Atrial Septal Defect With Severe Pulmonary Hypertension in Elderly Patients: Usefulness of Transient Balloon Occlusion

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BRIEF REPORT

In patients with an atrial septal defect and severe pulmonary hypertension, it is important to determine whether the latter is reversible before percutaneous or surgical closure. In addition to determining pulmonary resistance, one simple technique is to transiently occlude the septal defect using a balloon catheter and to evaluate the hemodynamic response. We defined a positive response as a $\geq 25\%$ reduction in mean pulmonary artery pressure during occlusion relative to the basal level, without a fall in systemic pressure or an increase in ventricular end-diastolic pressure. The study included five patients aged over 60 years with an atrial septal defect and severe pulmonary hypertension who were referred for percutaneous closure. In one patient, the test gave a negative result and closure of the atrial septal defect was not performed. In the remaining four, closure was indicated. In three patients, closure was performed percutaneously, while the fourth underwent surgery. The drop in pulmonary pressure observed during the test was maintained over the long term at a mean follow-up time of 22 months.

Key words: Pulmonary arterial hypertension. Atrial septal defect. Congenital heart disease. Cardiac catheterization.

INTRODUCTION

Atrial septal defect (ASD) is congenital heart disease which is more frequently diagnosed in adults. In some patients, chronic exposure to high pulmonary flow typically causes a progressive increase in pulmonary pressure and resistance, with an estimated prevalence of less than 10\%. Patients with severe pulmonary hypertension (PH) or Eisenmenger syndrome have a poor prognosis, but better than the prognosis of patients with idiopathic PH.1 In patients with Eisenmenger syndrome, defect closure may alter their prognosis to one more similar to patients with idiopathic PH. The difference between the 2 entities is thought to be the atrial defect which would function as a safety valve to unload the right ventricle thereby preserving its function and maintaining systemic output, although at the risk of hypoxemia and cyanosis.2

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Patients
Between January 2002 and December 2007, 51 adult patients with ASD were referred to our catheterization laboratory for potential percutaneous closure. In total, 5 of the patients presented severe PH, defined as right ventricular systolic pressure (RVSP) ≥70 mmHg as estimated by transthoracic echocardiography. These patients underwent balloon test occlusion to assess whether PH was reversible. Follow-up was conducted on an outpatient basis and included echocardiography at 1 month, 3 months, and 6 months and annually thereafter.

Statistical Analysis
The SPSS statistical package (version 15) was used. Serial echocardiographic and hemodynamic values were compared using the Wilcoxon nonparametric test.

RESULTS
The results of basal echocardiography and echocardiography at follow-up are shown in Table 1. All the patients were symptomatic women with a mean age of 65 (6) years. The mean size of the ASD as estimated by transeosophageal echocardiography was 25 (12) mm. The defect was occluded in 4 patients; this was done percutaneously in 3, and 1 underwent surgery due to the lack of suitable margins. Despite receiving treatment with bosentan and sildenafil, pulmonary hypertension remained unchanged at 5-year follow-up in the patient in whom closure was underestimated, although her New York Heart Association (NYHA) functional class changed from III to class II. In the patients in whom ASD closure was successful, the mean RVSP was 85 (17) mmHg before closure, falling to 52 (7) mmHg (P= .06) after a mean 22 (16) months of follow-up. In 2D-mode, a reduction was also observed in the size of the right ventricle from 37 (4) mm to 31 (5) mm (P=.06) and 2D-sonography showed a reduction in the outflow...
ppm; inhaled iloprost 10 µL) with negative results (Figure 1). In the patients in whom ASD was successfully closed, mean pulmonary pressure was 38 (5) mmHg, decreasing to 26 (5) mmHg after balloon test occlusion. Systemic pressure did not

tract from 54 (3) mm to 42 (6) mm (P=.06). Table 2 shows the hemodynamic values obtained during temporary balloon test occlusion. The patient in whom closure was underestimated also underwent tests with vasodilators (inhaled nitric oxide, 80

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<tbody>
<tr>
<td>PAP, mmHg</td>
<td>AoP, mmHg</td>
<td>PVR, uW/m²</td>
<td>Qp/Qs</td>
</tr>
<tr>
<td>1</td>
<td>70/30 43</td>
<td>160/61 76</td>
<td>6</td>
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<tr>
<td>2</td>
<td>52/20 31</td>
<td>165/85 108</td>
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<td>3</td>
<td>87/34 52</td>
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<td>4</td>
<td>70/20 38</td>
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<td>5</td>
<td>65/28 41</td>
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AoP indicates aortic pressure; PAP, pulmonary arterial pressure; PVR, pulmonary vascular resistance; Qp pulmonary output; Qs, systemic output.

Figure 1. Recording of pulmonary and systemic pressure during basal study, after temporary balloon test occlusion, and after nitric oxide (NO) inhalation in patient 3. The results of temporary balloon test occlusion were negative.
of pulmonary vascular resistance, its relation to systemic resistance and the size the shunt, balloon test occlusion can provide additional information on whether the pulmonary arterial hypertension is reversible or not in order to establish whether the defect is operable.

Formerly, in order to establish whether adult patients with severe PH and dominant left-right shunt were operable, pulmonary biopsy was conducted to determine whether the histological changes associated with PH in small pulmonary vessels were irreversible or reversible, mainly by vasoconstriction. Currently, different vasodilator agents, such as tolazoline, epoprostenol, nitric oxide, iloprost, or sildenafil, are used to assess the degree of reversibility. However, there is no evidence regarding their usefulness in predicting the response of PH to defect closure.

Two studies have demonstrated long-term reductions in pulmonary pressure in patients with ASD and PH who have undergone percutaneous closure. In a study by Suárez de Lezo et al, mean systolic pulmonary pressure was invasively measured in 29 patients. The initial value of 65 (23) mmHg decreased to 54 (21) mmHg after occlusion during approximately 21 months of follow-up. More recently, Balint et al studied 44 patients; after a mean follow-up of 31 months, pulmonary pressure fell from 58 mmHg to 44 mmHg.

Temporary balloon test occlusion offers a unique and unrivalled opportunity to assess the response of PH following definitive closure.

If there are elevated ventricular end-diastolic pressures or decreased systemic expenditure, defect closure could be detrimental to the right ventricle as this would close the safety valve that helps maintain its contractile function. The degree to which there is a reduction in pulmonary pressure during closure provides information on the percentage of hyperkinetic hypertension. In addition to the acute reduction due to the increase in pulmonary flow, there is a subsequent reduction in pulmonary pressure in the long term. This small long-term reduction may be due to reversible changes in pulmonary arteriolar vasculature. It is reasonable to assume that temporary balloon test occlusion may reveal the short-term changes in PH by reducing the pulmonary hyperflow and that acute vasodilator testing may show the long-term changes in vascular remodelling. A mean ≥25% decrease in pulmonary pressure relative to basal values is required to define the test as positive. However, this value is arbitrary, and larger studies are needed to define objective criteria by which the test can be considered positive and to assess its relationship to the long-term evolution of pulmonary pressure.
REFERENCES


