Mirizzi syndrome type IV: A case report

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Abstract:

Introduction: Mirizzi Syndrome (MS) is a rare complication of chronic cholelithiasis triggered by impacted calculus in Hartmann pouch or cystic duct. Type IV is complete destruction of the bile duct and represents the most uncommon type. We reported a case of Mirizzi syndrome type IV discovered during elective laparoscopic cholecystectomy.

Case report: A 35 year female was found during elective laparoscopic cholecystectomy to have abnormal anatomy. Intraoperative cholangiogram and conversion to open approach confirmed the diagnosis of Mirizzi syndrome type IV. Cholecystectomy and hepaticojejunostomy were done. Recovery was uneventful.

Conclusion: Surgeons performing laparoscopic cholecystectomy should be aware of the possibility of Mirizzi syndrome. They should have a low threshold for intraoperative cholangiogram and hepatobiliary surgeons consultation.

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I. Introduction

Mirizzi Syndrome (MS) is a rare complication of chronic cholelithiasis triggered by impacted calculus in Hartmann pouch or cystic duct, causing extrinsic obstruction of the common hepatic duct. Preoperative diagnosis on the basis of clinical presentation and investigations is not possible in majority of patients. Due to inflammation and complete obliteration of Calot's triangle, cholecystectomy, either by open or laparoscopic approach, poses a significant risk of bile duct injury.

Incidence of MS is reported to be less than 1%. Csendes originally classified MS into four types (figure 1): type I includes those with external compression of the common bile duct (CBD); type II is a cholecystobiliary fistula presenting with erosion of less than one third of the circumference of the bile duct; type III is a fistula involving up to two-thirds of the duct circumference; and type IV is complete destruction of the bile duct. MS type IV represents the most uncommon type.

We reported a case of MS type IV found intraoperatively during elective laparoscopic cholecystectomy.

Case report:

A 35 year old female patient was admitted for elective laparoscopic cholecystectomy for symptomatic cholelithiasis on April 15, 2016. The patient had no previous surgical or medical history. On physical exam the patient looked well, Afebrile, not tachycardic, pale or jaundiced. Her abdomen examination revealed mild right upper quadrant tenderness with negative Murphy sign and no palpable masses. Her laboratory tests showed normal white blood cell count, hemoglobin, bilirubin and liver enzymes. Her preoperative ultrasound showed only multiple gall bladder (GB) stones with no pericholecystic fluid and normal CBD. The procedure started...
with introducing 10mm optical trocar using open technique. Three more trocars were inserted under vision. After fundus retraction, Calot's triangle was not clearly visualized. After meticulous dissection, a structure supposed to be the cystic duct (CD) was identified and dissected free. Because we were in doubt about that structure decision was made to perform intraoperative cholangiogram through that duct which showed the dye passing directly into the duodenum without any visualization of proximal biliary system. Decision then was made to convert to open technique through right subcostal incision. We confirmed that the structure once supposed to be a CD was indeed the CBD and there was no any CD. The CBD distal part measured about 1.5 cm in length but proximally it completely emerged with the lumen of the GB forming as a continuation or part of it and at this moment diagnosis of MS type IV was made. Proximal to the GB-CBD complex the common hepatic duct was found to be full of stones. Hepatobiliary surgeons were consulted and they confirmed our diagnosis and perform cholecystectomy and hepaticojejunostomy. The patient was discharged on postoperative day 7 with uneventful recovery.

II. Discussion

The Mirizzi syndrome refers to common hepatic duct obstruction caused by an extrinsic compression from an impacted stone in the CD or Hartmann's pouch of the GB. Mirizzi reported his syndrome for the first time in 1940. The pathophysiological process leading to the subtypes of MS has been explained as an inflammatory phenomenon secondary to a pressure ulcer caused by an impacted gallstone which can cause first external obstruction of the bile duct and eventually erodes into the bile duct evolving into a cholecystocholedochal fistula with different degrees of communication between the GB and bile duct. Anatomical variations with long CD parallel to the common hepatic duct or a low insertion of the cystic duct predispose to development of this syndrome.

MS appears in 1% to 2% of patients with symptomatic cholelithiasis with higher incidence in Central and South America where the reported incidence is 4.7% to 5.7% with type IV the most uncommon type. Reverdito R et al reported 12 cases of MS type IV in 3,691 cholecystectomies performed from December 2001 to September 2013 with incidence of 0.3%. Kulkarni SS et al reported MS in 60 patients out of 4939 patients (1.21 % incidence) who underwent cholecystectomy over 6 years, none of these 60 patients had type III or IV MS. Ashok Kumar et al reported MS in 169 patients among 8000 cholecystectomies performed between 1989 and 2011, and only 10 of them had type IV. Testini M et al reported 18 cases of MS over 10 years with only one of them had type IV. Similarly, Kamalesh NP et al reported 20 patients with MS among 1530 cholecystectomies, only one patient had MS type IV. Among 8697 cholecystectomies, Xu XQ et al reported 27 patients with MS, none of them was type IV. Lastly, Cui Y et al found 198 cases of MS among 29 875 patients who underwent cholecystectomy for cholelithiasis, only 6 patients had type IV. The case we reported has been the first case of MS type IV during 8 year experience at our institution.

The clinical diagnosis of MS is difficult, since there are no pathognomonic patterns of presentation. Ultrasonography and CT scan were diagnostic in only 10% patients, with ERCP and/or MRCP to confirm the diagnosis. Despite of all these modern diagnostic tools, the problem may become apparent only during surgery. At this situation, as in our case, old is still gold and intraoperative cholangiogram can provide a clearer picture of the anatomy and hence reduce the risk of injury to CBD.

Surgical management of Csendes type IV MS is well established as the procedure of choice is cholecystectomy with Roux-en-Y hepatico-jejunostomy.

III. Conclusion

Surgeons should be aware of the possibility of MS during laparoscopic cholecystectomy. Difficulty in identifying structures at Calot's triangle, presence of abnormal anatomic relations or abnormal size CD should alert the surgeon to the presence of biliary anomaly. There should be a low threshold for intraoperative cholangiogram and consultation of experienced hepatobiliary surgeons.

References


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