A Rare Case of Biliary Atresia and Choledochal Cyst in a Premature Infant

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Abstract
Recent case reports have described variants in the classification of biliary atresia and choledochal cyst requiring the expansion of the classical system for describing these anomalies. This in effect, further obscures the line between these two previously separate entities while highlighting the difficulties of intraoperative differentiation of biliary atresia, cystic biliary atresia, choledochal cyst, and Caroli’s disease. In our case report, we describe a 34-week premature infant who developed a direct hyperbilirubinemia and was found to have an atypical presentation of choledochal cyst and biliary atresia. We believe this case demonstrates the difficulty in distinguishing and managing this disorder.

Keywords
Biliary atresia, Choledochal cyst, Caroli’s disease

Case Report
Our patient was a 34-week, 3.2 kilogram premature but otherwise healthy-appearing African American male infant with no significant family history. After a short hospital stay during which the patient was noted to be hyperbilirubinemic, he was discharged home on day of life (DOL) five. While at home the patient remained jaundiced and developed acholic stools. He was evaluated by Pediatric Gastroenterology and worked up with hepatobiliary iminodiacetic acid (HIDA) scan and ultrasound (Figure 1), which were consistent with choledochal cyst and possible biliary atresia. The patient was referred to Pediatric Surgery for definitive management.

Operative Approach
The patient was taken to the operating room on DOL 28. At exploration, the gallbladder was noted to be atretic with clear mucus. The liver was somewhat firm in texture and dark red-grey in color. There was no obvious cirrhosis or fibrosis. Liver biopsies were taken and sent for pathology. At the confluence of the cystic and hepatic ducts, a 2-3 cm cyst was noted with obliteration of the distal common bile duct as it extended posterior to the duodenum. Cholangiogram first performed through the gallbladder and then through the cyst did not demonstrate proximal filling of ductal structures. The cyst was mobilized and was sharply transected just below the bifurcation of the right and left hepatic ducts. Copious bile flow was noted from the proximal hepatic duct stump. The proximal hepatic stump appeared to be of normal ductal size measuring approximately 2-3 mm in transverse diameter. This tract was easily intubated and cholangiogram demonstrated abnormal, but patent right and left hepatic ductal system with radicals (Figure 2). Because the diagnosis of biliary atresia versus choledochal cyst had not been entirely eliminated, the decision was made to reconstruct the patient’s biliary outflow with a hepatocoduodenostomy in the event that it may be necessary to re-operate to perform a hepatopancreatoenterostomy [1]. A short segment of 3.5 mm feeding tube was used as a stent in the hepatic duct and a hepatocoduodenostomy created.

On histologic examination of the liver wedge biopsies taken at the initial operation, fibrosis in mainly a portal distribution was evident, and bile ductular proliferation could be seen in most of the portal tracts. The excess bile that accumulated secondary to downstream obstruction formed bile plugs in canaliculi and in the ductules. The resected cyst measured 1.5 × 1.2 × 0.8 cm with a wall thickness of 0.6 cm and a lumen devoid of contents (Figure 3A).

Postoperatively the patient did well. The patient had return of bowel function and tolerated advancing his diet. His intra-abdominal drain was discontinued. Total bilirubin however, remained elevated and steroids were started one week postoperatively in an effort to limit progressive fibrosis of the biliary tree [2,3]. His hyperbilirubinemia persisted. Given the persistent hyperbilirubinemia and pathologic findings consistent with biliary atresia, the decision was made to take the patient back to the operating room for a formal Kasai portoenterostomy.

Second Operative Course
The patient was brought back to the operating room on DOL 70.
His original incision was reentered and the hepaticoduodenostomy was taken down. The duodenotomy was closed transversely. Dissection was then carried down toward the porta hepatitis. A rather broad liver plate was identified and sharply incised at which point good bile flow was noted. A hepatoportoenterostomy was then created using a mobilized antecolic jejunal conduit. The liver biopsy performed at the time of the Kasai portoenterostomy demonstrated progression of the fibrosis, and the common right and left hepatic ducts resected were markedly narrowed, the central lumen obscured by fibrosis. Denudation of the epithelium proximally was followed in the distal region by focal absence of the lumen (Figure 3B).

