Case Report

Surgical Treatment for Kommerell’s Diverticulum Associated with a Right-Sided Aortic Arch and an Aberrant Left Subclavian Artery: Endovascular or Hybrid

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A right-sided aortic arch, associated with an aberrant left subclavian artery and a Kommerell’s diverticulum (KD), is a rare congenital anomaly. Surgical management is crucial before rupture or dissection occurs. Although the outcomes of the traditional open surgery have been favorable, the invasiveness of this treatment remains high. We report two cases of KD treated with endovascular and hybrid approaches.

Introduction

A right-sided aortic arch (RAA), associated with an aberrant left subclavian artery (ALSA) and a Kommerell’s diverticulum (KD), is a rare congenital anomaly. Surgical management is crucial before rupture or dissection occurs. Although the outcomes of the traditional open surgery have been favorable, the invasiveness of this treatment remains high. We report two cases of KD treated with endovascular and hybrid approaches.

Case Report

Case 1

A 53-year-old man presenting with chronic dysphasia and chest pain was admitted to our hospital. He had been treated with an unidentified complaint by a physician. Upon admission, his blood pressure was 128/62 mmHg symmetrically. His pulse was regular (72 beats/min). Neither carotid bruit nor cardiac murmur was reported. Lung examination revealed clear breath sounds. Laboratory evaluations did not show any abnormalities. Trans-thoracic echocardiography (TTE) revealed normal left ventricular function unassociated with congenital heart disease. Computed tomography angiography (CTA) demonstrated an RAA and an ALSA originating from a KD (Fig. 1A). Cross-sectional measurements of the KD revealed that the diameter of the diverticulum from the opposite aortic wall to the tip of that was 52 mm, and the diameter at the origin of the ALSA from the aortic arch was 27 mm. The aortic arch vessels arose in the following order: left common carotid artery, right common carotid artery, right subclavian artery, and ALSA. The fourth branch posteriorly passed to the esophagus from a bulbous diverticulum across the midline toward the left and upward. The left vertebral artery arose from the left common carotid artery. The patient was considered to be unsuitable for thoracic endovascular aortic repair (TEVAR) alone because...
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of endoleak with insufficient proximal landing zone of stent-graft. We decided to perform two-stage surgery with endovascular treatment after written informed consent was obtained.

To ensure revascularization of the bilateral upper extremities, the bilateral axillary arteries were first prepared to be anastomosed with an 8-mm ringed polytetrafluoroethylene graft. Through a median sternotomy, cardiopulmonary bypass was instituted in standard fashion with ascending aortic and bicaval cannulation. The KD and an adjoining ALSA in the surgical field was difficult to confirm. Under deep hypothermic circulatory arrest, both common carotid arteries were reconstructed using the arch-first technique with retrograde cerebral perfusion. The orifice of the right subclavian artery was doubly ligated. Antegrade cerebral perfusion was started through a four-branch Dacron graft. Distal anastomosis, located on the right side of the trachea, was created just behind the right common carotid artery without an elephant trunk. Subsequently, the tube graft anastomosed at the distal side was combined with the main graft to achieve cerebral perfusion using a stepwise technique. Resumption of the whole body was performed, and proximal anastomosis was achieved with the transected aorta just above the sino-tubular junction. According to the reconstruction between graft conduits anastomosed to the bilateral axillary arteries and the side branches of the four-branch Dacron graft retrieved through the thorax, total arch replacement was completed. The postoperative hemodynamics state was stable. The artificial conduit competed with the native left subclavian artery with regard to blood flow. Because there was a concern that the graft anastomosed to the left axillary artery might be occluded by a thrombus, an endovascular procedure was performed the next day. The time interval between prior total arch replacement and TEVAR was 16 h. The right femoral artery was exposed through an oblique incision. A 31 × 150-mm conformable GORE TAG stent-graft (W. L. Gore and Associates, Inc., Flagstaff, AZ, USA) was advanced in retrograde fashion across the angulated aortic arch and subsequently deployed just distal to the side branch of the four-branch Dacron graft that controlled the perfusion to the right arm. Because endoleak was detected using angiography, the 31 × 100-mm conformable GORE TAG stent-graft was additionally inserted to ensure the coverage of the ostium of the KD. The left brachial artery was percutaneously cuffed with a 6Fr introducer (Destination, Terumo, Tokyo, Japan). For the purpose of resolving dysphagia, we did not select means to completely occupy the KD by coils. To completely exclude the KD, the non-dilated origin of the ALSA was intentionally embolized with detachable coils (MICRUSFRAME; Johnson & Johnson, New Brunswick, NJ, USA) to accurately construct the frame at an early stage. Further, the space in the frame was packed with pushable coils (Tornade Embolization Coil; Cook, Bloomington, IN, USA) to reinforce the embolic effect. The completion angiography demonstrated a complete exclusion of the diverticulum. The postoperative course was uneventful without cerebrovascular failure, respiratory failure, or paraplegia. Postoperative CTA demonstrated good patency of the bypass grafts to the supra-aortic vessels, and complete exclusion of the KD by the stent-grafts and coils (Fig. 1B). The patient’s dysphagia resolved, and he is being followed up for medical treatment once a month after discharge from the hospital.

