Congenital pressure gradients between main pulmonary artery and its primary branches

A. Barrillon, G. Havy, L. Scebat, J. Baragan, and A. Gerboux

From Service de Cardiologie de l'Hôpital Boucicaut, Paris, France

126 cases have been found with pressure gradients between the pulmonary artery trunk and its branches in the course of 594 diagnostic cardiac catheterizations. They occurred either alone: 16 cases (group 1) or more often combined with congenital heart disease: 110 cases (group 2), about half of which were atrial septal defect. Angiographic findings and pressure gradients were correlated in 33 cases. A minor gradient across the junction of the main pulmonary artery with one of the branch pulmonary arteries was frequently present in patients with no angiographic evidence of branch stenosis. These pressure gradients corresponded with pictures of either hypoplasia or localized narrowing of the pulmonary artery branches only when the pressure difference amounted to or exceeded 15 mmHg, in those cases with absent or moderate shunting. This was supported by the finding that the rate of the pressure rise decreased by at least 50 per cent in the low pressure area.

During cardiac catheterization we often found a pressure difference between main and branch pulmonary arteries. The present study aims at assessing whether the finding of a lower pressure in the pulmonary artery branches than in the trunk always corresponds to organic stenosis.

Methods and results

In the course of 2433 consecutive catheterizations for congenital heart disease performed in 10 years at the Boucicaut Hospital, a continuous withdrawal curve from the pulmonary capillary to the main pulmonary artery (PA) was obtained in 594 cases. A systolic pressure gradient was demonstrated in 126 cases (22% of the withdrawal curves). These 126 cases concerned 53 male and 73 female patients whose age averaged 14.5 years (range 3.5 to 66 years). The patients were divided into two groups according to whether this pressure gradient was a lone anomaly (group 1: 16 cases) or combined with other congenital cardiovascular lesions (group 2: 110 cases).

Group 1

Pulmonary artery pressure (Table 1)

A bilateral pressure gradient was found in the 12 cases in which both pulmonary artery branches were explored. In 4 cases, the right branch only was catheterized. The systolic pressure was lower in one or both pulmonary artery branches than the main pulmonary artery by an average of 14 mmHg (range 5 to 90 mmHg). The systolic pulmonary artery pres-

<table>
<thead>
<tr>
<th>Case No.</th>
<th>Main pulmonary artery</th>
<th>Left pulmonary artery</th>
<th>Right pulmonary artery</th>
<th>Angiogram</th>
</tr>
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<tr>
<td>1</td>
<td>20/8</td>
<td>18/10</td>
<td>15/8</td>
<td>Normal</td>
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<tr>
<td>2</td>
<td>25/8</td>
<td>—</td>
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<td>Normal</td>
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<td>4</td>
<td>15/5</td>
<td>—</td>
<td>10/5</td>
<td></td>
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<td>—</td>
<td>17/8</td>
<td></td>
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<td>16/5</td>
<td>16/5</td>
<td></td>
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<td>18/9</td>
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<td>27/7</td>
<td>18/5</td>
<td>18/5</td>
<td></td>
</tr>
<tr>
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<td>27/7</td>
<td>21/7</td>
<td>15/5</td>
<td></td>
</tr>
<tr>
<td>12</td>
<td>24/6</td>
<td>11/4</td>
<td>15/6</td>
<td></td>
</tr>
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<td>13</td>
<td>35/10</td>
<td>18/10</td>
<td>18/10</td>
<td></td>
</tr>
<tr>
<td>14</td>
<td>29/8</td>
<td>—</td>
<td>11/4</td>
<td>Stenosis of both pulmonary artery branches</td>
</tr>
<tr>
<td>15</td>
<td>35/8</td>
<td>15/8</td>
<td>11/8</td>
<td>Multiple stenoses on both pulmonary artery branches</td>
</tr>
<tr>
<td>16</td>
<td>100/5</td>
<td>10/5</td>
<td>20/5</td>
<td>Left pulmonary artery branch hypoplasia</td>
</tr>
</tbody>
</table>

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sure was within normal limits in 13 cases, moderately increased from 30 to 50 mmHg in 2, and much increased to 100 mmHg in 1 case. In 1 case only was the 'pseudoventricular' pattern found in the main pulmonary artery pressure curve. The speed of the pressure rise on either side of the pulmonary artery branch gradient was measured in 14 of the 16 cases and expressed in mmHg per second. A 50 per cent or more diminution of the rate of pressure rise of the pulmonary artery branch as compared with that of the main pulmonary artery was found in 6 cases. In 8 cases, the rate of rise was diminished by less than 50 per cent.

**Standard x-ray signs**

The diameter of the pulmonary artery branches was often normal (11 cases), sometimes diminished (3 cases), and rarely increased (2 cases). The pulmonary vasculature was normal in 12 of 16 cases and no difference was observed between the two lung fields.

