



This form should be used for all taxonomic proposals. Please complete all those modules that are applicable (and then delete the unwanted sections). For guidance, see the notes written in blue and the separate document "Help with completing a taxonomic proposal"

Please try to keep related proposals within a single document; you can copy the modules to create more than one genus within a new family, for example.

MODULE 1: **TITLE, AUTHORS, etc**

Code assigned:	2016.014aS	(to be completed by ICTV officers)
Short title: Create 2 new species (<i>Parechovirus C</i> , <i>Parechovirus D</i>) in the genus <i>Parechovirus</i> (e.g. 6 new species in the genus <i>Zetavirus</i>)		
Modules attached (modules 1 and 11 are required)	2 <input checked="" type="checkbox"/> 3 <input type="checkbox"/> 4 <input type="checkbox"/> 5 <input type="checkbox"/> 6 <input type="checkbox"/> 7 <input type="checkbox"/> 8 <input type="checkbox"/> 9 <input type="checkbox"/> 10 <input type="checkbox"/> 11 <input checked="" type="checkbox"/>	

Author(s):

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List the ICTV study group(s) that have seen this proposal:

A list of study groups and contacts is provided at <http://www.ictvonline.org/subcommittees.asp> . If in doubt, contact the appropriate subcommittee chair (fungal, invertebrate, plant, prokaryote or vertebrate viruses)

Picornaviridae Study Group

ICTV Study Group comments (if any) and response of the proposer:

Date first submitted to ICTV:

15/06/2016

Date of this revision (if different to above):

ICTV-EC comments and response of the proposer:

MODULE 2: **NEW SPECIES**

creating and naming one or more new species.

If more than one, they should be a group of related species belonging to the same genus. All new species must be placed in a higher taxon. This is usually a genus although it is also permissible for species to be “unassigned” within a subfamily or family. Wherever possible, provide sequence accession number(s) for **one** isolate of each new species proposed.

Code	2016.014aS	(assigned by ICTV officers)
To create 2 new species within:		
Genus:	<i>Parechovirus</i>	Fill in all that apply. • If the higher taxon has yet to be created (in a later module, below) write “ (new) ” after its proposed name. • If no genus is specified, enter “ unassigned ” in the genus box.
Subfamily:		
Family:	<i>Picornaviridae</i>	
Order:	<i>Picornavirales</i>	
Name of new species:	Representative isolate: (only 1 per species please)	GenBank sequence accession number(s)
<i>Parechovirus C</i>	Sebokele virus 1 An/B/1227/d	HF677705
<i>Parechovirus D</i>	ferret parechovirus MpPeV1	KF006989

<p>Reasons to justify the creation and assignment of the new species:</p> <ul style="list-style-type: none"> Explain how the proposed species differ(s) from all existing species. <ul style="list-style-type: none"> If species demarcation criteria (see module 3) have previously been defined for the genus, explain how the new species meet these criteria. If criteria for demarcating species need to be defined (because there will now be more than one species in the genus), please state the proposed criteria. Further material in support of this proposal may be presented in the Appendix, Module 11
<p>Sebokele virus and ferret parechovirus are new picornaviruses. Sebokele virus was isolated by intracerebral inoculation of suckling mice from the crushed organs of African wood mice (<i>Hylomyscus sp.</i>) collected in Botambi, Central African Republic, in 1972. Ferret parechovirus was detected in rectal swabs of a household ferret (<i>Mustela putorius furo</i>) collected in the Netherlands.</p> <p>Both viruses share significant similarities to <i>Parechovirus B1</i>, i.e.</p> <p>(i) the Ljungan virus genome layout (compare Appendix Figure 1): $VP_g+5'UTR^{IRES-II}[1AB-1C-1D-2A1^{NPG\downarrow P}/2A2^{H-box/NC}-2B-2C^{Hel}/3A-3B^{VPg}-3C^{Pro}3D^{Pol}]3'UTR-poly(A)$;</p> <p>(ii) significant amino acid identities of capsid proteins (proposed <i>Parechovirus C</i>: >44%, proposed <i>Parechovirus D</i>: >37%) and 3CD (proposed <i>Parechovirus C</i>: >45%, proposed <i>Parechovirus D</i>: >40%) protein (compare Appendix Tables 1 and 2)</p> <p>(iii) clustering with <i>Parechovirus</i> in phylogenetic trees (compare Appendix Figures 2-5).</p>

MODULE 11: **APPENDIX**: supporting material

additional material in support of this proposal

References:

Smits SL, Raj VS, Oduber MD, Schapendonk CME, Bodewes R, Provacia L, Stittelaar KJ, Osterhaus ADME, Haagmans BL. 2013. Metagenomic analysis of the ferret fecal viral flora. PLoS One 8(8):e71595.

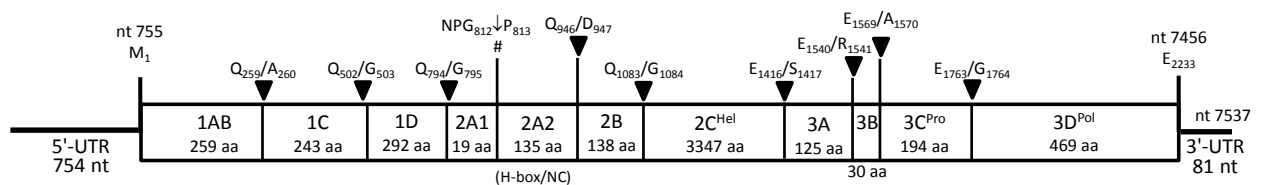
Joffret ML, Bouchier C, Grandadam M, Zeller H, Maufrais C, Bourhy H, Despres P, Delpeyroux F, Dacheux L. 2013. Genomic characterization of Sebokele virus 1 (SEBV1) reveals a new candidate species among the genus *Parechovirus*. J Gen Virol 94:1547-1553.

