



Molecular detection of canine bufaviruses in wild canids

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Abstract

Novel protoparvoviruses genetically related to human and non-human primate bufaviruses (BuVs) have been detected recently in respiratory and enteric specimens collected from dogs and cats. In this study, by molecular screening of archival collections of faecal samples from wolves and foxes, we detected BuVs with a rate of 17.1% (7/41) and 10.5% (9/86), respectively. Sequence analysis of a portion of the ORF2 gene region of nine positive samples showed that the viruses in these samples were closely related to BuVs (97.5–99.0% nucleotide sequence identity) found in domestic carnivores.

Introduction

Protoparvoviruses are small (20–25 nm), nonenveloped viruses with a single-stranded DNA genome of 4.5–5.5 kb flanked by two terminal hairpin structures [1, 2] and represent a distinct genus in the family *Parvoviridae*. The genus *Protoparvovirus* includes the species *Carnivore protoparvovirus 1* (canine parvovirus type 2 or CPV-2, feline panleukopenia parvovirus or FPV, racoon parvovirus, mink enteritis virus), *Eulipotyphila protoparvovirus 1* (megabat bufavirus 1), *Primate protoparvovirus 1* (human bufavirus 1a), *Primate protoparvovirus 2* (rhesus parvovirus 1), *Primate protoparvovirus 3* (human cutavirus), *Rodent protoparvovirus 1* (minute virus of mice), *Rodent protoparvovirus 2 and 3* (rat parvoviruses), *Ungulate protoparvovirus 1*, *Ungulate protoparvovirus 2* (porcine parvoviruses), and *Chiropteran protoparvovirus 1* (<https://talk.ictvonline.org/ictv-reports/>

[ictv_online_report/ssdnaviruses/w/parvoviridae/1045/genus-protoparvovirus/history](https://talk.ictvonline.org/ictv-reports/ictv_online_report/ssdnaviruses/w/parvoviridae/1045/genus-protoparvovirus/history)).

Novel protoparvoviruses were identified recently during an outbreak of canine infectious respiratory disease (CIRD) that occurred in a litter of mixed-breed 5-month-old puppies in Italy [3]. Also, by screening collections of faecal and respiratory specimens obtained from juvenile dogs, the virus was detected in stool samples of dogs with or without enteric disease and in nasal and oropharyngeal swabs of dogs with CIRD. Analysis of the complete genome sequences of three strains (ITA/2015/297, HUN/2012/126 and HUN/212/22) [3] showed that a novel canine parvoviruses, designated canine bufaviruses (CBuVs), exhibited the highest sequence identity in NS1 (66.9%–69.4% nucleotide [nt] and 47.2%–51.4% amino acid [aa]) and VP1 (66%–68.2% nt and 62.5%–67.2% aa) regions to human and non-human primate bufaviruses (BuVs) (*Primate protoparvovirus 1* and *Primate protoparvovirus 2*) [4–6], while identity to the well-known CPV-2 (*Carnivore protoparvovirus 1*) was 56% nt and 40.6% aa in NS1 and 42.6% nt and 33.4% aa in VP1. According to the criteria established by the International Committee on Taxonomy of Viruses (ICTV) for classification of parvoviruses into the same species (cutoff > 85% aa sequence identity in the NS1 protein), the canine protoparvovirus may be considered a member of a new species within the genus *Protoparvovirus*, for which the name “*Carnivore protoparvovirus 2*” is proposed.

More recently, evidence was obtained by molecular screening of respiratory specimens collected from cats with or without signs of upper respiratory tract disease (URTD) and stool samples from diarrhoeic animals that feline hosts

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are also susceptible to similar viruses [7]. Sequence analysis of the nearly complete VP2 coding region of three feline strains (ITA/2012/TE109, ITA/2015/BA509 and ITA/2017/BA291) revealed that the viruses displayed > 99.9% nt sequence identity to CBuVs detected in dogs [3]. These findings seem to indicate that genetically closely related parvoviruses circulate between dogs and cats, suggesting limited host-species restriction among domestic carnivores. In order to obtain a more complete picture of the ecology of this novel protoparvovirus, we investigated whether wild canids can also harbor this virus given the availability in our laboratory of two sample collections of enteric origin from wolves (*Canis lupus*) and red foxes (*Vulpes vulpes*).

