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

MODULE 1: TITLE, AUTHORS, etc.

MODULE 1: 111 LE, AUTHORS, etc						
Code assigned:	gned: 2013.020a-dB		(to be completed by ICTV officers)			
Short title: To create a new genus, <i>Pgonelikevirus</i> , within the family <i>Siphoviridae</i>						
(e.g. 6 new species in the genus Modules attached (modules 1 and 9 are required)	Zetavirus)	1 ⊠ 6 □	2 × 7 □	3	4 ☐ 9 ⊠	5 🗌
Author(s) with e-mail address(es) of the proposer:						
Evelien M Adriaenssens, eveli Andrew M Kropinski, kropins Rob Lavigne, rob.lavigne@biv John Nash, john.nash@phac-a	k@queensu.ca w.kuleuven.be	s@gmail.c	<u>om</u>			
List the ICTV study group(s) that have see	n this pro	posal:			
A list of study groups and contact http://www.ictvonline.org/subcomin doubt, contact the appropriate chair (fungal, invertebrate, plant, vertebrate viruses)	mittees.asp . If subcommittee					
ICTV-EC or Study Group co	omments and r	esponse o	of the pro	poser:		
Date first submitted to ICTV: Date of this revision (if different	ent to above):			2013 2014		

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 2013.020aB	(assigned by ICTV officers)
To create 12 new species within:	
Genus: Pgonelikevirus (new) Subfamily: Family: Siphoviridae Order: Caudovirales	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.
And name the new species:	GenBank sequence accession number(s) of reference isolate:
Mycobacterium phage Pg1	AF547430
Mycobacterium phage oline	JN192463
Mycobacterium phage rosebush	AY129334
Mycobacterium phage pipefish	DQ398049
Mycobacterium phage gadjet	JN698992
Mycobacterium phage athena	JN699003
Mycobacterium phage cooper	DQ398044
Mycobacterium phage chrisnmich	JF704094
Mycobacterium phage stinger	JN699011
Mycobacterium phage nigel	EU770221
Mycobacterium phage zemanar	JF704104
Mycobacterium phage acadian	JN699007

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.
 - o 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 9

We have chosen 95% DNA sequence identity as the criterion for demarcation of species in this new genus. Each of the proposed species differs from the others with more than 5% at the DNA level as confirmed with the EMBOSS Stretcher algorithm.

MODULE 3: NEW GENUS

creating a new genus

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

Code	201	3.020bB	(assigned by I	CTV officers)
To create a	a new	genus within:		Fill in all that apply.
Subfar	nily:			If the higher taxon has yet to be created
Fan	nily:	Siphoviridae		(in a later module, below) write "(new)" after its proposed name.
Or	rder:	Caudovirales		 If no family is specified, enter "unassigned" in the family box

naming a new genus

Code	2013.020cB	(assigned by ICTV officers)
To name the	he new genus: Pgonelikevirus	

Assigning the type species and other species to a new genus

Code	2013.020dB	(assigned by ICTV officers)		
To designa	ate the following as the type sp	pecies of the new genus		
Mycobacte	rium phage PG1	Every genus must have a type species. This should be a well characterized species although not necessarily the first to be discovered		
are being m		Please enter here the TOTAL number of species us will contain:		
12				

Reasons to justify the creation of a new genus:

Additional material in support of this proposal may be presented in the Appendix, Module 9

This genus was originally recognized by the Mycobacteriophage group ($\underline{www.phagesdb.org}$) as belonging to cluster B. Phages belonging to this genus share a comparable genome size (67 – 71 kb), a comparable GC content (66 – 69%), and a circularly permuted genome. Members of this genus also have a comparable morphology, with an isometric head and a long, non-contractile tail (Figure 1).

A ClustalW analysis of the complete genomes of this genus with all other *Mycobacterium* phages belonging to the *Siphoviridae* reveals that this genus is a clearly separate group (Figure 2s and 3). We propose a shared protein content of at least 40% with the type phage, *Mycobacterium phage PgI*. Based on the recognizable groups in Figure 2 within this genus, we performed a CoreGenes 3.5 analysis [1–3] with selected phages from each group against PG1 as the reference phage (Table 1). The CoreGenes analysis was also performed against the type species of other proposed genera of siphoviruses infecting *Mycobacterium* and the shared protein content was consistently below 40% (data not shown).

