



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"

MODULE 1: **TITLE, AUTHORS, etc**

Code assigned:	2012.010f-iP	(to be completed by ICTV officers)				
Short title: create a new genus, <i>Macanavirus</i> , in the family <i>Tombusviridae</i>						
Modules attached	1 <input checked="" type="checkbox"/>	2 <input checked="" type="checkbox"/>	3 <input checked="" type="checkbox"/>	4 <input type="checkbox"/>	5 <input type="checkbox"/>	
	6 <input type="checkbox"/>	7 <input checked="" type="checkbox"/>	8 <input type="checkbox"/>	9 <input checked="" type="checkbox"/>		

Author(s) with e-mail address(es) of the proposer:

D'Ann Rochon (dann.rochon@agr.gc.ca). On behalf of the Tombusviridae SG

List the ICTV study group(s) that have seen this proposal:

A list of study groups and contacts is provided at <http://www.ictvonline.org/subcommittees.asp> . If in doubt, contact the appropriate subcommittee chair (fungal, invertebrate, plant, prokaryote or vertebrate viruses)

Tombusviridae

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

Date first submitted to ICTV: 8 August 2011
 Date of this revision (if different to above): June 27 2012

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	2012.010fP	(assigned by ICTV officers)
To create 1 new species within:		
Genus:	<i>Macanavirus</i> (new)	
Subfamily:		
Family:	<i>Tombusviridae</i>	
Order:		
And name the new species:		GenBank sequence accession number(s) of reference isolate:
<i>Furcraea necrotic streak virus</i>		FJ768020

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:

Furcraea necrotic streak virus (FNSV) is the only member of the newly proposed *Macanavirus* genus. Also note that FNSV is not listed in the 2011 ICTV Master Species list

Background information:

FNSV has been known for many years as the agent of the damaging ‘macana’ disease of fiqué [fibre crop plants of the genus *Furcraea* (Agavaceae)] in Colombia (Dabek & Castano, 1978). Some preliminary serological and cDNA hybridization studies suggested that it might be a member of the genus *Dianthovirus*, family *Tombusviridae* (Morales et al., 1992). However, dianthoviruses have a bipartite genome and the complete genome of FNSV now determined by the same research group (accession FJ768020) shows it to have a monopartite genome. There is no publication describing the sequence but Dr. Morales has confirmed to the SG the sequence is indeed derived from the isolate described in the 1992 paper. Although the Genbank accession is still labeled as an unclassified dianthovirus, FNSV was listed as a possible member of the genus *Necrovirus* in the *Tombusviridae* chapter of the ICTV 9th Report. This was because comparisons of its replication protein sequence showed that it was more closely related to some necroviruses than to viruses classified in other genera of the family.

As described below, there is a current proposal, expected to be ratified this year, to revise the genus *Necrovirus* and to split it into two genera (*Alpha-* and *Betanecrovirus*) (2011.009a-mP) but both the polymerase sequence and coat protein properties of FNSV place it outside either of these and suggest it is a member of a separate genus (see Module 3 for further details).

Also, last year a proposal was put forth (2011.010a-fP.N.v1) to create the new genus *Gallantvirus* in the family *Tombusviridae* and to assign *Galinsoga* mosaic virus (GaMV) currently a species in the genus *Carmovirus* in the *Tombusviridae* as its type member and to create FNSV as a new species in the genus. This proposal was based on the fact that FNSV and GaMV share similar genome organizations, similar “necrovirus-like” polymerase sequences but the coat proteins have protruding domains, unlike the necroviruses (note that the presence or absence of a coat protein protruding domain is a genus demarcation criteria in the *Tombusviridae*). However, the Study Group was asked to “reconsider whether the two viruses should be placed in the same genus as the phylogenetic trees of the polymerase suggested that two genera could be justified”. The *Tombusviridae* Study Group has considered these comments and has agreed to create two separate genera for FNSV and GaMV.

