Arterial microembolisation: an unusual presentation of dilated cardiomyopathy

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Abstract

Systemic embolisation is common in patients with dilated cardiomyopathy. Microembolisation as a presenting sign of dilated cardiomyopathy, however, has not been reported before. A 37 year old woman in whom dilated cardiomyopathy presented as arterial microembolisation to the toes is described.

Dilated cardiomyopathy is a clinical syndrome characterised by ventricular dilatation and associated systolic dysfunction in the absence of pericardial, valvar, or coronary artery disease.1 The clinical presentation, course, and prognosis of dilated cardiomyopathy are highly variable. In most studies of the clinical presentation of dilated cardiomyopathy, embolism was identified as a frequent and serious complication of this disease.2 It is estimated that at least 11% of patients with dilated cardiomyopathy have one or more embolic events during the course of their illness.3 The timing and severity of thromboembolism in patients with dilated cardiomyopathy are highly variable. In some instances embolic events are clinically silent and are detected only at necropsy as peripheral infarcts in patients dying of progressive congestive heart failure. More commonly, embolism occurs long after the onset of clinical evidence of congestive heart failure.4 In a few patients embolic events precede all other cardiac symptoms and the clinical effects of peripheral infarcts may overshadow the primary cardiac disease and confuse the diagnosis. Retrospective evaluation of these patients, however, has generally shown that radiographic cardiomegaly or electrocardiographic abnormalities were present at the time of embolisation.5 6

The systemic emboli described in patients with dilated cardiomyopathy are generally large and affect, in order of decreasing frequency, the lungs, the kidneys, the spleen, and the brain.6 7 8 9 10 The occurrence of arterial microemboli as a presenting feature of dilated cardiomyopathy has not been reported before. We describe a patient in whom arterial microembolisation was the presenting sign of dilated cardiomyopathy.

Case report

A 37 year old white woman was admitted to the hospital with severe pain and discoloration in both fifth toes. She had been in good health until a month before admission when she noticed a painless “blush discolouration” of the left fifth toe followed by similar changes in the right fifth toe. Two weeks before admission she had severe intermittent pain in both discoloured toes particularly at night. The pain was improved by walking and was partially relieved by analgesics. The pain was not exacerbated by exposure to cold. The patient did not report any symptoms of congestive heart failure including exertional dyspnoea, orthopnoea, paroxysmal nocturnal dyspnoea, and peripheral oedema. The patient had no history of arthritis or cerebrovascular disease. No history of rash, or symptoms attributable to a coagulation disorder, connective tissue disease, or vasculitis were elicited. There was nothing relevant in the patient's family history. The patient had consumed two ounces of vodka daily since the age of 18 and had smoked 40 cigarettes a day for 15 years.

Physical examination

Examination showed: blood pressure 110/68 mm Hg, pulse 80 beats/min, respiration 16 breaths/min, and temperature 36.5°C. Fundoscopic examination did not show retinal haemorrhages, exudates, or other vascular abnormalities. Heart sounds were normal but there was a soft systolic ejection murmur along the left sternal border. The abdominal examination was entirely normal. The right and left fifth toes were cool, cyanotic, and exquisitely tender. Peripheral pulses were within normal limits. Both affected toes showed increased sensitivity to pin prick but the neurological examination was otherwise normal.

Blood chemistry values were normal. Complete blood count was normal except for a raised mean corpuscular volume of 112 fl. Prothrombin and partial thromboplastin times were normal. The erythrocyte sedimentation rate was 11 mm/h (Westergren). Antinuclear antibody, rheumatoid factor, and cryoglobulins were not detected. Antithrombin 3, protein C, and complement concentrations were normal.

