

CHA BUNDANG MEDICAL CENTER

# 혈변의 진단과 치료

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# 1. Classification and definition

# Classification and definition

## 1. Overt vs Occult

Overt : hematemesis, melena, black tarry stool, hematochezia

Occult : In the absence of overt bleeding - symptoms of blood loss or anemia (lightheadedness, syncope, angina, dyspnea),  
Iron deficiency anemia, positive stool occult blood

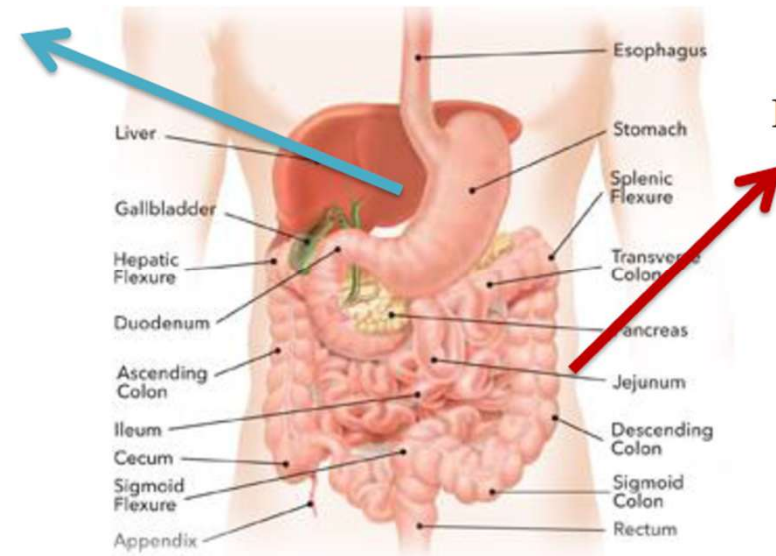
## 2. According to the site of bleeding

Upper GI bleeding : Esophagus, Stomach, Duodenum (Proximal to Treitz ligament)

Lower GI bleeding : Colon, small intestine

Obscure GI bleeding : unclear source

**Upper GI bleeding**



**Lower GI bleeding**

## 2. Initial assessment

# Initial assessment

## 1. History

Symptom:

- 1) Hematemesis (DDx. blood tinged vomit, dark green or tarry vomit)
- 2) Melena (DDx. black tarry stool, dark green or brown stool)
- 3) Hematochezia (DDx. anal bleeding, blood tinged stool or diarrhea)

Duration: Acute (3 days) or Chronic (Several days or more)

Associated symptom: Abdominal pain, Diarrhea, Nausea/Vomiting, Weight loss

Previous history:

- 1) Previous gastrointestinal bleeding
- 2) Medical history: Hypertension, DM, Vascular disease, Liver cirrhosis, Cancer, *H. pylori*
- 3) Surgical history: GI tract operation, Recent endoscopy (EMR/Polypectomy/ESD/Biopsy)

Drug history: Antithrombotics, NSAIDs or steroid, etc

Social history: Alcohol, Smoking, Travel, Stress

# Initial assessment

## 2. Physical exam and simple test – Add clues to the impression

Check exact color of suspicious bleeding: Rectal exam, Patient's photo, etc.

EGD or sigmoidoscopy for stable ambiguous patient

Chronic and low probability actual bleeding: CBC, Stool occult blood

Tenderness on abdomen: Indicated high probability of inflammatory disease

Rectal bleeding has been reported to be the sixth most common symptom prompting an outpatient clinic visit.<sup>1</sup> Anorectal causes account for from 3% to 10% of patients presenting with hematochezia.<sup>2-4</sup> In a study that used artificial neural networks, 12.5% of patients with acute lower GI bleeding had an anorectal etiology.<sup>5</sup> The years.<sup>7,8</sup> Mortality from lower GI bleeding (0.6%) is significantly lower than that from upper GI bleeding (2%).<sup>11</sup> However, outcomes data for anorectal causes of

### Editorials

#### Rectal bleeding in general practice:

new guidance on commissioning

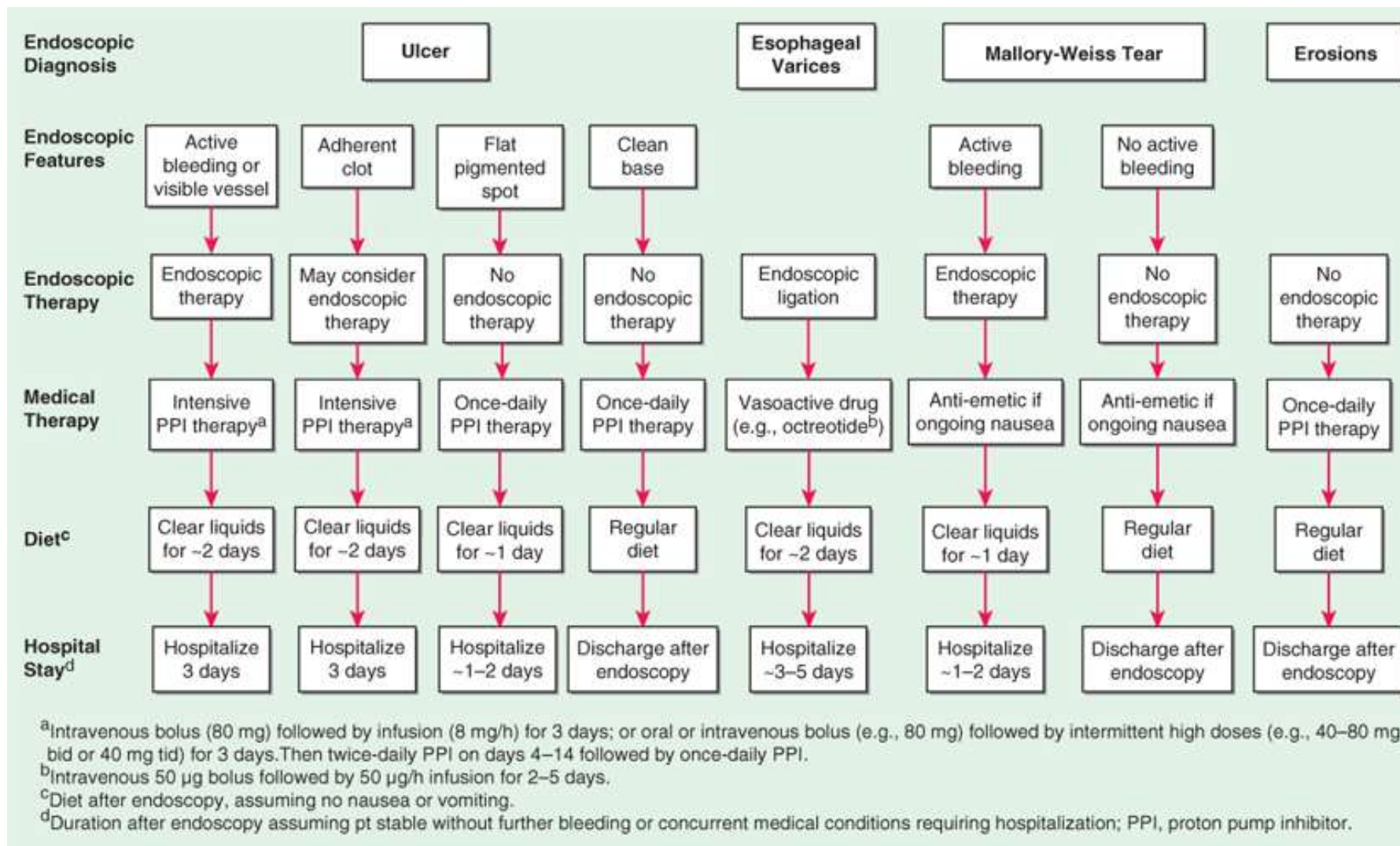
an underlying sinister cause. In the vast majority of patients in primary care the cause will be benign and most often a benign anorectal condition such as haemorrhoids and/or an anal fissure. However, rectal



### **3. Clinical guidelines**



# 1-1) Upper - Traditional Harrison's algorithm



Source: J.L. Jameson, A.S. Fauci, D.L. Kasper, S.L. Hauser, D.L. Longo, J. Loscalzo: Harrison's Manual of Medicine, Twentieth Edition. Copyright © McGraw-Hill Education. All rights reserved.

