When It Doesn’t Fit: Congenital Anomalies of the Choledochus

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Keywords
► choledochal cyst
► congenital choledochal malformation
► hepaticojejunostomy
► epidermoid cyst

Abstract

Introduction Congenital choledochal malformations (CCMs) are characterized by intra- and/or extrahepatic bile duct dilatation. Five basic types (1–5) are recognized in Todani’s classification and its modifications, of which types 1 and 4 typically have an associated anomalous pancreatobiliary junction and common channel (CC). We describe two cases with previously undescribed features.

Case Report 1 Antenatal detection of a cyst at porta hepatis was made in an otherwise normal girl of Iranian parentage. She was confirmed to be a CCM (20 mm diameter), postnataally, with no evidence of obstruction. Surgical exploration was performed at 12 weeks. She had an isolated cystic dilatation of the right-hepatic duct only. The left-hepatic duct and common bile duct (CBD) were normal without a CC. Histology of the resected specimen showed stratified squamous epithelium.

Case Report 2 A preterm (31 weeks of gestation) boy of Nigerian parentage was presented. His mother was HIV+ve and he was treated with nucleoside reverse transcriptase inhibitors following birth. He had persistent cholestatic jaundice and a dilated (10 mm) bile duct from birth. Although the jaundice resolved, the dilatation persisted and increased, coming to surgery aged 2.5 years. This showed cystic dilatation confined to the common hepatic duct, and otherwise normal distal common bile duct and no CC.

Result Both underwent resection with the Roux-en-Y hepaticojejunostomy reconstruction to the transected right-hepatic duct alone in case 1, leaving the preserved left duct and CBD in continuity, and to the transected common hepatic duct in case 2.

Conclusions Neither choledochal anomaly fitted into the usual choledochal classification and case 1 appears unique in the literature.

New Insights and the Importance for the Pediatric Surgeon

We describe two variants of choledochal malformations that do not fit the standard classifications. One case appears unique consisting of a unilateral cystic dilatation of the right-hepatic duct, surgically resected and anastomosed to a jejunal loop with preservation of the left-sided biliary drainage. The cyst was lined with nonkeratinising stratified squamous epithelium which is extremely rare in choledochal malformations.
Introduction

Congenital choledochal malformation (CCM) simply implies dilatation of the extra- and/or intrahepatic biliary tract in the absence of obstruction. The actual pathogenesis is not known for certainty, although most seem to be related to a degree of distal stenosis leading to proximal dilatation.\(^1\)

We have used our own Kings’ College Hospital classification, which is a simplification of Todani’s classification, a modification of previous classifications. The commonest are characterized as a cystic extrahepatic dilatation (type 1c), a fusiform extrahepatic dilatation (type 1f), and the combination of either with intrahepatic duct dilatation (type 4). In most of them, there is also an accompanying pancreaticobiliary junction malformation and long common channel (CC) which has the potential for free intermixing of bile and pancreatic juice.

We describe two patients with atypical morphology that did not conform to any of the usual descriptions.

Case Reports

Case 1: the first case concerns a female infant of Iranian parentage born at term. Diagnosis of a cyst at the porta hepatis was made on an antenatal scan just before birth after arrival in the United Kingdom. Previous scans from abroad were not available. She developed jaundice with a maximum total bilirubin of 380 µmol/L that settled spontaneously. Alpha feto-protein (AFP) level was 756 kIU/L. Ultrasound and subsequent magnetic resonance cholangiopancreatography (MRCP) showed a cyst with dimensions 20 mm \(\times\) 16 mm at the porta hepatis, without evidence of proximal intrahepatic biliary dilatation.

Surgical exploration and cholangiography was performed at 12 weeks and showed cystic dilatation confined to the right-hepatic duct with an entirely normal left-hepatic duct draped around and with a low insertion of a cystic duct into a normal-looking common bile duct (CBD; \(\Rightarrow\) Fig. 1A). Intraductal pressure was measured at 5 mm Hg (normal). There was no evidence of a CC on the cholangiogram. Bile amylase was 4 IU/L and CA19-9 was 89,400 kU/L.

Following cholecystectomy, the cyst was detached from the CBD and the transected proximal part with its draining segmental ducts anastomosed to a Roux’s loop (\(\Rightarrow\) Fig. 1B). The left duct and CBD were left intact. Histopathology showed a cyst lined by nonkeratinising stratified squamous epithelium. Follow-up up to 1.5 years had been unremarkable.

