INTRODUCTION

Screening endoscopy in asymptomatic populations is widely performed worldwide, and it has been reported to help prevent gastric cancer [1]. However, some examinees experience difficulties in tolerating gastroscopy. In some cases, all parts of the stomach cannot be observed for a sufficient time due to the examinee’s poor cooperation. Furthermore, poor cooperation may increase the risk of complications after endoscopy and lower the detection rate of gastric lesions.

To date, few studies have been conducted to identify the factors involved in poor cooperation during screening upper endoscopy. Only one previous study has been reported, which showed that age and anxiety significantly correlate with patient cooperation during endoscopy [2]. However, there may be a variety of factors affecting poor cooperation. Some people cannot suppress involuntary gagging even after pharyngeal anesthesia because of a sensitive gag reflex [3]. Others may fail to stifle a belch due to a large hiatal hernia [4,5]. Procedural sedation during endoscopy can maximize patients’ comfort and satisfaction [6,7]. The anterograde amnesic effect of midazolam makes the examinees un-
able to recall painful memories of the procedure. However, it may reduce the examinee’s cooperation during the procedure, because it is difficult to obey the operator’s commands under sedation, and sometimes, paradoxical reactions to midazolam can develop [8]. In addition, many other factors may impact on the examinee’s level of cooperation, including age, gender, underlying diseases, medication history, obesity, alcohol drinking, smoking, emotional stress, and depression.

Therefore, we investigated the factors that affect examinees’ cooperation during screening upper endoscopy. In addition, we examined the effect of the examinee’s cooperation on the completeness of the endoscopic examination and on complications after the procedure.

METHODS

Patients and questionnaires
We retrospectively reviewed the records of health examinations at Konkuk University Medical Center (Seoul, Korea), from September 2013 to April 2015. Among these, records of individuals over 18 years of age who underwent a gastroscopy were included in the study. To reduce the effect of variations between endoscopists, only gastroscopies carried out by a single experienced endoscopist (S.P.L.) were included. The study was approved by the Institutional Review Board (IRB) of Konkuk University School of Medicine, which confirmed that the study was in accordance with the ethical guidelines of the Helsinki Declaration (KUH1010664), and it was registered in the Clinical Research Information Service (CRIS) ID: KCT0001512.

All the examinees filled out a self-reporting questionnaire, which included the following items: underlying disease, medication history, smoking, alcohol intake, dyspepsia symptoms, and levels of emotional stress and depression. Subjects’ age, gender, height, weight, and endoscopic and histologic findings were also investigated. Heavy drinking was defined as consuming 15 drinks or more per week for men and eight drinks or more per week for women.

Endoscopic procedure
Gastroscopy was performed using a standard upper endoscope (GF-260, Olympus, Tokyo, Japan; or EPK-I, Pentax, Tokyo, Japan). Lidocaine spray was routinely used. All examinees provided written consent to undergo the endoscopic procedures. Midazolam for procedural sedation was selectively administered just before the start of the procedure in accordance with the examinee’s preference. The usual starting dose for producing moderate sedation was 1 to 3 mg based on age, gender, and medical conditions. The dose was increased to achieve an adequate level of sedation [6]. According to the American Society of Anesthesiologists guidelines, moderate sedation is defined as a state in which patients respond purposefully to verbal command and their cardiovascular function is maintained [9]. If a severe paradoxical reaction to midazolam occurred, it was immediately treated with flumazenil. If symptoms of midazolam overdose or midazolam side effects were noted, we tried to stabilize the vital signs of the patient first, and we considered the use of flumazenil. Symptoms of overdose or side effects included hypoxia, hypoventilation, hypotension, apnea, and central nervous system (CNS) depression. Oxygen supplementation was provided via a nasal cannula if the oxygen saturation level remained < 90%, despite verbal and physical stimuli. Hypoxia was recorded only when supplemental oxygen was needed. The incidence of acute complications (abdominal pain, nausea, and vomiting) and late-onset complications (persisting throat pain, bowel perforation, and gastrointestinal [GI] bleeding) were investigated through chart review.

Definitions and review of the endoscopic reports
The examinee’s cooperation and the completeness of endoscopic examination were evaluated by the operator’s subjective judgment and recorded directly after the examination. Information on the level of cooperation was collected through a review of the endoscopic reports. “Poor cooperation” meant lack of cooperation of the examinees during the procedure regardless of one’s will. The immediate causes of poor cooperation included severe belching, retching, coughing, and non-compliance of the examinees. “Belching” was defined as the forceful expulsion of gas from stomach through the mouth and “retching” was defined as an involuntary spasm of ineffectual vomiting. “Coughing” was defined as a sudden noisy expulsion of air from the bronchus or lungs, which is triggered by aspiration. “Non-compliance” referred to an action in contrast to bearing and
managing impulses to enable the operator to perform the procedure. The study was considered incomplete when some parts of the stomach could not be sufficiently visualized during the procedure. The causes of incomplete study included poor cooperation and a large amount of food materials in the stomach.

Endoscopic and histologic records were reviewed to identify the presence of digestive diseases, such as peptic ulcer disease, reflux esophagitis, GI tract masses, nodular gastritis, chronic atrophic gastritis, and metaplastic gastritis. The endoscopic findings were described based on the criteria of the Sydney classification [10]. A diagnosis of erosive esophagitis was made in cases with hyperemic streaks or mucosal breaks on the lower esophagus. Minimal change lesions of the esophagus were excluded from this diagnosis. Peptic ulcer disease was diagnosed as a deep mucosal defect with evidence of submucosal invasion, excluding ulcer scars. Chronic atrophic gastritis was diagnosed when the mucosal and submucosal vessels were visible without full air distension [11]. Nodular gastritis was defined as chicken-skin like mucosal changes or unusual goose-flesh appearance, and metaplastic gastritis was defined as uneven, irregular whitish patches or plaques [12]. Diagnoses of adenoma and carcinoma were made based on the results of tissue biopsy.

**Depression and emotional stress**

Depression and emotional stress were assessed using the Korean version of the Beck’s Depression Inventory (K-BDI) and the Brief Encounter Psychosocial Instrument-Korean version (BEPSI-K) [13,14]. The BEPSI-K consists of five self-administered items. A total score of 0 to 1.8 indicates a low level of stress; 1.8 to 2.8, a moderate level of stress; and more than 2.8, a high level of stress. The cut off value of 2.4 as a significant stressful status was set based on previous studies [15,16]. The K-BDI is a 21-item self-report questionnaire. A total score of 0 to 9 signifies no depression; 10 to 15, mild depression; 16 to 23, moderate depression; and 24 to 63, severe depression. We considered a total score of ≥ 10 as positive for depression.

**Definitions of functional dyspepsia**

Based on the Rome III definition, functional dyspepsia was diagnosed when the subjects had symptoms of dyspepsia without any evidence of reflux esophagitis, peptic ulcer disease, postoperative stomach, upper GI cancers, or any structural disease on upper GI endoscopy. Dyspepsia was defined as one or more of the following symptoms occurring at least once per week in the last 3 months: (1) bothersome postprandial fullness after ordinary-sized meals, (2) early satiety that prevents finishing a regular meal, and (3) epigastric pain or burning. Symptoms with an onset occurring more than 6 months prior to the survey were not asked about in our questionnaire.

**RESULTS**

**Baseline characteristics of the subjects**

A total of 4,422 subjects (56.5% male) were included in the study. The overall mean age and body mass index (BMI) were 45.19 ± 9.87 years (range, 18 to 86) and 23.57 ± 3.27 kg/m² (range, 14.6 to 39.7), respectively (Table 1). The number of subjects who received sedation (midazolam) during endoscopy was 3,164 (71.6%). Cooperation during the endoscopic procedure was poor in 358 subjects (8.1%). The other characteristics of the study population are shown in Table 1.

**The poor cooperation and good cooperation groups**

In the poor cooperation group, the cause of poor cooperation was not recorded in 214 cases. Of the patients with causes recorded, the causes of poor cooperation were belching in 114 subjects (79.2%), retching in 94 (65.3%), non-compliance in 18 (12.5%), and coughing in
Poor cooperation was more common among younger subjects and in women ($p < 0.001$ and $p = 0.003$, respectively) (Table 2). When broken down by age group, cooperation was poor in 9.6% of those in their 20s or younger (18 of 187), 10.5% of those in their 30s (123 of 1,174), 7.9% of those in their 40s (112 of 1,436), and 3.8% of those in their 50s or older (28 of 741). BMI, K-BDI, BEPSI-K, smoking, drinking, and dyspepsia symptoms were not properly measured in 23, 126, 495, 439, 196, and 250 subjects, respectively.

