

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:	2013.011a-dV			(to be completed by ICTV officers)		
Short title: Create a new species, <i>Mosavirus A</i> , in a new genus, <i>Mosavirus</i> , within the family <i>Picornaviridae</i> (order <i>Picornavirales</i> ) (e.g. 6 new species in the genus <i>Zetavirus</i> )						
Modules attached (modules 1 and 9 are required)		1 × 6 □	2 <u>×</u> 7 <u></u>	3 ⊠ 8 □	<b>4</b> ∐ <b>9</b> ⊠	5 📋
Author(s) with e-mail address(es) of the proposer:						
Nick J. Knowles ( <u>nick.knowles@pirbright.ac.uk</u> ) on behalf of the <i>Picornaviridae</i> Study Group						
List the ICTV study group(s) that have seen this proposal:						
A list of study groups and contact <a href="http://www.ictvonline.org/subcom">http://www.ictvonline.org/subcom</a> in doubt, contact the appropriate chair (fungal, invertebrate, plant, vertebrate viruses)	Picornaviridae Study Group					
ICTV-EC or Study Group comments and response of the proposer:						
Date first submitted to ICTV: Date of this revision (if differe	nt to above):		25/0	5/2013		

## **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 <b>201</b>	3.011aV	(assigned by ICTV off	icers)		
To create one new species within:					
Canus	Magazinea (nove)		l in all that apply.		
Genus: Subfamily:	Mosavirus (new) n/a	C	reated (in a later module, below) write (new)" after its proposed name.		
Family:	Picornaviridae		no genus is specified, enter		
Order:	Picornavirales		"unassigned" in the genus box.		
And name the	e new species:		GenBank sequence accession number(s) of reference isolate:		
Mosavirus A			JF973687		

## 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

## Virus discovery

A novel picornavirus (designation M-7) has been detected in the faeces of a canyon mouse (*Peromyscus crinitus*) in the USA in 2010 (JF973687; Phan *et al.*, 2011).

#### Growth in cell cultures

The virus has not been cultivated in cell cultures.

## **Untranslated regions**

Only 138 nt of the 5' untranslated region (UTR) has been determined and the internal ribosome entry site (IRESE) type is not known. The 3' UTR is 91 nt long and neither region has significant primary sequence identity with any other picornavirus.

# Genome organization/proteins

VPg+5'UTR[L/1A-1B-1C-1D-2A<sup>npgp</sup>/2B-2C/3A-3B1<sup>VPg1</sup>-3B2<sup>VPg2</sup>-3C<sup>pro</sup>-3D<sup>pol</sup>]3'UTR-poly(A)

- [], defines the long ORF encoding the polyprotein.
- /, Indicates primary polyprotein cleavages.
- -, indicates secondary cleavages mainly performed by the 3C<sup>pro</sup> polypeptide.

A 146 aa leader polypeptide precedes the capsid and has no sequence identity with any other picornavirus protein. The 2A peptide is either very short or is part of VP1 and ends with an

NPG↓P motif. VP0 (1AB) is predicted to be cleaved to VP4/VP2 and possesses a myristoylation signal at its amino-terminus (i.e. GxxxT/S as GGGES).

Mosavirus is predicted to encode in tandem two VPg's:

```
GPYCGTCRQKKPVLKKAVTE (VPg1)
||| | || ||
GPYTGIVKKAPKKLKKVVTQ (VPg2)
```

The only other picornaviruses with multiple VPg's are *Foot-and-mouth disease virus* which has three copies and *Aquamavirus A* which is predicted to possess two copies.

## Genetic relationships

The proteins of mosavirus are most closely related to members of the *Erbovirus*, *Cardiovirus* and *Aphthovirus* genera with the closest relationships being with *Equine rhinitis B virus* (37.4%), *Theilovirus* (36.4%) and *Equine rhinitis A virus* (35.5%) in the P1, P2 and P3 polypeptides, respectively.

## **MODULE 3: NEW GENUS**

creating a new genus

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

Code	201	3.011bV	(assigned by I	(assigned by ICTV officers)		
To create	a new	genus within:		Fill in all that apply.		
Subfa	mily:	n/a		• If the higher taxon has yet to be created		
Fa	mily:	Picornaviridae		(in a later module, below) write "(new)" after its proposed name.		
C	rder:	Picornavirales		If no family is specified, enter     "unassigned" in the family box		

naming a new genus

Code	2013.011cV	(assigned by ICTV officers)	
To name the new genus: Mosavirus			

Assigning the type species and other species to a new genus

Assigning the type species and other species to a new genus				
Code	2013.011dV	(assigned by ICTV officers)		
To designate the following as the type species of the new genus				
Mosavirus	A	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:  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

The relationships between mosavirus and other picornaviruses in the P1, P2 and P3 polypeptides are 37.4% (*Erbovirus*), 36.4% (*Cardiovirus*) and 35.5% (*Aphthovirus*), respectively. The *Picornaviridae* Study Group (PSG) guidelines state that members of different genera share less that 40%, 40% and 50% amino acid difference in P1, P2 and P3, respectively. We therefore suggest that the proposed species *Mosavirus A* is placed in a new genus named *Mosavirus*.

