This form should be used for all taxonomic proposals. Please complete all those modules that are applicable (and then delete the unwanted sections).



example.

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:	2013.008	Ra-dB		(to be co	mpleted by	ICTV
Short title: To create a new genus, Che8likevirus, within the family Siphoviridae (e.g. 6 new species in the genus Zetavirus) Modules attached $1 \boxtimes 2 \boxtimes 3 \boxtimes 4 \sqsubseteq 5 \sqsubseteq 6 \boxtimes 1$ (modules 1 and 9 are required) $1 \boxtimes 2 \boxtimes 3 \boxtimes 4 \sqsubseteq 5 \sqsubseteq 6 \boxtimes 7 \boxtimes 8 \sqsubseteq 9 \boxtimes 1$					5 🗌	
Author(s) with e-mail address	Author(s) with e-mail address(es) of the proposer:					
Evelien M Adriaenssens, eveli Andrew M Kropinski, akropin Rob Lavigne, rob.lavigne@biv	s@uoguelph.ca		<u>com</u>			
List the ICTV study group(s) that have seen this proposal:						
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 comments and response of the proposer:						
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.

	ccession number(s) for one isolate of each new species proposed.		
Code 2013.008aB (assigned by ICTV officers)			
To create 28 new species within:			
	•	Fill in all that apply.	
Genus:	Che8likevirus (new)	If the higher taxon has yet to be	
Subfamily:	Citeotite (iiew)	created (in a later module, below) write	
Family:	Siphoviridae	"(new)" after its proposed name. • If no genus is specified, enter	
Order:	Caudovirales	"unassigned" in the genus box.	
And name the	e new species:	GenBank sequence accession	
		number(s) of reference isolate:	
Mycobacterium		AY129330	
Mycobacterium		DQ398050	
	phage mutaforma13	JN020142	
_	phage dotproduct	JN859129	
Mycobacterium		JN398368	
Mycobacterium		DQ398045	
	phage spartacus	JQ300538	
Mycobacterium		FJ174692	
Mycobacterium		JF937098	
Mycobacterium		EF536069	
Mycobacterium phage taj		JX121091	
Mycobacterium phage wee		HQ728524	
Mycobacterium phage dorothy		JX411620	
Mycobacterium phage deadp		JN698996	
Mycobacterium phage SG4		JN699012	
Mycobacterium phage boomer		EU816590	
Mycobacterium		JF937093	
_	phage ardmore	GU060500	
l '	phage fruitloop	FJ174690	
Mycobacterium		JN020143	
Mycobacterium		JN542517	
Mycobacterium		FJ174693	
	phage rockyhorror	JF704117	
Mycobacterium		JF937102	
Mycobacterium phage shauna1		JN020141	
Mycobacterium phage Che9d		AY129336	
Mycobacterium phage avani		JQ809702	
Mycobacterium phage yoshi		JF704115	

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

- Explain how the proposed species differ(s) from all existing species.
 - o 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	3.008bB	(assigned by ICTV officers)		
To create	a new	genus within:		Fill in all that apply.	
Subfai	mily:			If the higher taxon has yet to be created	
Fai	mily:	Siphoviridae		(in a later module, below) write "(new)" after its proposed name.	
О	rder:	Caudovirales		If no family is specified, enter "unassigned" in the family box	

naming a new genus

Code	2013.008cB	(assigned by ICTV officers)
To name t	he new genus: Che8likevirus	

Assigning the type species and other species to a new genus

Code	2013.008dB	(assigned by ICTV officers)			
To design	To designate the following as the type species of the new genus				
Mycobacte	erium phage Che8	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). Please enter here the TOTAL number of species (including the type species) that the genus will contain:					
28					

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 (www.phagesdb.org) as belonging to cluster F. Phages belonging to this genus share a comparable genome size (56 – 58 kb), a comparable GC content (~61.5%), and defined genome ends with comparable 3' overhangs. 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 (Figures 2 and 3). We propose a shared protein content of at least 40% with the type phage, *Mycobacterium phage Che8*. Based on the phylogenetic tree (Figures 2 and 3), two groupings were visible within the genus and we performed a CoreGenes 3.5 analysis [1–3] with all phages to confirm their presence in the same genus (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 Che8

Reasons to justify the choice of type species:

The genus *Che8likevirus* is named after one of the first isolated and sequenced phages of this group, *Mycobacterium* phage Che8 [4].

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.

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. Pedulla ML, Ford ME, Houtz JM, Karthikeyan T, Wadsworth C, et al. (2003) Origins of highly mosaic mycobacteriophage genomes. Cell 113: 171–182.
- 5. 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.

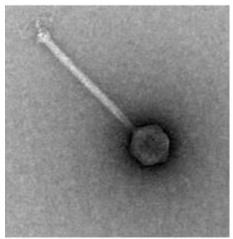


Figure 1: EM picture of phage Che8, the type species of the genus *Che8likevirus* (http://phagesdb.org/media/emPics/Che8.tif).

