Torque Teno Midi Virus/Small Anellovirus in Sera of Healthy, HIV/HCV and HIV Infected Individuals in Lorestan Province, Iran

Maryam Fatholahi¹ and Majid Bouzari¹,*

¹Department of Biology, Faculty of Sciences, University of Isfahan, Isfahan, IR Iran
*Corresponding author: Majid Bouzari, Department of Biology, Faculty of Sciences, University of Isfahan, P. O. Box: 8174673441, Isfahan, IR Iran. Tel: +98-3117932459, Fax: +98-3117932456, E-mail: bouzari@sci.ui.ac.ir

Received 2014 November 16; Revised 2015 February 8; Accepted 2015 March 17.

Abstract
Background: Torque Teno Midi Virus/Small Anellovirus (TTMDV/SAV) is a member of the Gammatorquevirus genus within the family Anelloviridae. It is detected in healthy, Hepatitis B Virus, Hepatitis C Virus and HIV infected individuals and also patients with acute respiratory disease in different countries, but its role in clinical diseases and its full geographical distribution is still unclear.

Objectives: The current study aimed to detect the frequency of infection with TTMDV/SAV in the sera of healthy blood donors, hepatitis C virus infected and HIV positive individuals in Lorestan Province, Iran; and also investigate the possible role of TTMDV/SAV virus in liver diseases.

Materials and Methods: Fifty two, 36, 4, and 110 serum samples from HIV positive, patients with HIV/HCV and HCV/HBV co-infections, and healthy individuals were collected in Khorramabad city, respectively. Nested-polymerase chain reaction was performed using SMa/SMAr primers to detect TTMDV/SAV DNA. Serum aminotransferases were measured.

Results: In the HIV/HCV, HIV/HCV/HBV, HIV, and control cases, 29 (80.5%), 3 (75%), 43 (82.7%), and 16 (14.5%) were positive for DNA of TTMDV/SAV, respectively. In the HIV/HCV infected cases and HIV positive cases the level of Alanine aminotransferase (ALT) and Aspartate aminotransferase (AST) were not significantly different in TTMDV/SAV infected and non-infected individuals (P > 0.05).

Conclusions: Although significant differences (P < 0.01) were observed in the frequency of TTMDV/SAV between healthy controls and each of the HIV positive and HIV/HCV co-infected individuals, no significant difference was observed between HIV positive and HIV/HCV co-infected cases, which may be due to HIV associated immunodeficiency. This is the first time that TTMDV/SAV is reported in HIV infected individuals worldwide. Interpretation of the high frequency of the virus (82.7%) in HIV cases needs more detailed studies.

Keywords: HCV, TTMDV/SAV, HIV

1. Background

The family Anelloviridae contains eleven genera of human and animal viruses that have a single-stranded, circular, negative sense DNA genome (1, 2). In 2007, Ninomyia et al. identified a novel DNA virus with a genomic organization resembling those of TTV and TTMV and designated this new virus as Torque Teno Midi Virus (TTMDV), they also showed that Small Anellovirus (SAV) and SAV2 were deletion mutants or artifacts generated during the amplification of TTMV (3). Therefore, these viruses were named TTMDV/SAV and then TTMDV viruses and were classified as Gammatorquevirus genus (including 15 species: Torque teno midi virus 1 - 15) of the family (1-4).

TTMDV/SAV DNA was found in 20% of plasma samples of French blood donors, which 33% of their saliva samples were positive too (5). TTMDV/SAV DNA was detected in 9.1% of serum samples of patients with hepatitis C infection and 8.6% of healthy individuals in Italy (6). In Korea, TTMDV/SAV was found in 34% of blood products and 40% of patients with chronic hepatitis (7). Also in another study, TTMDV/SAV was detected in the serum of healthy children, children with acute respiratory disease (ARD), hepatitis B and hepatitis C, and nasopharyngeal aspirates collected from children with ARD (8). TTMDV/SAV is reported, in 46.7% of nasopharyngeal aspirates collected from children with acute respiratory disease in Hungary (1) and in 50.4% of different cervical cancers of the female Iranian population (9). The role of TTMDV/SAV in clinical diseases and the full geographical distribution of the virus is unclear (1).

