A new device for endoscopic band ligation for colorectal diverticular bleeding

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ABSTRACT

Background and study aims Endoscopic band ligation (EBL) has been reported to be effective for colorectal diverticular bleeding. However, current EBL devices have some limitations, and we have developed a novel EBL device.

Methods This novel EBL device has a tapered hood attached to the tip of the endoscope and an outer cylinder that loads two EBL bands. Twelve EBL procedures were performed in a live porcine model alternately using a conventional EBL device (Group C) and the novel EBL device (Group N).

Results EBL was successful in all cases in both groups. There were no cases of perforation in the 10 days after EBL. After ligation, the mean major axis of the ridge was significantly larger in Group N than Group C (9.7 ± 1.4 mm vs 7.2 ± 1.4 mm, P = 0.011). Pathological examination revealed disruption of the muscularis propria at four of the six ligation sites in Group C and at five of the six ligation sites in Group N.

Conclusions Using this novel EBL device, it was possible to perform multiple ligation procedures in succession with a good field of view. No perforation was observed, but disruption of the muscularis propria was observed at approximately three-quarters of the ligation sites pathologically.

Introduction

Colorectal diverticular bleeding is now being encountered with increasing frequency in daily clinical practice, mainly because of population aging and the increasing use of antithrombotic agents. However, the endoscopic field of view is poor due to large amounts of bloody stool, and it is sometimes difficult to identify the bleeding source. Even if a bleeding diverticulum can be found, the exposed blood vessels cannot always be observed directly. Therefore, it is difficult to stop bleeding using a clip, and rebleeding after hemostasis remains a major problem [1–3].

Various hemostatic methods have been developed for diverticular bleeding, and endoscopic ligation, which has been used for hemostasis in patients with esophageal varices, has been reported to be effective [4]. Unlike when using a clip, bleeding can be stopped by suctioning, reversing, and ligating the bleeding diverticulum even if the exposed blood vessels cannot be observed directly. However, the field of view is poor when a conventional EBL device is used. Furthermore, the current EBL devices are single-use, and if ligation fails, it is necessary to remove the endoscope and attach an O-ring to the tip. Moreover, complete aspiration is not always possible when ligating a bleeding diverticulum. In collaboration with Asahi Intecc Co. Ltd (Aichi, Japan), we have developed a novel EBL device that allows multiple ligations to be performed in succession with a better field of view and a stronger suction force.
Methods

Description

The novel EBL device (Endoligar) consists of a tapered conical hood and an outer cylinder with an O-ring for ligation on the outside (Fig. 1). The tip of the hood is tapered for easier suction. Two O-rings are loaded, and it is possible to fire them separately depending on the amount of fluid injected from the outside (Fig. 2). The outer cylinder loaded with the O-ring is located behind the hood, so it does not interfere with the endoscopic field of view. The maximum diameter of this novel EBL device is 19 mm, and the tip diameter is 9 mm.

Experimental method

We investigated the feasibility of this novel EBL device in a porcine model at the Fukushima Medical Device Development Center. A 40-kg female pig was used. After intramuscular injection of a mixed solution of midazolam and medetomidine, tracheal intubation was performed and the experiment was conducted under general anesthesia. The experiment was approved by the Animal Experiment Committee of Toranomon Hospital.

Twelve ligation procedures were performed in the normal colonic mucosa alternately using the novel EBL device (Group N) and a conventional EBL device (Group C) (Sumitomo Bakelite Co. Ltd., Tokyo, Japan) between 30 cm and 16 cm from the anal margin. The distance from the anal margin to the ligation site was the same in both groups. Six ligation procedures were performed using three novel EBL devices (Video 1). In both groups, the ligation was performed after fully aspirating the target colonic mucosa. In Group N, a 10-mL syringe was used to perform a first injection of 5 mL of distilled water and a second injection of 10 mL of distilled water. In Group C, air was injected using a 5-mL syringe. A successful procedure was defined as swelling of the aspirated colonic mucosa and the residue of an O-ring at its base.

The pig was fasted on the day of the experiment with resumption of a normal diet on the following day. Ten days after the ligation procedure, the pig was euthanized after endoscopic observation under general anesthesia. The abdomen was then opened and the large intestine was removed for pathological examination. Each ligation site in the resected specimen was identified, and a residual O-ring was confirmed. Next, the size of the ridge at each ligation site was measured using a caliper. After fixing with 10% formalin solution, hema-
toxylin-eosin staining and Masson’s trichrome staining were performed (Fig. 3). Two pathologists determined the presence or absence of disruption of the muscularis mucosae, submucosal fibrosis, and disruption of the muscularis propria.

**Statistical analysis**

Data were analyzed using the Fisher’s test and Mann-Whitney U-test. A $P<0.05$ was considered significant. All statistical analyses were performed using SPSS version 20 (SPSS IBM statistics).

**Results**

The 12 EBL procedures were performed without any complications. In Group N, it was possible to fire the two O-rings separately, and all procedures were successful. No adverse events, such as fever or signs of abdominal pain, were observed during the 10-day observation, and food intake was good. The body weight was 42.2 kg on the day of ligation and 43.8 kg at the time of necropsy.

Endoscopic observation of the EBL site revealed that all O-rings remained (Table 1). No perforation was observed. The average major axis of the ridge was significantly larger in Group N than in Group C (9.7 ± 1.4 mm vs 7.2 ± 1.4 mm) ($P=0.011$).

