

A Rare Case of Choledochal Cyst complicated by Choledocholithiasis with Acute Pancreatitis

Zannatul Ferdous Sonia¹, ARM Luthful Kabir², Mahmuda Hasan³, Akmat Ali⁴, Richmond Ronald Gomes⁵, Masuma Khan⁶

¹Assistant Professor, Pediatrics, Ad-din Women's Medical College Hospital, Dhaka

²Professor, Pediatrics, Ad-din Women's Medical College Hospital, Dhaka

³Professor, Pediatrics, Ad-din Women's Medical College Hospital, Dhaka

⁴Professor, Hepatology, Ad-din Women's Medical College Hospital, Dhaka

⁵Associate Professor, Medicine, Ad-din Women's Medical College Hospital, Dhaka

⁶Assistant Professor, Pediatrics, Ad-din Women's Medical College Hospital, Dhaka

***Corresponding author:** Richmond Ronald Gomes, Associate Professor, Medicine Ad-din Women's Medical College Hospital, Dhaka, Bangladesh.

Citation of this article: Z Ferdous Sonia, ARM Luthful Kabir, Md Hasan, A Ali, R Ronald Gomes, M Khan. (2026) "The uses of pentoxifylline in cardiology and hypertension." *International Journal of Clinical Case Reports and Clinical Reviews* 1(1). **Doi:** IJCCRCR.CR.26.004

Submitted: April 19, 2026

Accepted: April 24, 2026

Published: April 29, 2026

Abstract:

Choledochal cysts are dilatations of biliary tree that causes a diversity of clinical symptoms & lead to several complications. Choledochal cysts are most usually diagnosed in childhood & frequently present with abdominal pain, jaundice and abdominal mass. Albeit the most concerning complication is malignancy. Other complications are stone formation, acute pancreatitis. Stricture may also occur & can lead to morbidity. Treatment goal is not only relieving patient symptoms but also decreasing long term cancer risk. Here we present a young child presenting with abdominal pain & vomiting secondary to type 1b choledochal cyst complicated by acute pancreatitis with choledocholithiasis.

Keywords: choledochal cyst; acute pancreatitis; choledocholithiasis; malignancy

Introduction:

Choledochal cysts are rare congenital dilatation of extra hepatic & or intra hepatic bile ducts. The incidence of choledochal cyst in Asian population has been reported to be 1 in 13,000 and 1:1000 in Japan [1]. Female are more frequently affected than male with a ratio 4:1 [2]. Although choledochal cysts are predominantly diagnosed in children, about 25% of patient's diagnosis is delayed until adulthood [3]. Choledochal cyst causes a variety of clinical presentation & can lead to several types of complications. It can be diagnosed by chance or serendipitously due to complications such as acute pancreatitis [4,5], choledocholithiasis. Early diagnosis & treatment of choledochal cyst are crucial as they carry malignant potential that increases with age.

Here in, we present a child presenting with abdominal pain & vomiting secondary to type 1b choledochal cyst complicated by acute pancreatitis with choledocholithiasis.

Case Description:

An 18 month old female child presented at pediatric department with the complaints of eruptive vomiting for 3 days and abdominal pain for same duration. Pain appeared to be cramping & intermittent, relived by lapping.

On physical examination, body weight 9 kg, height 79 cm, (WAZ: on 3rd centile, WHZ: on 3rd centile), icteric, irritable. Patient had significant sensitivity in abdomen.

Laboratory data showed the following values: Hb-8.4 g/l, microcytic hypochromic anemia, WBC- $8.2 \times 10^3/\text{mL}$, S. bilirubin was 3.9mg/dl with dominant direct bilirubin, SGPT 349 U/L, Anti HAV IgM: negative. Serum amylase 306u/l, S. lipase 2270 U/L, electrolyte was normal. Abdominal USG showed-fusiform dilatation of CBD measuring about (4.1×1.7cm) cystic duct dilated. There were multiple granules mobile echogenic structure without shadow seen in lumen of the CBD. MRCP showed fusiform dilatation (2.2cm) of proximal CBD involving common hepatic duct over 3.5 cm segment (Figure 1). Distal CBD was mildly (1cm) dilated containing 2 tiny flow void suggesting calculus (0.5cm) seen with lumen of distal CBD (Figure 2). Pancreatic duct was visualized & mildly dilated.

