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Efficacy and safety of EUS-guided gastroenterostomy for benign and malignant gastric outlet obstruction: a systematic review and meta-analysis





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ABSTRACT

Background and study aims Endoscopic ultrasound-guided gastroenterostomy (EUS-GE) is a novel endoscopic procedure designed to facilitate sustained luminal patency in patients with gastric outlet obstruction. The primary aim of this study was to evaluate the efficacy and safety of EUS-GE for treatment of gastric outlet obstruction.

Methods Searches of PubMed, EMBASE, Web of Science, and Cochrane Library databases were performed through April 2019. Patients with benign and malignant gastric outlet obstruction were included. Measured outcomes included: immediate technical and clinical success as well as rate of serious adverse events (AEs). Heterogeneity was assessed with Cochran Q test and I^2 statistics. Publication bias was ascertained by funnel plot and Egger regression testing.

Results A total of five studies (n=199 patients; 45.73% male) were included in this study. Four retrospective studies and one prospective study were analyzed. Mean age of patients that underwent the EUS-GE procedure was 64.52 \pm 1.37 years with a pooled mean follow-up period of 4.32 \pm 1.65 months. In 21% of patients (n=43), gastric outlet obstruction was due to benign causes. Immediate technical success was 92.90% (95% CI; 88.26 – 95.79; I^2 = 0.00%) and reported in all studies. The clinical success rate of EUS-GE was 90.11% (95% CI; 84.64 – 93.44; I^2 = 0.00%). Serious AEs occurred in 5.61% (95% CI; 2.87 – 10.67; I^2 = 1.67%) of cases and were related to peritonitis, perforation, bleeding, and abdominal pain. Re-intervention rate was 11.43% (95% CI; 7.29 – 17.46; I^2 = 17.38%).

Conclusions EUS-GE appears to provide an effective and safe minimally invasive alternative for treatment of benign and malignant gastric outlet obstruction.

Introduction

Gastric outlet obstruction is a clinical condition characterized by nausea and vomiting, postprandial fullness, as well as epigastric abdominal pain and weight loss as a pathophysiologic consequence of a mechanical obstruction within the gastrointestinal tract. Previously termed pyloric stenosis, gastric outlet obstruction is the preferred or standard terminology as mechanical obstruction may occur in the distal stomach, pylorus, or duodenum. The condition itself is typically classified based upon etiology with benign and malignant causes. Although the primary etiology over time has changed from peptic ulcer disease and other benign entities to a more common association with underlying malignancy, gastric outlet obstruction has

a significant effect on patient morbidity and overall quality of life [1-5].

Traditionally, surgical gastrojejunostomy has been the primary treatment for both benign and malignant gastric outlet obstruction although the procedure itself is associated with a high complication rate that approaches more than 40% [6,7]. Although surgical gastrojejunostomy is currently the standard of care for palliative treatment, the procedure is further limited by prolonged recovery times delaying chemotherapy for malignancy-related obstructions, delayed gastric emptying and gastroparesis, as well as substantial procedure-associated costs [7]. Given the significant morbidity of this surgical procedure, less-invasive endoscopic alternatives have been developed including endoscopic ultrasound-guided gastroenterostomy (EUS-GE).

EUS-GE has recently emerged as a procedure to treat patients with gastric outlet obstruction as an alternative to surgery or to standard endoscopy when endoscopic enteral stent placement is not possible [7,8]. Although EUS-GE was first described by Fritscher-Ravens et al. in the early 2000 s, it was only after the recent availability of a bi-flanged lumen-apposing metal stent (LAMS) that EUS-GE was clinically adopted. EUS-GE performed with placement of a LAMS has emerged as another treatment option that may provide long-term luminal patency without associated morbidity and complication rates of a surgical approach [9, 10]. While this novel procedure is designed to facilitate sustained luminal patency in patients with gastric outlet obstruction while avoiding the morbidity of a surgical procedure, the efficacy and safety is EUS-GE remains less clear.

The primary aim of this study was to perform a systematic review and meta-analysis to evaluate the efficacy and safety of EUS-GE for treatment of benign and malignant gastric outlet obstruction.

