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The use of hemostatic spray as an adjunct to the conventional hemostatic measures in high-risk nonvariceal upper GI bleeding (with video)

Rohit Sinha, MBBS, MRCP, PgDip, Khalida A. Lockman, MBChB, MRCP, MD, Nicholas I. Church, MBChB, MD, FRCP, John N. Plevris, MD, PhD, FRCPE, FEBGH, Peter C. Hayes, BSc, MBChB, MD, PhD, FRCPE

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Title:

The use of hemostatic spray as an adjunct to the conventional hemostatic measures in high-risk nonvariceal upper GI bleeding (with video)

Authors:
Rohit Sinha¹, Khalida A. Lockman², Nicholas I. Church¹, John N. Plevris¹, Peter C. Hayes¹

Affiliations:
¹Centre of Liver and Digestive Disorder, ²Acute Medicine Unit, The Royal Infirmary and The University of Edinburgh, 51 Little France Crescent, Edinburgh EH16 4SA, United Kingdom

Degrees:
R Sinha – MBBS, MRCP, PgDip
KA Lockman – MBChB, MRCP, MD
NI Church – MBChB, MD, FRCP
JN Plevris – MD, PhD, FRCPE, FEBGH
PC Hayes – BSc, MBChB, MD, PhD, FRCPE

Corresponding Author:
Dr Rohit Sinha
Centre of Liver and Digestive Disorder
The Royal Infirmary and The University of Edinburgh, 51 Little France Crescent, Edinburgh EH16 4SA, United Kingdom.
Email: rohit.sinha@nhs.net
Telephone: 0044 131 2421625
Fax: 0044 131 2421633
The use of hemostatic spray as an adjunct to the conventional hemostatic measures in high-risk nonvariceal upper GI bleeding (with video)

R Sinha¹, KA Lockman², NI Church¹, JN Plevris¹, PC Hayes¹

¹Centre of Liver and Digestive Disorder, ²Acute Medicine Unit,
The Royal Infirmary and The University of Edinburgh,
51 Little France Crescent, Edinburgh EH16 4SA, UK

Corresponding author:
Dr Rohit Sinha
Centre of Liver and Digestive Disorder
The Royal Infirmary and The University of Edinburgh,
51 Little France Crescent,
Edinburgh EH16 4SA,
United Kingdom.
Email: rohit.sinha@nhs.net
Telephone: 0044 131 2421625
Fax: 0044 131 2421633
Abstract

Background: Endoscopic management of nonvariceal upper gastrointestinal bleed (NVUGIB) can be challenging. Hemospray (Cook Medical, Winston-Salem, NC, USA) is a novel endoscopic hemostatic agent for NVUGIB. Its efficacy in attaining hemostasis in NVUGIB is promising, particularly with respect to technically difficult lesions. However, most of the currently available data are focused on its application as monotherapy. The aim of this study was to evaluate its efficacy as a second agent to adrenaline, or as an addition to the combination of adrenaline with either clips or a thermal device in NVUGIB.

Methods: Consecutive patients with Forrest 1a and 1b ulcer treated with hemostatic spray as an adjunct to conventional endoscopic hemostatic measures between July 2013 and June 2015 were included in this retrospective analysis. The endpoints were initial hemostasis, 7-day rebleeding, 30-day rebleeding, all cause, and GI-related 30-day mortality.

Results: A total of 20 (median age 75 years, 50% men, 60% Forrest 1a ulcer) were treated with hemostatic spray as a second agent to adrenaline, or as an adjunct to the combination of adrenaline with either clips or a thermal device. Hemostatic spray was used as a second agent to adrenaline in 40% and as a third agent to combined dual therapy in 60%. Initial hemostasis was attained in 95% with an overall rebleeding rate at 7-day of 16%. There was no difference between 7-day and 30-day rebleeding rate. The combination of hemostatic spray and adrenaline resulted in 100% initial hemostasis and 25% 7-day rebleeding. Similarly, initial hemostasis was achieved in 92% with 9% rebleeding rate when hemostatic spray was used as the third agent to 2 of the conventional measures. All-cause mortality was 15% with one GI-related death (3%).

