasoactive pharmaco-therapy is indicated in variceal hemorrhage.
- Subsequently, EGD facilitates an accurate diagnosis and endoscopic therapy.

Calès P, et al. Early administration of vapreotide for variceal bleeding in patients with cirrhosis. N Engl J Med 2001;344:23-28

Pharmacologic Therapy

- An attractive first-line approach in patients with probable variceal hemorrhage.
- Terlipressin:
 - Synthetic vasopressin analogue.
 - longer half-life has led to its successful use for variceal bleeding.
 - Terlipressin appears to be as effective as vasopressin or somatostatin.

Feu F, et al. Double-blind RCT comparing terlipressin and somatostatin for acute variceal hemorrhage. Gastroenterology 1996;111:1291-1299

Somatostatin

- Naturally occurring peptide, and its synthetic products octreotide and vapreotide.
- Stops variceal hemorrhage in up to 80% of patients.
- Octreotide works :
 - by preventing postprandial hyperemia or
 - by reducing portal pressure through effects on vasoactive peptides.
- Excellent safety profile.
- The addition of octreotide to EST or EVBL resulted in improved control of bleeding and reduced transfusion requirements.

 - Hasnain A. Shah, **Khalid Mumtaz**, et al.. Sclerotherapy Plus Octreotide Versus Sclerotherapy alone in the management of GOV Hemorrhage. *J Ayub Med Coll Abbotabad* 2005;17(1).

Abid S, Mumtaz K, et al. Efficacy And Safety Of Terlipressin Vs Octreotide As Adjuvant Therapy In Bleeding Esophageal Varices. Am J Gastroenterol 2009; 104:617–623;

- Consecutive cirrhotic patients with EV bleed were randomized to
- Consecutive cirriotic patients with EV bleed were randomized to Terlipressin (Group A, 163) or Octreotide (Group B, 161).

 Outcomes: Efficacy, safety, overall survival and length of hospital stay.

 Control of variceal bleed: 151 (92.63 %) in TERLI and 154 (95.6 %) patients in OCTREO (Cl: 0.22 1.5).

 Death: overall 16 deaths (3 failure to control bleed and 13 from other causes.

 - Death : Overlan is School 1. The causes); LOS: TERLI (108.40 ± 34.81) has shorter LOS as compared to OCT (126.39 ± 47.45 h), (P ≤ 0.001).
- CONCLUSION:
- The efficacy of TERLI was not inferior to OCTREO as an adjuvant therapy for the control of EV bleed and in-hospital survival.
- The length of hospital stay in the TERLI was significantly shorter.

Endoscopic Therapy

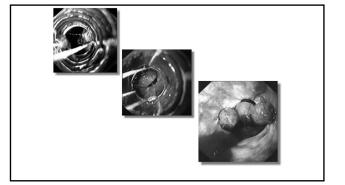
- Endoscopic Sclerotherapy:
 - It stops bleeding in 80 to 90% of acute variceal hemorrhage.
- The advantages of EST:
 - ability to establish definitive control of bleeding under direct vision.
- Drawbacks:
 - risk of local complications, including perforation, ulceration, thrombosis and stricture.

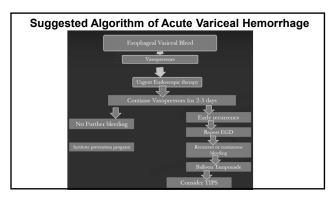
Endoscopic band ligation



- RCTs of acute variceal bleeding have shown that EBL is equivalent to sclerotherapy in achieving initial
- The complications associated with EBL are fewer and include superficial ulcerations and, rarely, the formation of strictures.

 - Khalid Muntaz, Hasnain Shah, et al. Comparison of EBL with EST in bleeding es (Abstract) No. M1263 Apr. 2004; 126(4): A728. Lo GH, et al. Emergency banding ligation versus scienotherapy for the control of a 1997;26:1101-1104.





Secondary Prophylaxis

- Secondary prophylaxis should be instituted after initial episode due to high risk of recurrent bleed.
- Variceal hemorrhage recurs in approximately 2/3 of patients.
- Endoscopic predictors of early recurrence:
 - active bleeding at the time of the initial endoscopy,
 - stigmata of recent bleeding and
 - large varices.

NSBB Therapy

 Reducing the portal pressure by > 20% from the base-line value results in a reduction in the cumulative probability of recurrent bleeding from 28% @ 1 yr, 39% @ 2 yr, and 66% @ 3 yrs to

4%, 9%, and 9%, respectively.

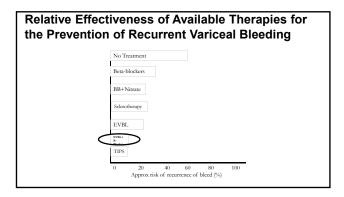
- Feu F, et al. Relation between portal pressure response to pharmacotherapy and risk of recurrent variceal haemorrhage. Lancet 1995;346:1056-1059
- Several RCTs, including a meta-analysis, have demonstrated that non-selective BB (Nadolol, Carvedilol) decrease the risk of recurrent bleeding and prolong survival.
 - Bernard B, et al. . B-adrenergic antagonists in prevention of GI rebleeding in patients with cirrhosis: a meta-analysis. Hepatology 1997;

Endoscopic Band Ligation

- EVBL is highly effective in obliterating varices.
- Ligation is associated with a lower risk of recurrent bleeding and fewer complications,
- EVBL is performed 2-4 weekly until varices are eradicated, which typically requires 3-4 sessions.
- Approaches that combine methods, usually including an endoscopic treatment and a pharmacologic treatment are effective.
 - Rosario Gonzalez, et al. Meta-analysis: Combination Endoscopic and Drug Therapy to Prevent Variceal Rebleeding in Cirrhosis. Ann Intern Med. 2008

Rosario Gonzalez, et al. Meta-analysis: Combination Endoscopic and BB Therapy to Prevent Variceal Rebleeding in Cirrhosis. *Ann Intern Med.*

- Study selection: RCTs comparing endoscopic plus BB therapy with either therapy alone.
- Data synthesis: 23 trials (1860 patients) included.
- Results: Combination therapy reduced overall rebleeding more than endoscopic therapy alone (RR: 0.68; Cl: 0.52 to 0.89) or beta-blocker therapy alone (RR: 0.71; Cl: 0.59 to 0.86).
- Combination therapy also reduced variceal rebleeding and variceal recurrence.
- Reduction in mortality from combination therapy did not statistically significantly differ from that from endoscopic (OR: 0.78; Cl: 0.58 to 1.07) or drug therapy (OR: 0.70: 0.46 to 1.06).
- Conclusion: A combination of endoscopic and drug therapy reduces overall and variceal rebleeding in cirrhosis more than either therapy alone.



Conclusion

- Bleeding from esophageal varices is dependent on severity of liver cirrhosis.
- Resuscitation is integral in management of EVB
- Vasopressors are helpful in initial stability of EVB
- Endoscopic band ligation is effective in securing initial active EV bleeding.
- Combination of repeated EBL and NSBB is effective for secondary prophylaxis.
- TIPS is needed in selective patients who don't respond to endoscopic intervention.