Methods

Literature search

A comprehensive search of the literature was performed to identify articles that examined the EUS-GE procedure. Systematic searches of PubMed, EMBASE, Web of Science, and the Cochrane Library databases were performed from 2001 through April 1, 2019. The following search terms included: "endoscopic ultrasound-guided gastroenterostomy (EUS-GE)," "endoscopic ultrasound-guided gastrojejunostomy (EUS-GJ)," "endoscopic ultrasound-guided gastroduodenostomy (EUS-GD)," "endoscopic ultrasound and gastric outlet obstruction," and "endoscopic ultrasound and pyloric obstruction." All relevant articles irrespective of year of publication, type of publication, or publication status were included. Titles and abstracts of all potentially relevant studies were screened for eligibility. The reference lists of studies of interest were then manually reviewed for additional articles by cross-checking bibliographies. Two reviewers (TRM and RG) independently screened the titles and abstracts of all the articles according to predefined inclusion and exclusion criteria. Any differences were resolved by mutual agreement and in consultation with the third reviewer (TR). In the case of studies with incomplete information, contact was attempted with the principal authors to obtain additional data.

Study selection criteria

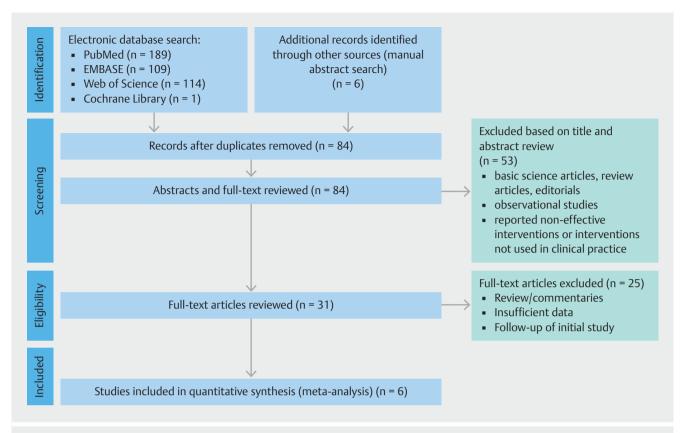
This study was prospectively submitted in PROSPERO, an international database of prospectively registered systematic reviews in health and social care. The Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) statement outline and Meta-Analysis of Observational Studies in Epidemiology (MOOSE) reporting guidelines for reporting systematic reviews and meta-analyses was used to report findings [11, 12]. Only studies investigating use of the EUS-GE modality for treatment of gastric outlet obstruction were included. Only human subject studies were considered in the analysis. A study was excluded if deemed to have insufficient data, as were review articles, editorials, and correspondence letters that did not report independent data. Case series and reported studies with fewer than 10 patients were excluded. Multiple published work from similar authors was evaluated for overlapping enrollment times to preserve independence of observations. Participants included patients of any age in whom presence of gastric outlet obstruction was suspected or confirmed based upon prior imaging studies.

Procedure characteristics

All patients included in this study for analysis underwent EUS-GE for treatment of gastric outlet obstruction. Multiple EUS-GE procedures (i.e., unassisted and assisted techniques) were also analyzed including: direct EUS-GE, balloon-assisted, EUS-guided double-balloon-occluded gastrojejunostomy bypass (EPASS), nasobiliary drain, and Natural Orifice Transluminal Endoscopic Surgery (NOTES). All sizes of available LAMS were included as well as both cautery-enhanced lumen-apposing metal stent (CE-LAMS) and non-cautery-enhanced lumen-apposing metal stent (NCE-LAMS). Both EUS-GJ and EUS-GD were included.

Outcome measures

The primary outcome measurement in this study was the feasibility, efficacy, and tolerability of EUS-GE in patients with gastric outlet obstruction. Efficacy and safety of the device were measured by immediate technical success rate (i.e., ability to perform the procedure without issue), clinical success as defined by individual study authors, and serious adverse events (AEs) reported during follow-up. Clinical success was also measured by level of oral intake or relief of symptoms (i. e., early satiety, nausea, vomiting) using the validated gastric outlet obstruction scoring system (GOOSS) [13]. This GOOSS is designed to provide an objective grade to determine patients' ability to eat before and after gastric outlet obstruction procedure (i.e., EUS-GE). Other measured outcomes included baseline patient characteristics (i.e., mean age, gender, benign or malignancy etiology of gastric out obstruction, and prior gastrointestinal treatment), procedural-related characteristics (i.e., EUS-GE technique, timing of procedure, and other AEs), and duration of follow-up.



