Fatal Gastrointestinal Bleeding Probably Caused by an Aortoduodenal Fistula Following Surgical Repair of an Inflammatory Abdominal Aortic Aneurysm during Postoperative Steroid Therapy

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Abstract

We describe a case of fatal gastrointestinal bleeding probably caused by an aortoduodenal fistula following open surgical repair of an inflammatory abdominal aortic aneurysm (AAA) during postoperative steroid therapy. A 71-year old man underwent a graft replacement for an inflammatory AAA. Prolonged elevated C-reactive protein and newly emerged ascites after surgery were improved by oral administration of predonine. The steroid therapy had been continued until 5 months after surgery, when he suffered from fatal gastrointestinal bleeding. We speculated that the gastrointestinal bleeding was caused by an aortoduodenal fistula and discuss the etiology of this condition.

Keywords: Inflammatory abdominal aortic aneurysm; Aortoduodenal fistula; Steroid

Introduction

Inflammatory abdominal aortic aneurysm (AAA) is characterized by markedly thickened aneurysmal walls and adhesion to adjacent organs. It is generally thought that the inflammatory process of the disease can be resolved by surgical treatment. Rarely, however, cases with progression of inflammation or newly developed complications in preoperatively uninvolved organs have been reported [1,2]. We herein report a case of open surgical repair of an inflammatory AAA, which complicated with prolonged elevated C-reactive protein (CRP) and newly emerged ascites after surgery. Although both disorders were improved by steroid therapy, our patient suffered from fatal gastrointestinal bleeding presumably due to an aortoduodenal fistula.

Case Report

A 71-year old man, who complained of abdominal pain, was referred to our hospital. Computed tomography (CT) revealed an infrarenal AAA with a maximal diameter of 80 mm at the level of the right common iliac artery. The density of fat around the aneurysm was high, which suggested the presence of inflammation (Figure 1A). Laboratory examination showed that the inflammatory response was elevated, with a white blood cell (WBC) count of 11000/μl and CRP of 19.3 mg/dl, which suggested the presence of inflammation (Figure 1A). Laboratory examination showed that the inflammatory response was elevated, with a white blood cell (WBC) count of 11000/μl and CRP of 19.3 mg/dl, though he had no fever. We performed emergent surgery because of the remnant aneurysm wall and peri-aortic region were still enhanced and high peri-aortic fat density remained, though there was no obvious sign of graft infection. Furthermore, 30-day postoperatively, the patient complained of loss of appetite and abdominal distension due to ascites (Figures 2A and 2B). The causes of transudative ascites, such as hypoalbuminemia, liver dysfunction, and portal hypertension, were not detected. Although an abdominal CT was performed again, there was no remarkable change, except the newly emerged ascites. Thereby, we considered that inflammation of the remnant aneurysmal wall and peri-aortic region were still persistent, and decided to start steroid therapy, which consisted of the oral administration of predonine at 20 mg/day. The steroid therapy drastically improved the patient’s condition and the dose of predonine was gradually decreased to 5 mg/day, however, it was returned to 10 mg/day postoperatively, the patient complained of loss of appetite and abdominal distension due to ascites (Figures 2A and 2B).

In the supine position, a full-laparotomy was made. The posterior peritoneum, which was red and thickened, widely adhered to the aneurysm (Figure 1B), though fortunately, the duodenum did not. A graft interposition using a Dacron graft (Gelsoft Plus 16 mm×8 mm, Vaskutek Ltd, Inchinnan, Renfrewshire, Scotland) with right internal iliac aneurysmorraphy was performed without any problems. Histological examination of the resected aneurysmal wall revealed that inflammatory infiltration of the lymphocytes existed in thickened adventitia with fibrosis, which was compatible with an inflammatory aneurysm (Figure 1C). Bacterial examination did not detect any microorganisms.

