Amplatzer occluders for refractory esophago-respiratory fistulas: a case series



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ABSTRACT

Background and study aims Endoscopic management of esophagorespiratory fistulas (ERF) is challenging and currently available options (stents, double pigtail, endoscopic vacuum therapy) are not very effective. We report the feasibility and efficacy of endoscopic placement of Amplatzer cardiovascular occluders for this indication.

Patients and methods This was a single-center, prospective study (June 2019 to September 2020) of all patients with non-malignant ERF persistent after conventional management with esophageal and/or tracheal stents. The primary outcome was the technical feasibility of Amplatzer placement. Secondary outcomes were clinical success defined by effective ERF occlusion and resolution of respiratory symptoms allowing oral food intake.

Results Endoscopic placement of Amplatzer occluders was feasible in 83% of patients (5/6), with a 50% (3/6) clinical success rate at 9 months. The mortality rate was 33% (2/6). **Conclusions** An Amplatzer cardiac or vascular occluder is a feasible and safe treatment option for refractory ERF, with a 50% short-term clinical success.

Introduction

Esophagorespiratory fistulas (ERFs) can be congenital or acquired, either in the context of prolonged mechanical ventilation, or mediastinal (esophageal or bronchopulmonary) tumor after surgery of chemoradiation. They are associated with poor quality of life, chronic respiratory symptoms, severe bronchopulmonary infections, and up to 27% short-term mortality[1]. Surgical management relies on soft tissue interposition or closure of the fistulous tract, with 83% efficacy, but up to 6% perioperative mortality [2,3]. Endoscopic therapy using esophageal fully covered self expandable metal stents (FCSEMS) aime at covering the esophageal defect, alone or combined with tracheal stents, only yields a 46% efficacy [1]. Indeed, stents neither completely seal the fistula, nor promote its healing. Other endoscopic techniques, such as occlusion (by clips or glue), have not shown better results [4].

A minimally invasive alternative using an Amplatzer cardiac or vascular plug septal occluder (AGA Medical, Plymouth, Minnesota, United States), intended for percutaneous vascular or septal defect closure or in cardiology, has been successfully used in endoscopy for ERF closure in humans [5,6]. The Amplatzer consists of two to three biocompatible self-expanding discs made of a nitinol mesh connected together by a thinner

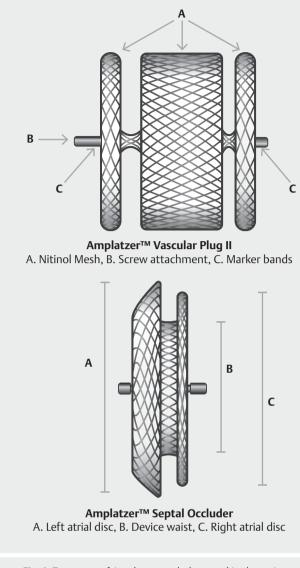


Fig.1 Two types of Amplatzer occluders used in the series.

waist, ensuring the stability of the device (**Fig.1**). We describe a case series of six patients with ERF treated with endoscopically placed Amplatzer occluders.

Patients and methods

Patients

Between June 2019 and September 2020, six patients had endoscopic closure attempts for chronic acquired ERF using an Amplatzer occluder at our center. All patients were selected after a multidisciplinary board discussion involving anesthesiologists, gastroenterologists, a pulmonologist, and gastrointestinal surgeons. The patients' written informed consent was obtained after explaining the modalities, benefits, and risks of this procedure.

Endoscopic procedures

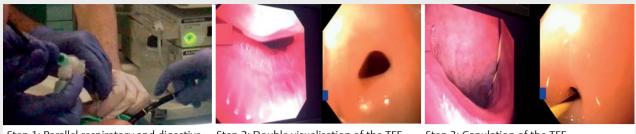
The endoscopic procedures were performed using a gastroscope (GIF-1TH190; Olympus, Tokyo, Japan) and a bronchoscope (BFQ 180, Olympus, Tokyo, Japan), with patients under general anesthesia with orotracheal intubation. Insufflation used air instead of CO2, considering the esophagorespiratory fistula. The ERF was cannulated using the cap-fitted gastroscope (12.4 mm, Model D-201-1104, Olympus, Tokyo, Japan) and a straight catheter (Guide Catheter 6 Fr 220 cm, Cook Medical) in which a guidewire (JagWire straight, 450 cm 0.035 in, Boston Scientific, Natick, Massachusetts, United States) was inserted. A bronchoscope placed in the trachea through the intubation canula allowed precise localization of the fistula and visualization of the guidewire, and facilitated correct positioning of the Amplatzer. The gastroscope was exchanged for a straight 7 Fr catheter (Destination Guiding Sheath, RSC08, Terumo), which enabled positioning of a Vascular Plug II Amplatzer (AGA Medical, Plymouth, Minnesota, United State), or a 9F catheter (Torqvue Delivery System, Abbott, Plymouth, Minnesota, United States), which enabled positioning of a Septal Occluder Amplatzer (Abbott, Plymouth, Minnesota, United States) depending on the size and shape of the ERF, under double endoscopic control. Successively, one disk was released in the trachea and one or two in the esophagus. Successful closure of the ERF was confirmed by injection of contrast medium into the esophagus showing the absence of passage in the tracheobronchial tree under fluoroscopy. The main steps of the procedure are presented in **Fig.2**.

