Pericardiobiliary Fistulation: A Rare Complication of Therapeutic ERCP in a Patient With IgG4-Related Sclerosing Cholangitis

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Abstract
A 70-year-old man presented with acute coronary syndrome 3 weeks after plastic stent insertion for hilar biliary stricturing secondary to IgG4-related sclerosing cholangitis (IgG4-SC). Imaging demonstrated haemopericardium due to proximal migration of the plastic biliary stent through the liver capsule and diaphragm into the pericardial sac. The stent was endoscopically removed and a pericardiocentesis was performed. The patient’s clinical condition rapidly improved. We illustrate an unusual but potentially serious complication that may arise from migration of a biliary stent and discuss a management strategy.

Introduction
IgG4-related disease is a steroid-responsive, multi-system inflammatory disorder. The most frequent extra-pancreatic manifestation of IgG4-related disease is IgG4-related sclerosing cholangitis (IgG4-SC), present in over 70% of patients with IgG4-related autoimmune pancreatitis (AIP). Endoscopic retrograde cholangiopancreatography (ERCP) is often required in such cases to obtain a tissue diagnosis and to treat strictures requiring a biliary stent.

Case Report
A 70-year-old Chinese man with a past medical history of hepatitis B virus infection (HBsAg negative) and glaucoma presented with pruritis and painless obstructive jaundice (bilirubin 300 µmol/L). Cross-sectional imaging revealed a 3-cm mass in the head of the pancreas abutting the superior mesenteric vein, portal lymphadenopathy, and a hilar liver mass with associated intrahepatic duct dilatation. ERCP revealed a long biliary stricture extending to the hilum, which was treated with a straight plastic biliary stent. Histological sampling from the ampulla, common bile duct (CBD), and the liver lesion all showed a lymphoplasmacytic infiltrate with a high expression of IgG4-positive plasma cells (>100 cells per high-powered field), consistent with a diagnosis of IgG4-related disease. Serum IgG4 was elevated at 4.36 g/L (normal: 0-1.3 g/L).

He started a tapering course of oral steroids and had an excellent biochemical and clinical response. His biliary stent was removed. After 6 months, steroid treatment was stopped. Three months later, he presented with jaundice and chills. Repeat ERCP demonstrated 2 short, dominant biliary strictures in the distal CBD and at the liver hilum (Figure 1), and a long, irregular, thin pancreatic duct (Figure 1). A 15-cm, 10-French, straight plastic biliary stent was placed across the liver hilum into the left main intrahepatic duct (Figure 1). The length of this stent was necessary to traverse both strictures and provide adequate biliary drainage. Steroids were restarted with intent of starting second-line long-term maintenance immunosuppression.
Twenty days following biliary stent insertion, he developed acute chest pain. An electrocardiogram showed widespread ST segment elevation; coronary angiography was normal. The following day, he developed a fever with dyspnea and supraventricular tachycardia. Chest radiography showed left lower lobe consolidation with air bronchograms and blunting of the left costophrenic angle. A biliary stent was noted on the inferior aspect of the film; its superior end was overlying the cardiac silhouette (Figure 2). He was treated with antibiotics for suspected pneumonia. A CT scan confirmed a large pericardial effusion, with the proximal tip of the plastic stent extending through the liver capsule and diaphragm into the pericardial sac (Figure 3). ERCP showed that the distal end of the stent was within the distal bile duct. The stent was mobilized with traction using a biliary extraction balloon and removed with stent grabbers. He remained hemodynamically stable during the procedure, but the pericardial effusion enlarged and was successfully treated with percutaneous pericardial drainage of approximately 300 mL of blood.

He made a swift recovery and his IgG4-related disease has since been remission with azathioprine and low dose prednisolone. Eighteen months later, no further biliary intervention has been required and he remains clinically well with normalization of his liver function tests and radiological improvement in biliary stricturing.

**Discussion**

IgG4-related disease is a multi-system inflammatory disorder and an emerging disease entity of unknown etiology. It comprises a collection of conditions that share clinical, serological, histopathological, and radiological features. It is characterized by a lymphoplasmacytic tissue infiltration with a predominance of IgG4-positive plasma cells and T-lymphocytes. Multiple organs may be involved, including the pancreas and biliary tree. Although the optimal treatment for IgG4-related disease has not been established, clinical and radiological response to steroids usually occurs within weeks of starting therapy.

Our patient had an established diagnosis of IgG4-related disease with biliary and pancreatic involvement. We hypothesize that rapid improvement in biliary stricturing in response to steroids reduced the traction force on the biliary stent, allowing it to migrate proximally, and that the 15-cm length of the stent may have been an additional factor influencing migration. An argument could be made for steroid treatment without biliary stenting, but there is not sufficient evidence at this stage to support that approach.

Proximal or distal migration of plastic biliary stents is a recognized complication of ERCP in 5-7% of patients. Migration
frequency is higher in patients with benign disease. Proxi-
mally migrated biliary stents can be removed successfully
using an endoscopic retrieval technique, the choice of which
depends on several factors including biliary duct dilation,
depth of stent migration, distal stent impaction, and biliary
stricture distal to the migrated stent. Common retrieval tech-
niques include biliary baskets, extraction balloons, snares,
forceps, and cholangioscopic stent retrieval.

Traumatic bronchopleurobiliary, hepaticogastric, and other
extrahepatic bile duct fistulae following migration of a plastic
biliary stent through the liver capsule have previously been
described, but we could find no previous case reports of in-
trapercardial migration. Our case illustrates the need for
clinical awareness and early recognition of this condition if
a patient presents with otherwise unexplained cardiorespira-
tory symptoms following biliary stent insertion.

Disclosures
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and is the article guarantor. D. Joshi contributed to the dis-
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images, and reviewed and edited the final manuscript.

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