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James P. Brinker
University of Massachusetts Medical School

Neil R. Blacklow
University of Massachusetts Medical School

Xi Jiang

See next page for additional authors

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Authors
James P. Brinker, Neil R. Blacklow, Xi Jiang, Mary K. Estes, Christine L. Moe, and John E. Herrmann
Immunoglobulin M Antibody Test To Detect Genogroup II Norwalk-Like Virus Infection

JAMES P. BRINKER, NEIL R. BLACKLOW, XI JIANG, MARY K. ESTES, CHRISTINE L. MOE, AND JOHN E. HERRMANN

Division of Infectious Diseases and Immunology, University of Massachusetts Medical School, Worcester, Massachusetts 01655; Center for Pediatric Research, Children’s Hospital of the King’s Daughters, Eastern Virginia Medical School, Norfolk, Virginia 23510; Division of Molecular Virology, Baylor College of Medicine, Houston, Texas 77030; and Department of Epidemiology, University of North Carolina, Chapel Hill, North Carolina 27599

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Sera obtained from adult volunteers inoculated with genogroup II Norwalk-like viruses (NLVs), Hawaii virus, and Snow Mountain virus and from patients involved in outbreaks of gastroenteritis were tested for genogroup II NLV Mexico-virus-specific immunoglobulin M (IgM) by use of a monoclonal antibody, recombinant Mexico virus antigen (rMXV)-based IgM capture enzyme-linked immunosorbent assay (ELISA). Sera from genogroup I Norwalk virus (NV)-inoculated volunteers and from patients involved in a genogroup I NLV outbreak were also tested. In sera from those infected with genogroup I NV or NLVs in volunteer and outbreak studies, only 3 of 25 were rMXV IgM positive; in contrast, 24 of 25 were IgM positive for recombinant NV (rNV). In sera from those infected with genogroup II NLVs in volunteer and outbreak studies, 28 of 47 were rMXV IgM positive and none were IgM positive for rNV, showing the specificity of each IgM test for its respective genogroup. In an outbreak of gastroenteritis not characterized as being of viral etiology but suspected to be due to NV, 7 of 13 persons had IgM responses to rMXV, whereas none had IgM responses to rNV, thus establishing the diagnosis as genogroup II NLV infection. The rMXV-based IgM capture ELISA developed is specific for the diagnosis of genogroup II NLV infections.

MATERIALS AND METHODS

Serum samples. Sera were obtained from patients in volunteer studies and in outbreaks of gastroenteritis. For sera from individuals known to be infected with genogroup I NLV, nine paired sera were from NV-inoculated volunteers who had been infected with the 8FIIa strain and were shown to be positive by seroconversion and by an IgM ELISA for NV (1). Sixteen paired sera from an outbreak of gastroenteritis in Erie County, New York, and originally diagnosed as NV positive by human reagent-based antigen and antibody assays were tested (10). By genotyping (1), the outbreak from which the sera used here were obtained was shown to be associated with a genogroup I NLV (V Ward J/90). Sera from patients infected with genogroup II NLVs included paired sera obtained in two HV-inoculated volunteer studies with two volunteers each (unpublished data), along with three convalescent-phase sera from SMV-inoculated volunteers (obtained from R. Dolin, University of Rochester). Both groups consisted of individuals who became ill. Sera from outbreaks included 21 paired sera from an outbreak of SMV in a New York City high school cafeteria in 1985 (16). This outbreak involved approximately 600 students and cafeteria workers. Acute-phase sera were collected 4 to 8 days after the onset of symptoms, and convalescent-phase sera were collected 2 weeks later. Two paired sera from individuals infected with Taunton virus (TNV) (2) were obtained from D. Lewis, Leeds, United Kingdom. Sera collected during investigations of two additional outbreaks were also tested. Five paired sera and eight single convalescent-phase sera were obtained from patients involved in an outbreak at a nursing home (University of Massachusetts Medical Center—University Commons [UMMC-UC]) in 1996 (unpublished data). During a 2-week period, 68 residents and staff members became ill with gastroenteritis. Routine examination for bacterial and parasitic agents by the University of Massachusetts Medical Center clinical microbiology laboratory yielded negative results. Stool samples from three patients were tested by reverse transcription-PCR for NLVs. The portion sequenced (57 bases) had 95% identity in the polymerase region with MD-V6, a genogroup II NLV involved in an outbreak in a Maryland nursing home in 1987 (28) (accession no. M70163), and 98% identity with Halifax NLV (unpublished data) (accession no. NLU37651). The second outbreak involved patients who developed gastroenteritis after a Rhode Island graduation banquet in 1986.

