Systemic lupus erythematosus presenting with cardiac tamponade due to a haemorrhagic pericardial effusion

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Summary: A 14 year old girl presented with cardiac tamponade due to a haemorrhagic pericardial effusion. Systemic lupus erythematosus was diagnosed. Pericardial stripping was performed due to recurrence of the effusion despite pericardiocentesis and steroid therapy.

Introduction

Pericarditis is the commonest cardiac complication of systemic lupus erythematosus (SLE). However, pericardial effusion causing cardiac tamponade is rare in SLE and it is even rarer for it to be the presenting feature of SLE. We present a case of SLE presenting with cardiac tamponade due to a haemorrhagic pericardial effusion.

Case report

The patient was a 14 year old girl of Turkish origin. She had been apparently well until 2 months before admission when she had transient arthralgia of her hands, elbows and knees. She subsequently developed a 'flu-like illness' which was associated with a low grade fever and a persistent non-productive cough. Three days before admission she became more breathless and was referred to us for further investigation.

On examination she was of thin build with pale conjunctivae. Her temperature on admission was 38°C. There were palpable lymph nodes in the neck, right axilla and both inguinal regions. She was not dyspnoic at rest. The pulse was 120/minute, blood pressure 120/70 mm Hg, with no paradoxical pulse. Jugular venous pressure was elevated. Examination of her chest revealed signs of bilateral pleural effusions and an audible pericardial rub.

Other investigations at this stage revealed a haemoglobin concentration of 8.6 g/dl, white cell count 11.4 \times 10^9/l, ESR 64 mm/h. Blood urea, electrolytes and liver and thyroid function tests were normal. There was microscopic haematuria and proteinuria; blood cultures proved sterile.

A diagnostic pericardial aspirate revealed a haemorrhagic effusion with a protein content of 49 g/l, polymorphs and mononuclear cells. Staining for alcohol and acid-fast bacilli was negative and culture revealed no growth.

A diagnostic pleural aspiration revealed straw coloured fluid with a protein content of 21 g/l, and the presence of polymorphs and lymphocytes. Culture proved negative as was the stain for alcohol and acid-fast bacilli.

Over the next few days she became more breathless and developed orthopnoea. The pulse rate was 150/minute and the blood pressure 100/70 mm Hg with a paradoxical pulse of 20 mm Hg. Chest X-ray showed an enlarging cardiac shadow and an echocardiogram confirmed the presence of large pericardial effusions anteriorly and posteriorly. The pericardium was not thickened. A therapeutic pericardiocentesis was performed and 500 ml of blood-stained fluid was aspirated.

At this stage the following results became available: anti-nuclear antibodies – positive (greater than 1:40), DNA binding 71% (normal 1–10%). Complement C3, normal, C4 decreased at 9. Tests for the following antibodies were all negative: rheumatoid factor (sheep cell agglutination), antimitochondrial, smooth cell and gastric parietal cell. Thyroid antibodies – positive 1:400 for microsomal antibodies but negative for thyroglobulin. Viral screen – negative.

A diagnosis of systemic lupus erythematosus was made and treatment commenced with prednisolone 40 mg/day. Despite this her condition deteriorated with evidence of re-accumulation of the pericardial effusion. It was therefore decided to excise the pericardium. The anterior pericardium was excised and the fluid aspirated. Two pericardial drains and a pleural

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drain on each side were inserted. She made a good recovery post-operatively. A post-operative echocardiogram showed no re-accumulation of her pericardial effusion. Histology of the pericardium was that of an organizing fibrinous pericarditis with no evidence of granuloma formation. She made an uneventful recovery. A year later she remained asymptomatic from her SLE but had some cushingoid features.

Discussion

The most common cardiac manifestation of SLE is pericarditis. The incidence is reported as between 20–30%. However pericarditis is the presenting feature in only 1–4% of patients with SLE. Dubois reported no cases of tamponade in 159 cases of lupus pericarditis. Estes & Christian in a prospective study of 150 patients with SLE found 29 patients with pericarditis of whom two developed cardiac tamponade.

In SLE the pericardial effusion is exudative in nature. It is usually clear or may be serosanguinous but is very rarely haemorrhagic. On microscopy, polymorphonuclear leucocytes are most numerous and monocytes too may be present. Examination of the centrifuged specimen may reveal LE cells which are then diagnostic of SLE. The pericardial fluid may also demonstrate anti-nuclear and anti-DNA antibodies.

In the absence of cardiac tamponade, lupus pericarditis may be treated with non-steroidal anti-inflammatory drugs. However, corticosteroids are more effective in resolving symptoms and eliminating the pericardial effusion. Should cardiac tamponade occur this needs to be drained urgently.

Our patient with SLE developed cardiac tamponade and the pericardial fluid was haemorrhagic in nature, both of which are rare features of lupus pericarditis. Despite pericardiocentesis and steroid therapy there was re-accumulation of pericardial fluid resulting in cardiac tamponade which led to surgical excision of her pericardium. Pericardial stripping or excision is not normally required to prevent recurrence of pericardial effusions in SLE once steroid therapy is initiated.

References

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