Coarctation of aorta with right aortic arch

Report of surgical correction in 2 cases: one with associated anomalous origin of left circumflex coronary artery from the right pulmonary artery

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Two patients with right aortic arch, anomalous left subclavian artery, and coarctation of the aorta, have had operations for the relief of coarctation, successful in one but only partially successful in the other: in one case, a 'dacron' patch was inserted into the narrowed segment, but a mild recurrence of coarctation was found a year later; in the other, the proximal segment of the anomalous left subclavian artery was used for the repair, with complete relief of the coarctation, both immediately and at late follow-up.

The first case had multiple associated congenital abnormalities, including anomalous origin of the left circumflex coronary artery from the right pulmonary artery, which was also corrected surgically: this anomaly has only been recorded once hitherto. The second had facial haemangiomata and a small ventricular septal defect.

Four previously reported cases of coarctation and right arch are reviewed. These 6 cases are considered in the light of current views on the development and classification of aortic arch anomalies.

The occurrence of coarctation in patients with right aortic arch is extremely rare: hitherto only 4 cases have been recorded. We have seen 2 further cases; in both of these, an attempt was made to correct the coarctation surgically.

Case reports

Case 1

A 13-year-old Arab boy from Libya was referred for treatment of coarctation of the aorta. He complained of fatigue and effort dyspnoea but his disability was slight. There was no dysphagia or stridor.

On examination, he was undersized. There was no cyanosis or clubbing. The left arm pulses were reduced; the blood pressure in the right arm was 190/130 mmHg (25.3/17.3 kPa) and in the left arm 110/70 mmHg (14.6/9.3 kPa). The carotid pulses were forceful and equal; both femoral pulses were grossly reduced and delayed. The cardiac impulse suggested left ventricular hypertrophy; there was a basal ejection systolic murmur.

FIG. 1 Case 1. Chest x-ray.
Intravenous pyelogram showed a normal left kidney, and no excretion on the right side. Chromosome studies gave normal results.

**Cardiac catheterization** (Table 1) Right heart pressures were normal, but there was a small left-to-right shunt at pulmonary artery level (flow ratio 1.2:1). An arterial catheter introduced from the right brachial artery entered the aortic arch and passed downwards through the coarcation, but was then manoeuvred into the ascending aorta and left ventricle. A second arterial catheter, introduced by right femoral arteriotomy, entered the left subclavian artery from below.

**Angiocardiography** (Fig. 3) Arch aortogram. Injection into the ascending aorta showed a right-sided aortic arch and descending aorta. The left common carotid artery, right common carotid artery, and right subclavian artery arose in turn from the arch; there was a moderately severe coarcation 2 cm in length just distal to the origin of the right subclavian artery (Fig. 3a). There was a very large and tortuous left main and anterior descending coronary artery, with later opacification of a vessel entering the pulmonary artery (Fig. 3b). Late films showed retrograde filling of the left subclavian artery from the left vertebral artery and thyrocervical trunk (Fig. 3c). The aortogram also showed a bicuspid aortic valve with minimal aortic regurgitation.

**Selective coronary arteriograms** showed a small but normal right coronary artery. The left coronary artery gave rise to a large anterior descending artery with diagonal and septal branches, anastomosing with distal circumflex branches; there was then retrograde filling of the main trunk of the circumflex artery, and puffs of contrast were seen to enter the posterior aspect of the right pulmonary artery.

**Abdominal aortogram** showed a normal left renal artery supplying a normal but large left kidney. No right renal artery was seen.

### TABLE 1 Preoperative cardiac catheterization data

<table>
<thead>
<tr>
<th>Case No., age, and sex</th>
<th>Pressures (mmHg)*</th>
<th>Oxygen saturation (per cent)</th>
<th>Flow (\text{l/min per m}^2) BSA</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>(\text{RA mean s/d})</td>
<td>(\text{PA wedge s/d})</td>
<td>(\text{LV s/d})</td>
</tr>
<tr>
<td>Case 1</td>
<td>13 yr M</td>
<td>3</td>
<td>24/14</td>
</tr>
<tr>
<td>Case 2†</td>
<td>1 yr 10 mth M</td>
<td>2</td>
<td>25/13</td>
</tr>
</tbody>
</table>

* Reference zero: mid-thorax.
† General anaesthesia. Inspired oxygen concentration 25 per cent.

