

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:	2009.002a-gB	(to be	completed	by ICTV o	officers)		
Short title: Create new genus named <i>Bcep781likevirus</i> in the family <i>Myoviridae</i> (e.g. 6 new species in the genus <i>Zetavirus</i>)							
Modules attached (modules 1 and 9 are	1 🖄	2 × 7 □	3 ⊠ 8 □	4 ☐ 9 ⊠	5 🗌		
Author(s) with e-mail address(es) of the proposer:							
	vigne@biw.kuleuven.be						
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Elizabeth Summer e	nzsum <i>w</i> tamu.edu						
Has this proposal has been seen and agreed by the relevant study group(s)? Please select answer in the box on the right						Yes	
ICTV-EC or Study	Group comments and r	espons	e of the p	roposer:			
[previous (EC41) de	ecision: inconsistent with r	naming	rules.]				
Date first submitted to ICTV:							
Date of this revision (if different to above):							

MODULE 2: NEW SPECIES

Part (a) to create and name one or more new species.

If more than one, they should be a group of related species belonging to the same genus (see Part b)

Code 2009.002aB (assigned by ICTV officers)

To create 5 new species with the name(s):

Burkholderia phage Bcep781

Burkholderia phage Bcep43 Burkholderia phage Bcep1

Burkholderia phage BcepNY3

Xanthomonas phage OP2

Part (b) assigning new species to higher taxa

All new species must be assigned to a higher taxon. This is usually a genus although it is also permissible for species to be "unassigned" within a subfamily or family.

Code 2009.002bB (assigned by ICTV officers)

To assign the species listed in section 2(a) as follows:

Genus:	Bcep781likevirus (new)
Subfamily:	
Family:	Myoviridae
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.

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.
- Provide Genbank accession numbers (not RefSeq accessions) for genomic sequences
- Further material in support of this proposal may be presented in the Appendix, Module 9

"Bcep" stands for <u>Burkholderia cepacia</u>, and phages with this designation infect bacteria belonging to the *B. cepacia* genomic complex. The proposed genus *Bcep781likevirus* comprises a group of virulent myophages of which the genome sequence of five members, Bcep781 (NC_004333), Bcep1 (NC_009015), Bcep43 (NC_005342), BcepNY3 (NC_009604) and *Xanthomonas* phage OP2 (NC_007710), is known. The Bcep781 phages are small viruses with distinctly shorter tails than P2, Mu, and BcepMu.

The genomes of these phages range from 46 to 49 kb in size and encode 66 to 71 proteins. The four Bcep phages encode a single tRNA each. They form a homogeneous phage group not just in terms of sequence, but also by their distinctive genome organization compared to other groups (Figures 1).

The genomes of the Bcep781 phages are divided into four gene clusters encoded on alternate strands such that, using Bcep781 as the example, genes 1 through 19 and 29 through 51 are present on the bottom strand while genes 20 through 28 and 52 through 66 are present on the top strand (Figure 2). Head genes are located in the first cluster and tail genes are located in the third cluster. The virion major capsid and decoration proteins, Bcep781 gp12 and gp13, were identified by protein sequencing and show some similarity to head proteins from the "PB1-like viruses". Several tail morphogenesis proteins, corresponding to Bcep781 gp29 through gp52, can be linked to P2 tail genes by PSI-BLAST. In contrast to structural genes, genes for DNA replication and lysis are scattered throughout the genome. The lysis genes of these phages are not organized into a cassette but instead overlapping Rz and Rz1 genes are separated from the endolysin and holin genes. A distinctive feature of these phages is the presence of highly, maybe completely, circularly permuted genomes. The terminases of these phages are strongly related to other pac-type phages that also have highly permuted genomes.

MODULE 3: **NEW GENUS**

Code		
	2009.002cB	(assigned by ICTV officers)
To crea	ate a new genus to contai	in the species listed below
Code	2009.002dB	(assigned by ICTV officers)
To nan	ne the new genus: Bcep78	81likevirus (new)
	g a new genus to higher ta	
Code	2009.002eB	(assigned by ICTV officers)
		lows: Ideally, a genus should be placed within a higher taxon, but if not
write "u	inassigned" in the box below	w.
Sı	ubfamily:	If any of these taxa has yet to be created
	Family: Myoviridae	(in module 4, 5 or 6) please write "(new)" after its proposed name.
	Order: Caudovirales	
assioni	ng type species and other s	species to a new genus
Code	2009.002fB	(assigned by ICTV officers)
To des	ignate the following as th	ne type species of the new genus Every genus must have a type species. This should
Burkholderia phage Bcep781		
	1 0 1	be a well characterized species although not
		necessarily the first to be discovered
Code	2009.002gB	
Code	2009.002gB	necessarily the first to be discovered
Code To assi	2009.002gB ign the following as additable and the place are place as a place as a place are place are place as a place are place and place are place are place as a place are	necessarily the first to be discovered (assigned by ICTV officers)
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Code To assi Burkho Burkho Burkho	2009.002gB ign the following as addited by the standard phage Bcep781 by the standard phage Bcep43	necessarily the first to be discovered (assigned by ICTV officers)
Code To assi Burkho Burkho Burkho Burkho Xantho	2009.002gB ign the following as addited as addited as addited as addited as addited as addited as a second as addited as a second as addited as a second as addited as addited as a second as addited	(assigned by ICTV officers) tional species of the new genus:
Code To assi Burkho Burkho Burkho Kantho Reasor	2009.002gB ign the following as addited the following as addited the following as addited the folderia phage Bcep43 olderia phage Bcep1 olderia phage BcepNY3 omonas phage OP2 ins to justify the creation of	(assigned by ICTV officers) tional species of the new genus:
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Code To assi Burkho Burkho Burkho Xantho Reasor Addition Based Origin	2009.002gB ign the following as additableria phage Bcep781 olderia phage Bcep43 olderia phage Bcep1 olderia phage BcepNY3 omonas phage OP2 ns to justify the creation of this on proteomic comparison	(assigned by ICTV officers) tional species of the new genus: of a new genus: proposal may be presented in the Appendix, Module 9 n (Figure 1)
Code To assi Burkho Burkho Burkho Xantho Reason Addition Based Origin	2009.002gB ign the following as additableria phage Bcep781 olderia phage Bcep43 olderia phage Bcep1 olderia phage BcepNY3 omonas phage OP2 ns to justify the creation of the new genus name:	(assigned by ICTV officers) tional species of the new genus: of a new genus: proposal may be presented in the Appendix, Module 9 n (Figure 1)