Case 2

An 81-year-old man complaining of syncope and dizziness was admitted to our hospital. He had been treated for brain infarction and hereditary tyrosinemia under adequate anticoagulation therapy with warfarin. At admission, his blood pressure was 120/70 mmHg symmetrically. His pulse rate was regular (58 beats/min). Laboratory evaluation showed no abnormalities. TTE revealed normal left ventricular function and moderate mitral regurgitation. CTA demonstrated a RAA, an ALSA originating from the KD, and a saccular left common iliac artery aneurysm (Fig. 2A). Endovascular aneurysm repair (EVAR) for a saccular left common iliac artery aneurysm...
was performed in combination with coil embolization to a left internal iliac artery. Seven months after EVAR, follow-up computed tomography showed an enlargement of the KD. Cross-sectional measurement of the KD in detail revealed that the diameter from the opposite aortic wall to its tip was 73 mm and the diameter at the origin of the aberrant subclavian artery from the aortic arch was 40 mm. Considering his medical history, the patient was deemed unsuitable for open surgery. The time interval between prior abdominal aortic aneurysm (AAA) surgery and TEVAR was 8 months. Because the proximal landing zone of the stent-graft above the KD was 2 cm in length, we decided to perform TEVAR after written informed consent was obtained.

To ensure revascularization to a left upper extremity after obliterating the KD, the bypass with a 7-mm ringed polytetrafluoroethylene graft was first prepared between the bilateral axillary arteries prior to TEVAR. The right femoral artery was exposed through an oblique incision. A 45 × 100-mm conformable GORE TAG stent-graft was deployed in retrograde fashion at the foundation of the upper descending aorta. Subsequently, the 45 × 100-mm conformable GORE TAG stent-graft was additionally inserted just distal to the right subclavian artery across the angulated aortic arch under right ventricular rapid pacing. To avoid the migration of the stent-graft, the touch-up via inflation of a balloon (tri-lobe balloon catheter, W. L. Gore and Associates, Inc., Flagstaff, AZ, USA) was performed only at the proximal side of the stent-graft. The left brachial artery percutaneously was cannulated with a 6Fr introducer. The left vertebral artery arose from the origin of the left subclavian artery adjacent to the KD. For the purpose of completely excluding the KD, we had to completely occupy the KD using detachable and pushable coils. Angiography revealed no endoleak. The completion angiography demonstrated a complete exclusion of the diverticulum. The postoperative course was uneventful without cerebrovascular failure, respiratory failure, or paraplegia. Postoperative CTA demonstrated good patency of the bypass grafts to the left subclavian artery, complete exclusion of KD, and the saccular left common iliac artery aneurysm by the stent-grafts and coils (Fig. 2B). The patient was being followed up for medical treatment once a month after discharge from the hospital. The patient was emergently admitted 21 months after TEVAR with syncope. Because electrocardiogram revealed complete atrio-ventricular block, a permanent pacemaker was implanted. The patient was transferred to the rehabilitation department of the hospital for disuse syndrome.

Discussion

KD is an outpouching aneurysm originating from aberrant subclavian arteries from the descending thoracic aorta. Embryologically, it is thought to be a remnant of the fourth primitive dorsal arch that does not regress. An RAA occurs in approximately 0.1% of the population.1) Half of the cases of RAA are associated with an ALSA. The left ligamentum arteriosus usually joins the root of the ALSA to the left pulmonary artery. RAA with an ALSA and a left ductus arteriosus sometimes forms a vascular ring. This type of RAA is seldom associated with congenital heart disease. Because this anomaly usually does not significantly compress the trachea and esophagus, it is mostly unnoticed or manifests as non-specific symptoms until adulthood. Clinical symptoms associated with the appearance of the KD, including dysphagia, wheezing, strider, shortness of breath, and chest pain, could be caused by the compression of neighboring structures as well as atherosclerotic changes of anomalous vessels. In a review of the literature concerning aneurysms arising in an aberrant subclavian artery, Austin and Wolfe reported that 19% of affected patients died from ruptures.2) Kouchoukos and Masetti reported that 20% of affected patients were associated with dissection.3) In a review of the literature concerning RAA with an aneurysm of ALSA, Cinà et al. reported that the incidence of either rupture or dissection was 53% in affected patients.4) Fisher et al. reported a case of a ruptured aneurysm arising from a right aberrant subclavian artery; the diameter at the orifice of the diverticulum measured only 2.0 cm.5) Because the entity is
Cinà et al. suggested aggressive treatment for aneurysms reported no hospital deaths with the standard surgical mortality in hybrid repair with endovascular technique. Therefore, given the higher incidence of rupture or dissection and characteristic pathological findings, it would appear appropriate to recommend early surgical management even in asymptomatic patients.

