Solitary cystic dilatation of the intrahepatic bile duct

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Abstract
A 31 year old man was hospitalised with general fatigue and epigastric pain. Abdominal ultrasonography, computed tomography, and magnetic resonance imaging showed a cystic lesion in the left lobe of the liver. Endoscopic retrograde cholangiopancreatography and percutaneous transhepatic cholangiography revealed a localised dilatation of the intrahepatic bile duct without any obstruction. However, a large mass of mucinous material was noted in the saccular intrahepatic duct and the common bile duct. There was no evidence of a choledochal cyst, anomalous pancreaticobiliary ductal union, or congenital cystic change of the kidneys. A possible diagnosis of mucinous cystic neoplasm of the intrahepatic bile duct was made and a left hepatectomy performed. Cholangiography of the resected specimen showed a non-obstructive solitary cystic dilatation of the left hepatic duct. Histologically, the dilated duct was lined with columnar biliary epithelium without any papillary proliferation and/or atypia. Neither malignancy nor hepatic fibrosis was observed; the term “solitary cystic dilatation of the intrahepatic bile duct” perhaps is more descriptive and the concept easier to understand.

Keywords: solitary cystic dilatation; intrahepatic bile duct; terminology

Recently, Terada and Nakanuma described two cases of solitary cystic dilatation of the intrahepatic bile duct without choledochal cysts, anomalous pancreaticobiliary ductal union, or congenital cystic change in the kidneys. To date, only 13 such cases have been reported. Several other terms have been used in the literature with regard to localised or non-obstructive dilatation of the intrahepatic bile duct. These include the localised form of Caroli’s disease, localised dilatation of a major intrahepatic duct, and a type V congenital bile duct cyst.

We report a case of solitary cystic dilatation of the intrahepatic bile duct as well as a review of the literature, and discuss the terminology concerning this entity.

Case report
In April 1995, a 31 year old man was admitted to Kawasaki Medical School Hospital because of general fatigue and epigastric pain. He presented with neither a pertinent family history nor a personal history of blood transfusion, tattooing, or drug abuse. On physical examination, there was a slight tenderness in the epigastrum and right hypochondrium, and the liver was slightly enlarged. Results of liver function tests were: total bilirubin, 2.1 mg/l (normal value 0.2–1.0); alkaline phosphatase, 127 IU/l (28–84); γ glutamyltranspeptidase 314 IU/l (4–30); aspartate aminotransferase 117 IU/l (7–20); and alanine aminotransferase 425 IU/l (7–28). Virus markers were negative for hepatitis A, B, and C. Abdominal ultrasonography, computed tomography, and magnetic resonance imaging showed a cystic lesion in the left lobe of the liver. Choledochal cysts and anomalous pancreaticobiliary ductal union were not demonstrated on endoscopic retrograde cholangiopancreatography. Percutaneous transhepatic cholangiography revealed a localised dilatation of the left intrahepatic bile duct. There was a large mass of mucinous material in the saccular intrahepatic bile ducts and the common bile duct. Under the diagnosis of a possible mucinous cystic neoplasm of the intrahepatic bile duct, we performed a left hepatectomy. A cholangiogram of the resected specimen showed a solitary cystic dilatation of the left hepatic duct (fig 1). Histologically, the
The term type V congenital bile duct cyst was proposed by Todani et al in 1977. Type V is an intrahepatic bile duct cyst and it may be single or multiple (our case was single). Todani et al suggested that type V might be closely related to intrahepatic stones and could be categorised as a pure form of Caroli’s disease without hepatic fibrosis.

Finally, solitary cystic dilatation of the intrahepatic bile duct is a term that was proposed by Terada and Nakamura in 1987. Although this is a rather descriptive term, this entity has some clinicopathological features such as female predominance and cystic dilatation involving the right or left hepatic duct near the hepatic hilus. In addition, there is no evidence of related conditions including choledochal cysts, anomalous pancreaticobiliary ductal union, or congenital cystic change in the kidney. As Terada and Nakamura pointed out, bile stasis and mucous hypersecretion are important for dilatation of the intrahepatic bile duct, our case may be explained by mucous hypersecretion as a large mass of mucinous material was seen in the intrahepatic duct.

To date, the above mentioned terms have been used for the same localised or non-obstructive dilated lesion of the intrahepatic bile duct. We prefer to use the last term—solitary cystic dilatation of the intrahepatic bile duct—because it is descriptive and the concept is easy to understand. Furthermore, it has clinicopathological features. Although this lesion is very rare and under-recognised, it should be considered by both pathologists and clinicians.