Origin of the new genus name:

Mycobacterium phage PG1

Reasons to justify the choice of type species:

The genus *Pgonelikevirus* is named after the first isolated and sequenced phage of this group, *Mycobacterium* phage PG1 (http://phagesdb.org/phages/PG1/). The number 1 in the phage name was replaced with 'one' to avoid confusion with the letter '1' that follows in the genus name, but not with 'una' to avoid confusion with the proposed genus *Pbiunalikevirus*.

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.

We have chosen 95% DNA sequence identity as the criterion for demarcation of species in this new genus. Each of the proposed species differs from the others with more than 5% at the DNA level as confirmed with the EMBOSS Stretcher algorithm.

As visible in Figure 2, there are many more isolates than proposed species. Table 2 lists the isolates belonging to the same species.

MODULE 9: APPENDIX: supporting material

additional material in support of this proposal

References:

- 1. Mahadevan P, King JF, Seto (2009) Data mining pathogen genomes using GeneOrder and CoreGenes and CGUG: gene order, synteny and in silico proteomes. Int J Comput Biol Drug Des 2: 100–114.
- 2. Mahadevan P, King JF, Seto D (2009) CGUG: in silico proteome and genome parsing tool for the determination of "core" and unique genes in the analysis of genomes up to ca. 1.9 Mb. BMC Res Notes 2: 168. doi:10.1186/1756-0500-2-168.
- 3. Zafar N, Mazumder R, Seto D (2002) CoreGenes: A computational tool for identifying and cataloging "core" genes in a set of small genomes. BMC Bioinformatics 3: 12. doi:10.1186/1471-2105-3-12.
- 4. Darling AE, Mau B, Perna NT (2010) progressiveMauve: multiple genome alignment with gene gain, loss and rearrangement. PLoS One 5: e11147. doi:10.1371/journal.pone.0011147.

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.

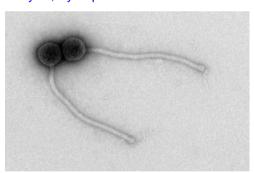


Figure 1: EM picture of Mycobacterium phage PG1 (http://phagesdb.org/media/emPics/PG1.tif).

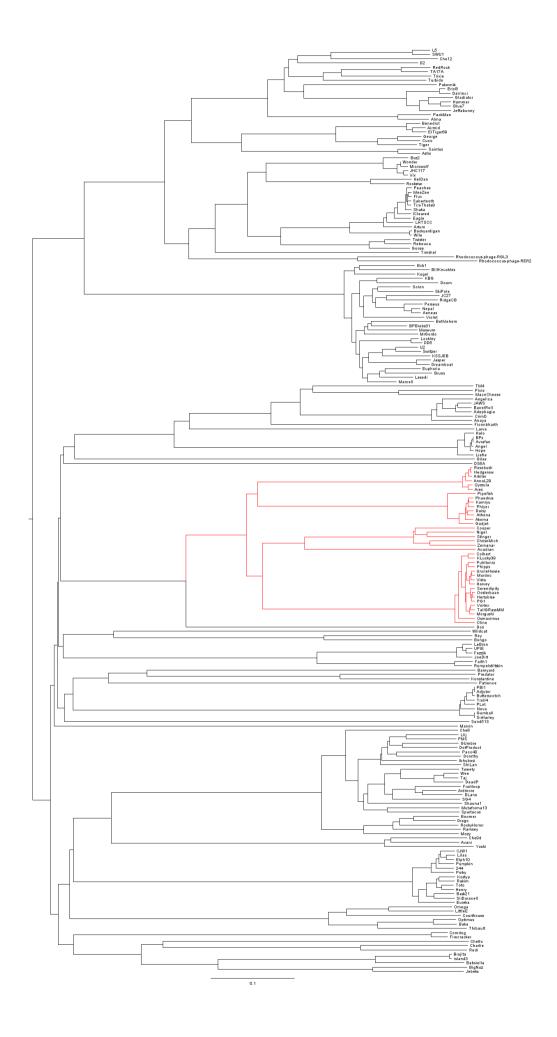


Figure 2: ClustalW phylogenetic tree (NJ) of complete genomes of all Mycobacterium siphoviruses in the NCBI database in November 2012. The proposed genus is colored in red.