The patient’s second postoperative course was protracted. While his bilirubin trended down gradually, he had slow return of bowel function and persistent abdominal distention. A HIDA scan demonstrated biliary drainage without evidence of anastomotic leak.

Several weeks after his second operation he was readmitted with fevers and feeding intolerance. On workup he was found to have new cavitory lesion consistent with abscess of the left hepatic lobe (Figure 4A). A percutaneous drain (Angiotech, Vancouver, BC) was placed and a study through the drain demonstrated continuity with the left hepatic duct (Figure 4B). This was therefore felt to be infected biliary abscess rather than bile lake formation. The patient subsequently developed an anastomotic stricture which was amendable to drainage catheter placement through the percutaneous drain (Cook Medical, Bloomington, IN). This drain was later internalized and upsized from 5-French to 6.5-French to maintain the patency of his anastomosis. After several weeks, the catheter was removed and drain study demonstrated patent intrahepatic ducts. All drains were therefore
removed. At the recommendation of gastroenterology, he continues to be on prophylactic Ursodeoxycholic acid and sulfamethoxazole/trimethoprim. He is now four years out from his last surgery. He is followed biannually by gastroenterology, and annually by pediatric surgery. He has had no episodes of jaundice, cholangitis, or changes in stooling pattern. His bilirubin levels have remained normal. He is at the 95th percentile for weight.

**Discussion**

Surgical correction is required in the treatment for pediatric biliary atresia and choledochal cyst. Age at initial operation has been shown to have the most significant effect on overall survival and subsequent need for liver transplant [4-6]. Recently, there have been multiple articles published, highlighting the difficulty in distinguishing between variants of biliary atresia and choledochal cyst in pediatric hepatobiliary surgery [4,5,7,8]. The adjunct of specific ultrasound findings have been described to help differentiate biliary atresia from choledochal cyst are supportive but not definitive [9]. Regardless, successful surgical management of this disorder requires accurate initial differentiation between choledochal cyst and biliary atresia. Attempts to establish radiographic diagnostic criteria have not
been highly successfully, and the mainstay of diagnosis still includes surgical exploration for pathology and cholangiography [4,10].

In our case report, the patient was diagnosed early in life with obstructive jaundice. However, as has previously been reported in other patients, imaging studies were inconclusive in differentiating between choledochal cyst and biliary atresia [10,11]. Despite this, our patient underwent operative management in a timely fashion. Liver parenchymal biopsies taken at the beginning of the operation were consistent with biliary atresia. Intraoperatively, a cystic structure was identified at the confluence of the cystic and hepatic ducts giving the appearance of a Type I choledochal cyst with distal atresia. This diagnosis was further reinforced whereupon transecting the proximal hepatic duct, good bile flow was noted and cholangiogram was able to demonstrate patent right and left hepatic ducts with radicals. Our patient failed reconstruction with hepaticoduodenostomy and eventually required formal portoenterostomy as he had progression of fibrosis. Kasai portoenterostomy remains first-choice procedure. In the long term, patients with biliary atresia may go on to require liver transplantation, even into adulthood [12].

Conclusions

Despite good evidence to support the diagnosis of Type I choledochal cyst with distal atresia, our patient failed reconstruction with hepatoduodenostomy and eventually required a second operation with formal Kasai portoenterostomy [13]. We believe that this case highlights the difficulties in making the diagnosis in the setting of congenital anomalies of the pediatric biliary tree and underscores the importance of surgical exploration and pathologic diagnosis. It is unclear whether the patient’s phenotype was a variant in the spectrum of biliary atresia and choledochal cyst or whether this infant’s prematurity created an evolution of postnatal progressive fibrosis, which ultimately led to failure of his initial reconstruction. Recent case reports have described the caveats of diagnosing biliary atresia in the setting of choledochal cyst, which correlate with our report [11,14,15]. Despite postoperative setbacks following his second operation, the patient is doing well and liver functions have stabilized near normal at four years of age.

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Disclosures

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References