Introduction and objectives. Aortopulmonary septal defect is an uncommon congenital cardiac anomaly. To date, approximately 300 cases have been reported. We present our experience, emphasizing the importance of early correction to avoid irreversible pulmonary hypertension.

Patients and method. Between 1979 and 2000, seven patients underwent surgical repair of this heart defect in our hospital. Two had type I (proximal), 4 had type II (distal) and 1 had type III (complete). Complex associated cardiac anomalies were present in 4 cases: type A interruption of the aortic arch in 2 cases, hypoplastic aortic arch in 1 and transposition of great arteries with ventricular septal defect in 1.

Four cases (57%) were diagnosed by echocardiography. In all patients diagnoses were confirmed by cardiac catheterization.

Patient records were reviewed retrospectively, with special attention to clinical, echocardiographic and hemodynamic data as well as surgical characteristics.

Results. No intraoperative deaths occurred. The patient with associated transposition of great arteries died 22 days after surgery as a result of severe pulmonary hypertension. The remaining patients are asymptomatic without treatment after a mean follow-up period of 69 months.

Conclusions. Even though aortopulmonary septal defect is a rare anomaly, it should be considered whenever the course of complex congenital heart disease includes early cardiac failure and pulmonary hypertension. Repair before 6 months will prevent irreversible damage of pulmonary vessels.

Key words: Aortopulmonary window. Diagnosis. Early correction.
differentiates it from truncus arteriosus (TA).

This rare anomaly represents 0.2%-0.6% of all congenital heart diseases. Since it was first described in 1830 until the present, only about 300 cases have been published, for the most part as isolated reports. Half of the patients have other more or less complex associated cardiac defects (complex windows) that make the diagnosis difficult.

Different modalities of APW exist and several classifications have been proposed. The classification most often used is that of Mori et al, which divides them in type I, or proximal (70%) (the defect is circular, located in a zone equidistant between the sigmoid valve plane and the pulmonary bifurcation); type II, or distal (25%) (of spiral form, it affects the trunk and origin of the right pulmonary artery), and type III (5%) (complete defect of the aortopulmonary septum).

In general, these malformations originate an important left-right shunt with congestive heart failure in the first days or months of life, and early development of severe pulmonary hypertension.

We report our experience with 7 patients diagnosed and treated surgically in our center.

PATIENTS AND METHOD

Between 1979 and 2000, 7 patients underwent surgery for APW. Four were females and 3 were males, the mean age at the time of the intervention was 8 months (range, 11 days to 30 months), and the mean weight was 6.2 kg (range, 2.7-14.8 kg).

All patients underwent a clinical study, chest radiograph, electrocardiogram, cardiac echocardiography, and catheterization (Table 1).

The first two patients were asymptomatic and had been referred to the hospital to study a systolic murmur of moderate intensity located in the upper left parasternal area. In the clinical examination, electrocardiogram, and chest radiograph, the findings were suggestive of left-right shunt. The third patient suffered heart failure at the 3 months of life precipitated by a pulmonary infection. The other

---

**TABLE 1. Echocardiographic and hemodynamic clinical data**

<table>
<thead>
<tr>
<th>Case</th>
<th>Age</th>
<th>Sex</th>
<th>Weight</th>
<th>Clinical presentation</th>
<th>Echocardiography</th>
<th>Hemodynamic study</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Window type</td>
</tr>
<tr>
<td>1/1984</td>
<td>30 months</td>
<td>M</td>
<td>14.5</td>
<td>Study of murmur</td>
<td>Ductus Dilation of left cavities</td>
<td>L (simple) 25/10 2.1 (52%) 1.32</td>
</tr>
<tr>
<td>2/1995</td>
<td>17 months</td>
<td>M</td>
<td>10</td>
<td>Study of murmur</td>
<td>Ao-P window</td>
<td>II (simple) 80/38 1.7 (41%) 3.81</td>
</tr>
<tr>
<td>3/1998</td>
<td>3 months</td>
<td>V</td>
<td>5.4</td>
<td>Heart failure</td>
<td>Large ductus</td>
<td>III (simple) 55/34 7.25(86%) 1.5</td>
</tr>
<tr>
<td>4/1993</td>
<td>7 days</td>
<td>M</td>
<td>2.5</td>
<td>Heart failure</td>
<td>D-TGV VSD</td>
<td>I (complex) D-TGV VSD 65/30 4.5 (84%) 3.4</td>
</tr>
<tr>
<td>5/1991</td>
<td>7 days</td>
<td>V</td>
<td>3.4</td>
<td>Heart failure</td>
<td>Ao-P window</td>
<td>II (complex) Type A AoAI 80/44 4.1 (75%) 4.6</td>
</tr>
<tr>
<td>6/1992</td>
<td>10 days</td>
<td>M</td>
<td>3.5</td>
<td>Cardiogenic shock</td>
<td>Ao-P window</td>
<td>II (complex) Type A AoAI 85/24</td>
</tr>
<tr>
<td>7/2000</td>
<td>2 days</td>
<td>V</td>
<td>3.4</td>
<td>Heart failure</td>
<td>Hypoplastic isthmus Ao RSA OS Pulmonary hypertension</td>
<td>II (complex) AоІ hypoplasia SA OS 60/24 3.73 (73%) 4.5</td>
</tr>
</tbody>
</table>

