Laparoscopic Management of Gallbladder Duplication: A Case Report

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Abstract

Laparoscopic cholecystectomy has become one of the most commonly performed abdominal surgeries worldwide. Several anatomic variations and congenital malformations in the gallbladder and biliary anatomy have been described. We reported the case of a middle-aged woman who presented with jaundice and abdominal pain. Her laboratory investigations revealed an elevated bilirubin level (mainly the direct component). The patient showed an improvement in the clinical and laboratory parameters after conservative management. Then, the patient was prepared for a laparoscopic cholecystectomy which revealed a duplication of the gallbladder with two distinct cystic ducts draining independently to the common bile duct. The procedure was completed uneventfully. This type of gallbladder duplication is among the least common types. The case highlights the importance of having a meticulous intraoperative evaluation of the biliary anatomy to avoid potential complications and injuries.

Introduction

The gallbladder is a pear-shaped hollow viscus that serves to store the bile produced by the liver and concentrates it to be released after meals [1]. It is formed from the caudal part of the foregut at six weeks of gestation [2]. Anatomic variations and congenital malformations of the gallbladder are well-recognized. However, they are rare and could be part of some syndromes [3]. The awareness of these variations is of paramount importance because of their direct clinical and surgical implications.

Gallstone disease is very common worldwide and it is considered the most common indication for abdominal surgical procedures in the United States [4]. Gallstones develop when there is an imbalance in the composition of the bile resulting in crystallized deposits of bilirubin and cholesterol [5]. Herein, we describe a case of gallbladder duplication that was incidentally identified during laparoscopic surgery for acute cholecystitis.

Case Presentation

A 54-year-old woman presented to the emergency department with a four-day history of right upper quadrant abdominal pain. She described the pain as constant and sharp in nature. The pain radiated to the right shoulder and was associated with nausea and two episodes of vomiting. She reported having previous similar episodes of this pain over the previous year that occur after fatty meals but the pain was milder and had spontaneous resolution. The patient noticed slight yellowish discoloration of her skin and eyes. The physical examination revealed no abnormalities. An ultrasonography of the abdomen was further performed to confirm the diagnosis of obstructive jaundice. A dilated common bile duct (8 mm) with the presence of a small stone was observed. The patient was started on conservative treatment and intravenous ceftriaxone 1 g daily. She demonstrated clinical and laboratory improvement within 48 hours. The patient was prepared for a laparoscopic cholecystectomy. The surgery was performed under...
general anesthesia and four ports were inserted to carry out the procedure. A diagnostic exploration was performed after establishing the pneumoperitoneum and introducing the trocars. It revealed the presence of a gallbladder duplication (Figure 1).

![Laparoscopic View](image1)

**FIGURE 1: Laparoscopic View**

Laparoscopic view demonstrating the presence of two gallbladders (arrows).

Meticulous dissection was performed. The cystic artery and both cystic ducts were ligated. Both gallbladders and their cystic ducts were resected (Figure 2).
FIGURE 2: Resected Specimen

A gross image for the resected specimen of gallbladders showing two distinct cystic ducts (arrows).

Intra-operative cholangiogram showed no evidence of choledocholithiasis. The total operation time was 90 minutes and the estimated blood loss was 50 ml. The patient tolerated the procedure well and was discharged on the fourth postoperative day. She remained symptom-free at one-month follow-up.

Discussion

We described the case of laparoscopic cholecystectomy in a patient with a gallbladder duplication. Gallbladder duplication is an unusual congenital anomaly with an estimated incidence of 1 per 5000 individuals [6]. Several classification systems have been proposed for a gallbladder supplication [7]. In Boyden classification, there are two main types which are bilobed gallbladder (vesica fellea divisa) and true duplication (vesica fella duplex) which is further classified into Y-shaped that has two cystic ducts uniting before entering the common bile duct while the H-shaped type has two cystic ducts entering the common bile ducts independently, as in our case, and is considered the most common type [8]. Harlaftis classification is a newer classification for gallbladder duplication that is based on embryological development and has two types [9]. In type 1 (split primordial group), there is a single cystic duct and this type is further subdivided into septated, V-shaped, and Y-shaped. In type 2, there are two distinct cystic ducts.

There are no unique presenting clinical features of gallbladder duplication. Each gallbladder may be involved in the pathologies that can affect the single gallbladder such as acute cholecystitis. Gallbladder duplication is usually discovered intraoperatively, as in our case, and could lead to surgical difficulties, including conversion to laparotomy and the risk of biliary injuries [10]. It is essential to recognize this condition intraoperatively since missing this condition could require the patient to have repeated surgery for the other gallbladder. It should be noted that it is recommended to resect both gallbladders even if the pathology is in one of them due to the risk of recurrence so that the patient does not need to undergo another operation for the second gallbladder [11].

Our patient responded with the conservative treatment and did not need to undergo further investigations. In cases of obstructive jaundice with failure to respond with conservative treatment, advanced imaging technique, including endoscopic retrograde cholangiopancreatography, magnetic resonance cholangiopancreatography, could be used to delineate the biliary anatomy and diagnose the duplication of gallbladder preoperatively. While the patient had undergone ultrasonography on presentation in the present case, this modality is largely user-dependent and gallbladder variations may not be recognized. The differential diagnosis of gallbladder duplication on ultrasound includes choledochal cyst, gallbladder fold, gallbladder diverticulum, and intraperitoneal fibrous bands [12].

Conclusions

Gallbladder duplication is a rare congenital anomaly. Careful evaluation of the vascular and biliary anatomy is crucial to avoid intraoperative complications, including common bile duct injuries. Laparoscopic surgery is a feasible option in the management of a gallbladder duplication with gallbladder disease.

Additional Information

Disclosures

Human subjects: Consent was obtained by all participants in this study. Imam Abdulrahman Bin Faisal University issued approval N/A. Case Reports are waived by the Institutional Review Board in our institution. Conflicts of interest: In compliance with the ICMJE uniform disclosure form, all authors declare the following: Payment/services info: All authors have declared that no financial support was received from any organization for the submitted work. Financial relationships: All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. Other relationships: All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

References