# 1-2) Upper – Korean Guidelines

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## REVIEW

*Gut and Liver, Vol. 14, No. 5, September 2020, pp. 560-570*

## Guidelines for Nonvariceal Upper Gastrointestinal Bleeding

### BODY

**Recommendation 1.** Risk stratification with an appropriate scoring system prior to upper gastrointestinal endoscopy in patients with NVUGIB can predict clinical outcomes.

Grade of recommendation: strong for recommendation

Level of evidence: high quality of evidence

**Recommendation 2.** Intravenous administration of PPIs prior to upper gastrointestinal endoscopy is recommended in patients with suspected NVUGIB.

Grade of recommendation: strong for recommendation

Level of evidence: high quality of evidence

# 1-3) Upper – American College of Gastroenterology

## ACG Clinical Guideline: Upper Gastrointestinal and Ulcer Bleeding

CME

**Table 1.** List of guideline statements with strength of recommendation and quality of evidence

### Risk stratification

1. We suggest that patients presenting to the emergency department with upper gastrointestinal bleeding (UGIB) who are classified as very low risk, defined as a risk assessment score with  $\leq 1\%$  false negative rate for the outcome of hospital-based intervention or death (e.g., Glasgow-Blatchford score = 0–1), be discharged with outpatient follow-up rather than admitted to hospital (conditional recommendation, very-low-quality evidence).

### Red blood cell transfusion

2. We suggest a restrictive policy of red blood cell transfusion with a threshold for transfusion at a hemoglobin of 7 g/dL for patients with UGIB (conditional recommendation, low-quality evidence).

### Pre-endoscopic medical therapy

#### *Prokinetic therapy with erythromycin*

3. We suggest an infusion of erythromycin before endoscopy in patients with UGIB (conditional recommendation, very-low-quality evidence).

#### *Proton pump inhibitor (PPI) therapy*

4. We could not reach a recommendation for or against pre-endoscopic PPI therapy for patients with UGIB.

### Endoscopy for UGIB

#### *Timing of endoscopy*

5. We suggest that patients admitted to or under observation in hospital for UGIB undergo endoscopy within 24 hr of presentation (conditional recommendation, very-low-quality evidence).

#### *Need for endoscopic hemostatic therapy for ulcers with active bleeding or nonbleeding visible vessels*

6. We recommend endoscopic therapy in patients with UGIB due to ulcers with active spurting, active oozing, and nonbleeding visible vessels (strong recommendation, moderate-quality evidence).

#### *Need for endoscopic hemostatic therapy for ulcers with adherent clot*

7. We could not reach a recommendation for or against endoscopic therapy in patients with UGIB due to ulcers with adherent clot resistant to vigorous irrigation.

#### *Choice of endoscopic hemostatic therapy for bleeding ulcers*

8. We recommend endoscopic hemostatic therapy with bipolar electrocoagulation, heater probe, or injection of absolute ethanol for patients with UGIB due to ulcers (strong recommendation, moderate-quality evidence).
9. We suggest endoscopic hemostatic therapy with clips, argon plasma coagulation, or soft monopolar electrocoagulation for patients with UGIB due to ulcers (conditional recommendation, very-low- to low-quality evidence).
10. We recommend that epinephrine injection not be used alone for patients with UGIB due to ulcers but rather in combination with another hemostatic modality (strong recommendation, very-low- to moderate-quality evidence).
11. We suggest endoscopic hemostatic therapy with hemostatic powder spray TC-325 for patients with actively bleeding ulcers (conditional recommendation, very-low-quality evidence).
12. We suggest over-the-scope clips as a hemostatic therapy for patients who develop recurrent bleeding due to ulcers after previous successful endoscopic hemostasis (conditional recommendation, low-quality evidence).

### Antisecretory therapy after endoscopic hemostatic therapy for bleeding ulcers

13. We recommend high-dose PPI therapy given continuously or intermittently for 3 d after successful endoscopic hemostatic therapy of a bleeding ulcer (strong recommendation, moderate- to high-quality evidence).
14. We suggest that high-risk patients with UGIB due to ulcers who received endoscopic hemostatic therapy followed by short-term high-dose PPI therapy in hospital continue on twice-daily PPI therapy until 2 wk after index endoscopy (conditional recommendation, low-quality evidence).

### Recurrent ulcer bleeding after successful endoscopic hemostatic therapy

15. We suggest that patients with recurrent bleeding after endoscopic therapy for a bleeding ulcer undergo repeat endoscopy and endoscopic therapy rather than undergo surgery or transcatheter arterial embolization (conditional recommendation, low-quality evidence for comparison with surgery, very-low-quality evidence for comparison with transcatheter arterial embolization)

### Failure of endoscopic hemostatic therapy for bleeding ulcers

16. We suggest patients with bleeding ulcers who have failed endoscopic therapy next be treated with transcatheter arterial embolization (conditional recommendation, very-low-quality evidence).



# 1-3) Upper – American College of Gastroenterology

1. We suggest that patients presenting to the emergency department with upper gastrointestinal bleeding (UGIB) who are classified as very low risk, defined as a risk assessment score with  $\leq 1\%$  false negative rate for the outcome of hospital-based intervention or death (e.g., **Glasgow-Blatchford score = 0–1**), be **discharged with outpatient follow-up** rather than admitted to hospital (conditional recommendation, very-low-quality evidence).

