Surgical correction of total anomalous pulmonary venous return in an adult patient

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Abstract

Background: Total anomalous pulmonary venous return (TAPVR) is rare congenital heart disease. Most TAPVRs require surgical corrections in the neonatal period and survival to adulthood without surgical correction is extremely rare. Most untreated patients with large atrial septal defects and no pulmonary venous obstruction have pulmonary vascular damage from pulmonary over circulation.

Case presentation: 44-year-old TAPVR patient admitted to our medical center. A snowman-shaped heart, including cardiomegaly and an increase in pulmonary blood flow, was seen in the chest X-ray. A large-sized (around 3 cm) atrial septal defect with dilated right atrium, right ventricle, and pulmonary artery was detected on echocardiography. Heart computed tomography was performed for further evaluation, and supra-cardiac type TAPVR without any obstructive lesion was identified.

Conclusions: TAPVR in an adult patient is extremely rare, and this patient was treated successfully with surgical correction and is doing well. A sinus rhythm and mild mitral valve regurgitation have remained during 2.5 years of outpatient follow-up.

Keywords: TAPVR, Adult congenital heart surgery, Vascular anomalies, Complex congenital heart disease

A 44-year-old man was admitted for acute onset palpitations and dyspnea. On physical examination, clubbed fingers were observed. The patient's oxygen saturation was 89% in room air, and atrial tachycardia was seen on electrocardiography. Blood chemistry evaluation showed elevated liver enzymes (aminotransferase/alanine aminotransferase: 2060/1649 U/L, total bilirubin/ direct bilirubin 4.41/0.84 mg/dL) and amino-terminal pro-brain natriuretic peptide (NT-proBNP) levels up to 6789 pg/mL. A snowman-shaped heart, including cardiomegaly and an increase in pulmonary vasculature, was seen in the chest X-ray (Fig. 1). Cardiomegaly on chest X-ray was detected 23 years ago and noted in his past medical history. However, there was no follow-up.

A large-sized (around 3 cm) atrial septal defect (ASD) with dilated right atrium, right ventricle, and pulmonary artery was detected on echocardiography. In addition, a D-shaped left ventricle (LV) and severe mitral valve regurgitation were also detected. Sinus rhythm recovered after digitalization and diuretic administration. The mitral valve regurgitation was improved to a mild degree on echocardiography, and the liver enzymes were also normalized. Cardiac computed tomography was performed for further evaluation, and supra-cardiac type total anomalous pulmonary venous return (TAPVR) without any obstructive lesion was identified. The pulmonary venous drainage course, including the superior vena cava was severely dilated. However, the left atrium and left ventricle sizes were adequate (Fig. 1). The pulmonary arterial pressure was 43/22 mmHg (Mean: 29 mmHg) on right heart catheterization (Table 1).

We performed a direct anastomosis of the pulmonary venous confluence to the left atrium and ASD patch.
closure by the transverse sinus approach without total circulatory arrest. The vertical vein was ligated. The cardiopulmonary bypass (CPB) and aortic cross-clamp times were 147 and 61 min, respectively. We also checked the mitral valve morphology. The mitral valve showed diffuse leaflet thickening at the anterior and posterior valves. However, the coaptation margin of the valve leaflet was good. Thus, we did not perform mitral valve surgery. The patient weaned off CPB easily. On postoperative TEE, no turbulence was observed across the anastomosis, and the mitral valve regurgitation was mild. The patient was extubated on postoperative day (POD) 0 and transferred to the general ward on POD 1. He was discharged from

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**Table 1 Preoperative cardiac catheterization data**

| Site                  | O₂ saturation (%) | Pressure systolic/diastolic pressure (mean), mmHg |
|-----------------------|-------------------|--------------------------------------------------|
| Superior vena cava    | 90.7              |                                                  |
| Right atrium          | 85.4              | 12/4 (7)                                         |
| Right ventricle       | 88.7              | 43/8 (20)                                        |
| Pulmonary artery      | 88.7              | 43/22 (29)                                       |
| Left atrium           | 89.6              | 12/5 (7)                                         |
| Aorta                 | 90                | 100/60 (73)                                      |
hospital on POD 8 without any symptoms. The patient is doing well. The cardiac rhythm remains in sinus rhythm and transthoracic echocardiography show mild mitral valve regurgitation after 2.5 years of operation (Fig. 2).

Discussion
TAPVR is a rare anomaly accounting for only 1.5% to 2.2% of congenital heart disease [1]. TAPVR is a cardiac malformation in which all pulmonary veins on either side are not connected to the left atrium, but directly to the somatic vein or the right atrium. Almost all patients with TAPVR present with symptoms of heart failure in the neonatal period, which require early surgical correction. It is unusual to encounter a adult patient with TAPVR due to low survival rate in untreated cases.

According to Talwar et al. [2] in group of 27 patients who underwent surgery for TAPVR at the age of 15–48 years, Unobstructed pulmonary venous drainage, a larger inter-atrial communication, maintenance of vascular resistance and pulmonary pressure closer to normal limits was important factors for survival in unoperated patients with TAPVR. Like this authors believes, Our patient also had mild pulmonary hypertension, supracardiaco type TAPVR with non restrictive ASD and absence of pulmonary obstruction. This condition is major factors for long term survival in adult TAPVR patients.

If patients not had pulmonary vascular disease, most patients are able to complete repair. But with severe pulmonary artery hypertension, Right heart decompression such as small interatrial communication, partial closure of the septal defect and unidirectional valve patch, should be performed [3]. All these techniques were left to permit right-to-left shunting during pulmonary hypertensive crisis. Alternate surgical option had been reported by India group about test protocol for selective ligation of vertical vein [4]. A patient vertical vein functions like an atrial septal defect, when too small noncompliant atrium to accommodate blood flow after surgical repair. But this technique was potential hazard to left-to-right shunt. Especially in pulmonary hypertensive crisis with pulmonary arteriolar constriction, Left-to-right shunt results in lower preload of left ventricle which exacerbates cardiac output reduction.

There are few adult TAPVR patients worldwide. The cases reported previously showed a favorable prognosis in the postoperative phase. However, approximately 10% to 15% of the patients have evidence of late pulmonary vein obstruction, which tends to be recurrent and progressive [5]. In the three years of follow-up after surgery, this patient did not show any complications, including pulmonary venous obstruction or mitral valve regurgitation. However, long-term surveillance and monitoring are required.

Abbreviations
NT-proBNP: NT-amino-terminal pro-brain natriuretic peptide; ASD: Atrial septal defect; LV: Left ventricle; TAPVR: Total anomalous pulmonary vein return; CPB: Cardiopulmonary bypass; POD: Postoperative day.

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Author contributions
WJ was the operator of this case and the corresponding author of this article. YB was a major contributor in writing the manuscript. KS made critical revisions to the manuscript and collect data. All authors read and approved the final manuscript.

Fig. 2 Postoperative computed tomography (CT). A Chest X-ray at discharge. B, C Coronal view on chest CT at 2.5 years after surgery. The PV was connected to the LA. PV, pulmonary vein; LA, left atrium. *ligated vertical vein
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Declarations

Ethics approval and consent to participate
All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

Competing interests
The authors declare that they have no competing interest.

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