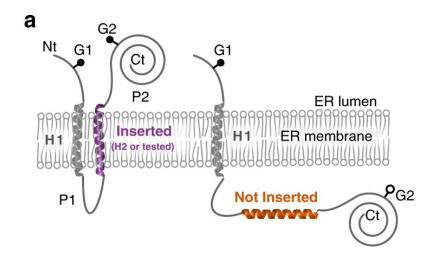


Research demonstrates the role of intramebrane interactions in viral the control of cell death

November 30 2020





b	Organism T	ested segment	Sequence	ΔG_{pred}	ΔG_{exp}
	E.coli	Lep H2	WLETGASVFPVLAIVLIVRSFIY	-0.01	-1.86
	HHV4	BHRF1 Ct	FSWTLFLAGLTLSLLVIW	-0.742	-1.16
	ннv8	ORF16 Ct	RRMTALLGSIALLATILAAVAM	-1.008	-1.30
	BoHV4	ORF16 Ct	SRWLFPMLAISGLVLTVGVAR	-0.168	-1.46
	VacV	F1 Ct	REYLKLIGITAIMFATYKTL	+0.663	-1.67
	MyxV	M11L Ct	ISVYLTAAVVGFVAYGILKWYR	-0.217	-2.17
	OrfV	ORF125 Ct	RALGVGAVVGVGMLLLGCRVLR	-0.725	-1.93
	H.sapiens	Tomm20 Ct	AIAAGVCGALFIGYCIYF	+0.312	-1.71
	H.sapiens	Tomm22 Ct	ALWIGTTSFMILVLPVVFET	+0.428	-1.34
	H.sapiens	Noxa Ct	FGDKLNFRQKLLNLISKLFCSGT	+8.054	+2.08

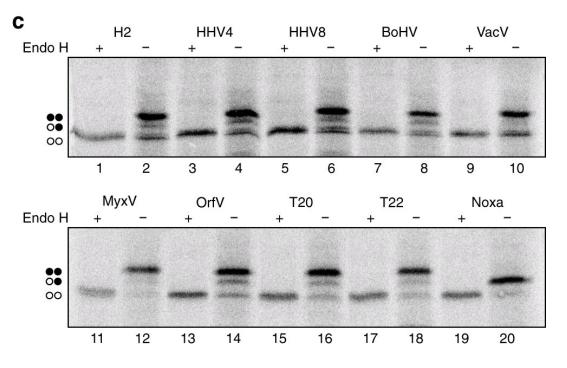




Fig. 1: Insertion of vBcl2 Ct hydrophobic sequences in eukaryotic membranes. From: Viral Bcl2s' transmembrane domain interact with host Bcl2 proteins to control cellular apoptosis

A research group from the Department of Biochemistry and Molecular Biology of the University of Valencia (UV), in coordination with the National Center for Biotechnology (CNB) of the CSIC, has studied the role of the interactions within the membrane of proteins of viral families Herpesviridae and Poxviridae in the control of programmed cell death. The work, published in *Nature Communications*, could have implications for the development of treatments for viral infection, as well as the prevention of cancers associated with them.

The results of the finding, led by Dr. Luis Martínez, Ph.D. in the Department of Biochemistry and Molecular Biology, would imply that interactions within the membrane between virus proteins and the host individual could be used as therapeutic targets for the treatment of some <u>viral infections</u>. An agent capable of blocking such interactions would not only reduce, or even inhibit, viral replication, but also slow down the possible development of cancer associated with such infections.

Cell apoptosis (programmed cell death) is an essential process in multicellular organisms, as it contributes to the balance between cell death, proliferation and differentiation, which is relevant for the development and proper functioning of living things. This makes it a highly regulated process involving many components, including the protein family known as Bcl2 (B-cell lymphoma 2).

In order to maximize their growth, viruses in the Herpesviridae and



Poxviridae families have developed mechanisms to modulate cell death in host individuals. Therefore, these viruses have proteins structurally similar to Bcl2 proteins, known as viral Bcl2, which have a <u>transmembrane domain</u> that allows the protein to be inserted into the target membrane to deregulate cell apoptosis.

"In this study we show that viral Bcl2 proteins have a transmembrane domain (TMD) that allows them to be anchored to the mitochondrial membrane. In addition, we observed that these proteins are able to interact with each other and with other Bcl2 proteins of host individuals through this domain. Our results also indicate that these interactions are key to controlling <u>cell death</u> after an apoptotic stimulus such as a viral infection", said Luis Martinez.

More information: Maria Jesús García-Murria et al. Viral Bcl2s' transmembrane domain interact with host Bcl2 proteins to control cellular apoptosis, *Nature Communications* (2020). DOI: 10.1038/s41467-020-19881-9

Provided by Asociacion RUVID

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