Quadricuspid aortic valve: case report and literature review

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ABSTRACT

Introduction: Quadricuspid aortic valve is a rare anatomical abnormality, its incidence ranging from 0.008% to 0.033%. Case Report: We describe the case of a female patient with a quadricuspid aortic valve. The direct visualization showed a type E quadricuspid aortic valve. After complete excision of the valve, it was replaced by a 23mm biological prosthesis. Postoperative recovery was uneventful. Discussion: Since the first case, described in 1862, fewer than 200 cases have been diagnosed so far in world literature. Given its frequent association with other cardiac malformations, it is important that physicians seek their preoperative diagnosis.

Keywords: Aortic valve. Congenital abnormalities. Diagnostic Techniques, Surgical.

INTRODUCTION

Quadricuspid aortic valve is a rare anatomical malformation, with few cases described in the literature. Its incidence ranges from 0.008% to 0.033% [1]. It may be presented as an isolated cardiac malformation, but in some cases there are other heart defects, such as coronary abnormalities (mainly ostial anomalies), atrial septal defect, or ventricular septal defect [2]. Diagnosis by transthoracic echocardiography is difficult, with reports of diagnosis only with intraoperative direct observation or by transesophageal echocardiography (either pre-op or intra-op) [3].

We describe the case of a 49-year-old patient diagnosed with aortic regurgitation, whose diagnosis was made only intraoperatively.

CASE REPORT

We present a case of a 49-year-old Caucasian female, who had been complaining of fatigue due to moderate exertion for 8 months, which had worsened to minor exertion for the last 2 months.

The patient had a previous diagnosis of aortic valve dysfunction (aortic insufficiency) with regular visits to her personal cardiologist for the last twelve years. During the follow-up period, the patient had remained asymptomatic until 8 months ago, when she started to complain of fatigue and dyspnea during moderate exertion. Two months before admission, dyspnea had begun to present on light exertion, associated with vertigo and sporadic tachycardia. There were no reports of chest pain, syncope, cough, expectoration, fever, or weight loss in the related period.

A transthoracic echocardiogram was performed, showing a tricuspid aortic valve with significant aortic regurgitation and with mild left ventricular dysfunction. A coronary angiography revealed normal coronary arteries without obstruction.

The patient underwent surgical treatment for aortic valve replacement under general anesthesia and access through median sternotomy. Cardiopulmonary bypass was initiated with cannulation of the aorta and inferior and superior vena cava. When moderate hypothermia at 28°C was achieved, aortic cross clamping was performed. After oblique aortotomy, a quadricuspid...
The presentation of four cusps in the aortic valve is a rare, probably underdiagnosed condition. From the various congenital abnormalities affecting this valve, the most frequent is the bicuspid aortic valve, occurring in approximately 2 percent of the general population [4]. Although very infrequent, an interesting piece by Leonardo da Vinci from between 1512 and 1513 was found, depicting a quadricuspid aortic valve [5]. The first case described in medical literature was presented in 1862 [5,6], and since then, fewer than 200 cases have been described so far [7,8].

In 1973, Hurwitz and Roberts [9] proposed an anatomical classification of quadricuspid semilunar valves, based on anatomic studies of 8 quadricuspid pulmonic valves and 2 quadricuspid aortic valves, after examination of 6,000 necropsies. The authors encountered seven different anatomic types, which were classified from A to G, in terms of the size of each cusp. The case described in our report could be classified into type E, with one bigger cusp and three smaller cusps. Feldman et al., in 1990 [1] reported an incidence of only 8 cases of quadricuspid aortic valve in more than 60,000 echocardiographic examinations analyzed, with an incidence rate of 0.013%. Often, due to the low incidence of this valvar anomaly, echocardiogram is unsuitable for definitive diagnosis.

Much like in the case described by Denker and Stagmo [10], our patient did not obtain a preoperative diagnosis via transthoracic echocardiography. It is probable that the unusual character of this anatomical variation leads to its low diagnosis through imaging methods, and even in some less careful necropsy analysis.

The presence of a quadricuspid aortic valve, although rare, may be associated with other cardiac malformations, which may require proper surgical correction. Therefore, it is important that physicians consider the possibility of such diagnosis prior to surgical intervention.