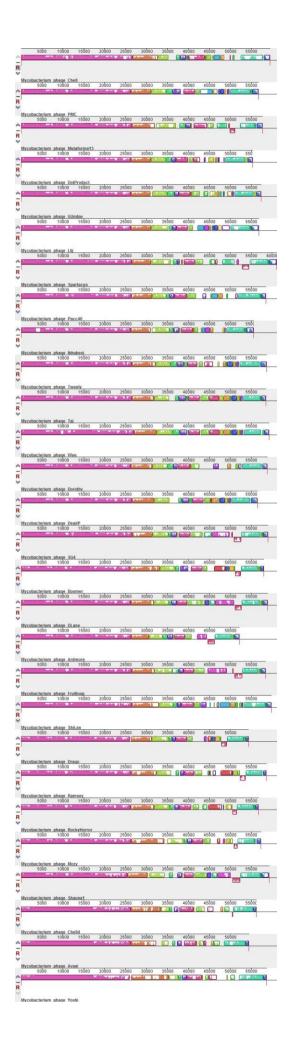


Figure 2: progressiveMauve alignment of the phage genomes belonging to the proposed genus (phages are depicted in the same order as in Table 1) [5]. Colored blocks indicate the regions of 1 to 1 best alignment with rearrangement breakpoints in a different random color. The degree of sequence similarity between regions is given by a similarity plot within the colored blocks with the height of the plot proportional to the average nucleotide identity.

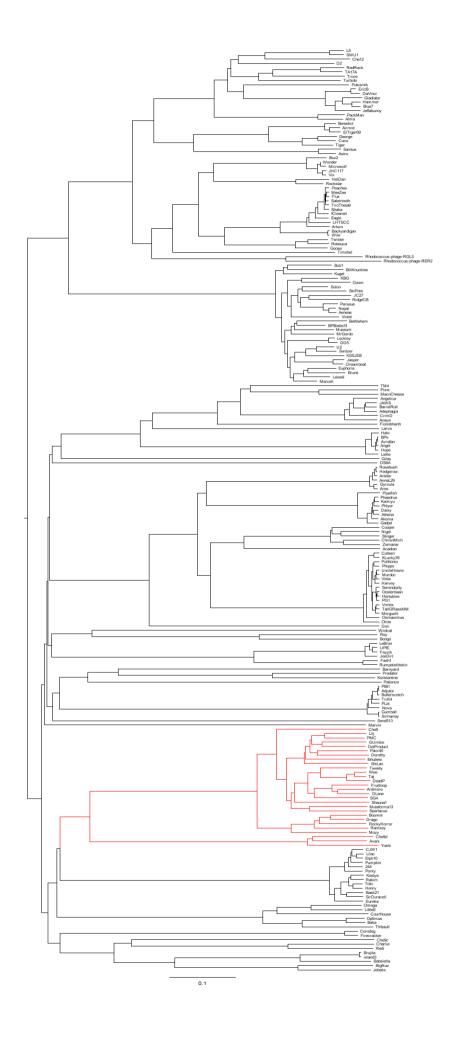


Figure 3: Clustal W phylogenetic tree (NJ) of complete genomes of all Mycobacterium siphoviruses in the NCBI database in November 2012.

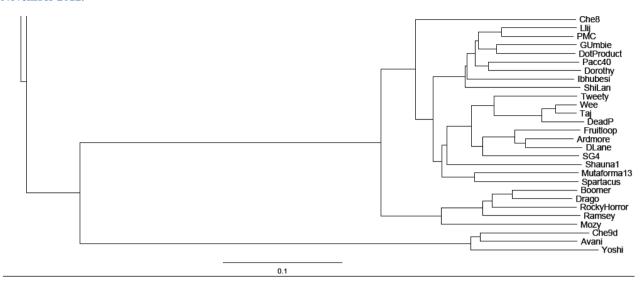


Figure 4: ClustalW phylogenetic tree of complete genomes of the isolates belonging to the genus *Che8likevirus*, excerpt of Figure 2.

Table 1: CoreGenes 3.5 and EMBOSS Stretcher analysis of the members of the proposed genus with the type phage, *Mycobacterium phage Che8*.

Phage name	% proteins in common with Che8	% of shared DNA with Che8
Mycobacterium phage PMC	69,6	74.3
Mycobacterium phage mutaforma13	67,0	72.4
Mycobacterium phage dotproduct	65,2	71.7
Mycobacterium phage gumbie	65,2	72.5
Mycobacterium phage Llij	67,0	72.8
Mycobacterium phage spartacus	67,9	70.3
Mycobacterium phage Pacc40	70,5	72.7
Mycobacterium phage ibhubesi	66,1	73.1
Mycobacterium phage tweety	65,2	72.0
Mycobacterium phage taj	61,6	70.4
Mycobacterium phage wee	60,7	69.4
Mycobacterium phage dorothy	67,0	71.2
Mycobacterium phage deadp	59,8	70.1
Mycobacterium phage SG4	63,4	68.3
Mycobacterium phage boomer	59,8	69.2
Mycobacterium phage dlane	60,7	68.1
Mycobacterium phage ardmore	59,8	68.1
Mycobacterium phage fruitloop	58,0	68.8
Mycobacterium phage shilan	68,8	69.6
Mycobacterium phage drago	59,8	68.0
Mycobacterium phage ramsey	62,5	67.4

Mycobacterium phage rockyhorror	63,4	69.2
Mycobacterium phage mozy	61,6	65.6
Mycobacterium phage shauna1	56,3	64.2
Mycobacterium phage Che9d	47,3	53.4
Mycobacterium phage avani	45,5	55.2
Mycobacterium phage yoshi	42,0	52.7