Very late transcatheter heart valve thrombosis

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Abstract

We describe a case of very late transcatheter heart valve (THV) thrombosis of a first-generation SAPIEN prosthesis (Edwards Lifesciences, Irvine, CA) implanted in a 64-year-old woman with severe symptomatic aortic stenosis. More than 54 mo after implantation, she presented with severe symptomatic prosthesis dysfunction (stenosis) which was successfully treated with oral anticoagulation. To our knowledge, this is the tardiest case of THV thrombosis ever reported. This case should increase clinical awareness for THV thrombosis even beyond the first two-year period following implantation.

Key words: Valve thrombosis; Transcatheter heart valve; Transcatheter aortic valve replacement

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Core tip: We describe the tardiest case of transcatheter aortic valve replacement (TAVR) thrombosis ever reported. A 64-year-old woman with severe symptomatic aortic stenosis underwent TAVR with a first-generation SAPIEN prosthesis. More than four years (> 54 mo) following implantation, she presented with a severe symptomatic prosthesis dysfunction (stenosis) which was successfully treated with oral anticoagulation.

INTRODUCTION

We describe a case of very late transcatheter heart valve (THV) thrombosis of a first-generation SAPIEN prosthesis (Edwards Lifesciences, Irvine, CA) implanted in a 64-year-old woman with severe symptomatic
aortic stenosis. More than 54 mo after implantation, she presented with severe symptomatic prosthesis dysfunction (stenosis) which was successfully treated with oral anticoagulation. To our knowledge, this is the third case of THV thrombosis ever reported. This case should increase clinical awareness for THV thrombosis even beyond the first two-year period following implantation.

CASE REPORT

A 64-year-old woman with severe symptomatic aortic stenosis and porcelain aorta underwent transcatheter aortic valve replacement (TAVR) using a first-generation balloon-expandable Edwards SAPIEN (Edwards Lifesciences, Irvine, CA) 23 mm THV in January 2011. The patient was discharged home with aspirin and clopidogrel for three months followed by low dose aspirin.

More than 54 mo following TAVR, she presented with progressive dyspnea, angina and dizziness on moderate exertion. Transthoracic echocardiogram (TEE) revealed a severe THV dysfunction with an aortic valve area (AVA) of 0.81 cm², peak and mean trans prosthetic gradients (TPG) of 76 and 39 mmHg respectively and a preserved left ventricular ejection fraction (Figure 1). Transesophageal echocardiogram (TEE) did not demonstrate abnormal leaflet function neither THV thrombosis (Figure 2) but revealed a moderate paravalvular leak (PVL) (Figure 1). Transesophageal echocardiogram (TEE) did not demonstrate abnormal leaflet function neither THV thrombosis (Figure 2) but revealed a moderate paravalvular leak (PVL) (Figure 1) in TPG (Figure 1). The patient was discharged home with aspirin, clopidogrel and warfarin for three months, followed by clopidogrel and warfarin. Angina was relieved immediately after PCI, but dyspnea on exertion improved over weeks. After three months, symptoms had resolved completely and TPG returned to their baseline values. They remained unchanged after nine months under chronic anticoagulation (Figure 1).

DISCUSSION

In two series, less than 30 cases of THV thrombosis have been reported with a median presentation time of 6 mo (range, 3-735 d) with only two cases occurring beyond one year[1-3]. A case of a Direct Flow Medical THV thrombosis three years following TAVR has also been reported recently[4]. To our knowledge, never a THV thrombosis has been described so late after implant (> 54 mo) as in this case. THV thrombosis is a new entity that needs to be recognized not only by TAVR specialists. Even though estimated incidence is low (0.61%), consequences can be catastrophic if appropriate therapy is not initiated promptly[5]. Only reported cases treated with anticoagulation had favorable outcome. Therefore, a sudden increase in TPG should trigger further investigation or therapies to exclude THV thrombosis. Valve hemodynamic deterioration (VHD), defined by an increase in mean TPG of more than 10 mmHg over time, was observed more frequently in patients with smaller THV (23 mm) and those not receiving oral anticoagulation[4]. Most of these patients did not have a progressive deterioration after the first year. VHD does not seem to be part of a continuum towards THV thrombosis and this later remains unpredictable. However, the pathophysiology of VHD may include some degree of sub-clinical leaflet thrombosis.

In our case, TEE evaluation was not diagnostic. In a series of 3 pathology-proven THV thrombosis, TEE was also negative in each cases[2]. However, Makkar et al[3] showed that 4D-CT and TEE had a diagnostic concordance of 100% in 10 patients presenting reduced leaflets motion following TAVR. Whether 4D-CT is the optimal imaging modality remains to be proven. CMR was chosen over CT scan to assess PVL severity but
also failed to identify the THV thrombosis. Therefore, negative imaging should not preclude an anticoagulation trial when the diagnostic is highly suspected.