After surgery, WBC was normalized, but CRP was not improved and remained at approximately 15 mg/dl. Because the patient had no fever, we did not test blood culture. CT showed that the remnant aneurysm wall was enhanced and high peri-aortic fat density remained, though there was no obvious sign of graft infection. Furthermore, 30-day postoperatively, the patient complained of loss of appetite and abdominal distension due to ascites (Figures 2A and 2B). The causes of transudative ascites, such as hypoalbuminemia, liver dysfunction, and portal hypertension, were not detected. Although an abdominal CT was performed again, there was no remarkable change, except the newly emerged ascites. Thereby, we considered that inflammation of the remnant aneurysmal wall and peri-aortic region were still persistent, and decided to start steroid therapy, which consisted of the oral administration of predonine at 20 mg/day. The steroid therapy drastically improved the patient’s condition and the dose of predonine was gradually decreased to 5 mg/day, however, it was returned to 10 mg/day postoperatively, the patient complained of loss of appetite and abdominal distension due to ascites (Figures 2A and 2B).

C. Histological examination of the resected aneurysmal wall. Inflammatory infiltration of lymphocytes observed in the adventitia.

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mg/day because of re-elevation of CRP (Figure 3). Finally, he was discharged with oral administration of predonine at 10 mg/day on the 43rd postoperative day. Four-month postoperatively, CRP decreased to 0.99 mg/dl with continuous administration of predonine at 10 mg/day, though the enhancement of the remnant aneurysmal wall and high peri-aortic fat densities were still observed on CT (Figures 2C and 2D).

Figure 2: Abdominal CT before and after steroid therapy.
A and B: Abdominal CT before steroid therapy. The remnant aneurysmal wall around the graft is enhanced and peri-aortic fat density is high. In addition, ascites is observed.
C and D: Abdominal CT after steroid therapy. (CRP indicated 0.99 mg/dl.) Ascites disappeared, but inflammatory signs of the remnant aneurysmal wall remained. The duodenum adhered to the remnant aneurysmal wall (Stars indicates this portion).

Figure 3: Relation between inflammatory response and dose of predonine.
Prolonged elevated CRP maintained after surgery, however, it was dramatically improved by steroid therapy. When dose of predonine was decreased to 5 mg/day, the value of CRP elevated to 14.4 mg/dl.

Figure 4: Pre and post-mortal abdominal CT.
A and B: Abdominal CT just before cardiac arrest. Peri-graft space is dramatically shrunken with bubble formation.
C and D: post-mortem abdominal CT. Alimentary tract is filled with a massive amount of blood.

Five-month postoperatively, however, the patient came to our emergency room complaining of fever, abdominal pain and diarrhea. Laboratory examination, revealed that inflammatory response was re-elevated (WBC: 9200/μl, CRP: 15.6 mg/dl). CT showed that the peri-graft space was dramatically reduced and was accompanied by bubble formation (Figure 4A and 4B). Although we immediately began antibiotic therapy and scheduled further examinations, the patient suddenly vomited a massive amount of blood and went into cardiac arrest. We performed cardiopulmonary resuscitation, but could not save his life. Post-mortal CT showed that the gastrointestinal tract was filled with a massive amount of blood and that there was no bleeding into the retroperitoneal space (Figures 4C and 4D). On the basis of pre- and post-mortal CT findings, we speculated that a fistula had formed between the duodenum and peri-graft space, which subsequently caused the graft infection leading to final fatal bleeding from the anastomosis into the duodenum.

Discussion
Inflammatory AAA, which was first reported by Walker et al., is characterized by marked thickening of the aneurysmal wall and strong adhesion to adjacent organs. Histologically, marked thickening of the adventitia with fibrosis and infiltration of inflammatory cells, mainly lymphocytes, are observed [3]. The inflammatory process can affect and obstruct the duodenum, ureters, left renal vein, inferior vena cava, and rarely, the small bowel or colonic mesentery [4]. The etiology of inflammatory AAA remains uncertain, though several theories, such as an autoimmune disease, a chronic viral infection, and a variation of the inflammatory process of atherosclerosis, have been previously discussed [5].

