Anomalous right subclavian artery and coarctation of the aorta

Surgical implications and the use of the right subclavian artery as a flap

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SUMMARY Coarctation of the aorta was repaired surgically in a 4 month old boy using the anomalous right subclavian artery as a flap. The role of the anomalous artery in providing collateral flow to the descending aorta should be determined so that injudicious clamping is avoided.

The anomalous right subclavian artery is the single most common abnormality of aortic arch development and is found at approximately one out of every 200 necropsies. Maude Abbott reported an incidence of 1% (2/200 patients) of an anomalous right subclavian artery associated with a postductal aortic coarctation. In our experience two out of 106 patients have had this anomaly; the first case has been reported previously and the second is reported here. To our knowledge, it is the first report of the use of the anomalous right subclavian artery as a flap to repair the coarctation.

Case report

A 7 week old boy had a heart murmur at a routine postnatal visit. The mother admitted that the child was breathless when feeding. On examination he was tachypnoeic but not cyanosed. The radial pulses were easily felt, but the leg pulses were not palpable. At the left sternal border an ejection systolic murmur and a loud pulmonary component of the second heart sound were heard. There was a short mid-diastolic murmur at the mitral area. The chest radiograph showed an enlarged heart and pulmonary plethora. The child was diagnosed as having a coarctation with a possible ventricular septal defect. Treatment for cardiac failure was started.

At 3 months of age cardiac catheterisation showed a non-restrictive ventricular septal defect and severe pulmonary hypertension (pulmonary artery pressure 95/30 mm Hg, pulmonary vascular resistance 2.7 units/m²). The aortogram showed the right carotid artery, left carotid artery, and left subclavian artery arising separately and in that order from the aortic arch. The right subclavian artery arose distal to the left subclavian and just proximal to the site of a pronounced coarctation. The distal descending aorta was normal in size. At 4 months of age repair of the aortic coarctation and banding of the pulmonary artery were undertaken. A left thoracotomy was made and the coarctation exposed. Because of the immediate proximity of the anomalous right subclavian artery to the coarctation we decided to use this vessel as a flap. The

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Fig. 1 Diagram showing the surgical technique. The descending thoracic aorta and anomalous right subclavian artery in the region of the coarctation are mobilised. The anomalous right subclavian artery is ligated as far distally as possible and an incision extended along the length of the right subclavian and across the coarctation. The flap is turned down and sutured.
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Fig. 2 (a) and (b): preoperative aortograms in a slight left anterior oblique view showing the right carotid, left carotid and left subclavian arteries filling sequentially followed by the anomalous right subclavian (arrow), which arises proximal to an aortic coarctation. (c) and (d): postoperative aortograms in the left anterior oblique view showing no filling of the right subclavian artery and repair of the coarctation.

Ligamentum arteriosum was ligated and divided. The aorta was displaced anteriorly and rotated to expose and allow dissection of the anomalous right subclavian artery, which passed medially and posteriorly to the oesophagus. The aorta was cross clamped and the right subclavian artery ligated distally. The subclavian artery was opened along its length and the incision extended across the coarctation into the area of poststenotic dilatation (Fig. 1). The flap of subclavian artery was turned down into the aortic incision and sutured in place with a continuous suture. The pulmonary artery was banded.

The postoperative course was uneventful, and the congestive cardiac failure was easily controlled. The child was recatheterised. There was satisfactory banding of the pulmonary artery, and a small gradient of 15 mm Hg across the site of coarctation. Fig. 2 shows the preoperative and postoperative appearances of the thoracic aorta.

Discussion

The normal right subclavian artery has three embryological components: the most proximal seg-
ischaemia—particularly of the spinal cord—are minimised during occlusion by the presence of collateral vessels. In patients with an aortic coarctation and an anomalous right subclavian artery undergoing surgical correction of the coarctation, cognisance should be taken of the role of the anomalous artery in providing collateral flow to the descending aorta. If the anomalous vessel lies proximal to the coarctation the length of aorta that may be cross-clamped may be limited because of the inadvisability of clamping both subclavian arteries that may be in close proximity. If the anomalous right subclavian artery arises distal to the coarctation it may act as the main collateral channel, and injudicious clamping should be avoided. In these patients there is an increased risk of postoperative complications such as paraplegia due to spinal cord ischaemia, and techniques such as atriofemoral bypass, ventriculofemoral bypass, jump grafts, or systemic hypothermia are advised to prevent this complication.

Aneurysms of the anomalous right subclavian artery in patients with coarctation have been reported, and the risk of creating a massive haemorrhage during dissection is increased. The anomalous subclavian artery tethers the aortic isthmus in a medial and posterior direction. This reduced mobility makes resection of the coarctation site and primary end to end anastomoses of the aorta more difficult. The subclavian flap method to correct coarctation of the aorta is becoming increasingly more common, as it is technically relatively easy to perform, and the morbidity and mortality is no higher than with other techniques and it has the potential of growth. This report shows that the anomalous right subclavian artery can be used as the flap, and in this situation it is probably the method of choice as in some patients the anomalous vessel may cause obstructive symptoms to the oesophagus and trachea and will require division. Using the anomalous vessel as the flap deals with both problems. The flap may be used when the anomalous vessel is proximal to the coarctation and when distal, provided that the vessel (subclavian artery) is of adequate calibre. If the anomalous vessel is distal to the coarctation due care should be taken to prevent spinal cord ischaemia. The only theoretical disadvantage of using the anomalous right subclavian artery as a flap is the possible development of a subclavian steal as the vertebral artery is not ligated. This complication is probably uncommon since it is rarely reported after division of the vessel for obstructive symptoms to the trachea or oesophagus.

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