**Biloma: An unusual complication in a patient with pancreatic cancer**

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**INTRODUCTION**

In 1979, Gould and Patel\(^1\) described the case of a 32-year-old man who was found to have a large encapsulated collection of bile outside the biliary tree, secondary to a tear in the right hepatic duct. This followed a traumatic insult to the upper abdomen. The patient proceeded to have this collection drained and a T-tube was inserted to permit bile drainage. Since the initial description, there have been a few documented cases of spontaneous biloma formation, usually in the context of choledocholithiasis. The current case is sufficiently exceptional with regard to location, etiology and mode of presentation to merit a report.

**CASE REPORT**

A 64-year-old man was referred to the gastroenterology clinic by his general practitioner (GP) with constant upper abdominal pain that was worse after eating. In addition, the patient had been suffering from alternating constipation and loose stools. He denied the passage of blood in his motions or any weight loss.

A full blood count revealed normocytic anemia (hemoglobin 123 g/L; mean corpuscular volume: 88.4 fL). Renal and liver function tests were normal. Given the features on presentation, a colonoscopy was suggested, however, he was reluctant to go ahead with the procedure. As an alternative, a computed tomography (CT) pneumocolon was arranged.

In the absence of significant polyps or other colonic abnormalities were seen, there was an irregular 3 cm × 4.5 cm mass arising from the neck/body of the pancreas, with multiple lymph nodes in the peripancreatic region. Further assessment with a dual-phase CT scan of the pancreas was performed (Figure 1). As a result of encasement
of the superior mesenteric artery and portal veins, the tumor was deemed inoperable. An ultrasound-guided biopsy of the lesion was undertaken, and histology confirmed adenocarcinoma of the pancreas.

The patient went on to complete three courses of gemcitabine chemotherapy but further staging CT demonstrated slowly progressive disease over a 3-mo period.

Six months later, the patient was referred by his GP to the on-call medical team with a week-long history of nausea, anorexia and new-onset jaundice. Upon examination, he was tender in the epigastrium and right upper quadrant, with no overt peritonism. His blood tests were as follows: bilirubin, 164 μmol/L; alanine aminotransferase, 84 IU/L; alkaline phosphatase, 5567 IU/L; albumin, 35 g/L; C-reactive protein (CRP), > 160 mg/L; prothrombin time, 16.3 s; and white blood cell count, 12.9 × 10^9/L.

Repeat CT of the pancreas was undertaken, which demonstrated a dilated common bile duct (CBD) with external compression from enlargement of the pancreatic tumor, and a large, intrahepatic cystic lesion that measured 12 cm × 9 cm, adjacent to the gallbladder (Figure 2).

Ultrasound-guided drainage was performed and yielded frank bile in keeping with an intrahepatic biloma. Microbiological analysis of the fluid did not reveal the presence of any pathogenic organisms. Endoscopic retrograde cholangiopancreatography (ERCP) demonstrated a smooth stricture of the distal CBD and a dilated common hepatic duct above. Contrast material was seen to flow within the biloma collection, which implied communication with the intrahepatic ducts (Figure 3). The stricture was stented and drainage from the biloma ceased within 48 h following insertion.

**DISCUSSION**

The first reported case of a biloma was reported by Gould and Patel in 1979[1]. They reported extrahepatic bile leakage post trauma to the right upper quadrant of the abdomen. The bile accumulated in an encapsulated form. Although originally described as a biliary collection outside the liver, the term biloma has been extended to include any such lesion that may be intrahepatic but anatomically outside the biliary tree[2]. The majority of bilomas are iatrogenic and follow transhepatic cholangiography, liver biopsy, ERCP or cholecystectomy. Biloma also has been recognized to arise following external trauma[3]. Spontaneous biloma is exceedingly rare, and the majority occur secondary to choledocholithiasis or cholangiocarcinoma[4]. Rarer causes have been reported in the context of sickle cell disease[5] or as a complication of hepatic infarction and abscess formation. To the best of our knowledge, biloma that occurs secondary to primary pancreatic malignancy has not been reported previously in the literature.

As diagnostic techniques continue to evolve, an increasing number of cases have been identified, but the exact mechanism behind spontaneous biloma formation is still unknown. Postulated pathogenic mechanisms are Sphincter of Oddi spasm, CBD tumor or calculus obstruction that results in increased intraductal pressure, bile duct necrosis and rupture of the bile duct. As a result of the relatively slow onset of ductal obstruction that occurs in the context of a pancreatic neoplasm, such an acute elevation in biliary pressure is unusual (cf. impacted CBD stone). The size and location of biloma is influenced by the cause of rupture, location and size of bile leak, and rate of absorption by the peritoneum.
Most bilomas are secondary to CBD rather than hepatic duct perforation\textsuperscript{6}. There is no difference in the incidence between males and females, but the condition is found more often in the sixth to seventh decades of life. The age predominance may reflect that of the underlying etiological factor rather than that of developing the complication. Presentation is nonspecific, with abdominal pain, usually in the right upper quadrant (although a few reported cases of bile migration to the left subphrenic space have been documented, which has given rise to a predominance of pain on the left side). Fever may be accompanied by jaundice and abdominal distension. Extreme cases that result in bilious ascites also have been reported\textsuperscript{7}. In our case, there was no history of recent hepatobiliary intervention (pancreatic biopsy had been performed 6 mo prior to presentation, but no evidence of biloma was seen on interval scanning). There were complaints of fever, rigors, anorexia and scleral icterus. Blood tests may show neutrophil leukocytosis, elevated CRP and obstructive liver function tests. Blood cultures may show Gram-negative bacteremia. Biloma can be picked up on ultrasound, CT or magnetic resonance imaging. Despite advancing imaging modalities, biloma may be difficult to differentiate from large cystic metastasis, seroma, angioma or lymphocele. Ultrasound becomes useful in this situation, with a definitive diagnosis being made following radiologically guided aspiration. Once fluid is obtained, microbiological testing is mandatory to exclude the presence of coexisting infection. ERCP is particularly useful in diagnosing an active leak; this may also allow therapeutic intervention\textsuperscript{8}. Precise location of the biloma allows for percutaneous drainage, which negates the need for surgical intervention. Endoscopic intervention includes sphincterotomy with stone extraction, if appropriate, to lower biliary pressure. Placement of a stent in more distal lesions is an option, as this reduces the pressure gradient into the duodenum and facilitates forward flow of bile. This also relieves obstruction from lesions that narrow the biliary tree. In our case, the latter approach was used to overcome the obstruction caused by the large pancreatic tumor.

Surgical management remains contentious but can be useful in cases of ongoing leakage despite endoscopic therapy. The goals are to halt abdominal contamination from bile by means of peritoneal drainage, surgical closure of active leaks, and T-tube drainage\textsuperscript{6,9}.

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