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Lumen-apposing metal stent placement for drainage of pancreatic fluid collections: predictors of adverse events

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MESSAGE

Although rare in occurrence, adverse events such as delayed bleeding and buried stent syndrome have been reported after lumen-apposing metal stents (LAMS) placement in patients undergoing endoscopic ultrasound (EUS)-guided drainage of pancreatic fluid collections (PFCs). In a prospective study, we observed delayed adverse events in 6.4% of 188 patients which occurred when the PFCs were 7 cm or smaller in size and the removal of LAMS was delayed beyond 4 weeks.

IN MORE DETAIL

LAMS are being increasingly preferred over double pigtail plastic stents for patients undergoing EUS-guided PFC drainage because their deployment is technically easy and the wide lumen facilitates quick drainage of cyst contents. However, delayed adverse events such as bleeding and buried stent syndrome have been reported after LAMS placement.¹⁻³ In this study, we attempted to identify predictors of adverse events by examining data that were collected prospectively in all patients undergoing EUS-guided PFC drainage using LAMS. By institutional protocol, after LAMS placement, the endoprotheses were removed at outpatient follow-up in 3–4 weeks and all patients were contacted by telephone call to obtain follow-up at 6 months. The data collected included 292 demographic, laboratory, radiological, technical, clinical and treatment outcome variables, with a minimum follow-up duration of 6 months (NCT02422095).

Patient details, PFC characteristics, disease severity and clinical outcomes were summarised as means with SD and medians with IQR for continuous variables and as frequencies and proportions for categorical variables. In order to identify the factors associated with incidence of delayed adverse events after LAMS placement, multiple logistic regression and reverse stepwise multivariate logistic regression analyses were performed. Also, penalised logistic regression with Firth's correction was performed to identify the factors associated specifically with the incidence of delayed bleeding after LAMS placement. All clinically relevant variables including patient demographics, PFC characteristics, procedure details, LAMS indwelling time and disease severity were included as predictor variables. Datasets were compiled using Microsoft Excel (Microsoft, Richmond, Washington, USA), and all statistical analyses were performed using

Stata 14 (Stata, College Station, Texas, USA). Statistical significance was established as $p < 0.05$.

A total of 188 patients underwent EUS-guided drainage of PFCs (pseudocysts 31.4%, necrotic collections 68.6%) using LAMS over a 5-year period between 2015 and 2019. **Table 1** shows the patient demographics, preintervention PFC characteristics and disease severity of the study cohort. Adverse events were observed in 12 patients (6.4%, 95% CI 3.3% to 10.9%) that included delayed bleeding in eight (4.3%, 95% CI 1.9% to 8.2%) and buried stent syndrome in four (2.1%, 95% CI 0.6% to 5.4%) (**figure 1**). Bleeding was observed in a branch of the gastroduodenal artery or the splenic artery in five patients that were managed by interventional radiology-guided coil embolisation; mucosal bleeding was observed in three others that did not require further treatment. Buried stent syndrome was observed in the proximal stomach in four patients, which were successfully removed using endoscopic techniques in two patients. However, the stents could not be removed endoscopically in two patients as they were embedded in the deeper layers of the gastric wall and were referred for surgical removal.

The median time to adverse events after LAMS placement was 41 days (IQR 17–68), the majority ($n=8$, 66.7%) due to non-compliance with timely follow-up. On multivariable logistic regression analysis, after adjusting for patient demographics, PFC characteristics, procedure details, LAMS indwelling time and disease severity, only stent removal after 4 weeks (OR 4.60, 95% CI 1.30 to 16.3, $p=0.018$) and PFC size of ≤ 7 cm in the anteroposterior dimension at computed tomogram (OR 4.33, 95% CI 1.10 to 17.0, $p=0.036$) were predictive of adverse outcomes (**table 2**). Importantly, the PFC size was specifically predictive of delayed bleeding (OR 42.4, 95% CI 2.28 to 787.8, $p=0.012$).