The treatment of inflammatory AAA involves surgical repair, endovascular stent grafting and steroid therapy. Surgical repair of inflammatory AAA, which is commonly a prosthetic graft replacement, is conventional. Although there are technical problems which are associated with adherence to adjacent structures, the results compare favorably with common atherosclerotic AAA [6]. In most cases, a prosthetic graft replacement not only prevents rupture, but also regresses the inflammatory process. However, there are a few cases with persistent inflammation after surgery or postoperative development.
of organ complications in preoperatively uninvolved structures, and it has been reported that postoperative steroid therapy improved complications caused by a persistent or progressive inflammatory process after surgery [1,2]. In our case, CT showed that the enhancement of the remnant aneurysmal wall and high peri-aortic fat density remained after surgery, which apparently suggested persistent inflammation of the remnant aneurysmal wall. On the contrary, no other abdominal organs showed inflammatory changes. Furthermore, our patient did not have other causes of transudative ascites. Thereby, we considered that the elevated CRP and ascites were caused by persistent inflammation of the remnant aneurysmal wall, and performed steroid therapy. Actually, ascites disappeared and the value of CRP fluctuated depending on dose of predonine (Figure 3). These facts suggested that postoperative steroid therapy was effective in our case.

Nevertheless, our patient died of gastrointestinal bleeding. Although an autopsy could not be performed, we speculated that a fistula between the duodenum and peri-graft space was formed, subsequently, the peri-graft space dramatically shrunk and the graft infection occurred. Finally, our patient suffered from fatal bleeding from the anastomosis to the duodenum.

Aortoenteric fistula, which often causes massive gastrointestinal bleeding, is divided into two groups, such as primary and secondary. Primary aortoenteric fistula is a communication between the native aorta and gastrointestinal tract. While secondary aortoenteric fistula arises from the suture line after abdominal aortic reconstructive surgery, especially using a prosthetic graft [7]. The etiology of secondary aortoenteric fistula after prosthetic aortic grafting includes direct contact with the prosthetic graft or the site of anastomosis, pseudoaneurysm formation caused by failure of anastomosis, and graft infection.

In our case, there were two possible causes of a fistula between the duodenum and peri-graft space. One was the local persistent inflammation of the remnant aneurysmal wall. On the basis of a series of CT findings, the prosthetic graft did not contact with the duodenum directly. Before the graft infection, the duodenum adhered to the enhanced aneurysmal wall located around the graft (Figures 2C). Such cases have never been reported, although cases of an aortoduodenal fistula following endovascular repair of inflammatory AAA have been reported [8]. Furthermore, primary aortoenteric fistula without aortic aneurysms is caused by inflammation of the aorta itself, such as septic aortitis [7].

The other potential cause was a duodenal ulcer which was one of the side effects of steroid, however, the risk of upper gastrointestinal complications induced by steroid is still debated [9]. In addition, we administered "low" dose of predonine. Our patient did not take NSAIDs, but took PPI and also the duodenal ulcer prefers to affect the bulbus. The cases with peptic ulcer induced aortoenteric fistula have been previously reported [10]. Most cases affected the esophagus and stomach, rarely duodenum, and had the gastrointestinal disorders except peptic ulcer, such as the hiatus hernia and after surgery of the gastrointestinal tract. The reports about an aortoduodenal fistula caused by only peptic ulcer are rare and outdated.

In conclusion, we experienced a case of inflammatory AAA complicated with fatal gastrointestinal bleeding probably caused by an aortoduodenal fistula. The pathological process of inflammatory AAA is rarely persistent after surgery, in such cases, steroid therapy is effective. However, it might not be able to completely resolve persistent local inflammation, attention should be given to any newly emerged complications, especially, in cases which CT shows the presence of inflammation of the remnant aneurysmal wall.

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