Ao-P indicates aortopulmonary; RSA, right subclavian artery; Co-Ao, coarctation of the aorta; AoAI, aortic arch interruption; AoI, aortic isthmus; L-R, left-right; OS, ostium secundum.
patients, who had complex windows, showed symptoms of severe heart failure in the first 10 days of life (case 6, cardiogenic shock).

In only 2 cases (3 and 4) was a continuous precordial murmur auscultated, which is described as somewhat lower than that corresponding to ductus and characteristic of the disease.

In 6 patients extracorporeal circulation (ECC) was used and the APW was approached by aortotomy and closed with an internal PTFE patch. In complex windows, the associated anomaly was corrected in the same surgical act. In case 4, closure was by direct suture without EEC, at the same time that the pulmonary artery was banded.

We evaluated the clinical aspects, echocardiographic diagnosis, and their correlation with the hemodynamic study that all patients underwent, as well as the incidents that occurred in the immediate postoperative period and later follow-up.

RESULTS

In Table 1, the clinical, echocardiographic and hemodynamic data of the patients are collected.

The diagnosis was initially made by echocardiography in 4 patients (57%). In case 7, a second echocardiography was made, which was indicated when the patient did not improve after a previous correction of severe coarctation of the aortic isthmus.

In all patients, the diagnosis was confirmed by hemodynamic study. Except for case 1, the patients presented severe pulmonary hypertension with pulmonary pressures that were practically the same as systemic pressures. The left-right shunt was voluminous in complex windows, reaching 86% of pulmonary flow in the absence of the entire aortopulmonary septum (case 3).

Two patients had type I APW, four type II, and one type III.

Four patients (57%) also presented an associated complex cardiac anomaly: type A (AoAI A) interruption of the aortic arch in 2 cases, hypoplasia of the aortic isthmus in one, and D-TGV with VSD in the one. These patients underwent surgery in the first month of life, when they had a mean weight of 3.6 kg (range, 2.7-3.9 kg).

No patient died during the intervention. Patient 4 died 22 days after surgery due to pulmonary hypertension refractory to treatment. Patients 3 and 6 presented minor complications in the immediate postoperative period.

After a mean period of follow-up of 6 years (range, 1 month to 16 years), the patients are asymptomatic without requiring medication and lead normal lives, except for patient 5, who has a moderate maturational delay. During the intervention, this patient had the longest aortic clamping time, reaching 75 min.

DISCUSSION

APW can result from incomplete fusion (type I), poor alignment (type II), or total absence (type III) of the right and left conotruncal cushions, which normally complete conotruncal septation between weeks 5 and 8 of intrauterine life. Recent studies indicate that it does not represent an earlier stage of TA, as had been generally accepted, but that the two anomalies have different origins. The origin of the latter malformation lies in a disturbance of the cells from the neural crest.

In contrast with reports in the literature,6,9 in our series the incidence of type II windows was greater (57%) than that of type I (28%). We had one type III case, which is exceptional.