**Angiocardiography**

Right-sided angiocardiography carried out in 5 of 16 cases did not show any narrowing in 2 cases (Cases 1 and 2 with pressure gradients of 5 mmHg). Stenosis was demonstrated in 3 patients.

*Case 13* Stenosis of both pulmonary artery branches (pressure gradient of 17 mmHg on the right and left sides).

*Case 15* Multiple stenoses on both right and left pulmonary artery branches (pressure gradient of 24 mmHg on the right side and 20 mmHg on the left) (Fig. 1).

*Case 16* Pronounced dilatation, almost aneurysmal, of the trunk with a hypoplastic left branch and a normal right branch; the upper and middle lobar divisions of the latter had a diameter which diminished gradually in size (pressure gradient of 90 mmHg on the left side and 80 mmHg on the right) (Fig. 2).

**Group 2**

110 cases of pulmonary artery branch pressure gradient combined with other congenital cardiovascular disease.

**Pulmonary artery pressures** (Table 2)

Both pulmonary artery branches were explored in 33 cases and a bilateral pressure gradient was found in 25 cases, and a unilateral one, right-sided only, in 8. In 77 cases one pulmonary artery branch only could be catheterized, which makes it impossible to provide information on bilateral pressure gradients. The pulmonary artery branch systolic pressure was found to be within normal limits in 36

![FIG. 1 A man aged 21 (Case 15 group 1): multiple stenoses on both right and left pulmonary artery branches. Pressure gradients 24 mmHg on the right side and 20 mmHg on the left.](image)
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**Fig. 2** A girl aged 20 (Case 16 group I): pronounced, almost aneurysmal, dilatation, of the main pulmonary artery with a hypoplastic left branch and a normal right branch. Pressure gradient of 90 mmHg on the left and 80 mmHg on the right side.

**Table 2** Average of systolic pulmonary artery pressures (mmHg) in 110 cases of group 2 (with combined cardiac anomalies): in brackets number of cases on which average was computed.

<table>
<thead>
<tr>
<th>Associated anomaly</th>
<th>No. of cases</th>
<th>Main pulmonary artery</th>
<th>Left pulmonary artery</th>
<th>Right pulmonary artery</th>
</tr>
</thead>
<tbody>
<tr>
<td>Atrial septal defect</td>
<td>59</td>
<td>31</td>
<td>25 (15)</td>
<td>20 (58)</td>
</tr>
<tr>
<td>Ventricular septal defect</td>
<td>12</td>
<td>26</td>
<td>12 (2)</td>
<td>20 (10)</td>
</tr>
<tr>
<td>Pulmonary valvar stenosis*</td>
<td>17</td>
<td>37</td>
<td>24 (10)</td>
<td>20 (16)</td>
</tr>
<tr>
<td>Obstructive cardiomyopathy</td>
<td>6</td>
<td>31</td>
<td>22 (1)</td>
<td>22 (5)</td>
</tr>
<tr>
<td>Coarctation of aorta</td>
<td>4</td>
<td>32</td>
<td>24 (1)</td>
<td>21 (4)</td>
</tr>
<tr>
<td>Partial anomalous pulmonary drainage</td>
<td>4</td>
<td>35</td>
<td>30 (2)</td>
<td>22 (4)</td>
</tr>
<tr>
<td>Persistent atrioventricular canal</td>
<td>3</td>
<td>33</td>
<td>24 (1)</td>
<td>21 (3)</td>
</tr>
<tr>
<td>Tetralogy of Fallot</td>
<td>1</td>
<td>35</td>
<td>—</td>
<td>25</td>
</tr>
<tr>
<td>Supravalvar aortic stenosis</td>
<td>1</td>
<td>25</td>
<td>—</td>
<td>20</td>
</tr>
<tr>
<td>Communication between right atrium and left ventricle</td>
<td>1</td>
<td>30</td>
<td>—</td>
<td>18</td>
</tr>
<tr>
<td>Single ventricle</td>
<td>1</td>
<td>45</td>
<td>29</td>
<td>24</td>
</tr>
<tr>
<td>Persistent ductus</td>
<td>1</td>
<td>33</td>
<td>—</td>
<td>11</td>
</tr>
</tbody>
</table>

* With superadded ventricular defect in 5 cases, atrial septal defect in 2, atrioventricular canal in 1 case.

X-rays

These frequently showed (60 cases) increased shadows of the pulmonary artery branches with congestion of both lung fields, easily accounted for by the presence of a shunt. In 77 cases, there was a difference in the vasculature between the two lung fields; in 6 cases decreased vascular shadows were noted on the same side as the gradient, but in 8 cases, larger vascular shadows were found on the same side as the pressure gradient; finally in 3 cases there were asymmetrical vascular shadows when the pressure gradients were identical on both sides.

