A case of right hepatic duct entering cystic duct successfully treated by laparoscopic subtotal cholecystectomy through preoperatively placed biliary stent

Hiroki Hirao1,2*, HiroHisa Okabe1, Daisuke Ogawa1, Daisuke Kuroda1, Katsunobu Taki1, Shinjiro Tomiyasu1, Masahiko Hirota1, Taizo Hibi2, Hideo Baba3 and Hiroki Sugita1

Abstract

Background: Laparoscopic cholecystectomy is a well-established surgical procedure and is one of the most commonly performed gastroenterological surgeries. Therefore, strategy for the management of rare anomalous cystic ducts should be determined.

Case presentation: A 56-year-old woman was admitted to our hospital owing to upper abdominal pain and diagnosed with acute cholecystitis. Magnetic resonance cholangiopancreatography suspected that several small stones in gallbladder and the right hepatic duct drained into the cystic duct. Endoscopic retrograde cholangiopancreatography confirmed the cystic duct anomaly, and an endoscopic nasobiliary drainage catheter (ENBD) was placed at the right hepatic duct preoperatively. Intraoperative cholangiography with ENBD confirmed the place of division in the gallbladder, and laparoscopic subtotal cholecystectomy was safely performed.

Conclusions: The present case exhibited rare right hepatic duct anomaly draining into the cystic duct, which might have caused biliary tract disorientation and bile duct injury (BDI) intraoperatively. Any surgical technique without awareness of this anomaly preoperatively might insufficiently prevent BDI, and preoperative ENBD would facilitate safe and successful surgery.

Keywords: Cystic duct anomaly, Right hepatic duct, Laparoscopic cholecystectomy, Endoscopic nasobiliary drainage catheter

Background

Variations of the bile duct before hepatobiliary surgery should be recognized before the hepatobiliary surgery to prevent postoperative complications. A rare cystic duct anomaly increases the risk of bile duct injury (BDI) [1]. However, congenital anomaly of the right hepatic duct (RHD) entering the cystic duct is very rare. Only 10 patients have undergone laparoscopic cholecystectomy on variant cystic ducts till date [2]. Here, we report a case of RHD entering cystic duct successfully treated with laparoscopic cholecystectomy using a preoperatively placed endoscopic retrograde biliary drainage tube inside the RHD.

Case presentation

A 56-year-old woman was admitted to our hospital owing to upper abdominal pain. She did not have fever. Her Murphy’s sign was negative. Laboratory data were as follows: white blood cell count, 72.0 × 10³/μL, and C-reactive protein, 0.23 mg/dL. Plain abdominal computed tomography (CT) revealed dilation of both the RHD and
gall bladder (Fig. 1a). She was admitted to our hospital for observation. On the next day, contrast-enhanced CT and magnetic resonance cholangiopancreatography (MRCP) were performed, with the latter detecting a congenital anomaly of the RHD entering into the cystic duct (Fig. 1b). Although several small stones in gallbladder were found, no cholecystolithiasis was confirmed to cause biliary obstruction. Endoscopic retrograde cholangiopancreatography revealed that the RHD entered the cystic duct. Therefore, an endoscopic nasobiliary drainage (ENBD) tube was placed inside the RHD. Laparoscopic cholecystectomy was planned. The cystic duct was carefully dissected, and RHD bifurcation was confirmed (Fig. 2a). Intraoperative cholangiography was performed using an ENBD catheter, after grabbing the gallbladder neck with forceps at 10 mm away from the confluence. Intraoperative cholangiography revealed that the catheter in the RHD was not grabbed by forceps (Fig. 2b).
The Hartmann’s pouch of the gallbladder was clipped and divided, and the ENBD catheter was removed. Laparoscopic subtotal cholecystectomy was safely performed. The patient was discharged without any complication. At 1 month postoperatively, the patient complained abdominal pain and high fever. Imaging analysis suspected biliary obstruction at the cystic duct due to choledocholithiasis which was also suspected at the first visit preoperatively. A plastic biliary stent was placed, resulting in immediate disappearance of symptom. The stent was removed 3 months postoperatively; however, abdominal pain recurred. Imaging analysis revealed that the RHD was dilated and cystic duct stenosis was suspected. The plastic stent was placed again, and further investigations for malignant diseases were performed. No malignant diseases were found. The patient was observed for 3 months, and the stent was removed again. One month after the last stent removal, MRCP revealed no RHD dilatation (Fig. 3). Sustained swelling of biliary tract after surgery might mainly be the cause of stenosis. The patient has been free from any symptom for a year after MRCP.