A total of 127 rectal swabs were collected between September 2009 and April 2017 by the National Reference Center for Wild Animal Diseases (CeRMAS) and submitted to the Faculty of Veterinary Medicine, University of Teramo, Italy. The faecal panel consisted of 41 samples from necropsized wolves found dead after traumatic events and 86 samples from red foxes shot during regular hunting seasons. Viral nucleic acids were extracted using a QIAamp UltraSens Virus Kit (QIAGEN GmbH, Hilden, Germany), following the manufacturer's protocol, and stored at -80°C until use.

The presence of CBuV DNA was assessed by real-time PCR (q-PCR) as described previously [3]. Viral DNA quantification was performed using TaqMan Fast Advanced master mix (Thermo Fisher Scientific, USA) in a 25- μl volume comprising 5 μl of extracted DNA and 20 μl of master mix. The primers CPPV-L3-For (5'-TGAACAAGAAATAGACAACATTGTCAT-3') and CPPV-L3-Rev (5'-AAAGAGCAGTTAGGTCATTGTTGT-3') and TaqMan probe CPPV-L3 (5'-Fam-CCAAACAAGGTACAGGACAGGAAGAAA CAACACAA-BHQ1-3') were used at concentrations of 200 and 100 nM, respectively. The DNA copy number was calculated based on a standard curve that was generated using a series of tenfold dilutions of standard plasmid vector pCR 2.1 (TA Cloning Kit, Invitrogen, Ltd, Milan, Italy) containing a 98-bp fragment of the VP2 region of the canine protoparvovirus 2 strain ITA/2011/297-15 (GenBank accession

no. MF198244) [3]. To minimize potential contamination, the plasmid was prepared in a separate room, and microtubes containing plasmid standards were not taken into the PCR setup room. Three negative controls were included in each run. Viral DNA was detected in 7 out of 41 (17.1%) faecal samples collected from wolves and in 9 out of 86 (10.5%) samples obtained from foxes. Viral loads ranged from 3.6×10^1 to 5.3×10^4 DNA copies/5 μl of template (mean, 5.4×10^3 DNA copies/5 μl , cutoff cycle threshold, 39.00). Using the specific primers CPPV 165 F (5'-CTGGTT TAATCCAGCAGACT-3') and CPPV 371 R (5'-TGAAGA CCAAGGTAGTAGGT-3') [3], which target a 207-bp region of VP2, viral sequences were obtained from five wolves and four foxes. BLAST (<http://www.ncbi.nlm.nih.gov>) and FASTA (<http://www.ebi.ac.uk/fasta33>) analysis showed that nine strains (GenBank accession nos. MK496165-73) shared 97.9–99.5% nt sequence identity with each other and displayed the closest identity to canine and feline BuV strains (97.5–99.0% nt sequence identity) [3, 7]. In order to obtain additional genetic information about the VP2 capsid gene, three positive samples with viral loads ranging from 10^3 to 10^4 DNA copies/5 μl were used as template to amplify viral DNA, using the primer pairs CPPV-1409 F/CPPV-1414 R and CPPV-165 F/CPPV-1571 R [7], which were designed based on alignments of the available sequences of Italian and the Hungarian canine strains (GenBank accession nos. MF198244-6) and were able to amplify two overlapping fragments of the VP2-coding region of 962 nt (CPPV-1409 F/CPPV-1414 R) and 1350 nt (CPPV-165 F/CPPV-1571 R) in length, respectively (Table 1). Despite several attempts, for one wolf faecal sample (130/Wolf/2015/ITA, GenBank accession no. MK496167), we obtained a sequence that was 500 nt in length, corresponding to the N-terminus of VP2, using a strategy of heminested PCR with external primers CPPV-1409 F/CPPV-1414 R and the internal primer CPPV-165 F. Sequence analysis showed that the wolf strain had 99.0–100% nt and 100% aa sequence identity to the Italian and Hungarian canine strains (GenBank accession nos. MF198244-6) and to the Italian feline strains (GenBank accession nos. MK030121-3) [3, 7]. Neighbor-joining

Table 1 List of primers used in this study. Nucleotide positions refer to the sequence of the canine protoparvovirus 2 strain ITA/2011/297-15 (GenBank accession no. MF198244)

