

# JAG clinical audit report

**Title: Acute upper gastrointestinal bleeding in over 16s: Management Audit**

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Aim(s) of Audit:

To review if GRHT manages Upper GI Bleeds (**UGIB**) are managed in accordance with **NICE recommendations for Acute upper Gastrointestinal bleeding in over 16s**.

NICE recommend patients should have the following diagnostic and therapeutic steps:

## **Risk Assessment**

- All patients should be given a Blatchford score at first assessment and the full Rockall score post endoscopy

## **Resuscitation and Initial management**

- Transfuse patients with massive bleeding with blood, platelets and clotting factors in line with local protocols for managing massive bleeding.
- Base decisions on blood transfusion on the full clinical picture, recognising that over-transfusion may be as damaging as under-transfusion
- Do not offer platelet transfusion to patients who are not actively bleeding and are haemodynamically stable.
- Offer platelet transfusion to patients who are actively bleeding and have a platelet count of less than  $50 \times 10^9$ /litre.
- Offer fresh frozen plasma to patients who are actively bleeding and have a prothrombin time (or international normalised ratio) or activated partial thromboplastin time greater than 1.5 times normal. If a patient's fibrinogen level remains less than 1.5 g/litre despite fresh frozen plasma use, offer cryoprecipitate as well.
- Offer prothrombin complex concentrate to patients who are taking warfarin and actively bleeding.
- Treat patients who are taking warfarin and whose upper gastrointestinal bleeding has stopped in line with local warfarin protocols.
- Do not use recombinant factor VIIa except when all other methods have failed.

## **Timing of endoscopy**

- Offer endoscopy to unstable patients with severe acute upper gastrointestinal bleeding immediately after resuscitation.
- Offer endoscopy within 24 hours of admission to all other patients with upper gastrointestinal bleeding.

- Units seeing more than 330 cases a year should offer daily endoscopy lists. Units seeing fewer than 330 cases a year should arrange their service according to local circumstances.

### **Management for non-variceal bleeding**

#### **Endoscopic treatment**

- Do not use adrenaline as monotherapy for the endoscopic treatment of non-variceal upper gastrointestinal bleeding
- For the endoscopic treatment of non-variceal upper gastrointestinal bleeding, use one of the following:
  - a mechanical method (for example, clips) with or without adrenaline
  - thermal coagulation with adrenaline
  - fibrin or thrombin with adrenaline.

#### **Proton pump inhibitors**

- Do not offer acid-suppression drugs (proton pump inhibitors or H<sub>2</sub>-receptor antagonists) before endoscopy to patients **with suspected non-variceal** upper gastrointestinal bleeding.
- Offer proton pump inhibitors to patients **with non-variceal** upper gastrointestinal bleeding and stigmata of recent haemorrhage shown at endoscopy.

#### **Treatment after first or failed endoscopic treatment**

- Consider a repeat endoscopy, with treatment as appropriate, for all patients at high risk of re-bleeding, particularly if there is doubt about adequate haemostasis at the first endoscopy.
- Offer a repeat endoscopy to patients who re-bleed with a view to further endoscopic treatment or emergency surgery.
- Offer interventional radiology to unstable patients who re-bleed after endoscopic treatment. Refer urgently for surgery if interventional radiology is not promptly available

#### **Management of variceal bleeding**

- Offer terlipressin to patients with suspected variceal bleeding at presentation. Stop treatment after definitive haemostasis has been achieved, or after 5 days, unless there is another indication for its use.
- At the time of publication (June 2012), terlipressin was indicated for the treatment of bleeding from oesophageal varices, with a maximum duration of treatment of 72 hours (3 days). Prescribers should consult the relevant summary of product characteristics. Informed consent for off-label use of terlipressin should be obtained and documented.
- Offer prophylactic antibiotic therapy at presentation to patients with suspected or confirmed variceal bleeding.

#### **Oesophageal varices**

- Use band ligation in patients with upper gastrointestinal bleeding from oesophageal varices.

- Consider transjugular intrahepatic portosystemic shunts (TIPS) if bleeding from oesophageal varices is not controlled by band ligation.

#### **Gastric varices**

- Offer endoscopic injection of N-butyl-2-cyanoacrylate to patients with upper gastrointestinal bleeding from gastric varices.
- Offer TIPS if bleeding from gastric varices is not controlled by endoscopic injection of N-butyl-2-cyanoacrylate.

#### **Control of bleeding and prevention of re-bleeding in patients on NSAIDs, aspirin or clopidogrel**

- Continue low-dose aspirin for secondary prevention of vascular events in patients with upper gastrointestinal bleeding in whom haemostasis has been achieved.
- Stop other non-steroidal anti-inflammatory drugs (including cyclooxygenase-2 [COX-2] inhibitors) during the acute phase in patients presenting with upper gastrointestinal bleeding.
- Discuss the risks and benefits of continuing clopidogrel (or any other thienopyridine antiplatelet agents) in patients with upper gastrointestinal bleeding with the appropriate specialist (for example, a cardiologist or a stroke specialist) and with the patient.

#### **Primary prophylaxis for acutely ill patients in critical care**

- Offer acid-suppression therapy (H<sub>2</sub>-receptor antagonists or proton pump inhibitors) for primary prevention of upper gastrointestinal bleeding in acutely ill patients admitted to critical care. If possible, use the oral form of the drug.
- As of September 2023, the use of proton pump inhibitors or H<sub>2</sub>-receptor antagonists other than ranitidine and cimetidine for this indication would be off label. Ranitidine is currently unavailable. See the [MHRA drug alert on ranitidine](#) for more information.
- Review the ongoing need for acid-suppression drugs for primary prevention of upper gastrointestinal bleeding in acutely ill patients when they recover or are discharged from critical care. Information and support for patients and carers

#### **Information and support for patients and carers**

Establish good communication between clinical staff and patients and their family and carers at the time of presentation, throughout their time in hospital and following discharge. This should include:

- giving verbal information that is recorded in medical records
- different members of clinical teams providing consistent information
- providing written information where appropriate
- ensuring patients and their families and carers receive consistent information.

