

Safety and outcomes of endoscopic ultrasound-guided drainage for malignant biliary obstruction using cautery-enabled lumen-apposing metal stent



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ABSTRACT

Background and study aims There is increasing evidence to suggest that EUS-guided biliary drainage (EUS-BD) is a safe and effective treatment alternative for patients with

malignant biliary obstructions (MBOs) after failed endoscopic retrograde cholangiopancreatography.

Patients and methods We performed a retrospective analysis of data prospectively collected from patients with MBO who underwent choledochoduodenostomy (CDS) or gallbladder drainage (GBD) between August 2016 and June 2020 using the electrocautery-enabled lumen-apposing metal stents (ECE-LAMS). The primary endpoint was technical and clinical success. Secondary endpoints were adverse events (AEs) and reinterventions.

Results A total of 60 patients were included in the study, with 56 CDS and 4 GBD. Median age was 76 years with 57% male (34/60). The most common indication for EUS-BD was pancreatic cancer (78%). Technical success was achieved in 100% of cases, with a clinical success rate of 91.7%. Mean total bilirubin pre-procedure was 202 $\mu\text{mol/L}$ (normal < 20 $\mu\text{mol/L}$) and 63.8 $\mu\text{mol/L}$ post procedure ($P < .001$). Twenty-one patients had bilirubin recorded at 2 weeks post EUS-BD with 20 of 21 patients demonstrating >50% reduction in bilirubin (mean bilirubin reduction 75%). AEs occurred in 12 of 60 patients (20%), all of which were mild. The reintervention rate was 11.7% (7/60). Stent occlusion occurred in 10 of 60 patients (16.7%) with a mean time to stent occlusion of 46.2 days (3–133). Stent patency of 83.3% was observed with a mean follow up of 7.9 months.

Conclusion EUS-CDS and GBD using ECE-LAMS are effective EUS-based techniques for managing patients with MBO. AEs are usually mild and resolved by reintervention.

Introduction

Malignant biliary obstruction (MBO) is conventionally treated with endoscopic retrograde cholangiopancreatography (ERCP) and placement of either plastic or metal stents. ERCP remains the gold standard in treatment of biliary obstruction, however, failed biliary cannulation is observed in up to 10% of cases, especially in patients with low performance status [1]. Common causes of inaccessible papilla include advanced locoregional obstruction or altered anatomy due to prior surgery.

Percutaneous transhepatic biliary drainage (PTBD) is an effective biliary drainage technique that provides an alternative treatment option when ERCP fails. However, PTBD is associated with rates of adverse events (AEs) between 13 to 60% [2,3]. This is due to the risks of accidental dislocation of an external drainage tube and recurrent infections, which have a significant negative impact on quality of life. From a patient standpoint, the possibility of internal drainage is appealing due to convenience and likely improved quality of life, which is supported by a multicenter survey in which more than 80% of patients preferred EUS-BD over PTBD [4]. Historical surgical biliary bypass is

seldom used as the preferred treatment option because it is associated with higher morbidity and mortality [5, 6].

Endoscopic ultrasound-guided biliary drainage (EUS-BD) is an advanced endoscopic technique that was first described in 2001 [7]. With the advent of the linear echoendoscope and various EUS-specific tools including lumen-apposing metal stents (LAMS), patients with failed ERCP have increasingly been offered EUS-BD to gain biliary access [8]. There are conceptual advantages associated with the technique, including the minimally invasive nature of the procedure and single-session ERCP/EUS procedure, thus reducing procedural duplicity and expediting post-procedural recovery. The advent of an electrocautery-enabled delivery system (ECE-LAMS) has further evolved the procedure into a single-step, exchange-free EUS-guided puncture and deployment. The aim of this study was to evaluate technical efficacy and clinical safety of EUS-guided biliary drainage using the ECE-LAMS.

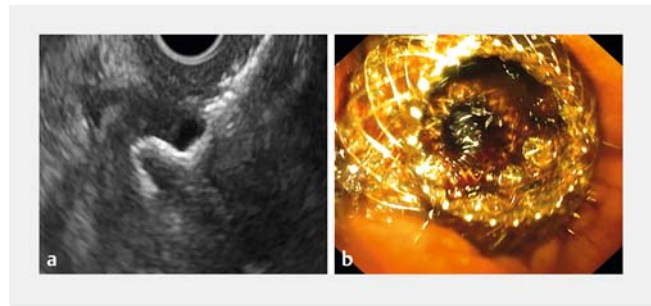
Patients and methods

We performed a retrospective analysis of prospectively collected data from all patients who underwent EUS-BD after failed ERCP from August 2016 through to June 2020. Patients underwent EUS-guided placement of a biliary stent, via a trans-duodenal approach, using the cautery-enabled LAMS system (Hot AXIOS, Boston Scientific). The procedures were classified as either choledochoduodenostomy (CDS) or cholecystoenterostomy for GBD. The study was approved by the institutional review board and granted exemption by the Health and Disability Ethics Committee.

