The efficacy of Hemospray in patients with upper gastrointestinal bleeding from tumor

Introduction

Tumor-related gastrointestinal bleeding is currently a challenging clinical problem [1]. A retrospective review of upper gastrointestinal bleeding in 55 patients with primary or metastatic gastrointestinal malignancy revealed that up to 20% were related to tumor invading gastrointestinal lumen and causing bleeding [2]. Currently, endoscopy is recommended as the main diagnostic tool to locate the site of bleeding [2,3]. Unfortunately, current endoscopic hemostatic methods including coaptation therapy, argon plasma coagulation, and mechanical hemostasis do not reliably control active bleeding, with rates of successful immediate hemostasis as low as 40% and a significant short-term rebleeding rate (up to 30%) [1,4,5]. Surgery, embolization, and radiotherapy can serve as salvage hemostasis because they are more effective (50%–100%) with lower rates of rebleeding (0–18%) [6–8]. However, a bridging endoscopic therapy is required during resuscitation and stabilization of patients. Hemospray is an inorganic powder not absorbed or metabolized by mucosal tissue. When in contact with blood, its adhesive properties result in a physical barrier that covers the bleeding site. The Hemospray barrier is stable because of the effect of accumulation of clotting factors, and consequently the bleeding point is not exposed to acid, allowing the healing process to continue [9]. Neither luminal nor systemic side effects have been reported with the product [9–11]. Recently, Hemospray has been proposed as a novel way of producing endoscopic hemostasis for active gastrointestinal (gastrointestinal) bleeding, mostly for ulcer bleeding [9]. To date, the study of Hemospray in tumor bleeding is limited as results have not focused on this particular patient group [9–15]. This study aimed to compare the hemostatic efficacy of Hemospray with that of conven-

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Patients and methods: Fourteen patients with active upper gastrointestinal bleeding from tumor were recruited. Hemospray was applied at the bleeding site until hemostasis was achieved. Four patients were excluded because they prematurely received definitive therapy to prevent further bleeding within 48 hours. Another 10 patients from historical control were matched based on the type of gastrointestinal tumors. The 14-day rebleeding rates, length of hospital stay (LOS) and mortality rate at 30-day follow up were assessed.

Results: Baseline characteristics including age, stage of tumor, and Blatchford score did not differ between the two groups. The 14-day rebleeding rate in the Hemospray group was 3 times lower than the control group but not statistically significant (10% vs. 30%; \( P = 0.60 \)). LOS was no different between the 2 groups (28.2 ±21.2 vs. 23.8 ±12.5 days; \( P =0.26 \)). The 30-day mortality rate in the Hemospray group was 3 times lower than that of in the conventional therapy group but not significant (10% vs. 30%, \( P =0.7 \)).

Conclusions: Hemospray is a promising therapy for initial hemostasis in upper gastrointestinal bleeding from tumor because it can achieve hemostasis during the first 14 days, thus potentially allowing sufficient time before appropriate definitive intervention is considered.

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License terms

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tional endoscopic treatment in patients who presented with upper gastrointestinal bleeding from tumor.

Patients and methods

Patients
Between January 2014 and January 2015, patients with a history of upper gastrointestinal bleeding from primary gastrointestinal malignancy or metastasis who presented at the King Chulalongkorn Memorial Hospital were enrolled. The inclusion criteria were as follows: 1) male or female patient, 18 years of age or older; 2) ability to provide written informed consent; and 3) presence of active bleeding from tumor during endoscopy. Patients were excluded if they had received any definitive treatment, such as surgery, embolization or radiation within the first 48 hours (unable to evaluate the hemostatic result of Hemospray). For the control group, the authors retrieved matched cases from the hospital electronic database during the previous 5 years using the key words “upper gastrointestinal bleeding AND cancer”. Tumor location was used as the criterion for matching and the patients were further divided into those with upper gastrointestinal tumor and those with hepatico-pancreatobiliary tumor corresponding to the Hemospray group.

In both groups, patient gender, age, type and stage of malignancy, Blatchford score, amount of blood transfusions, number of endoscopies needed, and requirement for additional interventions including endoscopy, surgery, adjuvant embolization, and radiotherapy during admission were analyzed. The 14-day rebleeding rates, lengths of hospital stay (LOS), and mortality rate at 30-day follow up were assessed. The study protocol and consent form were approved by the Chulalongkorn University Institutional Review Board (No.092/58).

Statistical analysis
For numerical variables, the results were expressed as a mean±SD, whereas other quantitative variables are expressed as percentages. Continuous variables were compared by the student t test. Discontinuous variables were compared by the chi-square (x2) test. SPSS version 17.0 (SPSS (Thailand) Co., Ltd., Bangkok, Thailand) for Windows systems was used for statistical analysis. Differences were considered significant at the level of 0.05.

Results
During the study period, 14 patients experienced blood oozing from tumor and in all cases, Hemospray was used as the first-line hemostatic therapy. Four of those patients were excluded because they received either chemoembolization or radiation therapy within 72 hours after Hemospray, although there was no evidence of rebleeding. Subsequently, 10 patients were eligible for analysis.

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Hemospray or TC-325 (Cook Medical, Winston-Salem, North Carolina, USA) was used as the only hemostatic method with the maximum dose of 20g (1 cartridge).

