Measurement of intragastric pressure: an objective method to ascertain whether gastric wall extension is sufficient for assessment of the non-extension sign*

ABSTRACT

Background and study aims The optimal intragastric pressure (IP) for strong gastric wall extension is unclear. We aimed to develop an accurate method to measure IP using endoscopy and determine the pressure required for strong gastric wall extension.

Methods An in vitro experiment using an endoscope with a rubber attached at its tip was conducted. The process of inserting the pressure measurement probe into the forceps channel was skipped, and the tube of the pressure measurement device was directly connected to the forceps channel. In vivo, the pressure in 51 consecutive patients at the time of strong gastric wall extension was measured. Strong extension of the gastric wall was defined as when the folds in the greater curvature were flattened as a result of sufficient extension of the gastric wall by insufflated air during upper gastrointestinal endoscopy. The IP at that time was measured.

Results In vitro, 20 mL of tap water was injected once into the forceps channel and then aspirated for 10 seconds. Pressure measurement after irrigation of the forceps channel as well as the measurement by inserting the probe procedure were accurately performed. In vivo, among the 51 included patients, the mean IP (range) was 14.7 mmHg (10–23). Strong extension of the gastric wall was obtained in 96.1 % of patients when the IP was 20 mmHg.

Conclusions We developed an accurate method to measure IP using upper gastrointestinal endoscopy. Strong extension of the gastric wall was obtained in almost all patients when the IP was 20 mmHg.

Introduction

During upper gastrointestinal endoscopy, the gastric wall can be easily extended by insufflated air. Sufficient extension of the gastric wall is important for endoscopic screening because insufficient extension of the folds in the greater curvature of the gastric corpus can result in blind spots. Conversely, suffi-
cient extension of the gastric wall leads to flattening of the folds in the greater curvature of the gastric corpus and disappearance of blind spots, thus enabling the detection of small and flat lesions [1]. In addition, when performing a preoperative examination, evaluation of the invasion depth of early gastric cancer is crucial to determine the indications for endoscopic treatment. We previously reported that the non-extension sign on endoscopy was useful for the diagnosis of submucosal invasive cancer [2, 3]. We also reported that strong extension of the gastric wall is required to assess the presence or absence of the non-extension sign [2, 3].

However, the optimal intragastric pressure (IP) for strong extension of the gastric wall is not clear. Therefore, the objectives of this study were to develop an easy and accurate method to measure IP through an in vitro study and to determine the IP required for strong extension of the gastric wall through an in vivo study. The time of strong gastric wall extension was defined as the time when the gastric wall was sufficiently extended by insufflated air to the extent that the folds in the greater curvature of the gastric corpus were flattened during an upper gastrointestinal endoscopy. The measurement was performed using the method developed in the in vitro study.

Methods
Part 1: Development of a method to measure intragastric pressure: an in vitro study
A rubber balloon was attached to the tip of an endoscope, and the balloon was inflated using a cuff pressure manometer (Fig. 1). After setting the pressure, the following experiments were performed (Fig. 2). It is reported that IP can be measured using an endoscope in which the probe was inserted into the forceps channel and advanced out of the tip of the endoscope [4]. However, it takes time and additional effort to insert the pressure measurement probe into the forceps channel during endoscopic examination. To enable the measurement to be easily performed in a shorter time, the tube of the pressure measurement device was directly connected to the forceps channel hole (Fig. 2). The pressure was measured using a pressure transducer AP-100 (Asahi Biomed Co. Ltd, Tokyo, Japan) connected to the forceps hole of the endoscope.

To apply this method in vivo, the following points are to be considered. In actual endoscopic examination, gastric juice with mucus and bubbles are aspirated from the forceps channel before starting the observation in order to optimize the examination. Therefore, if mucus is retained in the forceps channel of the endoscope, accurate measurement of IP may be difficult. Therefore, we injected 2% lidocaine jelly (Xylocaine Jelly; Aspen Japan, Tokyo, Japan) as a substitute for mucus, into the forceps channel of the endoscope. Thereafter, we performed an in vitro study under the following conditions in order to explore the most accurate method to measure the IP.