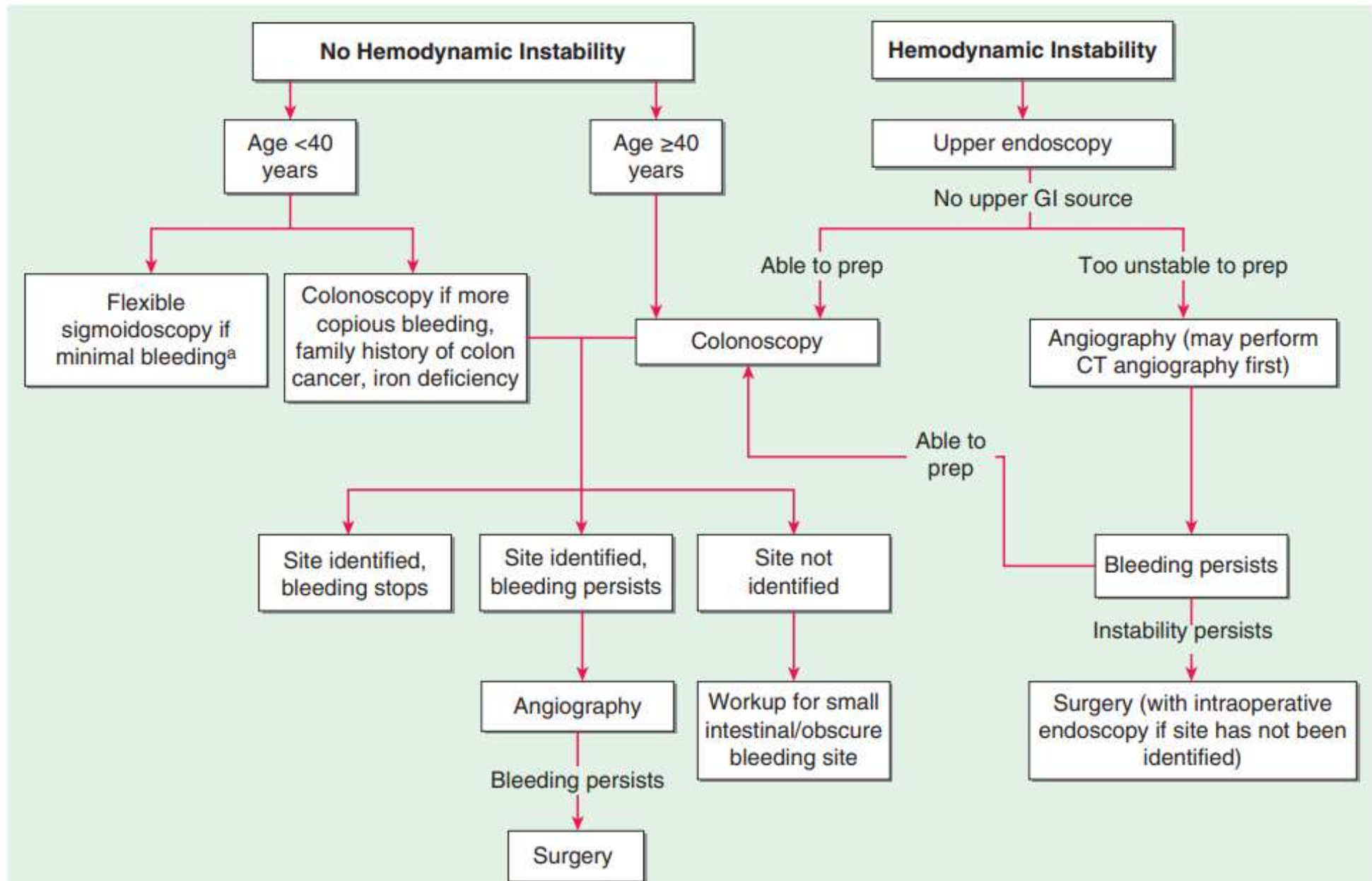
	Intervention or Death (N=100)	No Intervention or Death (N=150)
Glasgow-Blatchford Score >1	99 people with score >1 require intervention or die True Positive	100 people with score >1 do not require intervention or die False Positive
Glasgow-Blatchford Score 0-1	1 person with score 0-1 requires intervention or dies False Negative	50 people with score 0-1 do not require intervention or die True Negative
	<b>Sensitivity = 99/100 (99%)</b> The proportion of people who require intervention or die who have score >1	<b>Specificity = 50/150 (33%)</b> The proportion of people who do not require intervention or die who have score 0-1

**Figure 1.** Two-by-two table to determine sensitivity and specificity for hypothetical population of 250 patients presenting with upper gastrointestinal bleeding using Glasgow-Blatchford score cutoff of 1. The upper row includes patients with scores > 1, and the lower row includes those with scores of 0–1 (defined as very low risk). The left column shows the 100 patients who will require hospital-based intervention or die, with sensitivity calculated by the formula true positives divided by total number requiring intervention or dying (99/100 = 99%). The right column shows the 150 patients who will not require hospital-based intervention or die, with specificity calculated by the formula true negatives divided by total number not requiring intervention or dying (50/150 = 33%).

**Table 2.** Glasgow-Blatchford score

Risk factors at admission	Factor score
Blood urea nitrogen (mg/dL)	
18.2 to <22.4	2
22.4 to <28.0	3
28.0 to <70.0	4
$\geq 70.0$	6
Hemoglobin (g/dL)	
12.0 to <13.0 (men); 10.0 to <12.0 (women)	1
10.0 to <12.0 (men)	3
8.0 to <10.0 (men)	6
Systolic blood pressure (mm Hg)	
100 to <120	1
90 to <100	2
80 to <90	3
Heart rate (beats per minute)	
100 to <120	1
120 to <140	1
140 to <160	2
160 to <180	2
180 to <200	2