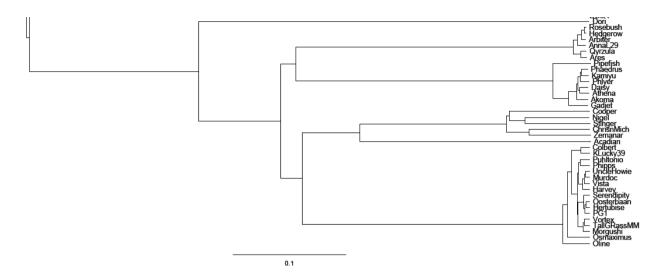


Figure 3: Clustal W phylogenetic tree of complete genomes of the isolates belonging to the genus Pgonelikevirus, excerpt from Figure 2.

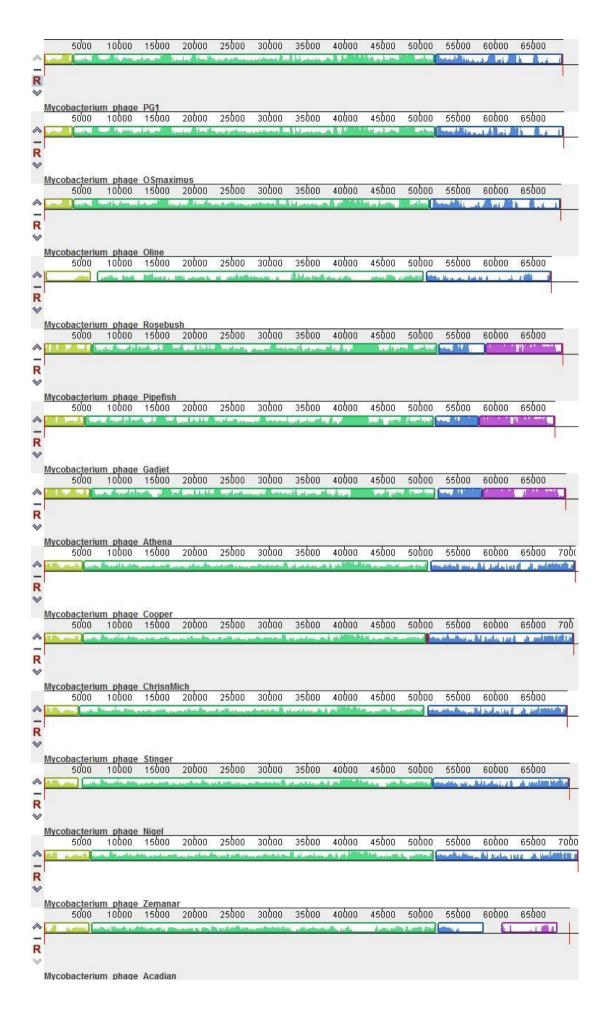


Figure 4: progressiveMauve alignment of the phage genomes belonging to the proposed genus *Pgonelikevirus* [4]. Colored blocks indicate the regions of 1 to 1 best alignment with rearrangement breakpoints in a different random color. The degree of sequence similarity between regions is given by a similarity plot within the colored blocks with the height of the plot proportional to the average nucleotide identity.

Table 1: CoreGenes 3.5 and Stretcher analysis of selected phages with the type phage of the genus, *Mycobacterium phage PG1*.

Phage Name	% proteins in common with PG1	% DNA similarity with PG1
Mycobacterium phage oline	95.0	93.8
Mycobacterium phage rosebush	48.0	53.5
Mycobacterium phage pipefish	49.0	53.5
Mycobacterium phage gadjet	50.0	54.2
Mycobacterium phage athena	51.0	53.9
Mycobacterium phage cooper	62.0	57.4
Mycobacterium phage chrisnmich	63.0	57.4
Mycobacterium phage stinger	61.0	57.6
Mycobacterium phage nigel	60.0	57.2
Mycobacterium phage acadian	61.0	56.7
Mycobacterium phage dori	33.0	56.9

Table 2: Isolates belonging to the same species based on a higher than 95% DNA identity calculated with EMBOSS Stretcher.

Species Name	Phage isolates belonging to species (>95% DNA identity)
Mycobacterium phage Pg1	PG1, Colbert, Klucky39, Puhltonio, Phipps, UncleHowie,
	Murdoc, Vista, Serendipity, Oosterbaan, Hertubise,
	TallGrassMM, Morgushi, OSMaximus
Mycobacterium phage rosebush	Rosebush, Hedgerow, Arbiter, AnnaL29, Qyrzula, Ares
Mycobacterium phage athena	Athena, Akoma, Phaedrus, Daisy, Phlyer, Kamiyu