Conclusion: In our single-center retrospective analysis, hemostatic spray appears promising as an adjunct to conventional methods for NVUGIB, although prospective controlled trials are needed to confirm.

Keywords:
Hemospray, nonvariceal gastrointestinal bleeding, hemostatic powder

Introduction

Mortality associated with acute upper gastrointestinal bleeding remains high at 10% 1, 2. This figure has not changed appreciably since the 1950s in part due to an aging population. In a recent United Kingdom-wide audit, 1 in 4 patients with acute upper GI bleeding were aged over 80 years 2. Because mortality from upper GI bleeding increases steeply with age, peptic ulcer bleeding will continue to pose a significant therapeutic challenge 3. Recent advances in endoscopic techniques have enabled effective hemostatic interventions to be deployed. The combination of adrenaline with a mechanical method (clips) or with a thermal device is the preferred strategy to achieve hemostasis 4, 5, 6. Indeed, monotherapy with adrenaline has largely been superseded by this approach due to an unacceptably high rate of rebleeding 7. However, in 10% to 20% of patients, rebleeding occurs despite initial hemostasis with combined therapy 1.
Hemostatic sprays are relatively new addition to the established hemostatic agents. Hemospray (Cook Medical, Winston-Salem, NC, USA is the first to be licensed in Europe, but it is not as yet FDA approved. It is a proprietary inorganic hemostatic powder approved for use in nonvariceal upper gastrointestinal bleed (NVUGIB). The powder is propelled through a carbon dioxide pressurized catheter fed through the working channel of an endoscope and sprayed at a distance of 1 to 2 cm from the bleeding site until a complete coating of the lesion and an obliteration of fresh blood is achieved. When put in contact with moisture (eg, blood or tissue) in the GI tract, the powder becomes cohesive and adhesive. Its hemostatic mechanisms are 2-fold. First, it exerts a tamponade effect by forming a confluent mechanical barrier at the bleeding site. Second, it promotes thrombus formation by enhancing platelet aggregation and increasing the concentration of clotting factors. Additionally, it has been shown to shorten coagulation time in vitro.

The effect of hemostatic spray on hemostasis in NVUGIB cases so far has been encouraging. Initial hemostasis was achieved in 81% to 100% with rebleeding rate of 5-39% . However, most of these data were focused on the use of Hemospray as monotherapy. Moreover, the study populations are heterogeneous with respect to the underlying etiology. Data on the use of hemostatic spray as a second-line hemostatic agent to adrenaline or as an adjunct to combined therapy are limited. Intuitively, these strategies could offer a significant advantage; hemostatic spray may augment hemostasis achieved by standard therapy. This approach is attractive in cases in which the application of dual therapy is challenging. Adrenaline injection occasionally masks the bleeding lesion. More frequently, position of the ulcer renders the effective application of clips or heater probe impossible. Finally, the fibrous nature of the ulcer base or large size of a visible vessel may mean that attempting to apply clips or heater probe would be unlikely to succeed or potentially dangerous.

In this study, we evaluated the effect of this hemostatic spray on initial hemostasis, rebleeding rate, and mortality when used in high-risk patients with peptic ulcer bleed. Here, hemostatic spray was used as a second agent to adrenaline, or as an adjunct to the combination of adrenaline with either clips or a thermal contact device.

**Methods**

**Study population**
This retrospective analysis reviewed 30 consecutive patients who had been treated with hemostatic spray for GI bleeding between July 1, 2013 and June 30, 2015 in a large tertiary unit. One case of hemostatic spray application to lower GI bleeding, one case with incomplete clinical information and 4 cases with the following underlying causes: gastroesophageal carcinoma, Mallory Weiss tear without visible vessel, angiodysplasia, and pangastritis, were excluded. Four patients with Forrest 2a and 2b ulcers were excluded due to the small sample size.