## 2-1) Lower - Traditional Harrison's algorithm

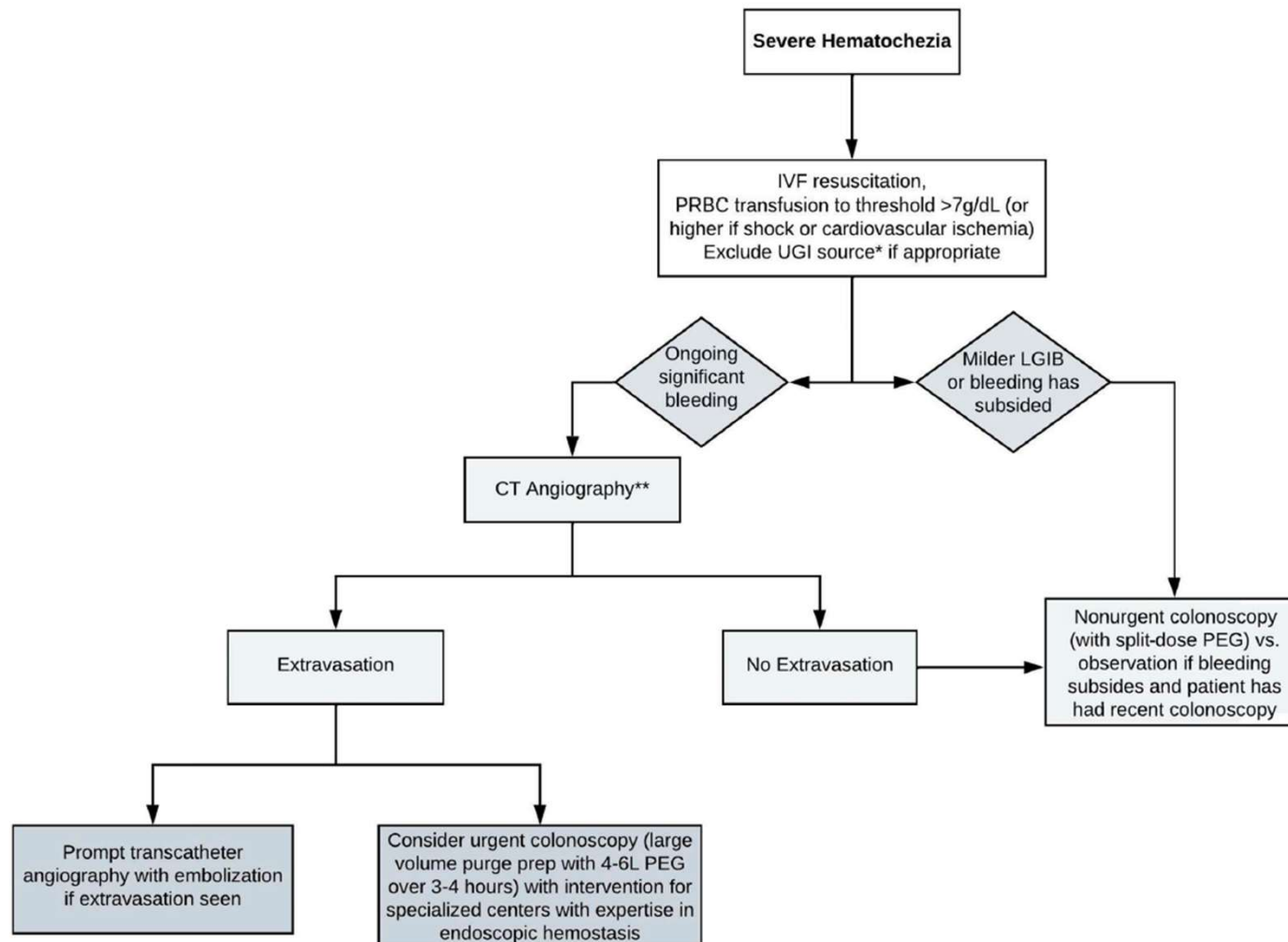


<sup>a</sup>Some suggest colonoscopy for any degree of rectal bleeding in patients <40 years as well.

## 2-2) Lower – US (2023) guideline

CME

# Management of Patients With Acute Lower Gastrointestinal Bleeding: An Updated ACG Guideline



## 2-2) Lower – Europe (2020) guideline

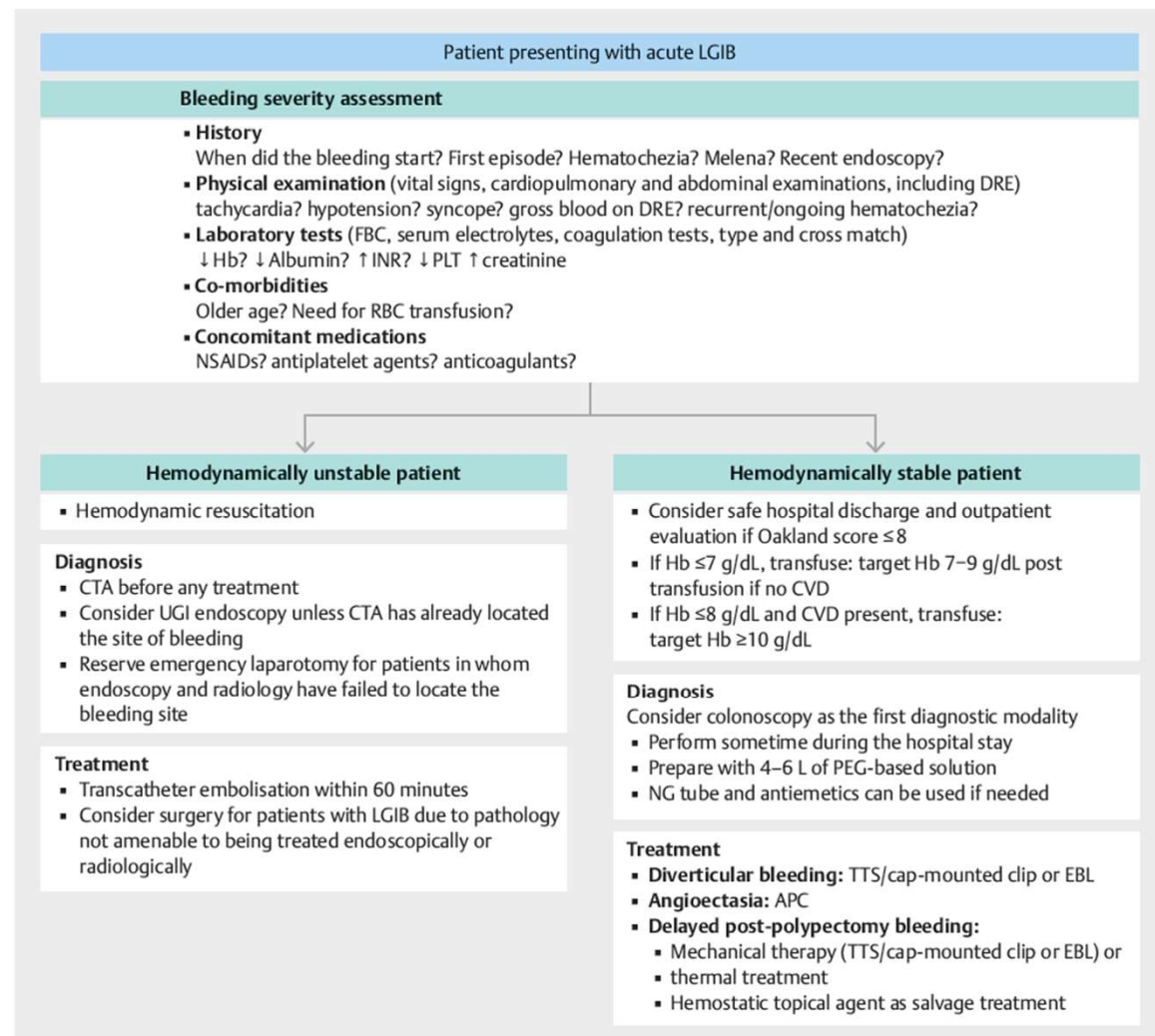
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### **Diagnosis and management of acute lower gastrointestinal bleeding: European Society of Gastrointestinal Endoscopy (ESGE) Guideline**



# 2-2) Lower – Europe (2020) guideline

## Diagnosis and management of acute lower gastrointestinal bleeding: European Society of Gastrointestinal Endoscopy (ESGE) Guideline



**Bleeding severity assessment**

- **History**  
When did the bleeding start? First episode? Hematochezia? Melena? Recent endoscopy?
- **Physical examination** (vital signs, cardiopulmonary and abdominal examinations, including DRE)  
tachycardia? hypotension? syncope? gross blood on DRE? recurrent/ongoing hematochezia?
- **Laboratory tests** (FBC, serum electrolytes, coagulation tests, type and cross match)  
↓ Hb? ↓ Albumin? ↑ INR? ↓ PLT ↑ creatinine
- **Co-morbidities**  
Older age? Need for RBC transfusion?
- **Concomitant medications**  
NSAIDs? antiplatelet agents? anticoagulants?

