Variable use of endoscopic haemostasis in the management of bleeding peptic ulcers

S Mahadeva, M Linch, M Hull

Background: Randomised controlled trials (RCTs) have shown that endoscopic haemostasis is beneficial for patients with a bleeding peptic ulcer. The relevance of such data to management outside of RCTs is unclear. Therefore we examined management of patients with a bleeding peptic ulcer in a UK teaching hospital.

Methods: All patients who underwent upper gastrointestinal (UGI) endoscopy for bleeding peptic ulcer between 1997 and 1999 were identified from an endoscopy database and the clinical records reviewed retrospectively.

Results: A total of 872 patients underwent UGI endoscopy for presumed acute UGI haemorrhage; 179 (21%) had an endoscopic diagnosis of bleeding peptic ulcer. Seventy nine patients had a peptic ulcer with stigmata of recent haemorrhage (SRH) but only 61 (77%) of these patients received endoscopic haemostasis (77% adrenaline, 23% combination therapy). Re-bleeding occurred in 24 patients with SRH in whom transfusion requirement was the sole predictor of re-bleeding. The re-bleeding rate among patients who received adrenaline was 25% (n=12), compared with 57% (n=8) in the combination group and 31% (n=4) in those who did not receive endoscopic haemostasis. Patients who received combination endoscopic haemostasis had an increased incidence of active bleeding (p=0.007) and an increased transfusion requirement (p=0.002). Eleven of 20 patients who re-bleed had repeat endoscopic haemostasis, with 45% eventually requiring surgery.

Conclusions: Results of endoscopic management of bleeding peptic ulcers in the unit studied differ markedly from those published by specialised centres. The data reported here suggest that increased standardisation of endoscopic haemostasis is required, especially in units with provision for emergency “out-of-hours” endoscopy, performed by several individuals of different grades.
or on a routine list, was also noted. The time that acute UGI endoscopy was performed; “out-of-hours” (between 6 pm and 8 am the next day), at the weekend
lesser curve gastric ulcer or posterior wall duodenal ulcer.
endoscopic haemostasis outcome, was defined as either high
ulcer location, which has been associated with suboptimal
either a clean ulcer base or a flat pigmented spot. Difficult
adrenaline, combination therapy) performed and if primary
haemorrhage (SRH; active bleeding, non-bleeding visible ves-
location of ulcer, presence and type of stigmata of recent
experience), or consultant (>5 years endoscopy experience),
senior registrar (3–5 years endoscopy experience), registrar (1–2 years
experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endoscopy experience), or consultant (>3 years endosco
received initial endoscopic haemostasis, 39 (64%) cases had active bleeding or a non-bleeding visible vessel at the time of endoscopy, compared with three (17%) in the group that did not receive any initial endoscopic haemostasis (fig 1). In contrast, 15 (83%) of the patients who did not receive initial endoscopic haemostasis had adherent clot compared with 22 (36%) patients who did receive initial endoscopic haemostasis (fig 1).

**Outcome of patients receiving different endoscopic haemostasis techniques**

Of the patients who received endoscopic haemostasis, 47 (77%) had adrenaline (1:10 000 dilution) injection only while 14 had combination endoscopic haemostasis (adrenaline plus heater probe, n=12; adrenaline plus thrombin injection, n=2) (fig 2). Patient characteristics, endoscopic findings, and blood transfusion requirements are documented in table 1. Complete haemostasis was achieved at the end of the endoscopic procedure in all but two patients (96%). Twelve of 47 patients (25%) re-bleed after adrenaline endoscopic haemostasis (fig 2). In the combination endoscopic haemostasis group, 8/14 patients (57%) re-bled after initial endoscopic haemostasis (fig 2). Univariate analysis showed that the transfusion requirement (p = 0.002) and the presence of active bleeding at initial endoscopy (p = 0.007) were significantly higher in the group of patients who received combination endoscopic haemostasis. A higher proportion of patients receiving combination endoscopic haemostasis had co-morbidity than the other two groups (table 2) but this did not reach statistical significance. Although differences in the grade of endoscopist between the three groups of patients with SRH were not statistically significant, combination endoscopic haemostasis was used more by senior registrars compared with other grades (table 1). Logistic regression analysis revealed that transfusion requirement was the only independent predictor of re-bleeding (p = 0.002). In the group of patients with SRH who did not receive initial endoscopic haemostasis, 4/18 (31%) had re-bleeding after initial diagnostic endoscopy. Only 2/84 patients who were reported to have no SRH re-bleed (which was managed surgically with one mortality).

