Prosthetic valve endocarditis after transcatheter CoreValve Evolut R bioprosthesis implantation

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Prosthetic valve endocarditis (PVE) affects 1–6% of prosthetic valve recipients and in patients treated with transcatheter aortic valve implantation (TAVI) has an incidence of 0.3–2.3% per patient-year [1–3].

A 79-year-old man who underwent TAVI with a Medtronic Evolut R 29 mm bioprosthesis 7 months earlier was admitted due to recurring episodes of fever up to 39°C and mild worsening of exercise tolerance lasting for 4 months. Other relevant medical history consisted of chronic obstructive pulmonary disease (COPD) and penicillin allergy. During the past months the patient was hospitalized twice (once due to fever, the second time due to syncope), but the symptoms were attributed to COPD exacerbation.

At admission, physical examination was remarkable for systolic murmur over the mitral valve. No fever, chest pain, cough, or meningeal signs were appreciated. Electrocardiogram (ECG) demonstrated sinus rhythm with left bundle branch block observed in previous ECGs. Chest X-ray revealed an enlarged cardiac silhouette. Transthoracic echocardiography (TTE) demonstrated mild mitral regurgitation (Figure 1 A) and 15 mm hyperechogenic structures on the mitral valve described as possible vegetations (Figure 1 B). Laboratory findings revealed a white blood cell count of 19 × 10³/μl, N-terminal prohormone of brain natriuretic peptide of 1000 pg/ml and C-reactive protein of 123 mg/l.

Due to high clinical suspicion of PVE, the patient was started on vancomycin, gentamicin and rifampin. Beforehand, three sets of blood cultures were collected, which were found to be positive for methicillin-sensitive Staphylococcus capitis. Transesophageal echocardiography (TEE) confirmed PVE demonstrating vegetations on the leaflet located in the vicinity of the native right coronary cusp (Figures 1 C and D). Further treatment with rifampin and gradually increased doses of vancomycin (gentamicin was discontinued after 2 weeks due to transitory worsening of the renal function) led to clinical improvement, whilst laboratory examinations showed C-reactive protein and white blood cell (WBC) normalization. Even though repeated TEEs demonstrated persistence of vegetations, clinically the PVE was deemed healed. After 6 weeks antibiotics were discontinued, blood cultures were negative and the patient was discharged.

A recent study [2] on 7944 patients found two independent factors associated with infective endocarditis (IE) prevalence – orotracheal intubation and use of self-expandable CoreValves. Among factors possibly responsible for PVE prevalence in the cohort treated with CoreValves, the following were suggested: larger stent frame, larger contact surface area between tissues and the frame, and paravalvular leak (PVL).

In another study [4] risk factors for post-TAVI infective endocarditis development were male sex, low implantation position, implantation of > 1 prosthesis, and vascular and bleeding complications.

The newly released Medtronic CoreValve Evolut R system shows promising features as compared to the first generation of CoreValves in terms of PVE prevention. Firstly, its height is 10% shorter, so both the stent frame and the contact surface between tissues and the frame are smaller. Secondly, it has an extended skirt designed to minimize PVL. Thirdly, the valve’s repositionability facilitates optimal implantation and may reduce the need for valve-in-valve procedures. Finally, its smaller insertion profile could minimize occurrence of vascular and bleeding complications and promote TAVI without orotracheal intubation.

In the case of our patient, potential risk factors included male sex, use of a self-expandable valve and orotracheal intubation. On the other hand, IE prophylaxis was administered (cefazolin), only one valve was used, the position was optimal, and neither PVL nor vascular and bleeding complications were present.
In conclusion, despite potentially protective modifications in design, PVE on the Evolut R valve may still occur and in order to accurately assess PVE prevalence larger registry-based studies are required. Furthermore, one must emphasize the need for repeated TEEs and blood cultures as well as using new imaging modalities such as cardiac computed tomography (CT), positron emission tomography (PET)/CT and single-photon emission computed tomography (SPECT)/CT in cases with a high level of clinical IE suspicion, especially in prosthetic valve recipients.

Conflict of interest
The authors declare no conflict of interest.
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