Intraductal Papillary Mucinous Neoplasm of the Pancreas

A 77-year-old man presented with jaundice, weight loss of 15 kg, right upper quadrant abdominal pain, fever and laboratory studies suggestive of biliary obstruction. Abdominal CT showed dilatation of the pancreaticobiliary tree, a large gallbladder filled with gallstones, and cystic appearing changes in the head of the pancreas, suggesting the possibility of a cystic pancreatic tumor. He was transferred to our hospital for an MRCP, which confirmed these findings but a definitive diagnosis was not made (Figure 1). ERCP revealed a distended ampulla of Vater bulging with mucin (Figure 2a) and multiple filling defects within the dilated pancreatic duct (Figure 2b). Biopsy and brushing of the pancreatic duct was performed and revealed a well-differentiated adenocarcinoma. The patient refused further treatment and was transferred back to his local hospital.

A mucous-secreting pancreatic cancer, now called intraductal papillary mucinous neoplasm, was first described in 1983. In this disease the pancreatic duct is lined by mucin-producing papillary neoplastic epithelium (1). Characteristic ERCP findings are diffuse or segmental dilatation of the main pancreatic duct and side-branches, intraductal filling defects and a dilated pancreatic duct orifice with bulging mucus (2). Magnetic resonance cholangiopancreatography (MRCP) is emerging as a new imaging method for the pancreaticobiliary tree. In a comparative study, similar diagnostic accuracies were reported for both MRCP and ERCP (3). However, in our case the diagnosis was based on the combined endoscopic, radiologic and histopathologic information obtained by ERCP. This example supports the view that, for now, ERCP should be the study of choice for suspected pancreaticobiliary diseases (4). With further improvements, widespread availability and expertise MRCP may become helpful in the better selection of patients for therapeutic ERCP.

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