Cholesterolosis of the gallbladder visualized by peroral cholecystoscopy using a SpyGlass probe

It is difficult to distinguish malignant gallbladder diseases from benign ones preoperatively, even though appropriate imaging techniques have been developed [1]. Therefore complementary techniques which facilitate direct visual assessment and visually guided tissue sampling are desirable. We describe the first case of gallbladder cholesterolosis successfully visualized and diagnosed by peroral cholecystoscopy using a SpyGlass probe (Boston Scientific, Natick, Massachusetts, USA) in a patient with pancreatobiliary maljunction.

A previously healthy 42-year-old woman presented at our institution with intermittent right upper abdominal pain. Abdominal ultrasonography and computed tomography showed a dilated cystic duct and polypoid lesions with circumferential wall thickness in the gallbladder. Endoscopic retrograde cholangiopancreatography (ERCP) revealed pancreatobiliary maljunction with congenital choledochal cyst. (CHD, common hepatic duct; CD, cystic duct; GB, gallbladder; CBD, common bile duct; PD, pancreatic duct.)

After a Tandem XL cannula (Boston Scientific) was advanced into the gallbladder, a SpyGlass probe was inserted through the catheter. Peroral cholecystoscopy showed numerous yellowish polypoid lesions resembling a strawberry in the gallbladder (Fig. 2 and Video 1). Transpapillary biopsy specimens of the gallbladder showed cholesterolosis, which was characterized by clusters of foamy macrophages in the lamina propria (Fig. 3).

It is difficult to distinguish malignant gallbladder diseases from benign ones preoperatively, even though appropriate imaging techniques have been developed [1]. Therefore complementary techniques which facilitate direct visual assessment and visually guided tissue sampling are desirable. We describe the first case of gallbladder cholesterolosis successfully visualized and diagnosed by peroral cholecystoscopy using a SpyGlass probe (Boston Scientific, Natick, Massachusetts, USA) in a patient with pancreatobiliary maljunction. A previously healthy 42-year-old woman presented at our institution with intermittent right upper abdominal pain. Abdominal ultrasonography and computed tomography showed a dilated cystic duct and polypoid lesions with circumferential wall thickness in the gallbladder. Endoscopic retrograde cholangiopancreatography (ERCP) revealed pancreatobiliary maljunction with cystic duct dilatation (Fig. 1). After a Tandem XL cannula (Boston Scientific) was advanced into the gallbladder, a SpyGlass probe was inserted through the catheter. Peroral cholecystoscopy showed numerous yellowish polypoid lesions resembling a strawberry in the gallbladder (Fig. 2 and Video 1). Transpapillary biopsy specimens of the gallbladder showed cholesterolosis, which was characterized by clusters of foamy macrophages in the lamina propria (Fig. 3).
The patient underwent resection of the extrahepatic bile duct and gallbladder with hepaticojejunostomy. A surgical specimen revealed that cholesterolosis had spread extensively to the gallbladder and bile duct, without there being a malignant lesion (Fig. 4).

There have been previous reports of peroral cholecystoscopy; however, the technique has not been widely accepted, because of technical difficulties [2]. The SpyGlass Direct Visualization System (Boston Scientific), which is a newly developed peroral cholangiopancreatoscopy system, provides improvements in the diagnosis and therapy of various pancreatobiliary diseases [3]. The SpyGlass probe can be used through a conventional ERCP catheter, so peroral cholecystoscopy can be performed easily even when the diameter of the bile duct or cystic duct is too small for conventional cholangioscopy. Thus this technique may expand the diagnostic possibilities in diseases of the gallbladder.

Endoscopy_UCTN_Code_CCL_1AZ_2AI

Competing interests: None

References


Bibliography

DOI http://dx.doi.org/10.1055/s-0031-1291495
Endoscopy 2012; 44: E145–E146
© Georg Thieme Verlag KG Stuttgart · New York
ISSN 0013-726X

Corresponding author

H. Isayama, MD, PhD
Department of Gastroenterology
Graduate School of Medicine
The University of Tokyo
7-3-1 Hongo
Bunkyo-ku
Tokyo 113-8655
Japan
Fax: +81-3-38140021
isayama-2im@h.u-tokyo.ac.jp