Case 2: The second case was a boy of Nigerian origin, born prematurely at 31 weeks, who was presented with transient conjugated jaundice and was found to have a dilatation (10 mm) of the proximal CBD. His mother was HIV + ve (low-risk retrovirus positive), and he was treated with a nucleoside reverse transcriptase inhibitor since birth. At birth, he had positive anti-HIV antibodies but DNA was not detectable. Images from the preoperative MRCP and ERCP are shown in \(\Rightarrow\) Fig. 2.

Jaundice resolved but the dilatation persisted and increased over time, so he underwent surgical exploration at the age of 2.6 years. This showed a cystic dilatation extending proximally from the junction of cystic duct to the common hepatic bile duct (CHD). Intracystic pressure was measured at 20 mm Hg (normal \(<\) 5 mm Hg). The remaining distal CBD appeared normal and there was no evidence of a CC. Bile amylase was 5 IU/L and CA 19.9 was 9,600 kU/L. Both cyst and gallbladder were resected and a hepaticojejunostomy en-Roux performed. Histopathology showed a dilated bile duct lined by columnar type biliary epithelium with mural fibrosis but with no significant inflammation.

He has had 2 years of follow-up with normal serial ultrasound scans.

Discussion

CCMs are rare with most infants and children (> 80%) presenting as types 1c, 1f, or 4. Most cases of extrahepatic dilatation are stereotypical with two basic morphologies. The classical choledochal cyst (type 1c) involves the entire CBD, and CHD ending abruptly in what can be a filamentous connection with the pancreatic duct. Alternatively, the dilatation appears fusiform, of smaller diameter, with a more gradual termination and junction with the pancreatic duct.

Unilateral cystic dilatation of the hepatic duct is exceptional with perhaps only two reported cases in the literature. Prekop et al reported a 14-month-old boy with a large “cystic
malformation of the right-hepatic duct,” with little more clinical
detail including nature of epithelial lining. Gidi et al reported a
56-year-old woman with cystic dilatation of the right-hepatic
duct who underwent a resection of the cyst with right hepatec-
tomy to achieve complete resection. Histopathology was similar
to our own, although it was described as extensive squamous
metaplasia with areas of columnar epithelium rather than de
novo stratified squamous epithelium characteristic of ours. As
normal bile duct consists of vascularized, innervated fibrous
tissue, lined by a single layer of tall columnar epithelium, we feel
that ours is likely to be congenital rather than acquired
metaplasia.

The lining and situation in the porta hepatis in our first
case do appear similar to that of the squamous-lined cystic
dilatation described by Chiu et al in a 6-month-old boy. This
malformation was antenatally detected and, at operation,
found to arise from the common hepatic duct with
separate entry of both right- and left-hepatic ducts. Finally,
Kwon et al described a diverticulum of the bile duct (type 2
CCM) with stratified squamous epithelial lining in a 63-
year-old man. Metaplastic squamous lining often associ-
ated with recurrent infection and stones can lead to
malignant change. Certainly, the age of our child implies
a congenital rather than metaplastic origin for the squa-
mosus nature of the epithelium.

The second case, while nominally a type 1c, is distinctly
unusual with sparing of the distal duct and no CC. The high-
intracystic pressure of 20 mm Hg suggests an obstructive
etiology, and there was no suggestion of extrinsic compression
as might be found with a healed spontaneous perforation. It is
interesting to note that this child was delivered from an
HIV + ve mother, though it is difficult to find real evidence
of cause and effect.

We have previously reported a series of CCM where we
measured CA 19.9 in bile to try and identify a cohort who
might have particular predisposition to dysplasia and per-
haps later malignancy. We found that it was invariably
raised and often in very-high concentration but bore no
relationship with simultaneous obtained bile amylase and
choledochal pressure. Indeed, it could be localized on stain-
ing to otherwise normal biliary epithelium. CA 19.9 levels in
both cases were raised, although whether that represents
normal biliary tract epithelium elsewhere or from the cyst is
not known.

Surgical reconstruction was bespoke in case 1 with preser-
vation of the left-sided biliary drainage but more typical in
case 2. Primary duct to duct anastomosis seemed possible in
that case but was rejected, as it was felt to be less safe than a
standard Roux’s loop.

Conclusion
In conclusion, neither case fits comfortably into the standard
CCM classifications and both present different perspectives
on possible etiological factors.

Conflict of interest
None.

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