COMMENTS

The present study demonstrates that adverse events can occur in up to 6% of patients undergoing EUS-guided drainage of PFCs using LAMS. This was observed when the endoprosthesis removal was delayed beyond 3–4 weeks when treating PFCs < 7 cm in size.

Unlike double pigtail plastic stents that tend to migrate out towards the gastrointestinal lumen when a PFC resolves, LAMS remain anchored in situ. It is postulated that the edges of the stent on persistent contact with vasculature adjacent to the



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Table 1 Baseline patient and pancreatic fluid collection characteristics

Age (years)	Mean (SD)	53.6 (14.2)
	Median (IQR)	55 (43.5–64.5)
Gender: n (%)	Female	59 (31.4)
	Male	129 (68.6)
Race: n (%)	Black	17 (9.0)
	White	152 (80.9)
	Other	19 (10.1)
Cause of pancreatitis: n (%)	Gallstones	39 (20.7)
	Alcohol	75 (39.9)
	Idiopathic	53 (28.2)
	Other*	21 (11.2)
Coexisting conditions: n (%)	Cardiovascular disease	23 (12.2)
	Pulmonary disease	17 (9.0)
	Renal disease	13 (6.9)
	Diabetes mellitus	49 (26.1)
ASA class: n (%)	I	13 (6.9)
	II	88 (46.8)
	III	78 (41.5)
	IV	9 (4.8)
CT severity index: n (%)	0–2	32 (17.0)
	4–6	58 (30.9)
	8–10	98 (52.1)
Type of pancreatic fluid collection: n (%)	Pseudocyst	59 (31.4)
	Acute necrotic collection	18 (9.6)
	Walled-off necrosis	111 (59.0)
Size of necrotic collection-AP axis (cm)	Mean (SD)	8.5 (3.7)
	Median (IQR)	7.8 (5.9–10)
Size of necrotic collection-transverse axis (cm)	Mean (SD)	11.0 (5.0)
	Median (IQR)	10 (7–14)
Percentage of necrosis: (%)	Mean (SD)	31.4 (22.6)
	Median (IQR)	30 (10–50)
Collection extending to lower abdomen/pelvis: n (%)		36 (19.1)
Disease severity: n (%)	SIRS	67 (35.6)
	ICU/high acuity care	73 (38.8)
Nutritional support: n (%)	Enteral feeding	38 (20.2)
	Parenteral feeding	9 (4.8)
	Oral diet	136 (72.3)
	Nil per os	5 (2.7)
Percutaneous catheter in situ prior to intervention: n (%)		9 (4.8)

*Other causes of PFC: Hypertriglyceridemia (n=8), malignancy (n=2), post-ERCP (n=2), post-trauma (n=9)

AP, anteroposterior; ASA, American Society of Anesthesiologists; ICU, intensive care unit; SIRS, systemic inflammatory response syndrome.

PFC cavity can cause erosion of vessels precipitating a bleeding episode. Likewise, when a PFC resolves, the mucosal overgrowth can bury the endoprosthesis deep within the gastrointestinal wall layers causing buried stent syndrome. Based on the results of a recent randomised trial, it has been suggested that the LAMS should be removed within 3–4 weeks, provided the PFC has resolved.⁴ However, it has been unclear as to which patient is particularly risk-prone and therefore will require close observation and follow-up.

The present study demonstrates that when a PFC is ≤ 7 cm in size, it is likely that the cyst contents drain rapidly after LAMS placement. In these patients, if the LAMS is not removed within a 3–4 week time frame, adverse events such as buried stent syndrome and particularly, delayed bleeding may occur. Therefore, patients with PFC less than 7 cm in size should be scheduled

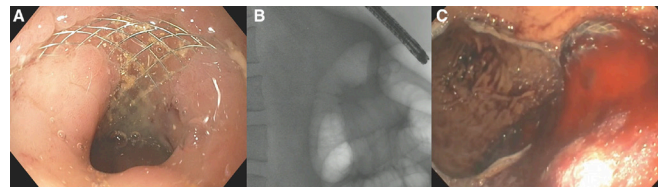


Figure 1 (A–C) Endoscopy view revealed mucosal overgrowth (A) over a buried LAMS, which was visualised at fluoroscopy (B). Endoscopic view of bleeding from LAMS (C). LAMS, lumen-apposing metal stents.

for follow-up imaging within 3–4 weeks and the LAMS must be removed in a timely manner if the fluid collection has resolved. Alternatively, small size PFCs may be treated using plastic stents in lieu of LAMS, particularly in patients with disconnected pancreatic duct syndrome who may benefit from an indwelling endoprosthesis to drain the upstream gland.

