Juvenile rheumatoid arthritis with rice bodies: light and electron microscopic studies

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SUMMARY  Rice bodies obtained from a young man with juvenile rheumatoid arthritis were found by light and electron microscopy to contain cells that appeared viable. The majority of these cells closely resembled type B synovial lining cells. Type A-like cells were also seen. The cells contained few mitochondria but often much lipid and glycogen, observations which suggested a dependence on anaerobic metabolic pathways in the avascular synovial fluid environment. Cells within the rice bodies lay in a matrix of collagen fibres, fibrin, and amorphous material. The source of the collagen appeared to be the cells themselves. The relatively normal appearance of the cells suggested that they were protected from many of the inflammatory stimuli present in rheumatoid synovia. This 'reversion' towards a normal appearance suggested that the stimuli inducing chronic rheumatoid inflammation might not originate in the synovial lining.

Rice bodies, so named because they resemble polished white rice, were extensively described almost a century ago by Riese (1895). In his classic study he reported the microscopic characteristics of rice bodies obtained from the synovial fluids of patients with tuberculous arthritis. There are no reports of the frequency of rice bodies in synovial fluids from patients with chronic synovitis or in association with specific diseases such as rheumatoid arthritis. In fact our clinical impression is that rice bodies are relatively uncommon. In 1965 Albrecht et al. (1965) described biochemical and ultrastructural details of rice bodies from adults with rheumatoid arthritis. The rice bodies from these cases were small and contained only cell remnants; thus, their description of cellular structure was necessarily limited. The present report describes the light and electron microscopic appearance of the cells and matrix of rice bodies obtained from a young man with juvenile rheumatoid arthritis.

Case report

The patient is a young white man who developed arthritis of the left knee at the age of 15 years in 1973. Swelling of the left calf followed, and an arthrogram showed rupture of a popliteal cyst. Numerous rice bodies were removed at synovectomy, and the synovium was described as hyperaemic and inflamed. The knee improved during the next year. The left ankle became swollen in the autumn of 1974 and the right knee in early 1975. The latter joint was aspirated at that time, and 50 ml of yellow turbid fluid was removed. The white cell count was 49 000/mm³ with 96% polymorphonuclear neutrophils. A few inclusion bodies were present. The mucin clot was poor. Cultures were negative. A consistent programme of aspirin was started at that time.

By June of 1975 the left knee had become massively swollen. A large effusion was aspirated and numerous rice bodies were evacuated as completely as possible through a 16 gauge needle before injection of intra-articular steroid. The fluid was nearly colourless but very turbid. The white cell count was 12 500/mm³ with 74% polymorphonuclear neutrophils. The mucin clot was fair. No crystals were seen. A few inclusion bodies were present. The rice bodies were cut readily with a razor, a finding which suggested a low collagen content.

Symptomatic relief was only temporary and a few rice bodies were obtained intermittently from the left knee. The right calf became swollen in the autumn of 1975 and a firm popliteal cyst enlarged...
behind the right knee. In October of 1976 moderate effusions of both knees, together with a small right popliteal cyst were still present. Full extension was preserved at both knees but flexion was limited to approximately 110°.

LABORATORY DATA
The Westergren erythrocyte sedimentation rate has varied between 40 and 53 mm/h. Tests for antinuclear antibodies and rheumatoid factors have been negative. The peripheral blood lymphocytes were negative for HLA B27 antigen. Ophthalmological examinations have given normal results. Radiographs of the left knee between 1974 and 1975 showed periosteal new bone formation along the medial and lateral femoral condyles and the posterior tibial metaphyseal area. Films have continued to show soft tissue swelling about both knees and the left ankle. A sub-Achilles bursitis with an erosion of the posterior tibial cortex was seen on the left in 1976. Frontal stereoscopic views of the sacroiliac joints have been negative.

Materials and methods
The rice bodies obtained at arthrocentesis were fixed immediately in 2% paraformaldehyde in 0.01 M phosphate buffer, pH 7.2. The bodies were diced into 0.5 to 1.0 mm cubes. After 3 washes in cold 0.01 M phosphate-saline buffer, the tissues were postfixed in Millonig's 1% osmium, pH 7.2, for 2 h at 4°C (Millonig, 1961). The specimens were dehydrated in a graded sequence of ethanols to 100% then infiltrated by and embedded in Spurr epoxy resin (Spurr, 1969).

Thick 1.0 μm and ultrathin sections were cut on a Reichert ultramicrotome. The thick sections were stained with 1.5% aqueous toluidine blue O. The ultrathin sections were mounted on Formvar and carbon-coated copper grids and stained with a saturated solution of uranyl acetate in 50% ethanol and with Reynolds's lead citrate solution (Reynolds, 1963). The grids were examined in a JEOL 100B electron microscope at 60 kV.