**Hemodynamically unstable patient**

- Hemodynamic resuscitation

**Diagnosis**

- CTA before any treatment
- Consider UGI endoscopy unless CTA has already identified the site of bleeding
- Reserve emergency laparotomy for patients if endoscopy and radiology have failed to locate bleeding site

**Treatment**

- Transcatheter embolisation within 60 minutes
- Consider surgery for patients with LGIB due to pathology not amenable to being treated endoscopically or radiologically

**Hemodynamically stable patient**

- **Consider safe hospital discharge and outpatient evaluation if Oakland score  $\leq 8$**

1. We suggest using risk stratification tools (e.g., Oakland score  $\leq 8$ ) to identify low-risk patients with LGIB who are appropriate for early discharge and outpatient diagnostic evaluation. Risk scores should be used to supplement but not replace clinician judgment. (Conditional recommendation, low-quality evidence)

- NG tube and antiemetics can be used if needed

**Treatment**

- **Diverticular bleeding:** TTS/cap-mounted clip or EBL
- **Angioectasia:** APC
- **Delayed post-polypectomy bleeding:**
  - Mechanical therapy (TTS/cap-mounted clip or EBL) or
  - thermal treatment
  - Hemostatic topical agent as salvage treatment

## 2-3) Lower – Risk stratification (Oakland score)

► **Table 3** The Oakland score for predicting the safe discharge of patients presenting with acute lower gastrointestinal bleeding (LGIB).

Variable	Score
Age, years	
▪ <40	0
▪ 40–69	1
▪ >70	2
Sex	
▪ Female	0
▪ Male	1
Previous LGIB admission	
▪ No	0
▪ Yes	1
Digital rectal examination findings	
▪ No blood	0
▪ Blood	1

Heart rate, bpm	
▪ <70	0
▪ 70–89	1
▪ 90–109	2
▪ >110	3
Systolic blood pressure, mmHg	
▪ 50–89	5
▪ 90–119	4
▪ 120–129	3
▪ 130–159	2
▪ >160	0
Hemoglobin, g/dL	
▪ 36–69	22
▪ 70–89	17
▪ 90–109	13
▪ 110–129	8
▪ 130–159	4
▪ >160	0



## 2-4) Lower – Key point in Europe and US guideline

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8 ESGE recommends temporarily withholding direct oral anticoagulants at presentation in patients with major lower gastrointestinal bleeding.

Strong recommendation, low quality evidence.

9 ESGE does not recommend withholding aspirin in patients taking low dose aspirin for secondary cardiovascular prevention. If withheld, low dose aspirin should be resumed, preferably within 5 days or even earlier if hemostasis is achieved or there is no further evidence of bleeding.

Strong recommendation, moderate quality evidence.

10 ESGE does not recommend routinely discontinuing dual antiplatelet therapy (low dose aspirin and a P2Y12 receptor antagonist) before cardiology consultation. Continuation of the aspirin is recommended, whereas the P2Y12 receptor antagonist can be continued or temporarily interrupted according to the severity of bleeding and the ischemic risk. If interrupted, the P2Y12 receptor antagonist should be restarted within 5 days, if still indicated.

Strong recommendation, low quality evidence.

5. We recommend against the administration of antifibrinolytic agents such as tranexamic acid in LGIB. (Strong recommendation, moderate quality evidence)

11a. We recommend discontinuing nonaspirin NSAIDs after hospitalization for diverticular hemorrhage. (Strong recommendation, low-quality evidence)

11b. We suggest discontinuing aspirin for primary cardiovascular prevention after hospitalization for diverticular hemorrhage given the risks of recurrent diverticular hemorrhage. (Conditional recommendation, low-quality evidence)

11c. We suggest continuing aspirin after hospitalization for diverticular hemorrhage for patients with an established history of cardiovascular disease given the benefits of reducing future ischemic events. (Conditional recommendation, low-quality evidence)

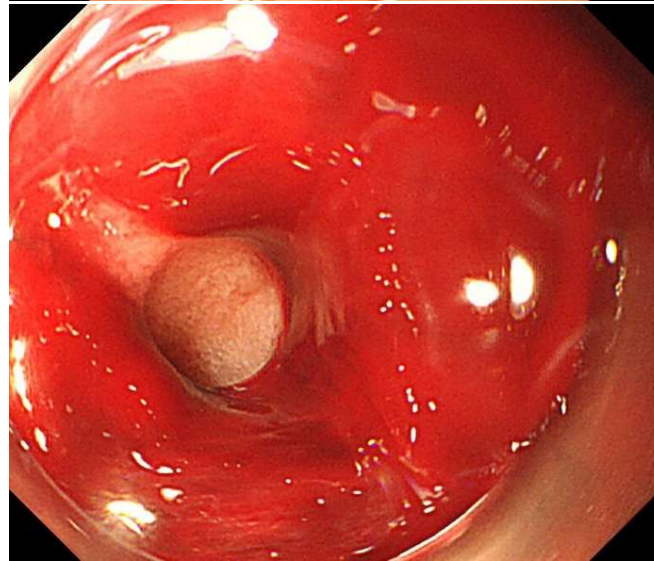
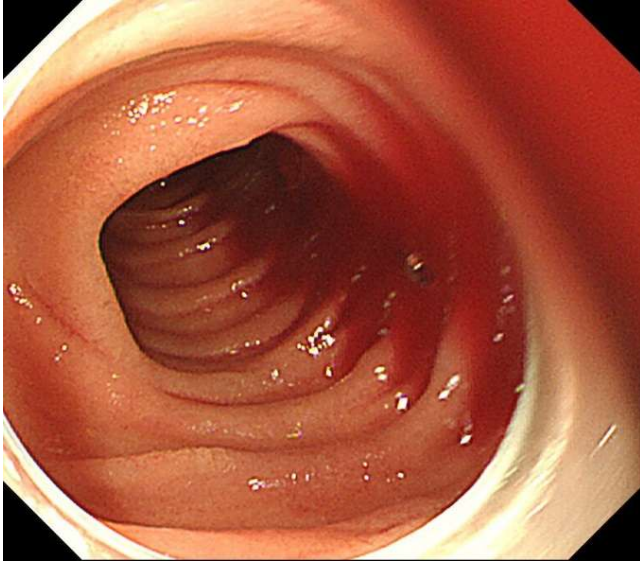
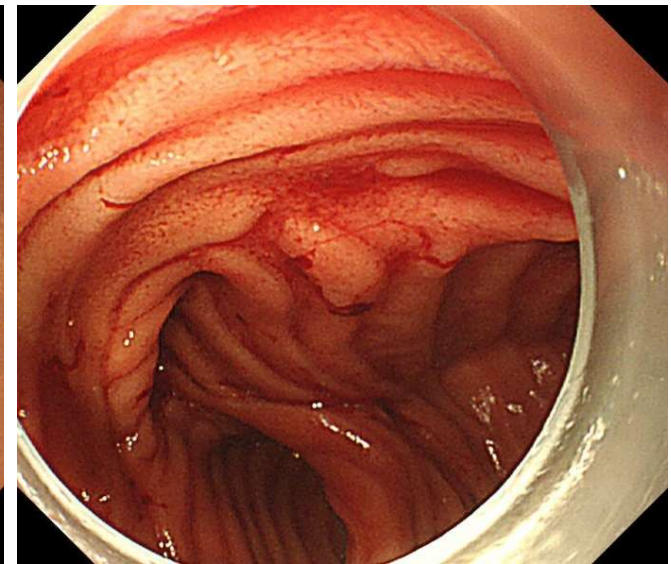
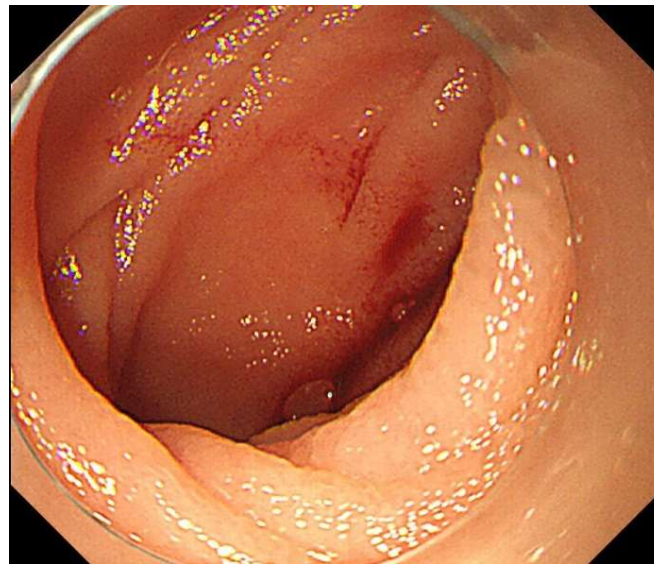
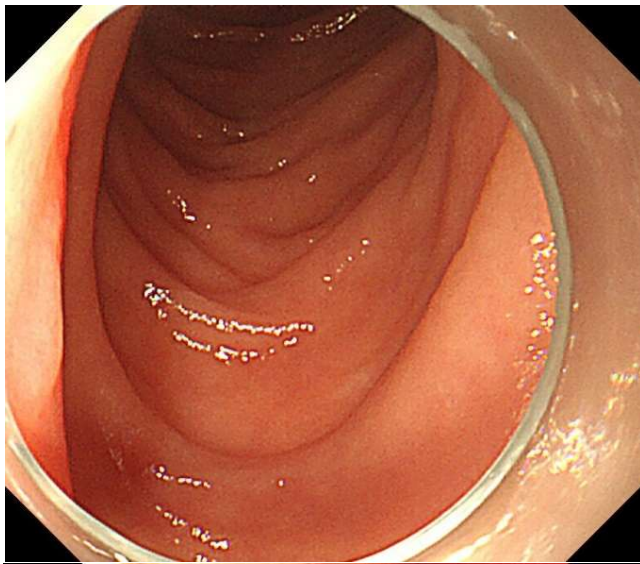
11d. We recommend that providers re-evaluate the risks vs benefits of continuing nonaspirin antiplatelets such as P2Y12 receptor antagonists in a multidisciplinary setting after hospitalization for diverticular hemorrhage given the demonstrated risks of recurrent diverticular hemorrhage. (Strong recommendation, low-quality evidence)

