

30. Tissot AC, Maurer P, Nussberger J, Sabat R, Pfister T, Ignatenko S, Volk HD, Stocker H, Muller P, Jennings GT, Wagner F, Bachmann MF. Effect of immunisation against angiotensin II with CYT006-AngQb on ambulatory blood pressure: a double-blind, randomised, placebo-controlled phase IIa study. *Lancet* 2008; **371**:821–827.
31. Weber MA, Black H, Bakris G, Krum H, Linas S, Weiss R, Linseman JV, Wiens BL, Warren MS, Lindholm LH. A selective endothelin-receptor antagonist to reduce blood pressure in patients with treatment-resistant hypertension: a randomised, double-blind, placebo-controlled trial. *Lancet* 2009; **374**:1423–1431.
32. Bhatt DL. Advancing the care of cardiac patients using registry data: going where randomized clinical trials dare not. *JAMA* 2010; **303**:2188–2189.
33. Kumbhani DJ, Steg PG, Cannon CP, Eagle KA, Smith SC Jr, Crowley K, Goto S, Ohman EM, Bhatt DL. Prospective assessment of the relationship between the use of evidence-based prevention therapies and long-term (4-year) outcomes in stable outpatients with established atherosclerotic disease: Insights from the international Reduction of Atherothrombosis for Continued Health (REACH) registry. *Circulation* 2011; **124**:A15957.
34. Osterberg L, Blaschke T. Adherence to medication. *N Engl J Med* 2005; **353**:487–497.
35. Kumbhani DJ, Bavry AA, Harvey JE, de Souza R, Scarpioni R, Bhatt DL, Kapadia SR. Clinical outcomes after percutaneous revascularization versus medical management in patients with significant renal artery stenosis: a meta-analysis of randomized controlled trials. *Am Heart J* 2011; **161**:622–630 e1.
36. Redon J, Campos C, Narciso ML, Rodicio JL, Pascual JM, Ruilope LM. Prognostic value of ambulatory blood pressure monitoring in refractory hypertension: a prospective study. *Hypertension* 1998; **31**:712–718.
37. Staessen JA, Beilin L, Parati G, Waeber B, White W. Task force IV: Clinical use of ambulatory blood pressure monitoring. Participants of the 1999 Consensus Conference on Ambulatory Blood Pressure Monitoring. *Blood Press Monit* 1999; **4**:319–331.

CARDIOVASCULAR FLASHLIGHT

doi:10.1093/eurheartj/ehs307

Online publish-ahead-of-print 17 September 2012

Regurgitation after Edwards SAPIEN valve implantation: truly paravalvular or 'supra-skirtal'?

Barbara E. Stähli¹, Catherine Gebhard¹, Volkmar Falk², Roberto Corti¹, Rolf Jenni¹, and Felix C. Tanner^{1*}

¹Department of Cardiology, Cardiovascular Center, University Hospital Zürich, Rämistrasse 100, 8091, Zürich, Switzerland and ²Clinic for Cardiovascular Surgery, Cardiovascular Center, University Hospital Zürich, Rämistrasse 100, 8091, Zürich, Switzerland

* Corresponding author. Tel: +41 44 255 99 97, Fax: +41 44 255 48 04, Email: felix.tanner@access.uzh.ch

This paper was guest edited by Frank Rademakers, University Hospital Gasthuisberg, Leuven, Belgium

A 73-year-old male was referred with severe mitral regurgitation 9 years after mitral valve annuloplasty with a 32-mm Edwards Lifesciences ring. On the basis of an anticipated high perioperative risk related to chronic kidney disease and left ventricular systolic dysfunction, mitral valve-in-ring implantation using a 29-mm Edwards SAPIEN prosthesis was performed by transapical access (Panel A). Post-procedural three-dimensional transoesophageal echocardiography revealed three paravalvular regurgitation jets at the 3, 7, and the 11 o'clock positions (Panels B and C).

In this patient, we observed a third type of regurgitation in Edwards SAPIEN prostheses, which is caused by the device design. Edwards SAPIEN prostheses are built up by bovine pericardial tissue leaflets fixed on a stainless steel frame. A polyethylene terephthalate skirt partially covers the steel frame. However, since the skirt covers the basal two-thirds of the frame only, regurgitation through its uncovered part may occur (arrow, Panel D; adapted from www.edwards.com/products/transcatheter/valve/Pages/sapienthv.aspx).

This type of regurgitation was seen in the patient presented here, since the skirt is positioned in the left atrium rather than the annuloplasty ring. The commissures cover the whole length of the steel frame, and the regurgitation jets are therefore located in between the commissures and positioned ~120° apart from each other. In patients with Edwards SAPIEN prostheses in aortic position, the same phenomenon may be observed with prostheses implanted too apically. However, as the left ventricular outflow tract is narrow, regurgitation jets are more difficult to visualize.

Hence, after Edwards SAPIEN prosthesis implantation, regurgitation is transvalvular or paravalvular or 'supra-skirtal'. As the technology of transcatheter valves is still evolving, constant improvement in the device design and proper placement of the prosthesis are important.

