Images of Anomalous Origin of The Right Coronary Artery From the Pulmonary Artery

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ABSTRACT

Anomalies of the coronary arteries -- especially their abnormal origin from the pulmonary artery (ARCAPA) trunk -- are among the least common. They're also the most dangerous of congenital heart defects with an incidence of 0.002% in the general population [Williams 2006]. The diagnosis exceedingly is difficult because anatomical abnormalities of the coronary arteries are subtle. We present a case of an anomalous origin of the right coronary artery.

INTRODUCTION

A 52-year-old male patient was referred to our hospital for gallbladder stones without symptoms associated with the cardiovascular system. Physical examination and X-ray revealed an enlargement of the cardiac boundary.

RESULTS

The patient’s initial electrocardiography (ECG) demonstrated sinus tachycardia without acute changes. His troponin I index was normal. Computed tomography angiography (CTA) examination was then used for more detailed evaluation. As shown in Figure 1, the left and right coronary arteries and their branches markedly were tortuous and thickened. The right coronary artery had an anomalous origin from the pulmonary artery, and the diameters of the left and right coronary artery orifice considerably were enlarged. The right coronary artery markedly was tortuous with beaded changes in the cross-section view, and collateral circulation had formed between the right coronary and left anterior descending coronary artery.

Transesophageal echocardiogram indicated a normal function of the left and right ventricles, and the ejection fraction was estimated by the range of 64.5% to 70%. There were no other structural cardiac abnormalities. The thickness of the left ventricular posterior wall and septal thickness was mildly increased. Multiple cardiac fistulas from the left and right coronary arteries ran into the right ventricular outflow tract, left atrium, left ventricle, and right atrium, respectively (Figure 2). The maximum flow rate of the right ventricular outflow tract was 2.7m/s. The maximum pressure gradient (PGmax) was 29mmHg, and the right ventricular systolic pressure was 35mmHg.

DISCUSSION

When it comes to anomalous coronary artery disease, 22% of patients have other congenital heart anomalies, such as aortopulmonary window, tetralogy of Fallot, and ventricular septal defect (VSD) [Maluf 1997]. The diagnostic modalities for
ARCAPA include cardiac computed tomography, cardiovascular magnetic resonance, and coronary angiogram, all of which comprise the Multi-Slice Computed Tomography (MSCT) coronary angiography, which is more reliable and plays an important role in accurate diagnosis. Patients with ARCAPA mostly are asymptomatic, but potentially life-threatening, surgical treatment of ARCAPA is highly recommended [Dodge-Khatami 2002]. The aim of surgical corrections is to eliminate “coronary steal” and other coexisting cardiac malformations. Since the risk of sudden cardiac death decreases with age, symptomatic relief is the main goal for treatment [Kautzner 1996].

In our patient, there were no symptoms of severe “coronary steal” and no other heart anomalies. Therefore, the patient refused to undergo surgical treatment and decided to return for a timely outpatient follow-up.

Figure 2. The views of transthoracic echocardiography. (A) A large fistula about 17.2mm diameter was seen at the left anterior wall of the right ventricular outflow tract near the pulmonary valve ring; (B) The long axis view of aorta showed that a 6.3mm diameter fistula was found in the left atrium near the mitral valve ring in the posterior wall of the left atrium; (C) Multiple abnormal blood flows were observed in the left ventricle, with a larger one about 2.2mm; (D) A 5.4mm wide fistula was observed on the right side wall of the right atrium near the tricuspid valve ring. LA = left atrium; LV = left ventricle; RA = right atrium; RV = right ventricle; TV = tricuspid valve. (Shown by arrow).

REFERENCES


