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Taxonomy proposal: One (1) new species, Pseudoalteromonas virus Cr39582, in the genus Corticovirus, in the family Corticoviridae

Leigh, Brittany A

International Committee on Taxonomy of Viruses (ICTV) 2018

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This Word module should be used for all taxonomic proposals. Please complete **Part 1** and:

either Part 3 for proposals to create new taxa or change existing taxa or Part 2 for proposals of a general nature.

Submit the completed Word module, together with the accompanying Excel module named in Part 3, to the appropriate ICTV Subcommittee Chair.

For guidance, see the notes written in blue, below, and the help notes in file Taxonomic_Proposals_Help_2018.

Part 1: TITLE, AUTHORS, etc

Code assigned:	2018.123B	(to be completed by ICTV officers)										
	new species, <i>Pseudoalterom</i> family <i>Corticovirida</i> e	onas virus Cr39582, in the genus										
(e.g. "6 new species in the genus Zetavirus")												
Author(s):												
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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 (there are six virus subcommittees: animal DNA and retroviruses, animal ssRNA-, animal ssRNA+, fungal and protist, plant, bacterial and archaeal)

Bacterial and Archaeal Viruses Subcommittee

ICTV Study Group comments (if any) and response of the proposer:

Date first submitted to ICTV: Date of this revision (if different to above): June 6, 2018

ICTV-EC comments and response of the proposer:

Part 2: NON-STANDARD

Template for any proposal regarding ICTV procedures, rules or policy, <u>not</u> involving the creation of new taxonomy.

Text of proposal:

Part 3: PROPOSED TAXONOMY

Name of accompanying Excel module: 2018.123B.N.v1.Corticovirus_sp

The taxonomic changes you are proposing should be presented on an accompanying Excel module, 2017_TP_Template_Excel_module. Please enter the file name of the completed module in this box.

Supporting material:

additional material in support of this proposal

Please explain the reasons for the taxonomic changes you are proposing and provide evidence to support them. The following information should be provided, where relevant:

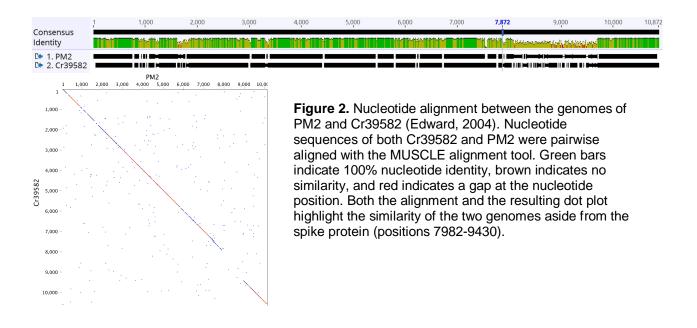
- **Species demarcation criteria**: Explain how new species differ from others in the genus and demonstrate that these differences meet the criteria previously established for demarcating between species. If no criteria have previously been established, and if there will now be more than one species in the genus, please state the demarcation criteria you are proposing.
- Higher taxa:
 - There is no formal requirement to state demarcation criteria when proposing new genera or other higher taxa. However, a similar concept should apply in pursuit of a rational and consistent virus taxonomy.
 - Please indicate the **origin of names** assigned to new taxa at genus level and above.
 - For each new genus a **type species** must be designated to represent it. Please explain your choice.
- **Supporting evidence**: The use of Figures and Tables is strongly recommended (note that copying from publications will require permission from the copyright holder). For phylogenetic analysis, try to provide a tree where branch length is related to genetic distance.

Species demarcation criteria: We propose to have 95% nucleotide sequence identity as the criterion for demarcation of species in the genus *Corticovirus*. The proposed species *Pseudoalteromonas virus Cr39582* differs from the type species *Pseudoalteromonas virus PM2* by ~15% at the nucleotide level as confirmed by the full genome alignment by Mauve (Darling et al., 2010) and MUSCLE (Edgar, 2004) nucleotide alignments (Leigh et al., 2018). *Pseudoalteromonas virus PM2* is currently the only species in the genus *Corticovirus*, the family *Corticoviridae* (Oksanen, 2017).

Supporting evidence: The Cr39582 and PM2 genomes are syntenous sharing sequence homology throughout their genome lengths (Figures 1 and 2) (Leigh et al., 2018). The only exception is the Cr39582 ORF (nucleotides 7982-9430) encoding for the putative spike protein which is not homologous with PM2 spike protein P1. The nucleotide sequence identity between Cr39582 and PM2 spike encoding genes is ~37%. Pairwise nucleotide identities between Cr39582 and PM2 in the 5' and 3' regions flanking the spike protein gene are ~91% and ~96%, respectively.

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Figure 1. Multiple genome alignment between the genomes of PM2 and Cr39582 (Mauve, Darling et al., 2010). Colored boxes indicate blocks of the genomes with high sequence identity, and regions lacking homology appear as white.



References:

Darling AE, Mau B, Perna N. 2010. progressiveMauve: multiple genome alignment with gene gain, loss and rearrangement. PLoS One 5:e11147

Edgar, R. C. 2004. MUSCLE: multiple sequence alignment with high accuracy and high throughput. Nucleic Acids Res. 32: 1792-1797.

Leigh BA, Breitbart M, Oksanen HM, Bamford DH, Dishaw LJ. 2018. Genome sequence of PM2-like phage Cr39582, induced from a *Pseudoalteromonas* sp. isolated from the gut of *Ciona robusta*. Genome Announc. 6:e00368-18.

Oksanen HM, ICTV Report Consortium. 2017. ICTV Virus Taxonomy Profile: *Corticoviridae*. J Gen Virol. 98:888-889.