Pancreas divisum: magnetic resonance cholangiopancreatography findings

ABSTRACT
Pancreas divisum is a common congenital abnormality of the pancreas that results from the lack of fusion between the dorsal and ventral pancreatic ducts during foetal development. In these cases, the dorsal duct becomes the main pancreatic duct and drains most of the pancreas. Pancreas divisum is mainly asymptomatic, but the prevalence of pancreas divisum is higher in patients with chronic abdominal pain and idiopathic pancreatitis. A study of 20 patients with pancreas divisum (12 men and eight women; aged 19–77 years; mean age 39 years) and who underwent magnetic resonance cholangiopancreatography (MRCP), was performed. In our series, pancreas divisum was clinically manifested as unexplained episodes of abdominal pain (mean duration 3.2 years) (60 percent), mild pancreatitis (30 percent) or incidentally (ten percent). MRCP demonstrated non-communicating dorsal and ventral ducts, independent drainage sites, a dominant dorsal pancreatic duct, and a small cystic dilatation of the dorsal duct at minor papilla (santorinocoele). In this pictorial essay, we review the most common MRCP features of pancreas divisum.

Keywords: magnetic resonance cholangiopancreatography, pancreas divisum, pancreatic anomaly, pancreatitis

INTRODUCTION
Pancreas divisum affects 5%–10% of the population and is considered to be a congenital anomaly of pancreatic ductal configuration. Embryologically, the ventral pancreatic bud and biliary system arise from the hepatic diverticulum, and the dorsal pancreatic bud arises from the dorsal mesogastrium. Under normal conditions, as the ventral bud rotates around the foregut, there is fusion of the dorsal pancreas and ventral pancreas, and the ventral pancreatic duct fuses with the dorsal pancreatic duct and form the duct of Wirsung. Drainage of a normal pancreas normally occurs through the ventral duct of Wirsung, which joins the common bile duct at the level of the major papilla. The dorsal duct often remains as duct of Santorini, and empties into the duodenum through the minor papilla. When the dorsal and ventral pancreatic ducts fail to fuse, pancreas divisum occurs; this means that most of the glandular parenchyma is drained by the dorsal duct through the minor papilla. The smaller ventral duct, however, drains some of the pancreatic head through the major papilla, including the uncinate process.

Although most people with pancreas divisum have no clinical disease, it has been suggested that pancreas...
Pancreas divisum may cause unexplained abdominal pain, recurrent episodes of acute pancreatitis or mild chronic pancreatitis, due to the presence of relative obstruction to the drainage of pancreatic secretions at the minor papilla. The inherently small diameter of the minor papilla causes increased pressures in the dorsal pancreatic duct. Pancreas divisum is found in 15%–20% of patients with unexplained pancreatitis.

Endoscopic retrograde pancreatography can be used to make a definitive diagnosis of pancreas divisum. If standard cannulation of the major papilla is performed, endoscopic retrograde pancreatography reveals only ventral duct opacification. The filling of the dominant dorsal duct can be achieved by cannulation of the minor papilla.

Magnetic resonance cholangiopancreatography (MRCP) is a non-invasive diagnostic technique that depicts the pancreatic ducts without injection of iotinated contrast material and has been shown to be highly sensitive and specific for pancreas divisum. Non-communication of the dorsal and central ducts, independent drainage sites and a dominant dorsal pancreatic duct can be seen using MRCP. Typically, the ventral duct is short and extremely narrow, while the dorsal duct is normally larger in calibre. Depiction of the pancreatic ducts can be improved using dynamic MRCP of the pancreatic duct after secretin stimulation. Secretin MRCP can improve the detection and classification of pancreas divisum. Abnormal persistent poststimulatory dilatation, a sign of ampullary dysfunction, can be shown with MRCP performed before and after secretin injection. Abdominal pain and recurrent pancreatitis associated with pancreas divisum is normally treated with minor papilla sphincterotomy.
CLINICAL FEATURES

Our series included 20 patients (12 men and eight women; age range 19–77 years; mean age 39 years). 12 of them presented with unexplained abdominal pain without hyperamylasaemia (four with first episode, eight with recurrent episodes, mean duration 3.2 years). Six patients presented with pancreatitis (three with first episode, three with recurrent episodes, mean 2.4 episodes) and two patients presented with symptomatic cholelithiasis.

The details of clinical history of some patients included:

- A 56-year-old patient with symptomatic cholelithiasis without hyperamylasaemia (Fig. 1).
- A 22-year-old man who presented with first episode of unexplained abdominal pain without hyperamylasaemia (Fig. 2).
- A 31-year-old man with abdominal pain (Fig 3).
- A 36-year-old woman with her fourth episode of mild pancreatitis (Fig. 4).
- A 24-year-old woman with her first episode of pancreatitis and incomplete pancreas divisum (Fig. 5).
- A 23-year-old man with episodes of unexplained pain (Fig. 6).
- A 54-year-old man with recurrent episodes of unexplained abdominal pain (Fig. 7). This patient refused surgery.