In half of the patients, APW is associated with other cardiac defects, the most frequent being: AoAI in 15-20% (principally type A), ductus arteriosus in 11%, VSD in 8%, coronary anomalies in 8%, tetralogy of Fallot in 5%, ect.10 AoAI or severe
preductal coarctation of the aorta (considered different stages of same organization) is very rare as an isolated disease. The high incidence found in association with APW could be attributed to the notable decrease in blood flow that the aortic isthmus suffers during the prenatal period in these cases. In TA, the association with type B AoAI is much more common, which coincides with an etiopathogenesis different from that of APW, as mentioned previously.

Until the moment, to our knowledge, only 3 cases of APW associated with TGV have been reported: the first two died, as did our patient, which is the fourth case communicated in the literature.

Both the type of associated lesion and size of the APW condition the clinical manifestations of patients. The diagnosis must be suspected in cases of early heart failure with signs of significant left-right shunt, such as dilation of the left cavities, particularly the left atrium, and/or functional mitral insufficiency with a morphologically normal valve, associated with severe early pulmonary hypertension. The continuous cardiac murmur characteristic of the disease is auscultated in fewer than half of the cases, as occurred in our cases.

The electrocardiographic and radiological findings are non-specific, which is why echocardiography has an important role in diagnosis. The cross-sectional parasternal planes over the aortic valve plane (Figure 1), coronal subcostal plane of both outflow tracts, suprasternal longitudinal and upper parasternal planes are used.

Some false positives are found, particularly when using equipment with scant lateral resolution, because there may be an artificial echo-loss phenomenon (dropout) in the region of the aortopulmonary septum, due to the alignment of the septum in the direction of the lateral resolution of the transducer. To differentiate this phenomenon from true APW, aside from exploring the septum in several planes, some authors resort to the T sign. This sign is nothing other than the greater refringence that the edge of the true defect acquires, which is perpendicular to the rest of the septum and adopts a T image.

The Doppler color study is of inestimable aid and reveals a low-speed bidirectional laminar flow in large, unrestricted defects, with pulmonary hypertension and continuous turbulent flow in the trunk and/or right pulmonary artery. A high-speed flow without pulmonary hypertension is found in small defects.

In our experience, neither the T sign nor Doppler color images provide a definitive diagnosis, as shown in Figure 1. In our series, the diagnostic performance of echocardiography was 57% (4/7), in consonance with reports: 53% (70% in simple APW and 37% in complex APW). In patient 1 (1984) we did not have Doppler color, and in patient 7 an echocardiographic diagnosis was made in the second study, which was performed when the patient did not progress well after a correction was made without a previous hemodynamic study of the associated defect. As reported in the literature, false diagnoses, both positive and negative, are not infrequent. Negative diagnoses are related mainly to the existence of associated complex anomalies that explain the symptoms of the patient. For this reason, we believe that the diagnosis should be confirmed by cardiac catheterization in all cases (Figure 2).

Due to the rapid development of irreversible pulmonary vascular disease, the malformation should be repaired when diagnosed, preferably before 6 months of age. When total pulmonary vascular resistance at the time of the intervention is <8 U/m², the long-term evolution is optimal, as occurred in all our surviving patients.

Since the first correction made by Gross, numerous techniques have been described, with or without ECC, and using a transaortic or transpulmonary patch. The transaortic approach is preferred because it provides better exposure of the window and ostium of the left coronary artery. Matsuki et al and Meissner later described the use of a pulmonary artery flap, later using autologous pericardium to repair the pulmonary artery. Di Bella et al also published a similar technique, but without using a patch to repair the pulmonary artery, which was closed using aortic adventitia, which produces excellent results when autologous tissues with normal growth potential are used.

Some cases of percutaneous closure with favorable results have been reported in specific situations, such as type I, small (3-4 mm) windows not associated with other anomalies, particularly in the origin of the
coronary arteries.\textsuperscript{22,23}

**CONCLUSIONS**

Echocardiography should be the diagnostic method of choice, but it does result in some false positive and negative diagnoses. For this reason, we still confirm the diagnosis by hemodynamic study.

Both the diagnosis and the correction must be made before the age of 6 months to avoid the development of irreversible pulmonary vascular disease.

**DEDICATION**

To Dr. Salud Ariza Almeida, Head of the Cardiology Section, Hospital Infantil Virgen del Rocío of Sevilla. \textit{In memoriam.} With affection from all of us who were once her residents.

**REFERENCES**