

Effectiveness of TC-325 (Hemospray) for treatment of diffuse or refractory upper gastrointestinal bleeding – a single center experience



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ABSTRACT

Background and study aims TC-325 (Hemospray, Cook Medical) is a powder agent for endoscopic hemostasis in patients with upper gastrointestinal bleeding (UGIB). Although most publications are based on case-reports and retrospective studies, data on efficacy are promising. Here we report our experience with TC-325 for diffuse or refractory UGIB.

Patients and methods Data on patients receiving TC-325 for endoscopic hemostasis from November 2013 to February 2017 at our center were analyzed retrospectively. Primary endpoints were technical success (successful immediate hemostasis) and clinical success (effective hemostasis and no recurrent bleeding). Secondary endpoints were recurrent bleeding within 3 and 7 days, hospital mortality and TC-325 associated complications. TC-325 was used for bleeding not amenable to standard endoscopic treatment (e.g. diffuse bleeding) or as salvage therapy after failure of conventional methods

Results Fifty-two patients received TC-325 treatment. Most of the patients were treated for peptic ulcer bleeding (18/52 patients, 34.6%) and post-interventional bleeding (13/52 patients, 25%). Hemospray was used in 23/52 (44.2%) patients as monotherapy and in 29/52 (55.8%) patients as a salvage therapy. Application of the powder on the bleeding source was successful in all patients with no therapy-related adverse events (AEs). Immediate hemostasis was achieved in 51/52 (98.1%) patients. Recurrent bleeding within 3 and 7 days was observed in 22/51 and 25/51 patients respectively (43.1% and 49%). The overall clinical success was 56.9% on day 3 and 51% on day 7. Total mortality was 15.4% (8 patients), bleeding associated mortality was 3.8% (2 patients). There were no therapy-related AEs.

Conclusions TC-325 showed a high technical success rate as monotherapy for bleeding sources not amenable to standard methods or as an “add-on” therapy after unsuccessful hemostasis. However, rebleeding was frequent in this cohort and further studies are warranted to exactly define a treatment algorithm for TC-325 use.

Introduction

Upper gastrointestinal bleeding (UGIB) is a common cause for hospitalization with an worldwide annual incidence of about 40 to 150 per 100.000 and significant morbidity and mortality [1]. The most frequent causes of UGIB are peptic ulcers (28–59%), mucosal erosive disease of the esophagus or stomach/

duodenum (1–47%), Mallory-Weiss-syndrome (4–7%) and malignancy (2–4%) [1].

Depending on the cause of bleeding and anatomical localization, management of UGIB can be challenging. Standard methods for endoscopic hemostasis include mechanical, injection or thermal therapy. The current ESGE guideline on diagnosis and management of non-variceal UGIB recommends a com-

bination of 2 modalities for management of peptic ulcer bleeding [1]. With a combination of these procedures, a hemostasis rate of 85% to 95% can be achieved, however, in about 5% to 10% of patients, rebleeding was reported [2].

TC-325 (Hemospray, Cook Medical) is a hemostatic powder approved for endoscopic hemostasis in the upper gastrointestinal tract. It consists of a mineral-based compound that after interacting with moisture (e.g. blood) becomes cohesive and adhesive, thus forming a tamponade covering the bleeding site. Through absorption of the fluid of the blood it leads to concentration of clotting factors and activates the clotting cascade [3]. Efficacy of TC-325 was initially shown in a pilot study in upper gastrointestinal peptic ulcer bleeding [4] and subsequently in various causes of gastrointestinal bleeding, ranging from cancer bleeding to acute variceal bleeding [5–9]. The SEAL-Study was the first multicenter prospective registry demonstrating successful immediate hemostasis in 85% and a rebleeding rate of 15% in non-peptic-ulcer bleeding [10], thus confirming data from other smaller cohorts [4, 7]. The overall primary hemostasis rate was otherwise reported to be in the range of 93% to 98% [2, 6, 7]. In the current ESGE guideline for management of non-variceal UGIB, the role of TC-325 has not been clearly defined. Available data suggest that Hemospray is safe and effective and may be best used in high-risk cases as a temporizing measure or a bridge toward more definitive treatment [1]. Herein we report on our single-center experience with TC-325 in a 38-month period in patients with UGIB that was not amenable to standard endoscopic therapy or had persistent bleeding after standard endoscopic hemostasis.

