Molecular diagnosis and genetic relationship of foot and mouth disease virus serotype Asia1/Basne/Sul/2015

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Abstract

Foot and Mouth Disease (FMD) is the most economically important viral-induced livestock disease worldwide. From April to May of 2015, tongue epithelial tissue samples were collected from 36 cattle in six villages, which share the border with Iran. Samples were screened using RT-PCR to amplify a conserved region in the VP1 gene, and phylogenetic tree analysis was performed based on the VP1 nucleotide sequence results. Furthermore, the nucleotide sequence was converted to an amino acid sequence in order to detect similarities between the studied samples and those previously published in GenBank (NCBI). Epidemically, based on the amino acid residues, genetic similarity, and amino acid substitutions, the VP1 nucleotide sequences were determined to be close to a novel group, group VII, with 94% identity. The VP1 amino acid sequence analysis revealed a close relationship to the Asia/BAL/PAK/iso-2/2011 isolate (Accession no. JX435109), with 95.7% identity. Analysis of the studied samples revealed that the FMDV serotype Asia1 causing the outbreak in the Basne district belonged to group VII, which was introduced from the Balochistan province of Pakistan through illegal movement of animals from this region.

Keywords: FMD virus, serotype Asia1, Genetic relationship, VP1

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Introduction

Foot and Mouth Disease (FMD) is an acute, highly contagious transboundary viral disease of cloven-hoofed domestic animals (1). The FMD virus (FMDV) belongs to the family Picornaviridae and the genus Aphthovirus. The infection is characterized by fever, lameness, and vesicular lesions of the feet, tongue, snout, and teats. In addition to high mortality rates, the disease has a serious impact on food security, rural income, and therefore imposes significant economic consequences on any country harboring the virus (2,3). The virus is non-enveloped and contains a positive-sense single-stranded RNA genome, approximately 8.3 kilobases in length. The RNA is surrounded by 60 copies of four structural proteins [VP1 (1D), VP2 (1B), VP3 (1C), and VP4 (1A)], which form a naked icosahedral capsid. The genome has three components, including the 5′ untranslated region (5′ UTR), a long open reading frame (ORF), which is the coding region, and the 3′ untranslated region (3′ UTR). The 5′ UTR is approximately 1300 nucleotides long and consists of a 360-bp S fragment at its 5′ end; a 150 to 250-bp poly(c) tract; a series of RNA pseudoknot structures; a cis-acting replication element (CRE), which is 55 nucleotides and contains a conserved motif (AAACA); and a 450-bp internal ribosome entry site (IRES), which is responsible for cap-independent initiation of viral protein synthesis (4). The virus also contains a single, long ORF encoding a polyprotein, which may generate precursors useful during the polyprotein processing. A series of precursor cleavages produces more than 14 mature proteins including the nonstructural proteins L, 2A, 2B, 2C, 3A, 3B1, 3B2, 3B3, 3C, and 3D and the structural proteins VP1, VP2, VP3, and VP4.

Among the structural proteins, VP1, which contains the highly conserved RGD motif, is exposed on the capsid surface and plays an essential role in forming the virus particles and is considered to be highly immunogenic. Hence, the genetic similarity of the FMDV strains and the transmission between susceptible livestock populations were studied using VP1 coding sequences (5-7). Seven distinct serotypes of FMDV have been recognized; these include the Eurasian serotypes A, O, C, and Asia1 and the South African territory serotypes SAT1, SAT2, and SAT3. SAT1, SAT2, and SAT3 are usually present in Africa, Asia1 is restricted to Asia, and serotypes O, A, and C are found in Africa, Asia, South America, and occasionally, Europe (5).

The Asia1 FMDVs belonging to group VII (Sindh-08) and currently dominant in west Eurasia and were detected for the first time in the Sindh province of Pakistan in 2008, and later detected in Iran, Afghanistan, and Turkey (8).

