Role of immune complexes in hypertrophic osteoarthropathy and nonmetastatic polyarthritis

MARK S. AWERBUCH AND PETER M. BROOKS
From the Department of Medicine, Repatriation General Hospital, Daw Park, South Australia 5041

SUMMARY Three case histories of patients with histologically proved malignant disease and an associated nonmetastatic symmetrical polyarthropathy are analysed. No evidence of a pathogenetic role for immune complexes was found in hypertrophic osteoarthropathy, whereas there is considerable evidence suggesting that the nonmetastatic polyarthropathy associated with malignant lymphoma is an immune complex phenomenon.

Hypertrophic osteoarthropathy (HOA) and nonmetastatic polyarthritis are relatively uncommon but well recognised syndromes seen in association with malignant disease. Despite the noninflammatory nature of the synovial fluid in the former,1-3 both may present clinically as an inflammatory arthropathy with a similar joint distribution. And where there exists no clinical or radiographic evidence of periosteal change and where clubbing is subtle or absent, these 2 arthropathies may be indistinguishable from one another or from other arthropathies.1 2 4 5 Their pathogenesis is unknown. We have examined 3 patients with histologically proved malignant disease and an associated non-metastatic symmetrical polyarthropathy in an attempt to determine the pathogenesis.

Material and methods

Circulating and intra-articular immune complexes were detected by the C1q binding assay.6 The results were expressed as micrograms of a preparation of heat aggregated human IgG (HAGG) per ml of serum. HAGG was prepared by heating Cohn fraction II gammaglobulin (20 mg/ml) at 63°C for 30 minutes and diluting in normal human serum.

Antinuclear antibody titres were performed by a standard indirect immunofluorescence technique.

Synovial fluid viscosity was judged by the length of ‘stringing’ of a falling drop of synovial fluid.7

Serum and intra-articular levels of C3 and C4 were measured by the automated immunoprecipitation method, Technicon.

Accepted for publication 14 November 1980. Correspondence to Dr M. S. Awerbuch, Repatriation General Hospital, Daw Park, South Australia 5041.
Role of immune complexes in hypertrophic osteoarthropathy and nonmetastatic polyarthritis

present. Three months later there was no obvious change in the arthropathy, but he complained of pain just proximal to both wrists. Radiographs were normal, but bone scintigraphy showed periosteal linear accumulation of isotope, characteristic of periosteal new bone formation.8

Case 2. The patient is a 66-year-old man who presented with a 10-week history of polyarthralgia and weight loss. On examination he was found to have severe clubbing of the fingers and toes, thickened and tender over the distal ends of the radius and ulna, and symmetrically warm and swollen ankles, knees, wrists, metacarpophalangeal joints and proximal interphalangeal joints of the fingers.

Investigations showed the following: Haemoglobin 12.6 g/dl, leucocytes 8·6 \times 10^9/l, ESR 76 mm/hour. Neither IgM rheumatoid factor nor ANF were detected. CICs in the blood were 500 \mu g/ml (normal less than 40 \mu g/ml), the serum C3 level was 110 mg/100 ml (normal range 55–120) and the serum C4 level 40 mg/100 ml (normal 20–50) (SI conversion: g/l = mg/100 ml \times 0.01). Synovial fluid analysis revealed fluid of reduced viscosity, leucocytes 0·75 \times 10^9/l, immune complexes 80 \mu g/ml, protein 3·1 g/100 ml (31 g/l), and normal complement levels for this protein concentration (C3 30 mg/100 ml and C4 5 mg/100 ml). No crystals or malignant cells were seen, and the synovial biopsy showed no metastases or inflammatory cell infiltrate, and immunofluorescence was negative. X-rays showed a small opacity in the left lung base and marked subperiosteal new bone formation over the distal ends of the femur, radius, and ulna, but no bone erosions or obvious metastatic deposits were seen. Skeletal imaging showed periosteal linear accumulation of the isotope characteristic of subperiosteal new bone formation, increased isotope uptake in involved joints, and no evidence of metastatic bone disease.

Bronchoscopy was followed by thoracotomy, and a neoplasm removed was found to be an adenocarcinoma. Within 24 hours of the operation joint symptoms had disappeared and swelling had subsided despite persistently high blood levels of CICs. One week postoperatively the patient developed painful swellings in the proximal left radius and distal right femur. These were found on skeletal imaging to be highly suggestive of metastatic deposits, and this was confirmed by open biopsy. When this scan was compared with the preoperative scan, a diminution of the pericortical linear accumulation of the tracer was noted. Repeat x-rays revealed no obvious change in the degree of periosteal new bone formation. Despite this metastatic recurrence, neither the arthropathy nor the pain and tenderness at the distal ends of the radii and ulnae recurred, and clubbing is continuing to resolve.