Annex:

Include as much information as necessary to support the proposal, including diagrams comparing the old and new taxonomic orders. The use of Figures and Tables is strongly recommended but direct pasting of content from publications will require permission from the copyright holder together with appropriate acknowledgement as this proposal will be placed on a public web site. For phylogenetic analysis, try to provide a tree where branch length is related to genetic distance.

Genome organization:

Proposed: *Parechovirus C*, Sebokelevirus [An/B/1227/d], GenBank acc. no. HF677705



Proposed: *Parechovirus D*, ferret parechovirus [MpPeV1], GenBank acc. no. KF006989

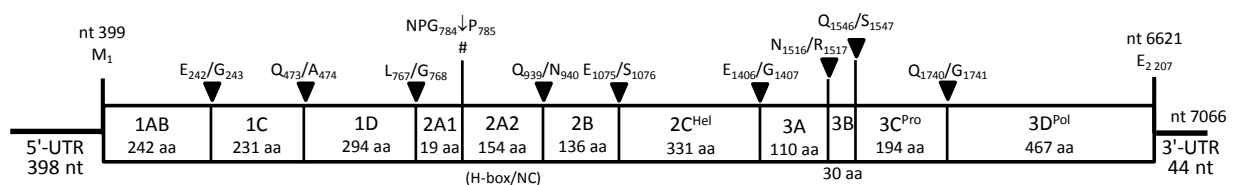


Figure 1: Schematic depiction of the parechovirus C and D genomes (*top*: Sebokele virus [An/B/1227/d], *below*: ferret parechovirus [MpPeV1]). The open reading frames are indicated by boxes. Positions of putative nt and aa cleavage sites and the lengths of the deduced proteins are shown as proposed by Smits et al., 2013. Triangles (▼) indicate the putative 3C^{pro} cleavage sites; the hash (#) indicates the ribosomal skipping site at the NPG↓P motif.