Experiment 1: The measurement was performed under a condition in which nothing was injected into the forceps channel.

Experiment 2: The measurement was performed under a condition in which lidocaine jelly was injected into the forceps channel.

Experiment 3: The measurement was performed under a condition in which lidocaine jelly was injected into the forceps channel and then 20 mL of tap water was injected once for irrigation.

Experiment 4: The measurement was performed under a condition in which lidocaine jelly was injected into the forceps channel and then 20 mL of tap water was injected once for irrigation; the tap water retained in the forceps channel was aspirated for 10 sec for further irrigation.

Each measurement was performed as follows. The intraballoon pressure was measured for 5 sec with a recording interval of 0.5 sec, and the mean pressure was calculated. The intraballoon pressure at the tip of the endoscope was set at 10 mmHg, 20 mmHg, 30 mmHg, and 40 mmHg using a cuff pressure manometer (Fig. 1). The pressure measurement using the endoscope channel was performed 10 times under each condition, and the mean pressure was calculated. In order to ensure the
accuracy and precision of the method to measure IP, the mean, standard deviation, absolute and relative errors, and coefficient of variation in pressures under each condition were calculated.

Absolute error = |true value – measured value|
Relative error = |true value – measured value| / true value
Coefficient of variation = standard deviation/average value

### Endoscopes

We used an electronic endoscopy system (EVIS LUCERA ELITE; Olympus Co., Tokyo, Japan) with high-resolution endoscopy (GIF-H290Z; Olympus) for the experiments.

### Part 2: Measurement of intragastric pressure during strong extension of the gastric wall: an in vivo study

### Patients

This study was approved by the Institutional Review Board of the Tobata Kyoritsu Hospital and was registered with the clinical trials registry (UMIN 000035500). All participating patients were explained about the study, and they provided written informed consent.

Consecutive patients scheduled for upper gastrointestinal endoscopy in the Department of Gastroenterology in Tobata Kyoritsu Hospital from January to March in 2019 were included in this study. Eligible patients were those who met all the following inclusion criteria and did not have any of the following exclusion criteria.

**Inclusion criteria**
1. Patients scheduled to undergo upper gastrointestinal endoscopy.
2. Patients with an Eastern Cooperative Oncology Group performance status score of 0 or 1 [5].
3. Patients with an age at registration ≤ 20 years.
4. Patients who provided written informed consent for study participation.

**Exclusion criteria**
1. Patients at risk of upper gastrointestinal bleeding.
2. Patients with previous gastric surgery and those with a gastric tube.
3. Patients with a serious underlying disease such as heart failure and ischemic stroke were excluded.
4. Patients who would have difficulty participating in this study due to psychiatric disorders or symptoms.
5. Patients deemed ineligible for participation by the investigator or associate investigator.

### Endoscopic procedures for measurement of intragastric pressure

All endoscopic procedures were performed using an electronic endoscopy system (EVIS LUCERA ELITE; Olympus Co., Tokyo, Japan) with high-resolution endoscopy (GIF-Q260, GIF-H260Z, GIF-H290Z; Olympus Co.) by certified specialists in gastrointestinal endoscopy (certified by the Japanese Gastroenterological Endoscopy Society) or an equally qualified endoscopist with > 10 years of experience in endoscopy.

All patients underwent optimum preparation and were required to drink a mixture of mucolytic and defoaming agents in water 30 minutes before the procedures [1]. The formula comprised 20,000 U pronase (Kaken Pharmaceutical, Tokyo, Japan), 1 g sodium bicarbonate, and 10 mL dimethylpolysiloxane (20 mg/mL; Horii Pharmaceutical, Osaka, Japan) added into 100 mL of water. Most patients (88%, 45/51) were sedated with 3 to 10 mg of diazepam (5 mg/mL; Takeda Pharmaceutical, Tokyo, Japan) or 2 to 8 mg of midazolam (5 mg/mL; SANDOZ, Tokyo, Japan) injected intravenously.

