A true vascular aneurysm of the hepatic artery proper as a rare cause of nonmalignant painless jaundice

Rare causes of painless jaundice include parasitic infections and lymphoma. To date, two cases of vascular pseudoaneurysm in acute cholecystitis and chronic pancreatitis have been reported [1, 2].

An 85-year-old man was diagnosed by contrast-enhanced computed tomography scan with a partially thrombotic aneurysm of the hepatic artery proper, which was compressing the biliary duct and left-sided segment branches (yellow arrow) (Fig. 1). An initial attempt to place an endoprosthesis via endoscopic retrograde cholangiopancreatography failed, and obstructive cholangitis developed (bilirubin 23.1 mg/dL, C-reactive protein 116 mg/L, leukocytosis 17800/µL), which required antibiotic treatment, resection, and/or a second problem-focused ERCP. Resection was discussed but was not considered to be feasible due to significant cardiovascular co-morbidity. Therefore, biliary tract decompression by ERCP was planned.

ERCP was particularly challenging. At a distance of 35 mm from the papilla, below the junction of the cystic duct, the vascular aneurysm caused a moderately severe smooth-walled stenosis (50%–90%), measuring at least 45 mm in length. The external compression resulted in a curved CBD with a right-angled kink (Fig. 2 and Fig. 3). After endoscopic papillotomy, widening of the stenosis was achieved by careful use of bougies (5–10 Fr). Subsequently, one double-pigtail endoprosthesis was placed in the right hepatic duct (7 Fr/16 cm) to serve as a splint for the second endoprosthesis, which had to be implanted around and over the aneurysm to finally reach the dilated biliary ducts of the left liver segments (10 Fr/12 cm; Fig. 4 and Fig. 5). Correct stent placement was confirmed by postinterventional ultrasound (Fig. 6).

Interventional occlusion of the aneurysm was not performed due to the risk of wide-ranging ischemia. Thus, only mechanical biliary drainage evidenced by decreasing cholestasis was able to circumvent the complications of this rare vascular cause of bile duct compression. In contrast to arterial pseudoaneurysms, which are a rare but established complication of ERCP [3, 4], this is, to our knowledge, the first case of a true vascular aneurysm leading to progressive cholangitis that required treatment by ERCP.

Fig. 1 The partially thrombotic aneurysm of the hepatic artery proper (yellow arrows) extended from the junction of the gastroduodenal artery up to the branching of the right and left hepatic artery (diameter 4.3 cm).

Fig. 2 At endoscopic retrograde cholangiopancreatography, the distal bile duct showed normal width, filling, and bile duct wall. However, approximately 35 mm above the papilla, the bile duct was bent at right angles and showed a 45-mm long stenosis in the middle and upper parts, extending upwards to the biliary hilus (white arrows). Of note, the descending biliary branch of the liver segment III also showed a termination of the duct (yellow arrow).

Fig. 3 At endoscopic retrograde cholangiopancreatography, the area without contrast media filling was measured as 39 × 47 mm, corresponding to the aneurysm of the hepatic artery proper, which compressed the biliary duct and left-sided segment branches (yellow arrow).

Fig. 4 An initial 7-Fr double-pigtail endoprosthesis (yellow arrows) was inserted into the biliary hilus and the right segment VII to serve as guide for the second endoprosthesis, which had to be inserted carefully around and over the aneurysm into the left-sided dilated segments.
Competing interests: None

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