Pathological examination revealed disruption of the muscularis mucosae at all ligation sites (Table 2). Fibrosis of the submucosal layer was also observed at all sites. Disruption of the muscularis propria was observed at five of six sites in Group N and at four of six sites in Group C. And the length of the disruption of the muscularis propria was 6.8 ± 3.5 mm in Group N and was 3.9 ± 3.6 mm in Group C (Fig. 4). Average length of the disruption of the muscularis propria did not significantly differ between the groups ($P=0.18$).

**Discussion**

We developed a novel EBL device for diverticular bleeding, and conducted a comparative experiment with a conventional EBL device. Using this novel EBL device, ligation procedures can be performed in succession with a good field of view. And the mucosal ridge after ligation by the novel EBL device was significantly larger compared to the conventional EBL device.

Diverticular bleeding is a common cause of acute lower gastrointestinal hemorrhage. It typically has a rapid onset without abdominal pain and can be distinguished from other acute lower gastrointestinal bleeding disorders, such as ischemic colitis and rectal ulcer. Diagnosis is more reliable when enhanced computed tomography (CT) is performed and outflow of contrast medium is observed from the blood vessel. Enhanced CT is useful for not only diagnosing diverticular bleeding but also identifying the bleeding site, which is usually in the ascending or sigmoid colon. An endoscopic finding of stigmata of recent hemorrhage involving a diverticulum is an indication for endoscopic treatment. However, it may be difficult to identify the diverticulum that is the source of the bleeding endoscopically because of the presence of blood and residual stool [5].

Various treatments have been used for diverticular bleeding. In the past, clipping was often performed but the rebleeding rate was high because of the difficulties involved in identifying the exposed blood vessels in the diverticulum [6–8]. More recently, EBL that ligates the entire diverticulum has been preferred, but its effectiveness varies depending on the site. It has been reported that EBL has a higher success rate with less risk of rebleeding in the ascending colon than in the sigmoid colon.

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**Table 1** Technical results of the study.

|                  | Group C | Group N | N.S. |
|------------------|---------|---------|------|
| n                | 6       | 6       |      |
| Success rate of ligation (%) | 100 (6/6) | 100 (6/6) |      |
| Perforation (%)  | 0 (0/6) | 0 (0/6) | N.S. |
| Residue of the O-ring 10 days after EBL (%) | 100 (6/6) | 100 (6/6) | N.S. |
| Average size of the ridge (mm ± SD) | 7.2 ± 1.4 | 9.7 ± 1.4 | $P=0.011$ |

EBL, endoscopic band ligation; SD, standard deviation.

**Table 2** Pathological results of the study.

|                  | Group C | Group N | N.S. |
|------------------|---------|---------|------|
| n                | 6       | 6       |      |
| Disruption of the muscularis mucosae (%) | 100 (6/6) | 100 (6/6) |      |
| Fibrosis of the submucosal layer (%) | 100 (6/6) | 100 (6/6) | N.S. |
| Disruption of the muscularis propria (%) | 66.7 (4/6) | 83.3 (5/6) | N.S. |
| Length of the disruption of the muscularis propria (mm ± SD) | 3.9 ± 3.6 | 6.8 ± 3.5 | $P=0.18$ |

SD, standard deviation.
The diverticulum may not be reversed even if the diverticulum is aspirated during EBL. If the O-ring is fired without the diverticulum being inverted by firm suction, ligation might fail. When ligation is inadequate using a conventional device, the endoscope needs to be removed and then reinserted after attachment of the O-ring. However, when using the novel EBL device, two O-rings are loaded and it is possible to attempt ligation again even if the first attempt has failed. Moreover, unlike a conventional EBL device, the tip of the hood is longer and tapered, so the suction power may be higher. In this experiment, the mucosal ridge after ligation was larger in Group N, suggesting that it may be effective for diverticula that cannot be aspirated by a conventional EBL device. A device that can be used to perform multiple ligation procedures for esophageal varices is available. Shiratori et al. have compared that device with the EBL device we used in our conventional group and have reported a similar postoperative course and pathological findings [15]. However, Shiratori et al. encountered problems, including a poor field of view and a lengthy insertion time.

This research and the device used have some limitations. First, the outer diameter of this novel EBL device is 19 mm, which is relatively large, and resistance may be encountered when it is inserted deep into the sigmoid colon. The sigmoid colon with diverticula can be stiff and this can be a major clinical problem. Therefore, further studies are needed to assess its safety in clinical practice. Second, the EBL procedures were performed in normal mucosa between the rectum and sigmoid colon, and the results may be different when performed for a bleeding diverticulum. Even in a normal colon wall, we confirmed disruption of the muscularis propria at five of six ligation sites in Group N and at four of six sites in Group C. In the experiment by Shiratori et al., the muscle layer was ligated at a high rate, but no perforation was observed. However, these studies were small, and large-scale prospective studies of the safety of these devices are needed in the future.

Conclusions

In conclusion, we developed a novel EBL device for diverticular bleeding. Our experimental findings suggest that the suction power of the novel EBL device may be superior to that of a conventional EBL system. Despite a high rate of disruption of the muscularis propria, there were no complications, such as perforation or fever. This novel device has the potential to be useful for treatment of diverticular bleeding.

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Competing interests

The prototype of the device described in this paper was manufactured by Asahi Intecc Co. Ltd. Drs. Kikuchi and Hayasaka and Asahi Intecc have signed a consulting contract for development of the device.

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