High levels of TTV DNA and TTMV DNA are reported in the bone marrow and spleen tissues of HIV positive individuals and HIV positive individuals who had progressed to acquired immune deficiency syndrome (AIDS) (10). TTV and TTMV loads increase in immune suppression and reduce the number of CD4⁺ T cells (10, 11). Viral hepatitis is common and considered as a serious public health problem (12, 13). Also, it is the cause of chronic hepatitis, cirrhosis, and hepatocellular carcinoma worldwide (14-16). As these viruses share the same transmission routes such as blood transfusion, co-infection of TTMDV/SAV and Hepatitis B Virus (HBV), or TTMDV/SAV; Hepatitis C Virus (HCV) is expected. However, the effect of TTMDV/SAV infection in patients with chronic HBV or HCV infection is unknown.

Copyright © 2015, Ahvaz Jundishapur University of Medical Sciences. This is an open-access article distributed under the terms of the Creative Commons Attribution-NonCommercial 4.0 International License (http://creativecommons.org/licenses/by-nc/4.0/) which permits copy and redistribute the material just in noncommercial usages, provided the original work is properly cited.
2. Objectives
Considering the possible role of other members of An-
elloviridae as etiological factors in hepatitis cases (17, 18) and that TTMDV/SAV virus is transmitted through blood products and can infect humans (8), the current study aimed to detect the frequency of TTMDV/SAV in the sera of HIV/HCV, HIV infected, and healthy individuals and their possible correlation, and also the possible role of TTMDV/SAV in the liver diseases was investigated.

3. Materials and Methods

3.1. Sampling and Serological Tests
Fifty-two, thirty-six, four, and one hundred-ten serum samples from HIV positive, patients with HIV/HCV and HIV/ HCV/HBV co-infected cases regardless of the presence of clinical signs or not, and healthy individuals (not positive for HIV, HBV and HCV, and also normal routine blood tests such as counting blood cells (CBC) and liver enzymes) were collected from pathobiology laboratories in Khorramabad from December 2010 to September 2011, respectively. Patients' characteristics, gender, and age were recorded. All samples were stored at -20°C until tested. The study was approved by the local University of Isfahan ethics committee. Different enzyme linked immunosorbent assay (ELISA) tests were used to detect anti-HIV antibody (Delaware Biotech, USA), HBs antigen (Delaware Biotech, USA) and anti-HCV antibody (DIA. PRO, Italy). Serum aminotransferases (AST and ALT) were evaluated by Biology Chemistry kit (Iran).

3.2. Nucleic Acid Extraction From Serum
For each sample 220 μL of serum was mixed with 6.5 μL of 0.25% Sodium Dodecyl Sulfate (SDS) and 10 μL of 0.2 M NaCl. Then, 12 μL of 10 mg/mL proteinase K (Roche, Germany) was added and thermo-mixed at 350 rpm for 2 hours at 65°C. Proteins were precipitated using phenol and chloroform. The supernatant phase containing DNA was collected and transferred to a microtube and 3 M sodium acetate was added (0.1 volume). Then, the samples were vortexed for 30 seconds. DNA was precipitated with the cold ethanol (100%) (Merck, Germany). The pellet was dried and dissolved in 50 μL of distilled deionized water or TE (Tris-HCL buffer). To determine the purity of the extracted DNA, absorbance of extracted DNA was read at wavelengths of 260 and 280 nm by the spectrophotometer.

3.3. Detection of TTMDV/SAV DNA by Nested-PCR
The SMAs1/SMa1 and SMAs2/SMa2 primer sets (5, 7) were used to perform nested Polymerase Chain Reaction according to Salmanizadeh et al. (9). The sequences of TTMDV/SAV virus specific primers were: SMAs1: GGCAGGGAGCCCCGAGGTGAGTGAC (sense), SMAr1: TTCGAAATAGTGCTGTTGCCATTTG (antisense), SMAs2: ACCGAGGTCTAGGGGCAATTCCG GCC (sense), and SMAr2: CGCGCCGTCTTTTCTGC CGCGCCGCA (antisense). A positive control was used, which was already sequenced with accession number of HQ259681 in the GenBank (9).