#### **Method:**

- Data of patients presenting with Upper GI Bleeding sourced from Medilogik
- Patient notes obtained on Sunrise Electronic Patient Record, Medilogik, ICE.
- Data analysed against NICE standards.

-50 patients were used in the analysis/reporting. Estimated patient group for first quarter of 2023; 452 patients (unable to filter out 2ww non acute referrals).  
-Patient age range 21-91  
-30 male patients, 20 female patients

### **Analysis of data for patient group 01/01/2023-24/02/2023**

#### **Risk Assessment**

-28 patients were given a Blachford score at first presentation, 3 were recorded after first assessment, 16 patients had no score recorded  
-Non-compliance with Rockall score post endoscopy (although recently added to EMS)

#### **Resuscitation and Initial management**

-25 patients required transfusion pre-endoscopy, all were within local guidance  
-All patients were monitored post endoscopy with 13 requiring further transfusion

#### **Timing of endoscopy**

-8 unstable patients were scoped immediately after resuscitation  
-27 patients were scoped within 24 hours  
-15 patients were scoped after 24 hours

#### **Analysis of patients scoped after 24 hours**

Most common cause of delay was unstable patient/ clinical judgement (10 patients).

Patient 1 required scope within 24 hours which was requested but issues with referral reported in notes, patient 2 ate on ward so procedure delayed, patient 3 stable suspected variceal bleed scoped after 24 (unsure of reason for delay), patient 4 suspected variceal bleed referred for OGD but was non-compliant until outside of recommended timeframe, remaining patients deemed stable and discharged home for outpatient procedure due to in-patient bed crisis, patient 5 was delayed as no space on the in-patient list.

#### **Endoscopic intervention: non- variceal bleeding, variceal bleeding and treatment after first or failed endoscopic treatment**

-12 patients required therapeutic treatment (1 x banding) and all were compliant with NICE guidance  
-4 patient were appropriately re-scoped as at high risk of re-bleed  
-2 patients rescoped due to further episode of UGIB after first endoscopy. Neither patient required interventional radiology but were further investigated with CT CAP

#### **Proton pump inhibitors**

-9 patients were taking PPI as outpatient prior to admission  
-15 patients with suspected non-variceal UGIB were given PPI pre-endoscopy in non-critical care  
-2 patients with suspected non-variceal UGIB were given PPI pre-endoscopy in critical care

#### **PPI in varices management**

-9 patients taking PPI as outpatient prior to admission  
-1 suspected variceal bleed had PPI stopped and re-started post endoscopy  
-3 suspected variceal bleed were not given PPI prior to endoscopy

-48 patients were treated with PPI/dose increased post endoscopy, 2 patients did not require treatment

**Management of variceal bleeding**

- All patients (4/4) with suspected variceal bleeding were given Terlipressin and prophylactic antibiotics pre-endoscopy
- 2 patients required banding of oesophageal varices

**Control of bleeding and prevention of re-bleeding in patients on NSAIDs, aspirin or clopidogrel**

- 2/2 patients taking NSAIDs had their medication held
- 4/4 patients continued to take aspirin and clopidogrel as advised
- 3 patients had aspirin or Clopidogrel held appropriately and re-started post endoscopy
- 1 patient taking aspirin and clopidogrel continued medication pre/post endoscopy. Clinical decision for this was not documented in notes

**Information and support for patients and carers**

- All patients had good verbal communication recorded in medical notes
- 38 patients had written discharge information as recorded in medical notes
- 12 patients had no record that written discharge information was given. 2 patients excluded from data as were deceased before discharge (non-endoscopy related)

**Discussion:**

- Efficient inpatient service allowing patients to be scoped within recommended timeframe.
- Therapeutic treatment always given within recommended guidance.
- Excellent documentation of verbal communication with patients.
- Compliant in treatment of patients presenting with suspected variceal bleeding.
- Compliance with local protocol for patients requiring transfusions.
- Appropriate management of patients taking NSAIDs, Aspirin and Clopidogrel.
- Compliance with PPI treatment post endoscopy.
  
- 56% compliant with Blachford score at first assessment.
- Non compliance with Rockall score post endoscopy.
- 15 patients given PPI treatment prior to endoscopy in non-critical care (not in line with NICE guidance).

**Recommendation:**

- Mandatory documentation of Blachford score on first presentation. Governance to discuss implementation
- Post Rockall score to be added to EMS (already actioned)
- Medical team to be reminded of PPI guidance pre-endoscopy for with suspected non-variceal UGIB and importance of clear discharge information
  
- Governance to discuss future management of UGIB audit; lead, medical involvement, large volume of patients to be distributed for analysis among a team of staff for large pool of data, method to ease analysis of data eg patient form in notes, upload to EMS from patient notes to record data.

**Audit presentation:**

**Date:** 26/01/2024

**Venue:**

Actions	Person responsible for implementing actions (name & designation)	Date for completion

DRAFT

Project Presentation(s)/Publications:

Date

Venue/Publication

*(Please attach copies of your presentation/article(s) to this form if possible)*

Do you wish to re-audit this aspect of care?

YES

NO

Recommended Re-audit date: