A novel wide viewing endoscope for upper gastrointestinal screening: a pilot study

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Introduction
Esophagogastroduodenoscopy (EGD) has become an important examination for screening and diagnosis of diseases of the upper gastrointestinal tract. Improved endoscopic screening was accomplished by the incorporation of a thinner caliber endoscope with the capacity to capture high definition images. However, the ability to visualize Vater’s papilla with front viewing endoscopes is less than satisfactory [1]. The duodenum has a narrow lumen; therefore, endoscopes with a side view are preferred for visualization of Vater’s papilla, although current side viewing scopes are not suitable for screening in EGD. In addition, switching to a side viewing endoscope may not be a viable option when considering a patient’s comfort and the availability of endoscopic equipment.

Full-spectrum endoscopy (FUSE; EndoChoice, Inc., Alpharetta, GA, United States) EGD is a newly developed video-endoscope that provides a 245-degree field of view with double imagers on the front and side of the endoscope tip. A wider view for endoscopic examinations, such as ultra-wide viewing scopes, has been reported to enhance diagnostic yield in the field of colonoscopy [2–4]. In the upper gastrointestinal, locations with a steep curve such as the squamo-columnar junction (SCJ) and the anal side of the pyloric ring may not be well observed by forward viewing endoscopes. Theoretically, the incorporation of double imagers in different directions has a possibility to produce a better view in these locations. However, there are no available data to determine whether a wider endoscopic view is efficient for upper gastrointestinal screening. The aim of this study was to evaluate the feasibility, usability, and safety of FUSE-EGD in human subjects. Feasibility was defined by the completion of taking images according to a standardized protocol within an acceptable time. Usability was defined by technical handling of the endoscope and quality of the images.

Methods and materials
The study included 21 healthy individuals (15 males and 6 females) with a median age of 35 years (range, 29–55 years) who underwent upper gastrointestinal screening using FUSE-EGD. The mean procedure time was 6.5 min, with a median of 91 images captured. The rate of successful visualization of Vater’s papilla was 90% (19/21). The whole circumference of the SCJ was observed with two video monitors in all cases. The anterior anal side of the pyloric ring was observed in 29% (6/21) of cases. However, the general impression of the usability of FUSE-EGD was that it was rather inferior to that of a standard front viewing endoscope. Although the usability requires further modification, FUSE-EGD provided excellent results for imaging Vater’s papilla and the SCJ.
Object drank 100mL of water containing pronase, sodium bicarbonate, and simethicone. The examination was performed under oral anesthesia with 1–5 sprays of 8% lidocaine, with no sedation. Five endoscopists who were board certified members of the Japanese Gastroenterological Endoscopy Society performed screening of the upper gastrointestinal according to a standardized protocol while capturing sequential images from the esophagus to the duodenum. The total duration of the endoscopic procedure was recorded using a stopwatch.

The primary end point of this pilot and feasibility study was the rate of successful visualization of Vater’s papilla. Additional end points included the duration of the procedure, successful visualization of the anal side of the pyloric ring and the SCJ, adverse events, and the endoscopists’ subjective evaluation of the usability of the FUSE-EGD procedure. The endoscopists’ subjective evaluation of the usability, field of view, and maneuverability of FUSE was assessed using a five-point Likert scale with the following answer options: unacceptable, difficult, acceptable, good, and excellent. In addition, the endoscopists’ subjective evaluation compared to the standard endoscope used in our hospital (9.8-mm, high definition, front viewing, video endoscope) was also evaluated.

Endoscopy platform
The FUSE platform consists of a video endoscope and a main control unit. The video images are presented to the endoscopist on two contiguous monitors. The FUSE-EGD (FSG-2500-ST) system is a 9.6-mm, flexible endoscope for repeated clinical use (diagnostic visualization and therapeutic intervention). The endoscope is equipped with a working channel (2.8 mm), air or CO2 insufflation options, suction, and forward waterjet irrigators. A wide 245-degree field of view is achieved using two lenses and groups of light-emitting diodes positioned on the front and side of the endoscope tip. This retrospective study was approved by the institutional review board of our hospital (No. 27-J69-27-1-3).

Results
Gastrointestinal screening using the FUSE-EGD system was successfully completed for all 21 subjects with a median procedure time (±SD) of 6.5 ± 1.6 min. No acute or delayed adverse events occurred. A median of 90.6 ± 24.9 endoscopic images was acquired. Successful visualization of Vater’s papilla was obtained in 90% (19/21) of the subjects (Fig. 1). The duodenal side of the pyloric ring (anterior wall only) was observed in 29% (6/21) of the subjects. The whole circumference of the SCJ was observed on the two video monitors in all cases (Fig. 2). The main finding detected among the subjects was esophageal hernia (100%, 21/21).

Subjective evaluation of the usability of FUSE-EGD was assessed by the five endoscopists, who reported that FUSE-EGD was inferior compared with the standard front viewing endoscope for all participants (Table 1). However, 80% of the participants answered that the FUSE-EGD system platform was feasible as a whole.

Discussion
This is the first study on humans utilizing the newly developed FUSE-EGD for upper gastrointestinal screening. The main end point of this study was to evaluate the feasibility of the new FUSE-EGD system. Successful visualization of Vater’s papilla was achieved in 90% of the study participants. In addition, no adverse events were recorded associated with this system. The duodenum has a narrow lumen and Vater’s papilla is not routinely identified during screening by EGD [1] or by capsule endoscopy [5]. A previous study reported that the detection rate of Vater’s papilla in whole or in part was only 57% using a routine imaging method, but it increased to 89% using a straightening endoscopic technique [1]. For surveillance of patients with duodenal polyposis, the use of a side viewing endoscope [6] or side and front viewing endoscopes at the same time under general
anesthesia has been reported [7]. However, these complex maneuvers may not be appropriate for upper gastrointestinal screening. The FUSE-EGD system is equipped with a side view imager on a front viewing endoscope that enables easier detection of Vater’s papilla with normal manipulation of the endoscope. The FUSE-EGD system allowed easier visualization of Vater’s papilla than the conventional endoscope.

The SCJ was visualized on both monitors in all subjects. Although Barrett’s esophagus is rare among the Japanese population, a clear view of the entire SCJ is greatly beneficial for the detection of early SCJ and Barrett’s adenocarcinoma. Vater’s papilla is an excellent marker to assess this system; however, the real clinical benefit could be at the SCJ.

The anal side of the pyloric ring was observed in 29% of the subjects. We expected that the side lens would visualize the anal side of the pyloric ring. In most cases, the mucosa contacted the side lens, and therefore it was difficult to keep a distance between the mucosa and the lens in the narrow duodenal bulb. The ability to visualize the anal side of the pyloric ring may be beneficial for gastric antral lesions extending to the duodenal bulb; however, these lesions are quite rare.

All of the endoscopists participating in this study reported that the usability of FUSE-EGD was inferior to that of the conventional endoscope. However, all procedures were performed safely and without difficulty with no sedation. Since none of the endoscopists had previous experience using FUSE-EGD, it may just be a matter of getting used to the maneuverability of the scope. The image quality and brightness of the view field are expected to be further improved in the near future.

A limitation to this study was the small sample size and limited number of participating endoscopists. However, the results of this first pilot and feasibility human study of FUSE-EGD are promising.

In conclusion, although its usability requires further modification, FUSE-EGD provided excellent results for imaging Vater’s papilla and the SCJ. FUSE-EGD has the potential to improve the efficacy of upper gastrointestinal screening.

**Competing interests:** None

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**References**