The final analysis included the remaining 20 cases of Forrest 1a and 1b ulcers. Of these, one patient had post-ampullectomy GI bleeding, but a spurting arterial vessel was identified on the index gastroscopy. Similarly, another patient who had a Mallory Weiss tear was included in this group because a visible vessel was treated. The nature of the lesions in these 2 cases was akin to that of a peptic ulcer and therefore were considered as Forrest 1a and 1b, respectively. In these cases, hemostatic spray was used either as a second agent to adrenaline or as an adjunct to the combination of adrenaline with either clips or a thermal contact device.
The outcomes were as follows: initial hemostasis (defined as adequate hemostasis confirmed on observation and to the satisfaction of endoscopist), 7-day and 30-day rebleeding (defined as (1) clinical presentation of hematemesis or melena; (2) hemodynamic instability; (3) drop in hemoglobin ≥2 g/L, blood transfusion of ≥4 units; necessitating further investigation such as repeat endoscopy or radiological intervention); all cause and GI bleed-related 30-day mortality; adverse event; and equipment failure relating to the use of hemostatic spray.

For comparison, we reviewed the outcomes of 20 consecutive patients with Forrest 1a and 1b ulcer who received conventional treatment before the introduction of hemostatic spray in our department between August 2010 and June 2013.

Patients were identified using the endoscopy logbook entries. Endoscopy report and clinical information were obtained from Unisoft (Unisoft Medical Systems, Enfield, UK) and Trakcare (Intersystem Corporation, Cambridge, Mass) respectively. Demographics, Rockall, and Blatchford scores, antiplatelet use, findings at endoscopy, endoscopic therapeutic interventions, use of proton pump inhibitor infusions, *Helicobacter pylori* status, and the study end points were recorded. All identifiable data were replaced with study identification numbers to preserve confidentiality. The use of these data was approved by the Confidentiality (Caldicott) Guardian for NHS Lothian.

This study was considered as a retrospective review by the local committee in our institution; hence, no formal ethical approval from the NHS Research Ethics Committee was necessary.

**Clinical procedure**

The choice of treatment was at the discretion of the attending endoscopist. All patients were adequately resuscitated before the endoscopic procedure. Hemostatic spray was used to achieve complete hemostasis if there was evidence of continued oozing after conventional therapy (adrenaline with either hemostatic clips or thermal devices). Similarly, hemostatic spray was used after adrenaline injection if the lesion was inaccessible for application of hemostatic clips or thermal devices. Hemostatic spray was delivered through a 10F catheter (Cook Medical, Winston-Salem, NC, USA) fed through the working channel of an endoscope (Olympus, Japan or FujiFilm, Japan). Adequacy of hemostasis was confirmed on observation of the treated lesion and to the satisfaction of the endoscopist. All patients received intravenous proton pump inhibitor infusion for 72 hours (Hong Kong regimen) after endoscopy.

**Statistical analysis**

Statistical analysis was performed using IBM SPSS for Macintosh version 22.0 (IBM Corporation, Armonk, NY, USA). Mann-Whitney U test was used to compare continuous variables between two populations. The Fisher exact test was used to compare categorical variables between 2 groups. Data are presented in median interquartile range (IQR) or frequency (percentage of the total study population). A 2-sided P value of less than .05 was considered to be statistically significant.

**Results**
Hemostatic spray group
A total 20 patients (median age 75 (IQR=11) years, men 50%) were included in the final analysis. Of these, 60% (12/20) had Forrest 1a and 40% (8/20) had Forrest 1b. The median Rockall score and Blatchford score were 8 (IQR = 2) and 14 (IQR = 6) respectively. Ten (50%) patients were on anti-platelet agents at the time of presentation.

Hemostatic spray was used as a second agent to adrenaline (AH) in 8 (40%) and as an addition to combined therapy (DH) in 12 (60%) patients (Table 1). The reasons for hemostatic spray application in the AH group were as follows: lesions in difficult site, suboptimal views, and concerns regarding perforation or torrential hemorrhage, all of which had precluded effective deployment of hemostatic clips or a thermal contact device. Hemostatic spray was used as a third agent with persistent oozing or in cases where the risk of rebleeding remained significant despite the conventional dual therapy. Descriptions of individual cases are provided in Supplementary Table 1.