1MODULE 3: **NEW GENUS**

creating a new genus

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

Code	2012.010gP	(assigned by ICTV officers)
To create a new genus within:		
Subfamily:		
Family:	<i>Tombusviridae</i>	
Order:		

naming a new genus

Code	2012.010hP	(assigned by ICTV officers)
To name the new genus: <i>Macanavirus</i>		

Assigning the type species and other species to a new genus

Code	2012.010iP	(assigned by ICTV officers)
To designate the following as the type species of the new genus		
<i>Furcraea necrotic streak virus</i>		
Please enter here the TOTAL number of species (including the type species) that the genus will contain:		
1		

Reasons to justify the creation of a new genus:

[Additional material in support of this proposal may be presented in the Appendix, Module 9](#)

Genera in the family *Tombusviridae* are distinguished by genome organization and by the properties of the replication protein and the coat protein. Necroviruses have a distinctive type of coat protein without a protruding domain and forming a single lineage but their replication proteins are of two different lineages. There is therefore a current proposal to split the genus *Necrovirus* into two genera, *Alphanecrovirus* and *Betanecrovirus* (2011.009a-mP).

The complete sequence of FNSV shows that this virus is most similar to alphanecroviruses in genome organization and in percent sequence identity in the replication protein and the two small movement protein ORFs (Tables 1-4). This is confirmed by phylogenetic analyses (Figs 3 and 4).

However, unlike necroviruses, the coat protein of FNSV contains a protruding domain (Fig. 1, 2) and is most closely related to the coat proteins of other members of the family with protruding domains in both sequence comparisons (Table 4) and phylogenetic analyses (Fig. 5).

To be consistent with genus demarcation criteria being applied in the family, a new genus is therefore needed to contain FNSV.

Origin of the new genus name:

From the common name of the disease “Macana” that is used for FNSV infected plants in Columbia, South America

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.

< 80% amino acid sequence identity in the polymerase;
<55% amino acid sequence similarity in the coat protein.

MODULE 9: **APPENDIX**: supporting material

additional material in support of this proposal

References:

Dabek, A.J. and Castano, J.J. (1978). The occurrence, symptomatology, transmission and virus aetiology of macana disease of figue (*Furcraea* spp.) in Colombia, South America. *Phytopath. Z.*, 92: 57-69

Morales, F.J., Castano, M., Calvert, L.A. and Arroyave, S. (1992). *Furcaea* necrotic streak virus: and apparent new member of the Dianthovirus group. *J. Phytopath.* **134**: 247-254.