Several options for surgical, endovascular, or hybrid techniques have been preferred as treatments for KD. However, there has been no established treatment strategy because of the rarity of the disease. Open surgical treatment has been described, including interposition grafts or endoaneurysmorrhaphy through thoracotomy with a left subclavian-to-carotid transposition, graft replacement with in situ reconstruction of the aberrant subclavian artery through thoracotomy, and total arch replacement through a median sternotomy and thoracotomy. Open repair of these aneurysms has a high mortality, ranging from 16% to 18%. Kouchoukos and Masetti recently reported no hospital deaths with total arch replacement and anatomical reconstruction of the subclavian artery without endovascular technique. Furthermore, Tsukui et al. reported no hospital deaths with total arch replacement and anatomical reconstruction of the subclavian artery without endovascular technique. Idrees et al. reported no hospital mortality in hybrid repair with endovascular technique. Cinà et al. suggested aggressive treatment for aneurysms ≥3 cm in diameter. Ota et al. suggested that the diameter of aneurysms >50 mm in symptomatic patients are indications for surgery. Idrees et al. recommended surgery if the diameter as cross-sectionally measured from the opposite aortic wall to the tip of KD is at least 5 cm, and the diameter as cross-sectionally measured at the origin of the aberrant subclavian artery from the aortic arch is at least 3 cm.

Endovascular treatment for the KD is a less invasive approach than open surgery, which requires a thoracotomy or a median sternotomy. Indications for endovascular treatment should be considered with respect to clinical situations and anatomical factors. The former involves clinical symptoms, hemodynamic state, comorbidities, and past medical history. The latter is related to the characteristic configuration of a sharply-curved distal arch, a distorted descending aorta, and anomalous supra-aortic branches. Total endovascular repair with a branched device has been reported to further reduce the invasiveness of treatment. Reconstruction of the left subclavian artery would be of utmost importance to prevent arm claudication in younger patients, subclavian steal syndrome in the elderly, and vertebrobasilar insufficiency. However, this remains controversial.

In Case 1, because there was no distance between the right subclavian artery and the top of the sharply-curved distal arch, the endovascular approach was not feasible because of the greater occurrence of endoleaks and migration. A thoracotomy provides satisfactory exposure of the ascending aorta, transverse arch, and descending thoracic aorta. However, this approach may give rise to lung injury by thoracotomy and brain infarction with difficulty in air removal during hypothermic circulatory arrest. At our hospital, the 30-day postoperative mortality rate of the 31 patients who underwent the aortic arch surgery between January 2014 and December 2018 was 3.2%. We adopted a hybrid repair, including initial total arch replacement without elephant trunk and subsequent endovascular exclusion of the KD with coil embolization of the left subclavian artery. Especially, when the graft inserted into the sharply-curved distal arch as an elephant trunk is transformed, it is anticipated to be difficult to deliver the stent-graft to its intended position in the subsequent TEVAR. Securing the distance between the back-most branch of the four-branch Dacron graft and the distal anastomosed site is crucial. We did not use the frozen elephant trunk because there was a possibility of an unexpected occurrence of paraplegia. The stent frame was designed as a stent-graft used as a frozen elephant trunk for the inner side and a stent-graft in TEVAR for the outer side. Endoleak might occur when the compatibility of overlapping varying characteristic stent-grafts is poor. There is a possibility of recurrent dysphagia because of tracheal and esophageal compression by the retained KD, so close follow-up of endoleak of the stent-graft and recurrent symptoms by the retained KD would be required in the future.

In Case 2, because the patient underwent EVAR with coil embolization to the left internal iliac artery before 8 months, it was predicted to increase the risk of spinal cord ischemia after TEVAR. Schlösser et al. reported that prior infrarenal AAA repair is associated with increased risk of spinal cord ischemia after TEVAR compared with patients without prior AAA surgery. It was mandatory to perform preventive management during TEVAR, including deployment of stent-grafts to make the length of aortic coverage as short as possible, maintaining the collateral network supplied from the covered left subclavian artery with revascularization, and meticulous attention to avoid intra-operative hypotension. Furthermore, we had to take care to deploy the stent-graft in a bird beak configuration on the lesser curvature of the arch. To strictly deploy stent-grafts, we usually perform right ventricular rapid pacing, when the proximal landing of the thoracic stent-graft is located in the range of zone 0 or zone 1 and supra-aortic branches anomalously exist. Cerebrospinal fluid drainage was omitted because of anticoagulation therapy. The technique, which bunched the next stent-graft upward toward the stent-graft deployed at the upper descending aorta as a
foundation, was thought to be useful to avoid the migration of the stent-graft with high accuracy.

Conclusion

Although the technical advancement of stent-grafts and the development of treatment strategies have been remarkable, long-term outcomes of endovascular or hybrid treatment for the KD are uncertain. Our findings in the two cases indicate that to ensure optimal treatment outcomes, it is mandatory to individually evaluate current treatment strategies in accordance with clinical symptoms, hemodynamic state, comorbidities, past medical history, and anatomical features of KD.

Disclosure Statement

All authors have no conflict of interest.

Author Contributions

Study conception: AM
Writing: AM
Critical review and revision: all authors
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