Pulmonary angiogram

This was normal in 17 of the 28 cases. In 11 cases the pulmonary artery branches were abnormal:
there was hypoplasia of the pulmonary artery tree in 5 cases, coarctation of both branches in 2 cases, coarctation of the right branch in 2 cases, hypoplasia of the left branch with stenosis of the division branches and stenosis with post-stenotic dilatation of the right branch in 1 case, and coarctation of both main pulmonary artery branches with narrowed distal branches in 1 case.

**Surgical findings**

Operation was attempted in 2 cases of pulmonary artery branch stenosis; in one case both branch stenoses were dilated with a dilatator; in one case a 'teflon' patch was inserted on the intermediate trunk of the right branch.

**Pathological findings**

One case aged 4 died on the day of catheterization, which showed an ostium primum with a single ventricle of the right type. There were no definite coarctations of the pulmonary artery branches but rather a hypoplasia of the distal branches which were thin, soft, and transparent, very different in texture from that of the main pulmonary artery. Yet on catheterization the systolic pressures were 45 mmHg in the pulmonary artery trunk, 29 mmHg

FIG. 3 Rate of rise of both proximal and distal pulmonary artery pressure pulses on either side of the 'stenosis' compared with the angiographic data in 29 patients. Left: cases with pulmonary artery branch stenosis or hypoplasia; right: normal angiograms. Slowing of the rate of rise of the pulmonary artery pressure pulse is much more pronounced in those cases with abnormal than with normal pulmonary artery branches. One case of the latter group was an exception.
Congenital pressure gradients between main pulmonary artery and its primary branches

Pressure gradients between the main pulmonary artery and its primary branches have been recorded with increasing frequency during catheterization for congenital heart disease (Eldridge, Selzer, and Hultgren, 1957; Bell et al., 1962; Franch and Gay, 1963; Gasul, Arcilla, and Lev, 1966; Latour et al., 1970). Axillary transmission of the systolic murmur, found in this series in 59 of 65 cases in which it was looked for, was the most reliable physical sign (Bouvrain, Bouroumieux, and Nezry, 1961; Oram, Pattinson, and Davies, 1964; Perloff and Lebauer, 1969). A difference in density between the two lung fields on the x-ray picture, obvious in only 17 of the 126 of this series, is an inconstant finding. From this, it is impossible to predict the site of a pressure gradient. Increased pulmonary vascular shadows were found on the same side as the pulmonary artery branch pressure gradient in 11 cases. Others (Gyllensward et al., 1957; Nadas, 1963; Fouron et al., 1967) have reported similar findings. According to Fouron et al. (1967) this change is related to age. Catheterization of the pulmonary circulation provides direct evidence of a pressure gradient in the pulmonary pathway. But selective angiography of the main pulmonary artery or the right ventricle is the only reliable procedure for demonstration of organic stenosis (Baum et al., 1964). A group of cases with an obstruction on the

Discussion

Pressure gradients between the branches and the trunk of the pulmonary artery have been recorded with increasing frequency during catheterization for congenital heart disease (Eldridge, Selzer, and Hultgren, 1957; Bell et al., 1962; Franch and Gay, 1963; Gasul, Arcilla, and Lev, 1966; Latour et al., 1970). Axillary transmission of the systolic murmur, found in this series in 59 of 65 cases in which it was looked for, was the most reliable physical sign (Bouvrain, Bouroumieux, and Nezry, 1961; Oram, Pattinson, and Davies, 1964; Perloff and Lebauer, 1969). A difference in density between the two lung fields on the x-ray picture, obvious in only 17 of the 126 of this series, is an inconstant finding. From this, it is impossible to predict the site of a pressure gradient. Increased pulmonary vascular shadows were found on the same side as the pulmonary artery branch pressure gradient in 11 cases. Others (Gyllensward et al., 1957; Nadas, 1963; Fouron et al., 1967) have reported similar findings. According to Fouron et al. (1967) this change is related to age. Catheterization of the pulmonary circulation provides direct evidence of a pressure gradient in the pulmonary pathway. But selective angiography of the main pulmonary artery or the right ventricle is the only reliable procedure for demonstration of organic stenosis (Baum et al., 1964). A group of cases with an obstruction on the
pulmonary artery pathway should be distinguished from a second group in which no such stenosis could be demonstrated.

