Aorto-esophageal fistula after thoracic endovascular aortic repair

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Abstract: Aorto-esophageal fistula (AEF) is a rare complication of Thoracic endovascular aortic repair (TEVAR). It is difficult to treat, and patients rarely survive without treatment. An 88-years-old woman presented with AEF and sepsis of Klebsiella pneumoniae. Since the open surgery was difficult, we implanted an esophageal stent to cover the fistula. It enabled her to eat and drink normally for 4 months. She died of hemorrhagic shock; however, the esophageal stent remained intact. Even though esophageal stent treatment is generally not totally complete, it is a favored treatment option for those who wish to eat and to live normal life as long as possible.

Keywords: Aorto-esophageal fistula (AEF); thoracic endovascular aortic aneurysm repair (TEVAR)

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Introduction

Thoracic endovascular aortic aneurysm repair (TEVAR) was first approved for use in the US in 2005, and it has been an appealing alternative to the traditional open aortic aneurysm repair because of the comparatively better early outcomes (1). Major complications of TEVAR that typically necessitate open distal aortic repair include endoleak, aortic fistula, infection, device collapse or migration, and continued expansion of the aneurysm sac (1). Among these complications, aorto-esophageal fistula (AEF) is a rare and life-threatening complication.

We experienced a case of AEF with sepsis. Because of patient’s age and poor general condition, she was not a candidate for open surgery. Here we report our unique solution to AEF using an esophageal stent to cover an opening in the esophagus to the fistula.

Case presentation

An 88-year-old woman visited our hospital presenting with appetite loss and chest pain. She was diagnosed with sepsis and was admitted. Seven months prior to admission, she had TEVAR for thoracic aortic aneurysm in another hospital. Her past history also included coronary artery bypass surgery and endovascular aortic repair for iliac artery. After TEVAR, she refused to visit the hospital for 6 months. Laboratory data at admission showed elevated white blood cells (WBC) counts of 12,400 μL and elevated C-reactive protein (CRP) of 19.29 mg/dL. Klebsiella pneumoniae was isolated in all 4 of 4 bottles of blood culture. Contrast chest CT scan revealed fluid congestion in mediastinum and free air around the stent-graft (Figure 1). Gastric fiberscopy revealed AEF in the middle intrathoracic esophagus, about 10 mm in diameter. The pus was flowing into the esophagus from the mediastinum (Figure 2). Klebsiella pneumoniae was also isolated from the pus. Complete repair by open surgery was high risk due to her advanced age and history of multiple cardiovascular surgeries, including coronary artery bypass surgery and endovascular stent grafting of the common iliac artery.

Oral intake was prohibited. She was on hyperalimentation and Ceftriaxone (CTRX) 2 g/day for 4 weeks. She was mentally alert and wished to eat her favorite foods. She underwent gastric fiberscopy again which revealed that the AEF had become more than double in size (Figure 2). Aortic stent-graft was prominent through such opening. In order to
grant her wish to eat and drink again, we implanted a Niti-S esophageal stent (Taewoong Medical, Seoul, South Korea) 18 mm × 100 mm partial covered type to cover the entry to the fistula. She underwent prandial rehabilitation for 5 weeks. As a result, she was able to eat normally again and was thus discharged from the hospital.

Seventeen weeks after the surgery she suddenly vomited blood and fell into hemorrhagic shock. Body CT scan showed blood in the stomach, however did not show any active bleeding (Figure 3). After blood transfusion she recovered from shock and became stable. Ten days after re-admission, gastric fiberscopy did not show any bleeding, and the esophageal stent was not dislocated (Figures 2,3). Fourteen days after re-admission she displayed sudden cyanosis followed by cardiac arrest and passed away. The hemoglobin level was 4.4, which suggested that she died of hemorrhagic shock.

