Complete agenesis of dorsal pancreas with pancreatic cyst: A case report

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ABSTRACT
Agenesis of the dorsal pancreas (ADP) is extremely rare disease with no specific symptoms and there is no clear pathogenesis. Approximately half of the affected individuals develop diabetes resulting from reduced islet cell mass secondary to lack of endocrine structures. In this case, we aimed to present a 17-year-old female patient with ADP accompanied by a pancreatic cyst.

Keywords: Agenesis of dorsal pancreas; magnetic resonance imaging; pancreatic cyst.

CASE REPORT
A 17-year-old female patient with resistant hypomagnesemia and insulin-dependent diabetes mellitus presented with intermittent epigastric pain for several months. The pain was not associated with fever or vomiting. Her laboratory tests revealed: fasting plasma glucose: 411 mg/dl, glycated hemoglobin/HbA1c: 7.8%, magnesium level: 1.02 mg/dl, and cholesterol: 222 mg/dl. Amylase and lipase levels were normal.

The patient was referred to the radiology department for abdominal ultrasound (US) examination. US revealed a cystic lesion on the head of pancreas (Fig. 1), whereas body and tail of pancreas could not be visualized due to bowel gas shadow. A contrast-enhanced abdominal computed tomography (CT) was performed for better evaluation of pancreas and cystic lesion. CT examination demonstrated the absence of neck, body, and tail of the pancreas (Fig. 2). Only the head and uncinate segment of the pancreas was visualized and the hypodense unilocular cystic lesion was revealed at the head of pancreas (Fig. 3). Furthermore, an accessory spleen was spotted. Abdominal magnetic resonance imaging (MRI) was ordered to evaluate the nature of the cyst. Contrast (gadolinium diethylene triamine pentaacetic acid) enhanced abdominal MRI with routine sequences revealed no contrast enhancement on the cyst and its wall. The cyst was hypointense in T1-weighted imaging and hyperintense in T2-weighted imaging (Fig. 4). These findings were compatible with dorsal pancreatic agenesis with pancreatic cyst.

DISCUSSION
Dorsal pancreatic agenesis is a rare congenital anomaly associated with abnormal embryogenesis [2]. The pancreas develops from ventral and dorsal endodermal outpouching of the duodenum. At the 6–7th week of gestation, the ventral and the dorsal parts of the pancreas fuses and then both of these buds form the main pancreatic duct. During the 7th week of gestation, the ven-
tral bud turns posteriorly to connect with the dorsal bud behind the duodenum to form the mature pancreatic gland [3, 4]. Developmental failures caused by abnormal embryogenesis may lead to partial or complete ADP [5]. In complete dorsal pancreatic agenesis, the body and the tail of the pancreas, the duct of Santorini, and the minor papilla are totally absent, whereas in partial agenesis, the minor papilla, the remnant of the accessory duct, and a small portion of the body of the pancreas can be seen [6].

The first case of dorsal pancreatic agenesis was reported as an autopsy finding in 1911 [7] and there have been around 100 cases published in the literature until today.

Most ADP patients are usually asymptomatic. In symptomatic patients, abdominal pain is the most common symptom. Half of the ADP patients may present with hyperglycemia due to the involvement and underlying diabetes mellitus [8, 9]. Acute and chronic pancreatitis may also be seen in ADP patients due to Oddi sphincter dysfunction, enzyme hypersecretion, and higher pancreatic duct pressures [9, 10]. Chronic pancreatitis can lead to pseudocyst formations [11]. Our patient did not have any symptoms which can be associated with acute or chronic pancreatitis.

Despite its extremely rare occurrence, it is crucial to distinguish solid pseudopapillary neoplasms (SPNs) combined with ADP from cyst and pseudocyst forma-
tions. Because SPN may appear cystic due to its necrotic areas [12–14]. In our case, there were no solid component or calcification in the pancreatic cyst that may cause suspicion of SPN.

Other developmental abnormalities may accompany ADP such as polysplenia syndrome, heterotaxy, ectopic spleen, bowel malrotation, coarctation of the aorta, trilalogy of Fallot, and atrioventricular valvular abnormalities [15–17].

Pancreatic carcinoma with proximal atrophy, autodigestion from chronic pancreatitis, pancreatic divisum, pancreatic masses, distal pancreatic lipomatosis, and pseudoagenesis can mimic ADP [18, 19]. Chronic pancreatitis may cause replacement of the pancreas by fat and it may lead to atrophy of the pancreatic body and tail which is called “pseudoagenesis.”

There are several useful modalities for the diagnosis of ADP. Initially, US is the most commonly used diagnostic modality for abdominal pain and other abdominal symptoms. On US, head of pancreas appears as a small hypoechoic structure in young population. At the junction of the head and neck of the pancreas, a hyperechoic line separates the hypoechoic pancreatic head from the more echogenic retroperitoneal fat. Nevertheless, US is often not adequate to visualize pancreas. Contrast-enhanced abdominal CT and MRI of the abdomen are better methods for ADP diagnosis. On CT study of abdomen, in the absence of pancreatic corpus and tail, small intestine or stomach can be seen within the distal pancreas area (dependent stomach or dependent intestine signs) [20]. MRI including Magnetic resonance cholangiopancreatography (MRCP) is a non-invasive method that can reveal the pancreatic ductal anatomy and confirm the absence of the dorsal duct system.

Treatment is usually symptomatic if there is not any suspicion for malignancy in pancreas.

Conclusion

ADP, although rare, sometimes could constitute a diagnostic challenge in routine practice. In our case, ADP was diagnosed evaluation of the patient for abdominal pain.

Informed Consent: Written, informed consent was obtained from the patient’s family for the publication of this case report and the accompanying images.

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