Results
LIGHT MICROSCOPY
The rice bodies after aspiration appeared as a mass of loosely aggregated material (Fig. 1). The individual rice bodies varied from 0.2 to 1.5 cm in length and were oval to slightly irregular in shape. They were white to slightly yellow and were contained in blood tinged synovial fluid. The toluidine blue O thick sections showed cells scattered in a blotchy, purple-blue amorphous matrix (Fig. 2). Fibrils were seen around the periphery. Cells of sizes 7 to 21 μm appeared viable, and most were surrounded by pale blue or clear zones. Three types of cells were seen. Some resembled synovial lining cells and sometimes contained granules; some were spindle-shaped and...
The predominant cell within the rice bodies was most like a type B synovial lining cell (Fig. 3) (Barland et al., 1962). Some of these were spindle-shaped and similar to fibrocytes of deeper areas of the synovium. All cells appeared viable and to be in active protein synthesis as shown by well developed rough endoplasmic reticulum. The lamellae of the endoplasmic reticulum were often dilated, and their contents were homogeneous in appearance. No intra-articular collagen fibril formation was noted. Most cells contained glycogen and sometimes large lipid drops (which appeared to correspond to the pale blue vacuoles of the toluidine blue stained sections). The lipid was nonmembrane bound and often surrounded by the glycogen granules. Mitochondria were sparse, markedly fewer than usual, and centrioles were occasionally seen. Dark staining lysosomal granules were present in moderate numbers in some cells and were few in others. Dark staining granules were also seen extracellularly close to cell surfaces and were often entwined in filopodia (Fig. 4). Some granules had the appearance of being phagocytosed; occasionally an intracellular granule was bound by a double membrane. Groups of free ribosomes were seen in the cytoplasm, and microfilaments were easily observed. Microtubules were not noted. Interestingly, Golgi apparatus was rarely observed within any cells. Nuclei generally were slightly irregular in shape and contained 1 or 2 nucleoli. No particles suggestive of virus or mycoplasma were seen within any cells. Cell membrane activity was marked as shown by many slender filopodia and micropinocytic vesicles (coated pits). Around most of the cells was a zone of collagen fibres, sometimes with fine whiskery material (Fig. 4, inset A). The collagen fibres were occasionally aligned in parallel with each other; however, most were short and without specific orientation. The periodicity of the collagen was 66 nm. The immediate area around a few cells was almost devoid of any amorphous material or fibres. Away from the zones of collagen a mesh of amorphous material and fibrin (sometimes with a periodicity typical of coagulated fibrin) made up the mass of a rice body (Fig. 4, inset B). One effete polymorphonuclear neutrophil was seen by electron microscopy. The nucleus was homogeneous and the cytoplasm was partially destroyed.

**Discussion**

Rice body formation occurs in a small number of patients with adult rheumatoid arthritis, and in this report we note this phenomenon in a young man with a persistent course of late-onset, oligoarthritic juvenile rheumatoid arthritis. The more usual finding in rheumatoid synovial fluids is that of clumps of fibrin, a byproduct of inflammation.
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Fig. 3 Part of a 'synthetic' or B type appearing fibroblast containing very well developed rough endoplasmic reticulum lamellae (rer). No mitochondria can be identified in this view, however, nonmembrane bound lipid (Lip) and glycogen (GLY) can be seen. The pericellular area contains small collagen fibres (Col). Fibrin (FIB) is seen well away from the cell. Filopodia (fil), lysosomes (Ly), nucleus (N), nucleolar material (n). The arrow points to a micropinocytotic vesicle (coated pit) (uranyl acetate, lead citrate, × 9600).

(Cohen et al., 1975). The morphology of the cells within these rice bodies suggested their synovial origin, presumably from sloughing lining cells. The cytoplasmic appearance of the rice body cells was closer to that of normal B lining cells than A cells (Barland et al., 1962). This suggested that the environment, though porous to nutrients, was a protected one, that the cells could proceed primarily with collagen and hyaluronate synthesis, and that some of the intense stimuli to primary lysosome formation which are present in the inflamed rheumatoid synovial lining were excluded. The differences from normal lining cells which were seen in these cells suggested that local environmental factors within the rice bodies produced them. The lipid and glycogen observed and the relative paucity of mitochondria suggested the greater use of anaerobic pathways of energy utilisation in this avascular environment of synovial fluid. To some extent the cells' appearances were reminiscent of fibroblasts grown in tissue culture, especially cultures maintained for longer than 1 week without trypsinisation (Wynne-Roberts and Castor, 1972).

The explanation for the extracellular granules,
Fig. 4  A more A-like cell showing marked filopodia formation (fil). The extracellular granules (G) may be in the process either of being extruded or of being ingested by the cell. The cell contains little rough endoplasmic reticulum (rer) and pleomorphic dark staining granules, some of which are most likely lysosomes (Ly), while others may be phagolysosomes or perhaps residual bodies. Free ribosomes (Rb) are scattered in the cytoplasm. Well marked micropinocytotic vesicles are arrowed. Nucleus (N), nucleolus (n), glycogen (GLY), microfilaments (mf) (uranyl acetate and lead citrate, \( \times 12000 \)). A: Small collagen fibres (Col) show periodicity which measures 66 nm (uranyl acetate and lead citrate, \( \times 34000 \)). B: Fibres from a rice body matrix show the appearance and periodicity typical of coagulated fibrin (FIB) (uranyl acetate and lead citrate, \( \times 34000 \)).

some apparently being phagocytosed, was not obvious. Their appearance was certainly similar to that of some intracellular lysosomes. Reasons for the pericellular clear zones might include an artefact of dehydration or that the cells secreted enzymes into their immediate vicinity, which destroyed fibrin. Possibly some of the extracellular granules might also play a role in fibrin destruction. Over a period of time the cells would then secrete collagen and mucopolysaccharide matrix into the pericellular area.

The rarity of inflammatory cells within these rice
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The authors conclude that rice bodies represent an end product of synovial inflammation, proliferation, and secondary degeneration.

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References