3.4. DNA Sequencing
PCR products of seven randomly selected samples from patients with HIV/HCV and HIV/HCV/HBV co-infections, and healthy blood donors were subjected to agarose gel electrophoresis (1%) and DNA was extracted according to the instructions of the DNA gel extraction kit K0513 (Fermentas, Germany). The DNAs were sequenced by Takapouzist Company, Iran. BLAST Nucleotide search of the determined sequences against the nucleotide sequence database (NCBI, national center for biotechnology information) was performed.

3.5. Molecular Evolutionary Analysis
Phylogenetic trees were constructed using neighboring method in MEGA4 (Molecular Evolutionary Genetics Analysis software version 4.0) based on the current study sequenced amplicons against sequences obtained from the GenBank (19).

3.6. Statistical Analyses
The Fisher exact test and paired t-test were employed for statistical analyses using the GraphPad Instat software version 3.05 (GraphPad software, USA). P value < 0.05 was considered statistically significant.

4. Results
In the gel electrophoresis expected 431 base pair product for SAV2 and 441 bp product for SAV1 (4, 8) were observed (Figure 1). Alignment of the seven obtained sequences (GenBank accession numbers: KC352905, KC352906, KC352907, KC352908, KC352909, KC352910, KC352911) revealed a high degree of genetic variability among the TTMDV/SAV DNA (homology ranging from 70% to 93%) with some sequences in the GenBank. As indicated in Figure 2, three of the sequences related to patients with HIV/HCV co-infection (KC352906, KC352908, and KC352910) were located in a separate cluster and had less similarity with other reported cases. Although the other four sequences (KC352905, KC352907, KC352909, and KC352911) showed differences, they were generally more similar to the other reported sequences. Also, as shown in Figure 3, there was high degree of genomic homology of the sequences with some SAV1 sequences (sequences with accession numbers DQ987836 and DQ409192) and SAV2 sequences (sequences with accession numbers DQ987853, DQ987851, and DQ987854) reported by Chung et al. (20).

Features and information of serum samples from blood donors are shown in Table 1. In the current study, healthy blood donors were used as the control group.
The frequency of TTMDV/SAV in different serum samples are shown in Table 2. The differences of the TTMDV/SAV frequencies in the control cases versus HIV positive and HIV/HCV co-infected cases were statistically significant ($P < 0.01$). The difference between HIV positive and HIV/HCV co-infected cases was not significant ($P > 0.05$).

There were not any significant differences in the means of the serum aminotransferases (ALT and AST) levels between TTMDV/SAV-positive and TTMDV/SAV-negative individuals in different groups ($P > 0.05$). Due to the low number of HBV infected samples, comparison was not possible (Table 3). Also in each age group the level of ALT and AST were not significantly different between the patients with and without TTMDV/SAV infection ($P > 0.05$). Comparisons of different age groups between the patients with and without TTMDV/SAV infection in healthy and HIV positive and HIV/HCV co-infected patients are shown in tables 4 and 5, respectively. About 46.5% of HIV positive patients and 55% of those with HIV/HCV infection, co-infected with TTMDV/SAV, were in the age group of 31 - 40 years. The highest prevalence of healthy blood donors was 56% which was in the age group of 20 - 30 years. Prevalence of TTMDV/SAV was not significantly different between males and females, in the healthy blood donors and the HIV positive groups ($P > 0.05$).
Figure 3. Phylogenetic Tree Constructed by Neighbor-Joining Method Implemented in MEGA 4.1 With 1000 Bootstrap Replicates to Compare the Current Study Sequences With Three Main Genotype Groups Reported From Korea (20)