Patients and methods

Since the introduction of TC-325 in November 2013, data has been collected on patients who have received TC-325 for endoscopic hemostasis at our center. Those data were analyzed retrospectively. This study was approved by the local institutional review board.

Indications for Hemospray use in our institution were: (1) UGIB that was not otherwise treatable (e.g. due to difficult anatomical situation or diffuse bleeding without definite source – in those cases, Hemospray was used as monotherapy); and (2) application of Hemospray in UGIB as a salvage therapy after failure of other endoscopic methods.

Primary endpoints were: technical success (defined as successful immediate hemostasis after TC-325 application) and clinical success (defined as effective hemostasis and no recurrent bleeding in 7 days). Recurrent bleeding was diagnosed if evidence of fresh blood or signs of bleeding in the treated site was seen on follow-up endoscopy. Otherwise recurrent bleeding was defined if one of these criteria had been met: (1) hematemesis or melena; (2) a drop in hemoglobin >2 mg/dl or transfusion of 4 or more blood packs; or (3) hemodynamic instability as previously described [4]. Secondary endpoints were recurrent bleeding until Days 3 and 7, hospital mortality, and TC-325-associated adverse events (AEs).

Complete Rockall-score was calculated for each patient to estimate the rebleeding risk as previously described [11].

Our center uses both standard and therapeutic endoscopes as part of standard protocol in acute bleeding situations. In patients with an expected major bleed, therapeutic endoscopes are used regularly. To avoid clogging of the working channel or kinking of the catheter, a 10F catheter was applied through a therapeutic endoscope to the TC-325 cartridges used in this study. Due to the strict indications for the application of Hemospray in our unit, therapeutic endoscopes were generally already in use and no further endoscope change was needed. Second-look endoscopy was performed in patients with high-risk bleeding (e.g. Forrest Ia-Forrest IIb ulcer bleeding) at the discretion of the endoscopist.

Statistical analysis

Patient data were collected prospectively in a database created with Microsoft Excel™. Statistical calculations were done with Microsoft Excel™ and with the Statistical Online Computational Resource (<http://www.socr.ucla.edu/htmls>). Descriptive statistics consisted of the mean, median, SD and range. Fisher's exact test was used to compare qualitative data. A *P* value <0.05 was considered significant.

Patient characteristics

Fifty-two patients with a total of 67 TC-325 applications in the upper gastrointestinal tract were documented from November 2013 until February 2017. TC-325 was used in 23 of these patients (44.2%) as a monotherapy in bleeding not amenable to standard endoscopic therapy and in 29 of the patients (55.8%) as a salvage therapy after unsuccessful hemostasis with other endoscopic treatment modalities.

The median complete Rockall-score was 7 points (range 3–10). Twenty-seven of 52 patients (51.9%) had a complete rockall-score of 7 or higher. Among the patients with a complete Rockall-score <7 points, 18 of 25 patients (72%) had a score of 6 points.

The overall median age of the patients was 69 years (range 32–92 years) and most of the patients were male (31 males: 21 females). The most frequent comorbidity was malignancy (16/52 patients, 30.8%), followed by atrial fibrillation and coronary heart disease (8/52 and 6/52 patients; 15.4% and 11.5%, respectively). Thirteen patients (25%) had salicylic acid as an antithrombotic therapy, whereas 13 patients (25%) were on anticoagulation with either warfarin, heparin, rivaroxaban or apixaban. Details of the demographics, medications and relevant comorbidities are stated in ► **Table 1**.