## **4. Actual clinical practice & cases**



# Surgical history is important

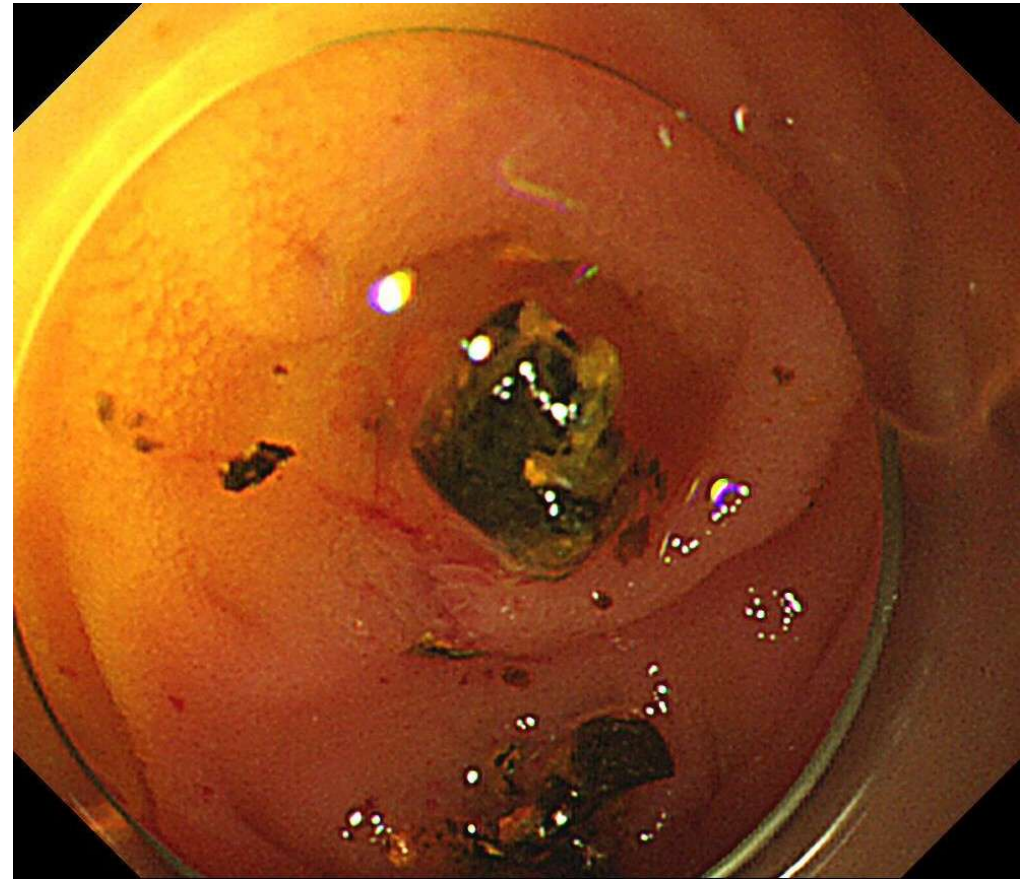
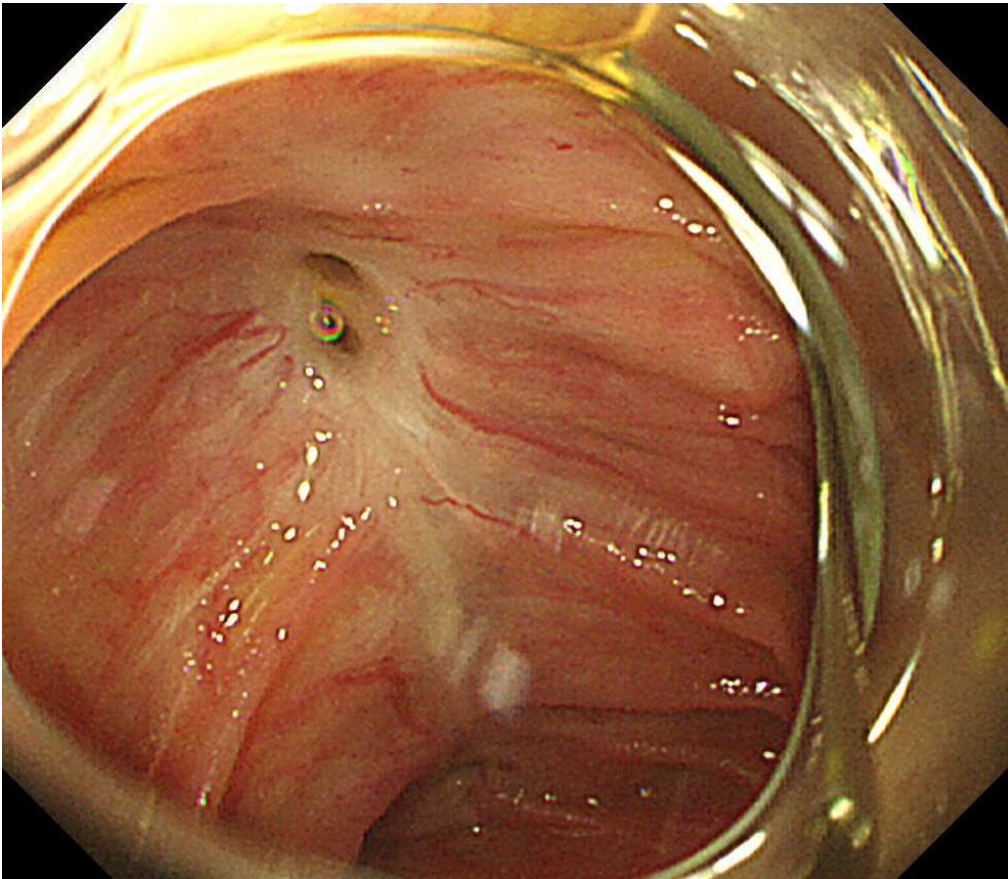
**Recurrent melena and Hb drop, partial hepatectomy due to CCC**





