Case reports

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Active chronic hepatitis and haemolytic anaemia associated with Rh-specific antibodies

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Active chronic hepatitis, a disease also known by the synonyms lupoid hepatitis, plasma cell hepatitis, chronic juvenile hepatitis and by a number of others, is now recognized as a not uncommon condition (Sherlock, 1968). None of these synonyms is entirely satisfactory but the name lupoid hepatitis has been used because in 15% of patients disturbance of the immunity mechanisms gives rise to a positive LE-cell test. Another manifestation of auto-immunity may be the development of a positive Coombs' test, or even a frank haemolytic anaemia (Read, Harrison & Sherlock, 1963), but this appears to be a rare complication of the disorder and these authors report only one case. We have been unable to find any references to patients with specific antibodies on the red cells or in the serum.

Case history

Mrs E. F., aged 62 years, without any previous illness of note, was referred to one of us (CDRP) on 22 February 1969. She had become jaundiced about 6 weeks previously. Her stools had been pale and urine dark. The illness had consisted of vague upper abdominal pain, nausea, heartburn and a little dysphagia referred to the lower end of the oesophagus. There was no history of alcoholism or of the taking of drugs. She had had two normal pregnancies and no miscarriages, and she had never received any blood transfusions or injections of blood. The illness was accepted by her general practitioner as being an example of ordinary viral hepatitis and the jaundice apparently improved, but had become worse a few days before 22 February. Liver function tests had shown the following results: bilirubin 3·6 mg/100 ml; alkaline phosphatase 34 KA units/100 ml; thymol turbidity 6·5 units; SGOT 148 IU/l at 25° C; SGPT 168 IU/l at 25° C; total protein 8·7 g/100 ml; albumin 3·2 g/100 ml; globulin 5·5 g/100 ml; electrophoresis showed marked increase in gamma-globulin.

On examination she looked generally well and was moderately jaundiced. Her liver was only slightly enlarged but very easily palpable and firm. Urine contained both bile and an excess of urobilinogen but was otherwise normal. There were no clinical signs of liver failure, no spider naevi and no palmar erythema. Her spleen was not palpable. Her blood pressure was 160/90 mmHg. She was admitted
to hospital for further investigation on 25 February 1969. On admission there were no new clinical features though after a few days her spleen became just palpable and her liver had enlarged to about 3.5 cm below the costal margin and was still very firm.

**Investigations**

ECG and X-ray chest normal. Serum electrolytes normal; blood urea 23 mg/100 ml; ESR (Wintrobe) 50 mm/hr, haemoglobin 11.6 g/100 ml, WBC 5700/mm³; differential count; polymorphs 40%, lymphocytes 40%, monocytes 6%, eosinophils 11% and basophils 3%; reticulocytes 2.4%; blood film; some anisocytosis only; platelets 280,000/mm³; mid-stream urine: no abnormal findings; liver function tests: bilirubin 5.4 mg/100 ml; alkaline phosphatase 29.5 KA units/100 ml with increase of intestinal isoenzymes suggestive of cirrhosis; thymol turbidity 6.0 units; SGOT 144 IU/l; SGPT 212 IU/l; protein 7.5 g/100 ml, albumin 2.9 g, globulin 4.5 g; immunoelectrophoresis: IgM 100 mg/100 ml (normal), IgG 2500 mg/100 ml (normal 60–1600), IgA 600 mg/100 ml (normal 150–450). Immuno-fluorescent tests for antinuclear factor (ANF), mitochondrial antibody and smooth muscle antibody all negative; no Australia antigen or antibody detected; thyroid auto-precipitin test, thyrotropic (anti-microsomal) C–F test and tanned cell agglutinating anti-thyroglobulin titre all negative; prothrombin level 100%; partial thromboplastin time normal; bleeding time (Duke) 2.5 min; serum cholesterol 150 mg/100 ml; blood group O Rhesus genotype R_1R_2 (cDE/cDE); occult blood tests on the stools negative or slightly positive; Rose-Waaler test negative; direct Coombs' test positive, serum noted to contain anti-e. Needle biopsy of the liver showed derangement of the normal architecture with a prominent mixed inflammatory cell infiltrate of lymphocytes, plasma cells, eosinophils and some polymorphs both in the portal tracts and in the lobules (Fig. 1). The appearances were those of an active chronic hepatitis with secondary micro-cirrhotic changes.

Blood from her daughters was tested by the Manchester Blood Transfusion Service and the probable Rh genotype of both was R_1R_2 (cDE/cDE).

**Early treatment and course**

Prednisolone 5 mg b.d. was started on 26 March and she was put on a high protein diet. Within a week she had clinically improved and her liver was less palpable and less firm. Her jaundice was less (bilirubin 4.8 mg/100 ml), but the other liver function tests and immuno-globulin levels were virtually unchanged. She was discharged on the same dose of prednisolone on 9 April 1969, at which time the haemoglobin was 10.8 g/100 ml, reticulocyte count 3%, and the total white count and differential count were normal.