Abbreviations: RA, right atrium; PA, pulmonary artery; LV, left ventricle; RV, right ventricle; SVC, superior vena cava; s, d, ed, systolic, diastolic, end-diastolic.

Conversion factor from Traditional Units to SI units: 1 mmHg ≈ 0.133 kPa.

Operation (1 May 1973) The chest was opened by a left lateral thoracotomy. The oesophagus was situated to the left of the descending aorta; there was post-stenotic dilatation of the aorta, from which arose the left subclavian artery; a left-sided ligamentum arteriosum connected the descending aorta to the left pulmonary artery. Retraction of the oesophagus revealed the coarcted segment 3 cm in length between the origins of the right and left subclavian arteries. This was exposed, and after cross-clamping the aorta proximally and distally, the coarcted segment was opened longitudinally. An onlay patch of woven ‘dacron’ was sewn on to the opened segment (Fig. 4A). Resection and end-to-end anastomosis was not feasible because the proximal limit of the coarctation lay in the right side of the mediastinum. The ligamentum was divided. A left superior vena cava was also noted.

Postoperative course Systemic hypertension required treatment initially with trimetaphan camyslate (Arfonad) infusion and intermittent intravenous chlorpromazine, and later with oral methylldopa. After the repair procedure the left leg pulses were normal, but in the right leg no pulse was palpable below the femoral. This was attributed to thrombosis at the site of preoperative femoral arteriotomy. After exploration and thrombectomy, an arterial catheter was passed retrogradely across the previously coarcted segment; no gradient was demonstrated and angiography confirmed successful relief of coarctation; a small false aneurysm was seen at the distal end of the repair (Fig. 3d). Subsequently all lower limb pulses were normal. Blood pressure averaged 140/90 mmHg (18.6/12.0 kPa).

He remained well, but was readmitted a year later for correction of the coronary artery anomaly. He was found to have signs of recurrence of coarctation. The femoral pulses were reduced and delayed. The right arm blood pressure was 170/100 mmHg (22.6/13.3 kPa), but in the left arm the pulses were reduced and blood pressure was 100/80 mmHg (13.3/10.6 kPa).

Arch aortogram (Fig. 3e) confirmed that there was now significant narrowing at the site of the repair. This was associated with a gradient of 45 mmHg (6.0 kPa) between the arch and descending aorta.

It was not thought feasible to repeat the onlay graft procedure; correction by insertion of a tube graft from ascending to descending aorta was considered as an alternative method of treatment, but this further extensive surgery was thought to be unjustified.

Second operation (5 July 1974) The heart was exposed through a median sternotomy. The anterior descending coronary artery was tortuous, but the circumflex artery was inaccessible. On routine cardiopulmonary bypass, the pulmonary trunk was opened in its long axis, and the orifice of the anomalous circumflex coronary artery was seen in the posterior wall of the first centimetre of the right pulmonary artery. This opening was closed by direct suture.

Postoperative course He made an uninterrupted recovery. The signs of coarctation remain. Electrocardiogram 5 days after operation showed slight peri-cardial ST elevation but no change in QRS pattern. Later tracings showed widespread postoperative T wave inversion, but these changes had disappeared completely within 3 months.

Case 2
An English boy aged 22 months was found to have a cardiac murmur on routine medical examination in infancy. His growth and development were normal, and there were no symptoms; there was no dysphagia or dyspnoea.

On examination he appeared healthy, and height and weight were normal for age. He had a large haemangioma on the right cheek and right side of the neck, and smaller ones on the lips and chin. The left arm pulses were reduced; the blood pressure was 160 mmHg (21.3 kPa) in the right arm and 80 mmHg (10.6 kPa) in the left arm by the Doppler shift method (Elseed, Shinebourne, and Joseph, 1973). The femoral pulses were weak and delayed. The heart was not enlarged; there was a pansystolic murmur (grade 4/6) at the left border of the sternum.

The chest radiograph showed moderate cardiac enlargement and pulmonary plethora; the main pulmonary trunk was prominent and the aortic knuckle invisible. The electrocardiogram was normal for age.

Cardiac catheterization (Table 1) Right heart pressures were normal. A left-to-right shunt at ventricular level was shown, with flow ratio 1.5:1. A right subclavian vein sample was 92 per cent saturated; this finding was attributed to the large haemangioma. An arterial catheter was passed retrogradely from the right brachial artery to the aortic arch and left ventricle, but the coarctation was not traversed.

