Endoscopic ultrasound characteristics of pancreatic lymphoepithelial cysts: A case series from a large referral center

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ABSTRACT

Background and Objectives: Lymphoepithelial cysts (LECs) of the pancreas are benign lesions that can mimic cystic neoplasms on imaging. Endoscopic ultrasound (EUS) features have not been well described. We aimed to describe the clinical and EUS characteristics of LECs and the present outcomes of management at a high-volume referral center.

Materials and Methods: We identified patients who underwent EUS and were found to have LECs based on fine-needle aspiration (FNA) cytology or surgical pathology from existing databases. EUS features, imaging characteristics, and pathology results were described. Results: Sixteen patients were found to have 17 LECs. The mean size was 33 mm ± 15 mm. Locations within the pancreas included 10 lesions in the tail, 3 in the body, 1 in the uncinate process; the remaining 3 were exophytic. Six lesions were anechoic, 6 were hypoechoic, and 5 had mixed echogenicity. Nine lesions had mixed solid/cystic components, 7 were purely cystic, and 1 was solid. Cyst fluid was thick or viscous in six cases and thin in three. Eleven patients had diagnostic cytopathology. Six patients ultimately underwent surgery due to symptoms, nondiagnostic FNA, or other clinical concerns for malignancy. Conclusions: Pancreatic LECs have variable morphology and echogenicity on EUS, but the appearance of a cyst with variable solid and cystic components combined with the appearance of thick, turbid, and viscous aspirate should raise suspicion for an LEC. The majority of patients with LECs at our center avoided surgery for LECs on the basis of diagnostic EUS-FNA.

Key words: Endoscopic ultrasound (EUS), lymphoepithelial cyst (LEC), pancreatic cyst

INTRODUCTION

Lymphoepithelial cysts (LECs) of the pancreas are benign lesions with no malignant potential. The imaging appearance can mimic other cystic neoplasms, and therefore distinguishing LECs and other benign lesions from those with malignancy or malignant potential is important. This is underscored by the fact that pancreatic cysts are...
increasingly detected due to the widespread use of cross-sectional imaging. However, literature describing imaging and clinical features of LECs is limited to case reports and small case series. Thus, features of such lesions may be less known compared to other lesions of the pancreas.

Clinical and imaging findings of LECs are typically nonspecific. Definitive diagnosis has traditionally relied on surgical histopathology following resection. It has been shown that benign cysts are often indistinguishable from cystic neoplasms on pre-operative imaging, and therefore the patients often undergo unnecessary resection. Carcinoembryonic antigen (CEA) levels may also be elevated in fluid aspirated from confirmed LECs, which may initially raise the suspicion for a mucinous neoplasm. Endoscopic ultrasound (EUS) with fine-needle aspiration (FNA) cytopathology may help diagnose LECs, but cyst cytology may be difficult to interpret due to contamination of the aspirate by intestinal tissue with mucinous and glandular epithelium. Additionally, the absence of classic cytological findings of LEC does not exclude the diagnosis. EUS may be able to further characterize such lesions based on morphology and cyst fluid analysis. While EUS features of LECs have been previously described, there remains an overall dearth of literature pertaining to EUS findings. Additional studies are needed to better characterize the EUS features of these lesions in order to lend support for their diagnosis and to help avoid unnecessary surgery. In this series, we describe the clinical and EUS characteristics of LECs confirmed on cytology and/or surgical pathology, and present outcomes of management at a high-volume referral center.

MATERIALS AND METHODS

At a single tertiary referral center, EUS, surgical, and cytology databases were accessed and retrospectively reviewed to identify the appropriate patients. All the patients who were diagnosed with pancreatic LECs based on definitive cytopathology or surgical resection pathology between 2004 and 2014 were included in this case series. Each EUS procedure was performed by one of six endosonographers. The EUS images of the LECs were pulled and retrospectively reviewed by an expert endosonographer (MAH) blinded to the original EUS reports to assess for the size, location, border, echogenicity, internal components, and presence of septations. Similarly, reports from computed tomography (CT) or magnetic resonance imaging (MRI) acquired before EUS were retrospectively reviewed by a radiologist to determine the location and other radiological features including exophytic appearance. The results of any imaging studies performed in patients managed nonoperatively were also reviewed to assess for changes in the size or morphology of the lesions. Cyst fluid physical characteristics as well as cyst fluid amylase and CEA levels were also retrieved when available. The Institutional Review Board of our medical center approved this study.

