Successful Surgical Treatment of an Infected Thoracoabdominal Aneurysm Accompanied with Leriche Syndrome

Masato Furui,1 Hirohisa Hirata,2 Bunpachi Kakii,1 Gaku Uchino,1 Mai Asanuma,1 Haruo Suzuki,3 Hiroaki Nishioka,2 and Takeshi Yoshida1

1Cardiovascular Surgery Department, Matsubara Tokushukai Hospital, 7-13-26 Amamihigashi, Matsubara, Osaka, Japan
2Surgery Department, Matsubara Tokushukai Hospital, 7-13-26 Amamihigashi, Matsubara, Osaka, Japan
3Cardiovascular Surgery Department, Kyoto Renaiss Hospital, 4-13, Suehirocho, Fukuchiyma, Kyoto, Japan

Correspondence should be addressed to Masato Furui; masatofurui@yahoo.co.jp

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1. Introduction

Leriche syndrome, or aortoiliac occlusive disease, usually presents with claudication, erectile dysfunction, and absence of a lower limb pulse and is often a result of arteriosclerosis obliterans in the lower limb. The study patient proved to be an extremely rare case of Leriche syndrome accompanied by an infected thoracoabdominal aortic aneurysm [1–5].

2. Case Presentation

A 56-year-old man was transported to our hospital because of massive melena and loss of consciousness. He was a current smoker with a history of hyperlipidemia. Physical examination revealed no inguinal pulse. He complained of mild chronic claudication despite having no impotence, and computed tomography revealed aortoiliac occlusive disease owing to atrophy and calcification of the terminal aorta and iliac arteries. His vital signs included a blood pressure of 135/70 mmHg and pulse of 90 beats/min. The white blood cell count was 16,300 cells/mm³, hemoglobin was 10.5 g/dL, and C-reactive protein level was 1.93 mg/L. A 35 mm thoracoabdominal aneurysm with intra-aneurysmal gas was seen on enhanced computed tomography (Figure 1), consistent with an infected thoracoabdominal aneurysm accompanied by Leriche syndrome.

Emergency surgery was performed under general anesthesia with the patient in the right semilateral position. The procedure began with a left thoracoabdominal incision through the seventh intercostal space. When we reached the aorta through careful retroperitoneal approach, we could not reserve the left inferior epigastric artery, running obliquely around the incision area (Figure 1). The color of aortic aneurysm wall changed from yellowish-white to black-brown. The aorta was clamped above the celiac trunk because the infection had spread around the superior mesenteric and bilateral renal arteries using a four-branched graft. The aortoduodenal fistula was resected after omental filling, and an enterostomy was performed for feeding. Intestinal reconstruction was performed in two stages. The patient was discharged on postoperative day 48 and was without evidence of recurrence at 23 months postoperatively.
aorta had a blind end because of Leriche syndrome, we used a
knitted graft with four branches (INTERGARD quarto 18-9
mm; Maquet Getting Group, Tokyo, Japan) (Figure 2(a)).
The SMA and right renal artery were reconstructed using
the right graft branch, and the left renal artery was recon-
structed with the left graft branch (Figures 2(b) and 3(a)).
The second left graft branch was passed to the left common
to femoral artery because of the left lower limb ischemia by
sacrificing the inferior epigastric artery. The omentum was
filled around the graft and debridement cavity. The proce-
dure time was 497 min and CPB time was 124 min. After
the vessel reconstruction, a horizontal segment of the duode-
num containing a 5 mm fistula was resected (Figures 2(a)
and 2(b)). The ends of the duodenum and proximal jejunum
were closed by an intestinal two-stage reconstruction; the first
stage includes removal of the aortoduodenal fistula and
stump of the edges, and the second stage includes a duodo-
jejunostomy. And the intestinal fistula was made for feeding
(Figure 2(c)). On postoperative day 1, the patient complained
of severe pain in the right lower limb on sitting. A Doppler
evaluation of the bilateral dorsalis pedis arteries revealed
right lower limb ischemia. Femorofemoral bypass was
performed using an ePTFE graft (Gore Propaten 8 mm;
WL Gore & Associates, Co. Ltd., Tokyo, Japan) attached to
the side of the leg graft branch (Figure 3(b)).

