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Sequence Analysis of the 3'-End of Feline Calicivirus Genome

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The nucleotide sequence of the 3'-end of the Japanese F4 strain of feline calicivirus (FCV) RNA was determined from a cloned cDNA of 3.5 kbp. We found three open reading frames (ORFs). The largest ORF encoded a 668-amino acid protein of 73,588 Da, which was presumably the capsid precursor protein of FCV and had significant amino acid sequence homology with the VP3 of picornaviruses. A small ORF at the extreme 3'-end was compared with that of the F9 strain of FCV, a vaccine strain originally from the U.S. Highly conserved amino acid sequences were shown, suggesting that this ORF might be functional and encode a putative 106-amino acid protein of 12,153 Da. The other ORF in the 5'-flanking region of the cDNA had consensus amino acid sequences conserved among the RNA-dependent RNA polymerases.

Caliciviridae is a family composed of single-stranded, positive-sense RNA viruses whose spherical virions are 35-40 nm in diameter and consist of a single major capsid protein of 60-70 kDa (1). Members belonging to this group are feline calicivirus (FCV), vesicular exanthema of swine virus (VESV), and San Miguel sea lion virus (1). Although the caliciviridae had been provisionally classified as a genus of the family Picornaviridae (2), they were reclassified as a new family Caliciviridae because of, in addition to their distinct morphology of the virion (3, 4) and possession of a single capsid protein (5-7), the presence of a subgenomic RNA found in calicivirus infected cells (8, 9).

Relatively few studies with regard to the molecular biology of the caliciviruses have been carried out. Neill and Mengeling (10) first cloned cDNAs from the CFI strain of FCV, which was isolated in the U.S. (11), and analyzed the FCV-specific RNAs formed in the virus-infected cells using the cDNAs as probes. They indicated that an 8.2-kb genomic RNA and three subgenomic RNAs of 4.8, 4.2, and 2.4 kb synthesized in the cells were nested, coterminal transcripts with common 3'-ends. Carter (12) reported the sequence of the 3'-terminal 505 nucleotides (nt) of FCV RNA using the F9 vaccine strain whose origin was the U.S. (13). The cDNA was used as a probe in Northern blot analyses to confirm and extend the observations of Neill and Mengeling (10). Recently, Neill (14) has determined the nucleotide sequence of the central region of the FCV genome and found a large open reading frame (ORF) which terminated 2.4 kb from the 3'-end of the genome. The deduced amino acid sequence of the region was shown to encode three picornavirus-like non-structural proteins, RNA-dependent RNA polymerase, cysteine protease, and 2C polypeptide. Although the data were not published, Neill (14) also mentioned the existence of another large ORF in the 3'-terminal 2400 bases of the genomic RNA, which was entirely encoded by 2.4 kb subgenomic RNA.

Previously, we have identified seven neutralizing epitopes of the F4 strain of FCV, the prototype strain of FCV in Japan, with the aid of an extensive panel of neutralizing monoclonal antibodies and neutralization-resistant variants (15, 16). Four of the seven epitopes were shown to exist on the capsid protein of the virus by immunoblot analysis (16). To further understand the antigenic structure of FCV, it is important to analyze the nucleotide sequence of the gene for the capsid protein.

In this paper we report the molecular cloning and the sequence analysis of a cloned cDNA representing the 3'-end of FCV F4 RNA. Within the sequence of the 3'-terminal 3516 nt we found three ORFs. The deduced amino acid sequences of the two ORFs at the extreme 3'-end and in the 5'-flanking region of the cDNA were compared with those of other FCV strains reported previously by Carter (12) and Neill (14). The other ORF which located between the two ORFs was
suggested to encode the capsid precursor protein having significant amino acid sequence homology with the VP3 of picornaviruses.

Crandell feline kidney (CRFK) cells were grown in Dulbecco’s modified Eagle’s medium supplemented with 8% fetal calf serum and antibiotics. The FCV F4 strain (7) was propagated in CRFK cells and plaque-purified to prepare working stocks of the virus without interfering particles. Following adsorption for 90 min at 37°, CRFK monolayers infected with the FCV F4 strain were further incubated for 24 hr, and the cell culture fluid was harvested. The cell debris was removed by centrifugation at 8000 rpm for 30 min at 4° (Sorvall GS-3 rotor, U.S.). The resulting supernatant was layered over 25% (w/w) sucrose in phosphate-buffered saline containing 0.1% Triton X-100 (PBS-TX) and centrifuged at 24,000 rpm for 3 hr at 4°C (Hitachi RPS25-2 rotor). The pellet was resuspended in PBS-TX, layered on a 5 to 35% (w/w) sucrose density gradient, and centrifuged at 24,000 rpm for 2 hr at 4°C (RPS25-2 rotor). Fractions were collected from the bottom of tubes. Infectivity and optical density at 260 and 280 nm of the fractions were measured. The peak fractions were pooled and then SDS was added to a final concentration of 0.5%.

Genomic RNA was extracted with phenol–chloroform from the pooled fractions and precipitated with ethanol. The RNA was placed on 15–30% (w/w) sucrose gradient in TNE buffer (10 mM Tris–HCl (pH 7.4), 100 mM NaCl, 1 mM EDTA) containing 0.5% SDS and centrifuged at 19,000 rpm for 18 hr at 22°C (Hitachi RPS40T rotor). The peak fractions in optical density at 260 nm were pooled and precipitated with ethanol. The purified viral genomic RNA was initially used for cDNA cloning using oligo (dT) as a primer for reverse transcription. cDNA cloning was performed using a recombinant plasmid (BRL, Gaithersburg, MD). The annealed molecules were used to transform Escherichia coli MC1061 and screened by plating on Luria broth agar plates containing 12.5 μg/ml tetracycline. The recombinant plasmids from tetra-resistant colonies were screened to determine whether such a protein is synthesized in FCV-infected cells.