The most frequent bleeding sources were peptic ulcers in 18 of 52 patients (34.6%) (5 gastric ulcer and 13 duodenal ulcer), post-interventional ulcers (i.e. post EMR, ESD, EFTR) in 14/52 patients (27%) and tumor bleeding in 10/52 patients (19.2%). The majority of the gastroduodenal ulcers were classified as Forrest Ib (7/18 patients; 38.9%) and Forrest IIa (6/18 patients, 33.3%). One patient initially had a Forrest-IIa ulcer, which bled repeatedly despite standard endoscopic treatment. The ulcer was eventually stable and became a Forrest-III ulcer. In repeated EGD the ulcer “deteriorated” from a Forrest III to a Forrest IIc ulcer and it was deemed as a sign of rebleeding. Because con-

► **Table 1** Patient demographic and complete Rockall score, important comorbidities and medication in the whole cohort, monotherapy and salvage therapy subgroups. NSAID: non-steroidal anti-inflammatory drugs.

	Total cohort (n = 52)	Monotherapy (n = 23)	Salvage therapy (n = 29)
Age, y, median (range)	69 (32–92)	68 (32–92)	69 (36–89)
Sex (m/f)	31/21	16/7	15/14
Platelet inhibition, n (%)	13 (25)	9 (39.1)	4 (13.8)
Anticoagulation, n (%)	13 (25)	3 (13)	10 (34.5)
NSAID, n (%)	8 (15.4)	1 (4.3)	7 (24.1)
Glucocorticoid, n (%)	1 (1.9)	0	1 (3.4)
Malignancy, n (%)	16 (30.8)	4 (17.4)	12 (41.4)
Atrial fibrillation, n (%)	8 (15.4)	0	8 (27.6)
Coronary heart disease, n (%)	6 (11.5)	3 (13)	3 (10.3)
Complete Rockall score, median (range)	7 (3–10)	7 (5–9)	7 (3–10)
▪ <7, n (%)	25 (48.1)	11 (47.8)	14 (48.3)
▪ ≥7, n (%)	27 (51.9)	12 (52.2)	15 (51.7)

ventional endoscopic therapies had been exhausted, TC-325 was applied.

The mean diameter of the gastroduodenal ulcers was 28 mm (range 15–50 mm). ► **Table 2** summarizes the bleeding etiologies and location. ► **Fig. 1** shows an example of TC-325's application in a duodenal ulcer.

The majority of the bleeding was located in the duodenum (29/52 patients, 55.8%). Of those, 25 were located either in the bulb or in the second part of the duodenum (25/29 bleedings in duodenum 86.2%). Four bleeds (4/29, 13.8%) were around the papilla vateri. In 16 of 52 patients (30.1%), bleeding source was located in the stomach, 10 bleeds were located in the lesser curvature and 11 bleeds were located in the body of the stomach. There were 7 bleeds in the esophagus (13.5%), all of which were in the distal third of the esophagus.

Results

Technical success (successful immediate hemostasis) was achieved in 51 of 52 patient (98.1%). In 1 patient with post-banding variceal bleeding, Hemospray was used as a salvage therapy and failed to achieve hemostasis; the bleeding was stopped with an OTSC-application.

After 3 and 7 days, the overall rebleeding rate was 43.1% (22/51 patients) and 49% (25/51 patients), respectively. The 3-day rebleeding rate for patients who received TC-325 as salvage therapy was 44.8% (13/29 patients) whereas patients with monotherapy had a rebleeding rate of 39.1% (9/23 patients), ($P=0.78$). After 7 days the rebleeding rates was 43.5% (10/23 patients) vs. 51.7% (15/29 patients) in the monotherapy and salvage therapy groups, respectively ($P=0.58$).

The overall rebleeding rate on Day 3 for patients with peptic ulcer, post-interventional ulcer and tumor bleeding was 44.4% (8/18 patients), 50% (7/14 patients) and 20% (2/10 patients),

respectively. On Day 7 the rebleeding rate was 55.5% (10/18 patients), 57.1% (8/14 patients) and 20% (2/10 patients) for peptic ulcer, post-interventional ulcer and tumor bleeding, respectively.