Microscopy to detect IgM to genogroup II NLVs (3, 30), and IgM responses to rMXV antigen in sera from persons infected with small round structured viruses have been reported (17). The purpose of this study was to determine the efficacy of a monoclonal antibody, rMXV-based IgM capture ELISA for the diagnosis of genogroup II NLV infections.

In a recent classification, the family Caliciviridae comprises four genera: Vesivirus, Lagovirus, Norwalk-like viruses (NLVs), and Sapporo-like viruses (14). In previous reports, viruses in the NLV genus have been categorized in two genogroups. Genogroup I includes the prototype Norwalk virus (NV) and related viruses, and genogroup II includes viruses such as Snow Mountain virus (SMV), Hawaii virus (HV), and Mexico virus (MXV) (23, 24, 26, 31, 36). NV has been the most extensively studied, although it currently does not seem responsible for most gastroenteritis caused by NLVs (4, 23, 26, 34, 37). The development of recombinant NV (rNV) has provided a highly purified antigen for detecting immunoglobulin G (IgG) antibodies against NV and genogroup I NLVs (11, 12, 15, 21, 22, 32, 33). More recently, recombinant capsid antigen has been developed from MXV, a genogroup II NLV (24, 25). This recombinant MXV (rMXV) antigen has been used in several studies to detect IgG antibodies against genogroup II NLVs (7, 17, 25, 33, 34).

Testing for seroconversion with either of these recombinant antigens requires an early acute-phase serum sample and a convalescent-phase serum sample to detect a minimum fourfold rise in IgG antibody levels required to diagnose infection. We recently described a monoclonal antibody, recombinant antigen-based IgM capture enzyme-linked immunosorbent assay (ELISA) for the detection of specific IgM antibodies to NV (1). We found that IgM antibodies to NV developed by 8 days after exposure and were not detectable in normal sera even if high titers of IgG antibodies were present. NV-specific IgM antibodies were not detected in sera from SMV- or HV-inoculated volunteers (1). Two studies have used immune electron microscopy to detect IgM to genogroup II NLVs (3, 30), and IgM responses to rMXV antigen in sera from persons infected with small round structured viruses have been reported (17). The purpose of this study was to determine the efficacy of a monoclonal antibody, rMXV-based IgM capture ELISA for the diagnosis of genogroup II NLV infections.
A solution was measured with a plate reader spectrophotometer. A positive test with the NV IgM test previously reported (1). In an outbreak of gastroenteritis not characterized with regard to viral etiology but suspected to be due to NV (the Rhode Island graduation banquet), 7 of 13 persons had IgM responses to rMXV, whereas none had IgM responses to rNV. These results established the diagnosis as genogroup II NLV. All four sera from patients in a documented MXV outbreak were positive in the rMXV IgM test and negative in the rNV IgM test.

Among four paired sera from children with astrovirus infections and among six paired sera from adults with rotavirus infections, none showed seroconversion or were IgM positive for either rMXV or rNV (Table 1). Eighty sera from noninfectious individuals ranging in age from 1 to 59 years were tested for IgG and IgM to rMXV (Table 2). The proportion of sera positive for rMXV IgG ranged from 45 to 90% for the different age groups. Two sera from children between 1 and 4 years of age were IgM positive for rMXV.

**RESULTS**

The results of the serological tests for MXV from volunteer studies and outbreaks are shown in Table 1. Among sera from those infected with genogroup I NLVs (combined volunteer and outbreak studies), only 3 of 25 were rMXV IgM positive; in contrast, 24 of 25 were IgM positive for rNV. In sera from those infected with genogroup II NLVs (combined volunteer and outbreak studies), 28 of 47 were rMXV IgM positive and none were IgM positive for rNV. These results show the specificity of the rMXV IgM test and confirm the specificity of the rNV IgM test previously reported (1). In an outbreak of gastroenteritis not characterized with regard to viral etiology but suspected to be due to NV (the Rhode Island graduation banquet), 7 of 13 persons had IgM responses to rMXV, whereas none had IgM responses to rNV. These results established the diagnosis as genogroup II NLV. All four sera from patients in a documented MXV outbreak were positive in the rMXV IgM test and negative in the rNV IgM test.

