Scientific letters

Heart Transplantation in Pediatric Patients With Pulmonary Hypertension

Trasplante cardiaco en pacientes pediátricos con hipertensión pulmonar

To the Editor,

Patients with congenital heart disease may have pulmonary hypertension secondary to increased pulmonary flow, persistent hypoxemia, or elevated left-side filling pressures. Persistently elevated pulmonary pressure causes pulmonary vasculature remodeling and pulmonary hypertension refractory to vasodilator therapy. Previous reports have described the anatomic-pathologic changes in pulmonary vasculature and their importance. Pulmonary hypertension may be a contraindication for heart transplantation. However, it is difficult to determine the pulmonary resistance value that should be used to contraindicate heart transplantation. Recommendations for pediatric patients are based on experience with adults, and the latest guidelines establish an upper limit of 6 UW/m² after the administration of pulmonary vasodilator therapy. Nevertheless, some authors defend the possibility of heart transplantation at higher values.

From December 2008 to December 2013, we performed 22 heart transplantations in pediatric patients, among them, 5 patients with severe pulmonary hypertension. The characteristics of these patients are described in the Table. All patients underwent catheterization prior to transplantation, except for 1 patient whose pulmonary pressure was estimated by echocardiography. Pulmonary resistances were calculated at baseline and after the administration of pulmonary vasodilator therapy (nitric oxide). Patient 4 was on the transplantation waiting list for 2 years, but had considerable clinical deterioration with the development of severe pulmonary hypertension (Table); hence, a decision was made to implant a left ventricular assist device and administer pulmonary vasodilator therapy. One month later, the catheterization was repeated and pulmonary resistances had dropped to 3.5 UW/m² and, therefore, the patient was put back on the transplantation waiting list.

One patient died in the acute phase of the postoperative period due to humoral rejection. All other patients are alive and progressing well. Two patients (40%) required mechanical assistance, 1 due to humoral rejection and the other due to right ventricular dysfunction. All had moderate-to-severe right ventricular dysfunction and required inotropic support and pulmonary vasodilator therapy. In the patients without pulmonary hypertension, right ventricular dysfunction was observed in 9 of 17 (53%; P < .05). Pulmonary vasodilator therapy was maintained at discharge (oral sildenafil), but all patients discontinued the drug during follow-up. Pulmonary biopsies were obtained in 2 patients (Figure) and showed the entire spectrum of vascular lesions characteristic of pulmonary hypertension, with involvement of precapilar and intraarterial arterial vessels, such as plexiform vasculopathy. A venous condition was also observed in the form of hypertrophy. In 1 patient (Figure A), there was a predominance of medial hypertrophy changes in precapilar vessels and plexiform vasculopathy. In the other patient (Figure B), these changes were less serious, but greater intimal thickening was observed, as well as venous involvement with lymph vessel dilatation.

Comparison of these patients with those without pulmonary hypertension showed no statistically significant differences in survival: 80% of patients with pulmonary hypertension survived compared with 88% of patients without hypertension, with a mean follow-up of 27 (10-62) and 29 (7-60) months, respectively (P > .5). We did observe a higher incidence of cellular (80% vs 24%; P = .02) and humoral (80% vs 12%; P < .01) rejection in patients with pulmonary hypertension, probably due to the greater complexity in this subgroup: 80% of patients with pulmonary hypertension compared with 29% in those without pulmonary hypertension underwent more than 1 cardiac surgery prior to transplantation, including placement of a ventricular assist device (P = .04). Only 2 patients, 1 in each group, had developed antihuman leukocyte antibodies (HLA) before transplantation.

In conclusion, it is difficult to establish a value of pulmonary resistance that could be used to contraindicate heart transplantation. Likewise, when referring to pulmonary resistances, the term irreversible should be used with caution because these resistances...
tend to drop over time and can even become normal. Additionally, the prognostic value of pulmonary biopsy is also unclear. The absence of fibrosis, as in our patients, may be a marker of reversibility. Pulmonary vasodilators and ventricular assistance have been shown to be useful as a bridge to eligibility in both adult and pediatric patients, as they allow transplantation in patients initially rejected due to pulmonary hypertension. This strategy may be preferable to cardiopulmonary transplantation.

Ferran Gran, a,b Dimpna Albert, c Joan Sanchez-de-Toledo, c Joan Balcells, c Joan Carles Ferreres, c and Raül Abella d

a Unidad de Cardiología Pediátrica, Hospital Universitario de la Vall d’Hebron, Universidad Autónoma de Barcelona, Barcelona, Spain
b Servicio de Cuidados Intensivos Pediátricos, Hospital Universitario de la Vall d’Hebron, Universidad Autónoma de Barcelona, Barcelona, Spain
c Servicio de Anatomía Patológica, Hospital Universitario de la Vall d’Hebron, Universidad Autónoma de Barcelona, Barcelona, Spain
d Servicio de Cirugía Cardíaca Pediátrica, Hospital Universitario de la Vall d’Hebron, Universidad Autónoma de Barcelona, Barcelona, Spain

*Corresponding author: E-mail address: fgran@vhebron.net (F. Gran).

Available online 14 June 2014

REFERENCES


http://dx.doi.org/10.1016/j.rec.2014.03.006

Nonvalvular Atrial Fibrillation: the Problem of an Undefined Definition

Fibrilación auricular no valvular: el problema de una definición indefinida

To the Editor,

The term nonvalvular atrial fibrillation (NVAF) is used with increasing frequency to describe patients who may benefit from new oral anticoagulants (NOACs). This is a cause for concern for us because the European guidelines for the management of atrial fibrillation, dated 2012, state that “no uniform or satisfactory definition of these terms exists.” The indication for NOACs is based on 4 pivotal studies. To clarify this concept, we have reviewed the inclusion criteria in the protocols of these studies in terms of native valve lesions:

• The RE-LY study2 did not include the term NVAF. Patients with “hemodynamically relevant valve disease” were excluded and, as far as we are aware, a more precise definition was not included.

• The ROCKET trial3 was the only study that included the term NVAF. However, the protocol only excluded patients with “hemodynamically significant” mitral valve stenosis. For the indication of rivaroxaban, atrial fibrillation with a valve lesion other than mitral valve stenosis would not be considered NVAF.

• The authors of the ARISTOTLE trial4 and ENGAGE AF-TIMI 48 trial5 did not use the term. Both these trials excluded only patients with moderate or severe mitral valve stenosis.

A patient with severe aortic stenosis or mitral valve regurgitation with atrial fibrillation would not have been excluded from 3 of the 4 pivotal trials due to valve lesions. It would appear striking and inconsistent to describe such a patient as having NVAF. With valve disease, generalizations are inappropriate. Thus, thromboembolism as a pathophysiologic mechanism for mitral stenosis cannot be considered similar to mitral valve regurgitation or pulmonary stenosis.

The use of a poorly defined term may lead to problems for certain therapeutic indications. To quantify the problem, we reviewed the echocardiography database of a secondary university hospital with no heart surgery facilities. In the last 6 months of