▶ Fig. 1 Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) flow chart of literature search results for EUS-guided gastroenterostomy for the treatment of gastric outlet obstruction.

Statistical analysis

This meta-analysis was performed by calculating pooled proportions. After appropriate studies were identified through systematic review, the individual study proportion was transformed into a quantity using the Freeman–Tukey variant of the arcsine square root transformed proportion. Then the pooled proportion was calculated as the back transform of the weighted mean of the transformed proportions, using inverse arcsine variance weights for the fixed effects model and DerSimonian–Laird weights for the random effects model [14–17].

Measured outcomes comparing the pre- and post-procedure GOOSS were also obtained. From this, standardized mean difference was calculated and transformed to the natural logarithm before pooling, and the variance was calculated. Fixed-effects models were applied to pre- and post-EUS-GE data to determine effect size and corresponding 95% confidence intervals (CIs). Tabular and graphical analyses were performing using Comprehensive Meta-Analysis software, version 3 (BioStat, Englewood, New Jersey, United States). Additional analyses were performed with the aid of the Stata 13.0 software package (Stata Corp LP, College Station, Texas, United States).

Risk of bias and quality assessment

Risk of bias and quality of observational studies was evaluated using the Newcastle-Ottawa Quality Assessment Scale [18]. Domains assessed were selection of cohort, ascertainment of

exposure/comparator, and assessment of outcome. Based upon this, criteria score ≥ 5 was consistent with high quality, 3 to 4 medium quality, and ≤ 2 was considered low quality. Quality of randomized clinical trials (RCTs) was assessed using the JADAD score [19]. Two authors (TRM and RG) independently extracted data and assessed risk of bias and study quality for each of the articles. Any disagreements were resolved by discussion and consensus, and in consultation with the third reviewer (TR).

Investigations of heterogeneity

Heterogeneity was assessed for the individual meta-analyses using the chi squared test and the l^2 statistic [20]. Significant heterogeneity was defined as P < 0.05 using the Cochran Q test or $l^2 > 50\%$, with values > 50% indicating substantial heterogeneity. To assess for publication bias, a funnel plot was created and visually inspected for asymmetry and quantitatively using Egger regression testing [21, 22].

Results

Baseline study information and patient characteristics

This meta-analysis included a total of five studies including years from 2016 to 2019 [7,23–26]. A PRISMA flow chart of search results is shown in ▶ Fig. 1. Four studies were multicenter with two single-center trials also included. Four included

studies were retrospective in nature with an additional prospective study analyzed. A total of 199 patients were included in this study. Forty-five percent of patients were male. In 21.61% of patient with gastric outlet obstruction (n=43), the condition was secondary to benign causes. Mean age of patients that underwent the EUS-GE procedure for both benign and malignant gastric outlet obstruction was 64.52 ± 1.37 years with a pooled mean follow-up period of 4.32 ± 1.65 months. Further baseline study and patient characteristics are highlighted in \blacktriangleright **Table 1**.

Procedure characteristics of included studies

Both assisted and unassisted EUS-GE techniques were included with an overall mean procedure time for all techniques of 43.49 ± 19.69 minutes. The most commonly employed technique was direct EUS-GE (n = 134), followed by balloon-assisted (n = 35), then EPASS (n = 20), next ultra-slim (n = 5), afterwards nasobiliary drain (n = 3), and finally NOTES (n = 2). All studies included the CE-LAMS device; however, one study by Chen et al. also utilized the NCE-LAMS in 7.04% patients (n = 14). A total of 99.50% of patients (n = 198) had placement of a size 15 mm×10 mm LAMS for EUS-GE, with 10 mm×10 mm LAMS placement in only one patient. Additional procedure-specific characteristics are described in **Table 1**.

