

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:	2010.017a-eP			(to be completed by ICTV officers)		
Short title: New genus Solend (e.g. 6 new species in the genus 2 Modules attached (modules 1 and 9 are required)	ovirus in the fa Zetavirus)	amily Caul 1 🖂 6 🗌	imovirida 2 □ 7 ⊠	ae 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 contacts is provided at <u>http://www.ictvonline.org/subcommittees.asp</u>. If in doubt, contact the appropriate subcommittee chair (fungal, invertebrate, plant, prokaryote or vertebrate viruses)

Caulimoviridae

ICTV-EC or Study Group comments and response of the proposer:

EC commented that a phylogenetic tree would be helpful. This has now been added (as a copy from a related proposal)

Date first submitted to ICTV:	04/06/2010
Date of this revision (if different to above):	03/07/2010

MODULE 3: NEW GENUS

creating a new genus

deally, a genus should be placed within a higher taxon.						
Code	2010.017aP (as		(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 (in a later reached halow) write "(reav)" 			
Fai	mily:	Caulimoviridae	(In a later module, below) write (new) after its proposed name			
0	Order:		 If no family is specified, enter "unassigned" in the family box 			

naming a new genus

1

Code	2010.017bP	(assigned by ICTV officers)
To name the new genus: Solendovirus		

Assigning the type species and other species to a new genus

	1					
Code 2010.0	17cP (assi	(assigned by ICTV officers)				
To designate the following as the type species of the new genus						
Tobacco vein clearing virus		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:						

Reasons to justify the creation of a new genus:

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

Tobacco vein clearing virus is currently a species in the genus *Cavemovirus* (family *Caulimoviridae*). However, it has a significantly different genome organization to *Cassava vein mosaic virus*, the type species of the genus *Cavemovirus* (see Fig. 1 in the Appendix). These differences are:

- 1. ORF1 of CsVMV encodes a protein containing the coat protein and movement protein precursors. These proteins are encoded by two ORFs in TVCV: ORF1 (coat protein) and ORF2 (movement protein).
- 2. The ORF1 protein of CsVMV has a 124 amino acid N-terminal extension of unknown function when compared to the ORF1 protein of TVCV.
- 3. CsVMV has an additional small ORF located between ORF2 (movement protein) and ORF3 (coat protein) in TVCV.

Other than differences in genome organization, the genetic distance (% amino acid identity) between CsVMV and TVCV in the polymerase protein is equivalent to that of different virus genera in the family *Caulimoviridae* (see Table 1 and Fig. 2 in the Appendix).

GenBank Accession for TVCV: AF190123 (= NC_003378).

Origin of the new genus name:

From Solanaceae endogenous virus

Reasons to justify the choice of type species:

This is a monotypic genus.

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.

Not applicable.

MODULE 7: REMOVE and MOVE

Use this module whenever an existing taxon needs to be removed:

- *Either* to abolish a taxon entirely (when only part (a) needs to be completed)
- Or to move a taxon and re-assign it e.g. when a species is moved from one genus to another (when BOTH parts (a) and (b) should be completed)

Part (a) taxon/taxa to be removed or moved

Code	201	0.017dP	(assigned by ICTV officers)				
To remo	To remove the following taxon (or taxa) from their present position:						
Tobacco	vein d	clearing virus					
The pres	The present taxonomic position of these taxon/taxa:						
G	enus:	Cavemovirus					
Subfa	mily:		Fill in all that apply				
Fa	mily:	Caulimoviridae	Thin in an that apply.				
C	order:						
If the taxon/taxa are to be abolished (i.e. not reassigned to another taxon) write "yes" in the box on the right							

Reasons to justify the removal:

Explain why the taxon (or taxa) should be removed

See module 3 for creation of the genus Solendovirus.

Part (b) re-assign to a higher taxon

Code	201	0.017eP	(assigned by IC	CTV officers)			
To re-as	To re-assign the taxon (or taxa) listed in Part (a) as follows:						
				Fill in all that apply.			
G	enus:	Solendovirus (new)		• If the higher taxon has yet to be			
Subfa	mily:			created write "(new)" after its			
Fai	mily:	Caulimoviridae		relevant module to create it.			
0	Order:			If no genus is specified, enter			
				"unassigned" in the genus box.			

Reasons to justify the re-assignment:

- If it is proposed to re-assign species to an existing genus, please 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.
- Provide accession numbers for genomic sequences
- Further material in support of this proposal may be presented in the Appendix, Module 9

See module 3 for creation of the genus *Solendovirus*.

MODULE 9: APPENDIX: supporting material

additional material in support of this proposal

References:

- Bousalem, M., Douzery, E., and Seal, S. (2008). Taxonomy, molecular phylogeny and evolution of plant reverse transcribing viruses (family Caulimoviridae) inferred from full-length genome and reverse transcriptase sequences. Archives of Virology 153, 1085-1102.
- de Kochko, A., Verdaguer, B., Taylor, N., Carcamo, R., Beachy, R.N., and Fauquet, C. (1998). Cassava vein mosaic virus (CsVMV), type species for a new genus of plant double stranded DNA viruses? Archives of Virology 143, 945-962.
- Geering, A.D.W., Scharaschkin, T., and Teycheney, P.-Y. (2010). The classification and nomenclature of endogenous viruses of the family Caulimoviridae. Archives of Virology 155, 123-131.
- Lockhart, B.E., Menke, J., Dahal, G., and Olszewski, N.E. (2000). Characterization and genomic analysis of tobacco vein clearing virus, a plant pararetrovirus that is transmitted vertically and related to sequences integrated in the host genome. Journal of General Virology 81, 1579-1585.