BMI, K-BDI, BEPSI-K, smoking, drinking, and dyspepsia symptoms were not properly measured in 23, 126, 495, 439, 196, and 250 subjects, respectively.

**Table 1. Baseline characteristics of the subjects ($n = 4,422$)**

| Variable                      | No. (%) | Mean ± SD | Median (range) |
|-------------------------------|---------|-----------|----------------|
| Age, yr                       | 45.19 ± 10.44 | 44 (18–86) |
| Male sex                      | 2,498 (56.5) |
| Body mass index, kg/m²        | 23.57 ± 3.27 | 23.31 (14.6–39.7) |
| ≥ 25                          | 1,339 (30.4) |
| ≥ 18.5 and < 25               | 2,882 (65.5) |
| < 18.5                        | 178 (4.1) |
| K-BDI level                   | 2.22 ± 4.05 | 0 (0–46) |
| No depression                 | 4,123 (96.0) |
| Mild                          | 96 (2.2) |
| Moderate                      | 61 (1.4) |
| Severe                        | 16 (0.4) |
| BEPSI-K level                 | 1.61 ± 0.83 | 1.4 (0–5) |
| Low level of stress           | 2,488 (63.4) |
| Moderate level                | 931 (23.7) |
| High level                    | 508 (12.0) |
| Smoking                       |         |           |                |
| Non-smoker                    | 2,131 (53.5) |
| Past smoker                   | 1,004 (25.2) |
| Current smoker                | 848 (21.3) |
| Alcohol                       |         |           |                |
| Non-drinker                   | 857 (20.3) |
| Social drinker                | 3,224 (76.3) |
| Heavy drinker                 | 145 (3.4) |
| Comorbidity                   |         |           |                |
| Diabetes                      | 196 (4.4) |
| Hypertension                  | 543 (12.3) |
| Congestive heart disease      | 68 (1.5) |
| Stroke                        | 5 (0.1) |
| Medication                    |         |           |                |
| Antiplatelet agent            | 76 (1.7) |
| Anticoagulant                 | 6 (0.1) |
| NSAID                         | 226 (5.1) |
| Antidepressant                | 22 (0.5) |
| Herb and health food          | 750 (17.0) |
| Symptoms of dyspepsia         | 691 (16.6) |
| Postprandial fullness         | 470 (11.3) |
| Early satiety                 | 280 (6.7) |

**Table 1. Continued**

| Variable                      | No. (%) | Mean ± SD | Median (range) |
|-------------------------------|---------|-----------|----------------|
| Epigastric pain/burning       | 218 (5.2) |
| Procedural sedation           | 3,164 (71.6) |
| Dose of midazolam, mg         | 3.69 ± 0.67 | 4 (1–7) |
| Poor cooperation during endoscopy | 358 (8.1) |

BMI, K-BDI, BEPSI-K, smoking, drinking, and dyspepsia symptoms were not properly measured in 23, 126, 495, 439, 196, and 250 subjects, respectively.

SD, standard deviation; K-BDI, Korean version of the Beck's Depression Inventory; BEPSI-K, The Brief Encounter Psychosocial Instrument-Korean version; NSAID, nonsteroidal anti-inflammatory drug.

**Figure 1.** Venn diagram of the causes of poor cooperation. In 214 subjects, the causes of poor cooperation were not recorded. After their exclusion, the causes of poor cooperation were belching in 114 subjects (79.2%), retching in 94 (65.3%), non-compliance in 18 (12.5%), and coughing in two (1.4%). Poor cooperation was often caused by multiple reasons (79 subjects, 54.9%). Belching and retching frequently occurred together.

Two (1.4%) (Fig. 1). Often, poor cooperation was caused by multiple reasons (79 subjects, 54.9%). Poor cooperation was more common among younger subjects and in women ($p < 0.001$ and $p = 0.003$, respectively) (Table 2). When broken down by age group, cooperation was poor in 9.6% of those in their 20s or younger (18 of 187), 10.5% of those in their 30s (123 of 1,174), 7.9% of those in their 40s (112 of 1,436), and 3.8% of those in their 50s or older (28 of 741). BMI, K-BDI, BEPSI-K, smoking, drinking, and dyspepsia symptoms were not properly measured in 23, 126, 495, 439, 196, and 250 subjects, respectively.