Medical resuscitation described elsewhere for upper gastrointestinal bleeding [16] was initiated in all eligible patients. Following medical resuscitation, therapeutic esophagogastroduodenoscopy (EGD) was performed by an experienced endoscopist (RP) certified in Hemospray endoscopic hemostasis. If tumor bleeding was identified, Hemospray tube was applied through a 10-Fr catheter in which the powder was sprayed onto the bleeding site until hemostasis was achieved, but no more than 1 cartridge was used. Successful initial hemostasis was defined as no further active bleeding seen at least 5 minutes before withdrawing an endoscope after Hemospray was applied (Fig. 1).

Then standard post-endoscopic care with a 72-hr proton pump inhibitor infusion was prescribed in all patients [16]. Re-EGD was done in patients with suspected rebleeding, which was defined as: 1) 3% of more drop in hematocrit level even after adequate blood transfusion; and 2) new development of hematemesis or hematochezia or melena.

Fig. 1 Pictures of active tumor bleeding (a) and post Hemospray (b).
of 10 patients in the Hemospray group did not undergo additional intervention during the first 14 days. Although there was no rebleeding, on Day 12, 1 patient with bleeding gastric cancer who had previously been scheduled for elective surgery underwent a partial gastrectomy (Table 2). Three of 10 patients (30%) in the control group re-bleed and required a rescue intervention during the first 14 days and another 4 patients (40%) underwent additional intervention according to a prescheduled plan (Table 3). In all 3 patients, rebleeding occurred quite early (within 48 hours) and results of treatment with Hemospray and with conventional endoscopic treatment.

Table 2 Detailed course of patients in the Hemospray group.

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<th>Rescue treatment</th>
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CA, carcinoma; CCA, cholangiocarcinoma; HCC: hepatocellular carcinoma; PRC, packed red cells; LOS, length of hospital stay; N/A, no data available

Discussion

Because most incidents of active tumor bleeding are difficult to control with conventional endoscopic hemostatic techniques such as coaptation, hemoclipping, and band ligation, it is necessary to use bridging therapy. Hemospray is well suited for treatment of upper gastrointestinal bleeding from tumor because it provides temporary hemostasis while allowing time to schedule a patient for more definitive treatment such as surgery, angiographic embolization, or radiation. The mechanism of tumor bleeding is explained by erosion of the raw surface of a malignant lesion. In addition, acidic content from the stomach cannot promote more bleeding because it dissolves clot and possibly digests tumor tissue that lacks a barrier of mucous and epithelium [4]. Although conventional endoscopic hemostatic techniques can provide immediate control for some tumor bleeding, it tends to recur in a short period of time [1, 4, 5]. It is possible that the conventional endoscopic treatments may fail to protect tumor tissue from digestion. In addition, if coaptation is selected, tumor necrosis may progress because of the effect of heat [12]. In contrast, Hemospray treatment can prevent tumor bleeding by providing immediate hemostasis and the powder that remains on the surface for a period of time may protect the tumor tissue from further erosion by gastric acid. In recent years, multiple studies have been conducted to prove the efficacy of Hemospray for many different types of gastrointestinal bleeding [9, 10, 13–15] but only 1 study focused on tumor-related gastrointestinal bleeding is available [10]. In 2012, Chen et al. reported a case series of upper gastrointestinal bleeding from different types of tumors which was successfully controlled by Hemospray [10]. In their series, there were 3 gastric cancers, 1 pancreatic cancer invading duodenum, and 1 metastatic breast cancer invading duodenum. Four of 5 patients (80%) subsequently received chemotherapy or radiotherapy within 5 days after initial hemostasis with Hemospray although there was no evidence of rebleeding during the 5-day observation [10]. Based on experience in a very limited number of patients, the authors concluded that Hemospray may be useful for both immediate hemostasis and as a bridging treatment for further adjuvant therapy. To our knowledge, this is the first study that compares the hemostatic effect of Hemospray with conventional treatment for upper gastrointestinal bleeding from tumor. To demonstrate the longer hemostatic effect of Hemospray in bleeding tumor, the current
study excluded patients who received definitive treatment within 72 hours. We could not, however, demonstrate statistical significance given the limited number of cohorts but we did demonstrate a trend toward lower rates of recurrent bleeding in 14 days (10% vs. 30%, P=0.6) and a lower mortality rate at 30 days (10% vs. 30%, P=0.7). In addition, no additional treatment was needed after Hemospray application in all except 1 patient during hospitalization whereas 30% of patients in the control group required additional treatment.

The limitation of the present study is a rather small number of cases, which made it impossible to make a pair match in other parameters apart from the location of tumor. In addition, we believe that more patients would have allowed the results to rise to statistical significance, as we showed a large gap in outcomes between the 2 groups. Furthermore, no patients had bleeding from a lower gastrointestinal tract tumor. Because little acid is involved in rebleeding from such tumors, the efficacy of Hemospray in those patients needs to be demonstrated and its differing mechanism of action in that setting explained.

Conclusions

Hemospray is a promising therapy for initial hemostasis in upper gastrointestinal bleeding from tumor because it can achieve hemostasis during the first 14 days, which may allow sufficient time before consideration of appropriate additional intervention.

Competing interests: None

References