The 3′-end (nucleotide positions 3152 to 3516 in Fig. 1) of pFCV 119 contained only four adenyllic acids at the end, the sequencing analysis of other clones indicated that the adenyllic acids were a part of a poly(A) stretch at the 3′-end of FCV F4 genome. Figure 2 shows a schematic diagram of the possible ORFs obtained from the predicted amino acid sequences. A large ORF (2004 nt) in frame 3 and a small ORF (318 nt) in frame 2 were found. There was another ORF (1143 nt) in the 5′-flanking region of frame 1, which remained open at the extreme 5′-end.

The short ORF in frame 2 and noncording region at the 3′ end (nucleotide positions 3152 to 3516 in Fig. 1) were compared with those reported by Carter (12). Both sequences were identical in nucleotide length and had ATG and TGA codons for the small ORF (nucleotide positions 1 to 316 in Fig. 3) at the same positions. In the coding region, there were 47 nt (14.8%) changes in 318 nt, but only 8 nt changes were accompanied by amino acid changes. The amino acid sequences in this ORF were highly conserved (92.5%) between the strains of different origins, suggesting that this ORF might be functional and encode a protein of 12,153 Da. For this ORF, a corresponding intracellular RNA of 550 bases was detected as a candidate band by Northern blot analyses (12). It will be interesting to determine whether such a protein is synthesized in FCV-infected cells.
When the deduced amino acid sequence of the ORF in the 5'-flanking region of frame 1 was compared with the RNA polymerase-like sequence of FCV reported by Neill (14), strong homology (89.5%) was found between the sequences of the Japanese F4 and the American CFI strains of FCV. It was also found that the sequence in this region of FCV had consensus amino acid sequences conserved among the RNA-dependent RNA polymerases.
dent RNA polymerases (22), including the KDEL, GM(L)PSG, YGDD, and FLKR sequences (boxed sequences in Fig. 1), as indicated by Neill (14). Therefore, the ORF in frame 1 was suggested to contain a part of the coding sequence for the RNA polymerase of FCV.

The largest ORF which could encode a protein of 668 amino acids was found in frame 3 (Fig. 2). The position of this ORF in the FCV genome was thought to locate in the region for the most abundant subgenomic mRNA of FCV, one of the mRNAs to form a 3' c-terminal nested set, whose size was reported to be 2.4 (10) or 2.7 kb (12). We also detected a 2.5-kb RNA by Northern blot analysis (data not shown) using the cDNA probe prepared from this ORF. This mRNA has been considered to produce the capsid protein of FCV (8–10, 23). If translation is performed from the first ATG in this ORF (nucleotide positions 1149 to 3152 in Fig. 1) to the termination codon, a protein with a molecular weight of 73,588 is expected to be produced. This molecular weight is similar to that of the 76-kDa capsid precursor protein of FCV estimated by sodium dodecyl sulfate–polyacrylamide gel electrophoresis (24). From these observations, it is strongly suggested that this ORF encodes the capsid protein of FCV F4. Preliminary experiments indicated that the capsid protein was blocked to Edman degradation. The presence of a blocked amino terminus was also indicated in attempts to dansylate the amino terminus of the capsid protein. The consensus sequence was calculated to represent areas where a majority of the sequences of many picornaviruses conserve specific residues or a series of very similar residues (20). The asterisks indicate the amino acid residues of FCV conserved in one of the two picornaviruses but not in the consensus sequence. The plus signs indicate amino acid residues that are present in all the sequences of the three viruses but not in the consensus sequence. The colons indicate amino acid residues of FCV conserved in one of the two picornaviruses.

A computer-aided search of known amino acid sequences as contained in the NBRF database revealed significant homology between the deduced amino acid sequences of the capsid precursor protein of FCV and the capsid protein VP3 of picornaviruses. The amino acid sequences of the VP3 of poliovirus Sabin 1 and foot and mouth disease virus A12, and an analogous region of FCV (an underlined amino acid sequence in Fig. 1) and VP3 of picornaviruses. The consensus sequence was calculated to represent areas where a majority of the sequences of many picornaviruses conserve specific residues or a series of very similar residues (20). The asterisks indicate the amino acid residues of FCV conserved in one of the two picornaviruses but not in the consensus sequence. The plus signs indicate amino acid residues that are present in all the sequences of the three viruses but not in the consensus sequence. The colons indicate amino acid residues of FCV conserved in one of the two picornaviruses.

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two representative picornaviruses suggested that these proteins have similar function(s) and have been diverged from a common ancestral protein. Similarities observed not only in nonstructural proteins between FCV and picornaviruses (14), but also in a structural protein between these viruses support the grouping of caliciviruses into the superfamily II, picornavirus-like families (27).

It has been suggested that the scheme of transcription of FCV is similar to that employed by the coronaviruses and the alphaviruses (10, 14, 28). Our nucleotide and amino acid sequence data appear to confirm the similarity in the genomic structure of FCV with that of the alphaviruses, since both viruses have the gene(s) for structural protein(s) at the 3′ region of their genome and the polymerase gene at the center of each genome. However, the junction sequences between the three ORFs of FCV were short (5 nt, nucleotide positions 1144 to 1148 in Fig. 1) or absent (1 nt overlapped, nucleotide position 3152 in Fig. 1). The junction sequences in the genomes of the coronaviruses and the alphaviruses are regarded to be of importance in transcriptions of their mRNA(s) (27, 29). The strategy of transcription in FCV replication may differ from that of the alphaviruses or the coronaviruses.

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