Oligonucleotide	Position	Sequence (5' to 3')	Sense	References
CCPV-L3 F	2938-2964	TGAACAAGAAATAGACAACATTGTCAT	+	Martella et al., 2018 [3]
CCPV-L3 R	3012-3035	AAAGAGCAGTTAGGTCATTGTTGT	-	Martella et al., 2018 [3]
CCPV-165 F	2872-2891	CTGGTTTAATCCAGCAGACT	+	Martella et al., 2018 [3]
CCPV-371 R	3060-3079	TGAAGACCAAGGTAGTAGGT	-	Martella et al., 2018 [3]
CCPV-1409 F	2411-2432	TCATATTCCTGGAGAAACATCA	+	Diakoudi et al., 2019 [7]
CCPV-1414 R	3351-3372	ATATGTCTGTTAGATTGCCAGT	-	Diakoudi et al., 2019 [7]
CCPV-1571 R	4201-4219	TTATAGAGTAATATTAGGC	-	Diakoudi et al., 2019 [7]

phylogenetic analysis was performed with a selection of partial capsid sequences representative of the genus *Protoparvovirus*. In the resulting trees (Fig. 1A and B), the wolf and fox sequences clustered tightly with the canine and feline strains. We propose that this lineage could constitute a novel species, tentatively named “carnivore protoparvovirus 2”.

Domestic and wild canids are known to be exposed to common viral pathogens, including CPV-2, canine adenoviruses (CAV1 and CAV2) and canine distemper virus (CDV) [8–11]. Furthermore, the advent of next-generation

sequencing has enabled the identification and characterization of several previously unknown viruses that can be transmitted between dogs and wildlife. These examples include canine kobuviruses (CaKoVs) [12–15], circoviruses [16, 17], and sapoviruses [12, 18]. Therefore, we hypothesized that the carnivore protoparvoviruses recently detected in dogs and cats [3, 7] could also circulate among wild canids. In this study, using specific primers and a probe, novel parvoviruses were found in wolves and foxes at a rate of 17.1% and 10.5%, respectively, suggesting that these viruses are

Fig. 1 (A) Phylogenetic tree based on a portion of the VP2 gene (~ 500 nt) of the wolf strain 130/2015/ITA (GenBank accession no. MK496167). (B) Phylogenetic tree based on a 200-nt region of the VP2 gene of the wolf and fox strains detected in this study (GenBank accession nos. MK496165–73). Both trees were generated using the neighbor-joining method and the Kimura 2-parameter model (MEGA software package, version 3.0), with statistical support provided by bootstrapping with 1000 replicates. The scale bar represents nucleotide substitutions per site. The labels denote the sequences detected in this study