**Management of re-bleeding after initial endoscopy**

Among the patients who re-bleed after receiving adrenaline endoscopic haemostasis, four patients had repeat endoscopic haemostasis (all combination), two died (one immediate death and the other did not have further intervention due to severe cerebrovascular disease), and six underwent surgery immediately (fig 2). Of those that re-bleed after combination endoscopic haemostasis, seven had repeat endoscopic haemostasis (all combination) and one patient had surgery immediately. Five of 11 (45%) patients who had repeat endoscopic haemostasis required salvage surgery for continued bleeding (fig 2). Nine of 11 (81%) patients who received repeat endoscopic haemostasis had co-morbidity compared with 3/7 (43%) who underwent surgery immediately for re-bleeding. The group of patients in whom repeat endoscopic haemostasis failed had 100% co-morbidity and a significantly higher mean transfusion requirement compared with those who had successful repeat endoscopic haemostasis for re-bleeding or immediate surgery (table 3). Two of 11 (18%) patients who had repeat endoscopic haemostasis suffered early mortality compared with 1/7 (14%) patients who had immediate surgery. Surgery for re-bleeding or secondary bleeding after failed endoscopic haemostasis consisted of under-running of the ulcer (n=6), under-running of the ulcer with pyloroplasty and vagotomy (n=3), excision of ulcer and vessel under-running (n=1), gastroenterostomy (n=1), and partial gastrectomy (n=1). Patients who had re-bleeding from an ulcer in a “difficult” location were more likely to be referred for immediate surgery (57%) than those whose ulcer was elsewhere (20%). The length of hospital stay after acute UGI bleeding was significantly longer in those patients who had repeat endoscopic haemostasis but eventually required surgery (table 3). However a similar duration of hospital stay was noted in those who underwent surgery or successful repeat endoscopic haemostasis for re-bleeding (table 3).

**Outcome of patients who did not receive endoscopic haemostasis**

In the group of patients with SRH who did not receive initial endoscopic haemostasis the outcome of the four patients that re-bleed were as follows: one suffered early mortality, two received endoscopic haemostasis (one of which required salvage surgery for continued bleeding but did not survive), and a further patient had surgery immediately.

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**Table 2** Co-morbidity of patients with SRH visible at initial endoscopy; values are number (%)

<table>
<thead>
<tr>
<th>Co-morbidity</th>
<th>Endoscopic haemostasis types</th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>No endoscopic haemostasis</td>
<td>Adrenaline</td>
<td>Combination</td>
</tr>
<tr>
<td>Absent</td>
<td>8 (44)</td>
<td>26 (55)</td>
<td>4 (29)</td>
</tr>
<tr>
<td>Present</td>
<td>10 (56)</td>
<td>21 (45)</td>
<td>10 (71)</td>
</tr>
<tr>
<td>Cardiovascular</td>
<td>3 (17)</td>
<td>9 (19)</td>
<td>4 (29)</td>
</tr>
<tr>
<td>Respiratory</td>
<td>1 (6)</td>
<td>2 (2)</td>
<td>2 (14)</td>
</tr>
<tr>
<td>Cerebrovascular disease</td>
<td>3 (16)</td>
<td>1 (6)</td>
<td>0</td>
</tr>
<tr>
<td>Cancer</td>
<td>2 (11)</td>
<td>3 (6)</td>
<td>2 (14)</td>
</tr>
<tr>
<td>Renal failure</td>
<td>1 (6)</td>
<td>0</td>
<td>2 (14)</td>
</tr>
<tr>
<td>Other</td>
<td>0</td>
<td>2 (5)</td>
<td>0</td>
</tr>
</tbody>
</table>

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**Figure 2** Outcome of patients who received endoscopic haemostasis.
Mortality of patients with bleeding peptic ulcer
The 30 day mortality rate for all patients with bleeding peptic ulcers was 11.6%. Subgroup analysis revealed 30 day mortality rates of 6% in the group of patients who received no SRH, 16.4% in the SRH group that received initial endoscopic haemostasis, and 22.2% in the SRH group who did not receive initial endoscopic haemostasis.

DISCUSSION
Use of endoscopic haemostasis for peptic ulcers with SRH
This retrospective study of the endoscopic management of bleeding peptic ulcers in a large teaching hospital has raised several important issues. Firstly, only 77% of patients with SRH at the ulcer base received endoscopic haemostasis at initial endoscopy. This was explained largely by reduced use of endoscopic haemostasis in cases with adherent clot at the ulcer base. Although endoscopic haemostasis has previously not been advocated for peptic ulcers with adherent clot alone at the base and many endoscopists are hesitant to disturb adherent clot when there is no active bleeding, recent evidence suggests otherwise. Variability of endoscopic haemostasis techniques
In cases where endoscopic haemostasis was applied, the type which was used was variable. The large number (n=19) and variable experience of endoscopists and the introduction of novel modalities, such as thrombin, during the review period is likely to have contributed to this lack of uniformity of endoscopic haemostasis.

Re-bleeding rates after endoscopic haemostasis
Another important point relates to the re-bleeding rate in patients who received endoscopic haemostasis. Although the re-bleeding rate after adrenaline endoscopic haemostasis (29%) was comparable to published RCT data, the re-bleeding rate in patients who received combination endoscopic haemostasis was particularly high (57%) in comparison with data from RCTs that have randomised only patients with actively bleeding ulcers. However, the size of the acute UGI bleed (measured by size of blood transfusion) in the unselected group of patients who received combination endoscopic haemostasis during our review period was higher than in published RCTs (reported between 3–5 units). This may explain the high re-bleeding rate in this group of patients as size of the acute bleed (measured by presence of shock, haemodynamic instability, and transfusion requirement) has been demonstrated to be an important predictor of re-bleeding. Another possible explanation which deserves further (prospective) study is that, outside of RCTs, combination endoscopic haemostasis may be less efficacious than adrenaline injection alone.

