Detection of human Bocavirus 1, 2 and 3 from patients with acute gastroenteritis

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ABSTRACT

Aim: We aimed to investigate the prevalence of Human Bocavirus (HBoV) isolates among Iranian patients suffering from acute gastroenteritis.

Background: Human Bocavirus is a new parvovirus that has been identified in association with gastroenteritis. Limited data are available about HBoV in Iran.

Patients and methods: Viral DNA was extracted from all 294-stool samples. HBoV DNA was detected in extracted samples by polymerase chain reaction (PCR) amplification of a 354 bp of noncapsid protein 1 (NP1) gene. In addition, all samples were also subjected to a nested PCR to amplify a 455 bp of nonstructural 1 (NS1) gene.

Results: The main clinical symptoms among HBoV positive patients were diarrhea (77.7%), fever (62.9%), vomiting (55.5%), and anorexia (59.2%). NP1 PCR was positive in 8 samples (2.72%), NS1 was positive in 16 patients (5.44%) and 3 samples had positive results in both regions (1.02%).

Conclusion: Our results suggest that HBoV could be considered as one of the important etiologic agents of acute gastroenteritis cases in Iran.

Keywords: Human Bocavirus, Acute gastroenteritis, Iran.

Introduction

Viruses, bacteria and parasites can cause gastroenteritis. Molecular epidemiology of enteric viruses in patients with acute gastroenteritis was previously investigated in Iran. The results showed Norovirus, Adenovirus and Sapovirus are responsible for 4 to 9 percent of sporadic gastroenteritis cases (1-4). Human Bocavirus (HBoV) was firstly detected in 2005 (5). This virus was first discovered in children with respiratory tract illness but also has been detected in stool samples from children with gastroenteritis (5, 6). Recently, several genotypes of this parvovirus, HBoV genotype 2 (HBoV2), genotype 3 (HBoV3) and genotype 4 (HBoV4) were discovered that were closely related to HBoV (7, 8). HBoV2 was firstly
detected in stool samples from children with flaccid paralysis in Pakistan (9) and followed in other countries. HBoV3 was detected in Australia (10) and HBoV4 was identified in stool samples from Nigeria, Tunisia and United States (11). Previous reports are demonstrated that there is a close taxonomic association between HBoV and bovine parvovirus, that it causes gastrointestinal disorders in cattle (12, 13). Bocaviruses like other parvoviruses are predominantly resistant to heat and detergent inactivation and due to this fact they could be stable in stool samples for long times (14). Distribution of HBoV infections is different during various seasons, and the highest prevalence is related to cold seasons (15). Due to considered persistence of HBoVs to different environmental factors and detergent material (16, 17), it is possible that fecal-oral transmission, in addition to transmission via respiratory droplets, could be important in interpreting the previous observations (6). The aim of this study was to determine the prevalence of HBoV infection and its dominant genotypes among Iranian patients with acute gastroenteritis.

**Patients and Methods**

From May 2008 to June 2010, 294 stool samples were collected from patients referred to Taleghani Hospital, Shohadaye Tajrish hospital, Mofid children’s hospital and children clinical center in Tehran, Iran. Samples were obtained from all age groups comprised of 227 children (<18 years of age, 77.2%, 132 male, 95 female) and 67 adults (>18 years of age, 22.8%, 36 male, 31 female). Informed consent was taken from all children’s parents or adult patients and the study protocol was reviewed and approved by the ethics committee at the Gastroenterology and Liver Diseases Research center. DNA was extracted from stool samples using the QIAamp DNA Mini kit (QIAGEN, Hilden, Germany). HBoV DNA was detected in extracted samples by polymerase chain reaction (PCR) amplification of a 354 bp of noncapsid protein 1 (NP1) gene using primers 188F and 542R as reported previously (5). In addition, all samples were also subjected to a nested PCR as described by Kapoor et al, to amplify a 455 bp of nonstructural 1 (NS1) gene (3). The BigDye Terminator v3.1 cycle sequencing kit (Applied Biosystems, Foster City, CA, USA) was employed for sequencing. All sequences were analyzed bidirectionally. To compare the frequency clinical symptoms between the HBoV positive and negative patients we conducted chi-square tests. Quantitative characteristics of the study population such as age were compared using student’s t-test. All statistical analyzes were performed using SPSS version 20 software (IBM SPSS Statistics, Chicago, IL, USA).

**Results**

Overall, of the 294 patients tested, 27 samples (9.18%) were positive (13 male and 14 female). Higher rate of positive result was observed in children group (24 of 227, 10.57%) in comparison with adult group (3 of 67, 4.48%) but this difference was not statistically significant (p=0.129 by $\chi^2$ test). Higher rate of infection were observed during autumn and winter seasons (16 cases). The main clinical symptoms among HBoV positive patients were diarrhea (77.7%), fever (62.9%), vomiting (55.5%), and anorexia (59.2%). Four HBoV positive patients reported cough and rhinorrhea accompanied with gastroenteritis symptoms (14.8%). No statistically difference in the prevalence of clinical symptoms between HBoV positive and negative patients was observed. NP1 PCR was positive in 8 samples (2.72%), NS1 was positive in 16 patients (5.44%) and 3 samples had positive results in both regions (1.02%). NP1 PCR products of 10 samples and NS1 products of 17 samples were subjected to direct sequencing. Phylogenetic analysis of NS1 nucleotide sequences revealed that 3 patients infected with HBoV genotype 1 (HBoV), 13 patients infected with HBoV2 and one patient infected with HBoV3 (Figure 1).
Figure 1. Phylogenetic analysis of the partial nucleotide sequences of non structural protein 1 (NS1) from Iranian isolates in comparison with different human Bocavirus genotypes. Three HBoV sequences (labeled with black squares), thirteen HBoV2 sequences (indicated with black triangles) and one HBoV3 sequence (labeled with black circle) were analyzed with thirty tree HBoV sequences of genotypes 1 to 4 (HBoV to HBoV4) available from GenBank as reference genes. Nucleotide sequences were aligned using CLUSTAL X software (www.clustal.org) and compared with corresponding regions of reference sequences from the GenBank database, genetic distances were calculated by a Kimura two parameter algorithms and phylogenetic trees were constructed by neighbor joining (NJ) method. These analyses were performed using the Molecular Evolutionary Genetics Analysis (MEGA) program, version 3.1 (www.megasoftware.net). The reliability of phylogenetic trees was confirmed by the bootstrap-resampling test (n=1000). Bootstrap values >70% are presented at the branching points.