Case 3. The patient, a 62-year-old man, who presented with a pyrexia of unknown origin and diffuse lymphadenopathy was found to have a poorly differentiated lymphocytic lymphoma, stage 3b. Two weeks after diagnosis he developed a widespread symmetrical polyarthritis involving the ankles, wrists, and elbows, a pericardial friction rub, and haematuria.

Investigations showed the following: Haemoglobin 14·2 g/dl, leucocytes 11 \times 10^9/l, ESR 93 mm/hour; IgM rheumatoid factor was not detected, the ANF was weakly positive, serum C3 was 95 mg/100 ml and C4 38 mg/100 ml (SI conversion: g/l = mg/100 ml \times 0·01) CICs in the serum were 400 \mu g/ml. Blood cultures were negative. Synovial fluid analysis revealed reduced viscosity, leucocytes 1·5 \times 10^9/l (90\% polymorphs), protein 4 g/1000 ml (40 g/l), and immune complexes of greater than 1500 \mu g/ml; immunochemical levels of complement were undetectable. There was no growth or crystals. Synovial biopsy showed no evidence of lymphomatous infiltration of the synovium. Unfortunately the specimen was misplaced and immunofluorescence was not done. Echocardiography confirmed a moderate sized pericardial effusion. Renal biopsy showed a mesangial proliferative glomerulonephritis with areas of acute tubular necrosis. Immunofluorescence showed deposits of C3.

The patient was started on immunosuppressive therapy (prednisolone, vincristine, and cyclophosphamide). Within 48 hours his arthropathy had clinically disappeared and the pericardial rub was no longer heard. The nephropathy resolved within 72 hours. Repeat serum levels of CICs taken 2 weeks later at a time when the patient was asymptomatic were normal. The patient has remained asymptomatic for 3 months.

Discussion

Patient 1 had a symmetrical polyarthritis with all the clinical signs of inflammation as well as marked clubbing of the digits. Analysis of the synovial fluid, however, showed it to be noninflammatory. This apparent dissociation between clinical signs and synovial fluid analysis is found in hypertrophic osteoarthropathy.1,2 Initially this patient had no evidence of periosteal new bone formation but later developed characteristic scintigraphic changes. Schumacher has emphasised that typical periostal changes may be absent in HOA of relatively recent onset.3

Patient 2 had classical HOA. The dramatic symptomatic improvement of the arthropathy following
tumour resection is well described and has recently been documented following chemotherapy. The failure of recurrence of the arthropathy with extrathoracic metastases has been previously reported. Of interest, metastases were not occurred despite in nature, inflammatory reduction in immunofluorescence and therefore case vascular deposits in the synovium of patients with glomeruli patients with lymphoma. Although material, synovium, although immune-complexes, although immune-complex mediated. Recently Vidal et al. testing HOA synovial tissue with fluorescein-conjugated antisera to immunoglobulins and complement components were also unable to find evidence supporting an antigen-antibody immune-deposit mechanism in the pathogenesis of HOA.

Patient 3 had a nonmetastatic polyarthritis associated with malignant lymphoma. The high levels of CICs in the blood, the even higher levels of immune complexes in the synovial fluid, plus the undetectable complement levels in the synovial fluid, followed by a fall in blood CIC levels to normal once the arthropathy had clinically resolved, are good evidence that in this patient the pathogenesis of the nonmetastatic polyarthritis was immune-complex mediated. In spite of being unable to perform immunofluorescence on the synovium all the other evidence together with the simultaneous presence of an immune-complex nephritis lends further support to this suggestion.

The relative rarity of the conditions reported makes the collection of a large series difficult. The evidence suggests that hypertrophic osteoarthropathy is unlikely to be an immune-complex-mediated condition. In the single patient with nonmetastatic polyarthritis associated with lymphoma the evidence does suggest an immune-complex mediated disease.

References

Role of immune complexes in hypertrophic osteoarthropathy and nonmetastatic polyarthritis.
M S Awerbuch and P M Brooks

doi: 10.1136/ard.40.5.470

Updated information and services can be found at:
http://ard.bmj.com/content/40/5/470

These include:

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/