Fig. 2
P1

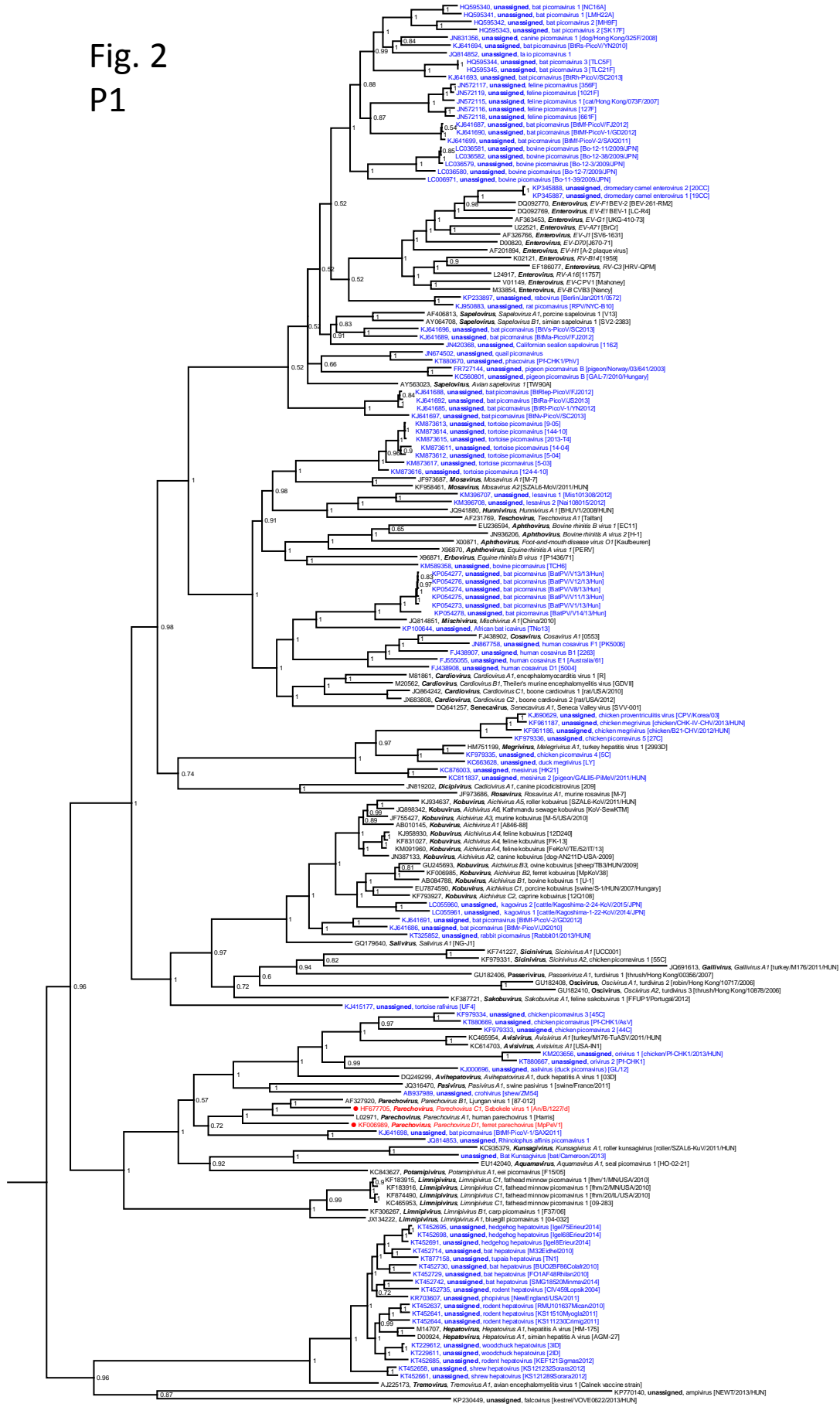


Figure 2: Phylogenetic analysis of picornavirus P1 gene region using Bayesian tree inference (MrBayes 3.2). 178 picornavirus sequences were retrieved from GenBank. Presented are GenBank accession numbers, *genus names*, *species names* and *types*. If available, designations of isolates are given in square brackets. Yet unassigned viruses are printed in blue. Proposed names are printed in red and indicated by a dot (●). Numbers at nodes indicate posterior probabilities obtained after 6,000,000 generations. The optimal substitution model (GTR+G+I) was determined with MEGA 5. The scale indicates substitutions/site.

Fig. 3
3CD

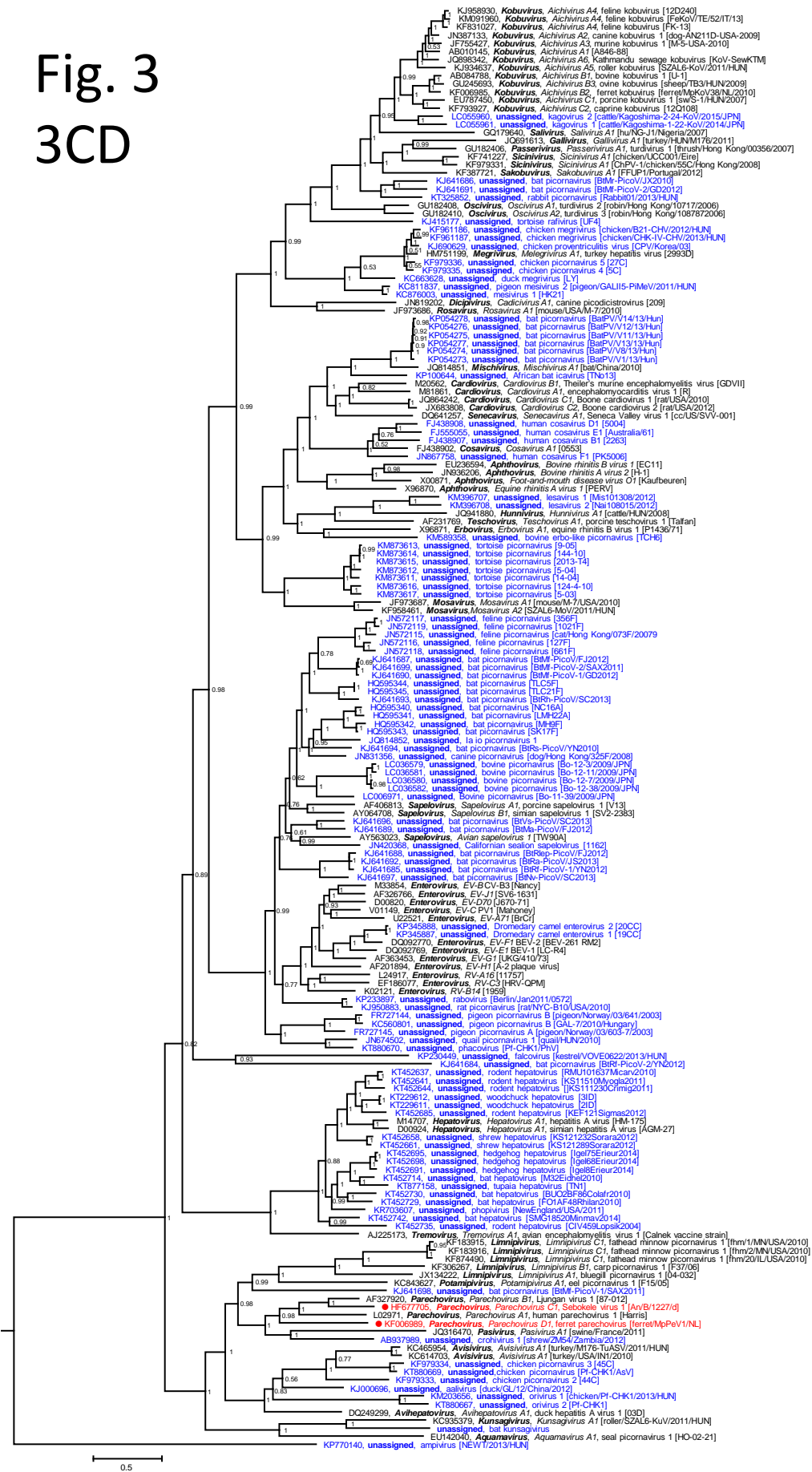


Figure 3: Phylogenetic analysis of picornavirus 3CD gene region using Bayesian tree inference (MrBayes 3.2). 178 sequences were retrieved from GenBank. Presented are GenBank accession numbers, *genus names*, *species names* and *types*. If available, common names and designations of isolates [in square brackets] are given. Proposed names are printed in red and indicated by a dot (●). Numbers at nodes indicate posterior probabilities obtained after 4,750,000 generations. The optimal substitution model (GTR+G+I) was determined with MEGA 5. The scale indicates substitutions/site.

