Infectious mononucleosis complicated by acute haemolytic anaemia with a positive Donath-Landsteiner reaction

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SYNOPSIS  A patient with infectious mononucleosis complicated by acute haemolytic anaemia is described. The patient had, in addition to non-specific cold agglutinins in the serum, a positive Donath-Landsteiner reaction in the absence of syphilis. The association of a positive Donath-Landsteiner reaction with haemolytic anaemia in infectious mononucleosis has been described only once before.

Acute haemolytic anaemia is an uncommon complication of infectious mononucleosis. In a recent survey of the literature on this subject only 53 cases were found (Worledge and Dacie, 1969). Hoagland (1967) stated that approximately 3% of the patients in his series of 500 cases had haemolytic anaemia.

In addition to the heterophil antibodies reactive with sheep red cells, which are a major criterion for the diagnosis of infectious mononucleosis, patients with this condition have often been found to have other hetero- and autoantibodies in the course of the illness. These include reagins, which give rise to false positive Wassermann reactions (Zarafonitis and Kent, 1954; Carter 1966), rheumatoid factor, antinuclear factor, an antibody similar to antithyroglobulin, and possibly leucoagglutinins (Carter, 1966). Autoantibodies to red cells are found not infrequently and may be associated with autoimmune haemolytic anaemia. Those most commonly described are cold agglutinins, either non-specific (Ellis, Wollenman, and Stetson, 1948; Green and Goldenberg, 1960; Fekete and Kerpelman, 1965) or with anti-i specificity (Rosenfield, Schmidt, Calvo, and McGinniss, 1965; Jenkins, Koster, Marsh, and Carter, 1965; Calvo, Stein, Kochwa, and Rosenfeld, 1965; Troxel, Innella, and Cohen, 1966). A number of patients have been investigated fruitlessly for cold haemolysins of the Donath-Landsteiner type (Green and Goldenberg, 1960; Kostinas and Cantow, 1966); only one case of infectious mononucleosis with haemolytic anaemia and a positive Donath-Landsteiner reaction has actually been recorded (Ellis et al, 1948). This paper presents a second case of this association occurring in the absence of syphilitic infection.

Case Report

The patient, a white male aged 17 years, was admitted to Fremantle Hospital on 20 July 1971, with a three-day history of dull, constant epigastric pain, anorexia, headache, weakness, tiredness, and passing dark urine. He was an English migrant and had arrived by air in Australia eight days earlier. Before leaving England he had been well and had had no known contact with patients suffering from hepatitis or infectious mononucleosis, nor any exposure to drugs or toxic chemicals.

He was mildly jaundiced and afebrile. There was mild, generalized, painless lymphadenopathy, particularly of the cervical, posterior cervical, and epitrochlear glands. The spleen was palpable 3 cm below the left costal margin and was tender. The liver was palpable 2 cm below the right costal margin and was also slightly tender. No other abnormalities were noted on examination.

Results of the most relevant investigations performed on admission and subsequently are set out in the table. The Paul-Bunnell test was performed by the State Health Laboratories, Perth, using the technique of Davidsohn (1937) as outlined by Wintrobe (1946). The Donath-Landsteiner test was performed using both the direct and indirect technique as set out by Boorman and Dodd (1970). Immunoglobulin levels were determined by radial immunodiffusion on Tripartigen plates (Behringwerke).

In addition to these, on presentation a screening

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### Table 1  Results of principal investigations

<table>
<thead>
<tr>
<th>Investigation</th>
<th>Date</th>
<th>21 July 1971</th>
<th>26-29 July 1971</th>
<th>6 August 1971</th>
<th>1 December 1971</th>
</tr>
</thead>
<tbody>
<tr>
<td>Haemoglobin (g/100 ml)</td>
<td></td>
<td>9-0</td>
<td>11-1</td>
<td>13-0</td>
<td>15-8</td>
</tr>
<tr>
<td>Haematocrit (%)</td>
<td></td>
<td>30</td>
<td>33</td>
<td>39</td>
<td>49</td>
</tr>
<tr>
<td>Reticulocytes (%)</td>
<td></td>
<td>3-3</td>
<td>9-8</td>
<td>3-0</td>
<td>1-7</td>
</tr>
<tr>
<td>White cells per mm³</td>
<td></td>
<td>11 200</td>
<td>6800</td>
<td>4100</td>
<td>6000</td>
</tr>
<tr>
<td>ESR (Westergren) (mm/hr)</td>
<td></td>
<td>9400¹</td>
<td>4600 (64% atypical)</td>
<td>2500¹</td>
<td>—</td>
</tr>
<tr>
<td>Direct Coombs test</td>
<td></td>
<td>Positive</td>
<td>—</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>Cold agglutinins</td>
<td></td>
<td>1 in 128</td>
<td>—</td>
<td>1 in 128</td>
<td>1 in 4</td>
</tr>
<tr>
<td>Donath-Landsteiner reaction</td>
<td></td>
<td>Positive</td>
<td>—</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>IM spot test (Denco)</td>
<td></td>
<td>Positive</td>
<td>—</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>Paul Bunnell reaction</td>
<td></td>
<td>1 in 112²</td>
<td>1 in 56</td>
<td>—</td>
<td>Negative</td>
</tr>
<tr>
<td>Wassermann reaction</td>
<td></td>
<td>Positive</td>
<td>—</td>
<td>Negative</td>
<td>—</td>
</tr>
<tr>
<td>Serum bilirubin (mg 100 ml)</td>
<td></td>
<td>2-3</td>
<td>0-8</td>
<td>—</td>
<td>0-8</td>
</tr>
<tr>
<td>(direct reacting)</td>
<td></td>
<td>(0-7)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Serum alkaline phosphatase</td>
<td></td>
<td>11</td>
<td>14</td>
<td>—</td>
<td>9-5</td>
</tr>
<tr>
<td>SGOT (normal range 0-35 IU/ml)</td>
<td></td>
<td>83</td>
<td>100</td>
<td>34</td>
<td></td>
</tr>
<tr>
<td>SGPT (normal range 0-35 Karmen U/ml)</td>
<td></td>
<td>82</td>
<td>138</td>
<td>—</td>
<td>10</td>
</tr>
<tr>
<td>Haemoglobins (mg/100 ml haemoglobin-binding capacity)</td>
<td></td>
<td>0</td>
<td>40</td>
<td>—</td>
<td>75</td>
</tr>
<tr>
<td>Immunoglobulins (mg/100 ml)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>IgG (Normal range 800-1600)</td>
<td></td>
<td>1160</td>
<td>1330</td>
<td>1250</td>
<td></td>
</tr>
<tr>
<td>IgA (Normal range 90-500)</td>
<td></td>
<td>315</td>
<td>315</td>
<td>100</td>
<td></td>
</tr>
<tr>
<td>IgM (Normal range 80-300)</td>
<td></td>
<td>140</td>
<td>243</td>
<td>140</td>
<td></td>
</tr>
</tbody>
</table>