The evolutionary distances were computed using the maximum composite likelihood model based on the substitutions per site. GenBank accession number: Small anellovirus 1 (SAV1 gp1 + SAV1 gp2 + SAV2: NC-007013), Small anellovirus 2 (SAV2 ORF1 + ORF2 + ORF3 + ORF4 + …: AY622909), SAV DF-VIPI (S. anellovirus DF-VIPI: DQ409191), SAV BEC-VIPI (S. anellovirus BEC-VIPI: DQ409192), SAV GAR-VIPI (S. anellovirus GAR-VIPI: DQ409196), SAV TER-VIPI (S. anellovirus TER-VIPI: DQ409199), SAV KR-BD (Korean blood donors: DQ987831-DQ987854), Torque teno mini virus (T. teno mini virus 8 PB4TL: AB038631), and our sequences Torque teno midi virus/small anellovirus Lorestan MFa1 (patients with HIV/HCV/HBV co-infection: KC352905), Torque teno midi virus/small anellovirus Lorestan MFa2, MFa3 (healthy individual: KC352907), Torque teno midi virus/small anellovirus Lorestan MFa4, MFa5, MFa6, MFa7 (patients with HIV/HCV infection: RC352906, RC352908-RC352911).

Table 1. Serum Samples Information

<table>
<thead>
<tr>
<th>Sample</th>
<th>Number</th>
<th>Gender (Male/Female)</th>
<th>Mean Age</th>
</tr>
</thead>
<tbody>
<tr>
<td>Healthy blood donors</td>
<td>110</td>
<td>Male/Female</td>
<td>25.8</td>
</tr>
<tr>
<td>HIV positive</td>
<td>52</td>
<td>26/26</td>
<td>38.25</td>
</tr>
<tr>
<td>HIV/HCV co-infected</td>
<td>36</td>
<td>35/1</td>
<td>38</td>
</tr>
<tr>
<td>HIV/HCV/HBV co-infected</td>
<td>4</td>
<td>4/0</td>
<td>42.75</td>
</tr>
</tbody>
</table>
Table 2. Frequency of Torque teno midi virus in the Serum Samples

<table>
<thead>
<tr>
<th>Sample</th>
<th>TTMDV/SAV Positive</th>
<th>TTMDV/SAV Negative</th>
</tr>
</thead>
<tbody>
<tr>
<td>Healthy</td>
<td>16 (14.5)</td>
<td>94 (85.5)</td>
</tr>
<tr>
<td>HIV positive</td>
<td>43 (82.7)</td>
<td>9 (17.3)</td>
</tr>
<tr>
<td>HIV/HCV co-infected</td>
<td>29 (80.5)</td>
<td>7 (19.5)</td>
</tr>
<tr>
<td>HIV/HCV/HBV co-infected</td>
<td>3 (75)</td>
<td>1 (25)</td>
</tr>
<tr>
<td>Total</td>
<td>91 (45)</td>
<td>111 (55)</td>
</tr>
</tbody>
</table>

aData are presented as No. (%).

5. Discussion

The current study results revealed a high prevalence of the virus in the population of the Lorestan province. The high frequency of the virus in the cases with hepatitis was much higher than the cases reported elsewhere. In the present study, high frequency of TTMDV/SAV in HIV positive individuals and those with HIV/HCV co-infection shows the possibility of association between TTMDV/SAV and blood-borne infections such as HIV and hepatitis C, and also indicates the main route of blood transfusion of the virus rather than via saliva, infected peripheral blood
mononuclear cells (5), the respiratory tract (8), nasopharyngeal discharge, urine, stool (1), and sexual transmission (9) already reported. According to Garcia-Alvarez et al. TTV and TTMV may cause liver diseases in patients co-infected with HIV and HCV (18). High frequencies of TTMDV/SAV among patients with HIV/HCV co-infection in comparison with low frequencies in healthy ones suggest that the virus is highly compatible with blood-borne viruses. This may lead to the greater stability of the virus in the patients with HIV/HCV co-infection.