Efficacy and clinical effectiveness of EUS-guided gastroenterostomy

The immediate technical success of EUS-GE as reported in all included studies was 92.90% (95% CI; 88.26 to 95.79; $I^2 = 0.00\%$) (\blacktriangleright **Fig.2a**). Four studies documented clinical success with a pooled success rate of 90.11% (95% CI; 84.64 to 93.44; $I^2 = 0.00\%$) (\blacktriangleright **Fig.2b**). Clinical success as determined by the GOOSS was only reported in one study [27]. Based upon this study by Itoi et al., the mean post-GOOSS score was significantly higher compared to the pre-GOOSS score (2.94±0.23 versus 0.6±0.75; P<0.001) (**Supplemental Fig.1**). The median post-GOOSS score was significantly higher than the pre-GOOSS score (0.00 versus 3.00; P<0.001).

Safety of EUS-quided gastroenterostomy

Serious AEs occurred in 5.61% (95% CI; 2.87 to 10.67; $I^2 = 1.67\%$) of cases and were related to peritonitis, perforation, bleeding, and abdominal pain (\blacktriangleright Fig. 3a). Overall, the EA rate for the EUS-GE procedure was 10.59% (95% CI; 6.74 to 16.25; $I^2 = 27.17\%$) (Supplemental Fig. 2a). Minor AEs were reported in 5.78% (95% CI; 3.13 to 10.41; $I^2 = 0.00\%$) of procedures and were most commonly due to stent migration (Supplemental Fig. 2b). Need for repeat procedure or conversion to open surgical intervention was reported in four studies. Repeat intervention was required for 11.43% (95% CI; 7.29–17.46; $I^2 = 17.38\%$) of EUS-GE procedures (\blacktriangleright Fig. 3b).

Risk of bias assessment

All studies were evaluated using the Newcastle-Ottawa Quality Assessment Scale scores with quality assessment for each study shown in ► **Table 1**. All included studies were considered to be of high quality with scores ≥ 5. Despite the limited number of

included studies, publication bias was assessed [28]. Based on visual inspection of the funnel plot as well as quantitative measurement using the Egger regression test, there was no evidence of publication bias (> Fig. 4).

Discussion

This systematic review and meta-analysis demonstrates that EUS-GE is a technically feasible and effective endoscopic procedure for management of patients with benign and malignant gastric outlet obstruction. With an immediate technical success rate of 92.90% and a low 5.61% rate of serious AEs, EUS-GE appears to be relatively safe and well-tolerated. Given the significant morbidity of surgical gastrojejunostomy and high incidence of recurrent gastric outlet obstruction after endoscopic luminal stenting, EUS-GE shows promise as a fledgling procedure with the potential to become a first-line minimally-invasive therapy for patients with benign and malignant gastric outlet obstruction

Evolution of EUS-GE

First performed in a porcine animal model by Fritscher-Ravens et al. more than a decade ago, EUS-GE provided a new method for stitching under flexible EUS control but required endoscope exchange and use of special devices, thus limiting adoption in clinical practice [8,10]. However, development of a lumen-apposing self-expandable fully covered metal stent (AXIOS Boston Scientific Corp., Marlborough, Massachusetts, United States) able to safely appose two juxtaposed luminal structures and form an endoscopic anastomosis has brought new insights into development of EUS-GE [8,29].

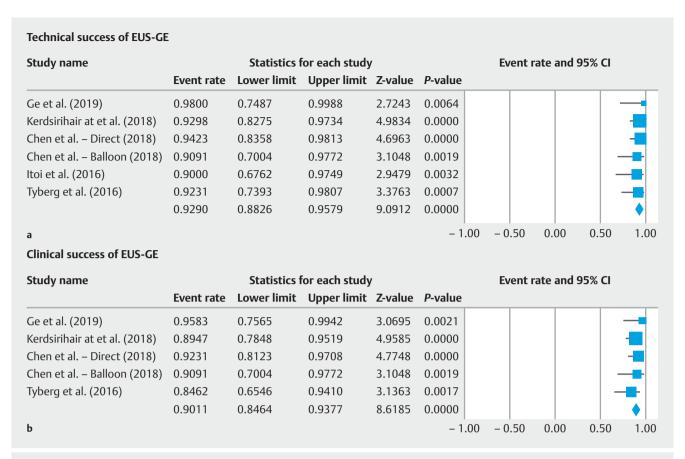
Although early animal models were successful, translation of these findings in humans has been challenging, represented by identification of the proper distal duodenal or proximal jejunal loop to be accessed from the gastric body to create the anastomosis [8, 30]. While a variety of EUS-GE techniques are in practice, addition of EUS allows for proper recognition of bowel structure, typically using a 19- to 22-gauge to puncture the loop, followed by contrast injection and guidewire placement [5,8]. Next, EUS-GE involves exchange with a CE-LAMS device to cautery puncture and dilate the tract, followed by distal flange deployment into the bowel lumen, withdrawal of the endoscope and the bowel loop toward the gastric cavity where the proximal flange is finally deployed. All included studies in this meta-analysis involved a CE-LAMS to perform EUS-GE, though one study included both CE-LAMS and NCE-LAMS [23].