Initial hemostasis and rebleeding
Initial hemostasis was achieved in 19 out of 20 (95%) cases. One patient (5%) who did not achieve initial hemostasis despite the use of hemostatic spray in the DH group subsequently had gastroduodenal artery embolization to achieve hemostasis. Seven-day rebleeding occurred in 16% (3/19; median day 0, IQR = 3). Rebleeding events in the AH and DH group were 25% (2/8) and 9% (1/11), respectively (Table 3). Two of the rebleeding cases were treated with gastroduodenal artery embolization. One patient was too frail for further intervention and subsequently died. There was no rebleeding recorded beyond day 7. Thus, the 30-day rebleeding remained at 15%.

Mortality
All-cause mortality at 30 days in the hemostatic spray group was 15% (3/20) (Table 3). GI bleed-related mortality at 30-day was 5% (1/20). Other causes of death were septicemia and aspiration pneumonia.

Rebleeding event and mortality according to the severity of bleed
Forrest 1a
Forty-two percent (5/12) of those with Forrest 1a lesions received hemostatic spray as a second line to adrenaline therapy (AH) (Table 3). In this group, initial hemostasis was achieved in 100%. However, rebleeding was noted in 40% (2/5). The 30-day mortality in this group was 20% (1/5). The remaining 58% (7/12) had hemostatic spray in conjunction with the combined therapy (DH). Of these, the initial hemostasis rate was 86% (6/7). Rebleeding occurred in 17% (1/6). There was no mortality in this group.

Forrest 1b
Thirty-eight percent (3/8) of patients with Forrest 1b lesions had AH therapy, with initial hemostasis of 100%, 0% rebleeding, and 0% 30-day mortality (Table 3). The remaining 62% (5/8) received DH therapy. Initial hemostasis was 100%, and there were no rebleeding episodes. However, 30-day mortality was 40% (2/5). One death was due to septicemia and another was due to aspiration pneumonia.

Other adverse events and technical failure with hemostatic spray
There was no significant hemostatic spray-related equipment failure. On occasion, it was necessary to use the second supplied catheter to achieve adequate therapy. This was due to blockage within the first catheter induced by moisture within the endoscope channel or the gut
lumen. The former can be minimized by vigorous flushing of the endoscope channel with air before catheter insertion. No other adverse event was observed.

**Conventional therapy group outcomes (August 2010 to June 2013)**

Baseline characteristics (age, gender, etiology, preceding anti-thrombotic use, Rockall and Blatchford scores) were similar in both groups (Table 2). Descriptions of individual cases are provided in Supplementary Table 2.

**Initial hemostasis and rebleeding**

Initial hemostasis with conventional strategies was achieved in 16 out of 20 (80%) and rebleeding occurred in 4 out of 16 (25%) and 5 out of 16 (31%) at 7 days and 30 days, respectively. Three out of 4 patients with failed initial hemostasis required emergency laparotomy. One died of ongoing bleeding. Three patients were treated with adrenaline monotherapy due to technical difficulties in deploying clips or thermal device. Rebleeding occurred in 2 out of 3 (67%) of these cases. Among those who received the standard dual therapy (adrenaline and hemostatic clips (AC) or adrenaline and a thermal device (AT), initial hemostasis was achieved in 14 out of 17 (82%). The 30-day rebleeding rate of 21% (3/14) was comparatively higher than that seen with the combination of hemostatic spray and standard dual therapy (9%).

**Mortality**

The overall 30-day mortality with conventional therapy was 6 out of 20 (30%), and GI bleed-related death occurred in 3 out of 20 (15%).