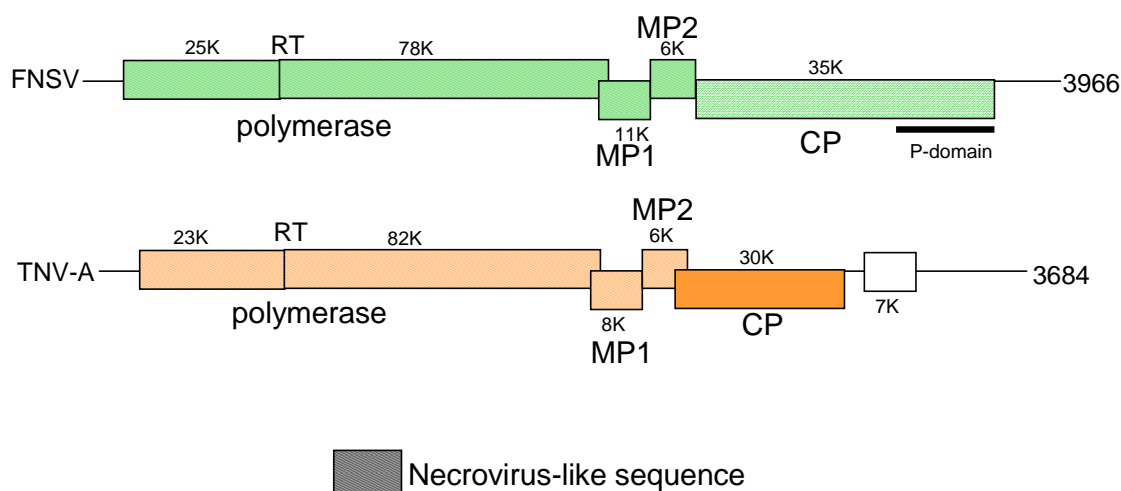
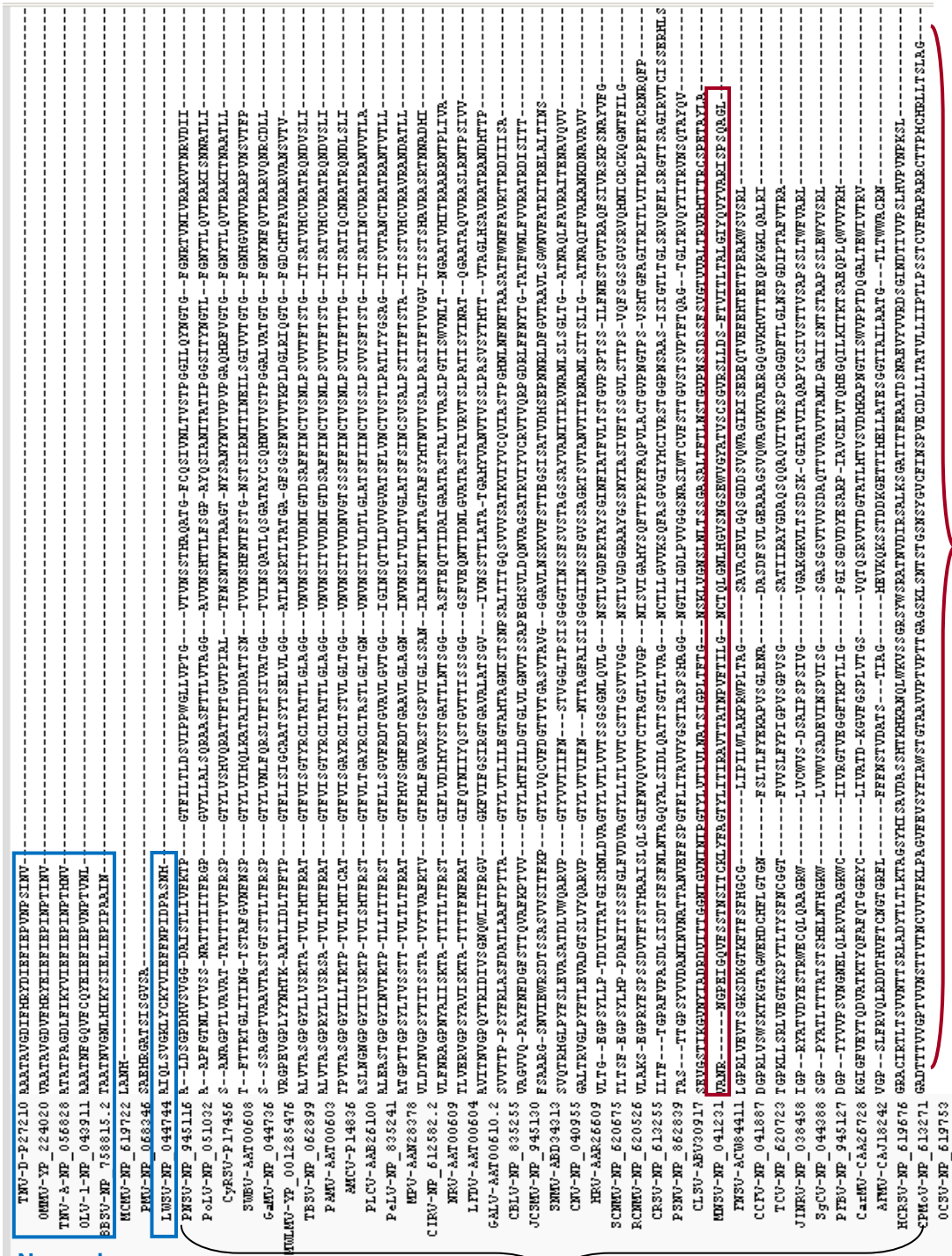


Figure 1. Genome organization of FNSV and comparison to tobacco necrosis virus A (TNV-A), the type member of genus *Necrovirus* (and proposed type member of the new genus Alphanecrovirus). Similar shading patterns indicate significant sequence identity. Note that the coat protein (CP) of FNSV, unlike TNV-A, contains a C-terminal protruding domain which is typical of currently classified viruses in the *Tombusviridae* with the exception of the necroviruses.



Coat protein protrusion

Figure 2. Alignment of the CP sequence of NNSV (boxed in red) with the CPs of the proposed alpha and betanecroviruses (boxed in blue) and remaining members of the *Tombusviridae*. Only the C-terminal region of the CP alignment is shown with the area corresponding to the P-domain being shown in brackets.