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.
- Provide Genbank accession numbers (not RefSeq accessions) for genomic sequences of new species

See module 2

MODULE 9: APPENDIX: supporting material

additional material in support of this proposal

References:

Summer EJ, Gonzalez CF, Bomer M, Carlile T, Morrison W, Embry A et al.: Divergence and mosaicism among virulent soil phages of the Burkholderia cepacia complex. Journal of Bacteriology 2006, 188: 255-268.

Inoue Y, Matsuura T, Ohara T, Azegami K: Sequence analysis of the genome of OP2, a lytic bacteriophage of Xanthomonas oryzae pv. oryzae. Journal of General Plant Pathology 2006, 72: 104-110.

Summer EJ, Berry J, Tran TA, Niu L, Struck DK, Young R: Rz/Rz1 lysis gene equivalents in phages of Gram-negative hosts. Journal of Molecular Biology 2007, 373: 1098-1112.

Casjens SR, Gilcrease EB, Winn-Stapley DA, Schicklmaier P, Schmieger H, Pedulla ML et al.: The generalized transducing Salmonella bacteriophage ES18: complete genome sequence and DNA packaging strategy. Journal of Bacteriology 2005, 187: 1091-1104.

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.

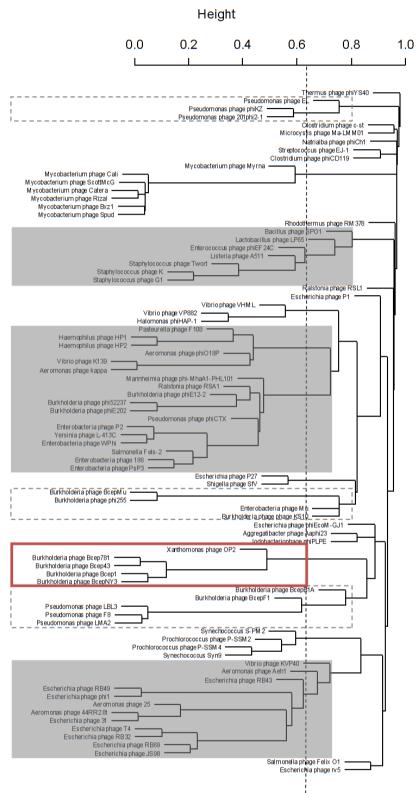


Figure 1: Hierarchical cluster dendrogram of the Myoviridae

The relative dissimilarity between the phage proteomes (between 0.0 and 1.0) forms the basis for the proposed groupings. The dotted lines reflects the cut-off value used for the establishement of genera, used consistently for all Myoviridae and the previously defined Podoviridae (Lavigne et al., 2008). Subfamily and tentative subfamily groupings are indicated in the grey and dotted boxes, respectively. The genus "Bcep781-like viruses" is highlighted in a red box

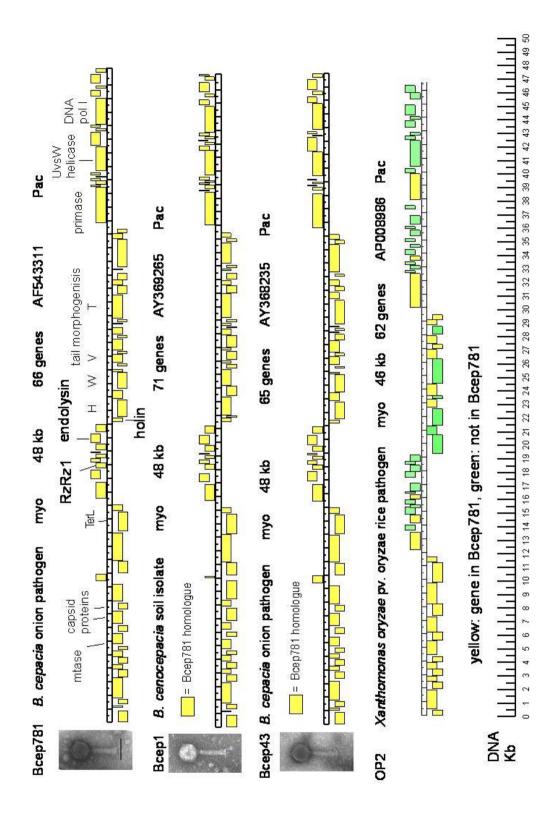


Figure 2: Genome organisation of the members of the "Bcep781-like viruses. Main predicted function are indicated and available EM micrographs have been added.