**Discussion**

Our study examined the efficacy of Hemospray as a second agent to adrenaline and as a third agent to the combined therapy in high-risk patients with nonvariceal upper GI bleed. These patients were truly high risk with median Rockall and Blatchford scores of 8 and 14, respectively. In this context, our rationale was to use hemostatic spray to maximize the hemostatic effect. Our data highlight 2 important points. First, using this approach, hemostatic spray was effective in attaining initial hemostasis as evident by the favorable rate of 95%. Second, hemostatic spray offers a good protection against rebleeding. Indeed, the overall rebleeding rate in our cohort was 16%, and when used in combination dual conventional treatment modalities, the rebleeding rate was a mere 9%.

In this study, hemostatic spray was used as a second agent to adrenaline in 40% of cases in which the deployment of clips or thermal contact devices was not technically feasible. Initial hemostasis achieved in combination with adrenaline was 100%, even in high-risk lesions. Importantly, the rebleeding rate of 25% in Forrest 1a and Forrest 1b ulcers with this combination was better than that seen with adrenaline monotherapy for similar lesions. When compared with our previous experience in a group of patients with Forrest 1a and 1b lesions before introduction of hemostatic spray in our unit, rebleeding occurred in 67% when the therapeutic option was technically limited to adrenaline monotherapy. Therefore, hemostatic spray may be a viable alternative for lesions inaccessible to hemostatic methods which require precise targeting. To this end, hemostatic spray can be applied without direct mucosal contact or en face view of a lesion. Prior injection of adrenaline can provide an immediate clear visualization for a targeted application of hemostatic spray and complement its hemostatic effect. Thus, hemostatic spray
offers ease of application, particularly in technically challenging cases. It obviates the need for more advanced techniques such as the use of a duodenoscope or a cap-assisted approach, for which the required skills may not be widely available.

Sustained hemostasis with a combination of hemostatic spray and adrenaline in our cohort was comparatively lower than that reported with conventional measures, particularly for Forrest 1a ulcers. It is of interest that although Hemospray has been licensed as monotherapy for NVUGIB, its efficacy in Forrest 1a lesions has not been adequately studied. Our center experience of hemostatic spray as monotherapy was limited to a single case where it was successfully used to control a significant bleeding episode from pangastritis. In a porcine model, hemostatic spray induces 100% hemostasis with 20% rebleeding within 24 hours. These findings were corroborated by subsequent clinical studies. For example, Sung et al. achieved 95% initial hemostasis with 85% sustained hemostasis at 72 hours in patients with Forrest 1a and 1b ulcers. Likewise, Smith et al. reported initial hemostasis in 85% and a rebleeding rate of 15% using hemostatic spray as monotherapy. However, in a subsequent case series, the rate of rebleeding with hemostatic spray monotherapy was considerable prompting authors to recommend either a second re-look or prolonged hospital observation. This raises the question as to whether hemostatic spray as monotherapy is a cost-effective approach. This was addressed in a recent study, which showed that the combination of hemostatic spray with conventional hemostatic measures was more cost-effective than hemostatic spray as monotherapy. In a recent case series, the use of hemostatic spray as monotherapy in Forrest 1a lesions was associated with 100% rebleeding necessitating further endotherapy or interventional radiology. Therefore, the available evidence currently suggests that hemostatic spray monotherapy is unlikely to replace conventional hemostatic therapy.

In our series, hemostatic spray was used as an adjunct to the standard dual therapy in cases where the risk of rebleeding was deemed to be significant. This triple therapy approach resulted in the overall initial hemostasis of 92% with the rebleeding rate of 9%. Even among those with Forrest 1a, the initial hemostasis and the rebleeding rate were encouraging. Furthermore, in Forrest 1a and 1b lesions, the rebleeding rate of 9% was comparatively less than the rebleeding rate reported with the conventional approaches for similar lesions. Indeed, the hemostatic effect with the triple therapy surpassed the initial hemostasis of 82% and the rebleeding rate of 21% previously observed with the conventional dual therapy in our institution. This effect might be explained by the ability of hemostatic spray to cover large areas and address multiple bleeding points from feeding vessels simultaneously. It could be surmised that the spray provides a sealant barrier that complements tamponade, vasoconstriction, and mechanical or thermal coagulation resulting from conventional combination therapy.