Left ventricular cineangiogram in the left anterior oblique position showed a small infracristal ventricular septal defect and a short coarctation proximal to the left subclavian artery. The abnormal position of the aorta and origin of the other brachiocephalic vessels were not clearly demonstrated and were not recognized.

Operation (20 July 1973) The chest was opened by a left lateral thoracotomy. The anatomical findings were similar to the first case: there was a right aortic arch with aberrant left subclavian artery arising distal to the coarcted segment; a left ligamentum arteriosum completed a loose vascular ring. The other brachiocephalic arteries arose in the same order, viz. left common carotid, right common carotid, and right subclavian artery, proximal to the coarctation. After cross-clamping, the aorta was opened in its long axis; the left subclavian artery was also incised along its long axis and laid into
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the coarcted segment to achieve an angioplastic repair (Fig. 4B). The left subclavian artery was divided and ligated distally, as also was the ligamentum.

Postoperative course His postoperative progress was entirely uneventful.

He remains well one year after operation. The blood pressure in the right arm is 120/70 mmHg (16.0/9.3 kPa), and in the right leg 110/70 mmHg (14.6/9.3 kPa), but in the left arm 70 mmHg (10.6 kPa). The ventricular septal defect murmur is unchanged.

Discussion

Right aortic arch: developmental considerations and classification

Stewart, Kincaid, and Edwards (1964) have proposed a classification of aortic arch anomalies. Their group III consists of patients with right aortic arch, and they subdivide this group according to the mode of origin of the brachiocephalic trunks. This may be the mirror-image of normal so that left innominate, right common carotid, and right subclavian arteries arise in turn from the aortic arch (Type IIIA); the left subclavian artery may be aberrant, arising from the descending aorta, with left common carotid, right common carotid, and right subclavian arteries arising in turn from the arch (Type IIIB); finally, the left subclavian artery may have no direct origin from the aorta (Type IIIC). Each type of right aortic arch may be associated with a left-sided ductus (subgroup 1), a right-sided ductus (subgroup 2), or bilateral ductus (subgroup 3). Blake and Manion (1962) in an analysis of 41 cases classify aortic arch anomalies in relation to abnormal sites of involution of the primitive aortic arch system. Involution of the left and persistence of the right dorsal aortic root give rise to Type IIIA of Stewart et al. (1974) with mirror-image branching. Involution of the left fourth arch and persistence of the left and right dorsal aortic roots give rise to Type IIIB with anomalous origin of left subclavian artery. Involution of the left ventral aortic root gives rise to a situation not recognized by Stewart et al. (1964) in which there is anomalous origin of a left innominate artery from the descending aorta.

The primitive aortic arch system is illustrated diagrammatically in Fig. 5A. The sites at which involution occurs to produce the observed patterns of branching in the normal subject and in patients with a right aortic arch are shown. The resulting postnatal right arch patterns are shown in Fig. 5C and D, with the normal for comparison in Fig. 5B.
TABLE 2  Coarctation and right aortic arch: anatomical features and associated lesions in 6 cases

<table>
<thead>
<tr>
<th>Authors</th>
<th>Age and sex</th>
<th>Type of right arch</th>
<th>Pattern of branching</th>
<th>Ductus or ligamentum (vascular ring + or −)</th>
<th>Upper descending aorta</th>
<th>Operation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Felson and Palayew</td>
<td>Child</td>
<td>‘Posterior’*</td>
<td>LCCA</td>
<td>Not stated</td>
<td>Not stated</td>
<td>Yes (no details)</td>
</tr>
<tr>
<td>Stewart, Kincaid, and Edwards</td>
<td>16 yr M</td>
<td>Anomalous left subclavian artery (IIB)</td>
<td>LCCA, RCCA, RSA</td>
<td>Left ductus (+)</td>
<td></td>
<td>Unknown (‘difficult’: no details)</td>
</tr>
<tr>
<td>Grossman and Jacoby</td>
<td>14 yr F</td>
<td>Anomalous left subclavian artery (IIB)</td>
<td>LCCA, RCCA, RSA</td>
<td>Unknown</td>
<td>Right</td>
<td>Not attempted</td>
</tr>
<tr>
<td>Price and Schieken</td>
<td>13 yr F</td>
<td>Mirror-image</td>
<td>L. innom. A, —</td>
<td>Not stated</td>
<td>Right</td>
<td>Successful resection and end-to-end anastomosis</td>
</tr>
<tr>
<td>Case 1</td>
<td>13 yr M</td>
<td>Anomalous left subclavian artery (IIB)</td>
<td>LCCA, RCCA, RSA</td>
<td>Left ligamentum (+)</td>
<td>Right</td>
<td>Partially successful (onlay graft) repair</td>
</tr>
<tr>
<td>Case 2</td>
<td>1 yr 10 mth M</td>
<td>Anomalous left subclavian artery (IIB)</td>
<td>LCCA, RCCA, RSA</td>
<td>Left ligamentum (+)</td>
<td>Left</td>
<td>Successful repair (angioplastc)</td>
</tr>
</tbody>
</table>

