Simultaneous presentation of upper lobe fibrobullossus disease and spinal pseudarthrosis in a patient with ankylosing spondylitis

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Abstract
A 51 year old man with a 20 year history of ankylosing spondylitis and pronounced thoracic gibbus presented with two simultaneous complications of longstanding ankylosing spondylitis, upper lobe fibrobullossus disease, and spinal pseudarthrosis. No neurological sequelae developed and treatment was conservative. Both these lesions mimic tuberculosis, and so it is important to determine them accurately to avoid unnecessary antituberculosis treatment. Both of these complications are reported to occur in longstanding ankylosing spondylitis and their simultaneous presentation may be more common than realized. This case is believed to be the first such report of their association.

There seems to be some discrepancy in the terms used to describe destructive vertebral lesions in ankylosing spondylitis. It is, however, recommended that the term 'spinal pseudarthrosis' is used to describe a destructive, localised lesion of vertebrae in ankylosing spondylitis, which is typically seen in the thoracolumbar region and leads to extensive bone resorption. The term 'spondylodiscitis' is confined to minor self-limiting eburnation of disc and end plates, which usually affects multiple levels simultaneously and occurs during the early inflammatory phase of the disease.

Upper lobe fibrobullossus disease is a well known manifestation of ankylosing spondylitis and mimics tuberculosis. About 100 cases had been reported up to 1977.

We describe a man with a 20 year history of ankylosing spondylitis who had simultaneous spinal pseudarthrosis and upper lobe fibrobullossus disease in both lungs. As far as we know, there has been no other published report of ankylosing spondylitis with the simultaneous presentation of these manifestations.

Case report
A 51 year old man who had had ankylosing spondylitis for 20 years was admitted to hospital in February 1987 because of apical shadowings in both his lungs. He had a history of hypertension and iritis. Except for breathlessness on effort he had no respiratory symptoms. Pains in the joints of the lower extremities, shoulders, and low- and mid-back were noted.

On admission he had muscle wasting, stiffened cervical, thoracic, and lumbar spine, and pronounced thoracic kyphosis. There was local tenderness over the thoracic gibbus. The movements of shoulder joints and right hip were restricted. No synovitis was found in peripheral joints. A slight systolic murmur was heard on cardiac auscultation and the breath sounds were faint. No neurological disturbances were found. Erythrocyte sedimentation rate was 77 mm/first hour, haemoglobin concentration was 127 g/l, leucocytes 12·2×10^9/l, thrombocytes 446×10^9/l and C reactive protein concentration 68 mg/l. A test for teichoic acid antibodies yielded negative results. A chest x ray picture showed apical shadowings of both lungs (fig 1) and changes in thoracic spine. A tomogram of thoracic spine showed a destructive process of D11 and D12 (fig 2). Computed tomography was used to assist the taking of a needle biopsy specimen from the destructive spinal area. The specimen showed necrosis without signs of inflammation. Antituberculosis treatment with a combination of four drugs was started. Culture of sputum specimens for tuberculosis, however, remained negative. After eight months of treatment the antituberculosis drugs were stopped when no changes in the roentgenological findings were found. It was concluded that the pulmonary spinal changes were associated with the patient's ankylosing spondylitis.

In May 1989 he had further back pain, but had not shown any signs of spinal cord compression during the two years of follow up. His right hip required total endoprothesis replacement.
Concomitant upper lobe fibroblulous disease and spinal pseudarthrosis in AS

Discussion
Based on the roentgenological picture, lung tuberculosis and tuberculous spondylitis were initially suspected in our patient. Indeed, the upper lobe fibroblulous disease of ankylosing spondylitis may simulate tuberculosis and the spinal process, infective spondylitis. In our case bacteriological and antibody evidence of an infective process were absent and no response to chemotherapy was found. Paravertebral soft tissue masses, a feature of infective spondylitis, were also absent.

Our patient was a typical case of spinal pseudarthrosis with longstanding ankylosing spondylitis and ankylosed spine with severe thoracic kyphosis. Chan et al. proposed that an initiating event for spinal pseudarthrosis is posterior element weakness of the spine, though the anterior element osteolysis may be the more prominent roentgenological feature. A posterior bone defect could also be seen in our patient. He had neither neurological sequelae nor intolerable pain, and conservative treatment was justified.

When patients with upper lobe fibroblulous disease also have longstanding ankylosing spondylitis its association with spinal pseudarthrosis (found in 15% of patients in a hospital series) may be fairly common.