¹Atypical lymphocytes of the type seen in infectious mononucleosis present.
²The Paul Bunnell reaction initially was also positive at 1 in 112 after absorption with guinea pig red cells, and negative at 1 in 14 after absorption with ox red cells.

About four months later he remained well with no jaundice or recurrence of symptoms. His spleen and liver were not palpable and there was no significant lymphadenopathy.

### Discussion

Dameshek (1943) first described a possible case of acute haemolysis complicating infectious mononucleosis. Since then at least 53 such cases have been reported (Worlledge and Dacie, 1969). The haemolysis is typically transient (Dacie, 1962) and in most instances develops one to two weeks after the onset of the illness, although in some cases haemolytic anaemia and infectious mononucleosis develop simultaneously (Dacie, 1962).

The patient described had a transient haemolytic episode lasting only a few days and apparently simultaneously with the onset of infectious mononucleosis. That the patient was actually suffering from infectious mononucleosis was confirmed by the typical morphological changes in his blood picture and by the presence of heterophil antibodies reactive with sheep red cells (the Paul-Bunnell test). A titre of 1 in 112, as in this case, is considered a ‘borderline titre’ (Davidsohn, 1937) but other causes of an elevated titre are ruled out in this case by the result of the differential absorption test (Davidsohn, 1937; Davidsohn, Stern, and Kashiwagi, 1951), and would be considered diagnostic by these authors.
The most interesting aspect of this case is the presence of a cold haemolysin in the serum in the absence of syphilis. Ellis et al (1948) described a patient with a history suggestive of virus pneumonia but with a blood picture suggestive of infectious mononucleosis and a heterophile antibody titre of 1:1024. Differential absorption titres were not performed. Cold agglutinins against human group O cells were present to a titre of 1 in 256 and agglutination also took place at room temperature, not dispersing completely at 37°C. The Donath-Landsteiner reaction was positive for at least seven weeks, although there was no serological evidence of syphilis. We were unable to find any other evidence in the literature of this finding in a case of haemolytic anaemia complicating infectious mononucleosis. Kostinas and Cantow (1966) studied Donath-Landsteiner reactions in 41 cases of infectious mononucleosis and all were negative. Green and Goldenberg (1960) reported a case of acute haemolytic anaemia complicating infectious mononucleosis with a positive direct Coombs test and cold agglutinins to a titre of 1:256. The Donath-Landsteiner reaction was negative.

The patient described in this report had a positive Donath-Landsteiner reaction both when the patient’s fresh serum was tested with his own red cells and with normal red cells, both with and without an added source of complement. It remained positive for at least eight days. This antibody is usually found only in association with syphilis (Becker, 1948; Dacie, 1962), but it has been described infrequently in the absence of syphilis (Sweetnam, Murphy, and Woodcock, 1952; Dacie, 1954).

This patient had a positive Wassermann reaction at the acute stage of his illness which reverted to negative without treatment. The other serological tests for syphilis performed (VDRL flocculation test and Reiter protein complement-fixation test) were negative at all times. There was no clinical evidence of syphilis at any time. Transient false positive Wassermann reactions have been described in infectious mononucleosis (Carter, 1966), and it would appear that this was the case in this patient. Of particular interest in this regard is the patient reported as case report no. 18 by Dacie (1954). The patient was a little girl who developed paroxysmal cold haemoglobinuria following an attack of measles. She had a positive Donath-Landsteiner reaction, and in the acute stage of her illness a doubtful positive Wassermann reaction. Other serology for syphilis was negative, and there was no clinical evidence of congenital syphilis.

The clinical course and investigation of this patient led us to conclude that he suffered from infectious mononucleosis, complicated by autoimmune haemolytic anaemia associated with cold autoagglutinins and a cold haemolysin of the Donath-Landsteiner type, and by mild hepatitis. While his Wassermann reaction was positive during the acute illness, there was no historical, clinical, or other serological evidence of syphilis.

He did not show the typical syndrome of paroxysmal cold haemoglobinuria; in the mild winter climate of Perth this is not surprising.

We wish to thank Dr J. D. Woods for allowing us to study the patient, and Dr G. A. Leyland, medical superintendent, Fremantle Hospital, for permission to publish this report.

References
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