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Lee SP, et al. Patient cooperation during gastroscopy

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https://doi.org/10.3904/kjim.2015.393

Table 2. Comparison between the poor cooperation and good cooperation group

| Variable                  | Poor cooperation group (n = 358) | Good cooperation group (n = 4,064) | p value |
|---------------------------|---------------------------------|-----------------------------------|---------|
| Age, yr                   | 43.40 ± 9.87                    | 45.35 ± 10.48                     | < 0.001b |
| < 40                      | 141 (39.4)                      | 1,220 (30.0)                      | < 0.001b |
| Female sex                | 183 (51.1)                      | 1,741 (42.8)                      | 0.003b  |
| Body mass index, kg/m²    | 24.17 ± 3.37                    | 23.52 ± 3.25                      | 0.003b  |
| ≥ 25                      | 130 (36.6)                      | 1,209 (29.9)                      | 0.010b  |
| Smoking                   |                                 |                                   | 0.049b  |
| Non-smoker                | 188 (52.7)                      | 1,943 (53.1)                      |         |
| Past smoker               | 64 (19.6)                       | 940 (25.7)                        |         |
| Current smoker            | 74 (22.7)                       | 774 (21.2)                        |         |
| Alcohol                   |                                 |                                   | 0.366   |
| Non-drinker               | 78 (22.5)                       | 779 (20.1)                        |         |
| Social drinker            | 266 (75.1)                      | 2,964 (76.4)                      |         |
| Heavy drinker             | 8 (2.3)                         | 137 (3.5)                         |         |
| Medication                |                                 |                                   |         |
| Antidepressant            | 0                               | 22 (0.5)                          | 0.252   |
| NSAID                     | 21 (5.9)                        | 205 (5.0)                         | 0.530   |
| K-BDI level ≥ 10          | 19 (5.4)                        | 154 (3.9)                         | 0.200   |
| BEPSI-K level ≥ 2.4       | 62 (19.4)                       | 710 (19.7)                        | 0.946   |
| First-time endoscopy      |                                 |                                   | 0.618   |
| Procedural sedation       | 316 (88.3)                      | 2,848 (70.1)                      | < 0.001b |
| Midazolam side effect     |                                 |                                   |         |
| Hypotension               | 0                               | 1                                 |         |
| Hypoxia                   | 15                              | 128                               |         |
| Use of flumazenil         | 10                              | 1                                 |         |
| Digestive diseases        |                                 |                                   |         |
| Functional dyspepsia      | 47 (13.9)                       | 595 (15.9)                        | 0.434   |
| Nodular gastritis         | 3 (0.8)                         | 44 (1.1)                          | 1.000   |
| Chronic Atrophic gastritis| 215 (60.1)                      | 2,350 (57.8)                      | 0.434   |
| Metaplastic gastritis     | 23 (6.4)                        | 240 (5.9)                         | 0.726   |
| Reflux esophagitis        | 22 (6.1)                        | 209 (5.1)                         | 0.456   |
| Peptic ulcer disease      | 5 (1.4)                         | 88 (2.2)                          | 0.353   |
| Postoperative stomach     | 1 (0.3)                         | 15 (0.4)                          | 1.000   |
| Gastric adenocarcinoma    | 0                               | 4                                 | 1.000   |

Values are presented as mean ± SD or number (%). NSAID, nonsteroidal anti-inflammatory drug; K-BDI, Korean version of the Beck’s Depression Inventory; BEPSI-K, The Brief Encounter Psychosocial Instrument-Korean version.

Table 2. Continued

| Variable                  | Poor cooperation group (n = 358) | Good cooperation group (n = 4,064) | p value |
|---------------------------|---------------------------------|-----------------------------------|---------|
| Gastric adenoma           | 0                               | 6                                 | 1.000   |
| Hiatal hernia             | 19 (5.3)                        | 118 (2.9)                         | 0.077b  |
| Complications             | 3 (0.8)                         | 12 (0.3)                          | 0.116   |
| Persisting throat pain    | 2                               | 8                                 |         |
| GI bleeding               | 0                               | 0                                 |         |
| Bowel perforation         | 0                               | 0                                 |         |
| Abdominal pain            | 1                               | 2                                 |         |
| Nausea/Vomiting           | 0                               | 2                                 |         |
| Incomplete study          | 36 (10.1)                       | 8 (0.2)                           | < 0.001b |
| Due to poor cooperation   | 36                              | 0                                 |         |
| Due to food materials     | 0                               | 8                                 |         |

Values are presented as mean ± SD or number (%). NSAID, nonsteroidal anti-inflammatory drug; K-BDI, Korean version of the Beck’s Depression Inventory; BEPSI-K, The Brief Encounter Psychosocial Instrument-Korean version.

