

MANAGEMENT OF INFANTILE CHOLEDOCHOLITHIASIS AND CHOLECYSTITIS: A CASE REPORT

General Surgery

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ABSTRACT

This case report presents the clinical management of a 6-month-old male infant presenting with fever, vomiting, and incessant crying. After initial conservative management, the patient was diagnosed with choledocholithiasis and cholecystitis. Due to the infant's weight, ERCP with CBD stenting was deemed inappropriate, and thus, a hepaticojunostomy procedure was performed. The postoperative period was uneventful, with resolution of symptoms and normalization of biliary parameters. This case underscores the challenges in diagnosing and treating gallstone-related conditions in infants, highlighting the importance of surgical intervention in certain cases.

KEYWORDS

Choledocholithiasis, Cholecystitis, Hepaticojunostomy, Infant, Case Report

INTRODUCTION:

Gallbladder diseases, though historically recognized since antiquity, continue to pose diagnostic and therapeutic challenges, especially in pediatric populations¹. While gallstones, or cholelithiasis, have been predominantly associated with adults, their occurrence in children, particularly infants, is relatively rare². Recent advancements in diagnostic imaging, such as ultrasound, have facilitated earlier detection of gallbladder diseases in pediatric patients. Additionally, various predisposing factors, including prematurity, hemolytic disorders, and medical interventions like total parenteral nutrition (TPN), contribute to the rising incidence of cholelithiasis in infants³.

This case report presents a 6-month-old male infant with choledocholithiasis, highlighting the diagnostic workup, therapeutic interventions, and postoperative outcomes. The case underscores the importance of prompt recognition and appropriate management of gallbladder diseases in pediatric patients. With the evolving landscape of pediatric healthcare, it is imperative for clinicians to remain abreast of the latest diagnostic modalities and treatment strategies to optimize outcomes for infants with gallstone-related conditions.

In this context, we discuss the risk factors, clinical presentation, diagnostic modalities, and therapeutic options pertinent to infantile choledocholithiasis and cholecystitis. The case also emphasizes the multidisciplinary approach required for comprehensive care, involving pediatricians, surgeons, and gastroenterologists. Through this case report, we aim to contribute to the existing literature on pediatric gallbladder diseases and underscore the importance of tailored management strategies in this vulnerable patient population.

Case Presentation:

A 6-month-old male infant was brought to the pediatric clinic with complaints of fever, vomiting, and incessant crying. Physical examination revealed icteric sclera and pale-colored stools. Laboratory investigations showed elevated liver enzymes and bilirubin levels. Ultrasonography (USG) revealed choledocholithiasis with biliary dilatation. Conservative management with antibiotics and ursodeoxycholic acid led to symptomatic improvement initially.

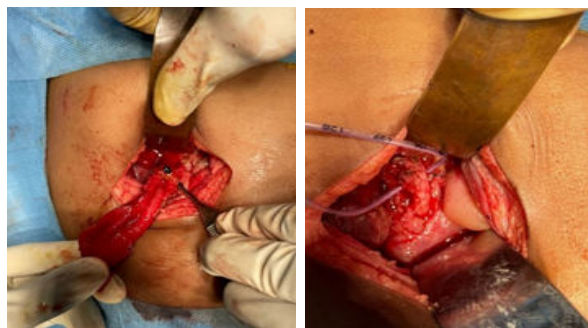
Preoperative Evaluation:

Four days later, the infant presented again with icteric eyes and pale stools. Blood investigations confirmed persistent biliary obstruction. USG indicated slow clearance from the liver, suggesting ongoing biliary obstruction. ERCP with CBD stenting was considered but deemed inappropriate due to the infant's weight being below the threshold for safe intervention. Therefore, a hepaticojunostomy procedure was planned.

	24/04/23	27/04/23
Total bilirubin	3.8	2.5
Direct bilirubin	2.9	1.8
Total count	28,000	36,000
CRP	104	307
Alkaline phosphatase	198	225

Operative Procedure:

Under general anesthesia, a right subcostal incision was made. Intraoperatively, the gallbladder was found to be contracted and inflamed and was subsequently removed, with ligation of the cystic duct. A small incision was made over the common bile duct (CBD), revealing multiple pigmented stones. Saline irrigation was performed, and a DJ stent was inserted into the CBD to ensure drainage into the duodenum.



Postoperative Course:

Postoperative imaging showed mild central intrahepatic biliary dilatation and the CBD stent in situ. There were no postoperative complications, and the patient recovered well. Follow-up assessments revealed resolution of symptoms and normalization of liver function tests.

