

Nightmare Case of Complex Neonatal Truncus Arteriosus

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Abstract

Neonatal management of severe quadricuspid truncal valve insufficiency is very challenging with high risk of postoperative mortality. Aortic valve reconstruction is the unique rescue and should be considered and previously planned in case of truncal valvuloplasty failure. Many extracardiac malformations can be associated to truncus arteriosus, therefore a cardiac and chest computed tomography scan must be performed before the elective surgical repair.

Key Words: neonatal truncus arteriosus; quadricuspid valve; severe regurgitation; aortic valve reconstruction; extrinsic airway compression.

Case report

A one-month-old term boy, weighing 3 kg, was referred to our institution with the diagnosis of truncus arteriosus.

Echocardiography revealed a TA type 1-2 according to Van Praagh classification (6-7); two symmetric ventricles, large outlet VSD, large truncus arteriosus, right aortic arch, 2 good sized pulmonary arteries and quadricuspid truncal valve with severe insufficiency and moderate stenosis.

An elective primary surgical repair according to Rastelli Procedure was performed; VSD closure, right ventricle (RV)-pulmonary arteries tube and truncal valve repair by tricuspidization.

The immediate postoperative echocardiography showed a persistent severe aortic regurgitation with moderate stenosis. We decided to return immediately to operating room for a new truncal valvuloplasty. An aortic valve reconstruction with fixed pericardium using Ozaki technique was performed [8].

The echocardiography performed after the second aortic valve reconstruction was found no ventricular dysfunction, mild to moderate aortic regurgitation, no aortic stenosis and a good coaptation of the new leaflets

Following a failure of weaning of mechanical ventilation after 2 weeks of surgery, a chest injected CT-scan was performed and showed an external compression of the trachea and the left and right main stem bronchus by a dilated ascending aorta (figure 2).

An aortopexy was performed and the baby was extubated 5 days after.

6 months later, the baby still free of respiratory signs with a good statural growth. The control of echocardiography showed a mild to moderate aortic regurgitation with no stenosis.

Discussion

Truncus arteriosus is considered as a congenital heart defect reported with many cardiac and extracardiac malformations.[9,10]

The management of aortic valve disease particularly in the neonatal period, still very challenging. In case of aortic valvuloplasty failure in infant, a rescue therapy is not achievable by a bioprosthesis substitution [11]. A mechanical valve replacement is not possible technically in neonates and a Ross procedure is a high-risk operation [12] and not possible in truncus arteriosus.

Despite improved surgical management of patients with truncus arteriosus, truncal valve regurgitation remains a risk factor for early and late morbidity and mortality [13].

Neonatal period and many others factors as dysplastic TV, severe TVI, TV stenosis, abnormal number of leaflets and dilated aorta still affect significantly the outcome of truncal valve repair.

Tricuspidization techniques for quadricuspid aortic valve repair was originally described by Imamura *et al.* [14] with a good result on TVI but, in their study, there was just one neonate among those who had a quadricuspid truncal valve with severe TVI without associated valvular stenosis. Myers *et al* [15] have shown that surgical creation of a tricuspid truncal valve seems to provide a best outcomes but, neonatal repair of truncal valve was a significant independent predictor of reoperation.

More recently, Cuttone et al, [16] have shown the feasibility and a good early postoperative result of aortic valve reconstruction following Ozaki procedure in children.

Currently, in neonatal period with severe truncal valve disease, the data available on techniques and outcomes still very limited.

Aortic valve reconstruction as described by Ozaki et al. [8] remains the unique option in case of failure of truncal valve repair in newborn. Questions remain regarding the durability of the fixed pericardium used and the late outcomes.

In this case, we report a tracheobronchial compression by a dilated ascending aorta results in ventilator dependency. It was diagnosed by a computed tomography (CT)

After TA repair.

The airway compression (AC) commonly occurs in patients with anomalies of the aortic arch system and vascular rings. However, several conditions cause AC in preoperative congenital heart diseases cases, including a left-to-right shunt causing dilatation of the pulmonary arteries, truncus arteriosus [17], tetralogy of Fallot with an absent pulmonary valve, and a malposition or dilated aorta

This associated abnormality to TA is rarely described in the literature [5, 17, 18], Its early detection and management is essential to reduce further morbidity and mortality.

Conclusion:

Aortic valve reconstruction should be considered in newborn with severe truncal valve disease. Preoperative meticulous analysis of truncal valve should be performed especially with 3D echocardiography. An aortic CT must be done systematically to eliminate any extrinsic airway compression and to carry out in the same time of TA repair.

Declaration of patient consent

The authors certify that they have obtained all appropriate patient consent forms. In the form, the legal guardian has given his consent for images and other clinical information to be reported in the journal. The guardian understands that names and initials will not be published and due efforts will be made to conceal patient identity, but anonymity cannot be guaranteed.

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Conflicts of interest

There are no conflicts of interest.

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