Reports have revealed subclinical leaflet thrombosis detected by TEE and 4D CT scan early after THV implantation\(^5\). Most of these cases were seen in patients who did not receive anticoagulation. The clinical significance of these findings remains unknown at this point. Studies are ongoing to evaluate the optimal antiplatelet/anticoagulant therapy after TAVR. The ARTE (clopidogrel; NCT01559298) and REAC-TAVI (ticagrelor vs clopidogrel; NCT02224066) trials are evaluating different antiplatelet strategies and the GALILEO (rivaroxaban; NCT02556203) and ATLANTIS (apixaban; NCT02664649) studies are looking at the effect of non-vitamin K oral anticoagulant following TAVR. Similarly, the POPular-TAVI trial (NCT02247128) is evaluating the effect of adding clopidogrel for 3 mo following implantation in patients with and without ongoing vitamin-K oral anticoagulation treatment.

This case is also particular because of the concurrent left main stenosis. Although thrombus migration in the left main could be considered an explanation, the angiographic appearance was more in favor of disease progression of a previous non-significant lesion seen on the pre-TAVR angiogram. Moderate PVL could be associated with a different flow pattern over the leaflets and in the left main that could lead to THV thrombosis as well as accelerated progression of atherosclerosis although this relationship is speculative.

This case should increase clinical awareness for THV thrombosis even beyond the first two-year period following implantation. Risk predictors for THV thrombosis and optimal antiplatelet/anticoagulant therapy post TAVR are still unknown and warrants clinical trials.

**COMMENTS**

**Case characteristics**

More than 54 mo following transcatheter aortic valve replacement, a 64-year-old woman presented with progressive dyspnea, angina and dizziness on moderate exertion.

**Clinical diagnosis**

Prosthesis dysfunction.

**Differential diagnosis**

Transcatheter heart valve (THV) degeneration; THV thrombosis; THV endocarditis.

**Imaging diagnosis**

Transesophageal echocardiogram (TEE) revealed a prosthesis dysfunction with a severe stenosis. TEE failed to revealed THV thrombosis as well as vegetations.

**Treatment**

Empiric treatment with unfractionated intravenous heparin followed by long-term anticoagulation with warfarin.

**Related reports**

Less than 30 cases of THV thrombosis have been reported with a median presentation time of 6 mo (range, 3-735 d) with only two cases occurring beyond one year.
**Experiences and lessons**

This case should increase clinical awareness for THV thrombosis even beyond the first two-year period following implantation. Even if TEE does not reveal THV thrombosis.

**Peer-review**

The paper is well written.

**REFERENCES**

1. Latib A, Naganuma T, Abdel-Wahab M, Danenberg H, Cota L, Barbanti M, Baumgartner H, Finkelstein A, Legrand V, de Lezo JS, Kefer J, Messika-Zeitoun D, Richard G, Stabile E, Kaleschke G, Vahanian A, Laborde JC, Leon MB, Webb JG, Panoulas VF, Maisano F, Alfieri O, Colombo A. Treatment and clinical outcomes of transcatheter heart valve thrombosis. *Circ Cardiovasc Inter* 2015; 8: e001779 [PMID: 25873727 DOI: 10.1161/CIRCINTERVENTIONS.114.001779]

2. De Marchena E, Mesa J, Pometi S, Marin Y Kall C, Marinicic X, Yahagi K, Ladich E, Kutz R, Aga Y, Ragosta M, Chawla A, Ring ME, Virmani R. Thrombus formation following transcatheter aortic valve replacement. *JACC Cardiovasc Interv* 2015; 8: 728-739 [PMID: 25946447 DOI: 10.1016/j.jcin.2015.03.005]

3. Regazzoli D, Ancona MB, Mangieri A, Agricola E, Spagnolo P, Mussardo M, Colombo A, Latib A. A Case of Very Late (3 Years) Transcatheter Heart Valve Thrombosis. *JACC Cardiovasc Interv* 2016; 9: e83-e84 [PMID: 27101920 DOI: 10.1016/j.jcin.2016.01.024]

4. Del Trigo M, Muñoz-García AJ, Wijeysundera HC, Nombela-Franco L, Cheema AN, Gutierrez E, Serra V, Kefer J, Amat-Santos IJ, Benitez LM, Mewa J, Jiménez-Quevedo P, Alnasser S, García Del Blanco B, Dager A, Abdul-Jawad Altisent O, Puri R, Campelo-Parada F, Dahou A, Paradis JM, Dumont E, Pibarot P, Rodés-Cabau J. Incidence, Timing, and Predictors of Valve Hemodynamic Deterioration After Transcatheter Aortic Valve Replacement: Multicenter Registry. *J Am Coll Cardiol* 2016; 67: 644-655 [PMID: 26868689 DOI: 10.1016/j.jacc.2015.10.097]

5. Makkar RR, Fontana G, Jilaihawi H, Chakravarty T, Kofoed KF, de Backer O, Asch FM, Ruiz CE, Olsen NT, Trento A, Friedman J, Berman D, Cheng W, Kashif M, Jelnin V, Klijer CA, Guo H, Pichard AD, Weissman NJ, Kapadia S, Bhatt DL, Leon MB, Søndergaard L. Possible Subclinical Leaflet Thrombosis in Bioprosthetic Aortic Valves. *N Engl J Med* 2015; 373: 2015-2024 [PMID: 26436963 DOI: 10.1056/NEJMoa1509233]

P- Reviewer: Grignola JC, Kettering K, Paradis JM, Said SAM, Vermersch P
S- Editor: Qiu S
L- Editor: A
E- Editor: Lu YJ