After completing the original indications for endoscopic examination, such as screening endoscopy or preoperative endoscopic assessment of early gastric cancer, we performed the following experiments specialized for this study. Air was insufflated until the gastric wall was strongly extended. Strong extension of the gastric wall was defined as that in which the greater curvature folds were flattened as a result of sufficient extension of the gastric wall by insufflated air (Fig. 3).

The intragastric pressure at the time of strong gastric wall extension was measured using the method developed in Part 1 (the in vitro study). CO2 was used for insufflation [6]. If adverse events including abdominal pain, Mallory-Weiss syndrome, and mucosal injury occurred during the experiments, the examination was immediately discontinued. After finishing the entire procedure, patients were asked whether they felt abdominal pain or discomfort.

### Endpoints

**Primary endpoint**

The primary endpoint of this study was determination of the IP at the time of strong extension of the gastric wall.

**Secondary endpoints**

There were several secondary endpoints evaluated in this study. First was the comparison of the IP at the time of strong extension of the gastric wall between patients with non-atrophic mucosa and those with atrophic mucosa on endoscopy [7, 8]. Second was the comparison of the IP at the time of strong extension of the gastric wall between patients with obesity (body mass index [BMI] ≥ 25 kg/m²) and those without obesity.
Finally, the safety was assessed by evaluating the frequency of adverse events (AEs). AEs included abdominal pain, Mallory-Weiss syndrome, and gastric fissures. Abdominal pain severity was graded using a numerical rating scale (NRS) from 0 to 10 (0, no pain; 10, worst imaginable pain) [10].

**Statistical analysis**

The mean values were compared using Student’s t-test. P<0.05 were considered statistically significant. SPSS version 21 (for Windows; SPSS, Chicago, Illinois, United States) was used for all statistical analyses.

**Results**

**Part 1: Development of a method to measure intragastric pressure: an in vitro study**

The standard deviation, absolute error, relative error, and coefficient of variation of pressures are demonstrated in Table 1, Table 2, Table 3, and Table 4. These values were high in Experiment 2 (Table 2) and Experiment 3 (Table 3), and they were higher than those in Experiment 1 (without any injection) (Table 1). Conversely, the mean pressures in Experiment 4 (Table 4) were similar to the intraballoon pressures that had been set, and the standard deviation, absolute error, relative error, and coefficient of variation were low. Thus, the measurements in Experiment 4 had high accuracy and reproducibility that were comparable to those in Experiment 1 in which nothing was injected into the forceps channel.

**Part 2: Measurement of intragastric pressure during strong extension of the gastric wall: an in vivo study**

**Clinical characteristics**

Fifty-four consecutive patients undergoing upper gastrointestinal endoscopy were included in this study. Of them, three patients were excluded because of difficulty retaining air in the stomach; the remaining 51 patients were included in the analysis. Overall, 31 (61%) patients were men and the mean age ± standard deviation (SD) was 64.7 ±20.3 years, while the mean BMI ± SD was 23.3 ±3.9 (Table 5). Atrophic changes were observed in 37 patients (73%) (Table 5) [7,8].

**Primary endpoint**

The mean IP ± SD at the time of strong extension of the gastric wall was 14.7 ±3.6 mmHg. The frequency of strong extension of the gastric wall under the condition of the IP of 20 mmHg or lower was 96% (49/51) (Fig. 4).

