A new endoscopic closure method for gastric mucosal defects: feasibility of endoscopic hand suturing in an ex vivo porcine model (with video)

Authors
Osamu Goto1, Motoki Sasaki1, Hiroyuki Ishii1, Joichiro Horii1, Toshio Uraoka1, Hiroya Takeuchi2, Yuko Kitagawa1, Naohisa Yahagi1

Background and study aims: More secure endoscopic closure techniques for iatrogenic gastric defects are required for safe endoscopic surgery. We developed a novel endoscopic suturing method, endoscopic hand suturing (EHS), of gastric mucosal defects and determined its feasibility and efficacy ex vivo.

Materials and methods: We created 24 mucosal defects (each 2 cm in diameter) by endoscopic submucosal dissection. The following three techniques were tested: EHS with a 3–0 barbed suture that was grasped with biopsy forceps (n=6) or a prototype through-the-scope needle holder (n=6) by endoscopy, looping with endoloops (n=6) by endoscopy, and clipping with hemoclips (n=6) by hand. The mucosal edges were attached to each other at three points. The closure strength was compared among the three groups, and the procedural duration was compared between the EHS and looping groups.

Results: All 12 lesions were completely closed by EHS. The median strength of the closure, measured with a spring scale, was significantly greater in the EHS group (0.74 kg) than in the looping group (0.33 kg, P=0.0012) or clipping group (0.07 kg, P=0.0009). The median procedural duration did not significantly differ between the EHS and looping groups (19.7 vs. 19.8 minutes, P=0.0009). The use of the needle holder significantly reduced the procedural duration compared with the biopsy forceps.

Conclusion: Mucosal defects can be firmly closed with EHS, which may be helpful for establishing a safer and more secure endoscopic surgery.

Introduction

Endoscopic submucosal dissection (ESD) is accepted, particularly in Asian countries, as a minimally invasive endoscopic approach to curing early stage cancer without organ resection [1–5]. Delayed bleeding is a major complication after gastric ESD that occurs in approximately 5% of patients [6–8]. Delayed perforation can also develop after successful resection [9]. Although the definitive causes of these complications are unknown, the exposure of an iatrogenic mucosal defect to gastric acid may be an important factor. When such defects are created in patients at risk for complications, closure of the defect with hemoclips or a detachable snare is sometimes attempted [10–12]. Suturing with a needle and suture thread can yield a secure closure when the alimentary tract is surgically anastomosed. If it is possible to firmly close an iatrogenic defect by suturing the mucosa endoscopically, delayed bleeding after ESD or anastomotic leakage after endoscopic translumen-
amount of indigo carmine was injected to create a submucosal cushion. A GIF-2TQ260M multibending scope with two working channels (Olympus Medical Systems Corp., Tokyo, Japan) was used for ESD and endoscopic closure. The closure methods described below were then applied by one endoscopist and one assistant (Fig. 1).