The electrocardiogram showed T wave inversion in the inferior and precordial leads. The chest x ray showed slight enlargement of the cardiac silhouette. An aortogram and arteriogram of the toes were performed from the femoral approach. This showed occlusion of the digital arteries supplying the left fifth and third toes (figure). There was partial revascularisation from a collateral circulation. There was no angiographic evidence of atherosclerosis or vasculitis in the aorta or more
Discussion

We report an unusual presentation of dilated cardiomyopathy in which systemic arterial microemboli preceded the symptoms and signs of congestive heart failure. Emboli to the lungs, kidneys, spleen, and brain are known to complicate the course of dilated cardiomyopathy and in some patients to preclude all other cardiac symptoms. Kyrle et al reported that peripheral embolisation complicates dilated cardiomyopathy. In this series, however, peripheral emboli occurred in patients with known cardiac dysfunction and were not the presenting sign that led to the diagnosis of dilated cardiomyopathy.

Ischaemia of the toes may be caused by peripheral vascular diseases including atherosclerosis, Buerger's disease, small vessel disease associated with diabetes mellitus, vasculitis, and microemboli. Angiography is helpful in identifying or ruling out these possibilities. In our case there were no angiographic changes of atherosclerosis, Buerger's disease, or vasculitis of large or medium sized arteries. Abrupt occlusion of the digital arteries was seen. This picture is compatible with embolisation, a vasculitis limited to the small arteries, or diabetic vascular disease. Vasculitis was believed to be an unlikely diagnosis because there were no constitutional symptoms to suggest a systemic inflammatory process. Nor did the laboratory data including a normal erythrocyte sedimentation rate, haemoglobin, and albumin, suggest an inflammatory vasculopathy. There were no other symptoms or signs to suggest a connective tissue disease and all serological testing was negative. The presence of diabetes mellitus was excluded by the absence of symptoms and retinal changes attributable to this disease and by the finding of a normal fasting blood glucose. By exclusion of other causes, embolisation was considered the most likely explanation for the digital artery occlusions.

The left ventricular mural thrombus seen on the cross sectional echocardiogram was the only identifiable source of the patient's microemboli.

The hypothesis that emboli originate from mural thrombi that form in the cardiac chambers as the result of blood stasis secondary to severely impaired systolic function is supported by the necropsy finding of cardiac thrombi in as many as 53% of patients with dilated cardiomyopathy. Other investigators have proposed that at least some embolic events in dilated cardiomyopathy are the manifestation of a generalised thrombotic tendency. This is suggested by a Japanese study of 36 patients with dilated cardiomyopathy. Four of these patients were young, suffered embolic events early in the course of their cardiac disease, which dominated the clinical picture, and died
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within a year of the onset of heart failure. Although a hypercoagulable state could not be conclusively ruled out in our patient, the normal platelet count and normal coagulation studies make the possibility of a generalised thrombotic tendency unlikely. Another possible cause of embolism in patients with dilated cardiomyopathy is arrhythmia. Our patient was in normal sinus rhythm and she had no electrocardiographic evidence of intermittent atrial fibrillation or complex ventricular arrhythmias.

Some investigators reported significant alterations in the activity of mitochondrial enzymes in the myocardial biopsy samples of patients with a history of excessive alcohol intake. On the basis of this finding it has been suggested that enzyme analysis of the biopsy samples may distinguish heart failure owing to alcohol from other causes of dilated cardiomyopathy. Biochemical analysis of the myocardial tissue was not performed in our patient. Three aspects of our patient's clinical picture, however, suggest that an unusual reaction to alcohol may have contributed to her cardiac dysfunction. These include a history of alcohol intake, the raised mean corpuscular volume, and the improvement of left ventricular function after she stopped drinking alcohol. Reversal of alcoholic cardiomyopathy has been previously reported by several investigators.

Another reversible cause of dilated cardiomyopathy is myocarditis. Myocardial inflammation was not seen on the patient's endomyocardial biopsy samples and she had no prodromes attributable to a viral infection. But because the diagnostic value of an endomyocardial biopsy specimen is limited by the possibility of sampling error, spontaneous resolution of myocarditis cannot be excluded in our patient.

Peripheral microemboli originating from a cardiac mural thrombus can be the presenting symptom of dilated cardiomyopathy. Thus this disease entity should be identified or ruled out in patients showing evidence of peripheral microembolisation even when there are no symptoms of congestive heart failure.