Table 2. Continued

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| Complications             | 3 (0.8)                         | 12 (0.3)                          | 0.116   |
| Persisting throat pain    | 2                               | 8                                 |         |
| GI bleeding               | 0                               | 0                                 |         |
| Bowel perforation         | 0                               | 0                                 |         |
| Abdominal pain            | 1                               | 2                                 |         |
| Nausea/Vomiting           | 0                               | 2                                 |         |
| Incomplete study          | 36 (10.1)                       | 8 (0.2)                           | < 0.001b |
| Due to poor cooperation   | 36                              | 0                                 |         |
| Due to food materials     | 0                               | 8                                 |         |

Values are presented as mean ± SD or number (%). NSAID, nonsteroidal anti-inflammatory drug; K-BDI, Korean version of the Beck’s Depression Inventory; BEPSI-K, The Brief Encounter Psychosocial Instrument-Korean version.

*Continuous variables were analyzed by the Mann-Whitney test. All other data were analyzed by the chi-square test.

bThe significant results.

among the subjects who received procedural sedation, 10 (3.2%) in the poor cooperation group and one in the good cooperation group were administered the antidote (flumazenil). In the good cooperation group, the reason for using the antidote was hypoxia that could not be alleviated with nasal cannula oxygenation. Three subjects in the poor cooperation group and 12 subjects in the good cooperation group developed complications after the procedure, with no significant difference between the two groups (p = 0.116). The endoscopic exam-
ination of 44 subjects was an “incomplete study.” The causes of incomplete study were poor cooperation in 36 cases and large amounts of food materials in the stomach in eight cases.

**Predisposing factors for examinees’ poor cooperation**

Logistic regression analysis revealed that that young age (< 40 years) and female sex were independent predisposing factors for an examinee’s poor cooperation during upper endoscopy (OR, 1.423; 95% CI, 1.125 to 1.801; \( p = 0.003 \)) (Table 3). In addition, high BMI (≥ 25), hiatal hernia, and procedural sedation were also independent predisposing factors for poor cooperation (OR, 1.501; 95% CI, 1.165 to 1.935; \( p = 0.002 \)) (OR, 2.235; 95% CI, 1.300 to 3.844; \( p = 0.004 \)) (OR, 2.918; 95% CI, 2.083 to 4.088; \( p < 0.001 \)). After logistic analysis, smoking history was not related with examinees’ poor cooperation.

**DISCUSSION**

Our study showed that 8.1% of examinees cooperated poorly with elective gastroscopy. This indicates that a considerable number of examinees have difficulty when undergoing gastroscopy. The main predisposing factor for poor cooperation was sedation using midazolam. The risk of poor cooperation in subjects who received sedation was three times higher than that in subjects who did not receive sedation. Sedation can assuage patients’ fears about receiving endoscopy and maximize patient comfort during the procedure [6,7,9]. However, a patient under sedation may not be able to obey the doctor’s commands. For example, when patients cannot suppress a belch during endoscopy, doctor can instruct them to hold their breath for a while. Likewise, doctor can direct not to swallow saliva to prevent coughing. However, it is difficult for patients under sedation to carry out the instructions because of altered level of consciousness. In addition, paradoxical reactions to midazolam can sometimes develop. Therefore the outcome of previous endoscopic examinations should be examined to decide whether sedation is performed. If

| Variable          | Univariate analysis | Multivariate analysis |
|-------------------|---------------------|-----------------------|
|                   | OR (95% CI)         | \( p \) value         | Adjusted OR (95% CI) | \( p \) value |
| Age < 40          | 1.515 (1.213–1.892) | < 0.001               | 1.423 (1.125–1.801)  | 0.003*       |
| Female sex        | 1.395 (1.124–1.732) | 0.003                 | 1.670 (1.216–2.294)  | 0.002*       |
| Body mass index ≥ 25 | 1.355 (1.081–1.355) | 0.008                | 1.501 (1.165–1.935)  | 0.002*       |
| Hiatal hernia     | 1.874 (1.140–3.081) | 0.013                | 2.235 (1.300–3.844)  | 0.004*       |
| Procedural sedation | 3.212 (2.312–4.493) | < 0.001              | 2.918 (2.083–4.088)  | < 0.001*     |
| Non-smoker        | 1.202 (0.956–1.511) | 0.116                | -                      | -             |

OR, odds ratio; CI, confidence interval.