Current standard of care and other available endoscopic treatments

Surgical gastrojejunostomy is the current standard of care for treatment of gastric outlet obstruction; however, the significant morbidity and high complication rate associated with surgery has necessitated a search for less invasive and better tolerated alternatives [6,7,9]. Endoscopic alternatives including placement of luminal enteral stents have been employed and have achieved comparable technical and clinical success rates with lower overall AE rates as compared to the traditional surgi-

Newcastle- Ottawa Quality As- sessment	5.5	6.5	7.0	7.0	5.0	0.9
No. of re-in- terven- tions	2/25	8/53	3/52	4/22	I	1/26
Minor adverse events	2/24	1/57	2/52	2/22	1/20	2/26
Serious adverse events	3/24	1/57	1/52	0/22	1/20	3/26
Clini- cal suc- cess	23/24	51/57	48/52	20/22	1	22/26
Tech- nical suc- cess	24/24	53/57	49/52	20/22	18/20	24/26
Pro- ce- dure time (min)	1	39	35.7	6.68	25.5	ı
No. of malig- nant ob- struc- tions	22	84	34	15	20	17
No. be- nign ob- struc- tions	0	0	18	7	0	o.
No. male- s (%)	o	28	22		10	Ξ
Mean age (years)	66.4	65	62.9	63.3		66.2
Follow- up period (mons)	1	6.5	3.8	3.8	3.3	÷.
No. pa- tients	22	57	52	22	20	26
Stent size, type	15×10mm CE-LAMS	15×10mm CE-LAMS	15×10mm CE-LAMS and 15× 10mm NCE-LAMS	15×10mm CE-LAMS and 15× 10mm NCE-LAMS	15×10mm CE-LAMS	and 10× 10 mm CE-LAMS
EUS-GE tech- nique	Direct $(n = 24)$	Direct (n=57)	Direct (n = 52)	Balloon- assisted (n = 22)	PASS (n = 20)	Balloon-assis- ted (n = 13); Ultra-slim scope (n = 5); Nasobiliary drain (n = 3); Direct (n = 3); NOTES (n = 2)
Study design	Single- center, retrospec- tive com- parator study	Multicenter, retrospective non-comparator study	Multicen- ter, retro- spective compara- tor study	Multicen- ter, retro- spective compara- tor study	Single- center, prospec- tive non- compara- tor study	Multicenter, retrospective non-comparator study
Year	2019	2018	2018	2018	2016	2016
Author	Ge et al.	Kerdsiri- hair at et al.	Chen et al. (Direct)	Chen et al. (Balloon)	Itoi et al.	Tyberg et al.
	Year Study EUS-GE tecch Stent size, No. Follow- No. be- No. be- No. of- Tech- Tech- Clini- Serious Minor No. of- design nique type period (years) s (%) ob- nant ob- dure suc- suc- events events terven- (mons) (mons) (years) s (%) ob- nant ob- dure suc- events events terven- (mons) (mons) tions tions (min) min) min) min) min)	Vear Study EUS-GE tech- design Stent size, tiones No. Follow- age Mo. be- male- mign No. be- male- mign No. be- male- mign No. be- male- malig- struc- stru	Figure F	Van Gesign Higher Higher	Figure Study Stu	1 1 1 1 1 1 1 1 1 1

CE-LAMS, cautery-enhanced lumen-apposing metal stent; NCE-LAMS, non-cautery-enhanced lumen-apposing metal stent; PASS, EUS-guided double-balloon-occluded gastrojejunostomy bypass.