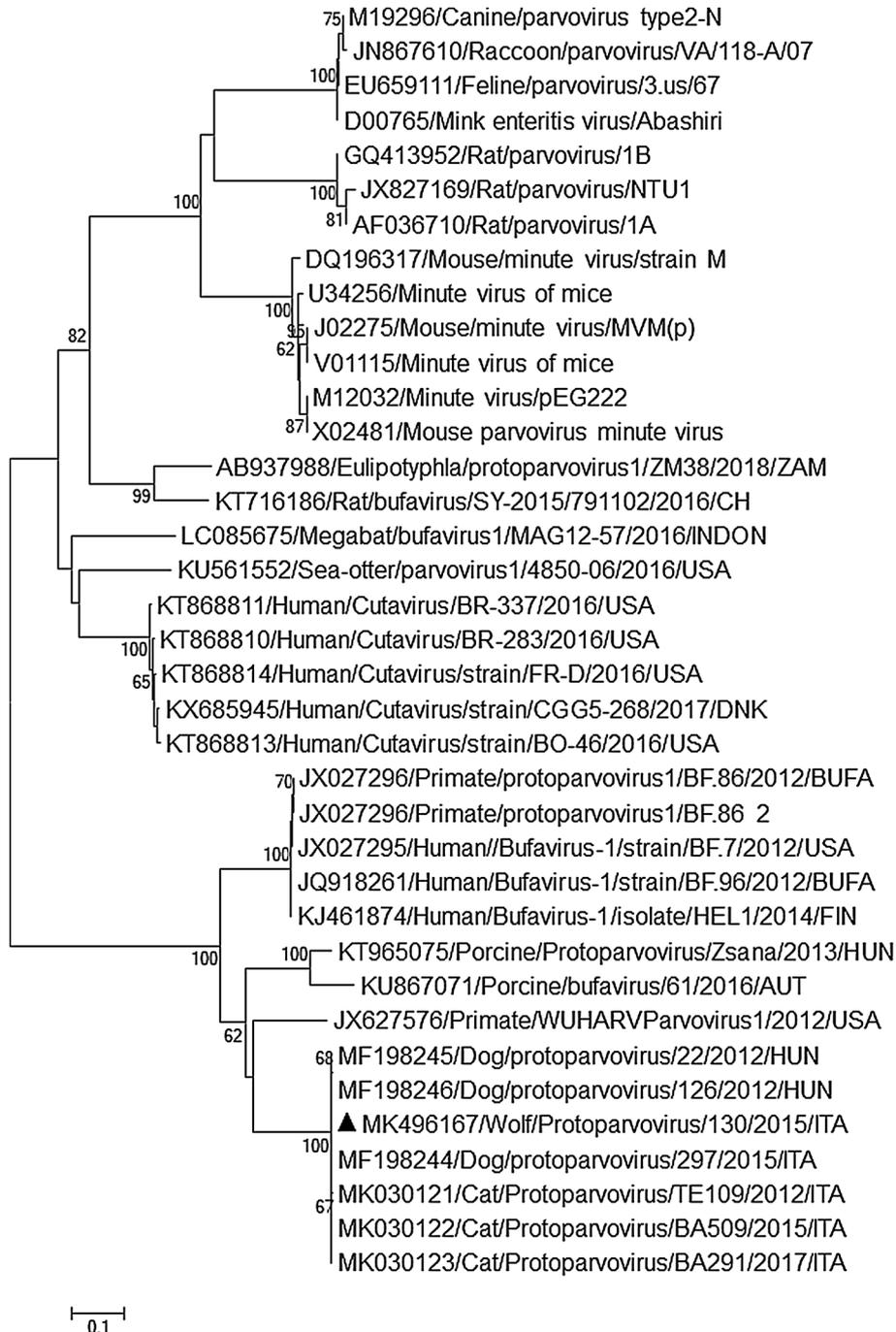
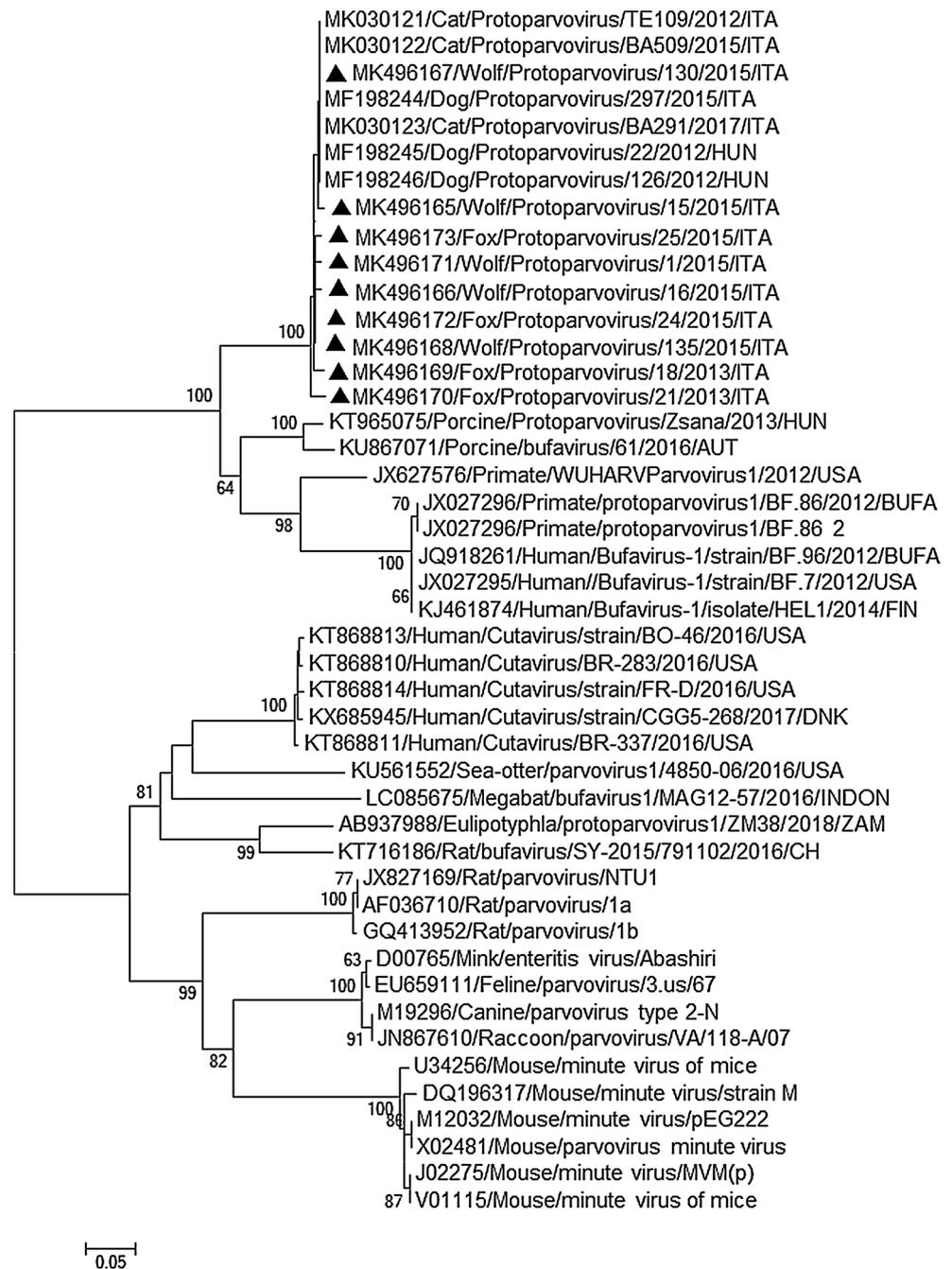