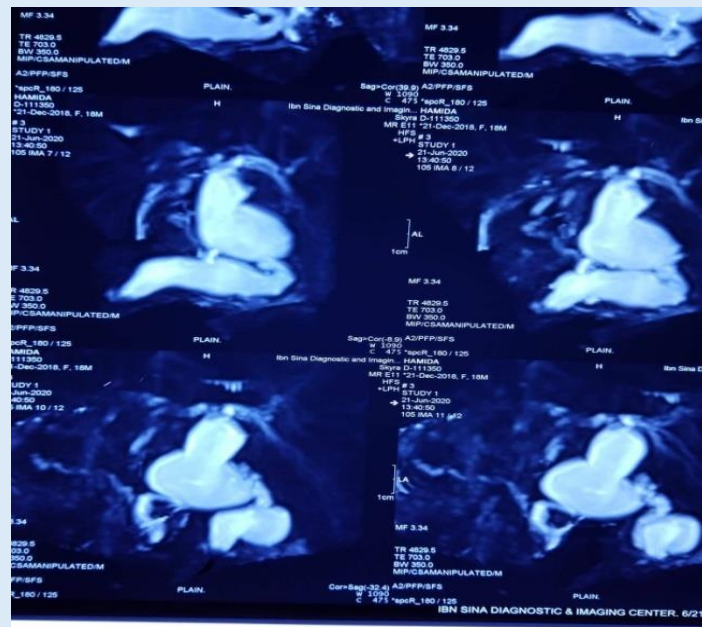


Figure 1: MRCP showing fusiform dilatation of proximal CBD involving common hepatic duct.

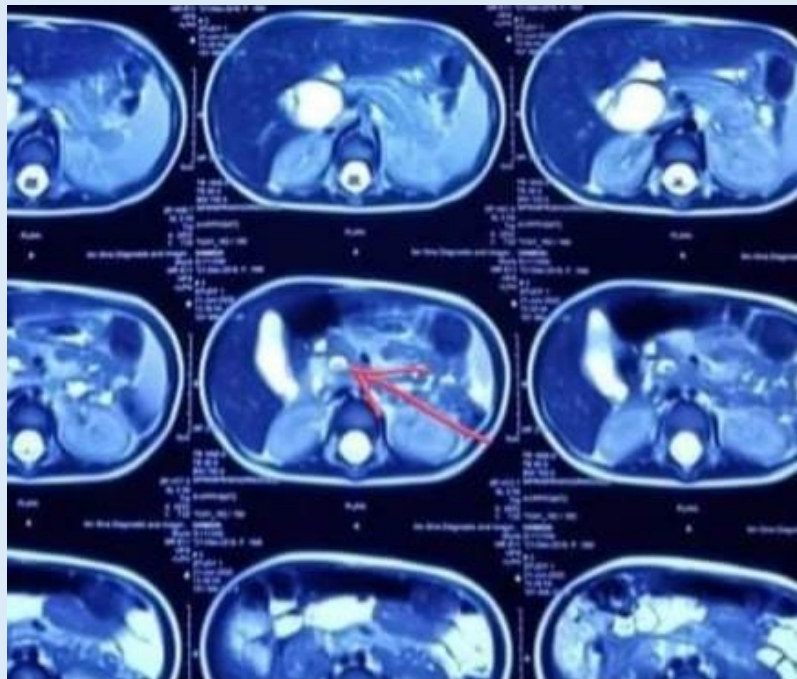


Figure 2: Distal CBD with signal void structure within suggestive of stones.

Thus the final diagnosis with acute pancreatitis with Choledocholithiasis caused by choledochal cyst type 1b was made. Fluid resuscitation was given initially due to profuse vomiting. Other treatment included parenteral antiemetic, proton pump inhibitor, opiate analgesic. After treatment, patient showed significant clinical improvement with decreased frequency of vomiting. Intensity of abdominal pain was also decreased. She was planned to undergo definite treatment by surgery. After complete resolution of 1st episode of pancreatitis the patient was operated 2 month later. The cyst was completely excised and cholecystectomy (Figure 3) and hepaticoduodenostomy were performed. Histopathology showed inflamed structure lined with biliary epithelium, free of any malignant lesion. The patient made an uneventful postoperative recovery.



Figure 3: Resected common bile duct cyst & gallbladder.