► Fig. 2 a Immediate success rate for EUS-guided gastroenterostomy for the treatment of gastric outlet obstruction. b Clinical success rate for EUS-quided gastroenterostomy for the treatment of gastric outlet obstruction.

cal approach [31]. Nevertheless, enteral stenting provides a less durable option for patients with a prolonged life expectancy, plagued by frequent need for reintervention due to stent migration or stent tumor ingrowth/overgrowth [6, 9, 32].

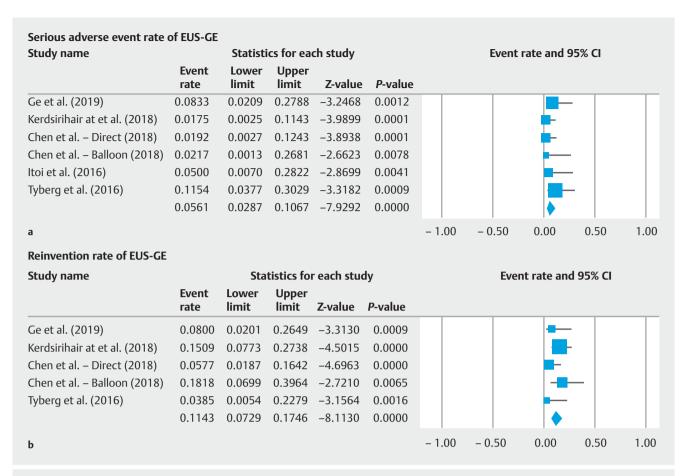
EUS-GE versus other treatments

Although evidence is limited, a recent study by Kashhab and colleagues comparing outcomes of EUS-GE to surgical gastrojejunostomy for treatment of malignant gastric outlet obstruction found the technical success rate was significantly higher in the surgical cohort (100% vs. 87%, P=0.009) [33]. While the surgical approach was more feasible and less technically challenging, the clinical success rate was no different between the surgical and EUS groups (90% versus 87%, P=0.18). There was also a favorable trend with lower recurrence of gastric outlet obstruction in the EUS-GE group (3%) as compared to the surgical group (14%); although this was not statistically significant (P=0.08). In another study by Manuel Perez-Miranda et al., technical success was nt different between an EUS-GE cohort and patients undergoing laparoscopic gastrojejunostomy (88% versus 100%, P=0.11), although EUS-GE was associated with a significantly lower rate of AEs (12% versus 41%, P=0.0386) [34]. Both of these studies were excluded from our systematic review and meta-analysis due to concerns regarding overlapping enrollment periods and duplication of patient data [26, 33, 34].

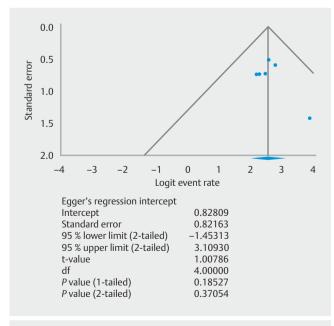
Another study by Chen et al. compared EUS-GE with endoscopic enteral stenting and demonstrated a significantly lower risk of recurrence and reintervention in the EUS-GE group compared to the enteral stenting group (4.3% vs 28.6%, P=0.015) [35]. Even on multivariable analysis, enteral stent placement was independently associated with need for reintervention (OR 12.8, P=0.027). These authors also found similar rates of technical success between the two strategies and a trend towards favorable clinical success among the EUS-GE group. While again not included in our meta-analysis due to concerns regarding overlapping enrollment periods, the results suggest EUS-GE provides longer-lasting luminal patency and reduces occurrence of stent obstruction, with a reasonable procedural risk, without the morbidity associated with a surgical procedure [8, 23, 35].

Clinical implications and future direction

Although these early data are promising and suggest that the procedure is safe and effective, there is not yet enough evidence at this time to endorse EUS-GE as the standard of care for gastric outlet obstruction [9]. Future, well-designed RCTs and prospective studies are needed to further validate these findings. Ultimately, more comparator studies are needed to



▶ Fig. 3 a Serious adverse events for EUS-guided gastroenterostomy for the treatment of gastric outlet obstruction. b Repeat interventions required after EUS-guided gastroenterostomy for the treatment of gastric outlet obstruction.



