Case report

Multiple microcrystal deposition disease in a patient with systemic lupus erythematosus

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SUMMARY  A 17-year-old female patient with systemic lupus erythematosus (SLE) developed chronic tophaceous gout, chondrocalcinosis and articular capsule calcification in several joints. Analysis of synovial fluid and tophi revealed the coexistence of monosodium urate, calcium pyrophosphate, hydroxyapatite, and cholesterol crystals.

Despite the high frequency of arthritis and soft tissue calcification in systemic lupus erythematosus (SLE) only few cases of crystal-induced synovitis have been recognised in this disease. Talbott and Yu reported the first case of coexisting gout and SLE in 1976. Since then only few cases have been described. Recently microcrystalline uric acid, calcium pyrophosphate dihydrate (CPPD), and hydroxyapatite crystals were found in synovial fluid from an SLE patient with acute synovitis, but so far cholesterol crystals have not been reported in this disease.

It is the purpose of this communication to report a very unusual case of a young female patient with SLE who developed chronic tophaceous gout, chondrocalcinosis, and articular capsule calcification in several joints. Analysis of synovial fluid showed monosodium urate (MSU) and CPPD crystals. Material extruded from tophi revealed MSU and cholesterol crystals. Apatite crystals were suspected on a clinical basis.

Case report

A 17-year-old female patient was first seen in November 1971 because of polyarthritis, malar rash, anasarca, asthenia, and high blood pressure. She denied a past history of drugs. Laboratory tests revealed leucopenia, reticulocyte cell count 4.2%, positive direct and indirect Coomb's test, false positive serological test for syphilis (confirmed by Treponema pallidum immobilisation test), positive LE cells, positive fluorescent antinuclear antibodies (FANA) test (diffuse pattern), anti-DNA 12.1% (normal range <5%), CH50 111 U/ml (normal range 150–250), hyaline and granular casts and red blood cells in the urine sediment, urinary protein excretion 1.8 g/24 h, and creatinine clearance 64 ml/min. A renal biopsy revealed proliferative glomerulonephritis with some epithelial crescents. A diagnosis of SLE was established, and treatment was initiated with prednisone 50 mg, frusemide 80 mg, and spironolactone 25 mg. In February 1972 she had episodes of convulsions, abnormal behaviour, haemoptysis, and chest pain. A chest x-ray showed cotton-wool-like infiltrates and pleural effusion. At that time she was on prednisone 30 mg/day, and was started on azathioprine 100 mg daily with good response. In August 1972 she began complaining of swelling in the left knee, wrists, metacarpophalangeal (MCP) and proximal interphalangeal (PIP) joints, which was successfully controlled with indomethacin.

In 1976 subcutaneous nodules were detected over the extensor surface of elbows, right knee, and MCP and PIP joints of the left middle finger. X-ray studies showed soft-tissue thickening, articular capsule calcification of several MCP and PIP joints (Fig. 1), fluffy calcification in the triangular ligament of the left wrist (Fig. 2), and linear calcification in both knees. Laboratory tests revealed serum uric acid 12...
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mg/dl (0.7 mmol/l), uricosuria 520 mg/24 h, and serum parathormone (PTH) 19 μEq/ml (normal range 2–10). A whitish paste-like material was obtained from an MCP joint nodule revealing abundant needle and rod-shaped crystals with strong negative birefringence on polarised light microscopy. A diagnosis of tophaceous gout was established, and allopurinol started at 300 mg/day.

During the last few years several acute inflammatory bouts have been superimposed on chronic articular manifestations, whereas SLE related symptoms have remained quiescent. In 1981 she was put on maintenance dialysis because of end-stage renal failure. Recent laboratory tests revealed serum uric acid 7.7 mg/dl (0.46 mmol/l), uricosuria 160 mg/24 h, proteinuria 2.6 g/24 h, creatinine clearance 8 ml/min, serum phosphaté 5.9 mg/dl, serum calcium 9.5 mg/dl (2.4 mmol/l), serum PTH 30 μEq/ml, and serum lipid profile within normal limits. She denied joint disease or known hyperuricaemia among her first-degree relatives, and levels of serum uric acid in her mother and 3 siblings were normal.

SYNOVIAL FLUID

Synovial fluid obtained from the left ankle in June 1978 showed a mildly inflammatory fluid with reduced viscosity, poor mucin clot, and leucocytes 8,700 × 10⁹/l. Abundant needle and rod shaped crystals with strong negative birefringence, suggestive of MSU crystals, were found on a wet preparation examined by compensated polarised light microscopy.

A drop of synovial fluid obtained from the PIP joint of the left middle finger in April 1981 revealed strongly negative birefringent needle shaped crystals, suggestive of MSU, and weakly positive birefringent rod shaped crystals suggestive of CPPD. Amorphous material and rod shaped crystals were strongly positive when stained with 2% alizarin red used as a rapid
material obtained from a nodule on the second MTP joint both revealed strongly negative birefringent needle shaped crystals, and large square and rectangular plate-like crystals with notched corners, suggestive of cholesterol (Fig. 3).