Discussion

Human Bocavirus (HBoV) is considered as major agent of several respiratory tract diseases. However, there are several reports that showed a relationship between HBoV and gastrointestinal disorders. To clear this relationship and due to the fact that the prevalence and molecular characteristics of HBoV in Iranian patients with acute gastroenteritis has not been well studied, we carried out a molecular detection and phylogenetic study of HBoV in Iran.

The present study identified a 9.18% HBoV DNA detection rate in children and adults with acute diarrhea in Tehran, Iran, during May 2009 to June 2010. According to the results of present study, Human Bocavirus (HBoV) was detected in 27 (9.18%) of 294 patients with acute gastroenteritis. NP1 and NS1 nucleotide sequences were highly conserved and no differences in nucleotide sequences of NP1 gene (354 bp) were observed between Iranian isolates, except one isolate. Nucleotide identity of Iranian HBoV NS1 sequences (435 bp) with other HBoV reference sequences (accession numbers were indicated in the figure 1) was 99%-100%. On the other hand, HBoV2 NS1 sequences share 97.4%-98.2% nucleotide sequence identity and 98.5%-99.2% deduced amino acid sequences identity with other HBoV2 reference isolates and 79.6%-81.5% nucleotide identity and 87.2%-91.8% deduced amino acid sequence identity with HBoV prototypes. Only one isolate was classified as HBoV3 based on phylogenetic analysis of NS1 region and this isolate showed 99.6%, 90.2% and 96.5% and 90.2% nucleotide sequence identity and 100%, 96.5% and 89.5% deduced amino acid sequence identity with HBoV3, HBoV and HBoV2 reference sequences, respectively.
than children group (67 versus 227 patients). The number of HBoV positive patients in the adult group was also less than children (4.48% vs. 10.57%). This may be due to fewer adults referral to hospitals or clinical centers especially when they don’t face with severe symptoms of gastroenteritis. Therefore, the study may underestimate the rate of HBoV infection among Iranian adult patients with acute gastroenteritis and it is necessary to perform further studies with larger adult populations to determine more accurate rate of HBoV infection among them.

Previous reports of HBoV prevalence around the world showed a variation from 1.5% to 19% (8, 18-21). In Iran, there were only two reports about HBoV infection. Naghipour et al. studied on 261 Iranian children with acute respiratory illnesses under 5 years of age in Rasht (a city located in north of Iran) during the winter of 2003 to 2004 and found HBoV positivity in 8% of studied children (22). Another study with lower sample size was performed by Nadji and his colleagues on 50 exacerbated asthma cases, 83 cases with acute respiratory illnesses and 47 patients with acute gastroenteritis and they reported 6%, 7.2% and 12.8% HBoV positivity in the studied groups, respectively (23).

We found some data from prevalence of other enteric viruses among Iranian patients with gastroenteritis. Nadji et al. in 2010 performed a small scale study on 47 stool samples of patients with gastroenteritis and reported that Norovirus and Sapovirus are found in 21.3% and 2.1% of samples (24). Our previous work has demonstrated that two genotypes (GI and GII) of Norovirus are involved in 9.8% of acute gastroenteritis cases (4). In another study among Iranian patients with acute gastroenteritis, we demonstrated that Sapoviruses could be responsible for 11.9% of acute gastroenteritis cases (2).

Higher rate of HBoV infection was detected in cold seasons in comparison with warm seasons of the year but this difference was not statistically significant. Although, Susanna et al. reported the seasonal epidemiological profile of HBoV with the highest incidence in fall and winter (25), but some other studies couldn’t find any obvious seasonal pattern (18, 23). The difference between results could be due to different sample collection periods and the age groups studied. The most common clinical symptoms of the disease in HBoV positive patients were diarrhea, fever, vomiting and anorexia. No significant differences in the rate of gastroenteritis symptoms were found between HBoV positive or negative patients, as well as between HBoV positive children and adults groups. HBoV nucleotide sequences of NP1 and NS1 genes are highly conserved and very low nucleotide or deduced amino acid sequence variability was observed between Iranian isolates, especially within HBoV1. Previous studies also reported low nucleotide sequence variability among HBoV1 isolates from different parts of the world. Kapoor et al. proposed that HBoV1 evolved more recently than other HBoV genotypes (11). It seems that different genotypes of HBoV are widespread in the world.

Our results suggest that different HBoV genotypes are circulating among Iranian patients and they must be considered as one of the etiologic agents of acute gastroenteritis cases in Iran. To confirm pathogenicity of HBoV in acute gastroenteritis further studies on mucosa biopsy samples from patients with disease, are suggested.

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References