Patients with peptic ulcer had a Day 3 rebleeding rate of 60% in the monotherapy group vs. 38.5% in the salvage therapy group (3/5 patients vs. 5/13 patients, $P=0.61$). The rebleeding rate for post-interventional ulcer on Day 3 was 25% in the monotherapy group vs. 60% in the salvage therapy (1/4 patients vs. 6/10 patients, $P=0.56$). The rebleeding rate for tumor bleeding on Day 3 was 28.6% vs. 0% (2/7 vs. 0/3 patients, $P=1$). On day 7 there were 2 further bleeding cases from peptic ulcer in the salvage therapy group and 1 more bleed from a post-interventional ulcer in the monotherapy group. There were no changes in the tumor rebleeding rate on Day 7. ► **Table 3** summarizes the overall rebleeding rate as well as rebleeding rate in the monotherapy and salvage therapy group after Days 3 and 7 for peptic ulcer, post-interventional ulcer and tumor bleeding. During further hospital stay, there was one patient with late rebleeding on Day 14 due to a mantle cell lymphoma.

Second-look endoscopy was performed in 42 of 52 patients (80.1%) and was done mostly on the first post-interventional day (mean 1.3 days, range 1–12 days). Rebleeding was seen in 19 patients (36.5%). In 13 of 19 patients (68.4%), Hemospray was applied once again. Acute hemostasis was again achieved in all these patients.

Mortality was 15.4% (8/52). Two deaths were bleeding-related: 1 patient had terminal cancer with malignant bleeding and refused further chemotherapy. The other died of hemorrhagic shock. Other causes of mortality are stated in ► **Table 4**. There were no technical difficulties during all TC-325's application or other procedure-related AEs.

► **Table 2** Bleeding characteristics.

Bleeding etiology	n (%)
▪ Peptic ulcer	18 (34,6)
▪ Post-interventional ulcer	14 (26,9)
▪ Tumor bleeding	10 (19,2)
▪ Endoscopic papillotomy	2 (3,8)
▪ Reflux esophagitis	2 (3,8)
▪ Post-variceal bleeding	2 (3,8)
▪ GAVE-Syndrome	1 (1,9)
▪ Dieulafoy's lesion	1 (1,9)
▪ Diffuse post-emesis bleeding	1 (1,9)
▪ Jejunal and duodenal ischemia	1 (1,9)
Bleeding	n = 52
▪ Distal esophagus	7
▪ Gastric	16
– Cardia	1
– Body	11
– Antrum	4
– Lesser Curvature	10
– Greater Curvature	5
▪ Duodenal	29
– Duodenal bulb	13
– Papilla	4
– Descending part of the duodenum	12
Localization of the peptic ulcer	n = 18
▪ Gastric Ulcer	5
▪ Duodenal Ulcer	13
Forrest classification for gastric and duodenal ulcer bleeding	n = 18
▪ Forrest Ia (spurting)	0
▪ Forrest Ib (oozing)	7
▪ Forrest IIa (visible vessel) + evidence of bleeding	6
▪ Forrest IIb (adherent clot)	4
▪ Forrest IIc (haematin on ulcer base)	1
Mean ulcer diameter 28 mm (Range 15–50 mm)	

Discussion

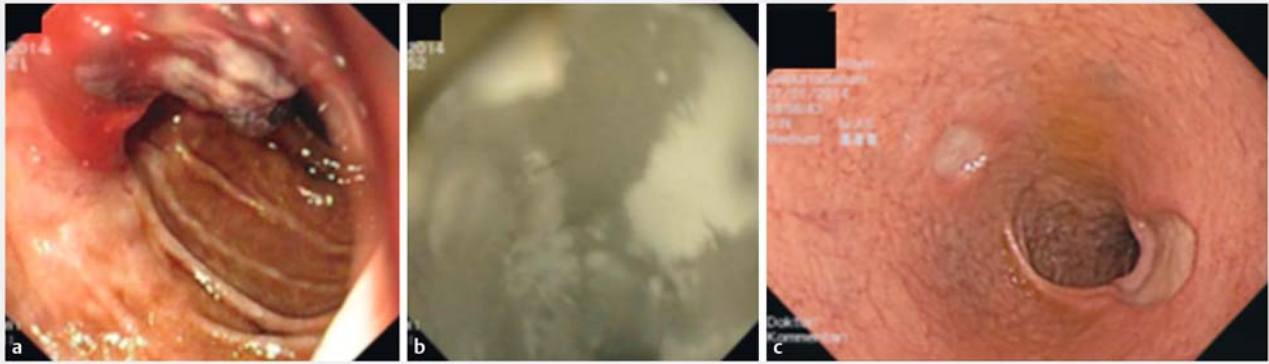
We report our experience with TC-325 for treatment of upper gastrointestinal bleeding in 52 patients over a 38-month period. In our cohort, TC-325 was used as monotherapy for bleeding sources not amenable to standard methods or as “salvage” therapy after failure of endoscopic hemostasis. To our knowledge this study represents one of the largest single-center retrospective series. The study showed excellent applicability and technical success (98.1%) but a high rate of rebleeding (43.1% and 49% on Days 3 and 7, respectively).