*Univariate and multivariate logistic regression analyses were performed and footnotes are placed at the significant results.
the examinee’s cooperation was very poor in a previous endoscopy procedure under sedation, endoscopy without sedation should be considered.

Young age and female sex were predisposing factors for poor cooperation. Young people may not be able to suppress involuntary gagging due to a more sensitive gag reflex [3]. Gender differences in pain sensitivity have become a focus of clinical interest in recent decades. Previous studies revealed that postoperative and procedural pain may be more severe among female than among male individuals [17]. In addition, women have lower visceral pain thresholds [18].

Hiatal hernia and obesity were also significantly associated with poor cooperation. People with a hiatal hernia may fail to stifle a belch during the endoscopic procedure, which can interrupt observation. Obesity can cause hiatal hernia [5,10,20]. However, regardless of the existence of hiatal hernia, obesity can increase intra-abdominal pressure, and the increased pressure can obstruct air inflation during endoscopy [19]. Thus it can prevent the stomach from expanding and interrupt observation. Although overall, the majority of females, young people, obese people, and people with hiatal hernias underwent endoscopy relatively well in our study, before performing the procedure in these patients, endoscopists should keep in mind that cooperation may be poor.

The causes of poor cooperation in this study were diverse, including belching, retching, non-compliance, and coughing. However, it was often caused by several factors, meaning that sometimes a cause cannot be determined. Belching and retching especially often occurred together. In practice, however, it is difficult to distinguish belching from retching in many cases. Non-compliance can be caused by anxiety related to fear or to paradoxical reactions to midazolam. If a paradoxical reaction was suspected during endoscopy, antidote was administered without delay. In our study, 10 examinees received flumazenil due to non-compliance, in all of whom the endoscopic examination was finished safely.

The manifestations of midazolam overdose or side effects include hypoxia, hypoventilation, hypotension, apnea, and CNS depression [6]. In our study, there were few severe symptoms of overdose or side effect, and the most common symptom that occurred was hypoxia. Most cases of hypoxia resolved with conservative treatment, with only one case requiring the use of flumazenil because of hypoxia.

Our study showed that about 10% of endoscopic examinations in the poor cooperation group were incomplete. In addition, cases of adenoma and adenocarcinoma were not detected in the poor cooperation group. Although the sample size was small, these results revealed that the examinee’s poor cooperation can interfere with adequate stomach observation.

This study has some limitations. First, the endoscopist’s judgment of poor cooperation is bound to be subjective. We therefore investigated the records of procedures performed by only one endoscopist to minimize bias and inter-individual variance of endoscopists. Second, some procedural complications may have been overlooked. Mild oral bleeding or throat pain after the procedure might not be included in a patient’s record, because the examinees can leave the hospital without complaint. Third, there were some missing values. The BMI of 23 subjects, K-BDI of 126 subjects, BEPSI-K of 495 subjects, smoking history of 439 subjects, drinking status of 196 subjects, and dyspepsia symptoms of 250 subjects were not measured or were not recorded. The cause of poor cooperation was also not recorded in 214 subjects. Lastly, although we measured the level of emotional stress and depression, which had no effect on the examinee’s cooperation, the anxiety level of the examinees was not measured.

In conclusion, cooperation during elective gastroscopy was poor in a considerable number of examinees. Poor cooperation did not lead to more frequent complications but could sometimes cause incomplete study. Endoscopists must keep in mind that cooperation is more likely to be poor in young people, obese people, women, those with hiatal hernia, and those who received sedation during the examination.

**KEY MESSAGE**

1. Cooperation during gastroscopy is poor in a considerable number of examinees and poor cooperation sometimes causes incomplete study.
2. Cooperation during gastroscopy is more likely to be poor in young people, obese people, women, those with hiatal hernia, and those who received sedation.
Conflict of interest
No potential conflict of interest relevant to this article was reported.

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