

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:	2013.031a-dB			(to be completed by ICTV officers)				
	title: To create a new genus, the <i>Phie125likevirus</i> , within the family <i>Siphoviridae</i> new species in the genus <i>Zetavirus</i>)							
Modules attached (modules 1 and 9 are required)		1 ⊠ 6 □	2 × 7 □	3 ⊠ 8 □	4 □ 9 ⊠	5 🗌		
Author(s) with e-mail address(es) of the proposer:								
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List the ICTV study group(s) that have seen this proposal:								
A list of study groups and contact								
http://www.ictvonline.org/subcom								
in doubt, contact the appropriate schair (fundal invertebrate plant)								
chair (fungal, invertebrate, plant, prokaryote or vertebrate viruses)								
ICTV-EC or Study Group comments and response of the proposer:								
Date first submitted to ICTV:			June	2013				
Date of this revision (if different to above): July				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 201	3.031aB	(assigned by ICTV officers)					
To create 3 n	ew species within:	-	Fill	in all that apply.			
Genus:	Genus: Phie125likevirus (new)			If the higher taxon has yet to be			
Subfamily:			created (in a later module, below) write "(new)" after its proposed name.				
Family:	Family: Siphoviridae		 If no genus is specified, enter 				
Order:	Caudovirales			nassigned" in the genus box.			
				GenBank sequence accession number(s) of reference isolate:			
	Burkholderia phage phie125			AF447491			
	phage phi1026b		AY453853				
Burkholderia	phage phi6442			CP000625			

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

BLASTN analyses reveal that these three *Burkholderia* temperate phages are related and distinct from any other phage. We have chosen 95% DNA sequence identity as the criterion for demarcation of species.

MODULE 3: NEW GENUS

creating a new genus

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

Code	201	3.031bB	(assigned by ICTV officers)		
To create	a new	genus within:		Fill in all that apply.	
Subfa	mily:			If the higher taxon has yet to be created	
Fa	mily:	Siphoviridae		(in a later module, below) write "(new)" after its proposed name.	
C	rder:	Caudovirales		 If no family is specified, enter "unassigned" in the family box 	

naming a new genus

Code	2013.031cB	(assigned by ICTV officers)			
To name the new genus: Phie125likevirus					

Assigning the type species and other species to a new genus

Code	2013.031dB	(assigned by ICTV officers)					
To design:	To designate the following as the type species of the new genus						
Burkholderia phage phie125 Every genus must have a type species. This is 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: 3							

Reasons to justify the creation of a new genus:

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

BLASTN analyses reveal that these three *Burkholderia* temperate phages are related and distinct from any other phage.

Burkholderia mallei-specific temperate, noninducible by UV light, phage phiE125 has an isometric head 63 nm in diameter and a long flexible tail (203 nm x 8 nm) (1). Phage phi1026b is also *B. mallei* specific and has similar morphology (head: 56nm; tail: 200x8nm) (2). DeShazer (2) provided evidence that phi1026b and phiE125 are related. Both of these phages use lipopolysaccharide as the cellular receptor. Phage phi644-2 has not been formally described but is *B. pseudomallei* specific.

Origin of the new genus name:

Burkholderia cepacia phage phiE125

Reasons to justify the choice of type species:

The original isolate of this group.

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.

MODULE 9: **APPENDIX**: supporting material

additional material in support of this proposal

References:

- 1: Woods DE, Jeddeloh JA, Fritz DL, DeShazer D. *Burkholderia thailandensis* E125 harbors a temperate bacteriophage specific for *Burkholderia mallei*. J Bacteriol. 2002 Jul;184(14):4003-17.
- 2: DeShazer D. Genomic diversity of *Burkholderia pseudomallei* clinical isolates: subtractive hybridization reveals a *Burkholderia mallei*-specific prophage in *B. pseudomallei* 1026b. J Bacteriol. 2004 Jun;186(12):3938-50.
- 3: Darling AE, Mau B, Perna NT (2010) progressiveMauve: multiple genome alignment with gene gain, loss and rearrangement. PLoS One 5: e11147
- 4: Rohwer F, Edwards RE (2002) The Phage Proteomic Tree: a genome-based taxonomy for phage. Journal of Bacteriology 184: 4529-4535

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.

Table 1. Phage genomes belonging to the proposed genus.