Technical Description

All EUS-BD cases were performed under general anesthesia in a single endoscopy session, by a single endoscopist with high level of experience in therapeutic EUS procedures and in using ECE-LAMS. CDS was the preferred technique unless it could not be performed safely, defined by either the lack of a vessel-free window due to intervening vasculature or common bile duct (CBD) diameter < 10 mm.

The bile duct was accessed with EUS localization, via a trans-duodenal approach. A Doppler signal was used to identify a vessel-free window and the initial puncture was made with an electrocautery tip (ERBE setting: Autocut 80–100, Effect 5). The device was also preloaded with a 0.035 guidewire to allow immediate placement of the guidewire after initial puncture. Once the catheter tip was fully inside the CBD, the distal flange was deployed under EUS guidance (► Fig. 1a). Then the catheter was retracted and “snugged” against the proximal wall of the CBD to create wall apposition. The proximal flange was deployed within the working channel of the echoendoscope, which was then slowly withdrawn with simultaneous gentle advancement of the catheter to ensure intra-duodenal deployment of the proximal flange. Finally, the correct position of the LAMS was endoscopically confirmed with visualization of bile flow through the stent (► Fig. 1b). EUS-GBD was performed if CDS was not technically feasible. This was achieved by local-



► **Fig. 1** a Deployment of the distal flange under ultrasonographic guidance. b Endoscopic confirmation of successful deployment of the proximal flange and visualization of bile drainage.

ization of the gallbladder as the target puncture site, with similar techniques applied as described for CDS.

Outcome measurement

Patient demographics were reviewed, including age, sex, referral specialty, procedural indication, size of LAMS, total bilirubin (pre- and post-procedure), early and delayed adverse events (AEs), need for reintervention, and overall survival during the study period. Primary endpoints included technical and clinical success. Technical success was defined as achieving successful transluminal placement of bilioenteric fully-covered stents. Clinical success was defined as a reduction in absolute bilirubin > 50% at any time point. In addition, early confirmation of effective biliary drainage was defined by a reduction in bilirubin of ≥ 50% within 2 weeks of EUS-BD. Secondary endpoints included rate of AEs, need for reintervention, and stent patency. Stent patency was defined as absence of recurrent biliary obstruction post EUS-BD, or until patient death secondary to underlying malignant disease. Early (< 30 days) and late (> 30 days) AEs included procedure-related bleeding, reflux or stent-related cholangitis, stent occlusion, and patient death.

Results

Patient demographics

A total of 60 patients underwent EUS-BD using an AXIOS LAMS after failed ERCP with a median age of 76 years. Men represented 57% of the cohort while women represented 43%. Indications for performing EUS-BD included primary and metastatic malignancies. Baseline characteristics are listed in ► Table 1.

Technical outcome

Fifty-six patients underwent EUS-CDS while four EUS-GBD were performed due to lack of “vessel-free” window. The mean diameter of CBD was 17 mm (10–33). An 8-mm LAMS was used in 71.7% of cases (43/60) while a 10-mm LAMS was used in 28.3% (17/60). Technical success was achieved in 100% of cases. One case of stent malposition was encountered during initial deployment and was salvaged immediately by stent-in-stent placement, resulting in successful biliary drainage via the LAMS without subsequent adverse clinical events. A total of 14 patients (23.3%) had concomitant insertion of duodenal stents

► Table 1 Baseline characteristics.

Age	
Median	76
Maximum	90
Minimum	52
Sex (%)	
▪ Male	34 (57%)
▪ Female	26 (43%)
Referral (%)	
▪ Surgery	52 (86.6%)
▪ Oncology	5 (8.3%)
▪ Hematology	1 (1.7%)
▪ Medicine	1 (1.7%)
▪ Palliative care	1 (1.7%)
Indication, n	
▪ Pancreatic cancer	47
▪ Cholangiocarcinoma	3
▪ Lymphoma	2
▪ Duodenal cancer	2
▪ Metastatic cancer	6
▪ Colon	1
▪ Breast	2
▪ HCC	1
▪ Lung	1
▪ Melanoma	1
HCC, hepatocellular carcinoma	

along with EUS-BD. A summary of technical characteristics is shown in ► **Table 2**.