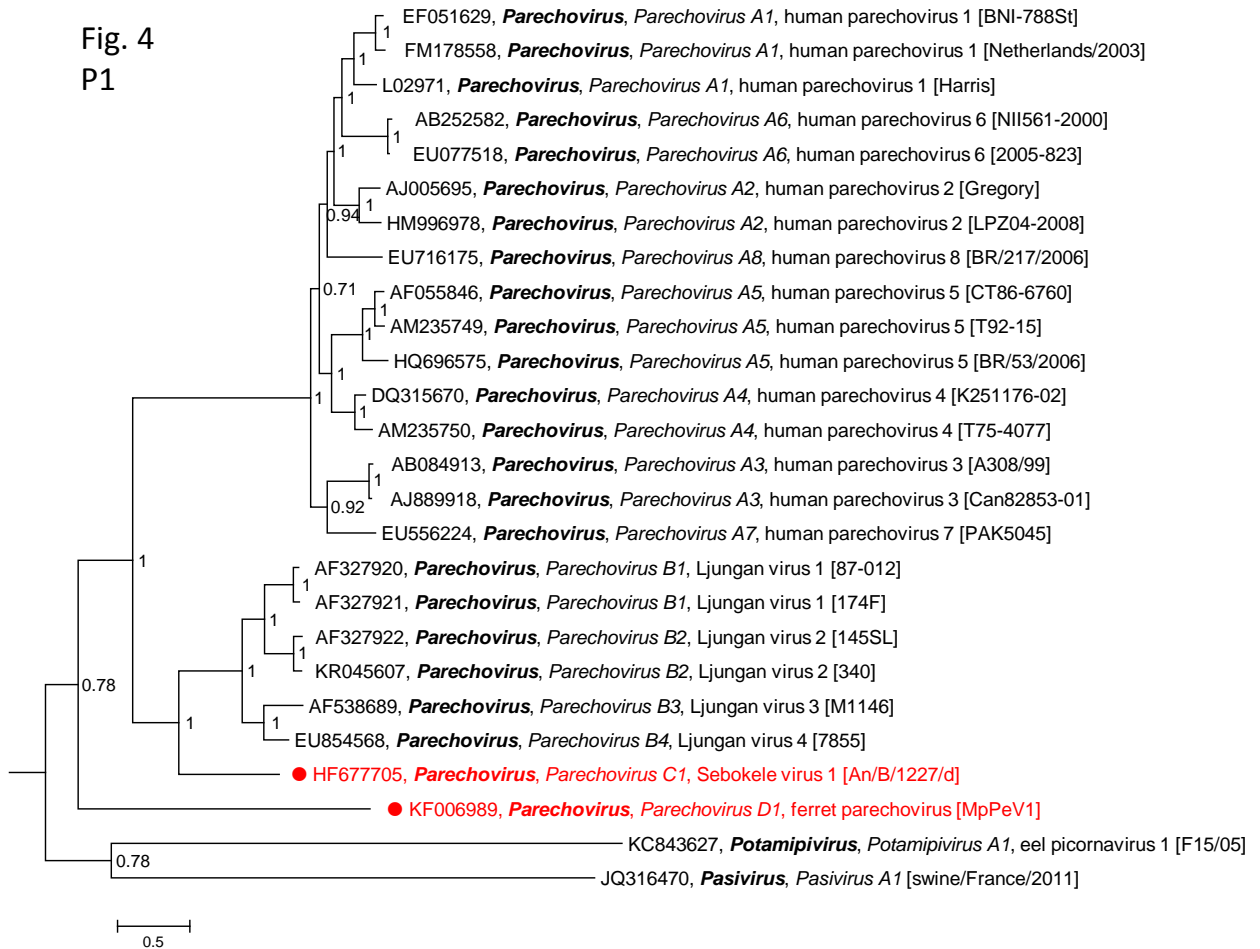


Figure 4: Phylogenetic analysis of parechovirus P1 gene region using Bayesian tree inference (MrBayes 3.2). 26 sequences were retrieved from GenBank. Presented are GenBank accession numbers, *genus names*, *species names* and *types*. If available, common names and designations of isolates [in square brackets] are given. Proposed names are printed in red and indicated by a dot (●). Numbers at nodes indicate posterior probabilities obtained after 1,000,000 generations. The optimal substitution model (GTR+G) was determined with MEGA 5. The scale indicates substitutions/site.