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**Table 1** Experiment 1: Pressure management in the forceps channel without any injection.

| Pressure in the balloon (mmHg) | Mean ± SD (mmHg) | AE (mmHg) | RE (%) | CV (%) |
|-------------------------------|------------------|-----------|--------|--------|
| 10                            | 10.4 ± 0.2       | 0.2       | 2.1    | 2.2    |
| 20                            | 20.4 ± 0.1       | 0.4       | 1.9    | 0.6    |
| 30                            | 30.1 ± 0.1       | 0.1       | 1.1    | 0.4    |
| 40                            | 40.2 ± 0.2       | 0.2       | 0.5    | 0.5    |

SD, standard deviation; AE, absolute error; RE, relative error; CV, coefficient of variation.

**Table 2** Experiment 2: Pressure management after injecting the forceps channel with lidocaine jelly.

| Pressure in the balloon (mmHg) | Mean ± SD (mmHg) | AE (mmHg) | RE (%) | CV (%) |
|-------------------------------|------------------|-----------|--------|--------|
| 10                            | 20.4 ± 8.9       | 9.4       | 93.9   | 43.5   |
| 20                            | 29.3 ± 3.5       | 9.3       | 46.5   | 11.8   |
| 30                            | 35.4 ± 4.7       | 6.1       | 20.4   | 13.2   |
| 40                            | 23.9 ± 8.9       | 11.6      | 29     | 30.9   |

SD, standard deviation; AE, absolute error; RE, relative error; CV, coefficient of variation.

**Table 3** Experiment 3: Pressure management after injecting the forceps channel with lidocaine jelly, injecting 20 mL of tap water once, and washing the forceps hole.

| Pressure in the balloon (mmHg) | Mean ± SD (mmHg) | AE (mmHg) | RE (%) | CV (%) |
|-------------------------------|------------------|-----------|--------|--------|
| 10                            | 10.3 ± 0.4       | 0.4       | 3.9    | 4.3    |
| 20                            | 20.2 ± 0.3       | 0.2       | 1.2    | 1.2    |
| 30                            | 30.1 ± 0.3       | 0.3       | 0.8    | 0.9    |
| 40                            | 39.9 ± 0.1       | 0.1       | 0.3    | 0.2    |

SD, standard deviation; AE, absolute error; RE, relative error; CV, coefficient of variation.

**Table 4** Experiment 4: Pressure management after injecting the forceps channel with lidocaine jelly, injecting 20 mL of tap water once to wash the forceps channel, and suctioning tap water from the forceps channel for 10 sec.

| Pressure in the balloon (mmHg) | Mean ± SD (mmHg) | AE (mmHg) | RE (%) | CV (%) |
|-------------------------------|------------------|-----------|--------|--------|
| 10                            | 12.1 ± 4.7       | 3.6       | 35.6   | 38.5   |
| 20                            | 21.1 ± 2.4       | 1.9       | 9.3    | 11.3   |
| 30                            | 26.5 ± 4.3       | 4.9       | 16.3   | 16.3   |
| 40                            | 37.9 ± 5.1       | 3.9       | 9.8    | 13.3   |

SD, standard deviation; AE, absolute error; RE, relative error; CV, coefficient of variation.
Secondary endpoints

The mean IP ± SD at the time of strong extension of the gastric wall in patients without atrophic mucosa on endoscopy was 14.6 ± 3.5 mmHg and that in those with atrophic mucosa on endoscopy was 15.0 ± 3.5 mmHg, with no statistically significant difference between the two groups ($p = 0.75$). The mean IP ± SD at the time of strong extension of the gastric wall in patients with a BMI $\geq 25$ kg/m$^2$ (N = 11) was 17.8 ± 3.7 mmHg and that in those with a BMI < 25 kg/m$^2$ (N = 40) was 13.9 ± 3.0 mmHg. Patients with obesity required significantly higher pressure to obtain strong extension of the gastric wall compared to those without obesity ($p = 0.007$ ($\uparrow$ Fig. 5). There were no serious AEs leading to discontinuation of the examination. Mild abdominal pain during the examination was observed in seven patients (13.7 %) ($\uparrow$ Table 5).