**DISCUSSION**

Several studies have shown the development of NV-specific IgM antibodies as a result of NV infection (1, 6, 9, 13, 29, 35), but reports on the development of IgM antibodies to genogroup II NLVs are limited. Two earlier studies (3, 30), with immune electron microscopy showed IgM reactivity to viruses that were later shown to be genogroup II NLVs, and a recent report with an ELISA showed IgM responses to genogroup II NLVs involved in outbreaks (17). Sera from genogroup I NLV infections were not tested in that study. Our results with volunteer and outbreak sera show that the rMXV-based IgM capture ELISA that we developed detects antibodies to genogroup II viruses such as HV, SMV, TNV, and related viruses. IgM antibodies to rMXV were not detected in sera from NV-inoculated volunteers. Neither the rMXV IgM test nor the rNV IgM test reacted with sera from patients with rotavirus or astrovirus infections, further demonstrating the specificity of these tests for NLV infections.

During NV infections, IgM to NV has been found to be more specific for NV than IgG to NV (1, 28, 34). It is well established that repeated stimulation by a given antigen usually increases IgG titers but decreases specificity. This situation should not occur with the IgM response, because IgM is not associated with anamnestic responses to repeated antigen exposure. Thus, exposure to several related NLVs could result in IgG with a broader specificity, as was seen in the seroconversion data from the Erie County outbreak involving a genogroup I NLV (V Ward 190). It has been shown in a previous study that there is a correlation between seroconversion and NLV genotype, but it was suggested in that report that the IgM response could be more specific (33).

The MXV IgM test was most useful when used in combination with the NV IgM test for the outbreaks at UMMC-UC, the Rhode Island graduation banquet, and in Erie County. The lack of fourfold or greater rises in IgG antibody titers to rMXV from patients in the outbreak at UMMC-UC was probably due
TABLE 1. Detection by ELISA of IgG seroconversion (sc) and IgM antibodies specific for rMXV and rNV in volunteers and in natural outbreaks of gastroenteritis

<table>
<thead>
<tr>
<th>No. of sera with the following result/no. tested:</th>
<th>Genogroup</th>
<th>Samples</th>
</tr>
</thead>
<tbody>
<tr>
<td>rMXV</td>
<td>rNV</td>
<td>rMXV</td>
</tr>
<tr>
<td>IgM* for:</td>
<td>IgG sc for:</td>
<td>IgM* and IgGsc for:</td>
</tr>
<tr>
<td>--------</td>
<td>--------</td>
<td>--------</td>
</tr>
<tr>
<td>I</td>
<td>NV-inoculated volunteers (paired sera)</td>
<td>0/9</td>
</tr>
<tr>
<td></td>
<td>Eric County outbreak (paired sera)</td>
<td>3/16</td>
</tr>
<tr>
<td>II</td>
<td>HV-inoculated volunteers (paired sera)</td>
<td>2/4</td>
</tr>
<tr>
<td></td>
<td>SMV-inoculated volunteers (convalescent-phase sera)</td>
<td>3/3</td>
</tr>
<tr>
<td></td>
<td>SMV outbreak (paired sera)</td>
<td>11/21</td>
</tr>
<tr>
<td></td>
<td>TNV outbreak (paired sera)</td>
<td>2/2</td>
</tr>
<tr>
<td></td>
<td>MXV outbreak (paired sera)b</td>
<td>4/4</td>
</tr>
<tr>
<td></td>
<td>UMMC-UC outbreak (paired sera)c</td>
<td>2/5</td>
</tr>
<tr>
<td></td>
<td>UMMC-UC outbreak (convalescent-phase sera)</td>
<td>4/8</td>
</tr>
<tr>
<td>Unknown</td>
<td>Rhode Island outbreak (paired sera)</td>
<td>7/13</td>
</tr>
<tr>
<td>Otherd</td>
<td>Rotavirus (paired sera)</td>
<td>0/6</td>
</tr>
<tr>
<td></td>
<td>Astrovirus (paired sera)</td>
<td>0/4</td>
</tr>
</tbody>
</table>

a A serum positive for IgM is one in which the IgM antibody titer is ≥1:25 in a single sample.
b The outbreak was confirmed to be due to MXV.
c Acute-phase sera were collected late.
d All sera from patients with rotavirus and astrovirus infections had IgG titers to rNV and rMXV in their acute- and convalescent-phase sera.

*NA, not applicable.

determine the time between exposure to NLVs and the appearance of specific IgM. In a previous study on the detection of IgM antibodies to genogroup I NLV infections (1), rNV-specific IgM was not detected in volunteers by 5 days but was detected by 8 days (sera from days 6 and 7 were not obtained). In outbreak sera, rNV-specific IgM was detected 6 to 7 days after the estimated time of exposure (1), and it is likely that similar times for the appearance of virus-specific IgM would be required for other NLVs.

rNV- and rMXV-based IgM capture assays can be used to determine whether an outbreak is due to NLVs and are useful for genogroup classification in epidemiological studies. IgM assays are especially useful if acute-phase sera are collected late or if paired sera are not available.

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REFERENCES


