Laparoscopic Cholecystectomy of Gallbladder Stone Presenting as Empyema in a 9 Year-Old Boy: A Case Report

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Abstract

Background: The gall bladder stones (GBS) are common disease related to diet habit, obesity, hematological diseases, diabetes mellitus and receiving total parenteral nutrition. GBS is a rare cause of hospitalization of pediatric patients but have been increasingly diagnosed in recent years due to widespread use of ultrasonography. The aim of this case report study is to determine the clinical presentation, risk factors, valuable diagnostic procedures and outcome of laparoscopic management of cholelithiasis in children.

Case report: Our case is a 9-year-old boy with empyemic gallbladder complaining of attacks of colicky epigastric pain associated with recurrent vomiting. On abdominal palpation, there is positive sign of deep tenderness at hypocondrium area and positive murphy sign. He has central obesity with BMI 29.9, skin fold more than 3.5 cm and waist circumference more than 90 cm. He has History of high cholesterol and fatty diet and fast food habit, prolonged intake of vitamin D and positive family history. While, no other associated risk factors were detected in our patient. All laboratory investigations were within normal ranges even for his lipid profile. Although, empyema was present, no leukocytosis was detected. Abdominal ultrasound showed the gall bladder wall is slightly thick surrounded by pericholecystic fluid, solitary gall bladder stone of 3 cm size and normal CBD. While, post-operative histopathological examination revealed thick wall gallbladder with multiple tiny stones of cholesterol type. The patient was treated with laparoscopic cholecystectomy with promising outcome and no complications.

Conclusion: Pediatric surgeons should consider cholelithiasis as from differential diagnosis of abdominal pain associated with vomiting in children. Empyema of gallbladder should be considered once murphy sign is positive in such child patients. Child obesity, fatty diet, vitamin D deficiency and family history are possible risk factors. Ultrasonography is the mainstay of diagnosis. Laparoscopic cholecystectomy is the appropriate management for symptomatic cholelithiasis in children.

Keywords: Gallstones; Empyema; Children; Boy; Laparoscopic cholecystectomy

Introduction

Gallbladder disease today is a common health problem affecting 10% to 15% of the adult population [1]. In contrast to adulthood, gallstones in childhood are rare. However, the incidence is increasing with the diagnosis of asymptomatic gallstones with the widespread use of ultrasonography [2]. Risk factors for gallstones in children include hemolytic disease, obesity, prematurity, sepsis, total parenteral nutrition (TPN), chronic liver disease, inflammatory bowel diseases, diuretic use, and ceftriaxone use [3]. Sickle cell disease is the most important cause of cholelithiasis in children. Pigment gallstones affect 15% of children with SCD younger than 10 years of age and more than 80% of those older than 30 years [4]. Gallstones found in adults are primarily cholesterol stones or mixed stones but in children pigment stones formed as a consequence of hemolytic diseases like sickle cell anemia, thalassemia and hereditary spherocytosis are more common [5]. In general, gallstones are uncommon in children, with patients under 15-year-old, comprising only 0.1-0.2% of the incidence of the disease [6]. Even so, define the association between the obesity and gall bladder stones in pediatric population were very few studies and taking long time on the PubMed. These few clinical reports suggest that hemolytic diseases are no longer the most frequent cause of pediatric gallbladder disease. Concomitant with the epidemic of childhood obesity and the shift towards extreme childhood obesity, the prevalence of gallstones in children and adolescents may be increasing due to childhood obesity [7]. In addition of risk of GBS in obese youngest children now days, the risk of choledocholithiasis had significantly higher with high BMI than those with simple cholelithiasis [8]. Cholelithiasis is sometimes diagnosed in patients incidentally or as silent stones. In some other cases, they are reported in
association with clinical symptoms such as cholecystitis and cholangitis [9,10]. However, there is little information about the epidemiology of GBS in children from Libya and no consensus among Libyan pediatric surgeons regarding management of gallstones in children. We want to report our experience of a rare case of 9 years-old with gall bladder large stone and sludge bile, complicated with empyema, operated straight forward by urgent laparoscopic cholecystectomy.

