Sequence analysis of open reading frames (ORFs) 2 to 4 of a U.S. isolate of porcine reproductive and respiratory syndrome virus

Brief Report

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Summary. The sequence of ORFs 2 to 4 of a U.S. isolate of porcine reproductive and respiratory syndrome virus (PRRSV), ATCC VR2385, was determined by analysis of a cDNA λ library. The cDNA clones containing PRRSV specific sequences were selected using a VR2385 ORF 4 specific PCR probe and sequenced. The ORFs 2, 3 and 4 overlapped each other and encoded polypeptides with predicted Mr of 29.5 kDa (ORF 2), 28.7 kDa (ORF 3) and 19.5 kDa (ORF 4), respectively. No overlap was found between ORFs 4 and 5, and instead there was a 10 bp sequence which separated these two ORFs. The nucleic acid homology with corresponding ORFs of the European PRRSV isolate Lelystad virus (LV) was 65% for ORF 2, 64% for ORF 3 and 66% for ORF 4. Comparison of the ORF 4 sequences of VR2385 with that of another U.S. isolate MN-1b revealed only 86% amino acid sequence homology and the presence of deletions in the ORF 4 of MN-1b. Our results further strengthen the observation that there is sequence variation between US and European PRRSV isolates.

Porcine reproductive and respiratory syndrome virus (PRRSV) belongs to the newly proposed virus family Arteriviridae, which also includes equine arteritis virus (EAV), lactate dehydrogenase-elevating virus (LDV) and simian hemorrhagic fever virus (SHFV). Porcine reproductive and respiratory syndrome (PRRS) was first described in the U.S. in 1987 [9]. A similar disease referred to as porcine epidemic abortion and respiratory syndrome (PEARS) was then reported in Europe [17]. PRRSV was first isolated in Europe and is believed to be widespread in swine population around the world [4, 21, 22]. All European isolates of PRRSV are antigenically and genetically related, whereas there are antigenic variations between US and European isolates as well as among US isolates [1, 16, 21]. The complete
nucleotide sequence of the genome of LV has been determined [14], but until recently
limited information was available about the molecular structure of the genome of
North American isolates of PRRSV [10–13]. We have previously reported the
cloning and sequencing of the ORFs 5 to 7 of a U.S. isolate of PRRSV VR2385 of
high virulence [12]. The 3' end of the genome of the VR2385 and the other U.S.
PRRSV isolates showed a striking difference when compared to the European
isolates [13]. In this study, we report on the cloning and sequencing of the ORFs
2–4 of the U.S. isolate VR2385.

For sequencing and characterization of the viral genome of VR2385 a cDNA
library was constructed. The CRL11171 cells were infected with VR2385 virus at a
M.O.I. of 0.1 and the total RNA from infected cells was isolated at 24 h post
infection by using a guanidinium thiocyanate method [18]. Polyadenylated RNA was
enriched, reverse transcribed and cloned into the λ-ZAP vector using the Uni-
Zap cDNA cloning kit (Stratagene, La Jolla, CA). A PCR probe generated by
ORF 4 specific primers DP585 (5'GCTTTGCTGCTCCTCAAG 3') and DP586
(5'GATGCCTGACACATTGCC 3') [11] were used to screen the library. Plaques
that hybridized with the probe were isolated and purified. The phagemids containing
viral cDNA inserts were rescued by in vitro excision using ExAssist helper phage
and E. coli SOLR cells (Stratagene, La Jolla, CA). Several recombinant phagemids
with virus specific cDNA inserts with sizes ranging from 2.3 to 3.9 kb were selected
and sequenced by Sanger’s dideoxynucleotide chain termination method [19] with an
automated DNA sequencer (Applied Biosystems, Foster City, CA). Universal,
reverse and specific internal primers were used to determine the sequence. At least 3
independent clones representing sequence of the ORFs 2 to 4 were sequenced. The
sequencing data was assembled and analyzed using Mac Vector (International
Biotechnologies, Inc., CT) and GeneWorks (IntelliGenetics, CA) computer
programs. The nucleotide sequence reported in this paper has been deposited in the
GenBank with the accession number U20788.

