A 54-year-old man was referred to our hospital because of severe excruciating acute colicky central abdominal pain. The pain was associated with nausea, postprandial vomiting, and constipation. He had history of 15 kg weight loss during the last two months. He was opium addict, and cigarettes smoker (20 packs-year) with a negative history of diabetes mellitus and hypertension. He underwent a surgical operation two months before, when he had a similar attack of abdominal pain. The surgery was then revealed small bowel necrosis for which resection of 30 cm of small bowel was performed. General physical examination was unremarkable. There was mild tenderness in deep palpation of periumbilical area. Laboratory data including serum amylase were reported to be normal.

The initial abdominal ultrasonography was normal. Upper gastrointestinal endoscopy and colonoscopy were also normal. On account of his severe abdominal pain, a multislice helical abdominal computerized tomography with contrast was requested (Figure 1).

What is Your Diagnosis?
See the next page for the diagnosis
Visceral artery aneurysms are found in only 0.2% of the general population. Among these, aneurysms of the superior mesenteric artery (SMA) are very unusual and account for only 5.5 – 8%.1 – 4 Most visceral artery aneurysms are located in either the splenic or hepatic arteries. SMA involvement is the third most common type of reported splanchnic artery aneurysm.1 – 4 SMA aneurysms, though uncommon, are lethal and must be treated expeditiously to avoid mortality and high incidence of ischemic small bowel complications. Risk of rupture or embolization is the impetus for their definitive treatment.1 – 4 Prompt diagnosis and treatment is essential to decrease the mortality and minimize the prevalence of intestinal infarction.5 These aneurysms are typically located distal to the origin of the SMA. This situation lends itself to interposition grafting as a means of both aneurysm repair and reestablishment of prograde SMA blood flow.1 – 4 Previous reports indicate that most SMA aneurysms were the result of bacterial seeding from endocarditis or septicemia.1 – 6 However, recent reports demonstrate atherosclerosis as the leading cause of this particular arterial dilatation.3 The SMA aneurysm in our patient appeared to be atherosclerotic in origin. Nonatherosclerotic visceral artery aneurysms are often multiple and their etiologies include arterial dissection, polyartritis nodosa, posttraumatic, inflammation from pancreatitis, Marfan’s syndrome, Ehlers-Danlos syndrome, Takayasu’s arteritis, α1-antitrypsin deficiency, and fibromuscular dysplasia.5 – 7 Symptoms of an SMA aneurysm are often subtle, and patients may present with chronic nonspecific or postprandial abdominal pain.1 – 4 Some patients present with acute complications of rupture or dissection.1 – 4 A pulsatile abdominal mass or bruit may be present in up to 50% of patients.2 The rate of diagnosis of asymptomatic aneurysms has been increased for use of CT and arteriography that aid in the diagnosis of vague abdominal complaints (Figure 2).1 – 4 The appropriate operative repair technique is based on anatomic features of the aneurysm, patency of the SMA, competency of visceral collateral circulation, and blood supply to the small intestine.

Most SMA aneurysms are not at the origin and can be surgically repaired by ligation/exclusion, resection with or without graft reconstruction, or endo-aneurysmorrhaphy. Revascularization is mandatory if bowel ischemia is present or a possibility. Otherwise, exclusion by simple ligation or by endovascular means may be an adequate treatment for the usual distal-based SMA aneurysm that is accompanied by a well-developed collateral circulation.1 – 4 In our patient, the SMA aneurysm was detected on CT scan and CT angiography. Abdominal pain in our patient became very severe after meal with a history of 15 kg weight loss during two months. During surgery, a thrombosed superior mesenteric aneurysm and stenosis of jejunal branches was detected. Thromboendarterectomy and patch graft of anterior wall of aneurysm with fugarty of jejunal branches were done. Postoperatively, abdominal pain subsided and the patient could tolerate food. He gained 7 kg during five months of follow-up.

References

4 Komori K, Mori E, Yamaoka T, Ohta S, Takeuchi K,


