Enteric Coronavirus in Ferrets, the Netherlands

To the Editor: Coronaviruses (CoVs) are enveloped, positive-sense, single-stranded RNA viruses that can cause acute and chronic respiratory, enteric, and central nervous system disease in a variety of animal species (1). Recently, a novel ferret enteric CoV (FRECV) was identified in domesticated ferrets (Mustela putorius) in which epizootic catarrhal enteritis had been diagnosed; the illness was characterized by foul-smelling green diarrhea with high mucus content, lethargy, anorexia, and vomiting (2). Another ferret CoV emerged in ferrets for which systemic pyogranulomatous inflammation, resembling the clinical and pathologic features of feline infectious peritonitis (FIP), was diagnosed (3–5).

In 2010, we investigated the prevalence of CoV antibodies in 85 asymptomatic ferrets obtained from 1 ferret farm in the Netherlands. Previous studies have shown that antibodies against different members of the α-CoVs show broad cross-reactivity (6). We used FIP virus (FIPV)–infected cells to screen for CoV antibodies in an indirect immunoperoxidase assay. Because 32% of the ferret serum had a titer >20 in this assay, we concluded that these animals most likely had been exposed to a CoV. To test for a CoV in these animals, we analyzed RNA extracted from rectal swabs with a degenerate set of primers to amplify a conserved region within open reading frame (ORF) 1 of CoVs (7). Remarkably, 36 (42%) of samples tested were PCR positive, suggesting excretion of a CoV by a substantial proportion of ferrets tested. To corroborate that the CoV detected in the rectal swabs was a ferret CoV (FRCrCoV), we amplified and sequenced the nucleocapsid protein by using primer pair 5′- TCCCC CGGGGGCTGGCAACGGACAA CTGTG-3′ and 5′-CCCAAGCTTTTATTTA GTT GACT AATA TTCCA A-3′. Phylogenetic analyses of 2 of the sequences obtained indicated a variant nucleocapsid that was similar to other FRCrCoVs described previously but that did not group with 1 of these sequences directly (Figure). Amino acid alignment of 1 of these sequences (FRCrCoV-511c) with FRECV-MSU2 demonstrated 91.8% identity and 95.7% similarity, whereas this virus shows 89.3% identity and 95.2% similarity to ferret systemic CoV (FRSCV-MSU1).

On the basis of obtained and published nucleocapsid sequences (2,4), we developed a TaqMan reverse transcription PCR to detect viral RNA using the following primers and degenerate probe: forward, 5′-TTGGAAAGAATG GTGCTAAAACGT-3′; reverse, 5′-CA TTAGCACGTATCCATCAAATT -3′; and probe, 5′-TACGACTCGT GGCCAACACCA-3′. Using this more specific and sensitive assay, we detected viral RNA in 63% of the rectal swabs tested; other CoVs including FIPV, severe acute respiratory syndrome–CoV, and human CoV NL-63, were not amplified by this assay (data not shown). All samples that had tested positive in the ORF1-CoV PCR were confirmed positive with this TaqMan assay. To analyze FRCrCoV in ferrets from geographically distinct sites, we tested fecal samples from 90 animals without signs of disease (including epizootic catarrhal enteritis) from 39 different locations in the Netherlands. FRCrCoV nucleocapsid TaqMan and ORF1-CoV PCR demonstrated that 61% of the fecal samples and 72% of the locations were positive. Multiple testing of fecal swabs at different times and use of FRCrCoV-specific antibody assays would probably further increase the FRCrCoV prevalence rate.

Further partial sequence analysis of the spike gene by using primers 5′-AARRTTATGAGTGTGTMGM DTC-3′ and 5′-CAACTCYYTTAA GCCARTCAAGG-3′ clearly showed that these viruses are more closely related to systemic FRCrCoVs than to FRECV (Figure). Amino acid alignment of 1 of these sequences (FRCrCoV-511c) with FRECV-MSU2 demonstrated 78% identity and 89% similarity, and FRCrCoV-511c shows 86% identity and 92% similarity to FRSCV-MSU1.

After identification of severe acute respiratory syndrome CoV in humans in 2003 and related viruses in civet cats and bats, an increase in CoV surveillance in different animal species resulted in identification and characterization of a broad range of previously unrecognized CoVs (8). Here we report an enteric FRCrCoV circulating in the Netherlands in a high percentage of asymptomatic ferrets. The ferrets tested did not have a previous record of foul-smelling green diarrhea described previously to be associated with FRECV, a virus detected in the United States and further characterized in 2006 (2). On the basis of the phylogenetic analysis of the spike sequences, FRECV-MSU2 might have evolved through recombination with some other unknown CoV. Alternatively, the viruses isolated in the Netherlands grouped more closely with FRCrCoVs causing systemic disease (e.g., FRSCV-MSU1). Thus far, no evidence indicates that the animals testing positive for the enteric CoV showed clinical disease that pointed to pyogranulomatous inflammation, necrosis with or without perivasculitis, and vasculitis in abdominal and visceral organs associated with the systemic variant. Further genetic characterization of these enteric FRCrCoVs variants might show genetic differences that could explain the apparent pathotypes of these 2 FRCrCoVs. FRCrCoVs might evolve through mutation or deletion into viruses that cause systemic
disease, or alternatively, different FRCoVs are circulating, analogous to the hypotheses put forward to explain the occurrence of FIP (9,10). Given the use of ferrets in testing efficacy of influenza virus vaccines and the propensity of CoVs to cross species barriers, further surveillance and investigation of the biology of these emerging FRCoVs is warranted.

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Author affiliations: Erasmus Medical Center, Rotterdam, the Netherlands (L.B.V. Provacia, S.L. Smits, B.E. Martina, V.S. Raj, P. v.d. Doel, G. v. Amerongen, A.D.M.E. Osterhaus, B.L. Haagmans); and Ferret Clinic Brouwhuis, Helmond, the Netherlands (H. Moorman-Roest, G. v. Amerongen).

LETTERS


Address for correspondence: Bart L. Haagmans, Department of Virology, Erasmus Medical Center, Dr Molewaterplein, 50, 3000 CA, Rotterdam, the Netherlands; email: b.haagmans@erasmusmc.nl

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