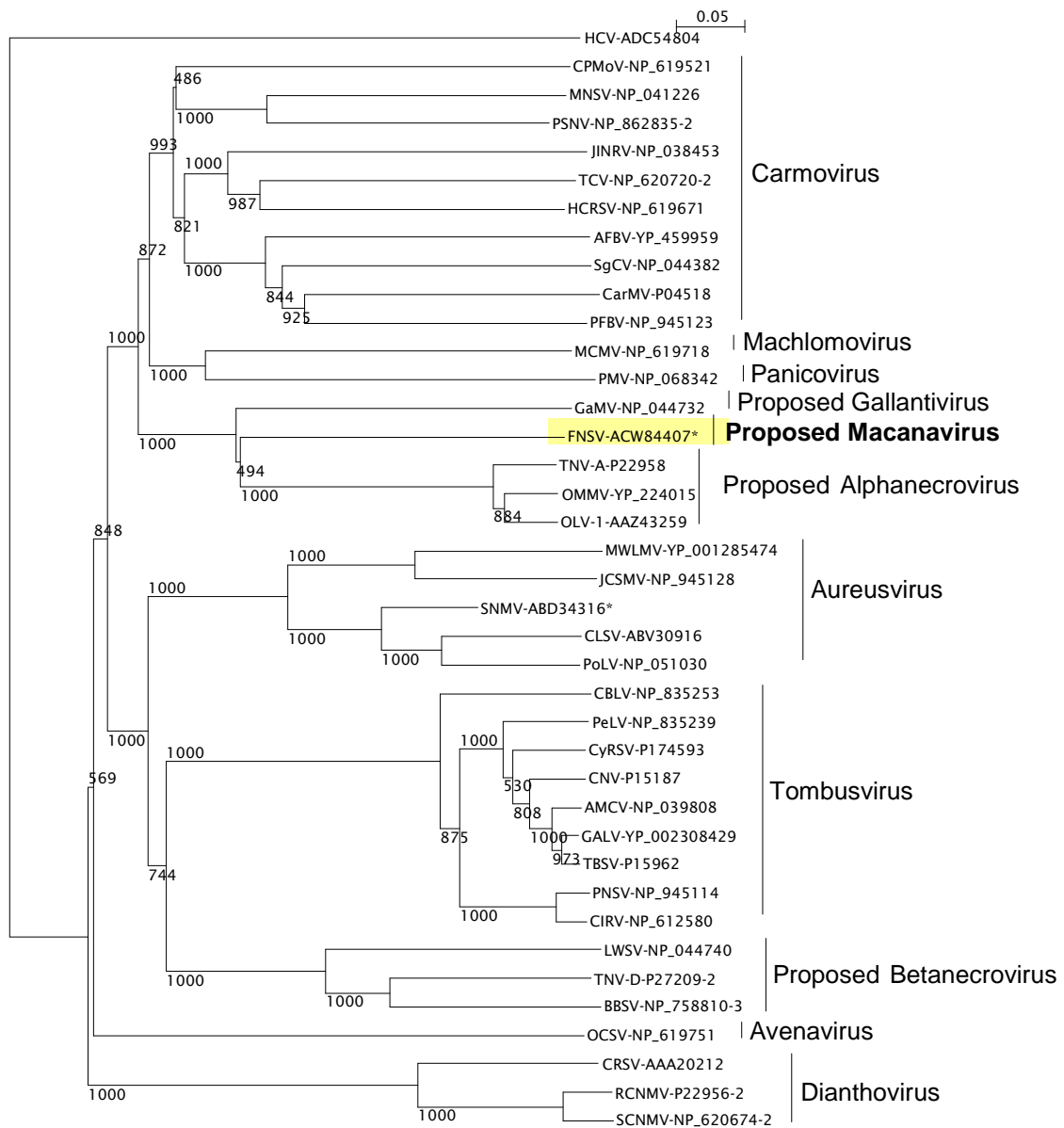


Figure 3. Phylogenetic (distance) tree of the polymerase of *Tombusviridae* members with the position of FNSV highlighted in yellow. Sequences were aligned using the ClustalX 2.1 algorithm and trees were generated by the Neighbor Joining method using 1000 bootstrap replicates. Hepatitis C virus (HCV) polymerase was used as an outgroup.