**Endoscopic hand suturing**

Defects were closed with 15-cm, 3–0 absorbable barbed V-Loc 180 (VLOCL0604) sutures (Covidien, Mansfield, MA, USA) (Fig. 2a). This type of unidirectional, knotless suture was originally developed to facilitate skin suturing, and it has subsequently been applied to gastrointestinal surgeries [13–15]. The short barbs are oriented along the suture to face away from the needle, preventing the suture from sliding backward after the tissues have been tightened. A loop on the tail and the continuous small part of the suture without barbs was cut and anchored with an EZ clip (HX-610–90; Olympus). In closing six of the lesions, the needle was grasped with biopsy forceps (FB-220U; Olympus). A 10-mm SD-210L-10 electrocautery snare (Olympus) was inserted into a separate working channel to tightly grasp the neck of the biopsy forceps for stabilization (Fig. 2b). For the next six lesions, a prototype through-the-scope needle holder (Olympus) was used (Fig. 2c–e) [16,17]. This device has one swinging lateral jaw and one fixed straight jaw, both of which firmly close when the handle slider is grasped. A button on the handle unlocks and opens the jaws. Irrespective of the shape of the endoscope, the tip of the device can simultaneously rotate as the handle rotates, which accelerates suturing, even inside the stomach. To insert the needle and string gently through the overtube, we grasped the tail of the needle and the suture near the anchor clip with biopsy forceps. We grasped the tail of the needle with the needle holder and placed the anchor clip into another working channel in a retrograde manner. The tip of the needle was ori...
iented toward the endoscope to avoid potential injury to the mu-
cosa. The needle was released in the stomach and grasped again,
and then the mucosal edges and part of the submucosa were con-
tinuously sutured to each other longitudinally, from the distal to
the proximal sides of the endoscope. After some amount of suture-
ing, the slack part of the suture was tightened by directly grab-
bining and pulling it with biopsy forceps or the needle holder. After
three sutures had been made, one mucosal edge slightly distal to
the third suturing point was sutured as a lock. The excess suture
and the needle were cut away with a loop cutter (FS-5L-1; Olym-
pus), the end of the suture was clipped to the mucosa, and the
needle was removed through the overtube by grasping the tip of
the needle with biopsy forceps or the needle holder (Fig. 1a).

Video 1 shows EHS with the needle holder.

**Looping**

An MAJ-340 endoloop, 20 mm in diameter (Olympus), was
opened above a mucosal defect and placed at each mucosal edge
with an EZ clip (HX-610–90 or HX-610–90L; Olympus) [11, 12].
Thereafter, the loop was slowly closed by heading a locking unit.
The loop was firmly closed and detached. The excess loop was cut
and removed with a loop cutter. We spaced three loops along the
line to divide the closure seam into four equidistant segments
(Fig. 1b).

**Clipping**

We applied three HX-610–90 or HX-610–90L EZ clips per lesion.
To stitch the mucosal edges to each other without slippage, the
procedure was performed by hand after the stomach had been
opened as described below. Clips were spaced along the line to di-
vide the closure seam into four equidistant segments (Fig. 1c).

We closed six experimentally created mucosal defects under each
condition (EHS with biopsy forceps, EHS with the needle holder,
looping, and clipping) (Fig. 3, Fig. 4a–c). We created two arti-
tficial lesions in the gastric corpus of nine stomachs for both the
EHS and looping groups. For a forward approach, one lesion was
located at the posterior wall, and for a retroflex approach, one le-
sion was positioned at the lesser curvature. An additional six le-
sions in total were created for the clipping group by using the re-
mainder of the two stomachs and one additional whole stomach.
Endoscopic closure of the lesion on the lesser curvature was initi-
ally achieved by a retroflex approach and repeated again on the
posterior wall to avoid potential mechanical injury to the sutured
lesion on the posterior wall while the endoscope was handled in
a retroflex approach. After closure by endoscopy, the stomach
was removed from the unit, cut along the greater curvature, and
opened with the mucosal surface facing upward to assess the
strength of closure. The clipping method was performed on two
of the opened stomachs. We also directly cut one additional
stomach and opened it for the clipping method after the creation
of four mucosal defects.

**Measuring outcomes and statistical parameters**

The primary outcome measured in the three groups was the
strength of closure, which was assessed with a digital spring scale
(Fig. 4d). The scale was attached to an 18-gauge needle that
was sewn parallel to the suture line on one side of the mucosa
and then pulled to the opposite side of the suture line while the
other side of the mucosa was held by hand (Fig. 4e). The weight
(in kilograms) indicated on the monitor when the closure
collapsed was taken as a surrogate marker of closure strength.
The secondary outcome measured was the procedural duration,
which we defined as the time elapsed from the first insertion of
the needle to the placement of a clip at the end of the suture in
the EHS group, and from the first clipping step to place the first
loop at the mucosal edge to cutting the excess of the third loop
in the looping group. Differences in closure strength and proce-
dural duration between the grasping devices used (biopsy for-
ceps or needle holder) and the approach used (forward or retro-
flex) were also analyzed in the EHS group.

