with complex heart diseases and alternative approaches: a) implantation of 1 to 3 subcutaneous electrodes as the high-energy electrode or the transvenous defibrillation electrode itself in a subcutaneous position; b) transatrial approach with implantation of the defibrillation electrode directly through the right atrium; and c) implantation of the defibrillation electrode in the pericardial sac.

Given the small number of patients and the limited follow-up, it is hard to predict the complications associated with these procedures in the medium term. Stephenson et al. proposed annual monitoring of the defibrillation thresholds, which can become fundamentally elevated with subcutaneous electrodes. Cannon et al. recommend performing an annual echocardiography when there are electrodes in the pericardial sac to detect the possibility of progressive strangulation of the myocardium. This technique is considered preferable in patients with larger body surface areas, as was the case in our patient.

Due to the increase in the population with congenital heart disease reaching adulthood, we will need to use this and other approaches increasingly often.

Migration and Percutaneous Implantation of a Second Aortic Prosthesis

Migración e implante de segunda prótesis aórtica percutánea

To the Editor,

We present the case of an 82-year-old woman with severe aortic stenosis who was admitted to the hospital due to heart failure with New York Heart Association class III. Given the high surgical risk (EuroSCORE, 24%) and the associated comorbidity, surgical replacement was ruled out and the decision was made to perform transcatheter aortic prosthesis implantation.

The procedure was carried out by femoral approach and involved the implantation of a 26-mm CoreValve prosthesis according to the standard technique. Moderate-to-severe aortic insufficiency was observed due to the low implantation of the prosthesis (Fig. 1A), and the attempt was made to relocate it by traction using a snare catheter (Fig. 1B).

After a few minutes of continuous traction, the prosthesis migrated toward the ascending aorta, where it remained fixed. Several angiographic images were obtained, and we confirmed that the prosthesis did not move and that the supraaortic vessels were patent (Fig. 1C).

A second prosthesis was implanted with no complications (Fig. 1D), the gradient disappeared and the residual aortic insufficiency was mild.

The echocardiographic follow-up confirmed the proper function of the prosthesis and its stability (Fig. 1E). Nine months after the procedure, the patient was diagnosed with a bone tumor with pulmonary metastases, a chest computed tomography revealed the position of both prostheses (Fig. 1F), which had not changed since their implantation.

Malposition and migration of aortic prostheses are complications that have been reported previously, attempting to retrieve it or move it to the descending aorta have been proposed as the most safe
alternatives. However, in the case presented here, we show that it is safe to leave a prosthesis in the ascending aorta if it is stable and does not compromise the flow in the supraaortic vessels.

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Intra-Hisian Block During Transcatheter Aortic Valve Implantation With the CoreValve Prosthesis

bloqueo intrahisiano durante el implante de la prótesis aórtica percutánea CoreValve

To the Editor,

Percutaneous aortic valve implantation is an alternative to surgery in the treatment of severe symptomatic aortic stenosis in patients at high surgical risk.1,2 The series published with the percutaneous implantation of the CoreValve® (Medtronic CV, Luxembourg) aortic valve prosthesis show a high success rate. However, a high incidence of conduction disorders has been described, with frequent occurrence of left bundle branch block and complete atrioventricular (AV) block in 20%-35% of cases.3,4

We report the case of an 83-year old patient diagnosed with severe symptomatic aortic stenosis who was rejected for surgery due to the high surgical risk involved. He was implanted with a CoreValve® percutaneous aortic valve prosthesis. An electrophysiological study was performed on the patient immediately before

REFERENCES


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