



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**

<b>Code assigned:</b>	<b>2016.024a-dM</b>	(to be completed by ICTV officers)
<b>Short title:</b> Four (4) new species in one new genus ( <i>Herbevirus</i> ) in the <i>Bunyaviridae</i> (to be renamed <i>Peribunyaviridae</i> ; see 2016.030M), order <i>Bunyavirales</i> (proposed in 2016.030M) (e.g. 6 new species in the genus <i>Zetavirus</i> )		
<b>Modules attached</b> (modules 1 and 11 are required)	6 <input type="checkbox"/> 7 <input type="checkbox"/> 8 <input type="checkbox"/> 9 <input type="checkbox"/> 10 <input type="checkbox"/> 2 <input checked="" type="checkbox"/> 3 <input checked="" type="checkbox"/> 4 <input type="checkbox"/> 5 <input type="checkbox"/>	

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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)

ICTV *Bunyaviridae* Study Group

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

The ICTV *Bunyaviridae* Study Group has seen and discussed this proposal, and agreed to its submission to the ICTV Executive Committee based on votes of support by individual Study Group members or the absence of dissenting votes.

Date first submitted to ICTV:

July 18, 2016

Date of this revision (if different to above):

September 21, 2016

**ICTV-EC comments and response of the proposer:**

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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	<b>2016.024aM</b>	(assigned by ICTV officers)
<b>To create 4 new species within:</b>		
Genus:	<b><i>Herbevirus</i> (NEW)</b>	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:	<b>unassigned</b>	
Family:	<b><i>Peribunyaviridae</i> (RENAMED, see see TP 2016.030M)</b>	
Order:	<b><i>Bunyvirales</i> (NEW, see TP 2016.030M)</b>	
<b>Name of new species:</b>	<b>Representative isolate: (only 1 per species please)</b>	<b>GenBank sequence accession number(s)</b>
<i>Herbert herbevirus</i>	Herbert virus (HEBV) C60/CI/2004	JQ659256–JQ659258
<i>Tai herbevirus</i>	Tai virus (TAIV) F47/CI/2004	KF590572–KF590574
<i>Kibale herbevirus</i>	Kibale virus (KIBV) P07/UG/2008	KF590575–KF590577
<i>Shuangao insect herbevirus 1</i>	Shuāngào insect virus 1 QSA02	KM817739, KM817679, KM817714

**Reasons to justify the creation and assignment of the new species:**

- Explain how the proposed species differ(s) from all existing species.
  - 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

The viruses listed above define a highly diversified monophyletic sister clade to all members of the accepted bunyaviral genus *Orthobunyavirus*. Maximal amino acid identities between proteins of the proposed novel genus *Herbevirus* and orthobunyaviruses range between 12 and 25%.

The mean genetic distances between HEBV, TAIV, and KIBV genome segments to those of orthobunyaviruses and tospoviruses are 71–79% and 81–86%, respectively (similar to the distances measured for between orthobunyavirus and tospovirus segments (81–86%).

HEBV, TAIV, and KIBV genomic segment termini contain only 7 reverse-complementary nucleotides that are identical to those of orthobunyaviruses, which have conserved termini of 10 nt in length.

HEBV, TAIV, and KIBV genomes contain an elongated L segment with a unique 500-nucleotide insertion absent in other bunyavirus genomes.

In contrast to orthobunyaviruses, HEBV, TAIV, and KIBV do not seem to encode NSs or NSm proteins.

The Gc proteins of HEBV, TAIV, and KIBV are truncated by about 480 amino acids at their N termini compared to those of orthobunyaviruses.

Finally, HEBV, TAIV, and KIBV seem to differ in their host range compared to orthobunyaviruses. Orthobunyaviruses replicate in insects and vertebrates. However, the novel viruses replicate only in mosquito cells and not in vertebrate cells. Thus, herbeviruses seem to have host range limited to insects.

Shuāngào insect virus 1 is distantly related to HEBV, TAIV, and KIBV. Whereas the S and L segments of Shuāngào insect virus 1 are similar in length to those of herbeviruses, the M segment is with 4.8 kb 2 kb longer as the M segments of herbeviruses and more similar to M segments of orthobunyaviruses. Shuāngào insect virus 1 was also predicted to encode an NSm protein in a similar coding strategy as found in orthobunyaviruses.

MODULE 3: **NEW GENUS**

creating a new genus

Ideally, a genus should be placed within a higher taxon.

