CASE REPORT

Percutaneous transcatheter closure of aorto-right ventricular fistula using the Amplatzer duct occluder

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Abstract The occurrence of aorto-right ventricular fistula after an aortic valve replacement is rare. If it remains untreated, this condition can result in heart failure and could thus significantly compromise patient survival. Surgical closure is the treatment of choice; however, transcatheter closure has been attempted with relatively acceptable results. Here, we report on a patient who presented with heart failure with an aorto-right ventricular fistula that was present for nine years following aortic valve replacement. Successful transcatheter closure of the fistula with the use of the Amplatzer duct occluder was performed, suggesting that the percutaneous approach is an efficient technique for the treatment of such fistulae.

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1. Introduction

Aorto-cardiac fistulas are relatively rare. They are often a complication of surgery, trauma or infective endocarditis. A 2 × 2 cm contact surface between the aorta above the right coronary cusp and the right ventricular outflow tract is the target area for an aorto-right ventricular fistula.1

2. Case presentation

A 73-year-old woman was admitted to our hospital for a scheduled percutaneous transcatheter closure of an iatrogenic aorto-right ventricular fistula. The patient had a history consistent with dyspnea at rest over the past three weeks. Nine years ago, she underwent aortic valve replacement with a bileaflet mechanical prosthetic valve.
for aortic stenosis of rheumatic etiology (Fig. 1). In the first post-operative echocardiogram, a communication between the aortic root and the right ventricle with continuous flow (Fig. 2C) was diagnosed. The patient was regularly followed up with for nine years, with no evidence of clinical or echocardiographic deterioration.

Upon admission, the patient was on atrial fibrillation; the physical examination was compatible with right heart
failure. Transthoracic (Fig. 2A) and two-dimensional (2D) (Fig. 2B)/three-dimensional (3D) (Fig. 2D) transesophageal echocardiography confirmed the communication from the aorta to the right ventricle at the level of the right coronary cusp. The right ventricular systolic pressure was over 40 mmHg. Aortography demonstrated the 4 mm-wide aorto-right ventricular fistula (Fig. 1A). Therefore, we performed percutaneous closure of the fistula using fluoroscopy, under regional anesthesia and sedation, with the use of an Amplatzer duct occluder (Fig. 1 — yellow arrow). In our case, and, to our knowledge, the first report, the aorto-right ventricular fistula was accessed in a retrograde manner: that is, the guide-wire of the duct occluder was promoted directly through the right ventricle to the fistula and the aortic root (Fig. 1 — white arrow). Right ventriculography (Fig. 1E) was performed to assess the occluder position.

Repeat aortography (Fig. 1F), transthoracic (Fig. 2E) and transesophageal (Fig. 2F) echocardiography showed an acceptable occlusion of the aorto-right ventricular fistula with minimal residual flow. Symptoms rapidly improved and the patient was discharged with explicit instructions and medication.

3. Discussion

Aorto-right ventricular fistulas are very rare. Most of cases are caused by rupture of the sinus of Valsalva or endocarditis but are also less frequently caused by trauma or in association with valve replacement. In the case of aortic valve replacement, iatrogenic injury of the septum, aggressive debridement of the annulus and inappropriate suturing of the mechanical valve onto the membranous portion of the ventricular septum may be reasons for fistula formation. There are also reports of primary suture failure (dehiscence) and infective endocarditis associated para-valvular leakages. Aortic dissection as a cause of aorto-right ventricular fistula has also been reported in the literature. Recently, the formation of such a fistula as a complication of transcatheter aortic valve implantation has been reported, indicating erosion by the mechanical prosthetic valve struts as the possible mechanism for fistula formation.

The clinical presentation of an aorto-right ventricular fistula depends on its etiology as well as on the size of the shunt. Upon clinical examination, a continuous murmur, a thrill or the combination of both may be present. Patients with a small aorto-right ventricular fistula may be completely asymptomatic. However, when the shunt is significant or comorbidities cause decompensation of a medium-size fistula, as in our case, the patient may present with manifestations of heart failure. Although aortography is the gold standard for the diagnosis of an aorto-right ventricular fistula, non-invasive modalities such as contrast-enhanced computed tomography, magnetic resonance imaging and especially transthoracic and/or transesophageal echocardiography are currently preferred.

The decision to close an aorto-right ventricular fistula depends on the size of the shunt and the clinical characteristics of the patient. Indications for fistula closure include right and/or left heart failure, pulmonary hypertension and hemolytic anemia. Surgical closure remains the treatment of choice; however, transcatheter closure has been attempted with relatively acceptable results. The devices used for a percutaneous transcatheter closure of aorto-right ventricular fistulas include: a coil, an Amplatzer atrial septal occluder, an Amplatzer muscular ventricular septal defect occluder, an Amplatzer vascular plug and, as in our case, an Amplatzer duct occluder.

In conclusion, aorto-right ventricular fistulas are an important complication of aortic valve replacement. Closure of such fistulae with the Amplatzer duct occluder in selected patients has been sporadically reported but could prove to be a useful procedure.

Conflicts of interest

None.

References