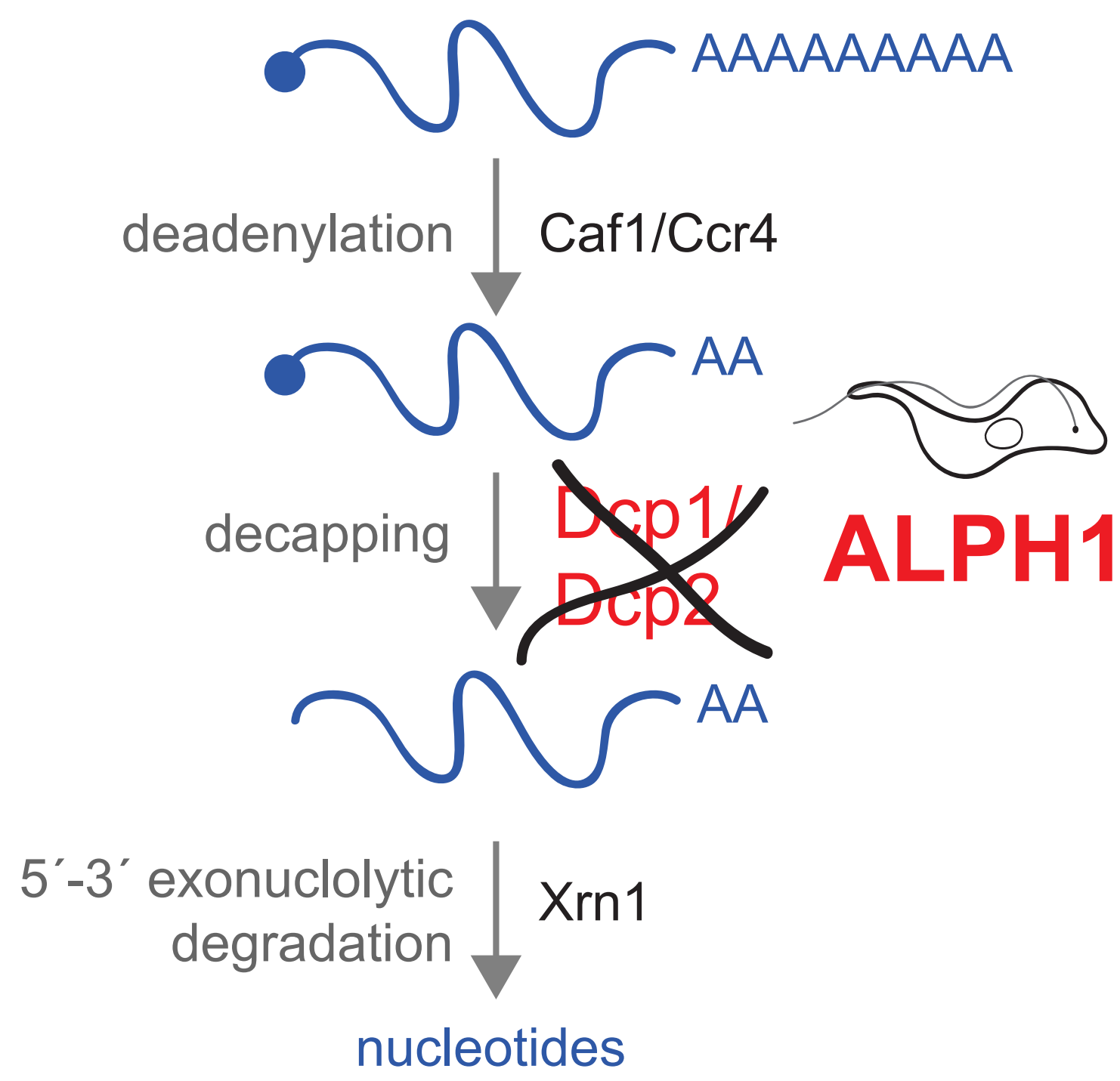


# mRNA decapping by an ApaH-like phosphatase in trypanosomes

Paula Andrea Castañeda-Londoño<sup>1</sup>, N Banholzer<sup>1</sup>, C M do Nascimento Moreira<sup>1</sup>, N Zhang<sup>2</sup>,  
M Zoltner<sup>3</sup>, M Field<sup>2</sup>, F B Holetz<sup>4</sup>, B Dallagiovanna<sup>4</sup>, N Karolak<sup>5</sup>, M W Górna<sup>5</sup>, S Kramer<sup>1</sup>

<sup>1</sup>Biozentrum, Lehrstuhl für Zell- und Entwicklungsbiologie, Universität Würzburg, Germany; <sup>2</sup>School of Life Sciences, University of Dundee, Dundee, UK;  
<sup>3</sup>Faculty of Science, Charles University, BIOCEV, Prague, Czechia; <sup>4</sup>Carlos Chagas, Fundação Oswaldo Cruz-Fiocruz, Curitiba, Brazil;  
<sup>5</sup>Biological and Chemical Research Centre, University of Warsaw, Poland

