Mirizzi syndrome type Va: A rare coexistence of double cholecysto-biliary and cholecysto-enteric fistulae

Pavlos Lampropoulos, Nikolaos Paschalidis, Athanasios Marinis, Spiros Rizos

Pavlos Lampropoulos, Nikolaos Paschalidis, Athanasios Marinis, Spiros Rizos, First Department of Surgery, Tzaneio General Hospital, Afentouli and Zanni STR, 18536, Piraeus, Greece
 Author contributions: Lampropoulos P and Marinis A drafted the manuscript; Paschalidis P operated the patient; Rizos R and Paschalidis N reviewed the manuscript and provided the final approval.

Correspondence to: Pavlos Lampropoulos, MD, First Department of Surgery, Tzaneio General Hospital, Afentouli and Zanni STR, 18536, Piraeus, Greece. pav.lampropoulos@gmail.com
 Telephone: +30-210-9949460 Fax: +30-210-4592491
 Received: June 20, 2010 Revised: July 16, 2010
 Accepted: July 23, 2010
 Published online: October 28, 2010

Via Tricomi 1, Palermo, 90100, Italy; Charles Bellows, MD, Associate Professor, Chief of General Surgery, Tulane University, Department of Surgery SL-22, 1430 Tulane Ave, New Orleans, LA 70112, United States

Lampropoulos P, Paschalidis N, Marinis A, Rizos S. Mirizzi syndrome type Va: A rare coexistence of double cholecysto-biliary and cholecysto-enteric fistulae. *World J Radiol* 2010; 2(10): 410-413 Available from: URL: <http://www.wjgnet.com/1949-8470/full/v2/i10/410.htm> DOI: <http://dx.doi.org/10.4329/wjr.v2.i10.410>

Abstract

Mirizzi syndrome is a rare cause of intermittent obstructive jaundice, where an impacted stone in the cystic duct or Hartmann's pouch mechanically obstructs the common bile duct (CBD). We report a rare case of double cholecysto-biliary and cholecysto-enteric fistulae, in a 75-year-old female patient, presenting with a right upper quadrant abdominal pain and intermittent obstructive jaundice. Endoscopic retrograde cholangiopancreatography suggested Mirizzi syndrome. Operative findings included erosions of the lateral wall of the CBD and the second portion of the duodenum due to impacted gallstones. The defects were reconstructed primarily and a Kehr tube was inserted. The patient had an uneventful postoperative course and was discharged on the 14th postoperative day.

© 2010 Baishideng. All rights reserved.

Key words: Mirizzi syndrome; Obstructive jaundice; Gallstone; Cholecysto-enteric fistula; Endoscopic retrograde cholangiopancreatography

Peer reviewers: Roberto Miraglia, MD, Department of Diagnostic and Interventional Radiology, Mediterranean Institute for Transplantation and Advanced Specialized Therapies (IsMeTT),

INTRODUCTION

Mirizzi syndrome is a rare cause of intermittent obstructive jaundice, where an impacted stone in the cystic duct or Hartmann's pouch mechanically obstructs the common bile duct (CBD). Its incidence varies from 0.7% to 1.4% of patients with gallstone disease. In this report, we describe a rare coexistence of cholecysto-biliary and cholecysto-enteric fistulae, corresponding to type Va Mirizzi syndrome.

CASE REPORT

A 75-year-old female patient, with a past medical history of type II diabetes mellitus and hypertension, was admitted to our Medical Center's Surgical Department with a right upper quadrant (RUQ) abdominal pain, which initially presented 1 wk before she visited our hospital and was accompanied with malaise and vomiting, without any associated fever or chills. The patient was slightly anxious but in no acute distress, with normal vital signs and body temperature. Physical findings included icteric scleras and RUQ abdominal tenderness in deep palpation, without neither rebound tenderness or abdominal muscle rigidity nor palpable masses. Laboratory tests demonstrated a normal white blood cell count and hematocrit, while liver function tests reported 8.00 mg/dL total bilirubin, 6.20 mg/dL direct bilirubin, 76 U/L serum glutamic oxaloacetic transaminase,

Figure 1 Abdominal magnetic resonance imaging in a sagittal plane demonstrating two stones in common bile duct (arrow) with a diameter of 2.2 cm and cystic duct, measuring 9 mm and 12 mm respectively, with dilatation of the proximal biliary tree.

Figure 2 Magnetic retrograde cholangio-pancreatography demonstrating stones in common bile duct and proximal dilatation of the biliary tree.

