Aneurysm of pulmonary artery with persistent ductus arteriosus and pulmonary infundibular stenosis

Fatal dissection and rupture in pregnancy


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A case of aneurysm of the pulmonary artery in a young pregnant woman is described. In addition, she had a persistent ductus arteriosus and congenital right ventricular infundibular muscular stenosis. At necropsy the wall of the pulmonary artery showed lesions of medionecrosis. Death occurred in relation to evacuation of the bowels. Presumably the increased intrathoracic pressure caused the final rupture of the aneurysm, which had probably already dissected extensively.

Aneurysms of the pulmonary artery are rare and may affect the main pulmonary artery or one of the major branches. An association with persistent ductus arteriosus has been noted in 20 per cent of cases (Deterling and Clagett, 1947). Occasionally ligation of a persistent ductus arteriosus may be complicated by the formation of a post-operative aneurysm (Thomas, 1960). Pulmonary aneurysm may also follow infective endocarditis of a persistent ductus arteriosus (Lillian, 1949). Occasionally Marfan’s syndrome (Tung and Liebow, 1952; Best, 1967) and long-standing pulmonary hypertension such as that accompanying a large ventricular septal defect (Johannsen and Connor, 1943; Richards and Cohn, 1954) may result in aneurysm of the pulmonary artery.

Aneurysms of the pulmonary artery are liable to rupture into the lung, resulting in fatal haemoptysis, or into the pericardium, with tamponade and sudden death. The diagnosis may be obvious on a straight chest film if considerable dilatation of the pulmonary artery is seen. If not, it may be proved by angiocardiography. Because of the rarity of aneurysm of the pulmonary artery, and because of the association of persistent ductus arteriosus and pulmonary infundibular stenosis in the present instance, we make this case report.

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2 Requests for reprints to K. S.

Case report

An 18-year-old Ugandan housewife, 28 weeks pregnant, was admitted on 2 May 1968. She had been under observation for one month at a district hospital, with pulmonary oedema and clinical features of severe pulmonary hypertension. She gave a history of having been somewhat dyspnoeic and also having suffered vague chest pain since childhood.

Examination revealed a well-built young pregnant woman who appeared restless as if from labour pains. She was aphyreal, not anaemic or cyanosed, and had no finger clubbing. The pulse was 85 a minute, regular, not collapsing, the blood pressure being 120/80 mm. Hg. The apex beat was located in the fifth left intercostal space in the anterior axillary line: the cardiac impulse was of left ventricular type together with a left parasternal heave. Auscultation revealed a long, grade 4/6, harsh systolic murmur, of ejection type, maximal over the left sternal edge in the third and fourth intercostal space, and associated with a systolic thrill; a high-pitched grade 2/6 early diastolic murmur was also present. The systolic murmur was also heard over the pulmonary area though less loudly, while the pulmonary component of the second sound was soft. There was no ejection click. At the apex there was a grade 2/6 pansystolic murmur radiating to the axilla. This was accompanied by a soft grade 2/6 mid-diastolic rumble and a third heart sound. No abnormality was observed at the aortic area. Coarse crepitations were heard in the lungs. The liver and spleen were not palpable. The fundi were normal. The chest x-ray (Fig. 1) showed a large heart with a cardio-
The necropsy

The pericardial sac was tense and contained 1000 ml. blood. The aorta, pulmonary artery, and large veins were normally positioned. There was a large fusiform aneurysm of the pulmonary trunk (Fig. 2), with a small oval rupture about 8 mm. in length communicating with the pericardial sac. On the intimal side of the aneurysm there were several semicircular areas of dissection. The aorta was connected to the main trunk of the pulmonary artery, just before its division, by a persistent ductus arteriosus, which was only a few millimetres in length and measured 1 cm. in diameter. There were some intimal plaques in the aorta around the opening of the ductus but no evidence of widespread atheroma. The intrapulmonary branches of the pulmonary artery showed plaques of atheroma, though none was present in the main pulmonary trunk.

The heart weighed 500 g., and the pericardium was covered with a thin fibrinous exudate. Both atria were dilated and trabeculated. The right ventricle wall thickness averaged 1 cm., 1.5 cm., and 1.7 cm. at the apex, middle, and infundibular parts, respectively, while the left measured 1.5 cm. in the middle. There was no ventricular dilatation. There was a muscular type of stenosis located in the infundibulum of the right ventricle. The tricuspid valve was thickened at its free edge, partly calcified and verrucated, but was neither stenosed nor incompetent. The mitral, pulmonary, and aortic valves were normal in all respects.

The lungs were heavy, firm, and congested. The uterus was bulky, its wall measuring 2 cm. All other organs were macroscopically normal.
Microscopical examination  The pulmonary artery and, to a lesser extent, the aorta, showed pools of mucopolysaccharide in the media, which stained strongly with alcin blue and Mayer's mucicarmine. The changes seen were diffuse in the pulmonary artery and resembled the lesions of aortic medionecrosis. The fragmentation of smooth muscle and elastic fibres was conspicuous in the pulmonary artery but minimal in the aorta. In some sections there was extravasation of blood in the walls of pulmonary artery. Sections of the pulmonary artery showed conspicuous changes due to accumulation of mucopolysaccharide with destruction of the elastica and smooth muscle. Both right and left ventricles showed hypertrophy of the myofibres. Aschoff's lesions were not seen in the ventricle or atria, and there was no evidence of myocarditis. The lungs showed thickened alveolar walls, medial hypertrophy of the pulmonary arterioles, and numerous macrophages laden with haemosiderin.

The cause of death was cardiac tamponade which had resulted from the dissection and rupture of the pulmonary aneurysm.

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References