## Surgical history is important

**Recurrent melena and Hb drop, partial hepatectomy due to CCC**





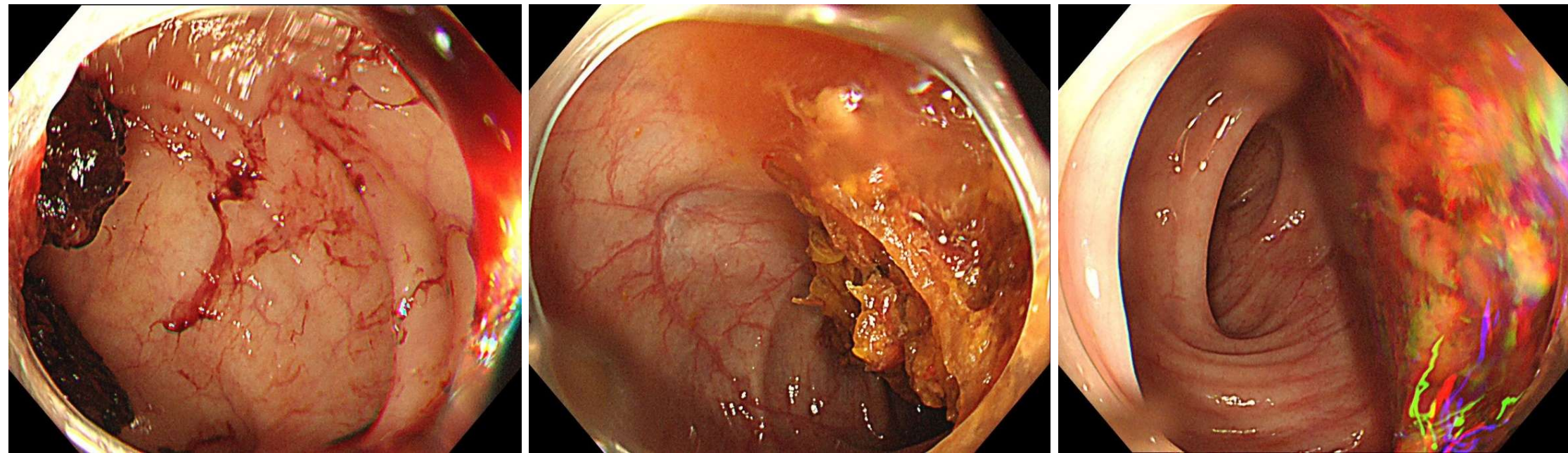
## Surgical history is important

**Hematochezia and anemia, s/p PPPD due to pancreatic cancer**





## Infection may cause hematochezia

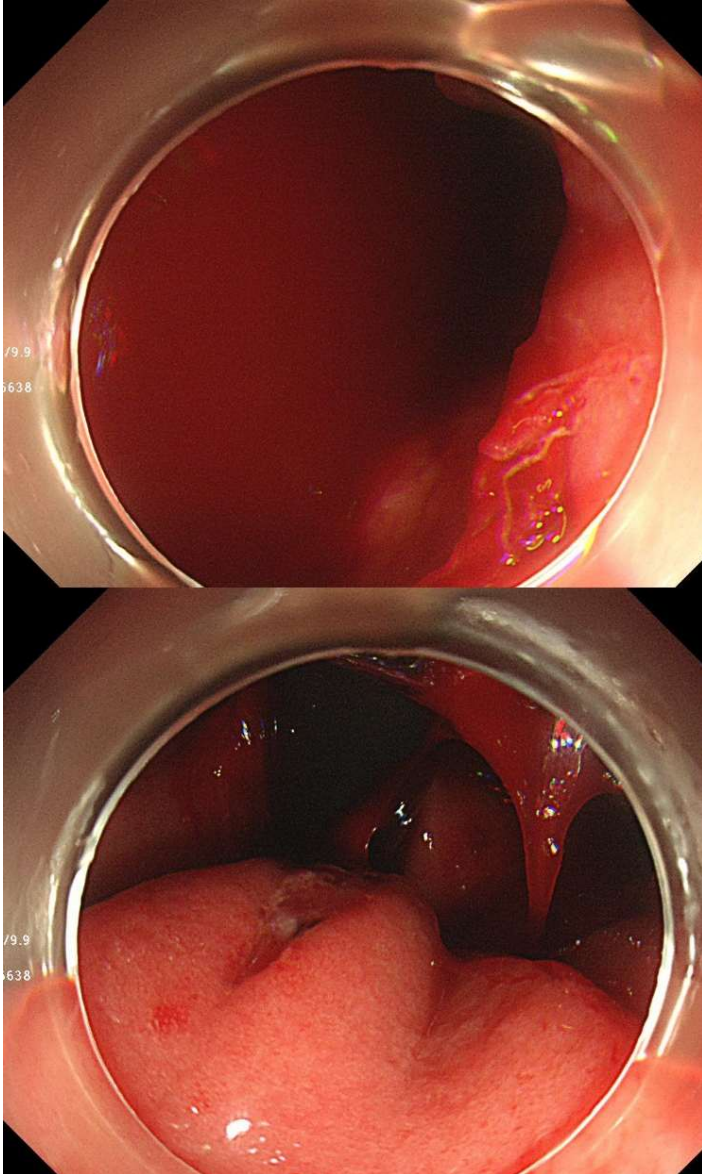


GIMD C.difficile Toxin Gene (Xpert) <Stool> ()

