

NONVARICEAL UPPER GASTROINTESTINAL HEMORRHAGE

INTRODUCTION

This ESGE Guideline focuses on the preendoscopic, endoscopic, and post-endoscopic management of patients presenting with acute nonvariceal upper gastrointestinal hemorrhage (NVUGIH), specifically peptic ulcer hemorrhage.

Pre-endoscopy management: Initial patient evaluation and hemodynamic resuscitation

RED BLOOD CELL (RBC) TRANSFUSION STRATEGY

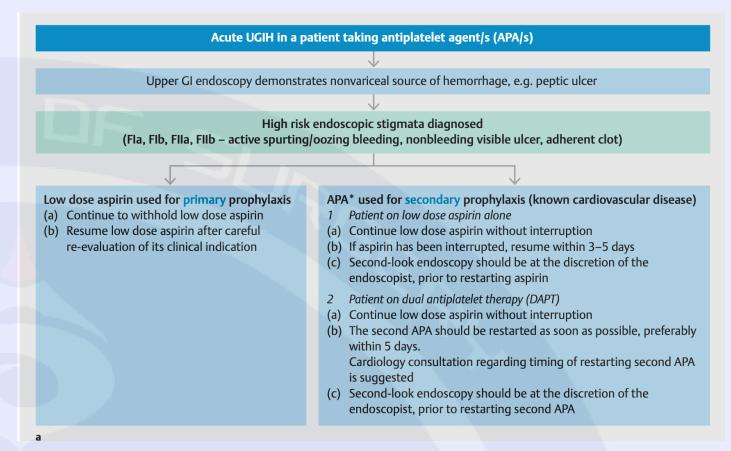
- Hemodynamically stable patients with no history of cardiovascular disease
 - hemoglobin threshold of ≤ 7 g/dL prompting RBC transfusion.
 - A post-transfusion target hemoglobin concentration of 7–9 g/dL is desired
- History of acute or chronic cardiovascular disease
 - a hemoglobin threshold of ≤8g/dL prompting RBC transfusion.
 - A post-transfusion target hemoglobin concentration of ≥ 10 g/dL is desired.

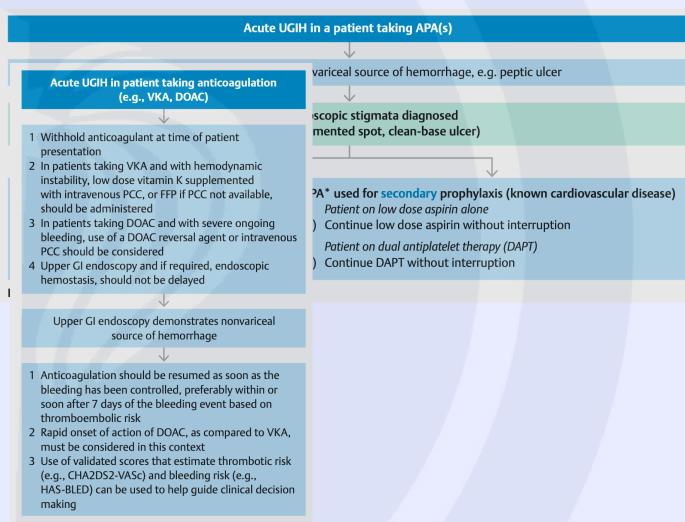
PATIENT RISK STRATIFICATION

| Risk factors at presentation | Threshold | Score |
|--------------------------------|--|------------------|
| 3lood urea nitrogen (mmol/I) | 6.5–7.9 8.0–9.9 10.0–24.9 ≥25.0 | 2 3 4 6 |
| Hemoglobin for men (g/l) | 120–130 100–119 <100 | 1 3 6 |
| Hemoglobin for women (g/l) | 100–120 <100 | 1 6 |
| Systolic blood pressure (mmHg) | 100–109 90–99 <90 | 1 2 3 |
| Heart rate (bpm) | >100 | 1 |
| Vielena | Present | 1 |
| Syncope | Present | 2 |
| Hepatic disease | Present | 2 |
| Cardiac failure | Present | 2 |

Total score (0–23). Patients with scores >0 are considered to be at high risk. Permission obtained from Elsevier Ltd © Blatchford, O. et al. Lancet 356, 1318–1321 (2000).

PRE-ENDOSCOPY MANAGEMENT OF ANTITHROMBOTIC AGENTS (ANTIPLATELET AGENTS AND ANTICOAGULANTS)





PRE-ENDOSCOPY PROTON PUMP INHIBITOR (PPI) THERAPY

 high dose intravenous proton pump inhibitor (PPI) therapy be considered in pa- tients presenting with acute UGIH, to downstage endo- scopic stigmata and thereby reduce the need for endo- scopic therapy;

SOMATOSTATIN AND SOMATOSTATIN ANALOGUES

 not recommend the use of somatostatin, or its analogue octreotide, in patients with NVUGIH.

PROKINETIC MEDICATIONS

 pre-endoscopy administration of intravenous erythromycin in selected patients with clinically severe or ongoing active UGIH.

