Painless isolated spontaneous dissection of the superior mesenteric artery

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DESCRIPTION
A man in his early 60s with a history of type 2 diabetes mellitus for more than 10 years and hypertension was admitted to the emergency room due to a sudden onset of nausea and vomiting from 2 days. On admission, he was haemodynamically stable, and no abdominal bruit was appreciated. Laboratory tests showed a slight increase in white blood cell count and C reactive protein, but no other abnormalities in biochemistry and coagulation ability were observed. Contrast-enhanced CT revealed an isolated dissection of the superior mesenteric artery (SMA) without aortic dissection or aneurysm (figure 1). There was an increase in SMA diameter (17 mm, maximum) extending to a length of 33 mm. Partial segmental contrast defect on the intimal flap was detected at 22 mm from the root of the SMA, suggesting an entry of the dissection. The true lumen was smaller due to compression by the false lumen. Both true and false lumens were contrast-enhanced and appeared to be cul-de-sac-shaped. This suggests the false lumen had an entry but not a re-entry and is classified as type 2 according to the original Sakamoto’s classification. The intimal flap of the SMA was identified by both CT scan and grey-scale ultrasound imaging (figure 2). On colour Doppler imaging, a swirling pattern was seen with the ‘yin-yang sign’ of the false lumen (figure 2), showing a to and fro motion of the false luminal blood flow. There was no suggestive sign of complete bowel ischaemia through the initial examinations, nevertheless, acute localised dissection of the SMA and a subsequent truncated upper bowel ischaemia possibly caused his primary symptoms such as nausea and vomiting. A transient ischaemia of the inferior pancreaticoduodenal artery, a branch of the SMA, may cause acute upper abdominal symptoms localised on the ascending and inferior parts of the duodenum.

The patient was treated conservatively with fasting and oral antihypertensive therapy. His nausea and vomiting completely disappeared on day 3, and oral intake was resumed. He was symptom-free within 6 months after discharge and the follow-up CT revealed partial remodelling of the patent false lumen.

In general, the most common symptom was abdominal pain, followed by nausea and vomiting, haematochezia and diarrhoea. In the previous case series, asymptomatic and painless isolated spontaneous dissection of the SMA were reported, accounting for 6.7%–35.7% and 19.7% of the total cases of SMA dissection, respectively.4 Due to a long history of diabetes in the present case, painless dissection with only digestive symptoms seemed to have occurred based on diabetic neuropathy as one of its chronic complications. Most cases without intestinal ischaemia are treated with conservative methods, ie, no endovascular or surgical interventions are performed.2,3 Although conservative treatment includes mainly intestinal rest by fasting, blood pressure control, anticoagulant, antiplatelet and

Learning points

- Isolated spontaneous dissection of the superior mesenteric artery (SMA) is relatively rare; nevertheless, it is still an important differential of acute abdomen.
- Asymptomatic or painless cases of the SMA dissociation are uncommon, but they do exist. These rare conditions may be overlooked in cases of hypaesthesia associated with metabolic diseases such as diabetes mellitus.
thrombolytic therapies, there is no clear, rigid protocol for the treatment of SMA dissection.

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Case reports provide a valuable learning resource for the scientific community and can indicate areas of interest for future research. They should not be used in isolation to guide treatment choices or public health policy.

REFERENCES


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