Discussion

Laparoscopic cholecystectomy is a well-established surgical procedure, and strategies to prevent BDI have been well-determined [3]. Although the critical view of safety is globally accepted as a standardized concept [4], the current cystic duct anomalies require exceptional strategies to prevent BDIs. In this case, a rare RHD anomaly entering the cystic duct was confirmed preoperatively, and the patient was successfully treated with endoscopic surgery using an ENBD that was preoperatively placed as a biliary guide. CT images during the first presentation showed a dilated RHD and a distended gall bladder. This finding implies the possibility of cystic duct anomalies that combined the right biliary branch draining together into the common hepatic duct (CHD), unless the gallbladder compressed the RHD, which is diagnosed as Mirrizi syndrome.

A well-known anomalous cystic duct requiring attention to prevent BDI during a laparoscopic cholecystectomy is characterized by the combination of the right posterior segmental duct (RPSD) draining into the CHD and cystic duct entering into RPSD. A previous investigation of cholangiograms on donors revealed that the rarity of RPSD draining directly into CHD was 6%, of which the anomalous posterior branch is close to cystic duct based on the anatomical positioning. Conversely, the rarity of RPSD or RHD entering cystic duct, as in the current case, is 2% or < 1%, respectively [5]. The real percentage of the cystic duct anomaly, as in the current case, is unclear due to its high rarity. Ten cases have been reported till date to be similar to the current case showing RHD entering cystic duct [2]. Intriguingly, one out of four cases (25%) showed RHD entering the proximal side of the cystic duct, causing a wide range of safe cut lengths of the cystic duct, implying that critical complications are less likely to occur intraoperatively. In contrast, three out of four cases (75%) showed RHD entering the distal side of the cystic duct, causing a very short range of safe cut length of the cystic duct, implying that unintended division of the bile flow in RHD is likely to occur intraoperatively (Fig. 4). In such cases, cutting cystic duct purposing to insert a catheter for intraoperative cholangiogram may cause serious complications even under laparotomy, because the cystic duct almost functions as the RHD and cannot be cut. Therefore, these cases, including our case, required subtotal cholecystectomy. Among the ten cases of anomalous RHD on the cystic duct, laparoscopic surgery was successfully performed in four [6–8]. The anomaly was confirmed by cholangiography using the percutaneous transhepatic gallbladder drainage catheter in two cases. Details of the intraoperative cholangiography procedure were not available in the other two reports. Based on these observations, intraoperative cholangiography should be carefully performed depending on the confluence position of the RHD on the cystic duct. We successfully performed laparoscopic cholecystectomy using an ENBD catheter placed during ERCP, which was aimed to diagnose an anomalous cystic duct preoperatively. This strategy decreases stress levels and prevents performance of technically difficult procedures intraoperatively [9].
Conclusions

The current case indicates that comprehension of cystic duct anomaly is mandatory; the rare anomalous cystic duct, as in the current case, might cause intraoperative disorientation, leading to BDI. This very rare anomaly that allows the RHD to enter the cystic duct should be carefully considered. Although confirming critical view of safety and performing bailout procedure might be enough for the current case, it is possible that an experienced surgical technique and knowledge of various possible anomalies of cystic duct could have been required, if cholecystitis was more challenging at surgery. If the careful investigation was required for a case suspected to have very rare anomaly of cystic duct, ERCP is performed to confirm the anatomy of biliary tract. It is not difficult to place the ENBD for such case. We suggest that preoperative placement of ENBD is a safe and certain strategy to make sure the biliary system anatomy during laparoscopic cholecystectomy.