81 U/L serum glutamic pyruvate transaminase, 327 U/L γ -glutamyltransferase, and 264 U/L alkaline phosphatase.

Abdominal ultrasound (US) and computer tomography (CT) showed a contracted gallbladder and stones in the CBD with an associated dilatation of the proximal biliary tree. Additionally, an abdominal magnetic resonance imaging (MRI) and retrograde cholangiopancreatography (MRCP) similarly demonstrated two stones in the CBD with a diameter of 2.2 cm and in the cystic duct, measuring 9 mm and 12 mm respectively, with dilatation of the proximal biliary tree (Figures 1 and 2).

Endoscopic intervention was decided and a retrograde cholangiopancreatography (ERCP) was carried out. A balloon catheter was inserted in the dilated CBD through a wide sphincterotomy. However, several attempts to clean the CBD did not reveal any bile duct stones, and an impression of an external CBD pressed possibly by the cystic duct was demonstrated. Surprisingly, when the patient was placed in an anti-Trendelenburg position, the dilated biliary tree subsided and the diagnosis of a Mirizzi syndrome was considered more possible (Figure 3).

Surgical treatment was decided and an exploratory laparotomy was performed through a Kocher incision. Intraoperative findings included a contracted gallbladder that was adherently attached to the duodenum and CBD.

Figure 3 Endoscopic retrograde cholangio-pancreatography showing gradual resolve of dilated biliary tree (A) and indirect signs of Mirizzi syndrome (B and C) when the patient was placed in an anti-Trendelenburg position.

During cholecystectomy, an impacted stone was found in the cystic duct which was firmly adhered to the CHD. After removal of the gallbladder, two defects were evident. One was on the lateral wall of the CHD involving less than 1/3 of the circumference, while the other was on the second portion of the duodenum. Both defects of the duodenum and CHD were sutured in one layer fashion. A choledochotomy was performed, the CHD was lavaged and finally a Kehr tube was inserted. The final diagnosis was a double cholecysto-choledochal and cholecysto-duodenal fistula. The patient had an uneventful postoperative course and was discharged on the 14th postoperative day.

DISCUSSION

Mirizzi syndrome is attributed to Pablo Mirizzi who was the first to describe it as a benign condition during

which a stone in the cystic duct or in the Hartmann's pouch impinges on the CBD, leading to mechanical obstruction by the stone itself or by secondary inflammation^[1]. He termed this phenomenon as "functional hepatic syndrome". In 1985, McSherry *et al*^[2] classified Mirizzi syndrome into two types based on ERCP findings: type I Mirizzi syndrome when there is extrinsic compression of the common hepatic duct and type II Mirizzi syndrome when the gallstone erodes the common hepatic duct wall, resulting in a cholecystocholedochal fistula. In 1989, Csendes *et al*^[3] proposed a new classification in order to clarify the definition of cholecystobiliary fistulae and to further establish an appropriate surgical treatment according to the underlying pathology. Thus, he defined the following four categories: type I: external compression of the CBD due to a stone impacted at the neck of the gallbladder or at the cystic duct (62.5%), type II: presence of a cholecystobiliary fistula (cholecystohepatic or cholecystocholedochal) due to erosion of the anterior or lateral wall of the CHD or CBD, respectively, with the fistula involving less than one-third of the circumference of the CBD (12.5%), type III: presence of a cholecystobiliary fistula with erosion of the wall of the CBD involving up to two-thirds of its circumference (12.5%), and type IV: presence of a cholecystobiliary fistula with complete destruction of the entire wall of the CBD (12.5%). In 2008, Beltran *et al*^[4] suggested that Mirizzi syndrome may not just end with a cholecystobiliary fistula but may further complicate the clinical picture with a superimposed cholecystoenteric fistula. Thus, a modified classification was proposed as follows: any aforementioned type plus a cholecystoenteric fistula without gallstone ileus (type Va), or with gallstone ileus (type Vb). Our patient had both a cholecysto-biliary fistula and a cholecysto-enteric fistula, without any gallstone ileus. Therefore, the patient had a Mirizzi syndrome type Va, according to the latest classification.