Discussion

In this study, the results suggested that an accurate and reproducible measurement of IP can be achieved by eliminating the influence mucus. Specifically, 20 mL of tap water was injected into the forceps channel of the endoscope for irrigation and was aspirated for 10 seconds; thereafter, the IP was measured. We aimed to develop an easy and accurate method to measure IP using endoscopy and to determine the IP required for strong extension of the gastric wall. During upper gastrointestinal endoscopy, blind spots can be avoided by sufficient extension of the gastric wall by air insufflation, which enables the observation of subtle changes in the color and surface (elevated or depressed lesions) of lesions [1]. In particular, strong extension of the gastric wall is essential to detect lesions in the folds. Therefore, observation at the time of strong extension of the gastric wall occurred in 96% of participants (49/51) when the gastric pressure was at most 20 mmHg.

$\uparrow$ Table 5 Clinical characteristics of 51 patients included in this study.

| Mean age ± SD, (years) | 64.7 ± 20.3 |
|------------------------|-------------|
| Sex                    |             |
| Male (%)               | 31 (61 %)   |
| Female (%)             | 20 (39 %)   |
| Mean body mass index ± SD, (kg/m$^2$) | 23.3 ± 3.89 |

| Endoscopic gastric mucosal atrophy |         |
|-----------------------------------|---------|
| Positive (%)                      | 37 (73 %) |
| Negative (%)                      | 14 (27 %) |

| Numeric rating scale | |
|----------------------|---------|
| 0 (%)                | 44 (86 %) |
| 1–2 (%)              | 7 (14 %) |
| 3–10 (%)             | 0 (0 %) |
| 3–10 (%)             | 0 (0 %) |

| Indications for endoscopy | |
|---------------------------|---------|
| Screening endoscopy       | 12 (24 %) |
| Abdominal pain            | 12 (24 %) |
| Appetite loss             | 1 (2 %) |
| Preoperative assessment for gastric cancer | 5 (10 %) |
| Surveillance after ESD for upper GI cancer | 14 (27 %) |
| Gastroduodenal ulcer      | 7 (13 %) |

SD, standard deviation; ESD, endoscopic submucosal dissection; GI, gastrointestinal.
gastric wall under certain conditions is important to establish the standard observation method. However, an objective method of measuring IP has not been established. Carbone et al. [4] and Suzuki et al. [11] reported methods to measure IP using a probe. In their studies, the probe was inserted from the forceps channel, and the IP was measured using a probe on the tip of the endoscope. However, it takes additional effort to apply this method during endoscopic examination. Therefore, we developed a simple and accurate method to measure intragastric pressure through an in vitro study. In our method, instead of inserting a probe from the forceps channel, the tube of the pressure measurement device was directly attached to the forceps channel hole. However, if this method is used in upper gastrointestinal endoscopy in clinical practice, the attachment of mucous in the forceps channel may prevent accurate measurement of the intragastric pressure because mucus intervenes between the IP measurement instrument and intragastric space by occluding the channel. There is a concern that retention of aspirated mucus in the forceps channel may prevent accurate measurement of IP. Therefore, we explored the optimal condition for measuring IP through experiments under various conditions.

Subsequently, we investigated the optimal IP for strong extension of the gastric wall in patients. The mean IP ± SD at the time of strong extension of the gastric wall was 14.7 ± 3.6 mmHg (10–23). Strong extension of the gastric wall occurred in 96% of patients (49/51) when the IP was at most 20 mmHg (Fig.4). These results suggest that the strong extension of the gastric wall can be obtained in almost all patients when the IP is at most 20 mmHg during upper gastrointestinal endoscopy.