**Haemolytic episode**

She had further improved when seen again in the OPD on 6 May, and had gained weight but was still jaundiced and her liver and spleen were still palpable. Hb 8.2 g/100 ml; WBC 7200/mm³; reticulocytes 10%. Sternal puncture showed an active cellular marrow with normoblastic erythropoiesis, the appearances being compatible with a haemolytic process. Urine contained an excess of both urobin and urobilinogen, but not bile. Serum haptoglobins showed 125 mg haemoglobin binding/100 ml.

She was re-admitted on 22 May with fast auricular fibrillation. Her haemoglobin was virtually unchanged at 7.9 g/100 ml but the reticulocytes had increased to 23.5%. Red cell survival with ^51Cr gave a half chromium time (T½) of 4.5 days, or a mean cell life span of about 13 days by Method A of Mollison (1956). Serological tests by the Manchester Blood Transfusion Service showed anti-C plus anti-e to be present acting in serum albumin.
at 37°C and in the indirect Coombs' test at 37°C but only when cells were washed with saline, cooled to 4°C and buffered to pH 8-2. The direct Coombs' test was positive, but again only when the cells were washed with saline as above.

Her jaundice persisted (bilirubin 5.8 mg/100 ml) but her other liver function tests showed improvement: alkaline phosphatase 13.5 KA units/100 ml; thymol turbidity 1.0 unit; SGOT 34 IU/l; SGPT 11 IU/l; albumin 3.8 g/100 ml, globulin 3.1 g/100 ml.

**Further course and treatment**

The prednisolone dosage was increased to 10 mg t.d.s. on 4 June. She was started on cyclophosphamide 100 mg/day, and her auricular fibrillation controlled with digoxin 0.25 mg b.d. The prednisolone dose was increased to 60 mg b.d. on 10 June. A blood transfusion of 2 units was given on 13 June 1969. Her haemoglobin gradually rose to 11.6 g/100 ml and it was soon possible to decrease the dosage of prednisolone again (Fig. 2), and the cyclophosphamide was reduced to 50 mg daily on 21 June. She was discharged on 24 July 1969, with a haemoglobin of 11.7 g/100 ml, reticulocyte count 3.5%, platelets 200,000/mm³, and WBC 2800/mm³. At that time she was taking prednisolone 5 mg b.d. and cyclophosphamide 50 mg daily. She has been reviewed at intervals. Her liver has become much smaller and the spleen impalpable. It has been possible to reduce her steroid therapy to 2 mg of prednisolone daily, but the cyclophosphamide has been continued in the same dose. A further needle biopsy of the liver carried out on 5 March 1970, showed basically the same appearances as before. The liver function tests have been normal since 25 February 1970. Immunoelectrophoresis (June 1969): IgM 100 mg/ml, IgG 1500 mg/ml, IgA 450 mg/100 ml. A recent blood count (November 1970) shows haemoglobin 12.9 g/100 ml, WBC 4000/mm³, platelets 168,000/mm³.

**Discussion**

Doniach *et al.* (1970) group three liver conditions in their concept of 'autoallergic' hepatitis. These are
biliary cirrhosis, active chronic hepatitis and certain forms of cryptogenic cirrhosis, particularly in women. All these conditions show a high incidence of positive autoimmune marker tests. In our patient although the ANF, mitochondrial and smooth muscle antibody tests were all negative, the LE-cell test was positive and an autoimmune haemolytic anaemia with demonstrable antibodies developed during the course of the active liver disease. The histological changes in the liver on three different occasions showed a picture of a chronic aggressive type of hepatitis with architectural lobular disorganization, the essential features required for a diagnosis of active chronic hepatitis (Scheuer, 1968).

The autoimmune haemolytic anaemia in our patient was associated with antibodies of Rh specificity (anti-C and anti-e) although her Rh genotype was R2R2 (cDE/cDE). The haemolysis was corrected with adrenocorticosteroid and cyclophosphamide therapy (Fig. 2). We have been unable to find any other comparable case in the literature, although Dacie & Worledge (personal communication) have seen one case of chronic hepatitis with haemolytic anaemia and an antibody of Rh specificity (anti-e).

The inhibitory effect of cyclophosphamide on immunity mechanisms is well known (Tripathy & Mackaness, 1969a, b; British Medical Journal, 1970). However, in our patient no improvement in the liver lesions was found. The active inflammatory process apparently continued unchanged in spite of the control of the immunological disorder. This is not surprising as the haemolysis is evidence of derangement of immunological balance which is not necessarily related to the underlying pathogenesis of the liver disorder.

The anti-e could have been the result of sensitization during either of her two pregnancies, but the anti-C could not have been accounted for in this way. It seems probable that both antibodies were entirely the result of a deranged immunological mechanism and that they arose at the time of her hepatitis and not before.

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