While the development of LAMS has significantly simplified the technique of PFC drainage and its utility for other applications such as gallbladder drainage, gastroenterostomy and biliary

Table 2 Multivariable logistic regression analysis examining factors associated with LAMS-associated adverse events

Variable	OR	95% CI	P value
<i>Multiple logistic regression analysis:</i>			
<i>Outcome variable=All LAMS-associated adverse events</i>			
Age: ≥ 60 vs < 60 years	1.23	0.33 to 4.63	0.762
Gender: Male vs Female	0.91	0.24 to 3.49	0.888
Race: Caucasian vs Non-Caucasian	0.96	0.18 to 5.28	0.966
PFC type: Necrotic collection vs Pseudocyst	0.79	0.13 to 4.86	0.802
PFC size: ≤ 7 vs > 7 cm	5.80	1.34 to 25.0	0.019
PFC location: Tail involved vs Tail not involved	1.15	0.21 to 6.40	0.871
Degree of necrosis: ≤ 40 vs $> 40\%$	1.52	0.30 to 7.61	0.608
Route of drainage: Proximal stomach vs Other	1.95	0.37 to 10.3	0.431
CT severity index: ≥ 8 vs < 8	1.07	0.18 to 6.20	0.943
ASA: III/IV vs I/II	2.80	0.63 to 12.4	0.174
Time to LAMS removal: > 4 vs ≤ 4 weeks	4.93	1.27 to 19.1	0.021
<i>Reverse stepwise multivariate logistic regression analysis:</i>			
<i>Outcome variable=All LAMS-associated adverse events</i>			
PFC size: ≤ 7 vs > 7 cm	4.33	1.10 to 17.0	0.036
Time to LAMS removal: > 4 vs ≤ 4 weeks	4.60	1.30 to 16.3	0.018
<i>Multiple penalised logistic regression with Firth's correction:</i>			
<i>Outcome variable=LAMS associated bleeding only</i>			
Age: ≥ 60 vs < 60 years	1.14	0.24 to 5.37	0.866
Gender: Male vs Female	0.34	0.07 to 1.74	0.197
Race: Caucasian vs Non-Caucasian	11.1	0.32 to 382.9	0.182
PFC type: Necrotic collection vs Pseudocyst	0.25	0.02 to 3.12	0.283
PFC size: ≤ 7 vs > 7 cm	42.4	2.28 to 787.8	0.012
PFC location: Tail involved vs Tail not involved	0.96	0.10 to 9.07	0.975
Degree of necrosis: ≤ 40 vs $> 40\%$	1.25	0.23 to 6.83	0.795
Route of drainage: Proximal stomach vs Other	5.76	0.79 to 41.9	0.084
CT severity index: ≥ 8 vs < 8	2.75	0.30 to 25.5	0.372
ASA: III/IV vs I/II	3.35	0.56 to 20.1	0.186
Time to LAMS removal: > 4 vs ≤ 4 weeks	1.54	0.35 to 6.78	0.566

ASA, American Society of Anesthesiologists; LAMS, lumen-apposing metal stent; PFC, pancreatic fluid collection.

bypass continues to evolve, a better understanding of the functionality and safety profile of the device is warranted. We believe that the observations detailed in this newsletter helps bridge some aspects of this knowledge gap.

Contributors JYB: study design, endoscopist performing procedures in the study, statistical analysis, interpretation of data, drafting of manuscript, critical revision of manuscript. SV: study concept and design, endoscopist performing procedures in the study, interpretation of data and drafting of manuscript and critical revision of manuscript. RH: endoscopist performing procedures in the study and critical revision of manuscript.

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