FMDV affects over 100 countries around the world. Sometimes incursions occur from endemic regions into countries that are normally disease-free, and this has an enormous economic impact (9). However, epidemiological studies to identify the source of the FMD outbreaks are an essential part of disease control and provide information to help take effective measures against reintroduction (10). To date, Iraq and Kurdistan governments have been unsuccessful in controlling transmission and eradication of FMDV. Geographically the villages of the Basne district share a border with Iran. Thus, this is one of the key places for illegal introduction of animals from Iran, Pakistan, and Afghanistan into Iraqi-Kurdistan regions, facilitating FMDV transmission.

This study, performed for the first time, focuses on the nucleotide sequence of VP1 and amino acid substitutions of FMD/Asia1 and uses the data to compare with FMDV sequences that were previously published in GenBank. This analysis led to the discovery of the source of the FMDV serotype Asia1, which is circulating among cattle in different villages in the Basne district.

Material and methods

Study area and sample collection

This research study was conducted from April 2, 2015, to May 26, 2015. Tongue epithelial tissue samples were collected from 36 unvaccinated cows, which clinically had blisters or vesicles on or in the nose, tongue, lips, teats, oral cavity, between the toes, and above the hooves. Lameness and reluctance to move were also found. Moreover, fever, depression, hypersalivation, loss of appetite and weight, and drop in milk production were also presented.

Basne district include 22 Villages, which are located in the governorate of Sulaimani in the Kurdistan region north of Iraq, which shares a border with Iran. The cattle in the Basne, Bewre, Marwe, Shiwakal, Boskan and Dere villages were suspected of being infected with FMD. Collected samples were stored on dry ice to maintain virus activity and shipped to the Molecular Diagnostic Laboratory of Sulaimani Veterinary Directorate. Annually, there are two FMD outbreaks in the Basne district, first in the spring and later in fall.

Sample preparation and RNA extraction

Epithelial tissue samples (25 mg) were transferred into a 1.5 ml microcentrifuge tube containing 400 μl of RB buffer and 4 μl of β-mercaptoethanol. A micropipette was used to
employing a universal primer (IF, IR) using the AccuPower PCR was performed in 0.2 ml tubes to detect the FMDV by Uniplex PCR according to the references. All of the primers were synthesized by Bioneer, Asia1 serotypes by amplifying a 911-bp cDNA fragment and the universal reverse primer, NK61, were used to detect 1C505F primer (5'–TACACTGCTTCTGACGTGGC-3’). As1-3’ were designed to detect serotype A by amplifying a 613-bp cDNA fragment within the VP1 sequence (12). (NK61: 5’–GACATGTCCTCCTGCATCTG-3’)) was designed for specific diagnosis of serotype O, by amplifying a cDNA fragment of approximately 290 bp, within the VP1 untranslated region (5’ UTR) of any FMDV serotype (11). The second, more specific primer set amplifies a 328-bp cDNA fragment within the coding sequences of VP1, a conserved region of the FMDV genome. Another primer pairs set ((P38F 5’-GCTGCCTACCTCCTTCAA-3’) and (NK61R 5’–GACATGTCCTCCTGCATCTG-3’)) was designed for detecting FMD: (IF 5’–GACATGTCCTCCTGCATCTG-3’). The above primer set was universal primers that were synthesized by Bioneer  and used for general FMDV detection and serotype detection in outbreak specimens. The first primer set was universal primers that detect FMD: (IF 5’–GACATGTCCTCCTGCATCTG-3’), and (IR 5’–CCAGTCCCCCTTCTCAGATC-3’). The above primer set amplifies a 328-bp cDNA fragment within the untranslated region (5’ UTR) of any FMDV serotype (11).

Reverse transcription of extracted RNA to cDNA
First strand complementary DNA (cDNA) was synthesized from all extracted RNA samples using the AccuPower RT Premix (Bioneer, Korea), according to manufacturer’s instructions. Five microliters of extracted RNA, 1 μl of IF and IR primers (see below), and 13 μl of RNase-free water were added to 0.2 ml tubes and incubated at 70°C for 5 min in a thermal cycler to remove any secondary structure and then incubated at 4°C for 5 min. The mixture was added to lyophilized RT premix and incubated at 42°C for 60 min, according to the manufacturer’s instructions. The temperature was then elevated to 95°C for 5 minutes to inactivate the reverse transcriptase.

