Aneurysm of the ascending aorta presenting with pulmonary stenosis

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The symptoms and signs of aneurysms of the thoracic aorta are due mainly to compression of surrounding mediastinal structures and depend on the size and location of the aneurysm. Compression of the trachea, bronchi, oesophagus, recurrent laryngeal nerve, bone, and superior vena cava is common, but pressure on the pulmonary artery has not been described in spite of its intimate relation to the ascending aorta.

The object of this paper is to describe a case presenting in this way and to discuss the particular surgical problems involved in the excision of such an aneurysm.

CASE REPORT

E. D., a man aged 52 years, was admitted to the Brompton Hospital in December 1963. He had a history of healed bilateral pulmonary tuberculosis, and he suffered from recurrent attacks of haemoptysis and bronchial infection. In January 1957 he had been admitted to hospital in acute cardiac failure due to coronary thrombosis. Since then he had been moderately dyspnoeic.

On examination he was a heavily built man. The pulse was in sinus rhythm, the blood pressure 120/80 mm. Hg. and the jugular venous pressure was normal. An abnormal systolic impulse was palpable in the second left intercostal space at the left sternal edge. There was a moderately loud, harsh, inspiratory pulmonary ejection murmur, and the pulmonary component of the second sound was delayed and reduced in intensity but moved on respiration. Bilateral basal crepitations were heard in the lungs, and the liver was palpable one finger’s breadth below the costal margin.

The electrocardiogram showed sinus rhythm with a normal PR interval. The mean frontal QRS vector was –40°, and there was right bundle-branch block with a terminal R' deflection in V1 of 13 mm. These changes were interpreted as indicating ischaemic heart disease.

The chest radiograph (Fig. 1) showed old bilateral apical tuberculosis, and both lungs were emphysematous. The emphysema was most marked in the left upper lobe with bullae present. A hemispherical mass, 9 by 6 cm., lay to the left and in front of the ascending aorta. Its border was slightly irregular, and showed flecks of linear calcification laterally and superiorly. The heart size was within normal limits.

At right heart catheterization difficulty was experienced in passing the catheter into the pulmonary artery. The right ventricular systolic pressure was 44 mm. Hg and the pulmonary artery systolic pressure was 18 mm. Hg. the systolic gradient being 26 mm. Hg., with a single change in pressure from arterial to ventricular configuration in the vicinity of the pulmonary valve. The angiogram (Fig. 2) showed compression and distortion of the right ventricular outflow tract, pulmonary valve, and main pulmonary artery by a mass lying above and in front of them. The aortic valve and lower part of the ascending aorta were normal. A saccular aneurysm, measuring 8 by 5·5 cm., arose from the upper part of the ascending aorta and contained clot. The neck of the sac was 2·5 cm. in diameter. The Wassermann, Price’s precipitin reaction, Treponema pallidum immobilization, Reiter protein complement fixation test, cardiolipin Wassermann, and fluorescent treponemal antibody tests were all negative.

Repair of the aneurysm was performed on 14 May 1964 with the aid of profound hypothermia by the Drew technique, using a bilateral transverse incision along the lower border of the third rib dividing the sternum. The circulation was arrested for 35 minutes at a temperature of 12° C. The aneurysmal sac was opened and the neck was seen to be 2·5 cm. in diameter, with strong fibrous margins. Direct suture of the defect in the aortic wall was performed with interrupted mattress sutures and reinforced with a continuous running stitch and a pericardial flap. Two thirds of the sac was removed and the remainder was obliterated by sutures. During rewarming of the patient there was no bleeding from the suture line.

The emphysematous bullae in the left upper lobe were obliterated by multiple catgut ligatures, a technique that avoided the air leak which is inevitable with excision and suture. The chest was closed and tracheostomy was performed.
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Artificial ventilation was required for three weeks, following which the patient made an uneventful recovery. The pulmonary ejection murmur disappeared immediately after operation.

When seen in July 1964 he was well, though still somewhat dyspnoeic. On examination the cardiac physical signs were normal except for a blood pressure of 145/100 mm Hg. The electrocardiogram was essentially unchanged from that before operation. His chest radiograph showed a normal aortic outline and diminution of the transverse cardiac diameter (Fig. 3).

DISCUSSION

Aneurysms of the thoracic aorta may present with signs of compression of the surrounding structures, trachea, bronchi, oesophagus, nerves, bone, and superior vena cava (Boyd, 1924; Brindley and Stembridge, 1956; Blakemore and Voorhees, 1954; Mills and Horton, 1938; Kampmeier, 1938). Compression of the pulmonary artery has not been previously described.

Systolic murmurs are frequently observed in patients with aneurysms of the thoracic aorta. Joyce, Fairbairn, Kincaid, and Juergens (1964)
reported that 50% of a group of 170 patients with aneurysms of the thoracic aorta had murmurs, but concluded that it was difficult to relate them to the aneurysm and that in most patients the murmurs 'seemed to be non-specific'. Mills and Horton (1938) found that 169 out of 339 patients with aneurysm of the thoracic aorta had systolic murmurs which were best heard in the aortic area in 71, at the apex in 52, and in the pulmonary area in 30 patients. In nine patients the systolic murmur was present in all areas and in seven the site of the murmur was not specified.

In the case reported here the systolic ejection murmur was unquestionably pulmonary in origin, as it was louder on inspiration, not conducted to the neck, and was associated with a delayed pulmonary component of the second sound. There was a systolic gradient of 24 mm. Hg between the right ventricle and the pulmonary artery, and compression of the outflow tract of the right ventricle and main pulmonary artery was confirmed by angiography and at operation. The clinical signs of pulmonary outflow compression were relieved completely by resection of the aneurysm. This suggests that some of the systolic murmurs described in association with aneurysms of the ascending aorta may be due to pulmonary outflow compression.

Obstruction of the right ventricular outflow may therefore be an additional factor in the production of the cardiac enlargement and failure that is common in cases of aneurysm, although these are usually due to associated syphilitic aortic regurgitation, ischaemic heart disease, or systemic hypertension (Brindley and Schwab, 1930; Mills and Horton, 1938; Brindley and Stembridge, 1956). Relief of pulmonary outflow compression in this patient resulted in diminution of the heart size.

Saccular aneurysms of the ascending aorta compressing the pulmonary outflow present special surgical problems. When the aneurysm has a narrow neck the use of cardiopulmonary bypass has disadvantages. Mobilization and occlusion of the aorta proximal to the aneurysm to allow the coronary arteries to be perfused by the beating heart is hazardous due to adherence of the aneurysmal sac to the right ventricle and the right coronary and pulmonary arteries. Clamping the aorta distal to the aneurysm necessitates coronary arterial cannulation for myocardial perfusion. This is difficult to do through the narrow neck of the aneurysm and may involve incising normal aortic wall above the aortic ring, which is often obscured by the aneurysm.

These problems do not arise with the use of profound hypothermia by the Drew technique (Drew and Anderson, 1959). With complete circulatory arrest at 12°C, proximal control and coronary perfusion are unnecessary. The aneurysmal sac is excised, the neck closed from inside, and the redundant sac removed, leaving the part adherent to the pulmonary and right coronary arteries.

The main hazard of profound hypothermia in the surgery of thoracic aneurysms is the absence
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REFERENCES