Streptococcus and Peptostreptococcus asaccharolyticus
were isolated from the aneurysm. Vancomycin and merope-
nem were administered for 1 month, and a duodenojejunos-
omy, as the second stage, was performed on day 30 after the
inflammation subsided (Figure 2(d)). Antibiotic was changed
to oral levofloxacin on day 43, and the intestinal fistula was
removed on day 44. The patient was discharged on day 48
and continued antibiotics at home for 6 months. To the best
of our knowledge, there has been no evidence of any com-
pliances such as graft infection and pseudoaneurysm for
23 months postoperatively.

This study was approved by the Ethics Committee
of Matsubara Tokushukai Hospital (approval number:
171202). The patient consented to the publication of
this report.

Figure 1: Preoperative enhanced computed tomography indicates an infected aortic aneurysm at the thoracoabdominal level. (a) Small
aneurysm with intra-aortic air (arrow), the superior mesenteric artery (star), and renal artery in the axial view. (b) Obstructed distal
abdominal aorta and common iliac arteries (arrows) with calcification in the coronal view. (c) Suspected aortoduodenal fistula
(arrowhead) in the sagittal view.
3. Discussion

Only 0.7%–3% of all aneurysms are infected, and those at the thoracoabdominal level are extremely rare [1–4]. The challenges of successful treatment include the revascularization in the infected territory, control of sepsis, and reconstruction of the digestive tract. Thoracic endovascular aortic repair for Leriche syndrome or infected aneurysms has been reported [5, 6]. In this case, open surgery was required because fibrosis and atherosclerosis of the iliac arteries made an endovascular approach impossible and debridement of the infected aorta was desirable. The revascularization of an infected aortic aneurysm by in situ or nonanatomical reconstruction is controversial. Yasuda et al. reported that extra-anatomical bypass was safe and reliable [2], and others have reported successful in situ reconstruction [1, 3, 5]. In this patient, the infected aortic aneurysm was successfully treated by debridement and omental filling, although we could not prepare a rifampicin-soaked graft owing to an emergency case.

Revascularization for the right lower limb was initially considered. We assumed bypass for the lower extremities as an option, but it was not performed because the patient complained of only mild claudication and had no trouble performing his daily activities [4]. Because femoral arteries were exposed for partial CPB and the patient required revascularization sooner or later, we should have grafted the bilateral femoral arteries in the first surgery.

Reports of intestinal reconstruction vary from case to case [7, 8]. Intestinal one-stage reconstruction to connect the duodenum and jejunum during the first procedure is simpler and easier than two-stage reconstruction; however, a minor leak following intestinal reconstruction can result in reinfection. We chose a two-stage reconstruction with initial resection of the fistula and an enterostomy for feeding, and a subsequent duodenojejunostomy after recovery, to ensure safety. Although the two-stage reconstruction requires a longer hospital stay than one-stage reconstruction, we believe it is more beneficial for patients in terms of

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**Figure 2:** Diagram of the surgical procedure. (a) Aneurysm, aortoduodenal fistula (arrow), and a 4-branched graft are shown. Oblique lines show the infectious area. Wavy lines are resected areas of the duodenum and jejunum. (b) Aortic repair and left leg revascularization are shown. The aortoduodenal fistula was subsequently resected. (c) Enterostomy was attached to the jejunum as the first stage of intestinal reconstruction. (d) The duodenojejunostomy was performed as the second stage of intestinal reconstruction.
decreasing morbidity and mortality in such unstable condition than one-stage surgery.

There is a room for further consideration about the order of procedures of an intestinal resection and revascularization by graft replacement. In the present case, revascularization preceded the intestinal resection similar to previous studies [9–11]. The patient actually recovered well. However, grafting in the irrigated surgical field after the intestinal resection may be an ideal order of procedure for decreasing the amount of bacteria around the graft. This order remains a debatable issue to be discussed with the digestive surgeons.

In conclusion, we reported on an extremely rare occurrence of an infected thoracoabdominal aneurysm associated with Leriche syndrome, which was successfully treated with an in situ four-branched vessel graft replacement, omental filling, and two-stage intestinal reconstruction. The prompt diagnosis and aggressive surgical treatment saved the patient with an infected aortic aneurysm and aortoduodenal fistula.

Conflicts of Interest

The authors declare that there is no conflict of interest regarding the publication of this article.

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