The high technical success rate in our study is in accordance with previous studies. Early studies and case series showed a constantly high technical success rate of 93% to 100% in various bleeding sources in the UGIB [5–8, 12, 13] and LGIB [14]. The SEAL-Study was the first multicenter study on TC-325. It included 63 patients and showed a hemostasis rate of 85%. The recently published French GRAPHE-Registry, which included 202 patients, revealed a high technical success rate of >95%, which confirmed results from early studies and case series [15]. Although our results confirm the effectiveness of TC-325 for immediate hemostasis, the high rebleeding rate after TC-325 application is nevertheless concerning. Early reports have shown an overall rebleeding rate of 12% to 15% in mixed cohorts of UGIB and LGIB [6, 7, 10]. If only UGIB was considered, the rebleeding rate was higher. Chen et al. reported a rebleeding rate of 29.4% in patients with high-risk non-variceal UGIB within the first 72 hours and a total of 47% after >72 hours [6]. The GRAPHE-Registry showed a rebleeding rate of 26.7% on Day 8 and 33.5% on Day 30 [15]. Another study from Yau et al. also revealed a rebleeding rate of 39% within 7 days in 19 patients with UGIB [2], meanwhile Dixon et al. reported a 33% rate of rebleeding within 30 days in patients with UGIB [16]. Despite no directly comparable report, our results with a rebleeding rate of 49% within 7 days are grossly comparable to the literature.

One explanation for the high rebleeding rate in our study is the selection of patients. TC-325 was applied only in patients not amenable to conventional endoscopic therapy or as salvage therapy for patients after unsuccessful hemostasis with conventional methods (e.g. peptic ulcers with large diameter and/or difficult anatomy). Fifty-two percent of our patients had a complete Rockall-score >7 and 86.5% had a score >6, indicating the high risk of re-bleeding in our cohort. The mean diameter of the gastroduodenal ulcers in our study was 28 mm (range 15–50 mm) and the majority of the lesions were in the duodenum. In a meta-analysis [17] and the current ESGE guidelines [1], ulcer diameter >20 mm and ulcer location in posterior duodenal wall are considered as predictors for endoscopic treatment failure [1, 17].

TC-325 becomes cohesive and adhesive after contact with blood or other moisture and mainly achieves hemostasis through formation of a mechanical barrier that adheres and covers the bleeding site. In the gastrointestinal tract with its constant peristaltic and mucus production, TC-325 is probably sloughed off the bleeding site within a few hours. When second-look endoscopy after 24 hours was performed, the powder was not visible in any patients in our study. The hemostatic effect of TC-325 thus is not as durable as other hemostasis methods such as the hemoclip.

Compared to other hemostasis topical powder such as EndoClot™, TC-325 confers a higher primary hemostasis rate in UGIB (64–100% [18, 19] vs. 85–98% [2, 6, 7, 10]). The rebleeding rate with EndoClot™, however, was reported to be low at 4.8% to 11%. Ankaferd BloodStopper™ (ABS), a topical hemostasis powder consisting a mixture of 5 different plant species, is available in Turkey and reportedly is effective for various casues of bleeding (e.g. diverticular bleeding [20], radiogenic esopha-



► **Fig. 1** An example of TC-325 application in peptic ulcer. **a** The patient has a Forrest-Ib bleeding from a duodenal ulcer with limited visibility, which was difficult to reach with mechanical hemostasis procedure or adrenalin injection. **b** Thus application of TC-325 was chosen as primary hemostasis method. **c** On second-look endoscopy, 2 duodenal ulcers with clean base (Forrest III) were visible without any further sign of bleeding.