Clinical outcome

Clinical success, determined by improvement of liver function in bilirubin, was observed in 91.7% of patients with a mean bilirubin of 202 $\mu\text{mol/L}$ (normal $<20\mu\text{mol/L}$) pre EUS-BD and 63.8 $\mu\text{mol/L}$ post EUS-BD, representing a reduction of 68.4%. Twenty-two patients had bilirubin value recorded on Day 3 (mean bilirubin 86 $\mu\text{mol/L}$) while 11 patients had Day 7 bilirubin (mean bilirubin 43.5 $\mu\text{mol/L}$). Twenty-one patients (35%) had bilirubin value recorded at 2 weeks after EUS-BD. Normal bilirubin value was observed in 6/21 (40%) at 2 weeks with 20 of 21 patients (95.2%) achieving $>50\%$ reduction in absolute bilirubin value at 2 weeks post EUS-BD and a mean reduction of 75% (► **Table 3**).

► Table 2 Technical characteristics.

Failed ERCP (%)	100
Procedure	(n=60)
▪ CDS	56
▪ GBD	4
Mean CBD diameter (mm)	17 (10–33)
Stent size (n)	
▪ 8 mm	43 (71.7%)
▪ 10 mm	17 (28.3%)
Reintervention	9 (15%)
Duodenal stenting	14 (23.3%)
ERCP, endoscopic retrograde cholangiopancreatography; CDS, choledochoduodenostomy; GBD, gall bladder drainage; CBD, common bile duct	

► Table 3 Clinical outcome.

Liver function improvement (n=60)	
▪ Yes	55 (91.7%)
▪ No	5 (8.3%)
Total bilirubin ($\mu\text{mol/L}$)	
▪ Pre EUS-BD	202 (30–719)
▪ Post EUS-BD	63.8 (3–323)
<i>(P<.001)</i>	
Day 3 bilirubin, (n)	
▪ Mean bilirubin ($\mu\text{mol/L}$)	86
Day 7 bilirubin, (n)	
▪ Mean bilirubin ($\mu\text{mol/L}$)	43.5
Week 2 bilirubin, (n)	
▪ Mean bilirubin ($\mu\text{mol/L}$)	46.7
Patient mortality	
▪ Mean survival time after EUS-BD	177 days
EUS-BD, endoscopic ultrasound-guided biliary drainage	

Adverse events

The total AE rate was 20% (12/60). Early AEs (<30 days) were observed in seven patients with one case of maldeployment and six cholangitis events. Stent-related cholangitis were due to stent occlusion, and in all patients, reintervention with placement of seven French \times 4 cm double pigtail stents within the LAMS to facilitate effective biliary drainage was successful. In addition, four patients developed stent occlusion as a delayed AE (>30 days), which also required reintervention. The overall stent occlusion rate was 16.7% with a mean procedure to stent occlusion time of 46 days (median 21 days; 3–133

► **Table 4** Adverse events.

Total AE, n	12
Early (<30 days)	7
▪ Stent related cholangitis	6
▪ Stent maldeployment	1
Late (>30 days)	5
▪ Stent occlusion – food impaction	4
▪ Stent migration	1
Stent occlusion, total	10
Time	
▪ Early (<30 days)	(6)
▪ Late (>30 days)	(4)
Type	
▪ Food impaction	(8)
▪ Stone/sludge	(2)
Mean time to stent occlusion, days	46 (3–133)
Reintervention	9
AE, adverse event.	

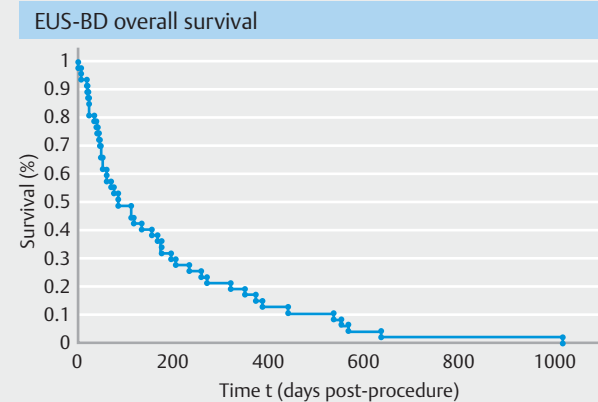
days). Nine patients underwent reintervention to rectify stent occlusion. Stent migration occurred in one patient with gall bladder drainage at 5 months after EUS-GBD. No significant stent-related bleeding had been recorded. ► **Table 4** provides a breakdown of the types of AEs encountered as well as A stent occlusion sub-analysis.