* Type of aortic arch abnormality not classifiable by the scheme of Stewart, et al (1964).
† Right subclavian artery arising at site of coarctation (case of Price and Schieken).
Abbreviations: LCCA, RCCA: left and right common carotid arteries; LSA, RSA: left and right subclavian arteries; SVC: superior vena cava.

The coexistence of a right arch and anomalous left subclavian artery passing behind the oesophagus with left-sided ductus or ligamentum results in the formation of a type of vascular ring, usually but not always a loose one (Stewart et al, 1964; Lincoln et al., 1969). It is noteworthy that in almost all cases, the mirror-image type of branching of a right arch is associated with cyanotic congenital heart disease, commonly Fallot’s tetralogy or persistent truncus arteriosus; on the other hand, in group IIB of Stewart et al. (1964), with right arch and aberrant left subclavian artery, intracardiac anomalies are unusual and coincidental.

**Right aortic arch and coarctation**

The association of right aortic arch with coarctation is very unusual. Four cases have been reported (Felson and Palayew, 1963; Stewart et al., 1964; Grossman and Jacoby, 1969; Price and Schieken, 1974) in addition to the 2 described here (Table 2).

Our 2 cases, the case of Stewart et al. (1964), and that of Grossman and Jacoby (1969) appear to be identical, in possessing right aortic arch, anomalous left subclavian artery arising from the descending aorta distal to the coarctation, and left common carotid, right common carotid, and right subclavian arteries arising in that order from the arch (Type IIB). This situation is the mirror-image of coarctation of a normal left arch, with anomalous right subclavian artery. Grossman and Jacoby present their case as a ‘unique anatomical variation’, though it appears to differ in no way from the case of Stewart et al. (1964). In Felson and Palayew’s case (1963) not only was the left subclavian artery aberrant but the left common carotid artery also arose distal to the coarctation; on embryological grounds one would expect these vessels to arise together as an aberrant left innominate artery (Blake and Manion, 1962), but this is neither stated nor denied. In the case reported by Price and Schieken (1974) the pattern of branching was the mirror-image of normal (Type IIIA) and was unusual in being unassociated with cyanotic congenital heart disease.
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Associated lesions

<table>
<thead>
<tr>
<th>Cardiovascular</th>
<th>Other systems</th>
</tr>
</thead>
<tbody>
<tr>
<td>None (?)</td>
<td>Not stated</td>
</tr>
<tr>
<td>Not stated (?)</td>
<td>Not stated (?)</td>
</tr>
<tr>
<td>None</td>
<td>None</td>
</tr>
<tr>
<td>Aortic valve stenosis (operation)</td>
<td>Turner’s syndrome</td>
</tr>
<tr>
<td>Anomalous origin of left circumflex coronary artery (operation), bicuspid aortic valve, bilateral SVC</td>
<td>Absent right kidney, skeletal anomalies</td>
</tr>
<tr>
<td>Ventricular septal defect</td>
<td>Facial haemangiomata</td>
</tr>
</tbody>
</table>

A left ductus or ligamentum was recognized in 3 of the cases; this resulted in the formation of a loose asymptomatic vascular ring. In the other 3 cases, it was either unknown or not stated on which side the ductus or ligamentum was situated. In our Case 1 and in 2 others, the upper descending aorta is right-sided, but in Case 2, the right arch, having passed behind the oesophagus, continues as a left-sided descending aorta; as a result the appearance at left thoracotomy differed so that in Case 1 the descending aorta was concealed by the oesophagus whereas the converse was true in Case 2.