## Background



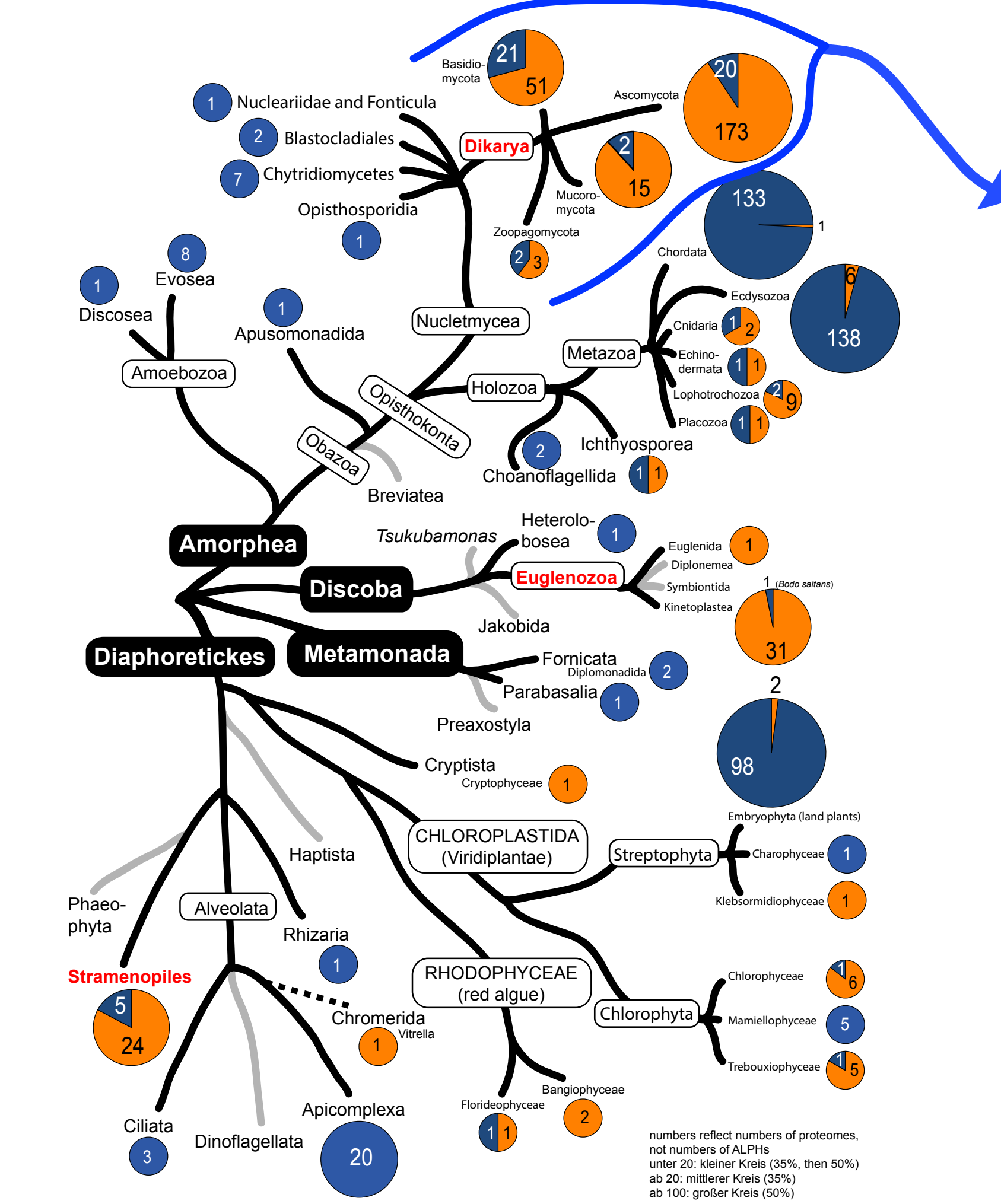
mRNA decapping is usually done by the nudix domain protein Dcp2 of the Dcp1/2 complex.

Kinetoplastida are the only eukaryotes with no homologues to Dcp1/2.