▶ Fig. 4 Funnel plot of publication bias and Eggers regression test for included studies to assess EUS-guided gastroenterostomy for the treatment of gastric outlet obstruction.

evaluate the efficacy and tolerability of EUS-GE as compared to enteral and surgical gastrojejunostomy as well as a comparison between various EUS-GE techniques and LAMS size. Additional limitations to mainstream use include the notion that the EUS-GE procedure is technically demanding, thus at present, it should be performed only by highly trained experts in interventional EUS [8]. It is important to note EUS-GE is not devoid of morbidity and AEs may be significant when they occur. Formal training and improved familiarity with the procedure overall will be required prior to seeing any meaningful clinical implication. Only then will the procedure likely see adoption into clinical practice.

Strengths and limitations

Specific limitations to this study include reliance largely on retrospectively collected data, differences in patient population across studies, lack of long-term follow-up periods, and inclusion of multiple EUS-GE techniques. While important to consider, heterogeneity of studies was low based on our analyses. In addition, while clinical success as determined by authors was reported in all studies, objective measures of clinical success (i.e. the GOOSS) were only reported in one study [27]. As the evidence stands now, there remains limited data regarding EUS-GE with sparse literature and numerous studies reporting

redundant patient information. Several studies in our literature search were excluded due to concerns regarding overlapping results from the same cohort of EUS-GE individuals. Although publication bias is not typically assessed with funnel plot asymmetry with fewer than eight to 10 studies in a meta-analysis, qualitative and quantitative publication bias was performed as pooled results of uncontrolled, retrospective studies carry a high risk of amplifying selection bias [28]. Despite this concern, our meta-analysis demonstrated no evidence of publication bias.

In addition, there remains a lack of data to compare EUS-GE directly with surgical gastrojejunostomy with only one study demonstrating similar efficacy among more complex patients [34]. With only five studies, subgroup analysis based upon EUS-GE technique or benign and malignant causes of gastric outlet obstruction, both of which would be highly clinically relevant, was not possible. An additional concern with any endoscopic procedure or technique is the learning curve or clinical expertise needed to perform an effective procedure [16]. It is possible the technical success achieved in these studies may be not be generalizable to centers with less familiarity or proven expertise.

Despite these limitations, this study has several strengths. Most importantly, our meta-analysis methodologically summarizes all available data to evaluate the feasibility, efficacy, and tolerability of the EUS-GE procedure. Overall, EUS-GE was demonstrated to be an effective device for gastric outlet obstruction in both short- and longer-term follow-up with impressive technical and clinical success rates and few serious AEs. Although we were unable to provide a subgroup analysis based upon gastric outlet obstruction etiology and EUS-GE technique, these findings provide an important step forward in proving the effectiveness and utility of further trials. With a relatively even distribution of benign and malignant gastric outlet obstructions included in this analysis, it is reasonable to assume EUS-GE is an effective modality for both etiologies. These data are pivotal to improve both symptom management and quality of life for individuals with gastric outlet obstruction, regardless of etiology, and may suggest an increasing role for EUS-GE.

Conclusions

Based upon our systematic review and meta-analysis, EUS-GE appears to be an effective and safe minimally invasive alternative for treatment of benign and malignant gastric outlet obstruction. Given the significant morbidity associated with surgical gastrojejunostomy reaching nearly 40%, less invasive options that provide durable and effective results will continue to be sought. Although EUS-GE remains a technically challenging procedure with limited evidence to date, it demonstrated high immediate technical and clinical success rates of 92.90% and 90.11%, respectively, with a low serious AE profile of 5.61%. While surgical gastrojejunostomy will continue to be the standard of care, this systematic review and meta-analysis suggests an increased role for EUS-GE in management of gastric outlet obstruction.

Competing interests

Dr. Thompson is a consultant for Boston Scientific, Olympus, and Apollo Endosurgery. Dr. Rustagi is a consultant for Boston Scientific and Cook Endoscopy.

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