Code	<b>2016.024bM</b>	(assigned by ICTV officers)
<b>To create a new genus within:</b>		
Subfamily:	<b>unassigned</b>	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 family is specified, enter “unassigned” in the family box
Family:	<b><i>Peribunyaviridae</i> (RENAMED, see TP 2016.030M)</b>	
Order:	<b><i>Bunyavirales</i> (NEW, see TP 2016.030M)</b>	

naming a new genus

Code	<b>2016.024cM</b>	(assigned by ICTV officers)
<b>To name the new genus: <i>Herbevirus</i></b>		

Assigning the type species and other species to a new genus

Code	<b>2016.024dM</b>	(assigned by ICTV officers)
<b>To designate the following as the type species of the new genus</b>		
<b><i>Herbert herbevirus</i></b>		Every genus must have a type species. This should be a well characterized species although not necessarily the first to be discovered
The new genus will also contain any other new species created and assigned to it (Module 2) and any that are being moved from elsewhere (Module 7b). <b>Please enter here the TOTAL number of species (including the type species) that the genus will contain:</b>		
<b>4</b>		

**Reasons to justify the creation of a new genus:**

Additional material in support of this proposal may be presented in the Appendix, Module 11

See justification for new species.
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**Origin of the new genus name:**

Derived from <u>Herbert</u> virus.
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**Reasons to justify the choice of type species:**

Herbert virus was the first discovered virus of this clade and is (together with Kibale virus) the best characterized virus of the proposed new genus.
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**Species demarcation criteria in the new genus:**

If there will be more than one species in the new genus, list the criteria being used for species demarcation and explain how the proposed members meet these criteria.

Until further herbeviruses are discovered, we propose to use the same species demarcation criteria for this genus as described for the proposed new genera “ <i>Goukovirus</i> ” and “ <i>Orthoferavirus</i> ” (see separate co-submitted proposals). Species demarcation criteria should be based on a ≈1 kb sequence fragment containing the core polymerase domain (premotif A to motif E) of the third conserved region of the L protein. These motifs can be aligned between all members of the
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proposed order *Bunyavirales* and would allow comparative species demarcation criteria for all genera of the entire family. Moreover, as the motifs are highly conserved between all bunyaviruses, amplification of this genome region from new viruses is facilitated. Species demarcation criteria of other viral families are also based on the replicative genes/domains and have been shown to be suitable criteria.

Species should be defined on the criterion that the  $\approx 1$  kb sequence fragment containing the core polymerase domain (premotif A to motif E) of the third conserved region of the L protein should be less than 90% identical on the amino acid level compared to that of any other described herbevirus.

This <90% aa identity threshold for the core polymerase domain is in agreement with the aa identity values for established bunyavirus species within the five established genera.

MODULE 11: **APPENDIX**: supporting material

additional material in support of this proposal

**References:**

**Li C.X., Shi M., Tian J.H., Lin X.D., Kang Y.J., Chen L.J., Qin X.C. Xu J., Holmes E.C. and Zhang Y.Z. (2015).** Unprecedented genomic diversity of RNA viruses in arthropods reveals the ancestry of negative-sense RNA viruses.  
**Elife** 4: e05378.

**Marklewitz M., Handrick S., Grasse W., Kurth A., Lukashev A., Drosten C., Ellerbrok H., Leendertz F.H., Pauli G., Junglen S.** 2011. Gouleako virus isolated from West African mosquitoes constitutes a proposed novel genus in the family Bunyaviridae.  
**Journal of Virology** 85: 9227-9234.

**Marklewitz M., Zirkel F., Rwego I.B., Heidemann H., Trippner P., Kurth A., Kallies R., Briese T., Lipkin W.I., Drosten C., Gillespie T.R., Junglen S.** 2013. Discovery of a Unique Novel Clade of Mosquito-Associated Bunyaviruses.  
**Journal of Virology** 87: 12850-12865.

**Marklewitz M., Zirkel F., Kurth A., Drosten C., Junglen S.** 2015. Evolutionary and phenotypic analysis of live virus isolates suggests arthropod origin of a pathogenic RNA virus family.  
**Proceedings of the National Academy of Sciences** 112: 7536-41.

**Junglen S. (2016).** Evolutionary origin of pathogenic arthropod-borne viruses — a case study in the family *Bunyaviridae*.  
**Current Opinion in Insect Science** 16: 81-86.

