

of severe symptomatic aortic stenosis (SSAS) offers little hope of survival and valve substitution surgery is the only effective treatment.² However, 30%-40% of patients with SSAS are not operated as they present too high an estimated surgical risk.³ Survival and quality of life are limited in these patients. Percutaneous balloon valvuloplasty was initially considered an alternative therapy but it offers very poor mid-term results.⁴ Recently, percutaneous implantation of prosthetic aortic valves (PIAP) has been introduced as a valid option for patients with SSAS and prohibitive surgical risk.⁵⁻⁸

One contraindication of PIAP is severe interventricular septal hypertrophy because the prosthesis may be displaced and migrate during implantation. We describe a simple modification to aortic prosthesis deployment that has enabled us to treat PIAP successfully (Edwards-Sapien prosthetic valve) in 2 patients with inoperable SSAS and severe interventricular septal hypertrophy.

Two patients with SSAS (men, aged 79 and 90 years) were referred for PIAP with Edwards-Sapien prostheses (Edwards Lifesciences Inc, Irvine, California, US). The patient selection process^{8,9} revealed no contraindication for the technique other than severe interventricular septal hypertrophy (18 mm and 19 mm, respectively). We decided to proceed with the intervention.

In both patients, we implanted an 26 mm Edwards-Sapien prosthesis, an expandable aortic valve balloon prosthesis delivered via a stainless steel stent, with 3 bovine pericardial valves. It is assembled in the interventional cardiology laboratory with a 30 mm long balloon. The 26 mm valve is 16 mm long, and its location in the middle of the balloon is recommended, leaving 7 mm of balloon on either side of the prosthesis. Severe interventricular septal hypertrophy is considered a contraindication for PIAP because when balloon inflation—and therefore, implantation—takes place, the prosthesis may be displaced by the septum and migrate towards the ascending aorta. In these 2 patients, we minimized contact between the balloon and the interventricular septum by positioning the prosthetic valve asymmetrically, more distal, leaving only 3-4 mm of balloon on the intraventricular side of the prosthesis (Figure). Thus, we were able to implant the prostheses successfully in both 2 patients, without displacement, with optimal expansion, and without periprosthetic insufficiency. Both patients were discharged without complications.

These 2 patients described show severe interventricular septal hypertrophy, not necessarily a contraindication for transcatheter aortic valve implantation (TAVI) of Edwards-Sapien prostheses. This technique can only be recommended for patients with severe interventricular septal

Severe Septal Hypertrophy: Is It Necessarily a Contraindication for the Transcatheter Implantation of an Edwards-Sapien Prosthesis?

To the Editor,

Aortic stenosis is the most frequent valvular heart disease in the developed world.¹ The natural history