This syndrome is a very rare cause of obstructive jaundice and occurs in 0.7%-1.4% of patients with gallstones^[3-6]. It is very difficult to establish a preoperative diagnosis of intermittent jaundice, as there is no specific clinical or laboratory presentation for this syndrome. However, jaundice, abdominal pain and alternating hepatic biochemical profile can suggest Mirizzi syndrome in 80% of the cases. Ultrasonography and abdominal CT disclose a dilated CBD above the gallbladder neck, an abrupt disruption of the CHD and a decompressed gallbladder with stones^[7]. Furthermore, laparoscopic US and CT scanning may rule out a tumor mass^[8,9]. MRCP and MRI are the useful methods for the diagnosis of Mirizzi syndrome as they can precisely demonstrate the presence and degree of dilated biliary tree, and can also reveal extrinsic narrowing of the CHD as well as anatomical variations and malformations^[10].

ERCP remains the gold standard for preoperative diagnosis of Mirizzi syndrome in jaundiced patients, although it cannot often document the presence of a fistula. ERCP provides indirect signs such as lateral filling gap of

the CHD, central dilatation of the biliary tree and insertion of the cystic duct at the point of obstruction and/or complete obliteration of the cystic duct^[11]. In our case, an interesting finding during ERCP suggesting indirectly Mirizzi syndrome, was the fact that biliary tree dilatation subsided when the patient was placed in an anti-Trendelenburg position. To our knowledge, such an indirect sign has never been reported in the English literature. Preoperative diagnosis of Mirizzi syndrome by ERCP is important to prevent complications^[12]. At last, a recent study demonstrated that MRCP as a noninvasive alternative to ERCP is a useful preoperative diagnostic modality for Mirizzi syndrome when combined with CT, with a higher accuracy and better inter-observer agreement compared with CT alone^[13].

Management of patients with Mirizzi syndrome is a real challenge, especially if a fistula is present. Several minimal invasive techniques have been proposed. Percutaneous treatment with electrohydraulic lithotripsy under cholangioscopic control with a mother-daughter endoscope system has been reported to successfully fragment and clear all stones in 38 cases^[14]. These advanced endoscopic approaches cannot be applied to all hospital settings but they are definite treatment modalities that might prove helpful in the future. Moreover, the impacted stone cannot be extracted endoscopically and therefore surgery is the only chance for cure. Although successful attempts of laparoscopic treatment have been reported, conversion to laparotomy is most commonly required with an overall conversion rate of 31.7%-100%^[15,16]. Thus, several open surgical procedures are described for the treatment of different types of Mirizzi syndrome. Surgical strategy, however, should be based on three basic principles: removal of the obstructing stone and gallbladder, restoration of the integrity of biliary tree and in case of type V, closure of the gastrointestinal tract defect.

More specifically, the usual surgical treatment of type I Mirizzi syndrome is laparoscopic or open cholecystectomy and exploration of the CBD. In case of type II Mirizzi's syndrome, most studies favor open techniques, although Binnie *et al*^[17] reported successful laparoscopic treatment of type II Mirizzi's syndrome after previous endoscopic biliary stenting. Choledochoplasty with gallbladder flap, cholecystoduodenostomy and Roux-en-Y hepaticojejunostomy are the possible surgical approaches to the treatment of type II Mirizzi's syndrome^[10]. Additionally, partial cholecystectomy with primary closure using the gallbladder remnant and T-tube insertion though the fistula opening are options for the treatment of type II Mirizzi syndrome^[18,19]. However, T-tube insertion into the CBD rather than the fistula opening reduces the risk of stenosis^[20,21]. Roux-en-Y hepaticojejunostomy is recommended as a safe approach in cases of wider CBD defects (types III and IV) in order to avoid complications such as biliary leakage and stricture formation^[3,9,18,20]. Finally, a surgeon must be aware that Mirizzi syndrome coexists with cancer in 27% of the cases^[22], as it requires a totally different approach.

The co-existence of double cholecysto-biliary and cholecysto-enteric fistulae is rare. Mirizzi syndrome should be highly suspected in cases of unexplained intermittent obstructive jaundice. Several imaging modalities for the diagnosis of Mirizzi syndrome exist and have been proved useful preoperatively. Treatment of Mirizzi syndrome is considered challenging and determined by the type of Mirizzi syndrome.