Follow up

The mean follow-up was 7.9 months. Stent patency was recorded in 83.3% of patients (50/60) at the end of the follow-up period. There was no observed procedure or stent-related mortality. During the study period, 48 patients died (80%) with a mean survival time of 177 days post EUS-BD. Overall survival is shown in the Kaplan-Meier curve in ► **Fig. 2**.

Discussion

In recent years, there has been an increasing body of evidence supporting high rates of safety and efficacy for EUS-BD in the short and medium term and technical success rates of 88% to 100% and clinical efficacy rates up to 97% [9–11]. The outcome of our study is consistent with other published studies using the same ECE-LAMS (Hot AXIOS, Boston Scientific, Massachusetts, United States). EUS-BD is associated with higher clinical success and lower AE and reintervention rates compared to PTBD [12]. However, there remains much debate about its role as first-line therapy rather than ERCP-BD. Recent meta-analyses that reviewed randomized trials comparing EUS-BD and ERCP-BD have shown no appreciable difference in technical and clinical success between the two treatment modalities [13, 14]. Fur-

► **Fig. 2** Kaplan-Meier curve of overall patient survival.

thermore, overall AE rates are lower for EUS-BD, including decreased risk of stent dysfunction (RR, 0.39), when compared with ERCP or PTBD [14]. One distinct advantage observed in EUS-BD is the significantly reduced rate of post-procedural pancreatitis, which is traditionally associated with ERCP (RR, 0.12) [14, 15]. Paik et al also reported lower rates of reintervention and higher stent patency, and therefore, improved patient-reported outcome measures in the EUS-BD cohort [15].

Our observational study has demonstrated high technical success (100%) and clinical efficacy (91.7%) for EUS-BD using ECE-LAMS in the setting of MBO and prior failed ERCP. We observed a mean bilirubin reduction of 68.4% across the study cohort. Furthermore, those who had bilirubin monitored experienced a reduction of >50% within the first 2 weeks, achieving a mean reduction of 75%. Therefore, our data demonstrate effective biliary drainage with both CDS and GBD. Most of the cases were performed with preloading of guidewire in the LAMS delivery system to allow for successful salvage procedure in case of potential maldeployment. This occurred in one case and rescue was immediate with placement of a tubular biliary stent across the newly created transduodenal fistula. An important technical consideration is site selection for stent deployment. The mean CBD diameter was 17 mm in our study (10–33 mm). A more dilated CBD means there is more space to allow the first flange of the stent to be deployed safely. Conventionally, the diameter of the CBD should be at least 10 mm to allow safe deployment of LAMS. In addition, we also found that “snugging” the deployed distal flange against the ductal wall, then deploying the proximal flange inside the delivery system prior to withdrawing the system ensures the correct positioning of the stent and minimizes maldeployment. One of the accepted indications for EUS-GBD is its application as an alternative to EUS-CDS in achieving endoscopic biliary drainage [16]. Technical success was achieved in all four patients who underwent GBD. EUS-GBD is a safe and effective endoscopic drainage option when CDS is not feasible.

The observed total AE rate in our study was 20% (12/60). This is similar to previous reports in various meta-analyses involving a heterogeneous cohort of stenting system [17, 18]. Im-

portantly, the AEs observed in our study were mild and did not result in prolonged hospitalization. This is favorable when compared to a surgical CDS with a morbidity rate of 30% [19]. One case of stent maldeployment was encountered in our study. This was due to maldeployment of the proximal flange with an 8-mm stent into a 10-mm CBD. This was immediately rectified by insertion of a 4-cm fully-covered, self-expanding stent through the lumen of the AXIOS stent to achieve effective biliary drainage, without further complications post procedure. It is worth noting that most of the AEs observed in our study were related to stent occlusion, either early (<30 days) or delayed (>30 days). The likely reason for cholangitis and delayed stent occlusion is related to the non-coaxial drainage achieved by the EUS-CDS. Historically coaxial placement of biliary stents with ERCP presented again with cholangitis due to stent occlusion (plastic vs self-expanding metal stent [SEMS] 21% vs 8%) [20]. Importantly, we did not encounter procedure- or stent-related hemorrhage in our cohort. Furthermore, there was no reported bile leakage or associated peritonitis in our study. The observed reintervention rate of 15% was predominantly related to occluded LAMS. Overall, the recorded mortality rate and survival time of just under 6 months in our study are consistent with other studies involving endoscopic management of MBO in patients with locally advanced or metastatic malignancies [21–23]. We did not observe any fatality related to EUS-BD.