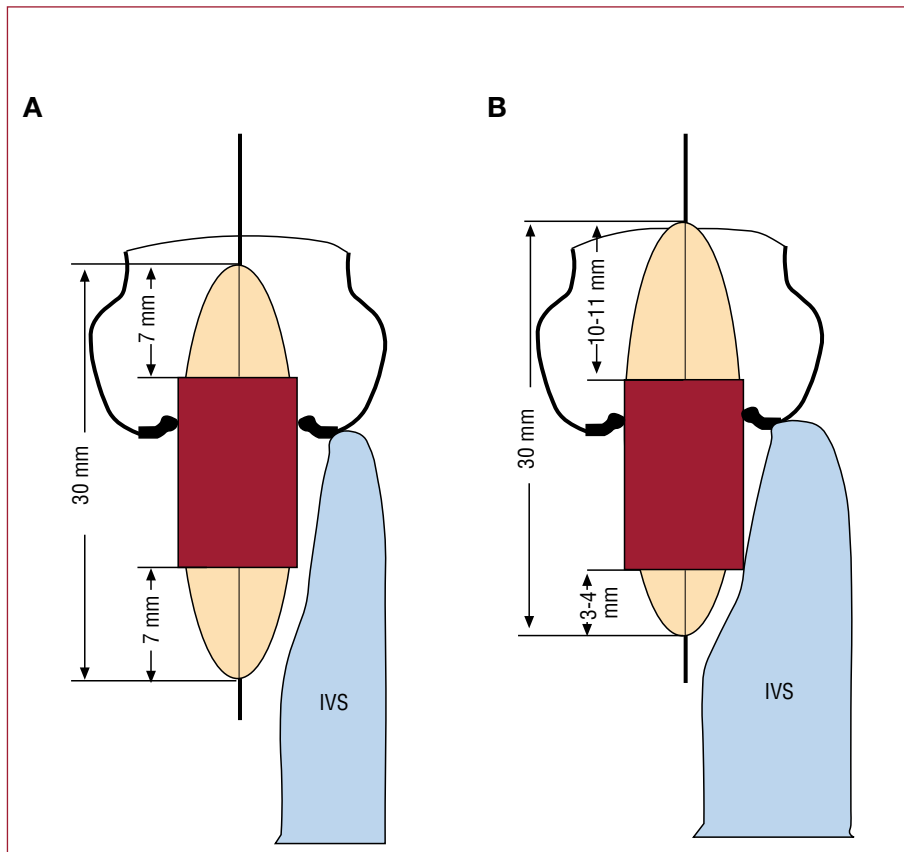


Figure 1. A: deployment of the prosthesis in a patient without severe interventricular septal hypertrophy. The prosthesis is positioned symmetrically in the balloon. B: patient with severe interventricular septal hypertrophy. To minimize contact between the balloon and the interventricular septum, the valve is located in the balloon asymmetrically, more distal, to leave only 3-4 mm of the balloon on the ventricular side of the prosthesis.

hypertrophy, given that the risk of prosthesis displacement in most patients is very low when the prosthesis is located symmetrically in the balloon.

Raúl Moreno,^a Luis Calvo,^a Eulogio García,^b
and David Dobarro^a

^aUnidad de Cardiología Intervencionista, Hospital Universitario La Paz,
Madrid, Spain

^bUnidad de Cardiología Intervencionista, Hospital Clínico San Carlos,
Madrid, Spain

REFERENCES

1. Nkomo VT, Gardin JM, Skelton TN, Gottdiener JS, Scott CG, Enriquez-Sarano M. Burden of valvular heart disease: a population-based study. *Lancet*. 2006;368:1005-11.
2. Task Force on the Management of Valvular Heart Disease of the European Society of Cardiology. Guidelines on the management of valvular heart disease. *Eur Heart J*. 2007;428:e1-39.
3. Iung B, Baron G, Butchart EG, Delahaye F, Gohlke-Bärwolf C, Levang OW, et al. A prospective survey of patients with valvular heart disease in Europe: The EuroHeart Survey on Valvular Heart Disease. *Eur Heart J*. 2003;24:1231-43.
4. McKay RG. The Mansfield Scientific Aortic Valvuloplasty Registry: overview of acute hemodynamic results and procedural complications. *J Am Coll Cardiol*. 1991;17:189-92.
5. Cribier A, Eltchaninoff H, Tron C, Bauer F, Agatiello C, Nercolini D, et al. Treatment of calcific aortic stenosis with the percutaneous heart valve: mid-term follow-up from the initial feasibility studies: the French experience. *J Am Coll Cardiol*. 2006;47:1214-23.
6. Webb JG, Pasupati S, Humphries K, Thompson C, Altwegg L, Moss R, et al. Percutaneous transarterial aortic valve replacement in selected high-risk patients with aortic stenosis. *Circulation*. 2007;116:755-63.
7. Grube E, Schuler G, Buellesfeld L, Gerckens U, Linke A, Wenaweser P, et al. Percutaneous aortic valve replacement for severe aortic stenosis in high-risk patients using the second and current third generation self-expanding CoreValve prosthesis. *J Am Coll Cardiol*. 2007;50:69-76.
8. Moreno R, Calvo L, Filgueiras D, López T, Sánchez-Recalde A, Jiménez-Valero S, et al. Implantación percutánea de prótesis valvulares aórticas en pacientes con estenosis aórtica severa sintomática rechazados para cirugía de sustitución valvular. *Rev Esp Cardiol*. 2008;61:1215-9.
9. García E, Pinto AG, Sarnago F, Pello AM, Paz M, García-Fernández MA, et al. Implantación percutánea de prótesis valvular aórtica: experiencia inicial en España. *Rev Esp Cardiol*. 2008;61:1210-4.