## Origin of the new genus name:

Mosavirus, from mouse stool-associated picornavirus.

## Reasons to justify the choice of type species:

The genus is proposed to contain only a single species.

## 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.

None, since there is only a single species.

# MODULE 9: **APPENDIX**: supporting material

additional material in support of this proposal

## **References:**

Phan, T.G., Kapusinszky, B., Wang, C., Rose, R.K., Lipton, H.L. and Delwart, E.L. (2011). The fecal viral flora of wild rodents. PLoS Pathog 7(9): e1002218.

#### 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.

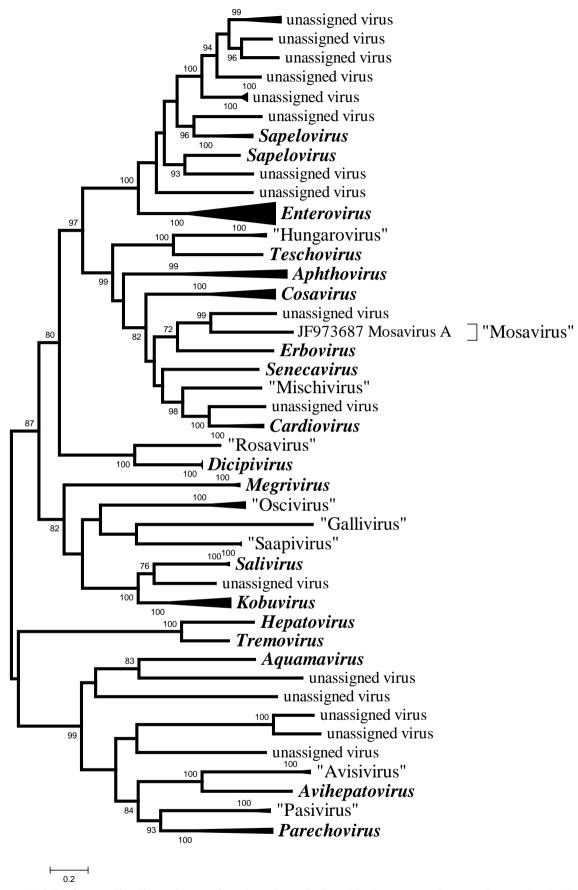


Figure 1. Maximum likelihood tree showing the relationship between picornaviruses in the P1 capsid. Sequences were aligned using MUSCLE and the tree constructed using MEGA 5.2.

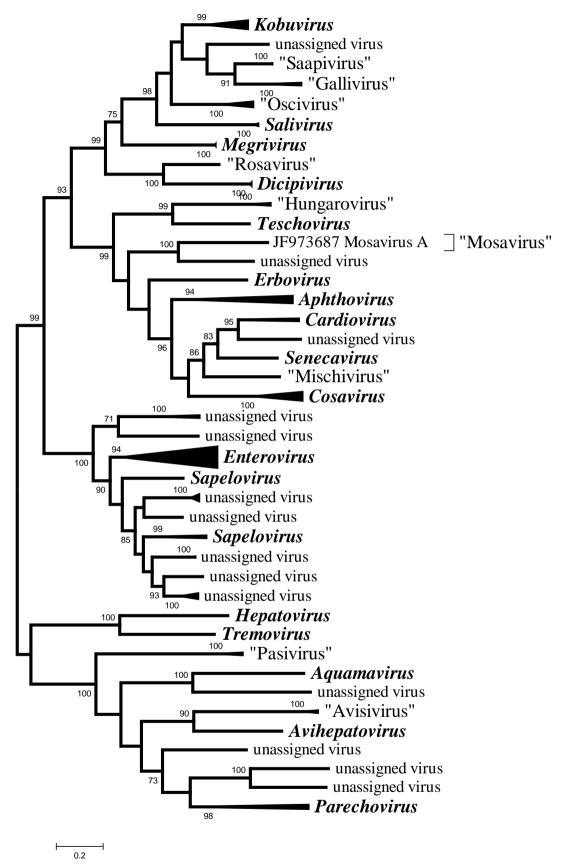


Figure 2. Maximum likelihood tree showing the relationship between picornaviruses in the 3D polymerase. Sequences were aligned using MUSCLE and the tree constructed using MEGA 5.2.