FNA cytology results were reviewed and confirmed by an expert cytopathologist. The presence of anucleated and/or nucleated squamous cells admixed with a variable number of lymphocytes in a background of keratinaceous and/or amorphous debris on FNA smears was considered diagnostic of an LEC [Figure 1]. In cases where FNA cytology was nondiagnostic, surgical resection pathology reports were reviewed to confirm the diagnosis of LEC.

RESULTS

Sixteen patients (12 males; mean age 57 ± 13 years) with 17 confirmed LECs underwent EUS (1 patient was found to have 2 distinct LECs) [Table 1]. Presenting symptoms included: Abdominal pain in six patients and weight loss in two patients; eight patients were asymptomatic with incidental lesions on cross-sectional imaging performed for unrelated indications. Reports from imaging obtained prior to EUS were available in 13 patients (CT in 6, MRI in 2, and both in 5 patients), accounting for 14 lesions. On CT or MRI, lesions were predominantly located in the tail (N = 8) or body (N = 5); nine of these lesions were exophytic.
Dalal, et al.: EUS Characteristics of pancreatic lymphoepithelial cysts

but originated from the tail or body of the pancreas (Figure 2 for representative CT images of case 5).

On EUS [Table 1], the mean LEC size was 33 ± 15 mm. Location of LECs within the pancreas were as follows: 10 were in the tail, 3 in the body, 1 in the uncinate process, and 3 were described as exophytic (2 in the body and 1 in the tail). Borders were well-defined in 16 and poorly defined in 1. Echogenicity was described as anechoic in 6, hypoechoic in 6, and mixed in 5 patients. Representative EUS images are shown in Figure 3.

Internal components within the lesion were also characterized by EUS. Nine lesions had mixed solid/cystic components (of which 2 were mostly solid and 1 had calcifications), 7 were purely cystic (2 had internal debris), and 1 was solid with no cystic components. Internal septations were noted in 6 (5 with thin or incomplete septations and 1 with thick septations); septations were suggested on imaging in two additional cases, but were not confirmed on EUS. No wall nodularity or pancreatic ductal dilation was noted on EUS, though a mural nodule was seen on MRI in one lesion.

When adequate cyst fluid was aspirated, its consistency was described as thick, mucoid, and/or viscous in six cases and thin in three cases; color was clear in three cases and turbid/opaque in three cases [Table 2]. Median cyst fluid CEA (N = 5) and amylase (n = 6) were 2.4 ng/mL (range 1.3-13,088) and 100 U/mL (range 5-1111), respectively. Eleven LECs (from 11 distinct patients) were diagnosed on the basis of characteristic FNA cytology [Table 2]. In these cases, cytologic exam of the aspirated cyst fluid revealed a combination of nucleated or anucleated squamous cells, keratinous debris, and a variable number of lymphocytes (ranging from scattered to numerous). In a diagnostic FNA case, cholesterol crystals were seen as well. Six cases from five patients had nondiagnostic cytology (five cases with findings of benign, indeterminate cysts based on FNA alone and one case with hypocellular aspirate); the diagnosis of LEC in these cases was established by subsequent surgical resection pathology [Figure 4].

Six patients accounting for seven cysts ultimately underwent surgical resection by laparoscopic or open distal pancreatectomy. Two of these patients (totaling 3 cysts) required surgery for abdominal pain attributed to the cyst (cases 7 and 10). In another patient (case 4), FNA cytology was diagnostic for LEC; however,
### Table 1. Imaging and EUS features of pancreatic LECs