Discussion:

Vater and Ezler (1723) defined Choledochal cyst as a congenital dilatation of extra & / or intrahepatic bile duct. Pathophysiology remains still mysterious. The most widely accepted hypothesis is abnormal biliopancreatic junction [6]. Our patient has no such abnormality.

Anomalous Pancreaticobiliary Ductal Union (APBDU) allows reflux of pancreatic enzyme into biliary tree with consequent biliary duct inflammation predisposing to abnormal dilatation [7,8]. Some studies reported that sphincter of oddi dysfunction may predispose to Choledochal cyst [9]. Choledochal cyst may be part of congenital anomaly of biliary system or multiple congenital anomaly proposed by some other expert [10,11,12].

According to modified Todani classification (Figure). Choledochal cyst are classified into 5 types [13]. Clinical feature is variable & may depend on patient age. Frequently observed symptom in infant is jaundice and abdominal pain as in adult [14]. Classic triad consisting of jaundice, abdominal pain and palpable mass would only be found in 15 to 25 % case [15]. Our case, patient had jaundice and abdominal pain.

Potential complications of this malformation include acute pancreatitis, ascending cholangitis, cholecystitis, cholelithiasis, biliary cirrhosis and cholangiocarcinoma have been reported [7]. Choledochal cyst in adult are often associated with complication compared to children has been reported [16].

The occurrence of pancreatobiliary complications were to be brought by biliary stasis. This causes stone & sludge formation that lead to bile duct & pancreatic duct obstruction and resultant pancreatitis [18].

Acute pancreatitis is a common complaint of adult patient. Acute pancreatitis is rare occurrence in patient with type 1 choledochal cyst & just a couple of cases have been reported in the literature. Only 9% of pediatric patient with choledochal cyst were reported to have choledolithiasis [18]. In this patient diagnosis of pancreatitis was based on abdominal pain and high level of amylase. To our Knowledge the current patient is the youngest person with concomitant acute pancreatitis with choledocholithiasis which is rare.

Different imaging technique USG, MRI(MRCP), CT are available for the determination of choledochal cyst [19]. MRCP is considered the gold standard for diagnosis of biliary cyst with sensitivity 90-100%²⁰.

Appropriate management consists of conservative treatment of complications followed by definitive surgery [16,17]. Saluja et al found that cyst excision with biliary reconstruction is necessary to prevent the recurrence of complication [16]. Now a days, complete resection of the cyst with cholecystectomy & biliary reconstruction (RYHJ or HD) is considered the treatment of choice [21]. This operation was successful in our patient with no post operative complication.

There are small numbers of studies in the literature on the pancreatic complication of choledochal cyst in children. Fujishiro et al [22]. stated that pre-operative pancreatitis was most commonly seen in children with choledochal cyst type a.

To date, couples of studies have discussed the optimal timing of surgery for choledochal cyst complicated with acute pancreatitis in children. Cho et al proposed that early cyst excision after development of acute pancreatitis in patient with choledochal cyst reduced patient symptoms, especially occurrence of pseudopancreatitis [23]. However deciding the timing of surgery in choledochal cyst complicated with pancreatitis is challenging. Here we report our experience in treating type 1b choledochal cyst with acute pancreatitis with choledocholithiasis.

Conclusion:

Congenital cystic dilatation of biliary tract is a rare anomaly that may stay quiet or uncovered by acute pancreatitis. The diagnosis is based on USG & MRCP. Early diagnosis and prompt treatment is required to prevent long term complications. Acute pancreatitis is a uncommon occurrence in patient with type 1 choledochal cyst. Our patient with choledochal cyst associated with pancreatitis with choledocholithiasis was treated surgically with uneventful clinical outcome.