Fig. 5
3CD

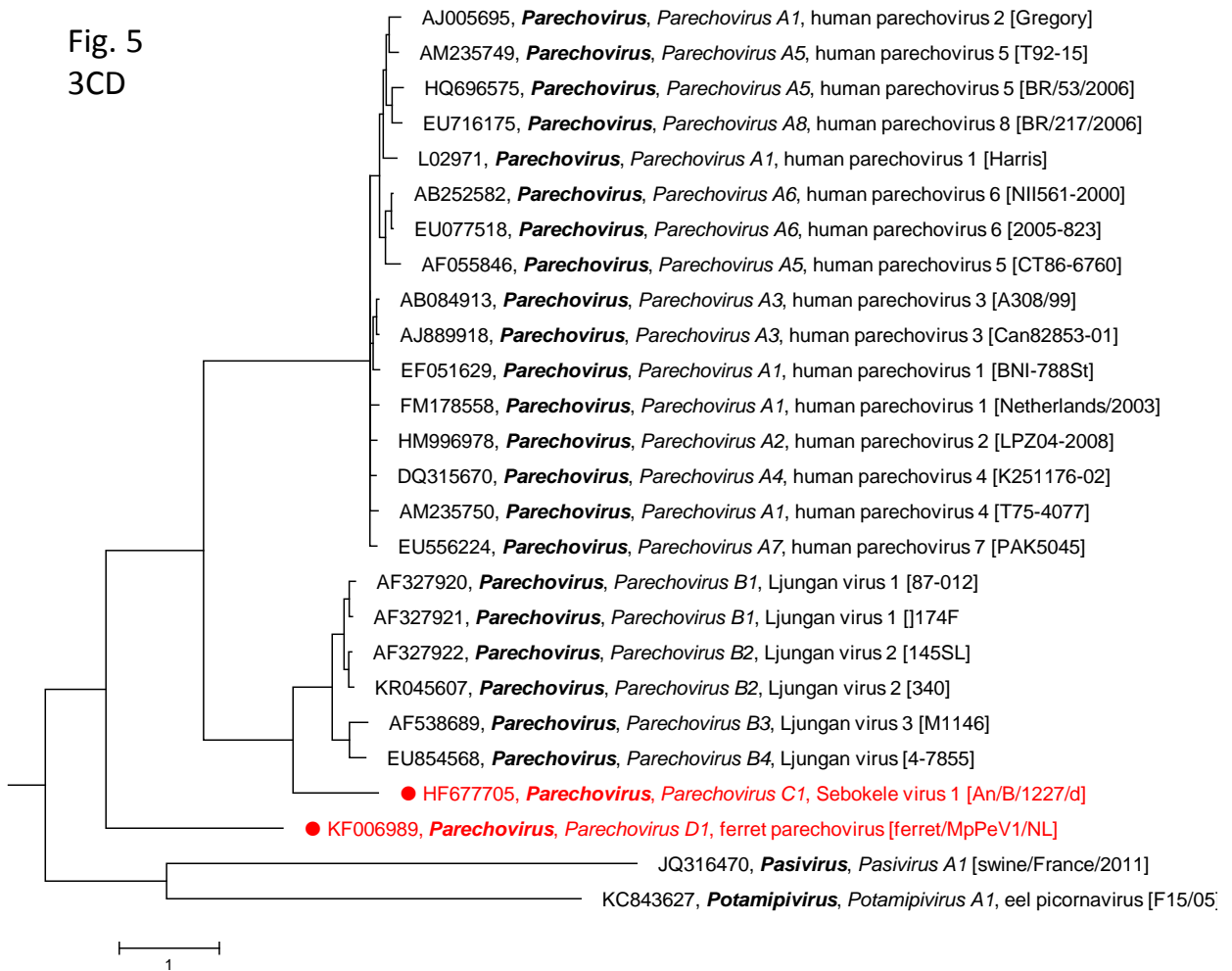


Figure 5: Phylogenetic analysis of parechovirus 3CD gene region using Bayesian tree inference (MrBayes 3.2). 26 sequences were retrieved from GenBank. Presented are GenBank accession numbers, **genus names**, *species names* and *types*. If available, common names and designations of isolates [in square brackets] are given. Yet unassigned viruses are printed in blue. Proposed names are printed in red and indicated by a dot (●). Numbers at nodes indicate posterior probabilities obtained after 3,000,000 generations. The optimal substitution model (GTR+G) was determined with MEGA 5. The scale indicates substitutions/site.

Table 1: Estimates of evolutionary divergence between P1 aa sequences:

```
[ 1] PeV-A1 L02971_Human_parechovirus_1_Harris
[ 2] PeV-A1 EF051629_Human_parechovirus_1_isolate_BNI-788St
[ 3] PeV-A1 FM178558_Human_parechovirus_1_Netherlands/2003
[ 4] PeV-A2 AJ005695_Human_parechovirus_2_Gregory
[ 5] PeV-A2 HM996978_Human_parechovirus_2_strain_LPZ04-2008
[ 6] PeV-A3 AB084913_Human_parechovirus_3_strain_A308/99
[ 7] PeV-A3 AJ889918_Human_parechovirus_3_isolate_Can82853-01
[ 8] PeV-A4 DQ315670_Human_parechovirus_4_isolate_K251176-02
[ 9] PeV-A4 AM235750_Human_parechovirus_4_strain_T75-4077
[10] PeV-A5 AF055846_Human_parechovirus_5_strain_CT86-6760
[11] PeV-A5 AM235749_Human_parechovirus_5_strain_T92-15
[12] PeV-A5 HQ696575_Human_parechovirus_5_strain_BR/53/2006
[13] PeV-A6 AB252582_Human_parechovirus_6_strain_NII561-2000
[14] PeV-A6 EU077518_Human_parechovirus_6_isolate_2005-823
[15] PeV-A7 EU556224_Human_parechovirus_7_PAK5045
[16] PeV-A7 EU716175_Human_parechovirus_8_isolate_BR/217/2006

[17] PeV-B1 AF327920_Ljungan_virus_1_87-012
[18] PeV-B1 AF327921_Ljungan_virus_1_strain_174F
[19] PeV-B2 AF327922_Ljungan_virus_2_strain_145SL
[20] PeV-B2 KR045607_Ljungan_virus_2_strain_340
[21] PeV-B3 AF538689_Ljungan_virus_3_M1146
[22] PeV-B4 EU854568_Ljungan_virus_4-7855

[23] PeV-C1 HF677705_Sebokele_virus_1_An/B/1227/d

[24] PeV-D1 KF006989_Ferret_parechovirus_isolate_MpPeV1

[25] PoV-A1 EelPV_F15-05
[26] PaV-A1 JQ316470_Swine_pasivirus_1
```