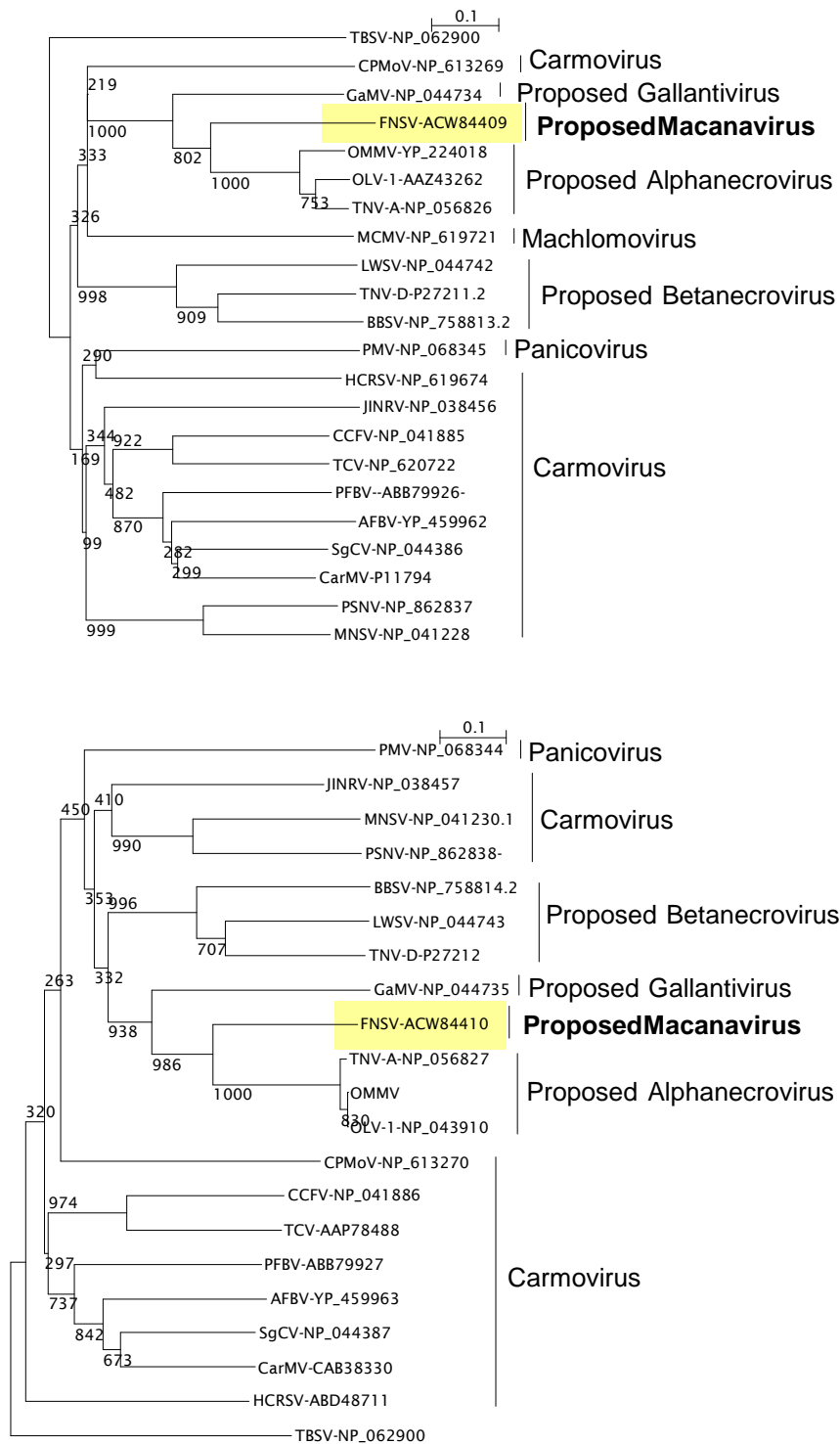