IMAGING FINDINGS

Pancreas divisum was detected in 15 patients. In 12 of those patients, this was the sole abnormality seen; whereas in the other three patients, the pancreas divisum was associated with the presence of small santorinocoele (small cystic dilatation of the dorsal duct). Incomplete pancreas divisum was depicted on MRCP images in two patients; one of them associated with santorinocoele. Other findings included two cases associated with dilatation of dorsal duct and side branches, and one case associated with significant focal dilatation of the duct with septa, probably intraductal papillary mucinous tumour (IPMT).

On SS-TSE images, the dorsal duct drained to the minor papilla, independent of common bile duct (Figs. 1a & 2). On axial HASTE images, the course of the duct was
horizontal in front of the common bile duct towards the minor papilla (Fig. 1b). Most cases of pancreas divisum in our series showed a normal diameter main duct without associated findings (Figs. 2 & 7). The main pancreatic duct and collateral side branches may be dilated (Figs. 3a & b). Sometimes stenosis of the main duct at the minor papilla may be evident (Fig. 3c). One case of incomplete pancreas divisum showed a small delicate side branch connecting the ventral with the dorsal duct, a finding that may be more evident on tomographic HASTE images than projectional SS-TSE images because of MIP limitations (Fig. 5). A santorinocoele is a small saccular dilatation of the dorsal duct at minor papilla,\(^5,8\) and we found it in 15% of our cases (Figs. 5 & 6). Its incidence may be higher on MRCP post-secretin stimulation. One case was associated with significant focal dilatation of the main duct in the tail with multiple septa, a finding that may be due to chronic pancreatitis or to IPMT of the main duct type (Fig. 7). The patient refused further investigation and was lost to follow-up. In three patients, we compared a respiratory-triggered 3D-TSE technique that showed pancreas divisum more clearly than the SS-TSE technique (Fig. 3c).

**CONCLUSION**

Although the majority of patients with pancreas divisum are asymptomatic, there is a subset of patients who have either pancreatic-type pain or recurrent episodes of acute pancreatitis. MRCP (particularly with secretin stimulation) may be the best way to diagnose and investigate patients with pancreas divisum because of its capacity as a non-invasive examination.

**REFERENCES**

Question 1. Pancreas divisum is:
(a) A common congenital variant of the pancreatic anatomy. ☐ ☐
(b) A rare form of annular pancreas. ☐ ☐
(c) Mainly asymptomatic. ☐ ☐
(d) Found rarely in patients with unexplained pancreatitis. ☐ ☐

Question 2. The following statements about pancreas divisum are true:
(a) It occurs when the ductal systems of the ventral and dorsal pancreatic ducts fail to fuse during foetal development. ☐ ☐
(b) It is characterised typically by intraluminal filling defects of the pancreatic duct in MRCP. ☐ ☐
(c) It is associated with the presence of choledochal cysts. ☐ ☐
(d) It is associated with cystic pancreatic neoplasm. ☐ ☐

Question 3. Regarding the recurrent attacks of pancreatitis in the presence of pancreas divisum:
(a) They occur more commonly in people with pancreas divisum than in the general population. ☐ ☐
(b) Mucinous pancreatic neoplasms are the predisposing notorious factor. ☐ ☐
(c) Recurrent episodes may be caused due to the presence of relative obstruction to the drainage of pancreatic secretions at the minor papilla. ☐ ☐
(d) Minor papilla sphincterotomy is the treatment. ☐ ☐

Question 4. Regarding pancreas divisum:
(a) Stenosis of the accessory papilla never occurs in cases of pancreas divisum. ☐ ☐
(b) No further therapy should be systematically proposed for patients with pancreas divisum. ☐ ☐
(c) Stress MRCP can reveal dilation of the duct of Santorini and a cystic dilation of the dorsal duct just before its insertion into the minor papilla in patients with pancreatic divisum. ☐ ☐
(d) Endoscopic retrograde pancreatography can be used to make a definitive diagnosis of pancreas divisum. ☐ ☐

Question 5. Parameters that can be assessed before and after administration of secretin in MRCP include the following:
(a) Complete ductal anatomy with presence or absence of pancreas divisum. ☐ ☐
(b) Classification of pancreas divisum. ☐ ☐
(c) Visualisation of side pancreatic branches. ☐ ☐
(d) Ampullary dysfunction. ☐ ☐

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