Case reports

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Biloculate false aneurysm of the right ventricle after cardiac surgery

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A case of a 12-year-old boy who had double false aneurysms of the right ventricle after incomplete closed pulmonary valvotomy six years earlier is presented. The aneurysms were successfully treated surgically, and the aetiology is discussed.

False ventricular aneurysms are a rare and potentially fatal complication of cardiac surgery. They are less common than true aneurysms (Hurst, Fine, and Keyes, 1963; Littler, Meade, and Hamilton, 1971; Rosenthal, Gross, and Pasternac, 1972; Van Tassel and Edwards, 1972) but are more likely to rupture (Hunter and Benson, 1933; Gobel, Visudh-Arom, and Edwards, 1971; Van Tassel and Edwards, 1972; Rosenthal et al., 1972). Both types of aneurysms are more frequent sequelae of operations on the left ventricle (Littler et al., 1971).

This report describes what we believe to be a unique case of a patient who, after a closed pulmonary valvotomy, developed a false right ventricular aneurysm. This aneurysm, in turn, had ruptured into the left pleural cavity, resulting in a second false chamber. The patient survived this event and surgical correction of the lesion followed successfully.

Case history

A 12-year-old West Indian negro boy was born normally at full term. His mother had had rubella in the first month of pregnancy, and a heart murmur was discovered when he was 3 months old. He failed to thrive and was prone to upper respiratory tract infection. In 1967, at the age of 5 years, the diagnosis of pulmonary stenosis was made, and a transventricular pulmonary valvotomy was performed through a left thoracotomy. Apart from transient early postoperative pulmonary collapse and infection he maintained a gradual progressive improvement. Three years later he experienced slightly reduced exercise tolerance. In May 1973 a routine chest radiograph disclosed a widened mediastinum and a large round opacity in the upper left lung field. A right ventricular aneurysm was suspected. Two months later he was referred to the Hammersmith Hospital. His only symptom was a slightly reduced exercise tolerance.

On examination he was a thin, fit young boy. His arterial pressure was 110/80 mmHg (14.6/10.6 kPa), his pulse rate 100 per minute and regular; the jugular venous pulse was 2 cm above the sternal edge of the sternoclavicular joint, with a prominent 'a' wave. There were distinct visible pulsations at the second and third left intercostal spaces. A grade 4/6 ejection systolic murmur was heard at the pulmonary area, with grade 3/4 pulmonary diastolic murmur. There were no signs of heart failure, and the previous thoracotomy was well healed, with a keloid scar. Routine laboratory investigations were normal. The chest radiograph showed a large right ventricle and slightly oligo-aemic lung fields. A large rounded shadow was superimposed on the main pulmonary artery and extended, just above it, into the upper left lung field. The lateral view showed anteroposterior extension of the mass (Fig. 1).

The electrocardiogram showed right axis deviation and right ventricular hypertrophy.

Cardiac catheterization disclosed a residual gradient of 60 mmHg (8.0 kPa) across the pulmonary valve: MPA pressure 30/13 mmHg (4.0/1.7 kPa), RV body 90/21 mmHg (12/0.28 kPa), pressure within aneurysm 89/15 mmHg (11.8/2.0 kPa).

The cineangiogram of the right ventricle showed a centrally placed mediastinal aneurysm above and anterior to the heart, which filled by a jet from the right ventricular infundibulum. This mediastinal sac, in turn, communicated with and filled a posterolaterally placed sac in the left pleura.
The main pulmonary artery was displaced backwards and was compressed by the mediastinal aneurysm. No communication could be demonstrated between the pulmonary trunk with its two branches and either of the two aneurysmal sacs. Injection of contrast medium into the superior vena cava disclosed the close proximity of the juxta-atrial part of the superior vena cava and of the right atrial appendage to the mediastinal aneurysm. The appearance was that of a fairly large right ventricular aneurysm communicating with a second aneurysmal chamber in the left pleura.

Operation (H.H.B.)
In view of the angiographic findings and the intimate relations of the superior vena cava and the right atrial appendage to the central aneurysm sac, the risks involved in using a median sternotomy approach were appreciable. The right femoral artery, therefore, was cannulated, and a right anterolateral thoracotomy was performed through the fourth intercostal space. This approach provided trouble-free access to cannulate both cavae. Total cardiopulmonary bypass was established, the mediastinal aneurysm was freed from the back of the sternum, and the sternum was then divided transversely. The aneurysm was a false sac formed by the pericardium. It was opened anteriorly and found to communicate with the right ventricular infundibulum through a 1.5 cm elliptical hole, presumably the site of the previous ventriculotomy. Another 1.5 cm hole, laterally placed in the wall of the central aneurysm, led to the second false aneurysm formed by the adherent left pleura (Fig. 2). Neither cavity contained any thrombus.

The pulmonary trunk was incised longitudinally through the posterior wall of the mediastinal aneurysm to relieve the residual obstruction. The pulmonary valve was thickened, the commissures were fused, and the cusps were rolled in. It was evident that the pul-
monary ring down to the ventriculotomy. Probably have a variety of causes. Part of the tricular gusset was inserted to enlarge partially excised and drained. The lateral aneurysmal valve was drained via its communication with water-seal-suction system. The patient came off bypass late to the development of ventricular aneurysm.


Hunter, W. C., and Benson, R. L. (1933). Rare form of saccular cardiac aneurysm with spontaneous rupture. American Journal of Pathology, 9, 593.


References


Conclusion

The majority of the recorded false aneurysms of the heart are related to myocardial infarction or surgery on the ventricles (Hurst et al., 1963; Littler et al., 1971; Fadali et al., 1974). False aneurysms of the right ventricle are less common and follow surgery or trauma. Usually a false aneurysm follows a leak from the right ventricular outflow patch in the surgical correction of Fallot's tetralogy (Payne and Kirklin, 1961; Bahnson et al., 1962; Wada et al., 1965), but, whether the ventriculotomy itself or the patch dehisces, the main aetiological factor is the persistence of residual outflow obstruction producing raised right ventricular pressure. This was certainly a factor in our case.

If the patient is to survive the right ventricular dehiscence, there must be coexisting pericardial adhesions to prevent fatal tamponade (Hurst et al., 1963; Stansel, Julian, and Dye, 1963; Fadali et al., 1974). A unique feature of this case is the further rupture of this false aneurysm into the left pleural space, which must have been limited by adhesions, a result of the previous thoracotomy – hence the development of the second false chamber. There is no way of knowing whether this second rupture occurred simultaneously with the ventricular dehiscence.

This patient had a regurgitant pulmonary valve and a chest infection after the initial valvotomy. Both these conditions have been suggested by Hipona and Bloom (1965) as being possible factors in aneurysm formation. Other factors, in childhood, such as residual left-to-right shunts and complete heart block (Payne and Kirklin, 1961), congenital defects of the myocardium, traumatic myocardial necrosis, and Chagás's disease have also been blamed (Bahnson et al., 1962). On the whole


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