Novel Management of an Aortoesophageal Fistula Prior to Definitive Surgical Repair

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ABSTRACT: An aortoesophageal fistula (AEF) is a rare but often fatal complication associated with the placement of an aortic stent graft such as those utilized in thoracic endovascular aortic repair (TEVAR) procedures. Definitive surgical repair of AEF is the treatment of choice, as conservative management is nearly always fatal. We present the case of an AEF in a 74-year-old male managed by a unique treatment method: an esophageal stent was deployed to cover the fistula as a temporizing measure prior to definitive surgical correction, thus allowing time for resuscitation and hemodynamic optimization. The use of esophageal stents in the setting of AEF following TEVAR has been previously reported in the literature as a palliative measure for patients deemed incapable of tolerating open repair surgery. Our case demonstrates a new and innovative approach to the management of AEF following TEVAR in which the use of esophageal stenting is expanded beyond the role of palliative care and should be considered as a means to optimize at-risk patients prior to definitive corrective surgery in the hopes of improving outcomes.

KEYWORDS: Aortoesophageal fistula (AEF), TEVAR, esophageal stent, thoracic aorta

Introduction
An aortoesophageal fistula (AEF) is a rare but life-threatening cause of upper gastrointestinal bleeding that results from a communication between the esophagus and aorta. Etiologies include foreign body ingestion, malignancy, infection, and thoracic aortic aneurysms.1 Rarely a fistula results after placement of an aortic stent graft, such as those utilized in thoracic endovascular aortic repair (TEVAR) procedures.1,2 Although less invasive than an open repair, TEVAR carries its own morbidity and mortality due to the risk of unanticipated complications.2 Aortoesophageal fistula is a rare but often fatal complication occurring in 1.9% of TEVAR procedures.2

We present a case of an AEF in a 74-year-old male managed by a unique treatment method whereby an esophageal stent was deployed covering the fistula as a temporizing measure prior to definitive surgical correction allowing time for optimization. Written authorization was obtained from the patient prior to publication of this case.

Case Description
A 74-year-old male with a past medical history significant for chronic obstructive pulmonary disease, cerebral vascular accident, hypertension, peripheral vascular disease, and type 2 diabetes mellitus presented to the emergency department with hematemesis versus hemoptysis 2 months after undergoing emergent TEVAR for a ruptured thoracic aortic aneurysm. A computed tomography (CT) angiogram illustrated an AEF.

The patient was admitted to our Surgical Intensive Care Unit (SICU) and resuscitated, then the patient was promptly taken to the operating room for flexible bronchoscopy and esophagogastroduodenoscopy (EGD). Bronchoscopy was negative for endobronchial lesions in all lung lobes. On endoscopy, a large thrombus was noted in the mid-esophagus and significant blood was present in the stomach. Following irrigation and suction, no active bleed was discovered in the stomach, and attention was given to the large esophageal thrombus believed to represent the connection to the aorta. The vascular surgery team who placed the initial endograft was consulted, and the decision was made to stent the esophagus as a temporizing measure to tamponade the fistula. Fluoroscopy was utilized to measure the length of the thrombus to determine the size of the stent needed to fully cover the area. The large clot measured around 5 cm, extending from the level of about 30 cm from the incisors to 35 cm. In order to ensure that the entire defect was completely covered, a 15-cm stent was chosen. The area was delineated with radiopaque markers, and under direct vision, a guidewire was placed through the endoscope which remained in the esophagus. Under fluoroscopic guidance, the 15 cm stent was deployed, extending above and below the region of the thrombus.

The interim between esophageal stent placement and definitive surgical repair allowed adequate time for medical resuscitation and optimization of the patient’s condition. Seven days after esophageal stent placement, the patient returned to the operating room for definitive surgical repair of the AEF including left muscle-sparing thoracotomy, partial esophagectomy and high cervical esphagostomy, and resection of the infected thoracic aortic stent graft under femoral-femoral cardiopulmonary bypass. The
thoracic aorta was reconstructed utilizing a rifampin soaked 24 mm × 30 cm gel weave tube graft. The surgery was completed without complication and the patient tolerated the procedure well.

Post-operatively, the patient was transitioned to enteral feeds via a J-tube. IV ampicillin-sulbactam was given for 6 weeks after tissue cultures grew Enterococcus faecalis. After a 22-day hospital course, the patient was discharged to a skilled nursing facility for further rehabilitation.

Discussion
The incidence of AEF following endovascular repair of the aorta has been reported to range from 1.7% to 1.9%, although this rate is expected to rise as TEVAR continues to be an increasingly utilized alternative to open surgery for emergent treatment of aortic conditions. Fistula formation usually occurs within 1 to 16 months following the original TEVAR procedure. The definitive mechanism by which fistula formation occurs has not yet been elucidated. Possible mechanisms that have been hypothesized include erosion of the stent graft through the aorta, endoleaks into the residual aneurysm sac, pressure necrosis on the esophageal wall due to progressive expansion of the endograft, and ischemic necrosis of the esophageal wall secondary to the stent graft compressing aortic vessels that supply the esophagus.

The classic presenting symptoms of an AEF have been described as the triad of mid-thoracic chest pain, sentinel hemorrhage with a small episode of hematemesis followed by a brief interval period, and eventual massive hematemesis and exsanguination. Constitutional signs and symptoms such as new onset fever, fatigue, anorexia, and elevated inflammatory markers are also commonly associated with this condition. Aortoesophageal fistula should be considered early and often in the differential for a patient with a history of TEVAR presenting with any of the aforementioned symptoms. Although many potential causes of hematemesis exist, AEF must be ruled out first in a patient prior to undergoing open repair. The use of esophageal stents in the setting of AEF following TEVAR has been reported previously, but mainly as a palliative measure for patients deemed incapable of tolerating open repair surgery. Our case demonstrates a novel management approach in which an esophageal stent was placed as a temporizing measure 7 days prior to definitive AEF repair. The stent was placed during the interval period after the sentinel bleed but prior to a potential massive hemorrhage, thus preventing potential exsanguination by tamponading the fistula. Stenting the esophagus also provided an ample time frame for resuscitation and hemodynamic optimization of the patient prior to undergoing open repair. The use of esophageal stents in the setting of AEF following TEVAR has been reported previously, but mainly as a palliative measure for patients deemed incapable of tolerating open repair surgery.

Our case indicates there may be a further role for esophageal stenting in AEF management. This management strategy is not without risk, however, as there is the possibility for stent migration or further vascular erosion from the friction between the endograft and esophageal stent. The stent in our case was only deployed for 7 days, which may have limited the potential for complications, but evidence from further cases utilizing this approach would be beneficial. The temporizing measure of stenting the esophagus provides adequate time to resuscitate and optimize at-risk bleeding.
patients prior to definitive corrective surgery in the hopes of improving morbidity and mortality outcomes (Figure 1).

Author Contributions
Conceived and designed the experiments: ES, AMJ, JJM; Analyzed the data: ES, AMJ, JJM; Wrote the first draft of the manuscript: ES, AMJ, JJM; Contributed to the writing of the manuscript: ES, AMJ, JJM; Agree with manuscript results and conclusions: ES, AMJ, JJM; Jointly developed the structure and arguments for the paper: ES, AMJ, JJM; Made critical revisions and approved final version: ES, AMJ, JJM; All authors reviewed and approved of the final manuscript.

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