We cannot suggest any plausible mechanism to account for the site of coarctation in these cases. This is neither juxta ductal coarctation nor the isthmal narrowing not uncommonly seen in left-to-right shunt situations (Rudolph, Heymann, and Spitznas, 1972; Shinebourne and Elseed, 1974), but partial involution of the terminal segment of the primitive right dorsal aortic root, distal to the origin of the seventh dorsal intersegmental artery (which gives rise to the right subclavian artery). This has resulted in the appearance of a coarcted segment distal to the origin of the right subclavian artery in all cases, and in 5 of the 6 cases proximal to an aberrant left subclavian artery (and left common carotid artery in the case of Felson and Palayew, 1963).

Associated lesions

In 1 case certainly and probably in 2 others there was no associated cardiovascular lesion. One who had Turner’s syndrome with characteristic chromosomal abnormality had aortic valve stenosis of sufficient severity to require surgical relief (Price and Schieken, 1974). Our Case 1 has a bicuspid aortic valve, bilateral superior vena cava with variable P wave axis on the electrocardiogram, and an unusual congenital coronary arterial anomaly. He also has minor skeletal anomalies, and congenital absence of the right kidney. Our Case 2 has a small ventricular septal defect. None of the cases had any form of conotruncal abnormality.

Surgical treatment

Surgical correction of coarctation was clearly indicated in both our cases, in the second of whom the presence of a right arch was unsuspected before operation. Because of the difficult access to the proximal end of the coarcted segment of the posterior aortic arch, resection with end-to-end anastomosis was not technically feasible. In both cases, the coarcted segment was opened longitudinally and its internal diameter increased, in Case 1 with a ‘dacron’ onlay graft, and in the other by using the proximal portion of the aberrant left subclavian artery for the same purpose. When Case 1 was re-examined a year after operation it was clear that coarctation had recurred and this was confirmed by pressure measurements and aortography; it was thought that any further attempt to correct the coarctation would be a difficult and hazardous procedure. In Case 2, normal femoral pulses and normal right arm and lower limb blood pressure 13 months after operation confirm the adequacy of the repair procedure.

Unfortunately no details of the operation are given in 2 of the other cases (Felson and Palayew, 1963; Stewart et al., 1964), but in Price and Schieken’s case (1974) resection and end-to-end anastomosis were successfully accomplished in a different anatomical situation. In the sixth case (Grossman and Jacoby, 1969), surgical repair was not attempted, as this was thought to pose technically insuperable problems.
Anomalous origin of left circumflex coronary artery

We have been able to find only one published report of anomalous origin of the circumflex branch of the left coronary artery from the pulmonary artery, in a patient in whom the anterior descending artery arose from the right artery (Effier et al., 1967: Case 7). In the Bland-Garland-White syndrome, the main left coronary artery arises from the pulmonary artery, and one artery (the right) arises from the aorta. This syndrome has been the subject of several reviews, but no reference is made to any other patient in whom the left anterior descending or circumflex artery alone arises from the pulmonary artery. The published reports on congenital coronary arterial fistula contain several cases in which there is a fistula into the pulmonary artery or one of its main branches. It is presumed, but not always stated explicitly, that there are normal proximal right, left anterior descending, and circumflex arteries. Possibly cases of anomalous origin of the left circumflex coronary artery may have masqueraded as coronary fistulae terminating in the pulmonary artery.

Our Case 1 had a small but normal right coronary artery. The left coronary artery had no circumflex branch but continued in the anterior interventricular groove, giving rise to normally situated but large diagonal and septal branches, from which drainage occurred into the circumflex territory, and thence into a single large vessel entering the right pulmonary artery. This lesion constituted a low pres-
sure run-off from the coronary circuit and gave rise to a small but measurable left-to-right shunt. There was no clinical or electrocardiographic evidence of impaired myocardial perfusion, but it was thought rational to correct the anomaly either by dividing the anomalous circumflex artery or by reimplanting it into the aortic root. The anomalous vessel was closed from within the pulmonary artery and, as would be expected, this procedure was well tolerated. The case of Effler et al. (1967) was also treated by division of the anomalous circumflex artery at its pulmonary arterial end (D. B. Effler, 1974, personal communication).

References

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