T1b primary remnant cystic duct cancer following cholecystectomy: A case report

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INTRODUCTION: Although primary cystic duct cancer is a rare entity, remnant cystic duct cancer is even more rare. We report a case of early cystic duct cancer following cholecystectomy.

PRESENTATION OF THE CASE: A 81 year-old man complained temporary loss of appetite. He had underwent cholecystectomy for acute cholecystitis 5 years prior. Contrast enhanced computed tomography, magnetic resonance image and endoscopic ultrasonography showed remnant cystic duct tumor with protrusion to common bile duct. Endoscopic retrograde cholangiography revealed defect of contrast medium around confluence of the remnant cystic duct and common bile duct. We performed step biopsy by using forceps which revealed adenocarcinoma. Based on these findings, extraperitoneal bile duct and remnant cystic duct resection were performed. The histopathology showed adenocarcinoma, pap > tub2, filling in remnant cystic duct, 30 mm in size but showed no lymphovascular or perineural invasion, no lymph node metastasis and negative surgical margin, and was classified as pT1bN0M0.

CONCLUSION: This is a rare case of primary carcinoma of remnant cystic duct cancer which is detected during computed tomography follow up for hepatic cell carcinoma recurrence. We confirmed remnant cystic duct cancer and its superficial extension to common bile duct with endoscopic ultrasonography and intraductal ultrasonography. Proper curative surgery was performed.

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and no malignant lesion from the hepatic hilar and the intrapancreatic bile duct. We considered to perform pancreateoduodenectomy. However, we performed distal bile duct resection in consideration for the patient’s age and comorbidities.

The histopathology showed adenocarcinoma, pap > tub2, filling in RCD with protrusion to CBD, 30 mm in size, that had invaded to the fibromuscular layer, but showed no lymph node metastasis, and negative surgical margin, and was classified as pT1bN0M0 according to the American Joint Committee on Cancer and Union for International Cancer Control manual, 8th edition [1]. (Figs. 4 and 5) He recovered without any complication. There was no recurrence 8 months after operation.

3. Discussion

We herein describe a case of T1b RCD cancer following cholecystectomy. On 1 year prior CT image, we can detect a nodule in CBD retrospectively, but its findings was overlooked because the main purpose was detection of HCC recurrence and there was no bile duct dilation. His symptoms such as loss of appetite and liver dysfunction led to diagnosis of RCD cancer. Cystic duct cancer is a rare entity. Tu-Nan Yu et al. reviewed extrahepatic cholangiocarcinoma diagnosed between 2006 and 2015, cystic duct cancer was only accounted 0.44% [2]. In 1951, Farrar proposed diagnostic criteria for cystic duct carcinoma. Cystic duce cancer was defined based on the following three criteria: (a) The growth must be restricted to the cystic duct; (b) There must be no neoplastic process in the gallbladder, hepatic or common bile ducts; and (c) A histological examination of the growth must confirm the presence of carcinoma cells” [3]. But Farrar’s criteria is only applicable to limited carcinoma and excludes advanced one originating from the cystic duct. In most cases, the tumor extends to gallbladder and CBD and the origin is difficult to determine. Ozden et al. proposed new working definition which describes a cystic duct adenocarcinoma as a gallbladder cancer whose center is located in the cystic duct [4].

Cystic duct cancer is rare but might be suspected in patients presenting with distended gallbladder or cholecystitis. However, RCD cancer doesn’t cause symptoms related to gallbladder swelling. If it advanced to CBD, obstructive jaundice will occur. There are limited reports about RCD cancer following cholecystectomy. To the best of our knowledge, only 9 cases of subsequent

Fig. 1. a the enhanced nodule was already detected on 1 year prior image in the common biliary duct (CBD). b the nodule expanded slightly when this patient complained sympoms. c a multi planar reconstruction image shows the enhanced lesion in the remnant cystic duct(RCD)(arrow) and protrusion to CBD.

Fig. 2. Endoscopic ultrasound sonography: Low echoic mass lesion in remnant cystic duct(arrow) protrudes to common hepatic duct(arrow head). Another nodule on opposite side wall of common hepatic duct as well.
Fig. 3. Endoscopic nasobiliary cholangiography depicts filling defect around the confluence of cystic duct and common hepatic duct, but cystic duct wasn’t shown. Intraductal ultrasound image: 1,3 no significant bile duct wall thickness. 2 low echoic mass protrudes to common hepatic duct from remnant cystic duct.

Fig. 4. Resected specimen (arrow head is duodenal surgical margin of side common hepatic duct). a nodules on common hepatic duct wall and papillary expanding tumor in remnant cystic duct(arrow). b solid lines: tumor spreading along to common hepatic duct wall. spotted line: tumor in remnant cystic duct.
remnant cystic duct cancer following cholecystectomy including one case of T1b cancer have been reported (Table 1; found by searching the PubMed databases by using the keywords “remnant cystic duct cancer” or “cystic duct cancer after cholecystectomy” all articles were written in English) [5–12].

All their cause of prior cholecystectomy were cholelithiasis or cholecystitis. Cases since 2003 were reported from Korea or Japan. This would be affected by worldwide prevalence of biliary tract cancer [13]. Although histological type of the RCD cancer case series were various, early stage cancer cases were tub1 or pap [12]. Our case is not only resected in early stage, but also 1 year has passed since it has detected firstly with CECT. This suggests that the tumor progressed slowly. The Japanese Society of Hepato-Biliary-Pancreatic Surgery proposed that extrahepatic cholangiocarcinoma could be classified as one of three types based on gross morphology: papillary type, nodular type or flat type types [14]. Papillary type is reported to significant tendency of noninvasiveness to deeper layers and better prognosis. Frequent occurrence of papillary type with early extrahepatic bile duct cancer may be explained by the fact that these tumor types grow and spread superficially along the bila duct mucosa and do not invade deeply into the bile duct wall [15–18].

To confirm RCD cancer is challenging if there isn’t jaundice or bile duct dilatation. Although this kind of cancer is extremely, careful radiographic image interpretation, EUS and IDUS may be helpful to diagnose early stage RCD cancer. In our case, RCD cancer was spreading to CBD. This enabled us to diagnose as malignant by ERCP preoperatively.

4. Conclusion

This is a rare case of primary carcinoma of RCD which is detected during CT follow up for HCC recurrence. We confirmed RCD cancer and its superficial extension to CBD with EUS and IDUS. Proper curative surgery was performed.

Declaration of Competing Interest

The authors declare that they have no conflict of interest.

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Ethical approval

This study is exempt from ethical approval in Kishiwada city hospital ethics review committee.

Consent

Informed consent was obtained from this patient to be included in the study.

Author contribution

All authors in this manuscript contributed to the interpretation of data, and drafting and writing of this manuscript. Muneji Yasuda is first author and corresponding author of this paper. Yuichi Tanaka, Shinji Miyajima, Toyokazu Fukunaga and Haruo Takaya were engaged in patient’s care in our hospital. Kozo Kajimura reviewed the final manuscript. All the authors read and approved the final manuscript.
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