References

1. O'Neill JA Jr. Choledochal cyst. Current Problem in Surgery.1992;29 (6):361-410.
2. Wong-Hoi S, Chung HY, Lan LCL, Wong KKY, Saing H, et al.(2009). Management of choledochal cyst: 30 years of experience and results in a single center. Journal of Pediatric Surgery.44:2307-2311.
3. Liu CL, Fan ST, Lo CM, Lam CM, Poon RT, Wang J. Choledochal cysts in adults. Arch Surg 2002;137:465-468.
4. Ahmed A., Nazlae Z , et al. " Kyste du choledoque rompu:revue de la literature." [Cyst of the broken common bile duct: literature review].The Pan African Medical Journal 33(2019).
5. Kabbaj N.,et al. "Apropos of a case of acute pancreatitis revealing cystic dilatation of common bile duct."Journal de radiologie 79.11 (1998):1393-1397.
6. Ponce J, Garrigues V, Sale T, Pertejo V, Benenguer J. Endoscopic biliary manometry in patients with suspected sphincter of oddi dysfunction and in patients with cystic dilatation of bile ducts.Dig Dis Sci. 1989;34:367-71.
7. Bouali O., et al. "Pertontite biliaire par rupture traumatique d'un kyste du choledoque [Biliary peritonitis after traumatic rupture of a choledochal cyst]". Archives of pediatrics 22.7(2015):763-766.
8. Babbitt DP. Congenital choledochal cysts:new etiological concept based on anomalous relationships of common bile duct and pancreatic bulb. Ann Radiol 1969;12:231-40.
9. Babbitt DP, starshak RJ, Clemett AR. Choledochal cyst:a concept of etiology. Am J Roentgenol Radium Ther Nucl Med 1973;119:57-62.
10. Ponce J ,Ganrigues V Sale T, Pertejo V, Benenguer J. Endoscopic biliary manometry in patients with suspected sphincter of oddi dysfunction
11. Goldman M, Pranikoff T. Biliary disease in children. Curr Gastroenterol Rep 2011; 13: 193-201.
12. El Mouhadi S, Arrive L. Choledochal cyst.Gastroenterol Clin Biol 2010;34:347. Epub 2010 May 27.
13. Brancatelli G, Federle MP, Vilgrain V,etal. Fibropolycystic liver disease: CT and MR imaging findings Radiographics 2005; 25: 659-70.
14. Todani T, Watanabe Y, Toki A, Morotomi Y. Classification of congenital biliary cystic disease: special reference to type 1c and 1Va cysts with primary ductal stricture. J Hepatobil Pancreat Surg 2003;10:340-4.
15. FAIK M., et al. "Dilatation kystique du choledoque a propos de 8 cas." 75 (1999): 23-27.
16. Bouali O.,et al. "Biliary peritonitis after traumatic rupture of a cholrdochal cyst". Archives of pediatrics 22.7(2015): 763-766.
17. Saluja SS, Nayeem M, Sharma BC, Bora G, Mishra PK. Management of choledochal cysts and their complications. Am Surg 2012;78:284-90.
18. Singham J, Yoshida EM, Scudamore CH. Choledochal cysts: diagnosis. Can J Surg 2009;52:506-11.



19. Matsumoto Y, Uchida K, Nakase A, Honjo I. Congenital cystic dilatation of the common bile duct as a cause of primary bile duct stone. *Am J Surg*.1977;134:346-352.
20. Yu J, Turner MA, Fulcher As, Halvorsen RA. Congenital anomalies and normal variants of pancreaticobiliary tract and the pancreas in adults: part 1, Biliary tract. *AJR Am J Roentgenol* 2006;187:1536-1543.
21. Park DH, Kim MH, Lee SK, Lee SS, Choi JS, et al.(2005). Can MRCP replace the diagnostic role of ERCP for patients with choledochal cysts? *Gastrointest Endosc* 62: 360-366.
22. Patil V, Kanetkar V, Talpalkar MC. Hepaticoduodenostomy for biliary reconstruction after surgical resection of choledochal cyst: A 25 year experience. *Indian journal of surgery*.2015;77 (suppl 2): 240-244. DOI: 10.1007/s12262-012-0783-2 [Epub Dec 18, 2012].
23. Fuishiro J, Masumoto K, Urita Y, Shinkai T, Gotoh c. Pancreatic complications in pediatric choledochal cysts. *J pediater Surg* 2013;48:1897-902
24. Cho MJ, Kim DY, Kim SC, et al. Delayed surgery for choledochal cyst with acute pancreatitis in children. *Hepatogastroenterology* 2011;58:709-12.



©The Author(s) 2026. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) 4.0 License

Benefits of Publishing with us:

- ✓ fast; convenient online submission
- ✓ rigorous peer review by experienced research in your field
- ✓ rapid publication on acceptance
- ✓ authors retain copyrights
- ✓ unique DOI for all articles
- ✓ immediate; unrestricted online access

Click Here:

[Submit Manuscript](#)