Haemorrhagic colitis and haemolytic-uraemic syndrome: False positive reaction with a rotavirus latex agglutination test

R P Bendall, J J Gray

Abstract

A stool sample from a child with haemorrhagic colitis and haemolytic-uraemic syndrome gave a positive reaction with the RotaScreen latex agglutination test in the absence of other evidence of rotavirus infection. When this test is performed on bloody specimens, positive reactions should be interpreted with caution and confirmed by other means.

Haemorrhagic colitis can be caused by an infection with verotoxin producing strains of Escherichia coli 0157. This in turn has been associated with the haemolytic-uraemic syndrome which is an uncommon condition, principally of children, and associated with significant morbidity and mortality. By contrast, rotavirus infection is a common, self-limiting condition and does not usually cause any long term sequelae. We describe a patient with haemorrhagic colitis and haemolytic-uraemic syndrome whose stool gave a false positive reaction in a latex agglutination test for detecting rotavirus.

A 4 year old girl was admitted to this hospital with a three day history of diarrhoea, becoming bloody on the day of admission. She was vomiting but not clinically dehydrated. She had an oral temperature of 37.3°C. Physical examination was normal and her white cell count was raised at 18.5 × 10^9/l with normal haemoglobin at 145 g/l and a platelet count of 404 × 10^9/l. Urea and electrolytes were all within normal limits (sodium 136 mmol/l, potassium 4.0 mmol/l, urea 3.2 mmol/l and creatinine 62 μmol/l).

A faecal sample was sent for bacteriological and virological examination; it was bloody and liquid. The RotaScreen latex agglutination test (Mercia Diagnostics Ltd, Surrey) for detecting rotavirus antigen was performed according to the manufacturer’s instructions on a centrifuged extract of this sample. It agglutinated the latex particles coated with rabbit anti-rotavirus antibody (test latex), but not the latex particles coated with normal rabbit globulin (control latex); this was interpreted as evidence of rotavirus infection.

By the third day after admission the patient had become oliguric with peripheral oedema noted on examination. The haemoglobin had fallen to 101 g/l, there was thrombocytopenia (platelet count of 54 × 10^9/l) and a white cell count of 11.6 × 10^9/l. The blood film contained numerous red cell fragments, spherocytes and Burr cells. There was evidence of renal impairment with urea at 15 mmol/l and creatinine at 279 μmol/l. Haemorrhagic colitis and haemolytic-uraemic syndrome were diagnosed.

The next day an organism, subsequently identified as a verotoxin producing strain of E coli serotype 0157, was isolated from the stool. No other enteric pathogens were isolated. Her renal function deteriorated further and she required peritoneal dialysis for 12 days.

Stool samples from days 1, 3, and 5 were retested with the RotaScreen test, examined by electron microscopy, and tested with a

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**Results of rotavirus antigen detection and electron microscopy**

<table>
<thead>
<tr>
<th>Day faecal sample taken</th>
<th>Latex agglutination</th>
<th>Rotavirus antigen ELISA</th>
<th>Electron microscopy</th>
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<tbody>
<tr>
<td></td>
<td>Test</td>
<td>Control</td>
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<tr>
<td>1</td>
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<td>Negative</td>
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<td>3</td>
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<td>Negative</td>
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<td>5</td>
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<td>Negative</td>
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Detection of Borrelia burgdorferi in patients with Lyme disease by the polymerase chain reaction

E C Guy, G Stanek

Abstract

*Borrelia burgdorferi*, the causative agent of Lyme disease, was detected in patients' serum by DNA amplification using the polymerase chain reaction (PCR). *B burgdorferi* was pelleted from serum samples by centrifugation (10 000 × g for 10 minutes) and lysed by treatment with ammonium hydroxide (100°C for 15 minutes). Two pairs of “nested” PCR primers complementary to the gene encoding a major outer surface protein (OSP A) of *B burgdorferi* were used in DNA amplification under standard PCR conditions (Perkin-Elmer Cetus). Two out of five patients with erythema migrans, the characteristic primary skin lesion associated with early Lyme disease, were positive by the PCR. This method could form the basis of a useful routine laboratory test in those cases of early Lyme disease where conventional serological testing commonly yields equivocal or false negative results.

The early diagnosis and treatment of Lyme disease can prevent progression to the later, more serious neurological and arthritic complications. Unfortunately, the serodiagnosis of early Lyme disease is difficult due to the absence of barely detectable antibody response in the first weeks of infection. Furthermore, isolation of *Borrelia burgdorferi*, the causative agent of Lyme disease, from specimens is notoriously slow and difficult. We report a method, based on the polymerase chain reaction (PCR), for the direct detection of *B burgdorferi* in the serum of patients manifesting the primary skin lesion of Lyme disease, erythema migrans.

Methods

Two pairs of “nested” PCR primers were constructed complementary to the gene encoding the *Osp-A* protein of *B burgdorferi* strain B31. Primer pair 1, *OspA-N1* (5'-GAGCTTAAAGGAACTTCGATAA-3') *OspA-C1* (5'-GTATTGGTTGACTGTAA-TTGT-3'), and primer pair 2, *OspA-N2* (5'-ATGGATCTGGAGTACCTGAA-3') *OspA-C2* (5'-CTTAAAGTAAAGTTCTCT-3'), correspond to nucleotide nos 334-356, 874-894, 362-381 and 693-713, respectively, of the *OspA* gene. The primers chosen are complementary to regions of the gene that are highly conserved between strains B31, ACA1,
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