The First Case of Successful Stenting of the Dissection of the Ascending Aorta and the Aortic Arch that Occurred During Surgical Correction of the Supravalvular Aortic Stenosis in a Child with the Williams Syndrome

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Abstract

Congenital supravalvular aortic stenosis (SVAS), in vast majority of cases is a manifestation of Williams syndrome. Surgical correction of this pathology is a "gold standard" for treatment in these patients. One of the most dangerous potential complications in surgical repair of this disease is acute dissection of the ascending aorta, arising in 0.7% of cases. This complication can be attributed to both surgical errors and or due to the anatomical features of the aortic wall. We report on a pediatric patient with Williams syndrome, 2.3 yr, 11.9 kg, who underwent surgical repair for supravalvar aortic stenosis using the Doty technique. Aortic dissection was diagnosed using aortography. This was managed by implanting two Valeo stents.

Key Words

Williams syndrome • Aortic dissection • Stenting

Introduction

Congenital supravalvular aortic stenosis (SVAS), in the vast majority of cases, is a manifestation of Williams syndrome or other pathological condition associated with mutation of the 7q11.23 gene [1-4]. In patients with Williams syndrome or SVAS, other obstructive vascular lesions are often present such as coarctation of the aorta and peripheral stenoses of the pulmonary arteries. Patients with Williams syndrome also often have a bicuspid aortic valve and mitral valve prolapse, as well as various heart rhythm disturbances [6-8].

SVAS is a complex pathology that may present in a wide variety of clinical and morphological forms. The most common form is aortic lumen obstruction localized directly above the aortic valve [5, 9]. The severity of stenosis is the main factor determining the severity of hemodynamic burden and thus the clinical manifestation of this pathology. In most cases, the clinical picture of the disease manifests during childhood or adolescence, and most patients require surgical correction of the defect, which is performed under cardiopulmonary bypass [1, 10, 11].

Although surgery is safe and successful in most cases, there is the rare potential complication of dissection of the ascending aorta after its surgical reconstruction. This complication can be attributed to both surgical errors and anatomical features of the aortic wall. The incidence of this complication is 0.6% [12-13]. Here, we describe a child with Williams syndrome...
and SVAS who underwent successful hybrid stenting of an extensive dissection of the ascending aorta extending to the aortic arch.

**Case Presentation**

Patient K. was 2.25 years and weighed 11.9 kg. He was diagnosed with Williams syndrome with severe SVAS. Echocardiography revealed the presence of concentric hypertrophy of the left ventricle with normal ejection fraction of 68%. The peak gradient across the ascending aorta was 100 mmHg. He also had moderate stenosis of the branch pulmonary arteries, with a peak gradient across the entire right ventricle outflow tract of 11 mmHg. He also had a patent foramen ovale. His echocardiogram showed left axis deviation and left ventricle hypertrophy. Chest X-ray demonstrated cardiomegaly, mainly due to left heart hypertrophy, with a cardiothoracic ratio of 67%.

Multislice computed tomography was performed to clarify the anatomy and determine the localization of the lesion and severity of the stenosis (Figure 1). We observed severe aortic stenosis at the level of the sino-tubular junction measuring 4.5 mm. The isthmus and aortic arch diameters were 6 mm. Brachiocephalic vessels originated from separate origins, with brachiocephalic trunk, left common carotid artery, and left subclavian artery diameters of 3.0, 2.8, and 3.8 mm, respectively.

Due to his clinical symptoms of fatigue associated with severe stenosis and left ventricle hypertrophy, we decided to perform surgical repair of the ascending aorta using the Doty technique. Median sternotomy was performed, and under cardiopulmonary bypass with moderate hypothermia (28°C), the ascending aorta was opened via a longitudinal incision toward the non-coronary sinuses of Valsalva almost to the aortic valve annulus. The second incision crossed the stenotic sinotubular zone, forming a reverse “Y” shape in the direction of the right coronary sinus anterior to the intracoronary commissure. Visually, we noticed a thickening of the aortic wall to a diameter of 3.5 mm. A reverse “Y”-shaped xenopericardial patch was made and fixed to the edges of the aortic incision using a premilene suture starting from the right coronary sinus. The suture site was reinforced with medical hemostatic glue [14].