Uricase digestion test. A part of a purified suspension of uricase (Sigma Chemical Co.) was added to 10 parts of material obtained from tophi, followed by incubation at 25°C for 2 h at pH 8.5. The needle and rod shaped crystals disappeared, while the plate-like crystals remained unaltered (Fig. 4).

Electron microscopy. One drop of tophus material was promptly placed on a clean glass slide. Six Formvar coated EM grids were floated on the drop for 10 s. The grids were then air dried and examined unstained in a Zeiss EM 10 TEM with a 60 kV beam. The needle and rod shaped crystals exhibited the characteristic ultrastructure of MSU crystals.

X-ray diffraction analysis. Chalky material obtained from tophi was treated with 2 drops of lyophilised hyaluronidase for 15 minutes at 20°C. After centrifugation for 20 min at 2500 rpm, 5 ml of 1% solution of 1:50 trypsin in deionised water was added to the residue, followed by incubation for 4 h at 37°C. The sediment was washed twice in saline and centrifuged at 2500 rpm for 10 min. The resulting sediment was then dried in an oven at 130°C for 30 min, and studied by x-ray diffraction with a Debye-Scherrer power camera (North American Phillips) of 114.6 mm diameter using chromium K alpha radiation with a vanadium filter. Measurements were based on wavelength 0.229092 nm. The specimens were exposed for 20 h at 60 kV and 20 mA. The patterns were characteristic of MSU and cholesterol crystals.

Discussion

Whereas athralgia and nondeforming arthritis are very common, only a few cases of crystal-induced synovitis have been recognised in SLE patients. To our knowledge this is the first case with articular manifestations due to concurrent deposition of MSU, CPPD, hydroxyapatite and cholesterol crystals. Although MSU, CPPD, and hydroxyapatite crystals have been previously identified in association with SLE, so far as we know cholesterol crystals have not been described before. Ryan et al. observed a lipid laden effusion in SLE, but did not report the presence of cholesterol crystals. These crystals have been identified in synovial fluid from patients with rheumatoid arthritis and other rheumatic conditions.

MSU crystals were identified in our patient by their typical strongly negative birefringence on polarised
light microscopy, a positive uricase test, and characteristic x-ray diffraction pattern. Some precipitating factors might be considered as explaining the development of gout in this patient. She was on diuretics and prednisone and had renal failure with increased serum PTH levels. Diuretics are a well known cause of hyperuricaemia and secondary gout, and Schumacher has suggested that corticosteroid therapy might predispose to unusual MSU crystal collections. Impairment of renal function may also contribute to hyperuricaemia, and high levels of uric acid have been recognised in hyperparathyroidism. It seems likely that the MSU crystal deposition resulted from a combination of several factors in our patient.

Our radiological finding of chondrocalcinosis in knees and wrists, and calcification around MCP and PIP joints in both hands is also interesting. Similar features have been described in mixed connective tissue disease, overlap syndromes, primary hyperparathyroidism, and chronic renal failure treated with periodic haemodialysis. However, our patient does not have the clinical criteria for any associated collagen tissue disease, and evidence of multiple crystal deposition antedated by several years her current dialysis programme. On the other hand, although the relationship between primary hyperparathyroidism and articular calcification is well established, we are unaware of any previous report concerning the coexistence of CPPD crystal deposition and secondary hyperparathyroidism. Thus our finding of CPPD-like crystals is intriguing. They might represent apatite crystals or other calcium phosphate compounds as suggested by the alizarin red stain. Rod shaped clumps of apatite, which may be mistaken for CPPD, have been described previously. Nonetheless, because of the small amount of MCP joint fluid obtained, crystals could not be further analysed, and therefore we could only suspect but not confirm CPPD crystals.

The presence of cholesterol crystals within the tophi in this patient is also interesting. Lichestein et al. described lipid material containing tophi, and Ryan et al. reported lipid laden effusions in patients with SLE, but these authors did not describe cholesterol crystals. Recently Reginato et al. have described acute monoarthritis associated with lipid microspheres, the significance of which remains unclear. Since our patient did not have systemic lipid abnormalities, and cholesterol crystals were found within the tophi but not in synovial fluid, we assume that local factors, such as release of lipids due to tissue breakdown, may have been responsible for cholesterol crystal formation.

The increasing number of reports on the coexistence of gout and SLE and on multiple crystal deposition suggest that there is some underlying common predisposition to crystal formation in the connective tissue of these patients. This report illustrates the complexity of such cases and the need to rule out other aetiologies during articular flare-ups in patients with SLE.

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References


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