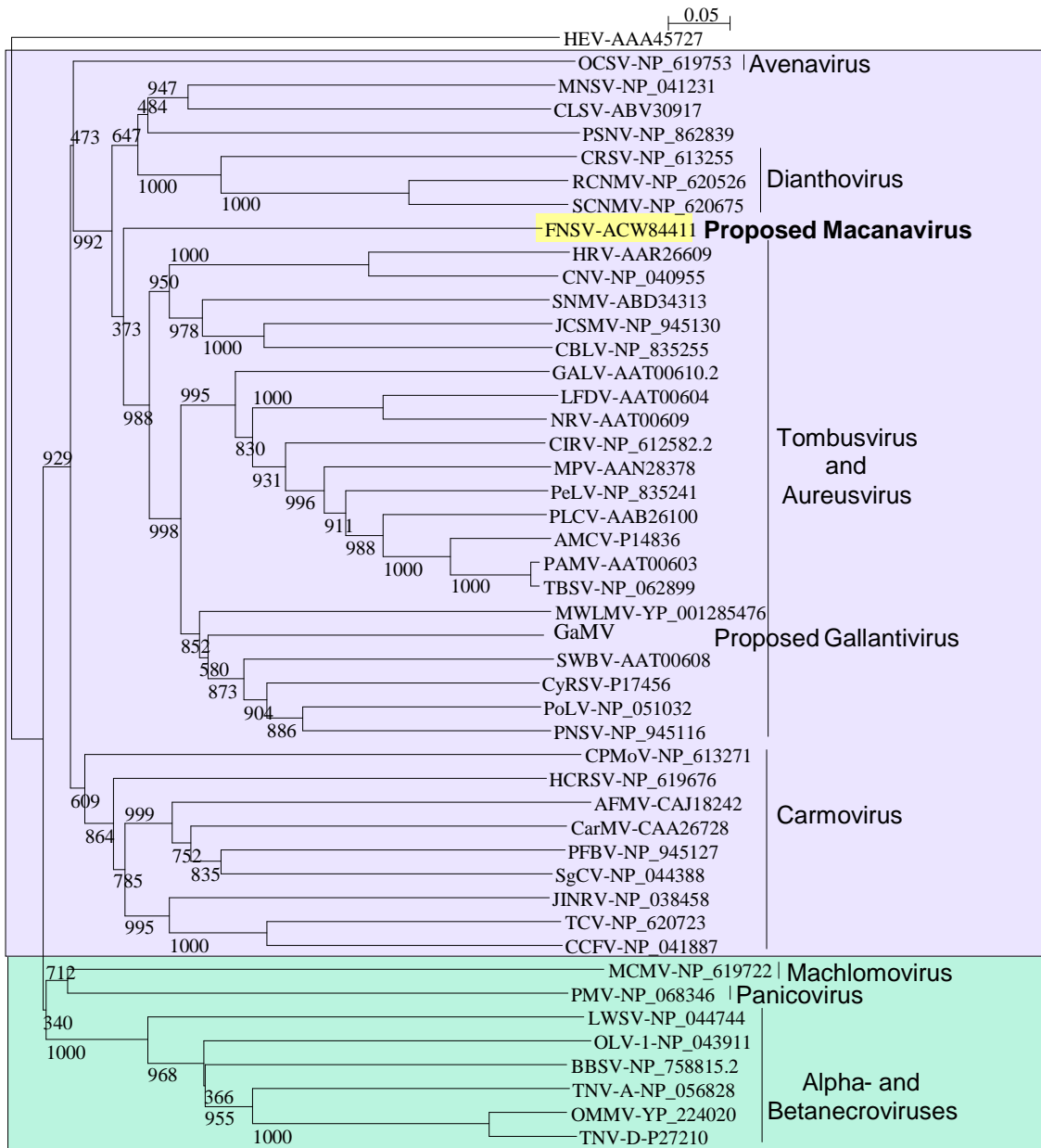