Patients were followed up clinically to ensure that cough resolved and no new episode of pneumonia had occurred. Thoracic computed tomography (CT) with oral contrast ingestion was performed before resuming food 1 to 2 days after the procedure. A double endoscopic control 2 to 4 weeks after the procedure was performed to check the absence of migration of the device.

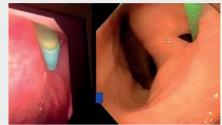
Results

Case 1 was an 80-year-old patient followed for several episodes of aspiration pneumonia and chronic cough after a prolonged intensive care admission for cardiac arrest. The patint was diagnosed with an esophago-tracheal fistula on esophagogram. At 19 cm from the dental arches, an 11-mm-wide esophago-tracheal fistula on the right side of the esophagus was seen at esophagogastroduodenoscopy (EGD). A 14-mm Amplatzer vascular plug was placed under double endoscopic control. The patient was discharged from the hospital the next day and has not had any respiratory symptoms since then. Follow-up endoscopy 4 months later showed the Amplatzer in place with no contrast medium leakage towards the trachea.

Case 2 was a 67-year-old woman who was admitted to the Intensive Care Unit after an Ivor Lewis esophagectomy following neoadjuvant chemoradiotherapy for treatment of a squamous cell carcinoma of the esophagus. The diagnosis of ERF was suspected based on recurrent aspiration with pneumonia. After confirmation of a 14-mm-wide esotracheal fistula 25 cm



Step 1: Parallel respiratory and digestive endoscopy



Step 4: Insertion of a straight catheter (left: tracheal side; right: esophageal side) side; right: esophageal side)



Step 5: Amplatzer release (left: tracheal



Step 6: Control by opacification



Step 7: Esophagoscopy at 2 weeks

Fig.2 Main steps in Amplatzer (vascular plug II) occluder placement in an esophagotracheal fistula.

from the dental arches on EGD, an esophageal FCSEMS was placed on the 12th postoperative day. The patient's respiratory symptoms persisted despite two other endoscopic procedures to replace the stent with larger (22-mm) stents, and we finally placed an 18-mm Amplatzer vascular plug under double endoscopic control 3 months after the surgery. Despite the technical success, the patient died from septic shock of pulmonary origin 3 days after the procedure.

Case 3 was a 60-year-old man who was treated with repeat surgery and chemoradiotherapy for medullary carcinoma of the thyroid and developed a 10-mm ERF, located 21cm from the dental arches. There was no evidence of local tumor recurrence. After failure of conventional management using repeat placement of an esophageal FCSEMS, a 14-mm Amplatzer vascular plug was deployed. Initially the procedure was clinically successful, with resolution of respiratory symptoms. However, cough and pneumonia recurred 9 months later and the patient reported new onset of dysphagia. EGD showed an esophageal stenosis located in the cervical esophagus related to a local tumor recurrence. EGD with a nasogastroscope permitted the stenosis to be passed and showed that the Amplatzer was still in place, but only partially occluding an enlarged and tumor-invaded fistula, with contrast leakage in the trachea. Repositioning of the Amplatzer was technically impossible due to the stenosis, and the patient was switched to long-term enteral nutrition.

Case 4 was a 71-year-old man in clinical remission from squamous cell carcinoma of the upper third of the esophagus after intial chemoradiotherapy. He was diagnosed with local and metastatic cancer recurrence 1 year later and an 8-mm ERF located 18 cm from the dental arches, which caused cough during oral food intake. Placement of a 10-mm Amplatzer vascular plug allowed him to resume a well-tolerated normal diet by mouth beginning on Day 3. However, the patient died 10 days later from massive aspiration pneumonia.

Case 5 was a 56-year-old patient who was diagnosed with an ERF on a CT performed for workup of a septic shock 12 days after an Ivor Lewis esophagectomy for esophageal adenocarcinoma. EGD showed a 14-mm fistula located 29 cm from the dental arches and bronchoscopy identified a fistula in the right primary bronchus. The fistula was initially treated with three successive esophageal FCSEMS, without fistula closure. Six months after the surgery, an 18-mm Amplatzer septal occluder was placed and allowed the patient to resume oral food intake with no respiratory symptoms. Follow-up endoscopy 5 months later found the Amplatzer in place with partial reepithelialization on the esophageal side and no contrast leakage towards the respiratory tree.

