Acquired right ventricular outflow tract obstruction

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It is not unusual for complicated aneurysms of the ascending aorta to affect left ventricular function. Processes causing or associated with an aneurysm may narrow the coronary artery ostia and produce myocardial ischemia or infarction; the aneurysm may distort the aortic valve ring and produce aortic insufficiency with subsequent left ventricular hypertrophy, dilatation, and failure. However, an unusual manner in which an aortic aneurysm can lead to heart failure is by right ventricular outflow tract obstruction. Here, the clinical picture often resembles that of pulmonic stenosis.

This report describes such a complication of an aortic aneurysm and focuses attention on the variety of mechanisms by which acquired right ventricular outflow tract obstruction may be produced.

Case report

A. D. (UMMC 084202), a 57-year-old Caucasian housewife, was admitted Sept. 14, 1964, because of chronic back and chest pain. She had sustained fractures of the left clavicle and the left lower extremity in an automobile accident 12 years earlier and since then had noticed an intermittent left subscapular ache which did not limit activity. In 1960, she was refused employment because of a chest x-ray abnormality. Five months prior to admission the pain increased in frequency and severity. It radiated to an area beneath the left breast and varied with position and respiration. It did not resemble typical angina pectoris. She also experienced dyspnea on mild exertion and three episodes of transient lightheadedness. She had not had syncope or hemoptysis. There was no history of rheumatic fever, venereal disease, or heart murmur.

The blood pressure was 130/80 mm. Hg and the pulse rate 86 with a regular rhythm. The trachea was in the midline and the neck veins were not distended in the supine position. The carotid artery pulses were normal. The lungs were clear. The heart was not enlarged. A prominent left parasternal lift was present, but there were no thrills. The second sound split physiologically. A Grade 3/6 harsh systolic ejection murmur was present over the entire precordium but was loudest in the third intercostal space at the left sternal border. It radiated to the neck, left axilla, and the lung bases. There were no diastolic murmurs. The peripheral pulses were normal and equal bilaterally. Cyanosis, clubbing, and edema were absent.

The hematocrit was 37 per cent and the white cell count was 4,700 with a normal differential. The serology was negative. A fasting blood sugar was 81 mg. per cent, the urea nitrogen 18 mg. per cent, and cholesterol 189 mg. per cent. The total serum protein was 7.8 mg. per cent, with an albumin-globulin ratio of 1.2.

X-ray examination (Fig. 1) revealed a 10 cm. calcified thoracic aneurysm which had increased 1 cm. in size since a chest film taken six months earlier. Electrocardiograms (Fig. 2) showed incomplete right bundle branch block, the S1Q3T3 pattern, and symmetrical T-wave inversion in Leads II, III, F, and V1-2. An intravenous angiogram was done with serial biplane filming. The right atrium was normal. The right ventricle was moderately enlarged. The pulmonic valve and the common pulmo-

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The left pulmonary artery did not visualize and the right pulmonary artery was compressed at its origin (Fig. 3). The caliber of the vessels on the right beyond the partial obstruction appeared increased. The pulmonary veins, left atrium, left ventricle, and ascending aorta appeared normal. There was a large saccular aortic aneurysm of the arch distal to the left subclavian artery which was almost completely filled with laminated clot. The remainder of the aorta was normal. A right heart catheterization was performed (Table I) and showed right ventricular and common pulmonary artery hypertension with a 60 mm. Hg pressure gradient at the bifurcation of the common pulmonary artery. Cineangiograms demonstrated slowing of the circulation through the right side of the heart. The common pulmonary artery appeared “kinked.” The remainder of the study confirmed the findings of aortography.

On the seventeenth hospital day a left thoracotomy was performed. The aneurysm, measuring 10 cm. at its greatest diameter and 4 cm. at its base, was dissected free of the lung surface. An attempt was made to isolate the aorta above and below the aneurysm. However, these manipulations further compromised the already partially obstructed pulmonary outflow tract and resulted in a decrease in systemic arterial pressure. It became necessary to use cardiopulmonary bypass during which diffuse bleeding began. When clot was evacuated from the aneurysm, the left pulmonary artery began to pulsate vigorously. The aneurysm was excised from the anterolateral aspect of the aorta and replaced by a Teflon graft. However, despite all measures, the diffuse bleeding became more severe and the patient died in the operating room.