Strength and procedural duration were analyzed with the
Mann-Whitney U test. Statistical significance was set at a P value
of <0.05. All data were statistically analyzed with JMP version
9.0 software (SAS Institute, Cary, NC, USA).

**Results**

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All 24 procedures were completed, including EHS procedures
with biopsy forceps (6 of 6, 100%) or the needle holder (6 of 6,
100%). We measured the median strength in the three groups
(Fig. 5a) and found that closures were significantly tighter in

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**Fig. 3** Endoscopic images of the closure methods. a Endoscopic hand suturing with biopsy forceps. b Endoscopic hand suturing with a prototype needle holder. c Looping.

**Fig. 4** a–e Endoscopic suturing for mucosal defects with loop clipping and the prototype needle holder.

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**Fig. 5** a Endoscopic hand suturing with biopsy forceps. b Endoscopic hand suturing with the needle holder.
the EHS group (0.74 kg [range 0.49 – 1.16 kg]) than in the looping group (0.33 kg [range 0.21 – 0.54 kg], P = 0.0012) or clipping group (0.07 kg [range 0.02 – 0.31 kg], P = 0.0009). Moreover, the strength was significantly weaker in the clipping group than in the looping group (P = 0.0200).

The median procedural duration did not significantly differ between the EHS and looping groups (19.7 minutes [range 12.5 – 27.1 minutes] vs. 19.8 minutes [range 15.5 – 37.4 minutes], P = 1.0000) (Fig. 5b). The median procedural duration for stitching in the EHS group (excluding time required for locking, cutting, and clipping the end of the suture) was 13.6 minutes (range 7.9 – 20.2 minutes), indicating a median time of 4.5 minutes (range 2.6 – 6.7 minutes) per stitch.

We found no significant difference in median strength between EHS with biopsy forceps or with the needle holder (0.77 kg [range 0.63 – 1.02 kg] vs. 0.74 kg [range 0.49 – 1.16 kg], P = 1.0000), whereas the median procedural duration was significantly shorter with the needle holder than with the biopsy forceps (17.0 minutes [range 12.5 – 21.2 minutes] vs. 22.7 minutes [range 16.8 – 27.1 minute], P = 0.0161) (Fig. 6). The median stitching durations with the biopsy forceps and the needle holder were 16.4 minutes (range 11.3 – 20.2 minutes) and 10.3 minutes (range 7.9 – 17.1 minutes), respectively. The time required per stitch tended to be shorter with the needle holder than with biopsy forceps, but the difference was not statistically significant (median 3.4 minutes [range 2.6 – 5.7 minutes] vs. 5.5 minutes [range 3.8 – 6.7 minutes], P = 0.0538). The median strength and median procedural duration did not significantly differ between the forward and retroflex approaches in the EHS group (0.72 kg [range 0.60 – 1.16 kg] vs. 0.79 kg [range 0.49 – 1.06 kg], P = 0.6884 and 21.6 minutes [range 12.5 – 27.0 minutes] vs. 17.5 minutes [range 12.8 – 27.1 minutes], P = 0.6884, respectively) (Fig. 7).

Discussion

Using an ex vivo porcine model, we found that EHS was feasible and offered firmer closure than conventional looping and clipping methods. Based on our findings, we speculate that EHS would make mucosal defect closure more secure than the other two methods.

Clipping is the most popular method in clinical practice because it is convenient and simple, particularly for patients at a high risk for delayed bleeding due to antiplatelet or antithrombotic medications [10]. However, the findings presented here imply that the mucosal edges attached by clipping are too fragile to stick together for long periods of time. Looping can be used particularly for large lesions that cannot be closed by clipping [11,12], although this method is somewhat tedious and more time-consuming.

Here, we found that looping was stronger than clipping, but EHS was more secure and did not take any longer than looping. Therefore, our data suggest that EHS is the most reliable of the three methods tested. Although in practice looping is accompanied by clipping, this combined process is still predicted to be weaker than EHS based on the data presented in this study. Because clipping is far weaker than looping, it would not significantly enhance the strength of looping.

