
Giant Aneurysm in a Coronary–Pulmonary Artery Fistula

Aneurisma gigante de fı́stula coronariopulmonar

To the Editor,

A 67-year-old woman, with no cardiovascular risk factors and asymptomatic from the cardiologic point of view, was referred to our unit for the study of a cystic mass adjacent to the cardiac outline, detected incidentally on a chest radiograph (Fig. 1). The transthoracic echocardiogram showed a large spherical mass with a hyperechogenic wall and heterogeneous content, situated at the level of the atrio-ventricular sulcus, adjacent to the aortic root (Fig. 1, Video 1). Multislice computerized tomography and coronary angiography showed a fistula between the right coronary artery and the pulmonary artery, which originated near the ostium of the right coronary artery and...
Figure 1. The left side of the image is the chest radiograph of the patient, showing a paracardiac cystic mass. The transthoracic echocardiogram shows a heteroechoic mass adjacent to the aortic root.

Figure 2. Multidetector computed tomography and invasive coronary angiogram: fistula of the right coronary artery to the pulmonary artery with a giant thrombosed aneurysm.
drained into the main pulmonary artery via a giant aneurysm, with parietal calcification and mural thrombus (Fig. 2, Video 2). Once the presence of a major left-to-right shunt and myocardial ischemia had been ruled out, a wait-and-see attitude was adopted.

Coronary fistulas are anomalous communications, either congenital or acquired, between a coronary artery and a cardiac chamber or blood vessel, present in 0.1%–0.8% of all coronary angiographies.1 The development of saccular aneurysms in coronary–pulmonary fistulas is even less common.2 Most arise from the right coronary artery or the left anterior descending artery, and about 90% drain into the venous circulation (right chambers, pulmonary artery, superior vena cava or coronary sinus).1 Its spectrum of clinical presentation varies, and depends on the severity of the left-to-right shunt. The entity is usually an incidental finding, though it may cause myocardial ischemia, arrhythmias, heart failure or sudden death.3 As shown by our case, multislice computerized tomography permits clear definition of the origin of these fistulas, their path and their distal site of drainage, as well as their relationship to other cardiac structures, and represents a very important advance in diagnostics, compared with coronary angiography. The main indications for closure of these fistulas are the development of clinical symptoms, especially myocardial ischemia, on the severity of the left-to-right shunt. The entity is usually an incidental finding, though it may cause myocardial ischemia, arrhythmias, heart failure or sudden death.3 As shown by our case, multislice computerized tomography permits clear definition of the origin of these fistulas, their path and their distal site of drainage, as well as their relationship to other cardiac structures, and represents a very important advance in diagnostics, compared with coronary angiography. The main indications for closure of these fistulas are the development of clinical symptoms, especially myocardial ischemia, arrhythmias, heart failure or sudden death.4

APPENDIX . SUPPLEMENTARY MATERIAL

Supplementary material associated with this article can be found in the online version available at doi:10.1016/j.rec.2010.08.002.

To the Editor,

Transposition of the great arteries (TGA) is the most common type of cyanotic heart disease in newborn infants. The atrial switching procedures (Mustard and/or Senning procedures) have permitted these infants to survive to adulthood.5 Sudden death is a substantial cause of late mortality. Presence of flutter or atrial fibrillation, syncope, or heart failure are predictors of sudden death in such patients.2 Ventricular arrhythmias are uncommon when ventricular dysfunction is not present. Implantation of an implantable cardioverter-defibrillator (ICD) can be a complex procedure and, although there have been reports of implantation using the subclavian venous approach, this might not always be feasible.

We present the case of an adolescent girl with TGA who underwent implantation of a atrio-biventricular ICD.

The 16-year-old patient had been born with interventricular shunting and persistent ductus arteriosus. She had undergone a Rashkind balloon atrial septostomy and, when 6 months old, a Mustard-type repair. For several years, she had asymptomatic paroxysmal complete ativoventricular block with narrow QRS interval and was in functional class II. She was admitted after cardiogenic syncope. An ejection fraction of 25% was observed in the echocardiogram and there were no obvious abnormalities in the baffles for redirection of atrial flow. We performed a catheterization study with induced syncopal ventricular tachycardia, with cardioversion at 200 J. It was decided to place an ICD and resynchronize because of the possible need for antibradycardia pacing as she had a history of CAVB with severe systemic ventricular dysfunction. Implantation by the subclavian approach was impossible as the electrode could not be advanced further than the upper area of the right atrium. It was decided to operate and place an epicardial pacing system. During opening of the sternum, rupture of the right ventricle occurred, requiring extracorporeal circulation and repair with sutures and Teflon. Epicardial pacing and sensing electrodes were placed on the right atrium and both ventricles. In addition, conventional 2-coil defibrillation electrodes were placed in the pericardial sac, around both ventricles, and were fixed with loose stitches. The electrodes were tunneled until reaching an infracavicular left subcutaneous bag and a high-energy (41 J) Guidant Contak Renewal 4 generator was implanted. It was confirmed that defibrillation was effective with 21 J and pacing thresholds less than 1 V in the 3 chambers. During the postoperative period, the patient developed bacteremia due to Serratia, which was successfully treated with a targeted antibiotic. The patient was discharged after 20 days. After 18 months of follow-up, she had not had any arrhythmic episodes and the device is currently functioning normally (Fig. 1).

The increased number of ICDs in pediatric patients with congenital heart diseases faces, among other problems, the limitations of percutaneous access in many patients, either because of small vessel size or the particular anatomy of the patient. The traditional placement of epicardial patches is associated with problems of pericardial restriction and increases in the defibrillation thresholds. Stephenson et al4 and Cannon et al5 both published series (22 and 8 patients, respectively)