Malignant transformation of a duodenal duplication cyst in a cat

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Abstract

Case summary A 15-year-old domestic shorthair cat presented for lethargy, vomiting and anorexia. Abdominal ultrasound showed a bi-lobed cystic duodenal mass. Based on ultrasonographic features, malignant transformation of a duodenal duplication cyst was suspected. A resection and anastomosis was performed. Histology of the mass was consistent with carcinoma and an intestinal mucosa was present along the inner surface of the mass, suggestive of a duplication cyst. The patient returned 3 months postsurgery with recurrence of clinical signs. Abdominal ultrasound showed a recurrent duodenal mass at the surgery site and body wall nodules. Fine-needle aspirates of these lesions showed epithelial neoplasia. Owing to the poor prognosis, the owners elected euthanasia.

Relevance and novel information Malignant transformation of duplication cysts in cats is rare but can be detected on ultrasound. The described ultrasonographic features can aid in prioritization of malignant transformation of duplication cysts as a differential diagnosis.

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A 15-year-old castrated male domestic shorthair cat presented to the Foster Hospital for Small Animals at the Cummings School of Veterinary Medicine for a 3 day history of lethargy and 2 day history of vomiting and anorexia. No abnormalities were noted on the referring veterinarian’s blood work. Radiographs performed by the referring veterinarian showed a round soft tissue opaque cranial abdominal mass that was superimposed on the pylorus, and the stomach contained a moderate volume of fluid. Abdominal ultrasound showed a bi-lobed cystic mass-like structure that was closely associated with the duodenum (Figure 1a). The walls were echogenic and variably thickened (up to 9 mm; see Figure 1a), and a hypechoic focus was seen within the wall of the mass. Focal loss of wall layering at the junction between the cavitated mass and the duodenum was identified. The adjacent duodenal segment had thickened walls, focal loss of wall layering and a narrowed lumen (Figure 1b). The duodenal papilla was not identified in the region of the mass. The pancreas also appeared to be displaced by the mass but was otherwise normal. The remainder of the abdominal ultrasound was within normal limits.

Based on the ultrasonographic appearance of the bi-lobed cystic mass, a duodenal duplication cyst was suspected. The loss of wall layering was supportive of neoplasia. Fine-needle aspiration of the wall of the mass was performed and fluid was removed from the cavity of the mass. Cytology of the mass revealed an epithelial neoplasm with squamous cell differentiation. The fluid portion of the mass revealed necrosis and cyst formation with chronic haemorrhage. Thoracic radiographs were performed to evaluate for evidence of metastasis. These showed a bronchial pattern, mild cardiomegaly and no evidence of pulmonary metastatic disease. An echocardiogram was performed and showed hypertrophic cardiomyopathy with no need for medical management. Exploratory laparotomy was considered but not performed at this time, as the patient was clinically improving with intravenous fluids and antinausea medications.

The patient returned 6 days later for recurrent anorexia and vomiting. Repeat ultrasound showed evidence of mechanical obstruction with a fluid-distended stomach and distended duodenum oral to the suspected malignant transformed duodenal duplication cyst.

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Exploratory laparotomy was performed. At surgery, a bi-lobed mass was seen associated with the duodenum. The duodenal papilla was located approximately 3 cm oral to the mass, and a resection and anastomosis of the cystic duodenal mass was performed with no complications (Figure 2). Histology of the mass revealed an adenocarcinoma with lymphatic invasion. Intestinal mucosa was present on the inner surface of the bi-lobed cystic region, supportive of a duplication cyst. Grossly, no connection between the intestinal lumen and cavitated portion of the duplication could be found. Focal mineralization was present within the wall of the mass on histology, which likely represented the hyperechoic focus seen on ultrasound. The patient did well postoperatively and went home. The owners elected not to pursue chemotherapy.

Approximately 3 months after surgery, the cat returned for recurrent vomiting. Ultrasound was performed and a 2.5 cm, poorly echogenic mass was seen at the duodenal surgery site, and loss of intestinal wall layering was noted at this level. Additionally, three abdominal wall nodules were seen. Fine-needle aspirates of the duodenal mass and wall nodules were compatible with epithelial neoplasia. Given the progression of disease, the owners elected for humane euthanasia.

Enteric duplications are rare congenital abnormalities reported in humans, cats and dogs. Enteric duplications are thought to be the result of abnormal recanalization of the gastrointestinal tract. The mucosal layer of enteric duplications is most often similar to adjacent intestinal mucosa. Duplication cysts can occur anywhere along the gastrointestinal tract, and can be round or tubular in appearance. In humans, duodenal duplication cysts are rare and account for approximately 5% of enteric duplications. Children and adults often present with signs of obstruction or a painful abdomen. In veterinary patients, clinical signs are often absent. One case report of a rectal duplication cyst in a cat resulted in tenesmus and constipation due to...
compression of the rectum.\textsuperscript{10} In humans and dogs, an ultrasonographic feature of duodenal duplication cysts is a cystic mass that is contiguous with an intestinal segment,\textsuperscript{9,11,12} thus, giving it the appearance of two non-communicative bowel segments adherent to one another. In humans and dogs, the presence of a muscular rim sign is suggestive of a duplication cyst.\textsuperscript{9,12} The muscular rim sign is seen as a hypoechoic muscularis layer that surrounds an inner echogenic mucosa.\textsuperscript{12}

In humans, malignant transformation of duplication cysts is suspected when there is loss of the muscular rim sign and thickening of the normally thin-walled cyst.\textsuperscript{11,12} This is similar to the present case, where the wall of the cystic mass was irregularly thickened with loss of wall layering (Figure 1a). Therefore, wall layering of duodenal duplication cysts should be carefully evaluated, as loss of wall layering is supportive of neoplasia. Additionally, there was concurrent loss of wall layering of the adjacent duodenum (Figure 1b). Loss of gastrointestinal wall layering on abdominal ultrasound is a primary feature of gastrointestinal neoplasia.\textsuperscript{13,14} However, cystic dilation of gastrointestinal masses has not been reported as a primary feature of gastrointestinal neoplasia.\textsuperscript{13,14} For this reason, we suspected the duodenal mass in the current case was a malignantly transformed duplication cyst. In the current case, we speculated that the clinical signs were likely due to the presence of carcinoma restricting the lumen of the duodenum and not the malignantly transformed duodenal duplication cyst, as intestinal carcinoma has been reported in cats to cause mechanical obstruction due to a constrictive lesion.\textsuperscript{13}

In humans, malignant transformation of duplications cysts is rare, with only 30 cases being reported in the literature.\textsuperscript{15} In the veterinary literature, duplication cysts have historically been thought of as an incidental finding. However, the current case presents the possibility of duplication cysts undergoing malignant transformation. For this reason, surgical removal of duplication cysts could prevent this rare occurrence.

Conclusions
Malignant transformation of a duodenal duplication cyst is possible in cats and can be detected on ultrasound. Malignant transformation of duplication cysts should be considered as a differential for a cystic mass with loss of wall layering associated with an intestinal segment or if clinical signs develop in a patient with a previously diagnosed incidental duplication cyst.

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