A case of hepatic intraductal papillary neoplasm of the bile duct

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Abstract

Introduction: Intraductal Papillary Neoplasm of the Bile ducts (IPNB) is a rare entity characterized by exophytic growth of the bile ducts.

Case presentation: In this report, we present a 57-year-old male with no prior medical history consulted for upper right abdominal pain, jaundice and pruritus. Abdominal Ultrasound (US) and magnetic resonance imaging/Biliary magnetic resonance were performed. They revealed that the circumferential parietal thickening of the common hepatic duct had extended approximately 4 cm with moderate dilatation of the left intrahepatic bile ducts. Computed tomography showed no evidence of distant metastasis.

Biopsy revealed a high-grade intraductal papillary neoplasm. After 40 days, the patient had left hepatectomy with resection of the main bile duct and the gallbladder.

Macroscopic examination of the surgical specimen showed a dilatation duct at the hilum with thickening of their walls.

The histopathology report revealed multiple intra-hepatic papillary neoplasms with high-grade dysplasia with an invasive carcinoma component in the left hepatic duct without extending to the biliary wall, classified as pT1N0.

Conclusion: This premalignant lesion has the potential to transform into invasive carcinoma if not properly diagnosed.

Our case illustrates how early identification can lead to potential surgical resection.

Introduction

Bile duct neoplasms are a diverse group of benign and malignant tumors that can develop anywhere along the biliary tree [1].

Traditionally, the term biliary papillomatosis has been used to describe both imaging and pathology features. The World Health Organization adopted the term intraductal papillary neoplasm in 2010 [2].

In 2019, the World Health Organization defined intraductal papillary neoplasm of the bile ducts (IPNB) as “a grossly visible premalignant neoplasm characterized by intraductal papillary or villous growth of biliary-type epithelium” [3]. As a result, IPNB is a precancerous lesion. It is distinguished by an intraluminal papillary tumor in the dilated bile duct with a fibrovascular core [4]. However, the malignant transformation rate of IPNB has been reported to be as high as 41% - 83% [5].

Because precancerous lesions take 1 - 2 years to transform into invasive carcinoma, surgical treatment is feasible [6].

We report the case of a patient who consulted for upper right abdominal pain and was diagnosed with IPNB, with an invasive carcinoma component.

Clinical presentation

A 57-year-old male with no prior medical history was consulted for upper right abdominal pain, jaundice and pruritus. His initial laboratory values were as follows:

- Total bilirubin (tbil): 139 mg/dl
- Direct bilirubin (dbil): 74.8 mg/dl
- Aspartate aminotransferase (AST): 300 IU/L
- Alanine aminotransferase (ALT): 135 IU/L

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had extended approximately 4cm with moderate dilatation of the left intrahepatic bile ducts (Figures 1 and 2). Computed tomography showed no evidence of distant metastasis.

Biopsy revealed a high-grade intraductal papillary neoplasm. After 40 days, the patient had left hepatectomy with resection of the main bile duct and the gallbladder. A manual biliary digestive anastomosis was performed. The surgical exploration showed the absence of ascites, carcinosis, and liver metastasis.

Macroscopic examination of the surgical specimen showed a dilatation duct at the hilum with thickening of their walls (Figures 3 and 4).

The histopathology report revealed multiple intra-hepatic papillary neoplasms with high-grade dysplasia with an invasive carcinoma component in the left hepatic duct without extending to the biliary wall, classified as pT1N0 (Figures 5 and 6).

The postoperative course was uneventful.
Discussion

The diagnosis of hepatic cystic masses is increasing as a result of the rapid advancements in imaging. In fact, it has been estimated that 20% of the general population has cystic masses in the liver [7]. While the majority of these masses are benign cysts, 3% – 5% of them are biliary cystic malignant tumors. The most common bile duct cystic tumors are IPNB and pancreatic mucinous cystic neoplasms (MCN). IPNB was found in 10% of all resected bile duct tumors [8].

The incidence of IPNB varies geographically, accounting for 7% - 10% of all bile duct tumors in Europe and North America and 10% - 40% in Asian cohorts [9]. IPNB patients are mostly men between the ages of 50 and 70, with recurrent and intermittent pain and cholangitis. IPNB is uncommon in children, and there is no evidence of a tendency for familial aggregation of cases [10,11].