Similar to the findings of a recent United Kingdom–wide audit, our cohort was elderly with median age of 75 years and the mortality observed was largely related to decompensation of comorbid conditions. One GI bleed-related death was attributable to a duodenal ulcer that had rebled. In this case, no further intervention was deemed appropriate due to significant frailty. In our study, there was no difference in mortality when hemostatic spray was used as an adjunct to adrenaline or incorporated into the standard combination therapy. Even though there have been reports of perforated viscus, biliary orifice obstruction, and splenic infarct associated with the use of hemostatic spray, we did not observe any adverse events in our cohort.

This study has several limitations. The retrospective nature precludes a comparison of the efficacy of hemostatic spray as monotherapy compared with its adjuvant use to dual therapy.
Cases were highly selected and treatment was at the discretion of the endoscopist rather than directed by a specific protocol. Data regarding the exact amount of hemostatic spray applied are unavailable. Our case series, albeit small, is the largest to date that reports the complementary role of hemostatic spray, either as a second agent after adrenaline injection or as a third agent after conventional dual therapy in high-risk nonvariceal upper GI bleeding.

In conclusion, our data support the use of hemostatic spray in high-risk peptic ulcer bleeding, particularly when the application of additional hemostatic agents after adrenaline injection may not be technically feasible. Its use as an adjunct can be considered when primary hemostasis is not achieved with standard combination therapy. Whether it can replace conventional measures remains to be seen and its role in additionally securing hemostasis in high-risk patients remains unclear. Future prospective randomized controlled trials comparing hemostatic spray alone and in various combinations with conventional therapies in high-risk bleeding lesions are required to establish its optimal place in the hemostatic armamentarium.

REFERENCES


Legends

Figure 1: Forrest 1a duodenal ulcer

Figure 2: Forrest 1a duodenal ulcer after a triple therapy (a combination of adrenaline, thermal device and hemostatic spray)

Video Legend
A 76-year-old woman with cardio-respiratory comorbidity had presented with major peptic ulcer bleeding 2 weeks previously. She remained in intensive care after a laparotomy and under-running of a duodenal ulcer after failure of endoscopic hemostasis. After further bleeding and cardiovascular instability developed, gastroscopy under general anesthetic was performed. The surgical team felt she would not survive a second laparotomy.

The site of bleeding was a visible vessel in a high lesser curve gastric ulcer previously obscured by fundal clot. Circumferential adrenaline injection was performed to a total of 28 mL. The needle can be seen to puncture the feeding vessel during circumferential injection. The heater probe was used at 30J power setting to cavitate around the visible vessel. It was decided not to treat the vessel directly due to the high risk of precipitating torrential hemorrhage. The vessel began to actively ooze during therapy. Clips would not have been an option due to the chronic fibrous ulcer base. Therefore, hemostatic spray was used to effect hemostasis. The patient underwent a semi-urgent left gastric artery embolization the following day and was successfully discharged home 2 months later.
**Table 1: Demographics and clinical characteristics of patients treated with hemostatic spray.**

*Others: Mallory Weiss with visible vessel, ampullectomy with arterial spurt. Data presented as median (IQR) or n.*
Clinical characteristics

<table>
<thead>
<tr>
<th>Clinical characteristics</th>
<th>Hemostatic spray group (July 2013- August 2015)</th>
<th>Conventional therapy (August 2010 – June 2013)</th>
<th>P value</th>
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<td>Age (years)</td>
<td>75 (11)</td>
<td>73 (21)</td>
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</tr>
<tr>
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<td>10</td>
<td>9</td>
<td>1.00</td>
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<tr>
<td>Rockall score</td>
<td>8 (2)</td>
<td>7 (3)</td>
<td>0.30</td>
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<tr>
<td>Blatchford score</td>
<td>14 (6)</td>
<td>14 (4)</td>
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<tr>
<td>Use of antithrombotic therapy</td>
<td>10</td>
<td>12</td>
<td>0.75</td>
</tr>
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</table>

Data presented as median (IQR) or n.