REFERENCES

- 1 **Mirizzi PL.** [Hepatic duct syndrome.] *Cir Cir* 1964; **32**: 641-664
- 2 **McSherry CK, Ferstenberg H, Calhoun WF, Lahman E, Virshup M.** The natural history of diagnosed gallstone disease in symptomatic and asymptomatic patients. *Ann Surg* 1985; **202**: 59-63
- 3 **Csendes A, Díaz JC, Burdiles P, Maluenda F, Nava O.** Mirizzi syndrome and cholecystobiliary fistula: a unifying classification. *Br J Surg* 1989; **76**: 1139-1143
- 4 **Beltran MA, Csendes A, Cruces KS.** The relationship of Mirizzi syndrome and cholecystoenteric fistula: validation of a modified classification. *World J Surg* 2008; **32**: 2237-2243
- 5 **Bower TC, Nagorney DM.** Mirizzi syndrome. *HPB Surg* 1988; **1**: 67-74; discussion 75-76
- 6 **Mishra MC, Vashishtha S, Tandon R.** Biliobiliary fistula: preoperative diagnosis and management implications. *Surgery* 1990; **108**: 835-839
- 7 **Lakhtakia S, Gupta R, Tandan M, Rao GV, Reddy DN.** Mirizzi's syndrome: EUS appearance. *Gastrointest Endosc* 2006; **63**: 322-323; discussion 323
- 8 **Callery MP, Strasberg SM, Doherty GM, Soper NJ, Norton JA.** Staging laparoscopy with laparoscopic ultrasonography: optimizing resectability in hepatobiliary and pancreatic malignancy. *J Am Coll Surg* 1997; **185**: 33-39
- 9 **Baer HU, Matthews JB, Schweizer WP, Gertsch P, Blumgart LH.** Management of the Mirizzi syndrome and the surgical implications of cholecystcholedochal fistula. *Br J Surg* 1990; **77**: 743-745
- 10 **Gomez D, Rahman SH, Toogood GJ, Prasad KR, Lodge JP, Guillou PJ, Menon KV.** Mirizzi's syndrome--results from a large western experience. *HPB (Oxford)* 2006; **8**: 474-479
- 11 **Hazzan D, Golijanin D, Reissman P, Adler SN, Shiloni E.** Combined endoscopic and surgical management of Mirizzi syndrome. *Surg Endosc* 1999; **13**: 618-620
- 12 **Yonetci N, Kutluana U, Yilmaz M, Sungurtekin U, Tekin K.** The incidence of Mirizzi syndrome in patients undergoing endoscopic retrograde cholangiopancreatography. *Hepatobiliary Pancreat Dis Int* 2008; **7**: 520-524
- 13 **Yun EJ, Choi CS, Yoon DY, Seo YL, Chang SK, Kim JS, Woo JY.** Combination of magnetic resonance cholangiopancreatography and computed tomography for preoperative diagnosis of the Mirizzi syndrome. *J Comput Assist Tomogr* 2009; **33**: 636-640
- 14 **Seitz U, Bapaye A, Bohnacker S, Navarrete C, Maydeo A, Soehendra N.** Advances in therapeutic endoscopic treatment of common bile duct stones. *World J Surg* 1998; **22**: 1133-1144
- 15 **Yeh CN, Jan YY, Chen MF.** Laparoscopic treatment for Mirizzi syndrome. *Surg Endosc* 2003; **17**: 1573-1578
- 16 **Schäfer M, Schneiter R, Krähenbühl L.** Incidence and management of Mirizzi syndrome during laparoscopic cholecystectomy. *Surg Endosc* 2003; **17**: 1186-1190; discussion 1191-1192
- 17 **Binnie NR, Nixon SJ, Palmer KR.** Mirizzi syndrome managed by endoscopic stenting and laparoscopic cholecystectomy. *Br J Surg* 1992; **79**: 647
- 18 **Karademir S, Astarcioğlu H, Sökmen S, Atila K, Tankurt E, Akpınar H, Coker A, Astarcioğlu I.** Mirizzi's syndrome: diagnostic and surgical considerations in 25 patients. *J Hepatobiliary Pancreat Surg* 2000; **7**: 72-77
- 19 **Corlette MB, Bismuth H.** Biliobiliary fistula. A trap in the surgery of cholelithiasis. *Arch Surg* 1975; **110**: 377-383
- 20 **Sharma AK, Rangan HK, Choubey RP, Thakur SK, Kumar A.** Pitfalls in the management of Mirizzi's syndrome. *Trop Gastroenterol* 1998; **19**: 72-74
- 21 **Balthazar EJ.** The Mirizzi syndrome, inflammatory structure of the common hepatic duct. *Am J Gastroenterol* 1975; **64**: 144-148
- 22 **Redaelli CA, Büchler MW, Schilling MK, Krähenbühl L, Ruchti C, Blumgart LH, Baer HU.** High coincidence of Mirizzi syndrome and gallbladder carcinoma. *Surgery* 1997; **121**: 58-63

S- Editor Cheng JX L- Editor Wang XL E- Editor Zheng XM