► **Table 3** Overall rebleeding rate and rebleeding rate of the most frequent bleeding etiologies on day 3 and day 7.

	Total cohort	Monotherapy	Salvage therapy	P value
Technical success, n (%)	51/52 (98.1)	23/23 (100)	28/29 (96.6)	1
Overall rebleeding on Day 3, n (%)	22/51 (43.1)	9/23 (39.1)	13/29 (44.8)	0.78
Overall rebleeding on Day 7, n (%)	25/51 (49)	10/23 (43.5)	15/29 (51.7)	0.58
Rebleeding from peptic ulcer on Day 3, n (%)	8/18 (44.4)	3/5 (60)	5/13 (38.5)	0.61
Rebleeding from peptic ulcer on Day 7, n (%)	10/18 (55.5)	3/5 (60)	7/13 (54)	1
Rebleeding from post-interventional ulcer on Day 3, n (%)	7/14 (50)	1/4 (25)	6/10 (60)	0.56
Rebleeding from post-interventional ulcer on Day 7, n (%)	8/14 (57.1)	2/4 (50)	6/10 (60)	1
Rebleeding from tumor bleeding on Day 3, n (%)	2/10 (20)	2/7 (28.6)	0/3	1
Rebleeding from tumor bleeding on Day 7, n (%)	2/10 (20)	2/7 (28.6)	0/3	1

► **Table 4** Causes of mortality.

Cause of mortality	n
Bleeding-related*	2
Sepsis	1
Cerebral Ischemia	1
Aspiration	1
Not stated	1
End-stage liver disease	1
Ventricular fibrillation	1
Total	8

* See further description in text

gitis [21] and dieulafoy's lesions [22]. Larger studies regarding ABS and EndoClot™ are lacking and needed before their widespread use.

In tumor bleeding the application of TC-325 as a hemostasis method is very promising due to its easy non-contact application and large coverage. Given diffuse bleeding and large bleeding area, conventional methods regularly have high bleeding rates ranging from 28% to 49% [23–25]. There were 10 tumor bleeds in this study, all of which had a 100% technical success rate and a clinical success rate of 80%. This is comparable to outcomes previously reported [6, 15, 26]. A prospective study comparing TC-325 and other conventional methods in tumor bleeding is warranted to further validate these findings.

Based on the high rebleeding rates shown in our and other studies and the lack of randomized controlled trials comparing TC-325 with standard endoscopic methods, TC-325 may only be recommended as first-line endoscopic treatment for bleeding sources not amenable to conventional methods or as a salvage therapy. The role of Hemospray for first-line endoscopic treatment for bleeding sources accessible to standard methods as well as the question of whether it should primarily be used as monotherapy or combination therapy remains unclear. Current ESGE-guidelines recommend use of topical-spray (e.g. TC-325) only if bleeding was persistent or in recurrent bleeding [1]. Giv-

en its easy application it may also be used in primary care units with limited experience in hemostasis as a bridge to definitive therapy in a next level care center.

Limitations of our study are the retrospective analysis and the heterogeneity of the patient cohort, as well as the lack of a control group. Furthermore, the results of this study can only be applied to the patient cohort described above and cannot be generalized for all patients with UGIB.

Conclusion

In conclusion, this study demonstrates easy application and high immediate efficacy of TC-325 as a monotherapy in UGIB not amenable to standard endoscopic therapy or as an add-on (“salvage”) therapy after failure of other methods. However, re-bleeding in this high-risk cohort was frequent. Randomized trials are warranted to compare TC-325 with standard endoscopic methods and to further define its role in the therapeutic algorithm.

Competing interests

None

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