Management of patients with re-bleeding after initial endoscopy
Eleven of 20 patients who re-bleed had repeat endoscopic haemostasis. Patients who received repeat endoscopic haemostasis had a higher prevalence of co-morbidity than those who underwent emergency surgery immediately. The position of the ulcer and “ease” of endoscopic haemostasis at initial endoscopy also appears to have been a factor in the decision whether to refer a patient for immediate surgery or repeat endoscopy as those cases with a difficult ulcer location are over-represented in the group who received immediate surgery. Forty five per cent of the patients who had repeat endoscopic haemostasis eventually required salvage surgery compared with 27% of patients who had repeat endoscopic haemostasis for re-bleeding in the RCT performed by Lau et al. Furthermore, the failed repeat endoscopic haemostasis group had a significantly higher transfusion rate and longer inpatient stay compared with those who had immediate surgery (table 3). Although mortality rates in the immediate surgery and repeat endoscopic haemostasis groups were similar, the duration of hospital stay was longer for those patients who received repeat endoscopic haemostasis (especially if that modality failed with subsequent surgery). However, the small number of patients does not allow a definitive conclusion to be drawn. Despite reviewing endoscopic data in a large unit over a three year period, we were still not able to analyse a sufficient number of cases of re-bleeding after endoscopic haemostasis.

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**Table 3** Patient characteristics, transfusion requirement, and outcome of patients who underwent surgery or repeat endoscopic haemostasis after re-bleeding

<table>
<thead>
<tr>
<th>Treatment modalities</th>
<th>Immediate surgery (n=7)</th>
<th>Repeat endoscopic haemostasis Failed (n=5)</th>
<th>Successful (n=6)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean age (range)</td>
<td>69 (47–89)</td>
<td>72 (58–81)</td>
<td>58 (33–87)</td>
</tr>
<tr>
<td>Co-morbidity [%]</td>
<td>43</td>
<td>100</td>
<td>67</td>
</tr>
<tr>
<td>Active bleeding [%]</td>
<td>57</td>
<td>60</td>
<td>50</td>
</tr>
<tr>
<td>Endoscopist grade (n)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Registrar</td>
<td>4</td>
<td>2</td>
<td>2</td>
</tr>
<tr>
<td>Senior registrar</td>
<td>3</td>
<td>2</td>
<td>3</td>
</tr>
<tr>
<td>Consultant</td>
<td>0</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>Initial endoscopic haemostasis type [%]</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Adrenaline</td>
<td>86</td>
<td>40</td>
<td>33</td>
</tr>
<tr>
<td>Combination</td>
<td>14</td>
<td>60</td>
<td>67</td>
</tr>
<tr>
<td>No [%] with difficult ulcer location</td>
<td>3 (57)</td>
<td>1 (20)</td>
<td>0</td>
</tr>
<tr>
<td>Mean (SEM) total transfusion (units)</td>
<td>11.5 (1.54)</td>
<td>15.8 (2.71)</td>
<td>9.5 (2.24)</td>
</tr>
<tr>
<td>Mean (SD) length of inpatient stay (days)</td>
<td>16 (3)</td>
<td>41 (21)</td>
<td>17 (4)</td>
</tr>
<tr>
<td>Mortality [n]</td>
<td>1</td>
<td>1</td>
<td>1</td>
</tr>
</tbody>
</table>
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ulcers with SRH. However, during our review period (1997–99), intensive acid suppression was not used after endoscopic haemostasis for bleeding peptic ulcer at St James’s University Hospital.

Our data have revealed a lower proportion of patients with a diagnosis of bleeding peptic ulcer than previous published reports of audits of acute UGI bleeding.1 Only one reason for this may have been bias due to the retrospective nature of the study. A more likely explanation is the less stringent criteria for evidence of gastrointestinal bleeding that was applied in this study when compared with the published literature.2

Although an endoscopic technique. However, the need for surgery or repeat endoscopic haemostasis for bleeding peptic ulcer at St James’s University Hospital was likely to have included a number of cases with less or no pathology and lowered the proportion of peptic ulcers presenting as an acute UGI bleed.

The results presented here have implications for the British Society of Gastroenterology working party report on provision of endoscopy related services which has recently been published.11 Our study suggests that specific guidelines for emergency endoscopic management should be incorporated in order to minimise variability of practice, especially for units where provision of emergency UGI endoscopy is dependent on multiple endoscopists. Our unit now uses a specific protocol for management of patients with a bleeding ulcer which incorporates specific guidelines on endoscopic haemostasis technique. However, the need for surgery or repeat endoscopic haemostasis is still left to individual gastroenterologists and surgeons in collaboration.

In conclusion, this study has demonstrated variability of endoscopic management of patients with bleeding peptic ulcer and higher re-bleeding rates after endoscopic haemostasis compared with those reported in RCTs. Although an increase in bleed severity and co-morbidity may account for higher re-bleeding rates, variability in operator experience and haemostatic techniques are also likely to be contributory.

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