The overall observed stent patency rate was 83.3% during the study period. It is worth noting that all of the stent occlusion events were related to food impaction. Six of the 10 cases of stent occlusion were associated with the 8-mm LAMS. We did not observe stent occlusion related to stent ingrowth, overgrowth, or migration. The encountered occlusions were effectively managed with insertion of plastic pigtail stents to facilitate ongoing biliary drainage. Most stent occlusions occurred in the early period of the study (<30 days). One US multicenter, retrospective analysis of EUS-CDS using ECE-LAMS demonstrated a significant reduction in rates of stent occlusion (50% to 11.8%) with preemptive placement of an axis-orienting stent through the LAMS [24]. This highlights the notion that LAMS can occlude early, and therefore, lends credit to prophylactic placement of axis-orienting stents through the LAMS lumen to prevent stent occlusion. There was also an association of LAMS obstruction in patients with concurrent duodenal obstruction [9]. In such scenario, one can also consider placement of a prophylactic axis-orientating double pigtail stent through the internal lumen of the LAMS at the time of initial deployment. We instituted routine prophylactic insertion of axis-orienting stents through the LAMS lumen in a more recent cohort of 10 patients undergoing EUS-BD. To date, we have yet to observe stent occlusion since this change in practice.

A unique clinical scenario in which EUS-BD may be preferentially selected as the primary treatment modality over ERCP-BD is in patients with concomitant gastric outlet obstruction. Though technically feasible, accessing the ampulla with a prior placed duodenal SEMS can be cumbersome, especially in type II obstruction. Achieving an appropriate endoscopic position can often be challenging, limiting the view of the ampulla. Previous studies have demonstrated significant variability in technical

success of ERCP performed through existing duodenal stents, with papillary cannulation rates between 34% and 85% [21, 25]. Conversely, single-session EUS-BD followed by duodenal SEMS deployment may be preferable and confers a higher success rate [9, 26, 27]. One of the major advantages of using ECE-LAMS is that it allows a single-step puncture to gain access to the biliary system, without the need for wire system exchanges, which confers a practical benefit in reducing the length of procedures. Anderloni et al reported reduced procedural time to 30 minutes in EUS-CDS while another study by Dollhopf et al reported a mean stent deployment time of just 3.1 minutes during EUS-GBD with direct access to the gallbladder [9].

To date, EUS-BD has been performed predominantly in patients with unresectable or advanced malignancy. However, the status quo is being challenged. Fabbri et al reported a small case series suggesting EUS-BD can be used safely as a bridge to surgical interventions such as pancreaticoduodenectomy (Whipple procedure) without significantly affecting outcomes [28]. Jacques et al also reported four patients who underwent successful pancreaticoduodenectomy after EUS-CDS and placement of LAMS [29]. We propose that selection of a first-line treatment modality should be based on clinical scenarios, ductal and luminal anatomy, and local expertise. Where possible, endoscopic interventions for MBO should be performed by expert endoscopists with significant experience in both advanced ERCP maneuvers and interventional EUS.

Our study does carry inherent limitations in that this was a retrospective analysis of patients who underwent EUS-BD in a single tertiary referral center. Our study was not designed to allow direct comparison between different treatment modalities including PTBD and EUS-guided biliary drainage using a non-LAMS stent. All procedures were performed by a single endoscopist with a high level of expertise in therapeutic EUS procedures and use of LAMS. Thus, our results may lack generalizability. However, it is our opinion that such a complex procedure should be performed only by experienced endoscopists in specialized centers with therapeutic EUS capability. We were not able to assess bilirubin response post procedure in every patient in a protocolized manner due to the retrospective nature of our study. A significant portion of our study population was referred from other institutions where the local referring physicians have varying practice in terms of monitoring for treatment response. These patients were transferred back to the referring institution once they recovered from the procedure.

Conclusion

In conclusion, EUS-BD performed with ECE-LAMS is a highly effective and safe treatment for patients who present with MBO. It can be performed in a single endoscopy session after failed ERCP. AE and reintervention rates are low and can be effectively managed conservatively. We believe that our findings will add to the currently growing evidence on the safety and efficacy of EUS-BD with ECE-LAMS, and will further support the use of EUS-BD as a safe and effective treatment option for a subset of patients with MBO in whom biliary cannulation cannot be achieved through conventional ERCP. However, larger random-

ized studies are needed to compare different drainage methods to produce more robust evidence and guide practice.

Competing interests

The authors declare that they have no conflict of interest.

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