Further studies regarding the usefulness and safety of the present method (IP of 20 mmHg) are required for clinical application. Regarding safety, Suzuki et al. [11] studied the changes induced by gastric stimulation using air inflation during endoscopy. They measured the IP at the visceral perception threshold in patients with functional dyspepsia/regurgitation esophagitis and controls and reported that visceral perception was observed in all groups when the IP was at least 22 mmHg. Additionally, incidents due to air inflation were not recorded. In the present study as well, there were no incidents or patients who complained of strong visceral perception (Table 5). However, no studies have investigated the safety of the measurement method in which the IP was maintained at 20 mmHg in patients undergoing endoscopy. We plan to conduct a prospective study on the safety of this method in which the IP is maintained at 20 mmHg (UMIN000039105). We previously reported that the non-extension sign with strong extension of the gastric wall was useful not only for endoscopic screening but also for preoperative endoscopic diagnosis of invasion depth of early gastric cancer [2,3]. In the future, we plan to conduct a prospective study on the optimal IP needed to detect the non-extension sign that is useful for the diagnosis of invasion depth in patients with early gastric cancer.

In the present study, the mean IP ± SD in patients with non-atrophic mucosa on endoscopy did not significantly differ from that in those with atrophic mucosa on endoscopy. The mechanism of dilatation of the stomach is considered as follows. In general, the stomach is dilated by release of relaxing substances such as nitric oxide and vasoactive intestinal peptide induced by physical stimulation (extension stimulation) associated with food intake [12,13]. In this study, it was considered that dilatation of the stomach was related to a gastric accommodation response associated with gastric wall extension by air insufflation; therefore, it was suggested that the presence or absence of atrophic mucosa might not have influenced the mechanism driving the dilatation of the stomach. Additionally, we compared the IP at the time of strong extension of the gastric wall between patients with and without obesity (BMI ≥ 25 kg/m² vs. <25 kg/m²) [9]. In the present study, the IP required for extension of the stomach was significantly higher in patients with obesity than in those without obesity (P = 0.007) (Fig.5). The potential reasons for this finding are as follows. First, as reported by Lambert et al. [14], obesity is associated with increased intraperitoneal pressure, suggesting that the pressure required for extension of the gastric wall is greater in patients with obesity because of their higher intraperitoneal pressure. Second, in this study, dilatation of the stomach was related to a gastric accommodation response, which was associated with gastric wall extension by air insufflation as described above. Weisen et al. reported that the autonomic responsiveness of the stomach was decreased in patients with obesity [15]. Therefore, it is suggested that the magnitude of stimulation required for strong extension of the gastric wall is greater in patients with obesity than in those without obesity.

The limitation of the in vitro experiment was that we did not perform a distribution-free test and calculated the standard deviation, absolute error, relative error, and coefficient of variation. Ten measurements may have been somewhat small for an accurate measurement. The limitation of the in vivo study...
was that it was an exploratory study of a small number of patients at a single center. With respect to the comparison of IP at the time of strong gastric wall extension between patients with endoscopic non-atrophic mucosa and those with endoscopic atrophic mucosa and between obese and non-obese patients as described in the secondary end-points, more number of cases are needed because the influence of other factors could not be excluded due to the small sample size. In addition, with respect to adverse events at the time of strong gastric wall extension, more cases are needed due to the small sample size. We are now planning to perform a clinical trial in a multicenter setting including a large number of patients.

**Conclusion**

In this study we developed an accurate, and reproducible method of measuring IP under the following conditions. To eliminate the factors that may negatively affect the accuracy of measurement (such as mucus in the forceps channel), 20 mL of tap water was injected into the forceps channel for irrigation and the tap water in the forceps channel was aspirated for 10 sec. Thereafter, the IP was measured. Our study also showed that strong extension of the gastric wall can be obtained in almost all patients when an IP of 20 mmHg is applied during upper gastrointestinal endoscopy. The quantitative measurement of IP may be useful for standardizing upper gastrointestinal endoscopy procedures, both for screening and for preoperative diagnosis of invasion depth of early gastric cancer.

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**Competing interests**

The authors declare that they have no conflict of interest.

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