In the looping and clipping groups, closure sites collapsed when the clips became detached as the mucosal edges parted. In contrast, closure sites dehisced when sutures in the EHS group tore
the mucosa. This finding suggests that sites closed with EHS will
remain intact unless affected by a strong force that is powerful e-
nough to tear the mucosa, whereas mucosal sites closed by loop-
ing or clipping may easily dehisce in the presence of less powerful
forces, such as food movement or peristalsis.

Many reports have addressed the management of delayed bleed-
ing, and an evidenced-based consensus is gradually forming [6
– 8, 18 – 20]; however, no definitive methodology can yet effective-
ly prevent delayed bleeding. Furthermore, delayed perforation
occasionally occur after a successful procedure [9]. The pre-
cise cause of delayed perforation is unknown, but excessive ther-
mal injury to the surface of a mucosal defect may be a risk factor.
Exposure of a mucosal defect to gastric acid after ESD can also be
a risk factor for delayed bleeding and delayed perforation be-
cause gastric acid slows the gastric ulcer healing process and
stimulates nonbleeding visible vessels or nonperforated thin
walls in an iatrogenic mucosal defect. EHS should prevent such
complications by preventing the exposure of mucosal defects to
gastric acid. Practically, antisecretory medicines, which keep the
risk for delayed bleeding low, are used to accelerate the healing of
mucosal defects. Therefore, EHS would be more effective in pa-
tients who have a high risk for delayed bleeding, such as those
who are taking antithrombotic agents, have a coagulopathy or
liver cirrhosis, or are being treated for chronic renal failure by he-
modialysis.

Over-the-scope clips [21] and the OverStitch Endoscopic Suturing
System (Apollo Endosurgery, Austin, TX, USA) [22], which were
principally developed to close transmural defects, can also effect-
ively close mucosal defects. A previous study showed compar-
able strength when wall defects were closed with over-the-scope
clips or surgical suturing [23]. The Eagle Claw (Olympus) [24] and
Flexible Endostitch (Covidien) [25] are also promising suturing
deVICES for wall closure, but the efficacy of mucosal closure with
these devices is unknown, and a comparison with EHS is warran-
ted. On the contrary, EHS may have potential application in the
treatment of perforation or anastomosis. Although full-thickness
suturing by EHS would be more difficult than mucosal suturing
because of collapse of the stomach during the procedure, it might
be worth trying in the future.

Our study has some limitations. EHS requires a certain amount of
suturing time, a skillful hand for endoscope maneuvering, and
some experience with the technique. In addition, extant needles
and sutures are not intended for endoscopic suturing of the mu-
cosa. Other limitations of our study are as follows: a relatively
small number of lesions closed per technique, the use of hand
clipping instead of endoclipping, and the use of porcine stomachs
ex vivo. The findings of this study should be considered prelimi-
nary at best. Thus, an in vivo porcine trial that included a histologic
assessment of healed sutured lesions would be desirable to con-

Fig. 5 Comparison of closure strength and procedural duration. a Endo-
scopic hand suturing is significantly stronger than looping and clipping.
b Procedural duration does not significantly differ between the endoscopic
hand suturing and looping groups.

Fig. 6 Comparison of closure strength and procedural duration with dif-
ferent grasping devices. a A prototype needle holder can maintain strength
at suture site. b Procedural duration is significantly shorter when a needle
holder is used instead of biopsy forceps.
firm the technical feasibility and safety of the EHS method. If successful, the porcine trial should be followed by a clinical trial to investigate the clinical benefits of EHS.

In conclusion, we have demonstrated the feasibility and utility of EHS in this ex vivo study. EHS may more effectively prevent complications after gastric ESD than current measures and may be more helpful in establishing a safe and secure endoscopic surgery. A future in vivo animal study and a clinical trial are desired.

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**References**