Case Presentation

Obese male child 9-year-old presented to our outpatient clinic after having attended several other facilities, complaining of attacks of colicky epigastric pain associated with recurrent vomiting 2 months back, aggravated by fatty meal, relieved by IV medications and vomiting. History taking revealed high cholesterol and fatty diet and fast food habit; long duration of Vitamin D intake and a positive family history of laparoscopic cholecystectomy (LC) for his mother and Uncle (his mother’s brother). No history of easily bruising, oral mucosa petechial rash, bleeding tendency, blood transfusion or previous operations. Physical examination revealed normal vital signs (temperature, pulse rate, blood pressure) were normal. The child was obese with abdominal skin fold 5 cm, axillary skin folds 3 cm, Height 140 cm, weight 58 kg, body mass index (BMI) is 29.9 (normal range in this age is between 14.2 and 19.4), waist circumference 93 cm, mid-arm circumference 29cm and mid-thigh circumference 55 cm. No jaundice, no pallor and no edema were present and oral mucosa and nostrils were completely normal. Per abdominal examination showed no distension, normal movement with breathing, no scratch markings, no visible mass. Generally, the abdomen was soft by palpation with positive sign of deep tenderness at hypochondrium area and positive murphy sign. Bowel sounds was normally heard.

Laboratory findings

Routine laboratory examination was done including WBC, full blood count, ESR renal function test, liver function tests, serum amylase and lipase, serum sodium and potassium, HBA1C and blood sugar. All the previous investigations were within normal limits. Total bilirubin was 0.2 mg/dl and Alkaline phosphatase was 302 U/L which is not considered to be high in this age group. Other was normal levels. In addition, lipid profile (triglycerides, cholesterol, HDL, LDL) was normal No other laparotomy investigations were indicated.

Radiological findings

Abdominal Ultrasonography showed homogenous fatty liver, the gall bladder wall is slightly thick surrounded by pericholecystic fluid, solitary single gall bladder stone of 3 cm size, Common Bile Duct is normal in diameter, spleen and otherwise are normal (Figures 1-4).

Figure 1: Abdominal US showing homogenous fatty liver.

Figure 2: Abdominal US showing slightly thick gall bladder wall with CBD normal in diameter.

Figure 3: A bdominal US showing thick gallbladder wall surrounded by pericholycystic fluid.
Abdominal US showing solitary single gallbladder stone of 3 cm size.

Laparoscopic cholecystectomy

The positive Murphy sign and the deep tenderness of the right hypochondria area regardless the afebrile status and normal level of WBC, let our thinking and decision are very clear and the patient was candidate for elective laparoscopic cholecystectomy. After appropriate preoperative investigations and discussion with anesthesia team, critical view Laparoscopic cholecystectomy was done under general anesthesia, the time of operation was one hour and twenty minutes, standard four laparoscopic ports inserted, after pneumoperitoneum achieved at 12 mmHg and camera 300 inserted, the view was difficult due to fatty omentum, the gall bladder was empyemic and large in size with significant adhesions and edematous wall, release of adhesions performed and critical view applied, The Calot’s triangle has been skeletonized, was found normal. Cystic duct and cystic artery were identified, clipped and ligated, no gall bladder bed oozing is noted and finally cholecystectomy done (Figure 5).

Recovery of the patient was smoothly without any complications, patient kept in the ward nil per mouth with iv fluid maintainace, Antibiotics IV, perfelgan IV and ranitidine IV. 12 hours post-operative, the general condition was normal, patient mobilized, his vital signs were within normal, afebrile, the bowel sound early regain to function, diet initiation with no vomiting and nausea. He has been discharged DAY 2 post operation with well general condition; no complain with only few cc serousanguinous discharge in the drain. Stitches has been removed one week after operation. Regular follow up applied at 2 months, 6 months and 1 year. The patient was symptomatic free with no GIT or Cholecystectomy related complications.

Histopathological findings

After cholecystectomy, the biopsy was sent to histopathology. The result showed that lumen contains tiny yellowish lucent stones, ulcerated mucosa, moderate wall fibrosis and transmural infiltrations of lymphocyte and eosinophil cells (Figures 6-9).

Figure 6: Gallbladder containing tiny yellowish lucent stones.

Figure 7: Gallbladder showing ulcerated mucosa.
The relation of cholesterol gall bladder stones in obese child could be related to cholesterol metabolism or related to the compression effect of fatty omentum and intestine on the stasis of biliary radicles (lithogenic effect). Hypersaturation of bile caused by either increased hepatic cholesterol uptake or increased cholesterol synthesis. Second, dysmotility and impaired contraction of the gallbladder is caused by the direct influence of cholesterol at the cellular level onto the plasma membrane of smooth muscle cells in the gallbladder wall [8].

Tsai performed prospective study of abdominal adiposity and gallstone disease in US men, the conclusion was, the collected data for abdominal obesity in relation to the incidence of symptomatic gallstone disease suggested presence of a significant association between abdominal adiposity and the incidence of symptomatic gallstone disease. As measures of abdominal adiposity, abdominal circumference and waist-to-hip ratio predict the risk of developing gallstones independently of body mass index [17].

Several risk factors for gallstones in adults are well-established, including age, female sex, Hispanic ethnicity, obesity, use of female sex hormones, pregnancy, sedentary lifestyle, and a family history of gallstones. However, scant information is available regarding risk factors for gallstones in the pediatric population [18].