Figure 4. Phylogenetic (distance) tree of the MP1 (top) and MP2 (bottom) sequences of *Tombusviridae* members with the position of the FNSV MPs highlighted in yellow. Sequences were aligned using the ClustalX 2.1 algorithm and trees were generated by the Neighbor Joining method using 1000 bootstrap replicates.



- Virus coat proteins with protruding domains
- Virus coat proteins without protruding domains

Figure 5. Phylogenetic (distance) tree of CP sequences of *Tombusviridae* members with the position of the FNSV CP highlighted in yellow. Sequences were aligned using the ClustalX 2.1 algorithm and trees were generated by the Neighbor Joining method using 1000 bootstrap replicates. Hepatitis E virus (HEV) coat protein was used as an outgroup.

ABBREVIATIONS USED IN THE PROPOSAL

Tombusviruses

<i>Artichoke mottled crinkle virus</i>	AMCV
<i>Carnation Italian ringspot virus</i>	CIRV
<i>Cucumber Bulgarian latent virus</i>	CBLV
<i>Cucumber necrosis virus</i>	CNV
<i>Cymbidium ringspot virus</i>	CyRSV
<i>Pear latent virus - Italy</i>	PeLV*
<i>Grapevine Algerian latent virus</i>	GALV
<i>Havel river virus</i>	HRV
<i>Lato river virus</i>	LRV
<i>Moroccan pepper virus</i>	MPV
<i>Neckar river virus</i>	NRV
<i>Pelargonium leaf curl virus</i>	PLCV
<i>Pelargonium necrotic spot virus</i>	PNSV
<i>Petunia asteroid mosaic virus</i>	PAMV
<i>Sikte waterborne virus</i>	SWBV
<i>Tomato bushy stunt virus</i>	TBSV
<i>Limonium flower distortion virus</i>	LFDV

*PeLV is an isolate of *Eggplant mottle crinkle virus*

Dianthoviruses

<i>Carnation ringspot virus</i>	CRSV
<i>Red clover necrotic mosaic virus</i>	RCNMV
<i>Sweet clover necrotic mosaic virus</i>	SCNMV

Aureusviruses

<i>Cucumber leaf spot virus</i>	CLSV
<i>Johnsongrass chlorotic stripe mosaic virus</i>	JCSMV
<i>Maize white line mosaic virus</i>	MWLMV
<i>Pothos latent virus</i>	PoLV
<i>Sesame necrotic mosaic virus</i> (tentative species)	SNMV

Avenavirus

<i>Oat chlorotic stunt virus</i>	OCSV
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Carmoviruses

<i>Angelonia flower break virus</i>	AFBV
<i>Cardamine chlorotic fleck virus</i>	CCFV
<i>Carnation mottle virus</i>	CarMV

<i>Cowpea mottle virus</i>	CPMoV
<i>Galinsoga mosaic virus</i>	GaMV
<i>Hibiscus chlorotic ringspot virus</i>	HCRSV
<i>Japanese iris necrotic ring virus</i>	JINRV
<i>Melon necrotic spot virus</i>	MNSV
<i>Pea stem necrosis virus</i>	PSNV
<i>Pelargonium flower break virus</i>	PFBV
<i>Saguaro cactus virus</i>	SgCV
<i>Turnip crinkle virus</i>	TCV

Necroviruses

<i>Beet black scorch virus</i>	BBSV
<i>Leek white stripe virus</i>	LWSV
<i>Olive latent virus 1</i>	OLV-1
<i>Olive mild mosaic virus</i>	OMMV
<i>Tobacco necrosis virus A</i>	TNV-A
<i>Tobacco necrosis virus D</i>	TNV-D

Proposed Macanavirus

<i>Furcraea necrotic streak virus</i>	FNSV
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Panicovirus

<i>Panicum mosaic virus</i>	PMV
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Machlomovirus

<i>Maize chlorotic mottle virus</i>	MCMV
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Proposed Zeavirus

<i>Maize necrotic streak virus</i>	MNeSV
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Table 1. Percent amino acid sequence identity between the polymerase of FNSV and those of the proposed alpha- and necrovirus genera and other *Tombusviridae* members

	% aa sequence identity in polymerase		
	Alpha-necrovirus	Beta-necrovirus	Other Tombusviridae
FNSV	52-52	32-42	25-38

Table 1. Percent amino acid sequence identity between the MP1 of FNSV and those of the proposed alpha- and necrovirus genera and other *Tombusviridae* members

	% aa sequence identity in MP1				
	Alpha-necro	Beta-necro	Carmo-	PMV	MCMV
FNSV	47-49	12-20	8-20	8	20

Table 1. Percent amino acid sequence identity between the MP2 of FNSV and those of the proposed alpha- and necrovirus genera and other *Tombusviridae* members

	% aa sequence identity in MP2			
	Alpha-necro	Beta-necro	Carmo-	PMV-
FNSV	57	21-25	7-23	23

Table 1. Percent amino acid sequence identity between the CP of FNSV and those of the proposed alpha- and necrovirus genera and other *Tombusviridae* members

	% aa sequence identity in coat protein								
	Alpha-necro	Beta-necro	PMV	MCMV	Tombus	Aureus	Carmo	OCSV	Diantho
FNSV	13-17	13-18	14	13	25-31	27-32	15-31	20	25-28

- Virus coat proteins with protruding domains
- Virus coat proteins without protruding domains