Phage	GenBank Accession No.	Genome size (bp)	Mol%G+C tRNA	Termini	% DNA sequence identity (a)	% Shared proteins (b)
Burkholderia phage phiE125	AF447491.1	53373	61.2	unknown	100%	100%
Burkholderia phage phi1026b	AY453853.1	54865	60.7	10-base 3' single-stranded extensions (5'- CGCCCGCTTC- 3')	81.3%	80.6%
Burkholderia phage phi644-2	CP000625.2	48674	60.4	unknown	74.1%	72.2%

- (a) Calculated using EMBOSS Stretcher (relative to phiE125)
- (b) Calculated using CoreGenes 2.0

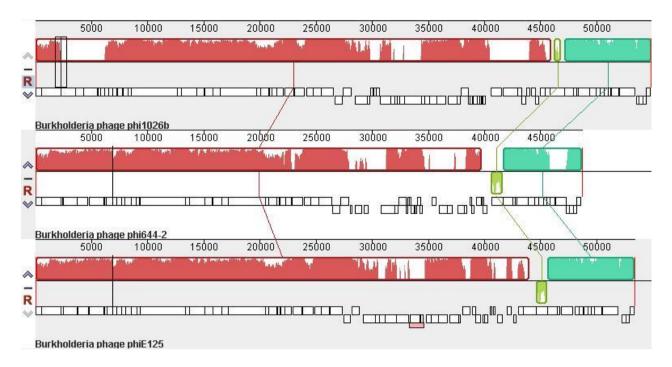


Figure 1. progressiveMauve alignment of the phage genomes belonging to the proposed genus (full genome represented by its annotated ORFs in white blocks) (3). Colored blocks indicate the regions of 1 to 1 best alignment with rearrangement breakpoints in a different random color. The degree of sequence 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.

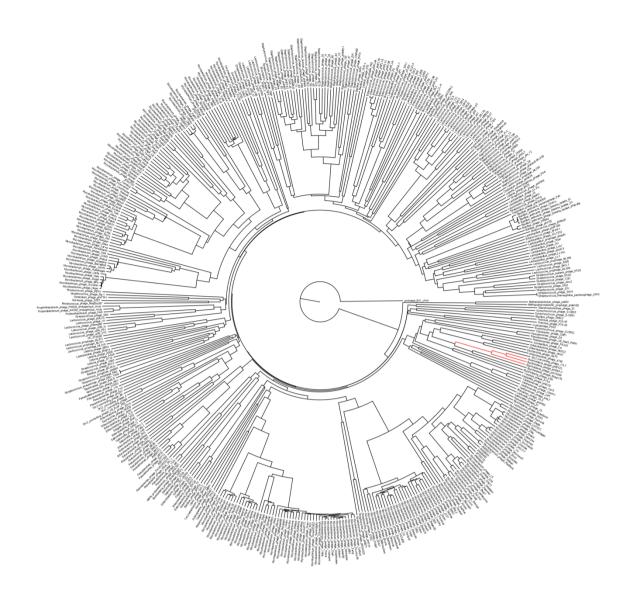


Figure 2: Phage Proteomic Tree (Rohwer & Edwards, 2002) of all the *Siphoviridae* phages in the NCBI database November 2012. Briefly, all predicted proteins sequences are compared with all others and a length-corrected protein distance matrix was calculated based on CLUSTALW alignment of sequences with a BLASTP e value < 0.001, with missing protein penalties of 10 and gap extension penalties of 0.20 (4). The tree was generated using FITCH. The proposed genus is in red. The scale bar represents protein distances of 2.0.

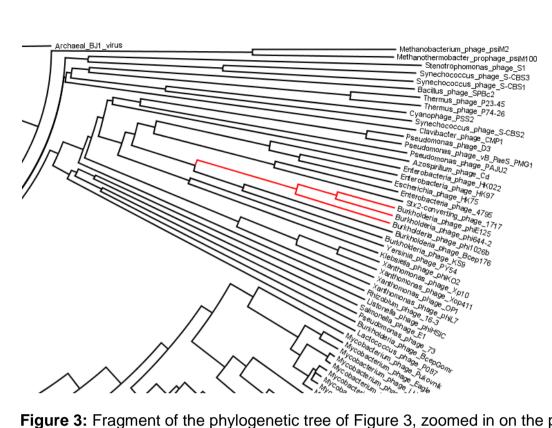


Figure 3: Fragment of the phylogenetic tree of Figure 3, zoomed in on the proposed genus.