Analysis of the nucleotide sequence identified three partially overlapping ORFs.
The ORF 2 extended from nucleotide 28 to 795, ORF 3 from 651 to 1412, and ORF 4
from 1196 to 1729. There was an overlap of 144 bp between ORFs 2 and 3, and 216 bp
between ORFs 3 and 4. Surprisingly, no overlap was found between ORFs 4 and 5.
The start codon of ORF 5 was located 10 bp downstream of the stop codon of ORF 4.
However, the ATG start codon of ORF 5 and TGA stop codon of ORF 4

| ORF | VR2385 | LV |
|-----|--------|----|
|     | size (bp) | predicted Mr of product (kDa) | potential N-glycosylation sites | size (bp) | predicted Mr of product (kDa) | potential N-glycosylation sites |
| 2   | 768    | 29.5 | 2 | 747    | 28.4 | 2 |
| 3   | 762    | 28.7 | 7 | 795    | 30.6 | 7 |
| 4   | 534    | 19.5 | 4 | 549    | 19.3 | 4 |
overlapped by only 1 bp in LV [5, 14]. The sequence at the region of the ORF 4 and ORF 5 junction of LV is ATATGA. We sequenced the corresponding region of 5 additional independent clones of VR2385 and in all cases the sequence of this region of VR2385 was ATTTGA. The point mutation from A to T in VR2385 and a

VR2385 ORF 2 MNPGLC--K----AFLKLALN-FLNLGELSRSSHCPFPICLIGLVTWVFPLASLPQGQWSFASSWAPFSPSTLVNLFTPL 69
LV ORF 2 MNPQHSGVRASCSNTKLSSSTLLWVHSL S----LFLGELSRSSHCPFPICLIGLVTWVFPLASLPQGQWSFASSWAPFSPSTLVNLFTPL 65

VR2385 ORF 2 SNTRTYAPLSLQCQSLTPTQCHHHNLMLHKLKVSTILDEHRVRMKYIQAQQVQVUEATLSLSSLD 143
LV ORF 2 PNTFRYAEELPNPQFPQVPHKGMSNHHLWVHSL---LFLGELSRSSHCPFPICLIGLVTWVFPLASLPQGQWSFASSWAPFSPSTLVNLFTPL 140

VR2385 ORF 2 VVAPFRQQAALEATCTYKLAESLHMLPMGQTVSSTLQYVTFPTKVPSKLFQQSLTVASHSF 218
LV ORF 2 LTVHPRQQALAEEKNCSLPSESRMLVHSLAV---QQNPQETNNTPLDLCLIFPTKLPQVRPGLISLVAFAT 215

b

VR2385 ORF 3 MANSCTFLYFLCQFLSPCGLAVAGNATYCPEFPELVGGRNPSPFELTVNTYCFPCLTRQQAAYAEFPQSSSNVLCNGC 50
LV ORF 3 MHQCARFHPFLDQICTYISRLASGSSSLGQWPSLLMLALNIIPTPMLTTECHFCTGTSQARQLPSGRNNVLCNGC 50

VR2385 ORF 3 RGGGSCDSEHDELQFVPSGLSHSEGLTSAAYLQAPLSFYTAQFPHEFGNVTQYDIQPICAVGDG 150
LV ORF 3 RGGGSCDSEHDELQFVPSGLSHSEGLTSAAYLQAPLSFYTAQFPHEFGNVTQYDIQPICAVGDG 149

VR2385 ORF 3 QNMTLSHEDNAGVTGYYQTVSDDQSVHFLPRLPFAAASVGGVSWAPVPSVRSFTVPSQQA 225
LV ORF 3 HNSTVSGTGRSISAFLYAYHTHQDOSWPHFELRLPSLSSVILVHSLWNPFLRSVEFVSFRTPQILAPFRPLSV 224

