A healthy 68-year-old white woman of eastern European origin who had lived her entire life in Canada presented with ascending cholangitis. She had undergone cholecystectomy many years earlier and had suffered recurrent bouts of abdominal pain over the last few years, for which multiple investigations demonstrated only common bile duct (CBD) dilatation without a clear etiology. CT of the abdomen (Fig. 1) now showed dilated left-sided central intrahepatic bile duct, and endoscopic retrograde cholangiopancreatography (ERCP) confirmed the presence of multiple small stones in those segments. A diagnosis of recurrent pyogenic cholangitis (RPC) was made. Because the large number of stones could not all be retrieved by ERCP and because of the recurrent nature of her disease, surgical management was indicated.

The patient underwent a left hepatectomy. Multiple, small, friable, pigmented stones were found in the left intrahepatic bile duct; intraoperative cholangioscopy confirmed the absence of any disease in the right biliary tree. A Roux-en-Y hepaticojejunostomy was performed with pexy of the blind-ended limb to the anterior abdominal wall for future percutaneous cannulation of the biliary tree in the event of recurrent stones or strictures (a modification of the technique described by Hutson and colleagues1 for the treatment of recurrent biliary strictures in sclerosing cholangitis). The excised specimen contained multiple intrahepatic pigmented stones. The biliary mucosa showed marked chronic inflammatory and fibrotic changes with a minimally inflamed hepatic parenchyma (Fig. 2).

Discussion

Recurrent pyogenic cholangitis, also known as Hong Kong disease and oriental cholangiohepatitis, is characterized by recurrent bouts of abdominal pain, jaundice and fever secondary to intrahepatic ductal dilatation and strictures, and subsequent intrahepatic stone formation. Most prevalent in the East, it is seen increasing in the West mainly owing to immigration.2,3 This is the ninth case of RPC in a non-Asian and only the third documented case in North America. Western doctors should be familiar with this disease as it differs significantly both in diagnosis and...
management as compared with biliary disease in that region of the world. If left untreated, RPC is associated with a high morbidity and mortality as it progresses to biliary cirrhosis and liver failure.

Recurrent pyogenic cholangitis is thought to occur in patients suffering from chronic infestation of the biliary tree by *Clonorchis sinensis* and *Ascaris lumbricoides*, liver flukes endemic to Southeast Asia that may obstruct the intrahepatic biliary ducts with resultant bile stasis, pigment stone formation and bacterial superinfection. The *Clonorchis* parasite has a lifespan of 20 years and can therefore manifest itself many years after initial infestation. Patients with RPC most often present with symptoms of cholangitis, and bile cultures are almost always positive and usually yield *Escherichia coli*, *Enterococcus fecalis*, or other enteric bacteria, but *Clonorchis* or *Ascaris* parasites or ova are recovered from the stool in only 25% of cases. Biliary disease involves the CBD and left hepatic lobe in 37% of cases and includes the right hepatic lobe in 31%. It is confined to the CBD in 27%, and only rarely confined to the right lobe. The gallbladder is usually disease-free.

Acutely, RPC is treated with broad-spectrum antibiotics and ERCP decompression. CBD exploration and T-tube drainage are reserved for failure of medical management. Further attacks may be prevented by elective CBD exploration with cholecystoscopy for stone retrieval, irrigation of the biliary tree and treatment of strictures. Biliary-enteric bypass is required for distal extrahepatic strictures; however, this is associated with cholangitis and intrahepatic abscess in up to 30% of patients. Hepatic resection should be reserved for patients with significant hepatic atrophy and fibrosis, multiple liver abscesses or concurrent intrahepatic cholangiocarcinoma, which may occur in 3%–5% of patients treated conservatively.

**Conclusions**

Recurrent pyogenic cholangitis should be suspected in anyone with a history of recurrent right upper quadrant pain despite past cholecystectomy. Its diagnosis and treatment require the cooperation of surgeon, endoscopist, and diagnostic and interventional radiologists. Surgical interventions still carry a high morbidity and must be tailored to the individual patient, depending on the severity of disease.

**Competing interests:** None declared.

**References**