	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	
[1] PeV-A1	0.0000																										
[2] PeV-A1	0.0852	0.0000																									
[3] PeV-A1	0.0865	0.0220	0.0000																								
[4] PeV-A2	0.1853	0.1701	0.1740	0.0000																							
[5] PeV-A2	0.1824	0.1686	0.1712	0.0453	0.0000																						
[6] PeV-A3	0.2516	0.2634	0.2660	0.2643	0.2575	0.0000																					
[7] PeV-A3	0.2464	0.2595	0.2634	0.2643	0.2549	0.0272	0.0000																				
[8] PeV-A4	0.2199	0.2137	0.2163	0.2176	0.2083	0.2584	0.2532	0.0000																			
[9] PeV-A4	0.2199	0.2254	0.2241	0.2267	0.2173	0.2610	0.2545	0.0245	0.0000																		
[10] PeV-A5	0.2408	0.2319	0.2383	0.2578	0.2406	0.2737	0.2685	0.1869	0.1985	0.0000																	
[11] PeV-A5	0.2382	0.2396	0.2435	0.2655	0.2484	0.2815	0.2763	0.1959	0.2010	0.0410	0.0000																
[12] PeV-A5	0.2546	0.2468	0.2494	0.2610	0.2516	0.2640	0.2627	0.1938	0.1990	0.0542	0.0606	0.0000															
[13] PeV-A6	0.1932	0.2052	0.2026	0.2195	0.2192	0.2660	0.2608	0.2306	0.2332	0.2642	0.2630	0.2714	0.0000														
[14] PeV-A6	0.1879	0.2000	0.1987	0.2156	0.2153	0.2647	0.2595	0.2280	0.2306	0.2604	0.2591	0.2675	0.0103	0.0000													
[15] PeV-A7	0.2573	0.2542	0.2595	0.2656	0.2588	0.2250	0.2250	0.2601	0.2666	0.2770	0.2809	0.2738	0.2881	0.2855	0.0000												
[16] PeV-A8	0.2275	0.2196	0.2235	0.2389	0.2360	0.2732	0.2680	0.2451	0.2516	0.2608	0.2673	0.2614	0.2490	0.2477	0.2588	0.0000											
[17] PeV-B1	0.5323	0.5267	0.5309	0.5350	0.5288	0.5234	0.5275	0.5369	0.5437	0.5402	0.5437	0.5370	0.5450	0.5436	0.5386	0.5268	0.0000										
[18] PeV-B1	0.5323	0.5267	0.5322	0.5336	0.5274	0.5234	0.5275	0.5383	0.5451	0.5416	0.5451	0.5384	0.5436	0.5422	0.5358	0.5254	0.0039	0.0000									
[19] PeV-B2	0.5378	0.5364	0.5391	0.5405	0.5370	0.5344	0.5358	0.5410	0.5451	0.5498	0.5505	0.5425	0.5490	0.5477	0.5441	0.5323	0.1539	0.1526	0.0000								
[20] PeV-B2	0.5351	0.5336	0.5364	0.5391	0.5356	0.5317	0.5331	0.5396	0.5437	0.5484	0.5492	0.5411	0.5463	0.5450	0.5413	0.5296	0.1526	0.1513	0.0079	0.0000							
[21] PeV-B3	0.5378	0.5322	0.5364	0.5487	0.5466	0.5358	0.5399	0.5314	0.5355	0.5539	0.5560	0.5479	0.5463	0.5463	0.5441	0.5378	0.2648	0.2635	0.2582	0.2582	0.0000						
[22] PeV-B4	0.5158	0.5185	0.5226	0.5350	0.5260	0.5152	0.5193	0.5150	0.5178	0.5348	0.5383	0.5315	0.5286	0.5286	0.5262	0.5172	0.2582	0.2582	0.2372	0.2372	0.1312	0.0000					
[23] PeV-C1	0.5467	0.5369	0.5396	0.5452	0.5389	0.5461	0.5447	0.5374	0.5416	0.5450	0.5429	0.5431	0.5373	0.5373	0.5559	0.5384	0.4144	0.4144	0.4184	0.4171	0.4177	0.4123	0.0000				
[24] PeV-D1	0.6087	0.6145	0.6159	0.6145	0.6194	0.5910	0.5968	0.6118	0.6176	0.6110	0.6162	0.6093	0.6273	0.6230	0.6041	0.5959	0.5939	0.5939	0.5953	0.5981	0.5961	0.5919	0.6064	0.0000			
[25] PoV-A1	0.7156	0.7208	0.7223	0.7230	0.7204	0.7128	0.7128	0.7139	0.7183	0.7202	0.7257	0.7175	0.7397	0.7353	0.7173	0.7162	0.6871	0.6885	0.6956	0.6956	0.6909	0.6909	0.7086	0.6921	0.0000		
[26] PaV-A1	0.6809	0.6780	0.6809	0.6809	0.6799	0.6667	0.6667	0.6780	0.6794	0.6784	0.6766	0.6742	0.6732	0.6704	0.6809	0.6728	0.6749	0.6749	0.6762	0.6776	0.6799	0.6799	0.6662	0.7028	0.7457	0.0000	