VR2385 ORF 3 LLSKTSV---AGLQATRPLRPA------KS---LEAEKR 254
LV ORF 3 SSWFPSTSTVSDLTSQQRERFSEPESRHSPVKEPSVPLSTSR 265

c

VR2385 ORF 4 MAALGFLSLYGPCLLVQSAQAPCHQFSSLSDKXTMDTAAAGFAVLQDISLCHRR--NSASEAIR--KVPQRCRT 71
LV ORF 4 MAALGFLSLYGPCLLVQSAQAPCHQFSSLSDKXTMDTAAAGFAVLQDISLCHRR--NSASEAIR--KVPQRCRT 75
MN1b ORF4 MAALGFLSLYGPCLLVQSAQAPCHQFSSLSDKXTMDTAAAGFAVLQDISLCHRR--NSASEAIR--KVPQRCRT 66

VR2385 ORF 4 ADOFVYTNTVANHTDIYSLNLLSCLLSCLFLYVAEMERKGPVFQPVGFVSITVRACUNFSTYSQVRQRPFTQSL 146
LV ORF 4 ADOFVYTNTVANHTDIYSLNLLSCLLSCLFLYVAEMERKGPVFQPVGFVSITVRACUNFSTYSQVRQRPFTQSL 140
MN1b ORF4 ADOFVYTNTVANHTDIYSLNLLSCLLSCLFLYVAEMERKGPVFQPVGFVSITVRACUNFSTYSQVRQRPFTQSL 141

VR2385 ORF 4 VVDH--VLLHMSMMFTQEVATLACLPTILLAI 178
LV ORF 4 LVIDHRHLLHLLSPAMNATACLCPLATTAL 183
MN1b ORF4 DFRVHLMH---PBMQMNATVLCPLATTAL 171

Fig. 1. Alignment of predicted amino acid sequence of ORFs 2 (a), 3 (b), and 4 (c) of PRRSV VR2385 and LV. The ORF 4 sequence of another U.S. isolate MN-1B was also included in the alignment. The sequences for LV were reported by Meulenberg et al. [14], and the ORF 4 sequence for MN-1B was reported by Kwang et al. [10]
probably some other unidentified changes in this region of VR2385 made the ORF 5 ATG start codon 10 bp downstream of the stop codon of ORF 4, and a 10 bp non-coding region appeared in the ORF 4 and 5 junction of VR2385.

The characteristics of ORFs 2 to 4 of VR2385 are summarized in Table 1. The ORF 2 encodes a 256 amino acid polypeptide with a predicted size of 29.5 kDa. The carboxy and amino terminus of the predicted protein are hydrophobic (data not shown) and there are two potential N-glycosylation sites in the ORF 2 protein. ORF 3 encodes a protein of 254 amino acids and contains 7 potential N-glycosylation sites. The amino terminus of the ORF 3 protein is extremely hydrophobic. ORF 4 encoded a 178 amino acid protein with a predicted size of 19.5 kDa. The amino and carboxy termini and 4 regions within the protein are highly hydrophobic. Comparison of the nucleotide sequences of VR2385 and LV showed extensive variations. Nucleotide sequence identity between VR2385 and LV is 65% for ORF 2, 64% for ORF 3 and 66% for ORF 4. Alignment of the predicted amino acid sequences of ORFs 2–4 of VR2385 and LV is presented in Fig. 1. Amino acid identity between VR2385 and LV is 58% for ORF 2, 56% for ORF 3 and 67% for ORF 4. We also compared the sequence of VR2385 ORF 4 with that of MN-1b, another US isolate of PRRSV [10]. The ORF 4 of VR2385 is 21 bp longer and shares an 88% nucleotide sequence homology with MN-1b. The amino acid homology between the ORF 4 of VR2385 and MN-1b is 86% (Fig. 1c). Several deletions were found in the ORF 4 of MN-1b compared to VR2385.

