Idiopathic calcified myocardial mass

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PATTERSON, D., GIBSON, D., GOMES, R., MCDONALD, L., OLSEN, E., PARKER, J., and ROSS, D. (1974). Thorax, 29, 589–594. **Idiopathic calcified myocardial mass.** Myocardial calcification can be subdivided into three groups—metastatic, dystrophic or an extension inwards from the pericardium. This case in which the calcified myocardial mass was initially delineated by radiography and by echocardiography and subsequently removed does not fit into any subdivision and has been termed idiopathic.

Myocardial calcification has been subdivided by Finestone and Geschickter (1949) and Gore and Arons (1949) into three groups—metastatic, dystrophic or an extension inwards from the pericardium. In the metastatic group hypercalcaemia is usually present and calcium salts are deposited in tissues which were previously normal. The dystrophic calcification is related to the laying down of calcium salts in dead, degenerated or devitalized myocardial tissue.

This case in which the calcified mass was successfully removed does not satisfactorily fit into any subdivision and it has therefore been termed idiopathic.

**CASE REPORT**

Miss M.D. aged 52 first developed symptoms at the age of 42 when she was suddenly awakened by breathlessness and palpitations. She subsequently felt tired and lacked energy; there was no associated chest discomfort and no expectoration of sputum. She was found to be in atrial fibrillation. Her symptoms improved spontaneously although she remained tired. At age 48 she suddenly developed a right lower nasal visual field defect that has persisted. No other neurological deficit was noted. At age 50 she noticed increasing shortness of breath which improved with diuretic therapy but one year later she became increasingly short of breath on exertion and orthopnoeic, and developed paroxysmal nocturnal dyspnoea. Her ankles and abdomen swelled and she again noted palpitations. No chest discomfort was associated with these symptoms. Digoxin, furosemide, and anticoagulants were started with some improvement in her symptoms.

Four months later she developed a saddle embolus while still on anticoagulant treatment; this necessitated bilateral femoral thrombectomy.

Her past medical history revealed that diphtheria was questioned at the age of 23 but never substantiated. She had no history of rheumatic fever.

Her father died aged 72 of a myocardial infarction and her mother died aged 72 of a cerebrovascular accident. One brother aged 50 had had a myocardial infarction; five sisters are alive and well.

She gave up smoking cigarettes at the age of 51. Her menopause occurred at age 49. She had never been abroad.

Physical examination revealed her to be in no acute distress. She was normotensive. There was no arcus and no xanthomata. There was no evidence of heart failure. There were no pulses palpable below the femoral arteries but there were no trophic skin changes. The apex beat was not displaced. The heart sounds and murmurs suggested mild aortic regurgitation and mild mitral valve disease. The remainder of the physical examination was normal.

**INVESTIGATIONS** An electrocardiogram (Fig. 1) showed atrial fibrillation with a controlled ventricular rate. The QRS axis and QRS duration (0-10 sec) were normal. Voltage criteria of left ventricular hypertrophy were present with ST segment and T wave abnormalities. There was no evidence of old myocardial infarction.

A chest radiograph (Fig. 2) showed enlargement of the left atrium and some pulmonary venous congestion. There was dense intracardiac calcification situated in the interventricular septum and remote from the mitral valve ring. There was no pericardial calcification.

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CARDIAC CATHETERIZATION There were normal right heart pressures with a mean pulmonary arterial wedge pressure of 16 mmHg rising to 25 mmHg after exercise. The end-diastolic indirect mitral valve gradient was 0 mmHg at rest rising to 10 mmHg after exercise. No systolic aortic valve gradient was present. Left ventricular end-diastolic pressure was 15 mmHg rising to 32 mmHg after angiography.

Angiography The left ventriculogram revealed that contractility of the left ventricle was severely impaired by the calcified mass, and numerous lobulated calcified excrescences encroached into the left ventricular cavity. The intracardiac calcification was confirmed to have its main concentration towards the apex but extended into the interventricular septum. Slight mitral regurgitation was present.

Selective coronary arteriography revealed normal coronary arteries.

The aortogram confirmed mild aortic regurgitation.

OPERATION A median sternotomy was performed in March 1972. The left atrium was enlarged and the intracardiac mass was readily palpable in the diaphragmatic and lateral walls of the ventricle. Opening the left atrium revealed a mitral orifice of about 1 cm² which was of rheumatic appearance. The mass in the ventricle was then approached through the diaphragmatic surface of the heart by an incision parallel to the posterior descending artery. The calcified mass was located 3–4 mm from the epicardium and had irregular projections extending into a healthy looking muscle. It was found to extend into the left ventricular cavity and up the diaphragmatic surface of the ventricle laterally, just across onto the septum about 2 cm below the aortic valve and about 2 cm from the mitral valve. It was enucleated by sharp dissection (Fig. 5). The mitral valve was replaced with a Starr–Edwards prosthesis and the ventriculotomy was closed with Teflon support.