Table. Estimates of Evolutionary Divergence between Sequences

The number of amino acid differences per site from between sequences are shown. The analysis involved 26 amino acid sequences. The coding data was translated assuming a Standard genetic code table. All ambiguous positions were removed for each sequence pair. There were a total of 840 positions in the final dataset. Evolutionary analyses were conducted in MEGA5 [1].

1. Tamura K., Peterson D., Peterson N., Stecher G., Nei M., and Kumar S. (2011). MEGA5: Molecular Evolutionary Genetics Analysis using Maximum Likelihood, Evolutionary Distance, and Maximum Parsimony Methods. Molecular Biology and Evolution 28: 2731-2739.

P1:	intra-typic	observed aa diversity: <9%	⇒	aa identity: >91%
	inter-typic (within species)	observed aa diversity: 17-29%	⇒	aa identity: 71-83%
	between species	observed aa diversity: 41-62%	⇒	aa identity: 38-59%
	between genera	observed aa diversity: >66%	⇒	aa identity: <34%

Table 2: Estimates of evolutionary divergence between 3CD aa sequences:

```

Title:
Description
Analysis
  Analysis ----- Distance Estimation
  Scope ----- Pairs of taxa
Estimate Variance
  Variance Estimation Method ----- None
Substitution Model
  Substitutions Type ----- Amino acid
  Genetic Code Table ----- Standard
  Model/Method ----- p-distance
  Substitutions to Include ----- All
Rates and Patterns
  Rates among Sites ----- Uniform rates
  Pattern among Lineages ----- Same (Homogeneous)
Data Subset to Use
  Gaps/Missing Data Treatment ----- Pairwise deletion
  Codons Included ----- 1st+2nd+3rd+Non-Coding
No. of Sites : 684
d : Estimate

[ 1] PeV-A1 L02971_Parechovirus_HPeV-1_Harris
[ 2] PeV-A1 EF051629_Human_parechovirus_1_isolate_ENI-788St
[ 3] PeV-A1 FM178558_Human_parechovirus_1_Netherlands/2003
[ 4] PeV-A2 AJ005695_Human_parechovirus_2_Gregory
[ 5] PeV-A2 HM996978_Human_parechovirus_2_strain_LFPZ04-2008
[ 6] PeV-A3 AB084913_Human_parechovirus_3_strain_A308/99
[ 7] PeV-A3 AJ889918_Human_parechovirus_3_isolate_Can82853-01
[ 8] PeV-A4 DQ315670_Human_parechovirus_4_isolate_K251176-02
[ 9] PeV-A4 AM235750_Human_parechovirus_4_strain_T75-4077
[10] PeV-A5 AF055846_Human_parechovirus_5_strain_CT86-6760
[11] PeV-A5 AM235749_Human_parechovirus_5_strain_T92-15
[12] PeV-A5 HQ696575_Human_parechovirus_5_strain_BR/53/2006
[13] PeV-A6 AB252582_Human_parechovirus_6_strain_NII561-2000
[14] PeV-A6 EU077518_Human_parechovirus_6_isolate_2005-823
[15] PeV-A7 EU556224_Human_parechovirus_7_PAK5045
[16] PeV-A8 EU716175_Human_parechovirus_8_isolate_BR/217/2006

[17] PeV-B1 AF327920_Parechovirus_LV-1_87-012
[18] PeV-B1 AF327921_Ljungan_virus_1_strain_174F
[19] PeV-B2 AF327922_Ljungan_virus_2_strain_145SL
[20] PeV-B2 KR045607_Ljungan_virus_2_strain_340
[21] PeV-B3 AF538689_Ljungan_virus_3_M1146
[22] PeV-B4 EU854568_Ljungan_virus_4-7855

[23] PeV-C1 HF677705_Ljungan-like_Sebokele_virus_1_strain_An_B_1227_d
[24] PeV-D1 KF006989_Ferretetechovirus_strain_ferret/MpPeV1/NL
[25] PoV-A1 KC843627_Potamipivirus_A1_Eel_picornavirus_F15-05
[26] PaV-A1 JQ316470_Pasivirus_A1_swine/France/2011