The patient was weaned off bypass without much difficulty. However, there was a significant hemodynamic difference in systolic pressure between the ascending aorta and the radial arteries to the right and left of 110 mm Hg. Dissection of the aorta was suspected. To confirm this, using a mobile angiographic unit (OEC 9900, GE Healthcare, Chicago, IL, USA), we performed ascending aortography using the right femoral arterial access. The systolic pressure gradient between the ascending and descending aorta was 177 mm Hg. Aortography revealed the presence of aortic dissection, with an intimal flap distal to the aortic patch on the ascending aorta with extension to the aortic arch and brachiocephalic vessels (Figure 2).

Given a high risk of surgical correction under cardiopulmonary bypass, a hybrid approach was proposed. We decided to perform stenting of the dissected part of the aorta. A 6-F Mullins sheath (Cook Medical, Bloomington, IN, USA) was placed in the ascending aorta. A standard diagnostic guidewire was exchanged for a 0.035” Amplatz super-stiff guide wire with a 1-cm soft tip (Boston Scientific, Marlborough, MA, USA) to deliver the stent into the ascending aorta so that it would completely cover the zone of dis-
section to the point of origin of the brachiocephalic vessels. A 18-mm Valeo stent (Bard, Murray Hill, NJ, USA) was attached to a 8-mm balloon deployed to 10 atm. After stent implantation, however, there was no significant change in systolic pressure gradient. Therefore, a second 26-mm Valeo stent on a 8-mm balloon was implanted covering the whole surface of the aortic arch. Repeat aortography revealed proper implantation of both stents covering the entire zone of dissection. Patency of the brachiocephalic arteries was preserved (Figure 3). Invasive pressure measurement after stent implantation showed minimal residual systolic pressure of 15 mm Hg. The child left the operating room on adrenaline at a dose of 0.1 μg/kg/min.

The child was extubated after 20 hours. Anticoagulant therapy was initiated using heparin at a rate of 200 U/kg/day for 3 days followed by aspirin at a dose of 50 mg/day. The patient was discharged home on postoperative day 13. Unfortunately, the parents refused postoperative follow-up consultation in our center due to a stated good clinical status of their child and distant place of residence.

**Discussion**

Surgical correction of SVAS in patients with Williams syndrome is safe and effective. However, a rare acute complication of dissection of the ascending aorta can lead to an unfavorable prognosis. There are no clear recommendations or algorithms for action in cases of dissection. In adult patients with acute dissection, prosthetic material can be used to repair the dissection. In children, however, this is more difficult and sometimes not feasible. Therefore, in such cases, using endovascular techniques to eliminate acute aortic dissection is potentially a promising solution. We did not find descriptions of similar clinical cases in young children in the literature. Available reports discussed planned endovascular and hybrid interventions to eliminate residual stenosis of the ascending aorta and aortic arch after previously performed reconstructive surgical interventions [15-16].
Despite good immediate results in our case, a future increase in systolic pressure gradient is possible due to growth of the child and neointimal proliferation [15, 16]. This complication often occurs after stenting of vessels, especially in children with Williams syndrome or other genetic defects accompanied by connective tissue dysplasia, which are characterized by a tendency to increased proliferation of the endothelium [17]. However, in our case of hybrid management of an urgent obstruction of the ascending aorta and aortic arch, we used Valeo stents, which have the capacity for further expansion to an adult size if and when needed [18]. Therefore, our case demonstrates the possibility of intraoperative diagnosis and elimination of a formidable complication such as acute aortic dissection using endovascular technologies in pediatric cardiac surgery.

Conflict of Interest

The authors have no conflict of interest relevant to this publication.

References


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