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.





Fig. 1. Comparison of the genome organisation of *Tobacco vein clearing virus* and *Cassava vein mosaic virus*. Genome maps are linearised and following convention, numbering begun at the first nucleotide of the tRNA^{met} binding site. Conserved motifs are marked with the following symbols: movement protein (\clubsuit), zinc finger (*), aspartic protease active site (\diamondsuit) and reverse transcriptase active site (\diamondsuit). Dotted lines mark homologous parts of the different virus genomes. Protein molecular weights (kDa) are provided under each ORF label. Arrows denote untranslated regions.

Fig. 2. Evolutionary relationships of viruses in the family Caulimoviridae based on pol gene nucleotide sequences. The tree was generated using the maximum likelihood method implemented using the online version of PhyML version 3. Acronyms are: Drosophila melangaster Gypsy virus (DmeGypV), Saccharomyces cerevisae Ty3 virus (SceTy3V), Petunia vein clearing virus (PVCV), Cassava vein mosaic virus (CsVMV), Tobacco vein clearing virus (TVCV - proposed new genus Solendovirus), Soybean chlorotic mottle virus (SbCMV), Cestrum yellow leaf curling virus (CmYLCV), Blueberry red ringspot virus (BRRV), Peanut chlorotic stunt virus (PCSV), Strawberry vein banding virus (SVBV), Carnation etched ring virus (CERV), Lamium leaf distortion virus (LLDV), Figwort mosaic virus (FMV), Dahlia mosaic virus strain D10 (DMV-D10), Mirabilis mosaic virus (MiMV), Dahlia mosaic virus strain Holland (DMV-Holland), Horseradish latent virus (HRLV), Cauliflower mosaic virus (CaMV), Commelina yellow mottle virus (ComYMV), Banana streak MY virus (BSMysV), Banana streak GF virus (BSGFV), Banana streak VN virus (BSVNV), Kalanchoe top-spotting virus (KTSV), Banana streak OL virus (BSOLV), Pineapple bacilliform ER virus (PBERV), Pineapple bacilliform CO virus (PBCOV), Dioscorea bacilliform SN virus (DBSNV), Dioscorea bacilliform AL virus (DBALV), Citrus vellow mosaic virus (CiYMV), Cassava swollen shoot virus (CSSV), Bougainvillea chlorotic vein banding virus (BCVBV), Taro bacilliform virus (TaBV), Sugarcane bacilliform IM virus (SCBIMV), Sugarcane bacilliform Mor virus (SCBMorV), Oryza sativa virus sequence cluster A (OsatV-A), Rice tungro bacilliform virus strains type and West Bengal (RTBV- type and -WB).



Table 1. Mean nucleotide (below diagonal) and amino acid (above diagonal) distances between virus genera in the family *Caulimoviridae*. Figures provided are the number of nucleotide substitutions per site (Kimura two-parameter distance; plain font), percent nucleotide difference (bold font in brackets), number of amino acid substitutions per site (Poisson correction distance; plain font) and percent amino acid difference (bold font in brackets).

	1	r	r	r				
	Orendovirus	Tungrovirus	Badnavirus	Soymovirus	Caulimovirus	Solendovirus	Cavemovirus	Petuvirus
Orendovirus		0.653	0.671	0.995	0.772	0.843	0.896	0.974
		(48.0)	(48.9)	(63.0)	(53.7)	(57.0)	(59.2)	(62.3)
Tungrovirus	0.619		0.798	1.124	0.968	0.978	1.087	1.091
-	(41.7)		(55.0)	(67.5)	(62.0)	(62.4)	(66.3)	(66.4)
Badnavirus	0.706	0.761		1.075	0.928	1.014	1.028	1.119
	(45.5)	(47.6)		(65.8)	(60.4)	(63.7)	(64.2)	(67.3)
Soymovirus	0.795	0.883	0.975		0.764	1.053	1.104	1.076
-	(48.9)	(51.8)	(54.4)		(53.4)	(65.1)	(66.8)	(65.9)
Caulimovirus	0.691	0.837	0.881	0.718		0.884	0.924	0.932
	(45.2)	(50.3)	(51.6)	(46.0)		(58.7)	(60.3)	(60.2)
Solendovirus	0.721	0.765	0.926	0.778	0.775		0.702	1.127
	(46.3)	(47.9)	(52.9)	(48.3)	(48.2)		(50.5)	(67.6)
Cavemovirus	0.694	0.802	0.948	0.827	0.783	0.545		1.099
	(45.3)	(49.2)	(53.7)	(50.0)	(48.5)	(38.7)		(66.7)
Petuvirus	0.832	0.925	0.976	0.919	0.831	0.938	0.891	
	(50.2)	(53.1)	(54.3)	(52.9)	(50.1)	(53.5)	(52.1)	

Note: The region of comparison is the conserved reverse transcriptase and RNase H1 domains of the virus replicase and comparisons used a representative of each recognised (or proposed) species in each genus. The entry 'Orendovirus' is for sequences that are probably replication-defective endogenous copies of DNA in the rice genome. The study group is still discussing whether to propose this as a new genus.