ENDOSCOPIC MANAGEMENT

| la | Ib | lla | IIb | IIC | III |
|-----------------------------|--------------|-----------------------------|---------------|---------------------------|---------------------|
| Spurting bleed | Oozing bleed | Non-bleeding visible vessel | Adherent clot | Flat spot in ulcer crater | Clean base ulcer |
| | | | | | |
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Performance of upper GI endoscopy¹

High risk endoscopic stigmata Fla (active spurting, pulsatile arterial bleeding) Flb (active oozing) FIIa (nonbleeding visible vessel)

Perform endoscopic hemostasis

Fla and Flb stigmata Combination therapy using dilute epinephrine injection + a second hemostasis modality (thermal², mechanical or sclerosant injection³)

Flla stigmata Thermal², mechanical, or sclerosant injection³ as monotherapy or in combination with dilute epinephrine injection

- High dose PPI (intravenous bolus + continuous infusion or minimum twice-daily intravenous bolus dosing for 72 hours or oral dosing)
- May start clear liquids soon after endoscopy
- Test for Helicobacter pylori at index endoscopy, treat if positive; document *H. pylori* eradication
- If negative H. pylori test at index endoscopy, repeat testing within 4 weeks following the acute bleeding episode to confirm initial test was true negative

If clinical evidence of rebleeding, repeat endoscopy with endoscopic hemostasis if indicated; If endoscopic hemostasis still unsuccessful, refer for TAE if locally available, otherwise refer for surgery

Performance of upper GI endoscopy¹

Consider performing clot removal followed by endoscopic hemostasis of underlying high risk stigmata4

Medical management with high dose PPI (intravenous bolus + continuous infusion for 72 hours or minimum twice-daily intravenous bolus dosing for 72 hours or oral dosing)

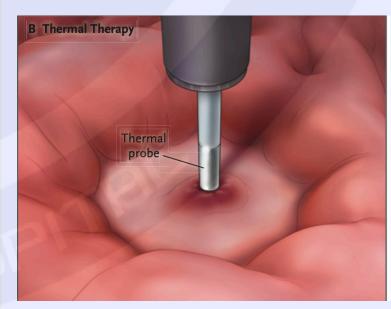
If clot removal/endoscopic hemostasis performed:

- of clot followed by clot removal using cold polyp snare guillotine technique
- removal, apply endoscopic hemostasis as described for Fla, Flb, Flla stigmata
- infusion or minimum twice-daily intravenous bolus dosing for 72 hours or oral dosing)
- May start clear liquids soon after endoscopy
- eradication
- If negative *H. pylori* test at index endoscopy, repeat testing within 4 weeks following the acute bleeding episode to confirm initial test was true negative

If clinical evidence of rebleeding, repeat endoscopy with endoscopic hemostasis if indicated;

if locally available, otherwise refer for surgery

THERMAL THERAPY



INJECTION THERAPY

hollow-core needle (19-25 gauge)

and around the ulcer base)

Ulcer

base

Diluted epinephrine (1:10 000 or 1:20 000 with

Endoscopic injection is performed using needles

normal saline inject ted in 0.5-2-ml aliquots in

which consist of an outer sheath and an inner

Injection

Visible

vessel



Performance of upper GI endoscopy¹

Low risk stigmata FIIc (flat pigmented spot) FIII (clean base)

No endoscopic hemostasis required In select clinical settings, these patients may have expedited hospital discharge

- Start oral PPI
- Start regular diet
- Test for *H. pylori*, treat if positive; document *H. pylori* eradication
- If negative H. pylori test at index endoscopy, repeat testing within 4 weeks following the acute bleeding episode to confirm initial test was true negative

FIIb (adherent clot)

- Dilute epinephrine injection circumferential to base
- If underlying high risk stigmata identified after clot
- High dose PPI (intravenous bolus + continuous
- Test for H. pylori, treat if positive; document H. pylori

If endoscopic hemostasis still unsuccessful, refer for TAE

POST-ENDOSCOPY MANAGEMENT

PROTON PUMP INHIBITOR THERAPY

- High dose proton pump inhibitor (PPI) therapy for patients who
 - receive endoscopic hemostasis
 - for patients with FIIb ulcer stigmata
 (adherent clot) not treated endoscopically.
- (a) PPI therapy should be administered as an intravenous bolus followed by continuous infusion (e.g., 80mg then 8mg/hour) for 72 hours post endoscopy.
- (b) High dose PPI therapies given as intravenous bolus dosing (twice-daily) or in oral formulation (twice-daily) can be considered as alternative regimens.

MANAGEMENT OF RECURRENT BLEEDING

- Repeat upper endoscopy, including hemostasis if indicated.
- failure of this second attempt at endoscopic hemostasis
 - transcatheter angiographic embolization (TAE) should be considered.
 - Surgery is indicated when TAAE is not locally avail- able or after failed TAE.
- Recurrent peptic ulcer hemorrhage
 - Capmounted clip should be considered.

HELICOBACTER PYLORI

- Patients with NVUGIH secondary to peptic ulcer, investigation for the presence of Helicobacter pylori in the acute setting (at index endoscopy)
- initiation of appropriate antibiotic therapy when H. pylori is detected.
- Re-testing for H. pylori in those patients with a negative test at index endoscopy.