	05/05/23		02/11/23
Total bilirubin	1	Total bilirubin	0.7
Direct bilirubin	0.7	Direct bilirubin	0.2
Total count	10,000	Total count	8,000
CRP	28	CRP	5
Alkaline phosphatase	168	Alkaline phosphatase	156

DISCUSSION:

Throughout history, diseases of the gallbladder have been recognized, with gallstones even discovered in Egyptian mummies⁴. While gallstone formation, known as cholelithiasis, is more commonly associated with adults, its occurrence in children, particularly infants, is a relatively rare phenomenon⁵. The prevalence of cholelithiasis in pediatric populations has been reported to range from 0.13% to 0.22%⁶. Recent advancements in diagnostic imaging techniques, such as ultrasound, have contributed to the increased detection of gallbladder diseases in children⁷. Additionally, factors like total parenteral nutrition (TPN), obesity, and prematurity have been implicated in the rising incidence of cholelithiasis in pediatric patients. Given the potential implications of gallbladder disease in children, healthcare professionals must remain informed about the latest diagnostic and treatment modalities^{2,8}.

Risk Factors and Etiology:

Various factors contribute to the formation of gallstones in children,

including prematurity, hemolytic disorders, gastrointestinal conditions like intestinal resection and Crohn's disease, hepatobiliary issues, pulmonary conditions, and certain medications⁸. Approximately 70% of infants with cholelithiasis have identifiable risk factors, most commonly associated with heart surgery, TPN, and medication use¹. However, in some cases, the cause of gallstones remains unknown. Congenital biliary abnormalities and hemolytic disorders, such as sickle cell anemia and hereditary spherocytosis, are significant predisposing factors for cholelithiasis in infants. Furthermore, factors such as prolonged TPN use, ileal resection, phototherapy, and infections have been implicated in gallstone development^{2,9}.

Clinical Presentation and Diagnosis:

Gallstones in infants are often asymptomatic but may occasionally manifest as cholestatic jaundice, transient alcoholic stools, sepsis, or abdominal pain. Notably, symptomatic infants are more likely to have common bile duct stones rather than gallbladder-only stones³. Ultrasonography is the primary diagnostic modality for detecting gallstones, with a sensitivity and specificity of approximately 95% for gallbladder cholelithiasis. However, its sensitivity for choledocholithiasis is lower, ranging from 50% to 75%. Gallstones in children may be radiopaque in 20–50% of cases¹⁰.

Management and Treatment:

The management of gallstones in children depends on various factors, including the patient's age and clinical symptoms. Spontaneous resolution of cholelithiasis occurs in 35–60% of patients, with a low recurrence rate¹¹. Conservative management is often recommended, especially for asymptomatic infants and toddlers. Surgical intervention, such as cholecystectomy, is typically reserved for symptomatic cases, persistent asymptomatic cholelithiasis, or radiopaque calculi^{12,13}. Alternative medical therapies, such as ursodeoxycholic acid and cholesterol-lowering medications, have not been proven effective in dissolving existing stones and are not routinely recommended¹⁴.

In our case, the decision to perform a hepaticojejunostomy was based on the infant's weight and the inappropriateness of ERCP. Hepaticojejunostomy is a surgical procedure commonly used to bypass biliary strictures or obstructions, providing an alternative route for bile drainage. Despite its effectiveness, the procedure carries inherent risks, particularly in pediatric patients, including anastomotic leaks, strictures, and bile reflux¹⁵. However, in our case, the postoperative course was uneventful, with successful resolution of symptoms and normalization of biliary parameters.

CONCLUSION:

Infantile choledocholithiasis and cholecystitis present unique challenges in diagnosis and management. While gallstones in children, particularly infants, remain relatively rare, advancements in diagnostic imaging have led to increased detection rates. Prompt recognition and appropriate intervention are crucial for optimizing outcomes in affected infants. While conservative measures may suffice in some cases, surgical intervention such as hepaticojejunostomy may be necessary in others, particularly when endoscopic options are not feasible. This case highlights the importance of a multidisciplinary approach involving pediatricians, surgeons, and gastroenterologists in the comprehensive care of infants with gallstone-related conditions.

Conflict of Interest:

The authors declare no conflict of interest.

Funding:

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Ethical Approval:

Ethical approval was obtained from the institutional review board.

Informed Consent:

Informed consent was obtained from the patient's parents for publication of this case report.

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