The pathogenesis of IPNB is not clear, but it may be related to repeated infection of the biliary tract and cholestasis. However, bile duct stones, \textit{Clonorchis sinensis} infection, primary sclerosing cholangitis and biliary malformation are considered risk factors for IPNB.

IPNB development and progression from low, high-grade dysplasia to invasive carcinoma follow a sequential acquisition of molecular alterations involving common oncogenic pathways, including \textit{KRAS}, \textit{CDKN2A} and \textit{TP53} genes [12].

IPNB patients frequently present with recurrent episodes of abdominal colic, intermittent obstructive jaundice and acute cholangitis.

Medical imaging doctors have an important role in the diagnosis of IPNB. They should be aware of the importance of capturing image findings of IPNB because this disease has a high potential for malignancy. IPNB may be radiologically undetectable, but in most cases, cholangiography shows filling defects in the biliary tree, caused by an intraductal mass, with dilatation of either proximal or distal bile ducts.

This diagnosis should always be considered in the presence of focal duct dilatation filled with fluid, mucin, or soft tissue, in both symptomatic and asymptomatic patients.

The prevalent location of IPNB (intrahepatic versus extrahepatic bile ducts) is highly variable among studies [10].

Surgery is considered the cornerstone of treatment, with partial hepatectomy (for localized disease) and liver transplantation along with pancreaticoduodenectomy (for extensive disease or recurrent IPBN) being the therapeutic options.

In this situation, the patient had a left hepatectomy with resection of the main bile duct and the gallbladder. A manual biliary digestive anastomosis was performed.

On macroscopic examination, IPNB appears as a papillary, villous, or polypoid, red-colored, soft mass, predominantly growing in a dilated bile duct lumen, which may be cystic or fusiform. Lesions might be single and isolated or multiple and mucus hypersecretions can be seen. An invasive component can appear as a nodule lesion [9].

On histological examination, IPNB is characterized by dilated bile ducts filled with papillary or villous structures with fine fibrovascular cores covered by biliary epithelial cells. Tubular or glandular components can be seen but not an ovarian-type mesenchymal stroma. Epithelial cells are cuboidal or columnar and may show intestinal, biliary, oncocytic and gastric-type differentiation, based on morphology and immunophenotype [10]. The biliary subtype is the most common in Western countries, while the oncocytic and gastric types are very rare. Mixtures of different subtypes are observed in about 50% of all IPNBs, so their classification is based on the most prevalent component [13].

Epithelial cells lining IPNB may show variable cytoarchitectural atypia, classified into low or high-grade dysplastic lesions. IPNBs with high-grade dysplasia are more frequent in extrahepatic bile ducts.

The extension of the neoplastic process into peribiliary glands, particularly in the hilar zone, is quite common.

Regarding our patient, he was diagnosed with IPNB with high-grade dysplasia to the left extrahepatic duct with an invasive carcinoma component.

Complete resection is a significant prognostic factor, reported in great numbers. The presence or absence of metastases to lymph nodes is a major prognostic factor along with complete resection [14].

IPNB has been known to recur in the bile duct following surgical resection. Despite that the initial Resection stump of the bile duct tested negative, intraductal papillary tumor presenting with tissue image similar to the initial resection recurred. IPNB recurs after more than 10 years in some cases, appearing as if it were new carcinoma. This suggests a multicentric growth of IPNB [15]. As a result, clinical course monitoring is required, with postoperative intraductal recurrence in mind.

Conclusion

IPNB is a relatively rare disease that should be considered in the differential diagnosis of hepatic cystic tumors.

IPNB can present diagnostic challenges. Each imaging modality has limitations in detecting IPNB because of its small tumor size, tumor multiplicity and tendency to spread superficially. The mainstay of treatment of IPNB is surgical resection, so an accurate preoperative evaluation of tumor extent is crucial. Thus, a multi-modality approach is needed.
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to diagnose IPNB and establish an optimal treatment plan for these patients. This premalignant condition has the potential to transform into invasive carcinoma if not properly diagnosed.

Our case illustrates how early identification can lead to potential surgical resection.

References