**Discussion**

AEF is a rare post-TEVAR complication. It has been reported that the incidence of AEF post TEVAR is 1.5–1.9% (2-4). Aortic fistulas are difficult to treat, and patients rarely survive without treatment. The mechanism by which AEF develops is not known. Coselli *et al.* wrote in their review that fistulas develop after TEVAR because of the radial pressure the endograft exerts on nearby structures, coverage of arteries that provide blood flow to the esophagus (which could cause ischemic necrosis), a perforation, an infection, or erosion due to either an endoleak or infection-related pseudoaneurysm (1). Our case patient had refused to visit the hospital after TEVAR for 6 months, even though she was instructed by her previous hospital to do so. Therefore it is not known how she developed AEF. Since she presented with severe mediastinitis and sepsis, it is highly likely that AEF was worsened by the infection. The question was how she developed mediastinitis. The mediastinitis was caused by either aortic stent-graft infection or by esophageal perforation. Aortic stent-graft infection is rare, and the incidence is only 0.5–1% (5). It is more likely that the mediastinitis was caused by esophageal perforation as the result of swallowing of fish bone, chicken bone, or some other such hard item.

Esophageal stent is commonly used for esophageal stenosis such as esophageal cancer. It is typically constructed of nickel and titanium. Potential bleeding due to the friction between the two metal stents was of concern. Another concern was possible dislocation or

![Figure 1 Contrast CT scan on admission. Fluid congestion in mediastinum and free air around the stent-graft (arrows) is observed. (A) Axial plane; (B) sagittal plane.](image-url)
migration of the stent, because the patient's esophagus had an elastic surface unlike esophageal cancer. However, the patient and her family strongly desired to allow oral intake and for her to go home. Esophageal stent was subsequently inserted after obtaining informed consent. The patient would not have survived without such esophageal stent intervention, since the opening in the esophagus had grown significantly larger even though she did not eat for 4 weeks, and the mediastinitis was not controlled with antibiotics. It became necessary to repair the significantly enlarged opening in order to prevent the esophageal mucus or saliva from flowing into the mediastinum.

There are only three case reports in the medical literature that utilized esophageal stent to repair AEF developed after TEVAR. Onodera et al. used a covered self-expanding esophageal stent to repair AEF which developed 2 years after TEVAR, but the patient died of sepsis 34 days later (6). Tao et al. used Niti-S stent to tamponade the bleeding point of AEF secondary to TEVAR, but the stent migrated because of continuous vomiting so that they had to insert another stent (7). This patient died of myocardial infarction 8 months later. Eggebrecht et al. also used self-expanding esophageal stent for two patients of AEF secondary to TEVAR, but they died of mediastinitis in 5 weeks and 10 months (8). We used Niti-S stent with distal flare to prevent migration, and it was successfully inserted without dislocation. Nonetheless, none of the cases survived beyond one year with esophageal stent. It is thus only a temporal repair to close the fistula so that the patient can enjoy eating again, living a normal life at home with family, and significantly improve quality of life (QOL).

The cause of patient's death points to hemorrhagic shock since the hemoglobin level dropped to one-third. This further points to the patient having aortic bleeding since she had sudden cyanosis and cardiac arrest. It is thus uncertain as yet whether the bleeding occurred due to friction between the two stents, or because the artery was damaged.

Figure 2 Gastric fiberscopy. (A) On admission. Opening of fistula is observed in the middle intrathoracic esophagus. The pus is flowing into the esophagus from the mediastinum; (B) 4 weeks after admission. The fistula has grown, and the aortic stent is prominent; (C) esophageal stent is implanted; (D) 10 days after re-admission. Bleeding is not observed in the stent.
by prolonged mediastinitis.

Conclusions

Esophageal stent is shown to be an alternative treatment option for patients of AEF secondary to TEVAR who cannot undergo open surgery. Although esophageal stent treatment is generally not totally complete, it is shown a favored treatment option for those patients who wish to enjoy eating and drinking normally, enjoy and live a normal life, with improved QOL as long as possible.

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None.

Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

Informed Consent: Written informed consent was obtained from the patient for publication of this manuscript and any accompanying images.

References


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