**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.

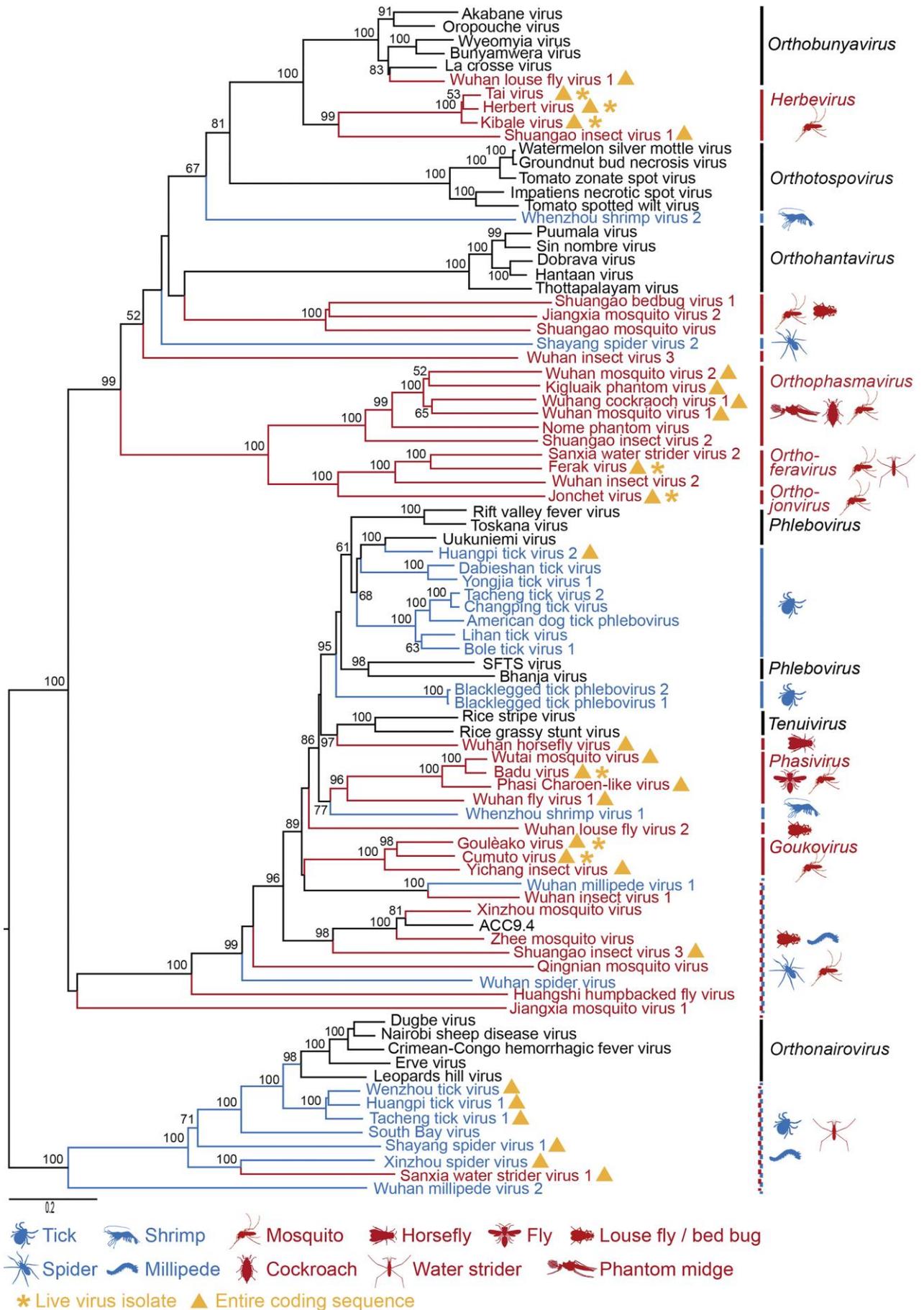


Figure: Phylogenetic relationship of bunyaviruses. Phylogenetic analyses were based on RdRp proteins. Complete RdRp proteins were aligned using MAFFT (E-INS-I algorithm). Alignment columns were stripped to 10% gaps in Geneious. Maximum likelihood (ML) analyses were performed on a 508 amino acid alignment guided by the Blosum62 amino acid substitution matrix with 4 gamma categories and a gamma shape parameter of 1. Confidence testing was performed by 1000 bootstrap replicates. Only bootstrap values over 50 are shown.