| Case no. | Sex | Age at EUS (yr) | Pre-EUS Imaging | Location on imaging | Size on EUS (mm) | Location in pancreas on EUS | EUS findings |
|----------|-----|----------------|-----------------|---------------------|-----------------|-----------------------------|--------------|
| 1        | F   | 43             | CT              | tail*               | 50×30           | body*                       | well-defined, anechoic, cystic |
| 2        | F   | 84             | n/a             | body*               | 40×32           | body*                       | well-defined, anechoic, thin outer wall, cystic, and thin/incomplete septations |
| 3        | F   | 44             | CT              | body                | 30×18           | tail                        | well-defined, anechoic, cystic with internal solid debris |
| 4        | F   | 66             | n/a             | body                | 18×8            | tail                        | well-defined, mixed echogenicity, heterogenous, calcified, mixed solid/cystic |
| 5        | M   | 65             | CT              | body                | 22×18           | body                        | well-defined, anechoic, cystic, thinly septated with two compartments |
| 6        | M   | 48             | CT/MRI          | tail                | 28×28           | body                        | well-defined, hypoechoic, heterogeneous, cystic |
| 7        | M   | 65             | MRI             | body                | 29×23           | body                        | well-defined, hypoechoic, cystic with internal debris, thin/incomplete septations with few compartments |
| 8        | M   | 56             | CT              | uncinate process    | 50×30           | uncinate process            | poorly defined, hypoechoic, heterogenous, mixed solid/cystic, incomplete septations |
| 9        | M   | 50             | CT/MRI          | tail                | 36×33           | tail                        | well-defined, mixed echogenicity with hyperechoic foci, heterogeneous, cystic with mostly solid components |
| 10†      | M   | 37             | CT              | tail                | 24×20           | tail                        | well-defined, mixed echogenicity, mixed solid/cystic |
| 11†      | M   | 37             | CT              | tail                | 25×20           | tail                        | well-defined, mixed echogenicity, mixed solid/cystic |
| 12        | M   | 73             | MRI             | tail                | 23×9            | tail                        | well-defined, hypoechoic, cystic component not seen |
| 13        | M   | 54             | CT/MRI          | tail                | 28×24           | tail                        | well-defined, hypoechoic, heterogenous, mixed solid/cystic but mostly solid, thin septations |
| 14        | M   | 40             | CT/MRI          | tail                | 6×7             | tail                        | well-defined, anechoic with hyperechoic attenuation deep to cyst, cystic |
| 15        | M   | 58             | CT/MRI          | body                | nos              | tail                        | well-defined, hypoechoic, mixed solid/cystic, thick septations with multiple small cystic lesions |
| 16        | M   | 62             | CT/MRI          | body                | 58×42           | body                        | well-defined, anechoic, complex mass, mixed solid/cystic with internal debris |
| 17        | M   | 52             | n/a             | body                | 60×45           | tail                        | well-defined, mixed echogenicity, mixed solid/cystic |

*Exophytic on imaging/EUS, †Denotes same patient, n/a: Not available, nos: Not otherwise specified

### Table 2. FNA and cytology features

| Case no. | Aspirate description       | Cytologic description                                                                 | Diagnosed by | Surgical resection |
|----------|----------------------------|--------------------------------------------------------------------------------------|--------------|--------------------|
| 1        | blood-tinged, thin, colorless | Scattered benign lymphocytes                                                           | FNA          | No                 |
| 2        | opaque, blood-tinged, thin  | Numerous lymphocytes                                                                  | FNA          | No                 |
| 3        | clear, yellow, thin         | Benign lymphocytes and benign epithelium                                             | FNA          | No                 |
| 4        | nos                        | Fragment of squamous epithelium and underlying lymphoid cells                        | FNA          | Yes*               |
| 5        | clear, white, slightly viscous | Benign nucleated and anucleated squamous epithelial cells, keratinous debris, rare scattered lymphocytes | FNA          | No                 |
| 6        | nos                        | Keratin debris, lymphocytes, squamous epithelium                                      | FNA          | No                 |
| 7        | thick, turbid, white        | Occasional acute inflammatory cells, abundant proteinaceous debris, scattered cholesterol crystals | Surgical pathology | Yes |
| 8        | bloody                     | Abundant nucleated squamous cells, scant lymphocytes                                   | FNA          | No                 |
| 9        | liquid, serosanguinous, somewhat thick, mucoid | Abundant anucleated squamous cells, macrophages, crystalline and scattered inflammatory cells | FNA          | No                 |
| 10†      | turbid, white, slightly viscous | Anucleate squames                                                                     | Surgical pathology | Yes |
| 11†      | nos                        | Anucleate squames                                                                     | Surgical pathology | Yes |
| 12        | nos                        | Hypocellular, fragments of contaminating epithelium                                   | Surgical pathology | Yes* |
| 13        | nos                        | Hypocellular, amorphous debris                                                       | Surgical pathology | Yes* |
| 14        | clear, slightly viscous     | Numerous lymphoid cells, macrophages; no epithelial component                         | FNA          | No                 |
| 15        | mucoid-type material        | Abundant squamous epithelial cells                                                    | Surgical pathology | Yes* |
| 16        | nos                        | Abundant debris and anucleate squamous cells                                          | FNA          | No                 |
| 17        | nos                        | Low cellularity, anucleate squamous cells, keratinaceous debris, cholesterol crystals, scattered lymphocytes | FNA          | No                 |