Case 6 was a 57-year-old man who underwent an lvor Lewis esophagectomy for an esophageal adenocarcinoma and was diagnosed on the 9th postoperative day with an esophago-tracheal fistula. Initial management consisted of repeated placement of esophageal FCSEMS over a 2-year period. Given the absence of fistula closure, with a remaining 25-mm esophago-tracheal fistula 24 cm from the dental arches, placement of a 24mm Amplatzer septal occluder was attempted. It did not provide a complete occlusion of the fistula, given the presence of an irregular, nodular mucosa on the tracheal side, and was immediately retrieved and replaced with another esophageal stent. The patient remained symptom-free at last follow-up, 4 months after the last endoscopic procedure. Currently, an en-

► Table 1 Characteristics of the six patients with esophagorespiratory fistulas managed with amplatzer occluders.

Case	Context of ERF	Location (cm from the DAs)	Size of the esophageal fistula (mm)	Amplat- zer type	Techni- cal suc- cess	Outcome	Follow-up (months)
1	Prolonged mechani- cal ventilation	19	11	Vascular Plug II, 14mm	Yes	Clinical success	15
2	lvor-Lewis esopha- gectomy	25	14	Vascular Plug II, 18 mm	Yes	Death 3 days after the procedure from pneu- monia	-
3	Chemoradiotherapy, no residual tumor	21	10	Vascular Plug II, 14mm	Yes	Cancer recurrence, re- currence of respiratory symptoms after 9 months	11
4	Chemoradiotherapy, residual tumor	18	8	Vascular Plug II, 10mm	Yes	Death 8 days after the procedure from pneu- monia	-
5	lvor-Lewis esopha- gectomy	29	14	Septal oc- cluder, 18 mm	Yes	Clinical success	5
6	lvor-Lewis esopha- gectomy	24	20	Septal oc- cluder, 24 mm	No	Technical failure	4

ERF, esophago-respiratory fistula; DA, dental arches.

dotracheal ablation procedure is scheduled to flatten the tracheal mucosa and correctly place an Amplatzer.

Patient characteristics and outcomes data are summarized in **> Table 1**.

Discussion

This case series found an 83% (5/6) technical success rate and a 60% (3/5) short-term clinical success rate with ERF occlusion using endoscopically placed Amplatzers. Because the recommended management of ERF relies on esophageal stents [7], we only included patients with chronic non-malignant ERF refractory to initial management with conventional stents. Thus, we selected cases with well-organized, fibrotic, large fistulas for which clipping, either directly during or after submucosal dissection [8] was not possible. Indeed, the stability of the device requires a regular-shaped and large fistula. Therefore, an Amplatzer is not likely to be an effective treatment for malignant ERF, which is best managed with esophageal, tracheal, or combined stenting; esophageal or tracheal stenoses often increase the stability of the stents [9]. Conversely, the long time between the onset of the ERF and the Amplatzer placement led to the inclusion of an extremely frail patient population, explaining the 33% (2/6) mortality rate in our series.

The use of Amplatzer occluders, initially designed for cardial septal defect occlusion, in the treatment of non-malignant refractory ERF has been reported in 10 case reports (11 patients) since 2006 [5,6,10–17]. In four of 11 cases, the ERF occurred after esophagectomy, and in four of 11 patients after prolonged mechanical ventilation. Clinical success was obtained in 45% of the cases (5/11), with heterogeneous follow-up duration, ranging from 1 month to 1 year. The mortality was high in these 11 extremely frail and malnourished patients, reaching 40% (4/11). In our case series, the mortality rate was 33%, illustrating the severity of the comorbid conditions of the patients, possibly accounting for the limited efficacy of the Amplatzer occluders.

Although Amplatzer occluders can improve the quality of life of a proportion of patients with refractory non-malignant ERF, the device suffers several limitations. First, although long-term follow-up for cardiovascular indications shows an excellent safety profile and endothelial regrowth [18], we only observed moderate epithelial growth on the esophageal side, and never on the tracheal side. This indicates that tracheal obstruction due to granulation tissue will not be an issue, but suggests that the device is only suspensive, and that ERF might recur in case of Amplatzer migration. Second, the size of the intratracheal flap, particularly with the vascular plug-type Amplatzer (for smaller-diameter fistulas), might result in airway obstruction.

Erosion of the respiratory mucosa by the Amplatzer occluder might also constitute a limitation of the use of this device, particularly in the bronchi, considering their limited diameter, because a case of fatal hemoptysis after closure of a gastrobronchial fistula was reported recently [19]. Use of novel occluders specifically designed for the endoscopic treatment of refractory ERF has been reported in a recent animal study [20]. Finally, the outcomes of the patients in case studies 3 and 4 suggest that the stability and the efficacy of the Amplatzer occluder can be compromised in cases of active malignancy: therefore, the use of these devices is probably advisable in patients with nonmalignant ERF.

Conclusions

Endoscopic occlusion of chronic, refractory, non-malignant ERF is feasible, with clinical success in half of such patients. Amplatzer occluders should be considered as a rescue therapy in selected cases that can facilitate avoidance of highly morbid surgical management.

Competing interests

The authors declare that they have no conflict of interest.

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