Fig. 1 (continued)



probably as common in wild canids as they are in domestic dogs. Analysis of a short fragment (207 nt) of the ORF2 gene, obtained from nine samples, revealed a high degree of genetic relatedness to canine and feline BuVs [3, 7]. This sequence conservation was also found when analyzing a longer region of the capsid gene (0.5 kb) of the wolf strain 130/Wolf/2015/ITA. However, a more precise and definitive characterization should rely on analysis of the complete genome sequence or at least the full-length capsid gene. In our analysis, all of the positive stool samples were collected from wolves and foxes that were apparently healthy at the

time of sampling, and we did not analyze specimens from animals with intestinal lesions or signs. Furthermore, all of the samples analyzed in this study had previously been screened for CPV-2, CadV1 and CadV2, canine coronavirus, CDV, caliciviruses and CaKoV [10, 14], and none of the protoparvovirus-positive animals had mixed infections.

To date, it is unclear whether members of this novel species of parvoviruses play a role in enteric or extra-enteric diseases in carnivores. Molecular screening of canine stool samples from dogs with or without enteritis has revealed that BuVs are a common component of the faecal virome,

although there was no significant difference in terms of prevalence between dogs with and without enteric signs. However, screening of canine respiratory samples revealed a positive association between the presence of the virus and respiratory signs [3]. More recently, canine BuVs that were closely genetically related to those identified in Italian and Hungarian canine samples were detected in serum specimens from dogs in China. Of interest, the positive sera were collected from animals showing signs of CIRDC [19]. Conversely, this association was not observed when testing nasal and oropharyngeal swabs of cats with or without signs of URDC [7], although the virus was about 5 to 6 times more common in respiratory samples (9.5% to 12.2%) than in diarrhoeic specimens (2.2%) [7]. Accordingly, it has been hypothesized that these protoparvoviruses may have a preferential tropism for the respiratory tract [3, 7]. In our study, we were able to identify the virus in the enteric tract of wild animals. It is therefore possible that faecal shedding of BuVs represents a strategy of virus persistence in animal populations.

In conclusion, the molecular data collected in our study demonstrate that similar BuVs circulate in dogs, cats and wild canids, extending the host range of these BuVs. This finding is intriguing, as it mirrors what has been observed over several decades of investigations for carnivore protoparvovirus 1 [8, 10, 20].

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Compliance with ethical standards

Conflict of interest All authors declare that there are no financial or other relationships that might lead to a conflict of interest. All authors have seen and approved the manuscript and have contributed significantly to the work.

Ethics standards All applicable international, national, and/or institutional guidelines for the care and use of animals were followed.

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