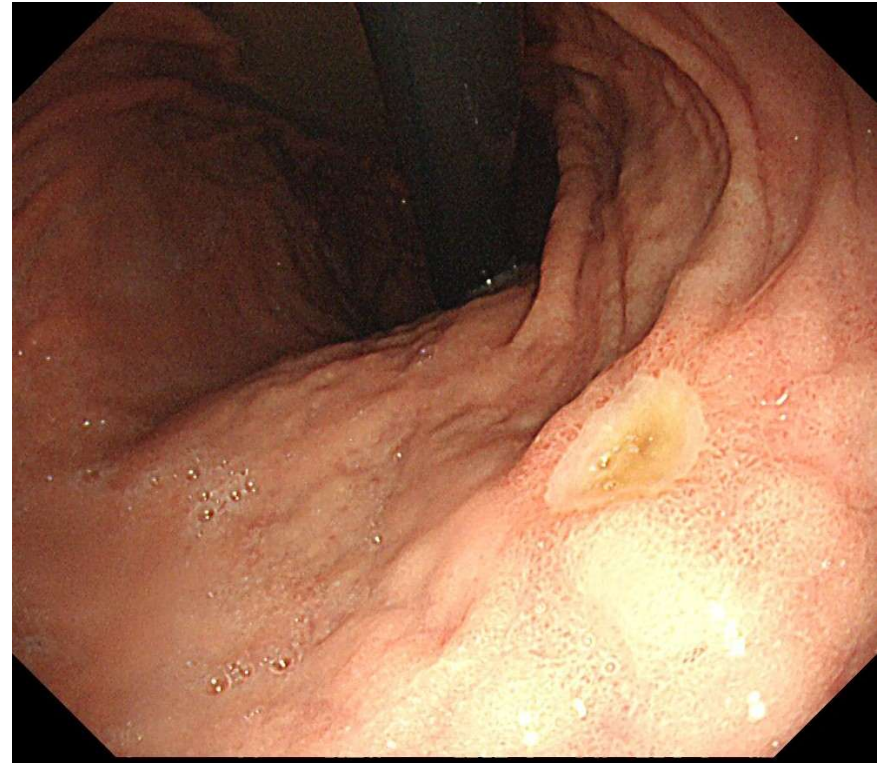
Positive

# Fitfalls of hemoglobin

Hb – 14.0g/dL



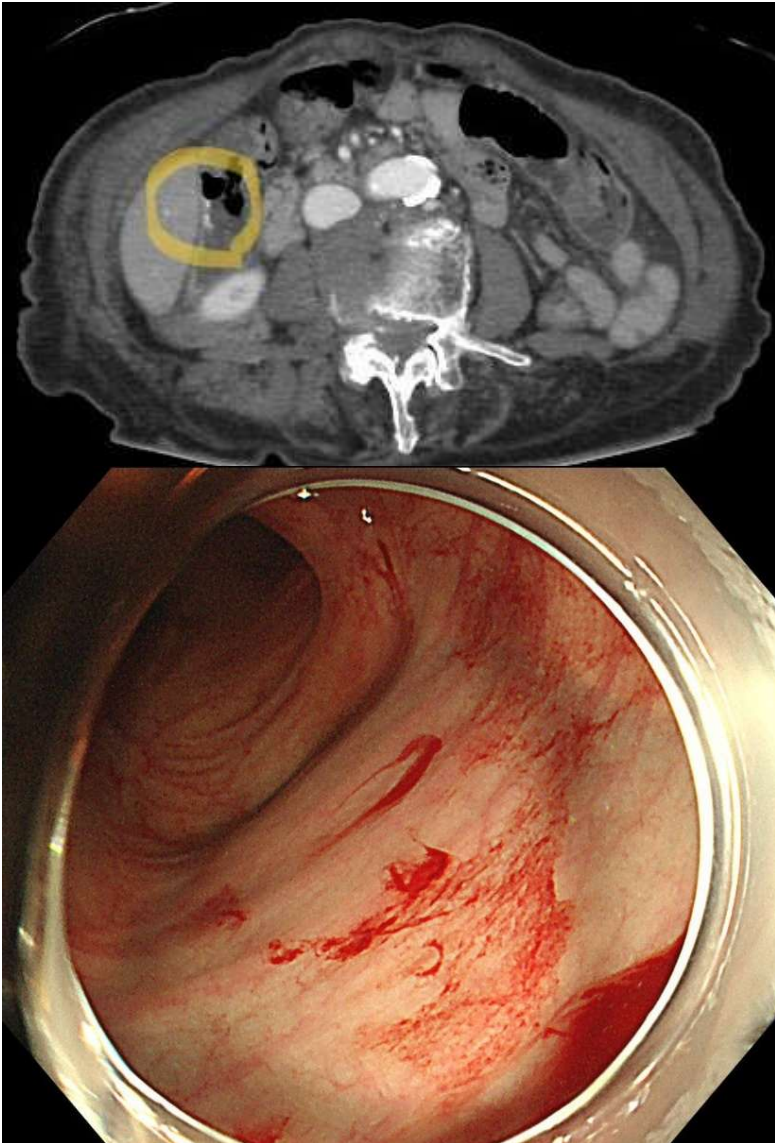
Hb – 8.1g/dL



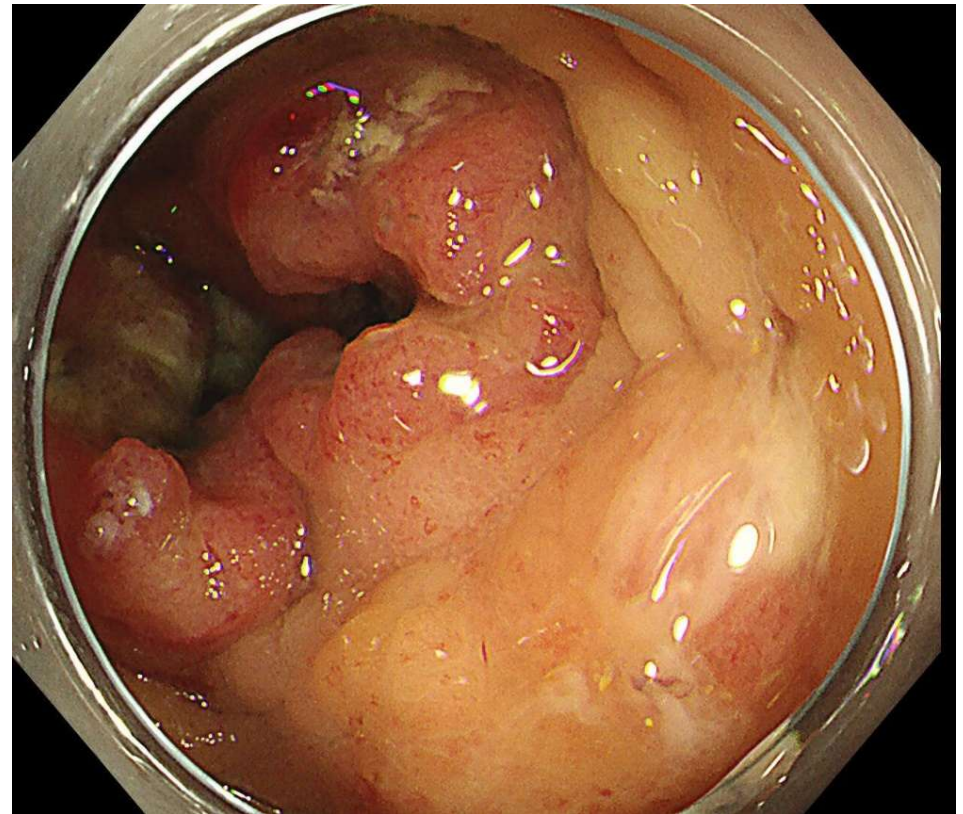


# Fitfalls of hemoglobin

Hb – 11.5g/dL



Hb – 6.3g/dL



# 5. Summary

## Summary

1. Careful history taking is **most important** in determining whether or not the bleeding is significant.
2. Risk stratification may reduce **unnecessary** emergency room visits
3. Discontinuation of antithrombotic agents should be done **cautiously**.
4. Since gastrointestinal bleeding occurs for a **variety of reasons**, its treatment varies from case to case.