```

	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26
[1] PeV-A1	0.0000																									
[2] PeV-A1	0.0377	0.0000																								
[3] PeV-A1	0.0513	0.0392	0.0000																							
[4] PeV-A2	0.0392	0.0498	0.0588	0.0000																						
[5] PeV-A2	0.0362	0.0271	0.0332	0.0422	0.0000																					
[6] PeV-A3	0.0422	0.0271	0.0362	0.0528	0.0241	0.0000																				
[7] PeV-A3	0.0422	0.0256	0.0377	0.0528	0.0226	0.0121	0.0000																			
[8] PeV-A4	0.0377	0.0241	0.0287	0.0452	0.0166	0.0211	0.0226	0.0000																		
[9] PeV-A4	0.0392	0.0271	0.0347	0.0498	0.0256	0.0241	0.0226	0.0196	0.0000																	
[10] PeV-A5	0.0362	0.0377	0.0498	0.0347	0.0332	0.0332	0.0362	0.0317	0.0347	0.0000																
[11] PeV-A5	0.0407	0.0558	0.0618	0.0468	0.0483	0.0543	0.0513	0.0483	0.0513	0.0437	0.0000															
[12] PeV-A5	0.0302	0.0362	0.0422	0.0377	0.0317	0.0332	0.0377	0.0287	0.0332	0.0256	0.0407	0.0000														
[13] PeV-A6	0.0302	0.0362	0.0437	0.0332	0.0287	0.0377	0.0347	0.0302	0.0302	0.0226	0.0407	0.0256	0.0000													
[14] PeV-A6	0.0317	0.0377	0.0452	0.0362	0.0302	0.0392	0.0392	0.0317	0.0347	0.0256	0.0437	0.0271	0.0045	0.0000												
[15] PeV-A7	0.0422	0.0302	0.0392	0.0483	0.0256	0.0287	0.0332	0.0256	0.0317	0.0317	0.0543	0.0377	0.0362	0.0377	0.0000											
[16] PeV-A8	0.0362	0.0437	0.0513	0.0392	0.0377	0.0407	0.0437	0.0332	0.0377	0.0317	0.0452	0.0271	0.0362	0.0392	0.0407	0.0000										
[17] PeV-B1	0.5068	0.4992	0.5083	0.5008	0.5053	0.5038	0.5038	0.5068	0.5008	0.5068	0.5173	0.5038	0.5053	0.5053	0.5083	0.5038	0.0000									
[18] PeV-B1	0.5068	0.4992	0.5068	0.5008	0.5053	0.5038	0.5038	0.5068	0.5008	0.5068	0.5173	0.5023	0.5053	0.5053	0.5098	0.5038	0.0150	0.0000								
[19] PeV-B2	0.5083	0.4992	0.5083	0.5023	0.5053	0.5023	0.5053	0.5068	0.5008	0.5083	0.5189	0.5053	0.5053	0.5053	0.5083	0.5053	0.0180	0.0269	0.0000							
[20] PeV-B2	0.5098	0.5008	0.5098	0.5038	0.5068	0.5038	0.5068	0.5083	0.5023	0.5098	0.5204	0.5068	0.5068	0.5068	0.5098	0.5068	0.0269	0.0344	0.0090	0.0000						
[21] PeV-B3	0.5264	0.5173	0.5294	0.5204	0.5234	0.5204	0.5189	0.5264	0.5189	0.5249	0.5339	0.5264	0.5234	0.5234	0.5249	0.5294	0.1517	0.1517	0.1517	0.1517	0.0000					
[22] PeV-B4	0.5339	0.5279	0.5339	0.5294	0.5309	0.5294	0.5264	0.5324	0.5264	0.5324	0.5385	0.5309	0.5309	0.5309	0.5324	0.5309	0.1517	0.1532	0.1502	0.1502	0.0691	0.0000				
[23] PeV-C1	0.5408	0.5257	0.5408	0.5408	0.5393	0.5332	0.5363	0.5378	0.5317	0.5363	0.5483	0.5423	0.5378	0.5363	0.5347	0.5378	0.3404	0.3373	0.3389	0.3389	0.3479	0.3539	0.0000			
[24] PeV-D1	0.5897	0.5957	0.5957	0.5897	0.5927	0.5897	0.5897	0.5957	0.5897	0.5927	0.5942	0.5973	0.5881	0.5881	0.5897	0.5927	0.5697	0.5697	0.5682	0.5712	0.5833	0.5894	0.5742	0.0000		
[25] PoV-A1	0.7043	0.6982	0.7058	0.7027	0.7027	0.6982	0.7012	0.7027	0.7012	0.7043	0.7043	0.7027	0.7012	0.7012	0.6982	0.7012	0.6550	0.6535	0.6535	0.6550	0.6520	0.6520	0.6505	0.6606	0.0000	
[26] PaV-A1	0.7189	0.7174	0.7236	0.7174	0.7158	0.7158	0.7189	0.7174	0.7158	0.7174	0.7174	0.7143	0.7220	0.7220	0.7174	0.7158	0.6961	0.6977	0.6946	0.6961	0.6930	0.6992	0.6930	0.6817	0.7176	0.0000

Table. Estimates of Evolutionary Divergence between Sequences

The number of amino acid differences per site from between sequences are shown. The analysis involved 26 amino acid sequences. The coding data was translated assuming a Standard genetic code table. All ambiguous positions were removed for each sequence pair. There were a total of 684 positions in the final dataset. Evolutionary analyses were conducted in MEGA5 [1].

1. Tamura K., Peterson D., Peterson N., Stecher G., Nei M., and Kumar S. (2011). MEGA5: Molecular Evolutionary Genetics Analysis using Maximum Likelihood, Evolutionary Distance, and Maximum Parsimony Methods. *Molecular Biology and Evolution* 28: 2731-2739.

3CD:	intra-typic	observed aa diversity: <6%	⇒	aa identity: >94%
	inter-typic (within species)	observed aa diversity: 3-15%	⇒	aa identity: 85-97%
	between species	observed aa diversity: 49-60%	⇒	aa identity: >40%
	between genera	observed aa diversity: >65%	⇒	aa identity: <35%