Table 2: Clinical characteristics of patients with Forrest 1a and 1b ulcers in hemostatic spray and conventional therapy
AH, adrenaline plus hemostatic spray; DH, conventional dual therapy plus hemostatic spray; A, adrenaline alone; D, conventional dual therapy (a combination of adrenaline with hemostatic clips or a thermal device); * 30-day rebleeding.

Table 3: Initial hemostasis, rebleeding events, and 30-day mortality stratified according to the Forrest classification in hemostatic spray and conventional treatment groups
<table>
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<th>No</th>
<th>Age (years)</th>
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<th>Blatchford score</th>
<th>ATT</th>
<th>Etiology</th>
<th>Forrest classification</th>
<th>Endoscopic treatment</th>
<th>Initial hemostasis</th>
<th>Rebleeding</th>
<th>Outcome</th>
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<td>1</td>
<td>81</td>
<td>M</td>
<td>10</td>
<td>7</td>
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<td>1b</td>
<td>DH</td>
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<td>no</td>
<td>survived 30 days</td>
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<td>2</td>
<td>70</td>
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<td>9</td>
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<td>Duodenal ulcer</td>
<td>1a</td>
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<td>No; embolized</td>
<td>no</td>
<td>survived 30 days</td>
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<td>DH</td>
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<td>4</td>
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<td>Yes</td>
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<td>17</td>
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<td>DH</td>
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<td>Yes; Day 6 - embolization</td>
<td>survived 30 days</td>
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<td>AH</td>
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<td>no</td>
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<td>DH</td>
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F, Female; M, Male; ATT, Antithrombotic therapy; AH, adrenaline plus hemostatic spray; DH, conventional dual therapy plus hemostatic spray; GI, gastrointestinal

Supplementary Table 1: Endoscopic treatment and outcomes of 20 consecutive patients with Forrest 1a and 1b ulcers treated with hemostatic spray from July 2013 to August 2015.
<table>
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<th>No</th>
<th>Age (years)</th>
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<th>Blatchford score</th>
<th>ATT</th>
<th>Etiology</th>
<th>Forrest classification</th>
<th>Endoscopic treatment</th>
<th>Initial hemostasis</th>
<th>Rebleeding</th>
<th>Outcome</th>
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</tbody>
</table>

F, Female; M, Male; ATT, Antithrombotic therapy; A, adrenaline only; AT, adrenaline plus a thermal device; AC, adrenaline plus hemostatic clip; GI, gastrointestinal

Supplementary Table 2: Endoscopic treatment and outcomes of 20 consecutive patients with Forrest 1a and 1b ulcers from August 2010 to June 2013 before the introduction of hemostatic spray.
### Supplementary Table 3: Previous studies on hemostatic spray as a monotherapy in high-risk NVUGIB.

<table>
<thead>
<tr>
<th>No. of subjects</th>
<th>Lesion</th>
<th>Initial hemostasis (%)</th>
<th>7-day re-bleeding (%)</th>
<th>30-day mortality (%)</th>
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*In animal (pig) model; NVUGIB, nonvariceal upper gastrointestinal bleed; ±Unsustained primary hemostasis; NA, not available
Authors' contribution
Contribution: RS collected, analysed the data and wrote manuscript. KAL and NIC advised on manuscript, presentation and DICOM file contribution. JNP and PCH conceptualized the idea and edited the manuscript.

Conflict of interest: The authors do not have any conflict of interest to declare.

Acknowledgement: Authors acknowledge and thank Mrs Anita Talbot for identification of cases through endoscopy log-book record.
List of Acronyms

AH     Adrenaline plus Hemospray®
DH     Dual therapy plus Hemospray®
Fr     French
GI     Gastrointestinal
IQR    Interquartile Range
NHS    National Health Service
NVUGIB Non Variceal Upper Gastrointestinal Bleed