This is the first time that TTMDV/SAVs are detected in HIV positive individuals in Iran and worldwide. Therefore it was not possible to compare the obtained results with any similar data. Although significant differences (P < 0.01) in frequency of TTMDV/SAV were observed between the healthy controls and each of the patients with HIV and HIV/HCV co-infections, no significant difference was observed between the cases with HIV and HIV/HCV co-infections. Since there was no information about the stage of clinical disease, its correlation with immunodeficiency caused by HIV remained unclear. The frequency of the virus in healthy individuals was 14.5% that was lower than frequencies reported in healthy individuals in Korea (34.5%) (8) and France (20%) (5), and higher than the reported frequency of 8.6% in Italian blood donors (6).

In the current study, the prevalence of the virus in patients with HIV and HIV/HCV co-infections was much higher than the prevalence of the virus in various diseases from other countries including acute respiratory diseases (46.7%) (1), Kawasaki syndrome (50%) (8), and cervicitis (50%) (9). Further studies are needed to determine the reasons for the high prevalence of TTMDV/SAV in the current study compared with those of the other ones, which may be due to the difference in the samples tested or the presence of some factors in the serum samples, which supports the presence and replication of the virus (possibly synergism in co-infection with two or three viruses and also the sensitivity of the assays which should be investigated).

The prevalence of the virus in the HIV positive and patients with HIV/HCV co-infections was 86% and 75% in the age group of 20 - 40 years, respectively, indicating that the individuals are most at risk of co-infection with TTMDV/SAV in this age group. It can be probably related to the same transmission route. On the other hand, co-infection with HIV/HCV may reduce the immune response and clearance of TTMDV/SAV from the body. There were not any significant differences in the means of ALT and AST levels between the patients with and without TTMDV/SAV infection in the studied groups; therefore, no association was expected between viral infection and altered levels of these enzymes. It is noteworthy that the amount of AST and ALT was higher than normal in fifty-two HIV positive individuals. Whether the virus (TTMDV/SAV) was involved in the raise of the levels of these enzymes in these cases or not, remains unknown and further investigations are required.

As shown in Figure 2, high genomic homology was observed between the obtained sequences and some of the TTMDV/SAV isolates in Korea (8). The sequences of HIV/HCV (MFa5, MFa7) and HBV/HCV/HIV (MFa5) in the co-infected patients were similar to the isolates of HCV (KR-HC-0005 ORF1, KR-HC-0023 ORF1) and HBV (KR-HB-0140 ORF1) in the co-infected individuals in Korea, also the sequence of healthy individual (MFa3) was similar to the isolate of healthy individual in Korea (SAV-KR-CT-0017 ORF1) and isolate of a cervicitis case in Iran (HQ259681 = SAV Isfahan Sal F1). Phylogenetic analysis of SAV in Korea proposed the three main groups of genotypes including G1, G2, and G3. The most prevalent genotype was G1, whose members showed high homology to SAV1 and Genotype 2 included SAV2 (20).

Comparison of the obtained sequences with the ones reported by Chung et al. (20) (MFa1, MFa2, MFa4, MFa5, and MFa6 sequences) indicated that they were placed in the same cluster with group one viruses (like SAV KR-BD-0105 Orf1 and SAV KR-BD-0078 Orf1) and M Fa3 and M Fa7 sequences were placed in the same cluster with group two viruses (like SAV KR-BD-0265 Orf1 and SAV KR-BD-0277 Orf1) (Figure 3). It seems that the SAV-like viruses are more prevalent in the sampled area and consistent with the genotype distribution of SAV in Korea. Since there were only a few individuals with HIV/HCV/HBV co-infection in the current study, interpretation of their role was not possible. According to the findings of the study, the studied region is at risk of TTMDV/SAV virus. Since TTMDV/SAV was discovered a few years ago, few studies have been done and there is a poor understanding of its molecular properties and pathogenic potential.

Footnotes

Authors’ Contribution: Majid Bouzari: developing the original idea and the protocol, analyzed, and interpreted the data; Maryam Fatholahi: performing lab works and preparing manuscript draft.

Funding/Support: The current study was granted by the research and postgraduate office of the University of Isfahan.

References