The immediate postoperative course was uneventful and there was a marked symptomatic improvement. The postoperative electrocardiogram showed no increase in the QRS duration but demonstrated a slight intraventricular conduction disturbance. Chest radiographs showed no change in heart size and confirmed that virtually all the visible myocardial calcified mass had been removed. An echocardiogram (Fig. 6) showed that the intracavitary echoes were no longer present and that the left ventricular cavity remained large. Septal movement was abnormal, which is characteristic of a normally functioning mitral Starr–Edwards valve and reflects a reduced rate of early diastolic left ventricular filling.

Seven months after the operation the patient suddenly collapsed and died. No necropsy was performed.

PATHOLOGY The specimen submitted for examination consisted of three pieces of calcified tissue, measuring 1·6, 1·9, and 0·9 cm in the longest diameter respec-
FIG. 2. Chest radiographs PA (a) and lateral (b) showing an enlarged left atrium, pulmonary venous congestion, and dense intracardiac calcification.
FIG. 3. Preoperative echocardiogram showing an enlarged left ventricular cavity with normal movement of the posterior wall and interventricular septum. Abnormal echo demonstrated 1 cm behind septum in the left ventricular cavity.

FIG. 4. Preoperative echocardiogram of the mitral valve showing impaired diastolic mobility of the anterior cusp.

FIG. 5. Calcified mass removed at surgery.

FIG. 6. Postoperative echocardiogram showing the absence of the intracavitary echoes. Left ventricular cavity remained large, and abnormal septal movements were demonstrated.

The mitral valve cusp showed thickening predominantly by collagen tissue particularly at the end distal to the line of closure of the valve leaflets. Some recent fibrin was also present on the deformed atrial surface of the valve leaflets. Vascularity was slightly increased, compatible with a rheumatic aetiology.

A portion of the posterior mitral valve leaflet was available for serial sectioning. Every tenth section, cut at 5 microns thickness, was examined. There was no calcification in any of the sections examined.

tively. In two of these pieces myocardial tissue could be identified with the naked eye. There were numerous large areas of calcification embedded and surrounded by myocardial fibres which apart from mild hypertrophic changes showed no abnormality (Fig. 7). In addition to this, an occasional area of fibrous tissue rich in capillaries was present. Surrounding the calcific areas a scanty chronic inflammatory infiltrate, patchily distributed, was present.
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DISCUSSION

The clinical features in the present case were not helpful in establishing the diagnosis, since systemic emboli, increasing breathlessness, and fluid retention might all have been due to mitral valve disease with associated atrial fibrillation. Chest radiography was of value in demonstrating intramyocardial calcification although its precise delineation required left ventriculography. The presence of the mass within the left ventricular cavity was also demonstrable by echocardiography and shown to be distinct from the mitral valve, with movement parallel to the posterior wall rather than to the septum. Although echocardiography is now a standard method of diagnosing left atrial tumours causing interference with mitral valve function, this appears to be the first time that the method has been used to demonstrate an intraventricular tumour. Although calcification extended into the lower part of the septum, there does not seem to have been involvement of the conduction system, since the QRS duration was normal.

Mitral ring calcification with involvement of adjoining myocardium is well recognized in 'older' women (Pomerance, 1970). The primary lesion may be due in part to calcification of organized thrombus in the subvalvar angle of the mitral ring. This mechanism is unlikely in this patient as the calcification was remote from the mitral valve ring and there was no histological evidence of calcification in the available sections of the mitral valve ring which were removed at surgery.

It has been suggested by Ernstene and Hazard (1951) that deposition of calcium may occur during the healing phase of a severe toxic or septic myocarditis. The patient had rheumatic involvement of the mitral and aortic valves, and it can be postulated that a severe myocarditis was associated with her rheumatic fever. However, scanty inflammatory infiltrate is not uncommon around areas of calcification, and elsewhere the myocardium was normal; this suggests that a past myocarditis is unlikely. The size that the calcified mass attained also makes a 'dystrophic' mechanism unlikely.
The commonest cause of dystrophic calcification in this country is coronary heart disease. The patient's family history, smoking habit, and lipoprotein abnormality are 'risk factors' in the development of coronary heart disease. However, there was no history of myocardial infarction nor of angina pectoris and the coronary arteriograms were normal.

Rarer causes of myocardial calcification such as hypokalaemia, renal failure, hydatid disease and endomyocardial fibrosis can be excluded.

Metastatic calcification is also unlikely in the presence of normal serum and urinary calcium levels, the absence of any destructive lesion of bone, and the absence of abnormal calcification in other parts of the body.

Calcified primary cardiac tumours are extremely rare. Shapiro et al. (1963) could find only five in the literature and two of these were myxomas. Geha, Weidman, Soule, and McGoon (1967) describe two children in whom large calcified intramural ventricular fibromas were successfully removed. Although there is no histological evidence of a tumour in this case, a calcified tumour cannot be completely excluded as a possibility.

Calcified endocardial granulomata secondary to histoplasmosis were suggested by Dean, Pamukcoglu and Roberts (1969) to be the cause of some small calcified nodules found in the right ventricular cavity of a male patient. There was no evidence of histoplasmosis in this patient, however.

The cause of this calcified myocardial mass remains unknown. It does not fit into the three recognized subdivisions of myocardial calcification and is therefore termed 'idiopathic'.

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REFERENCES


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