Abbreviations
ENBD: Endoscopic nasobiliary drainage catheter; BDI: Bile duct injury; RHD: Right hepatic duct; CT: Computed tomography; MRCP: Magnetic resonance cholangiopancreatography; CHD: Common hepatic duct.

Acknowledgements
None.

Authors’ contributions
HH described and designed the article. HO, ST, and HS edited the article. TH, HB, and HS supervised the edition of the manuscript. Other remaining co-authors collected the data and discussed the content of the manuscript. All authors read and approved the final manuscript.

Funding
This study was not funded.

Availability of data and materials
All data and materials are available.

Ethics approval and consent to participate
This is not applicable for this case report.

Consent for publication
Written informed consent was obtained from the patient for publication of this case report.

Competing interests
The authors declare that there is no conflict of interests.

Author details
1 Department of Surgery, Kumamoto Regional Medical Center, S-16-10 Honjo, Chuo-ku, Kumamoto 860-8811, Japan. 2 Department of Pediatric Surgery and Transplantation, Kumamoto University Graduate School of Medical Sciences, 1-1-1 Honjo, Chuo-ku, Kumamoto 860-8556, Japan. 3 Department of Gastroenterological Surgery, Graduate School of Medical Sciences, Kumamoto University, 1-1-1 Honjo, Kumamoto 860-8556, Japan.

Received: 22 July 2020 Accepted: 18 September 2020
Published online: 25 September 2020

References
1. Suhocki PV, Meyers WC. Injury to aberrant bile ducts during cholecystectomy: a common cause of diagnostic error and treatment delay. Am J Roentgenol. 1999;172:955–9.
2. Abe T, Ito S, Kaneda Y, Suto R, Noshima S. Main right hepatic duct entering the cystic duct: a case report. Surg Case Rep. 2019;5:46.
3. Altieri MS, Brunt LM. Elimination of bile duct injury in cholecystectomy. Adv Surg. 2019;53:145–60.
4. Strasberg SM, Hertl M, Soper NJ. An analysis of the problem of biliary injury during laparoscopic cholecystectomy. J Am Coll Surg. 1995;180:101–25.
5. Choi JW, Kim TK, Kim KW, Kim AY, Kim PN, Ha HK, et al. Anatomic variation in intrahepatic bile ducts: an analysis of intraoperative cholangiograms in 300 consecutive donors for living donor liver transplantation. Korean J Radiol. 2003;4:85–90.
6. Hashimoto M, Hashimoto M, Ishikawa T, Iizuka T, Matsuoka M, Watanabe G. Right hepatic duct emptying into the cystic duct: report of a case. Surg Endosc. 2002;16:359.
7. Matsumoto N, Hasegawa H, Shiroko T, Sakamoto E, Komatsu S, Kurumiya Y, et al. A case of laparoscopic cholecystectomy for acute cholecystitis associated with the left-sided gallbladder and an aberrant hepatic duct. Jpn J Gastroenterol Surg. 2008;41:1704–9.
8. Nomura T, Shirai T, Sasagawa M, Wakai T, Hatakeyama K. Anomalous insertion of the right hepatic duct into the cystic duct: report of a case diagnosed before laparoscopic cholecystectomy. Surg Laparosc Endosc Percutan Tech. 1999;9:211–2.
9. Kayahara M, Oyama K, Kitagawa H, Ohta T, Miwa K. Right hepatic duct opening into the cystic duct: the role of pre- and intraoperative cholangiography. Hepatogastroenterology. 2005;52:719–21.