**Discussion**

The pulmonary artery arises from the infundibulum of the right ventricle behind the sternal end of the second left intercostal space. It is about 5 cm. in length and 3 cm. in diameter in the adult and is entirely enclosed within the fibrous pericardium. It passes upward and backward while curving through the mediastinum from a position in front of the ascending aorta to a position left of the aorta. Anywhere along this path a vascular, neoplastic, or inflammatory lesion can compromise the outflow tract, masquerade as pulmonic stenosis, and produce right ventricular failure.

The early clinical finding of acquired right ventricular outflow tract obstruction is a systolic ejection murmur accompanied by a thrill over the upper left sternal border. The murmur may be continuous (machinery-like) and resemble that of patent ductus arteriosus. Such a murmur in the presence of a thoracic aortic aneurysm is suggestive of acquired right ventricular outflow tract obstruction. The electrocardiogram is helpful if it shows increasing right axis deviation and other signs compatible with right ventricular hypertrophy. Chest pain and dyspnea are common presenting symptoms.

The most common lesion producing acquired right ventricular outflow tract obstruction is an aortic aneurysm, particularly a syphilitic aneurysm. The concave surface of the ascending aorta is a common site for the aneurysm which may enlarge anteriorly and to the left and deform the lumen of the pulmonary artery. In 1839, Hope described a patient with an aortic aneurysm which ruptured into the pulmonary artery. Peacock in 1868 and Kap-
Fig. 2. Electrocardiogram demonstrating findings which may be the result of right ventricular hypertension (incomplete right bundle branch block, $S_3Q_3T_3$ pattern, and symmetrical T-wave inversion in Leads II, III, F, and V1-a).

Fig. 3. Intravenous angiogram revealing a large saccular aortic aneurysm of the arch distal to the left subclavian artery totally obstructing the left pulmonary artery and compressing the origin of the right pulmonary artery.

In 1907 each reviewed the literature and reported 19 and 32 similar cases, respectively. The signs and symptoms of the rupture of an aortic aneurysm into the right ventricular outflow tract were summarized by Porter and Nicholson and each of these authors presented the clinical picture as a syndrome associated with syphilitic aortitis. Brill and Jones cited 87 patients of whom 84 (96.5 per cent) came to medical attention at the time of rupture of a syphilitic aneurysm into the pulmonary artery. Only 3 (3.5 per cent) of this group had symptoms due only to compression of the right ventricular outflow tract. A non-syphilitic aortic aneurysm presenting in the latter fashion is an even rarer occurrence. Yacoub and associates presented one such case and another is reported in this paper.

Neoplasms rank second in frequency among the lesions producing right ventricular outflow tract obstruction. Primary intracardiac tumors which have been observed to compromise the right ventricular outflow tract included rhabdomyosarcoma, fibrosarcoma, chondrosarcoma, reticulum cell sarcoma, myxoma, pleomorphic sarcoma, and mesothelioma of the pericardium. The only reported metastatic intracardiac lesion which has produced this type of obstruction is a malignant argentafinoma. Extracardiac tumors which have mechanically obstructed the pulmonary artery include teratoma, Hodgkin's disease, thymoma, and bronchial carcinoma.

An annular constricting pericardial band is the most frequently encountered inflammatory lesion producing acquired obstruction. Gouley described five instances of rheumatic pericarditis which resulted in
supravalvular pulmonic stenosis. Mounsey and McGaff and associates have reported patients with subvalvular obstruction due to a pericardial band. Barros and Gomez described a patient with a pericardial band constricting the right ventricular outflow tract and noted that a partial pericardiectomy had antedated the acquired obstruction in his patient and in those mentioned above. Seymour and coworkers discussed a similar patient with acquired obstruction in his patient and in partial pericardiectomy had antedated the tricuspid outflow tract and noted that a cardiac band constricting the right ventricular outflow tract due to a pericardial band. Barros and Gomez described a patient with a pericardial band without previous pericarditis. Schrire, Beck, and Barnard, C. N.: Aneurysm of the ascending aorta obstructing right ventricular outflow and producing severe "pulmonary stenosis," Amer. Heart J. 23:396, 1963.

Although acquired right ventricular outflow tract obstruction is an unusual cause of right ventricular failure, it is not such a rare postmortem finding that the clinician should not keep the possibility high on the list of conditions to be considered when he is faced with the inexplicable and gradual onset of right ventricular failure—particularly when this is associated with a "new" ejection murmur along the upper left sternal border.

REFERENCES

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