Out of the 33 cases in which pulmonary angiography was carried out, 14 demonstrated a pulmonary artery branch hypoplasia (5 cases) or localized narrowing with coarctation of one or two branches (9 cases), a fact reported by Arvidsson, Karnell, and Moller as early as 1955 (Arvidsson et al., 1961). The alleged specificity of post-stenotic dilatation (Nadas, 1963) was not confirmed in this series. Two cases only were operated upon. This is possible when the stenosis is proximal (Baxter, Booth, and Sirak, 1961; McGoon and Kincaid, 1964) and desirable when the main pulmonary artery pressure is increased (Dubost, Cachera, and Vigano, 1967) (62 and 37 mmHg in the cases operated on). The severity of obstruction to pulmonary blood flow does not increase with age (Eldredge et al., 1972). Necropsy is exceptional, as this anomaly is usually well tolerated. Performed once in this series, it showed an abnormal structure of the elastic wall responsible for a vessel wall thinning. Other lesions have been described as intimal hyperplasia of the main branches simulating atheromatous plaques (Oppenheimer, 1938; Esterly and Oppenheimer, 1967). The arguments in favour of organic pulmonary artery branch stenosis are as follows.

**Clinical picture**—History of German measles in the mother combined with a reduced auditory acuity was found in 2 cases of this series, cataract was found in 1 further case (Oram et al., 1964; Esterly and Oppenheimer, 1967; Rowe and Mehrizi, 1970). Combined supravalvar aortic stenosis may be suspected by the peculiar facies and mental retardation with a history of hypercalcaemia (Beuren, Apitz, and Harmianz, 1962; Bourassa and Campeau, 1963).

*Pseudoventricular* pattern of the main pulmonary artery pressure curve was thought by Agustsson et al. (1962) to be pathognomonic of bilateral pulmonary artery branch stenosis; the more pronounced, the tighter the stenosis. It is characterized by a rapid systolic rise followed by a diastolic fall to low early diastolic pressure, followed by a mid or late diastolic plateau. In this series, 16 of the 49 cases with this pattern had angiograms which demonstrated pulmonary artery stenosis in 9 cases only. In this series this pressure curve pattern was thus absent in cases with severe stenosis while it was recorded in cases with mild gradients but without angiographic evidence of stenosis. According to Delaney and Nadas (1964) and Nadas (1963) this should be considered a suggestive though unreliable sign.

**Speed of pulmonary artery pressure rise** on either side of the pressure gradient appears to be a better sign of organic stenosis of the pulmonary artery branches. Rios et al. (1969) have shown this pressure rise in a series of 17 cases to be 50 per cent slower in the post-stenotic area than in the proximal area. In the absence of organic stenosis, these speeds were identical on either side of the pressure gradient. This sign was found in the present series in 30 of 126 cases. Of the 29 cases with angiograms, this sign coincided commonly with organic stenosis (10 of 15 cases). Conversely, stenosis was found only rarely (1 of 14 cases) in the absence of this sign. This 50 per cent slowing of the post-stenotic speed of the pulmonary artery pressure rise seemed to be a reliable sign (Fig. 3) and deserves to be retained as a good indication of organic stenosis.

**Magnitude of pressure gradient**—The presence of branch stenosis on the angiograms coincided with pressure gradients of 15 mmHg or more, in the absence of shunt (group 1). However, in the presence of left-to-right shunts, pressure gradients larger than 15 mmHg may be found in the absence of stenosis (Williams, Lange, and Hecht, 1957). In other words, a pressure gradient below 15 mmHg is unlikely to be caused by stenosis, while in the cases in which it exceeds 15 mmHg a true stenosis is a possibility which requires to be proved by an angiogram (Fig. 4).

In the remaining 19 cases (58%) the angiograms did not reveal any stenosis. To account for the presence of pressure gradients in the absence of an angiographic picture of stenosis, three hypotheses may be put forward:

**Increase of the pulmonary artery flow resulting from a left-to-right shunt** (Deyrieux, Tartutilier, and Tourniaire, 1961): On the one hand, pressure gradients have been found in heart diseases without shunts (group 1), and on the other, when these pressure gradients occur in heart diseases with a left-to-right shunt (group 2), the latter is often mild (pulmonary artery output/systemic output ratio below 2) (Fig. 4). Surgical closure of the shunt resulted in abolition of the systolic murmur in only 21 of 55 cases. Finally, numerous cases with large left-to-right shunts never demonstrated such gradients.

**Presence of mild stenosis invisible to angiography.**—Persistence or aggravation of physiological pressure differences between main and branch pulmonary arteries in infants (Danilowicz et al., 1972). Fouron et al. (1967) and Rowe and Mehrizi (1970) have demonstrated in the normal infant a pressure gradient of 3 to 7 mmHg between the main pulmonary artery and its branches. Danilowicz et al. (1972) have shown the disappearance in infants of systolic

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1 In 4 cases with angiograms, the speed of the pressure rise could not be measured.
pressure differences of 6 to 45 mmHg. Increase of the pulmonary artery blood flow, especially in the presence of a left-to-right shunt, would result in elastic tissue changes, with consequent pressure gradients between the main and branch pulmonary arteries, with or without organic stenosis.

References


Requests for reprints to Dr. A. Barrillon, Service de Cardiologie de l'Hôpital Boucicaut, 78 rue de la Convention, 75015 Paris, France.
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