*Malignancy suspected based on indeterminate clinical or imaging findings, †Denotes same patient, nos: Not otherwise specified
clinical suspicion for a neuroendocrine tumor remained high due to symptomatic hypoglycemia, thus this patient underwent surgical resection. Surgical pathology confirmed LEC in this patient. The remaining three patients underwent surgery for suspected malignancy based on indeterminate clinical and imaging findings and nondiagnostic FNA cytology; one underwent exploratory surgery after imaging suggested a malignant right adrenal lesion with liver invasion (case 12); one was found to have a hypermetabolic focus in and adjacent to the pancreatic cyst (case 13); one had benign FNA cytology, although mutational analysis on the aspirated fluid was performed at a commercial laboratory and revealed several allelic imbalances suggestive of malignancy (case 15). Of the 10 patients who did not have surgical resection, 4 had subsequent imaging available a median 39 months (range 19-62) after EUS; all 4 showed stable (3 cases) or decreased (1 case) cyst size.

DISCUSSION

EUS-FNA is generally recommended for the evaluation of pancreatic cysts detected on imaging to better characterize these lesions and to avoid unnecessary surgical resection of benign cysts. In the current EUS-based case series, which is the largest to date, we found that the majority of patients with LECs at our center avoided surgery for these benign lesions on the basis of diagnostic EUS-FNA.

In our series, the majority of patients were middle-aged males, with a male:female ratio of 3:1 and a mean age of 57 years. Our patient demographic was similar to that described by Yanagimoto et al., who reviewed 106 cases of LECs and found the mean age of presentation to be 56 years, with a male:female ratio of 4:1. The majority of our patients were asymptomatic, while abdominal pain was the most common presenting symptom as has been previously described by others.

Four patients underwent surgery for suspicion of malignancy including one with diagnostic FNA for LEC and three with nondiagnostic cytology. Asymptomatic patients with nondiagnostic cytology who have EUS and cytology features suggestive of an LEC may benefit from observation rather than surgical resection.

We found that EUS morphology of LECs varies considerably from a predominantly solid to purely cystic or a mixed solid/cystic appearance. Lesions were mostly well defined, predominantly anechoic or hypoechoic, and often exophytic on pre-EUS imaging, which should raise the pre-EUS probability of LEC. In our series, the majority of the lesions were localized to or exophytic from the tail of the pancreas, whereas previous studies have reported equal distribution throughout the pancreas.

In a prior EUS-based case series of nine patients, Nasr et al. similarly demonstrated diverse sonographic characteristics of LECs on EUS. The majority of the LECs in that series appeared hypoechoic and solid, with subtle posterior enhancement suggestive of a cystic component. This agrees with our findings of several cases with mixed solid/cystic components. Also, while the majority of our cases had hypoechoic or anechoic cysts on EUS, five cases were noted to have mixed hypo or hyper-echogenicity and another three cases were noted to have echogenic internal debris. Hypoechoic lesions with hyperechoic content has been described in cases outside our series, thus internal debris within the cyst likely account for mixed echogenicity seen in our cases.

On the basis of gross cyst fluid appearance, thick, turbid, and viscous appearing fluid was suggestive of LECs, as reported in 6 out of 10 cases in this series. Similarly, Nasr et al. showed that a thick milky, creamy, or frothy aspirate should raise suspicion for LECs. Cyst fluid CEA and amylase levels were highly variable in our series, as has been previously demonstrated. Therefore, these may not be useful markers.

In a single patient in our series, two distinct LECs were ultimately diagnosed based on surgical pathology, though preoperative FNA cytology described a differential diagnosis for a benign cyst including a dermoid or epidermoid cyst (cases 10 and 11). EUS reports of pancreatic dermoid cysts are rare, with isolated case reports describing a multilocular hypoechoic appearance or complex, honeycomb lesion with histiocytes, benign epithelial cells, and lymphocytes seen on cytology. The ability to discern LEC from other benign cysts on EUS, particularly dermoid and epidermoid cysts, can be difficult and requires further studies.

With the frequent detection of pancreatic cysts on imaging, it is likely that LECs will be increasingly reported and characterized in the literature. We anticipate that EUS will continue to play an important role in characterizing these lesions.
role in conjunction with cross-sectional imaging and FNA cytology to differentiate the benign lesions from malignant ones, and help avert surgery in asymptomatic patients. The appearance of an exophytic cyst with variable solid and cystic components combined with the appearance of thick, turbid, and viscous aspirate should raise suspicion for an LEC.

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Conflicts of interest
None relevant to this manuscript.

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