Oligonucleotide primers
Two sets of primers, which amplify conserved regions of the virus, were synthesized by Bioneer and used for general FMDV detection and serotype detection in outbreak specimens. The first primer set was universal primers that detect FMD: (IF 5’-GACATGTCCTCCTGCATCTG-3’) and (IR 5’-CCAGTCCCCCTTCTCAGATC-3’). The above primer set amplifies a 328-bp cDNA fragment within the untranslated region (5’ UTR) of any FMDV serotype (11). The second, more specific primer set amplifies a cDNA fragment within the coding sequences of VP1, a conserved region of the FMDV genome. Another primer pairs set ((P38F 5’-GCTGCCTACCTCCTTCAA-3’) and (NK61R 5’–GACATGTCCTCCTGCATCTG-3’)) was designed for specific diagnosis of serotype O, by amplifying a cDNA fragment of approximately 290 bp, within the VP1 sequence (12).

Primers (p87: 5’-GTCATTGACCTCATGCAGAC(C/T) CAC-3’) and (NK61: 5’-GACATGTCCTCCTGCATCTG-3’) were designed to detect serotype A by amplifying a 613-bp cDNA fragment within the VP1 sequence (12). As1-1C505F primer (5’-TACACTGCTTCTGACGTGGC-3’) and the universal reverse primer, NK61, were used to detect Asia1 serotypes by amplifying a 911-bp cDNA fragment (13). All of the primers were synthesized by Bioneer, according to the references.

Uniplex PCR
Following the first strand cDNA synthesis, the uniplex PCR was performed in 0.2 ml tubes to detect the FMDV by employing a universal primer (IF, IR) using the AccuPower PCR Premix (Bioneer). Five microliters of cDNA template, 1 μl of each universal primer (IF, IR), and 13 μl of DEPC-H2O were added to the lyophilized master mix. After initial confirmation of the FMDV, the uniplex RT-PCR reaction was performed to detect three virus serotypes (O/A/Asia1). During this study, 1 μl of specific forward primer and 1 μl of universal primer, NK61, were used. The PCR reaction mixture for both steps was subjected to one cycle at 94°C for 2 minutes, followed by 35 cycles of denaturation at 94°C for 30 s, annealing at 55°C for 30 s, and extension at 72°C for 5 min. A final extension step of one cycle at 72°C for 5 min was performed.

Gel analysis of RT-PCR amplification products
Eight microliters of the RT-PCR cDNA amplicon was analyzed by mixing it with 2 μl of loading dye in a 0.2 ml microfuge tube and then loading on a 1% agarose gel. A 100-bp DNA ladder was loaded on the agarose gel to identify the PCR product size. The agarose gel was run in 1X TBE buffer (89 mM Boric Acid, 2 mM EDTA, pH 8.3) at 90 V for 60 min. The results were visualized under UV light and were recorded by a gel documentation system.

Sequencing the PCR products
In order to confirm the PCR results, 25 μl of PCR product of one sample positive with the NK72 and As1-1C505F primers were sent for Sanger sequencing (Macrogen, Seoul, South Korea). The result was published in GenBank (NCBI) as Asia1/Basne/Sul/2015, with accession number KT861787. The name is based on the serotype, district; three letter provinces code (Sul), and the sampling year.

Computer analysis of genetic relationships
The partial VP1 nucleotide sequence of the FMD Asia1/Basne/Sul/2015 isolate (Accession no. KT861787) was aligned and compared with FMD/Asia1 from other countries, which were previously published in the GenBank database (Fig. 1). The ClustalW algorithm in BioEdit software version 7.2.5 was used and sequences were edited to remove unreliable sequences or regions (14). Phylogenetic tree analysis was performed based on the nucleotide sequences (Fig. 2) using MEGA 6.0.6 and the unweighted pair group with arithmetic mean (UWPGM) method with the Evolutionary distances were computed by using Kimura 2-parameter substitution model and are in the units of the number of nucleotide substitutions per site. The robustness of the tree topology was assessed with 1,000 bootstrap replicates, as implemented within the program, and the gaps or missing data were treated with pairwise deletions (15). and remaining data was divided into four groups. The nucleotide sequences were translated into amino acid sequences to discover any amino acid substitutions. Moreover, similarity percentages among 17 sequences were obtained in the BioEdit 6.0.6 software (14).
Figure 1: Alignment of amino acid sequence of partial VP1 FMDV serotype Asia-1 from Basne with other amino acid sequences showing difference (similarities shown as dot).

