Spontaneous asymptomatic rupture of pseudocyst into stomach

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DESCRIPTION

A 27-year-old woman presented with complaints of pain in the epigastric region radiating to back for 20 days. She was treated initially at local hospital and was diagnosed as acute pancreatitis as her amylase and lipase levels were >1000 U/mL. She was managed with analgesics and intravenous fluids and was discharged in 4 days. She then presented to our emergency, with complaints of epigastric pain, awareness of lump in the epigastric region and non-passage of stool or flatus for past 3 days. She also gave history of fever and multiple episodes of bilious vomiting. Patient denied previous history of similar episodes in past and is non-alcoholic. On abdominal examination, a lump of size 20×15 cm was palpable in the epigastric region extending into the umbilical region. On ultrasonogram abdomen, a cystic swelling was seen along with multiple gall stones in the gall bladder. Contrast-enhanced CT (CECT) abdomen was suggestive of a large cystic lesion with air-fluid levels occupying hepatoduodenal ligament lying anterior to the stomach and compressing anterior wall of stomach and duodenum (figure 1). A nasogastric tube was placed that drained around 800 mL bilious output on first day. Next morning, the nature of nasogastric output changed to purulent, and the lump disappeared. The following evening, the patient passed stool and flatus and underwent upper GI endoscopy the next day, which depicted a fistulous opening on the anterior wall of stomach with active pus discharge (figure 2). At 6 months of follow-up, patient is asymptomatic and doing well.

Pancreatic pseudocysts are cystic cavity lined by fibrous or granulation tissue containing amylase rich pancreatic juice with or without pancreatic duct communication. They have been defined by Atlanta classification as ‘an encapsulated collection of fluid with a well defined inflammatory wall usually outside the pancreas with minimal or no necrosis’. On CECT, they appear as well-defined, well-circumscribed cavities containing homogeneous fluid density. Spontaneous rupture of a pseudocyst occurs in less than 3% of the cases. Sites of drainage include oesophagus, stomach, duodenum, small intestine, colon, peritoneal cavity, portal vein, splenic vein, retroperitoneum, pleural cavity, pericardium or bile duct. Most of these ruptures are associated with acute abdominal pain, gastrointestinal bleeding or sepsis.

Although the gastric communication was clearly evident on upper GI endoscopy, the evidence of gastric fistula can be made out on CECT abdomen as well (figure 1). Air-fluid levels seen in the pseudocyst cavity, close abutment and compression of the anterior wall of the stomach by pseudocyst point towards possibility of communication with hollow viscus. Several mechanisms of rupture of pseudocysts have been proposed such as: raised intracystic pressure, gradual dissolution of the wall of the cyst by pancreatic enzymes that are activated by enterokinase following a fistulous communication with the bowel, increased intra-abdominal pressure and minor abdominal trauma.

More than half of the patients with pseudocysts do not need any intervention for these cysts. Spontaneous resolution of a cyst depends on the size,
duration, wall thickness, communication of the cyst with pancreatic duct and associated pancreatic duct stricture.4

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REFERENCES

Learning points
► Patients with asymptomatic, uncomplicated, stable or decreasing pseudocysts in size can be observed.
► Symptomatic, complicated and unstable patients with increasing pseudocyst size need an intervention in the form of percutaneous, endoscopic or surgical drainage.
► Conservative management seems appropriate in cases of silent, spontaneous, internal drainage, as in the present patient. However, one must remain cautious to detect complications early.