Immunoglobulin M Antibody Test To Detect Genogroup II Norwalk-Like Virus Infection

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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 seroconversion with small round structured viruses have been reported (17). 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 immunoglobin M (IgM) 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 (19). 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.

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 NLVs, 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 1/90). 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-U]; 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 polymersase region with MD-V6, a genogroup II NLV involved in an outbreak in a Maryland nursing home in 1987 (28) (accession no. M9170613), and 98% identity with Halifax NLV (unpublished data) (accession no. N0137651). The second outbreak involved patients who developed gastroenteritis after a Rhode Island graduation banquet in 1986
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 this 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 anamnestic responses to related viruses. IgM antibodies to rMXV were not detected in sera from patients with rotavirus or astrovirus infections, further demonstrating the specificity of these tests for NLV infections.

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.

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 noninfected 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.
to the late collection of the acute-phase sera. In paired sera from two patients at the Rhode Island graduation banquet, there were fourfold increases in IgG antibody titers to rMXV in both patients and to rNV in one patient. Both outbreaks were diagnosed by use of a combination of rNV- and rMXV-based IgM capture assays. The larger sample size of the SMV outbreak associated with eating in a school cafeteria also demonstrates the utility of the rMXV-based IgM test compared with seroconversion. Of the 13 persons found positive by either seroconversion or IgM, only 2 were missed by the rMXV-based IgM test, whereas 9 were missed by seroconversion.

Among 80 normal sera tested, sera from two children between 1 and 4 years old were rMXV-specific IgM positive. Occasional positive results may occur in the rMXV-based IgM test because high levels of IgG antibodies to MXV have been detected in young children (5, 7, 24, 34) and the IgM-positive sera could represent a recent infection. We did not have sufficient information available on volunteer or outbreak sera to 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