Figure 2. Phylogenetic tree based on the nucleotide sequencing VP1 of FMDV/Asia1/ recent outbreak in Basne district of Sulaimani province, which is showed relationships with the other isolate sequences were retrieved from GenBank/database.
Results

The collected samples were tested by RT-PCR, which was performed in two steps. First, RT-PCR amplification of the conserved region in the 5’ UTR was performed using a universal primer set (IF and IR). A positive result was observed for all FMDV serotypes, in which a 328-bp band was detected (11). In the second step, a uniplex PCR was used with primers to detect FMDV serotypes (O, A, and Asia1). The amplicon was confirmed by agarose gel electrophoresis, where a single 911-bp band was detected, revealing that the primer (As1 IC-505/NK61) specific for Asia1 amplified the correct size VP1 product. No amplicon band appeared with the other primer sets, which were specific for serotypes O and A. The results of uniplex PCR showed that the serotype Asia1, was the causative agent for FMD that circulated in the villages of the Basne district.

Phylogenetic tree and sequence analysis

Based on the VP1 gene sequence, a phylogenetic tree was constructed to clarify the genetic relationship of isolated FMDV serotype Asia1 with the FMDV strains from Pakistan, Afghanistan, Iran, India, and Bhutan (Fig. 2). The Phylogenetic tree revealed genetic relationships, showing that the Basne/Sul/2015 isolate circulating in the Basne district was closely related to the group VII virus, which fell into four lineages isolated in Balochistan (2011) and Sindh (2008-2009) in Pakistan. Members of this group, which had been isolated four to seven years earlier, had 97% to 99% nucleotide identity with the viruses in this study.

The partial VP1 amino acid sequences of the Basne/Sul/2015 isolate were compared with the 16 sequences previously published in the GenBank and showed 38 variable amino acids and (8-24) amino acid substitutions with all obtained virus sequences (Fig. 2).

The eight and ten amino acid substitutions of the Basne isolates were S42P, H43Y, V119A, S122P, V133I, E153D, N181D, H182R, and sequences were compared with the JX435109/BAL/PAK/iso-2/2011 isolate. Sequences A31T and A134V were compared with the JX435110/BAL/PAK/iso-1/2011 isolate. In addition, samples with 10 and 11 amino acid substitutions included the following substitutions: A9T, K25R, N28Q, S43P, H44Y, V119A, S122A, T139A, N181D, and H182R, and were compared with the HQ439189/SIN/PAK/L2810/2009 isolate. Those with the V133I substitution were compared with the HQ439187/SIN/PAK/L5/2008 isolate. In addition, samples with 10 and 11 amino acid substitutions included the following substitutions: A9T, K25R, N28Q, S43P, H44Y, V119A, S122A, T139A, N181D, and H182R, and were compared with the HQ439187/SIN/PAK/L5/2008 isolate. In the current study, it was noticed that the Basne/Sul/2015 isolate had eight amino acid substitutions (Fig. 1) and showed higher similarity (95.7%) to the group VII cluster (JX435109/BAL/PAK/iso-2/2011) circulating in the Balochistan province of Pakistan in 2011. Therefore, according to the nucleotide and amino acid sequence analysis, the Asia1/FMD/Basne/Sul/2015 virus isolated in the Basne district showed the highest similarity with group VII, particularly with JX435109/BAL/PAK/iso-2/2011 isolate, which was from clinically diseased animals in Balochistan, Pakistan.
Discussion