In our patient, there was positive family history and history of long duration of vitamin D intake. While, many other risk factors were absent as no hemolytic diseases, no diabetic, no history of biliary atresia or choledochal cyst.

Generally, the incidence and prevalence of cholelithiasis are influenced by age, gender, genetics, and race. Epidemiological studies have indicated the involvement of genetic factors in the formation of cholelithiasis. The effect of a gene on incontinencia pigmenti chromosome has been confirmed in the formation of cholelithiasis. In fact, patients with ABCB11 mutations are at a higher risk of cholelithiasis [19].

While, Onal and his collages in their study revealed that vitamin D deficiency is suggested to be associated with gallbladder stasis, and a role for vitamin D supplementation is thought to have potential to prevent gallstones in this special population [20]. This may explain the occurrence of gallbladder stones in our patient as a result of vitamin D deficiency in early childhood, and that was not due to increased intake of vitamin D as it seems to be. Our patient takes vitamin D to replace its deficiency and the deficiency of vitamin D itself is responsible for gall bladder stasis with development of stones in our case. In other studies, long-term use and high dose use of ceftriaxone was founded to be associated with increase the probability of occurrence of gallstones [21].

Our patient was complaining of attacks of colicky epigastric pain associated with recurrent vomiting. On abdominal palpation, there is positive sign of deep tenderness at hypochondrium area and positive murphy sign. The results of laboratory investigation was mostly normal in our case, even WBCs was of normal level despite the presence of empyema and that may be explained as a result of the previous empirical...
antibiotic course administrated in other clinics before the case attended our outpatient clinic. Abdominal ultrasound was the method of choice in diagnosis of the reported case. While, radiological investigations other than ultrasound (i.e., MRCP) was not indicated, so it was not performed. Cholelithiasis in children have been increasingly diagnosed in recent years due to better medical imaging (especially ultrasonography) and its usage in investigating children with unexplained abdominal pain [11]. So, we should consider the Cholecystolithiasis as from the differential diagnosis of unexplained abdominal pain associated with vomiting in children and we have to confirm that by conducting abdominal US, to avoid the complications of GBS as acute cholecystitis, empyema, biliary pancreatitis and sepsis. In addition, empyema of gallbladder should be considered once murphy sign is positive in such child patients even with absence of leukocytosis.

Because The youngest age of the patient, making the final decision of laparoscopic cholecystectomy (L.C.) is delayed from other clinics, to considerations of exclusion of some risk factors as hemolytic diseases. For this reason, he was operated lately. The patient was treated with laparoscopic cholecystectomy with prolonged time but with clear anatomy.

This is in agreement with the recommendations of other studies, which stated that the laparoscopic cholecystectomy is safe and preferable in pediatric patients. L.C. have better results, short post-operative hospital stays and with low complication rates in particular, zero bile duct injuries were noted [22,23].

Post-operative histopathological examination revealed thick wall gallbladder with multiple tiny stones of cholesterol type. Our findings were compatible with that of Agostino Di Ciaula who investigated several pathogenic mechanisms in literature review article, the Major pathogenetic factors for cholesterol gallstones include a genetic background, hepatic hyper-secretion of cholesterol and supersaturated bile which give life to precipitating cholesterol crystals that accumulate and grow in a sluggish gallbladder [24]. While, our study is in contrary to our other studies stated that pigment stones containing bilirubin salts are more common in the pediatric population. Nevertheless, our explanation is that these types of bilirubin stones are associated with hemolytic disorders, most commonly sickle cell anemia, which no of them were identified in our patient.

Conclusion

Despite cholelithiasis is rare in children, pediatric surgeon should consider the cholecystolithiasis as from differential diagnosis of abdominal pain associated with vomiting in the youngest age group even with absence of positive laboratory investigations, to avoid the complications of GBS as acute cholecystitis, empyema, biliary pancreatitis and sepsis. In addition, empyema of gallbladder should be considered once murphy sign is positive in such child patients even with absence of leukocytosis. Ultrasonography is the mainstay of diagnosis and it is very useful in detecting these cases. Child obesity, fatty diet, and family history were the most frequent risk factors in our patient. Vitamin D deficiency is suggested to be associated with gallbladder stasis and may increase the risk of cholelithiasis. Laparoscopic cholecystectomy is the appropriate management for symptomatic cholelithiasis in children, as it can be performed safely using technique of single-incision. We emphasize the importance of further researches to explore the epidemiology of gallbladder disease in children for better diagnosis and management of such cases.

Informed Consent

An informed consent was obtained from our patient’s parents for publication of this case report and the accompanying images.

Competing Interests

The authors declare they have no competing interests.

References