The ORFs 6 and 7 of PRRSV are predicted to encode the viral membrane glycoprotein and the viral nucleocapsid protein, respectively [12, 13]. Analysis of predicted amino acid sequences encoded by ORFs 2–5 of LV, LDV and EAV showed that all of these proteins share features of membrane associated proteins [5, 6, 8, 14]. The EAV ORF 5 product was identified as the main envelope glycoprotein.

| Table 2. Potential leader-mRNA junction regions in the genome of VR2385 and Lelystad virus |
|-----------------------------------------------|
| **ORF** | **VR2385 sequence** | **position** | **Lelystad virus sequence** | **position** |
| 1 | - | - | UUAACC | - |
| 2 | UGAACC | 20 | UAAACC | 38 |
| 3 | GUAACC | 83 | UUGACC | 11 |
| 4 | CCAACC | 35 | - | - |
| 5 | UUGACC | 230 | CAGACC | 44 |
| 6 | - | - | UCAACC | 83 |
| 7 | UAAACC | 17 | ACAACC | 32 |

*aSequence for VR2385 ORFs 2–4 is presented in the study, ORFs 5–7 was reported by Meng et al. [12] and LV ORFs 1–7 was reported by Meulenberg et al. [14].

*bDistance in nucleotides between proposed junction motif and AUG start codon of downstream ORF.
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[7]. Our data indicates that the proteins encoded by ORFs 2–4 of VR2385 possess characteristics similar to those of LV and probably are envelope or membrane associated glycoproteins because of their hydrophobicity and presence of potential glycosylation sites. Further work is necessary to determine the roles of these proteins. The variability found in the ORF 4 sequence between the two U.S. isolates correlate with the findings that ORF 4 protein of the MN-1b expressed in E. coli reacted with only 65% of PRRSV positive sera by Western blot analysis [10].

A nested set of subgenomic mRNA is formed during replication of PRRSV and other members of the arterivirus group [5, 6, 8, 12, 14]. All subgenomic mRNAs contain a common leader sequence derived from the 5' noncoding region of the viral genome. The site of the leader-mRNA junction is similar and located upstream of the start codon of each ORF. The consensus leader-mRNA junction sequence of the six subgenomic mRNAs of LV was determined to be (U/A)(C/U/A)(A/G)ACC [15]. Similar sequences were also found as leader-mRNA junction regions for LDV [3]. The potential leader-mRNA junction motifs of ORFs 2 to 4 of VR2385 was proposed and compared with those of LV (Table 2). The last four nucleotides of the motif for ORFs 1, 2, 4, 5, 6 and 7 in LV are AACC, and for ORF 3 is GACC. The AACC motif has been found upstream of ORFs 6 and 7 of VR2385 [12] as well as ORFs 2 and 3. There are two potential junction regions for ORF 3, 83 bp and 35 bp upstream of the ORF 3 start codon, respectively (Table 1). No AACC motif was found upstream of VR2385 ORFs 4 and 5. However, the sequences UUGACC and CAGACC upstream of ORF 4, UUGACC and GAGACC upstream of ORF 5, may be the leader-mRNA junction regions for the mRNAs 4 and 5 of VR2385. Multiple potential leader-mRNA junction sites suggest that polymorphism of subgenomic mRNAs may exist among PRRSV isolates. Experiments to determine the exact locations of leader-mRNA junction regions are now in progress.

The sequence variations observed in this study between a U.S. and a European PRRSV isolate, as well as between two North American PRRSV isolates, indicates the heterogenetic nature of PRRSV isolates and the need for further characterization of additional PRRSV isolates. Whether this genetic variation between VR2385 and LV reflects the observed difference in virulence needs to be further studied.

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