Early diagnosis of FMDV is a vital tool for disease control and it has been established that quick control of FMD is the foremost way to reduce dissemination of the causative virus to other non-infected regions (16,17). In this study, uniplex RT-PCR was used to diagnose FMDV using the generic reverse and forward primers (IF/IR) which recognize a highly conserved region in the 5’ UTR of the FMDV. Reid et al. (11) in the UK, previously found the IF/IR set was sensitive in RT-PCR in the primary diagnosis of FMD. However, IF/IR primers cannot be used to identify specific serotypes (O, A, C, and Asia1) since the 5’ UTR does not contain serotype-specific information. The nested RT-PCR was an additional improvement in the primary diagnosis of FMD (18) allowing for the successful detection of the Asia1 serotype, which is consistent with previously reported results using the same primer set (13).

During this study, the conserved region of the VP1 gene from the FMDV was used for detecting different serotypes. This region of the gene has the most variable regions among the capsid polypeptide sequences. Conventional RT-PCR was successful for amplification in this region. Moreover, VP1 contains important immunogenic sites, including amino acid residues within the G-H loop and the C-terminus. The G-H loop includes an arginine-glycine-aspartic acid (RGD) motif that is necessary for attachment of the virus to integrin receptors on the host cell (19,20). Phylogenetic analyses based on the VP1 nucleotide sequence have been used widely to deduce evolutionary dynamics, understand genetic relationships among the genetic lineages, and trace the origin and movement of outbreak strains (21,22). Additionally, researchers have used the VP1 coding sequence for molecular epidemiology in an effort to determine the origin of the Asia1 serotype outbreaks circulating in Pakistan, Afghanistan, Tajikistan, Kyrgyzstan, and Hong Kong (11,23,24).

Using BLAST in NCBI, the VP1 nucleotide sequence of the Asia1 FMDV serotype responsible for FMD outbreaks in the Basne district in 2015 were compared with other isolates in GenBank, indicating that the virus in this study was closely related to the following isolates: Asia/BAL/PAK/sio-1/2011, Asia/BAL/PAK/sio-2/2011, Asia/SIN/PAK/L2954/2009, and Asia/SIN/PAK/L2810/2009. Their overall sequence identity was 94% and this close relationship was confirmed by phylogenetic tree analysis, as shown in (Fig. 1). Surprisingly, the isolated FMDV from outbreaks in the Basne district genetically belonged to a novel group, group VII, which had been detected for the first time in the Sindh province of Pakistan in 2008-2009 and was later identified in Afghanistan, Turkey, Iran, and west Eurasian regions (8).

For additional confirmation, the partial amino acid sequence of VP1 was used for analysis and revealed that the FMDV in the Basne district was closely related to the Asia/BAL/PAK/sio-2/2011 isolate (Accession no. JX435109), which was responsible for the FMDV outbreak in the Balochistan province in Pakistan in 2011. It was determined that the sequence identity between the two was 95.7%, as shown in Table 1, and there was an even greater similarity between the two at the amino acid level. This finding suggested that the Asia1 FMDV circulating in Basne, epidemiologically originated in Pakistan, although this strain has been reported previously in the middle east (25). The isolates of FMDV serotype Asia1 were collected from 2003 to 2007 from several Asian countries and were classified into six groups, I–VI, based on the nucleotide divergence, with a 5% cut-off (26). Later, Jamal et al. (9) identified a novel group, group VII, from Pakistan, which was collected from 2008 to 2009.

Conclusion

This study revealed that the FMDV serotype Asia1 circulating in the Basne district genetically belonged to the group VII, which was imported into this region from the Balochistan province in Pakistan. Furthermore, using of an appropriate vaccine, such as that for group VII strains in this case, is critical. However, further studies will be required to confirm these findings.

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Competing interests

The authors report no conflicts of interest in this work.

Ethical approval

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Contributorship statement

Jeza Muhamad Abdul Aziz "Conception and design, Acquisition of data, Analysis, and interpretation of data."
Salih Ahmed Hama "Drafting the article, Critical revision of the article". Hawre Kamel Faraj "Final approval of the version to be published".

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