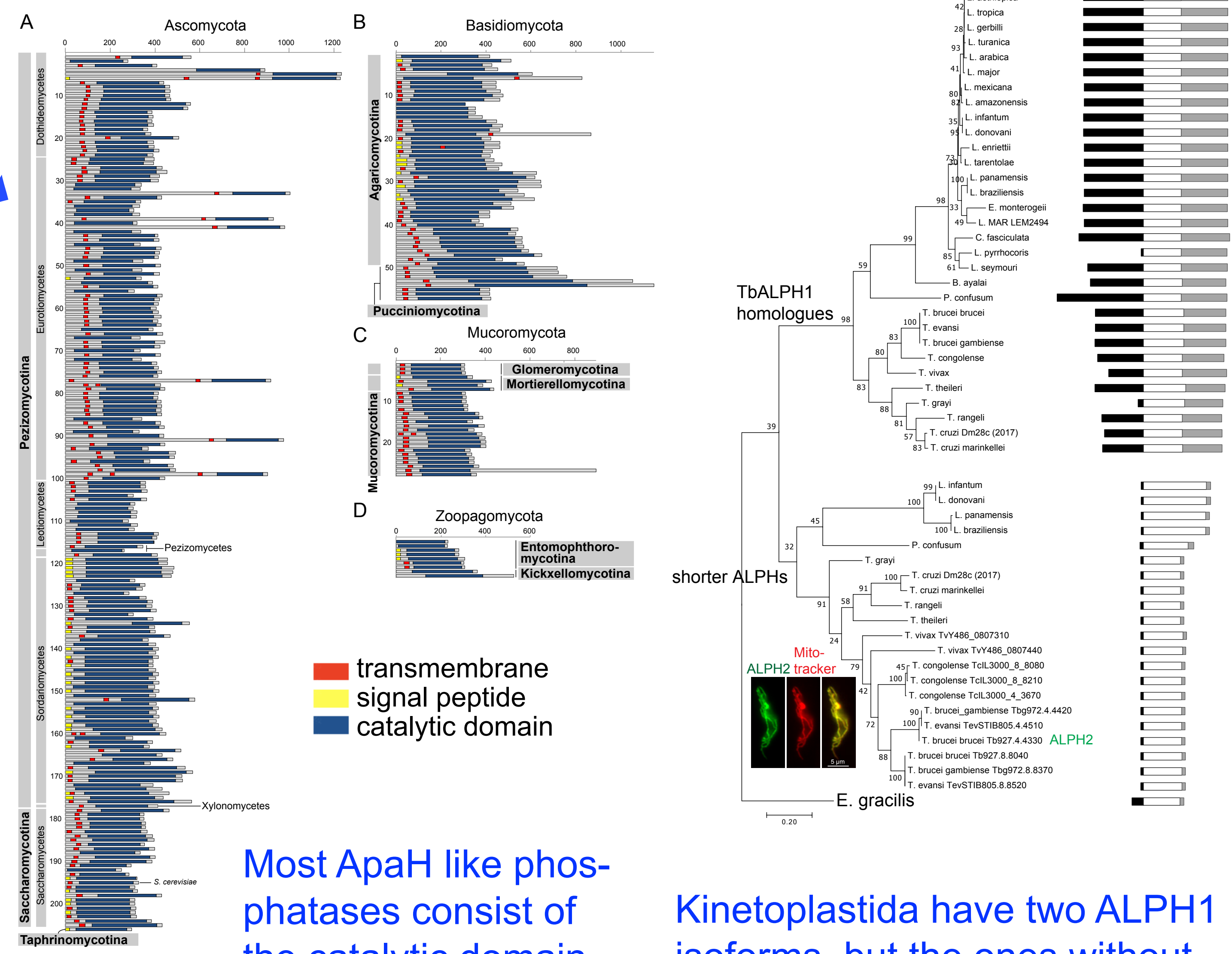
Instead, they use an **ApaH like phosphatase (ALPH1)**, a protein of an ancient protein family from bacteria, closest related to the PPP type of protein phosphatases (Kramer, S. PLoS Pathog 13, e1006456 (2017)).

## The usage of an ApaH like phosphatase as mRNA decapping enzyme appears restricted to trypanosomes.

organisms with ALPH  
organisms without ALPH



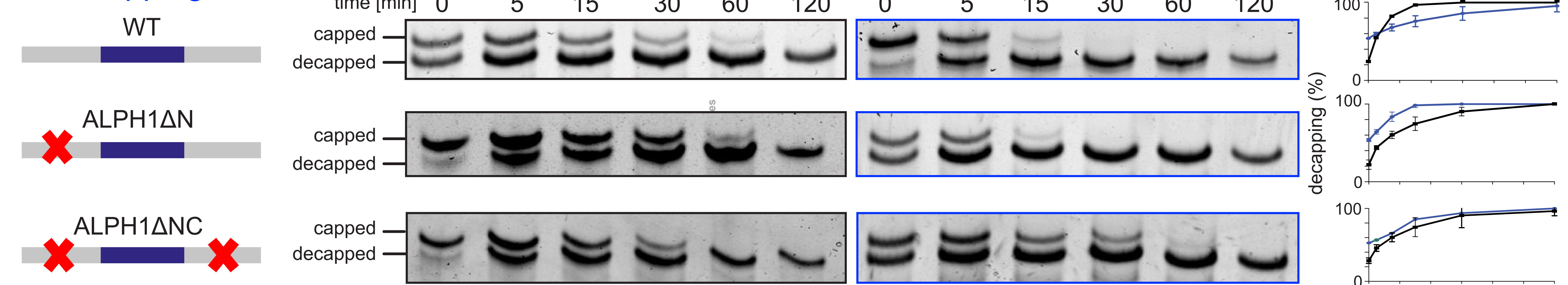
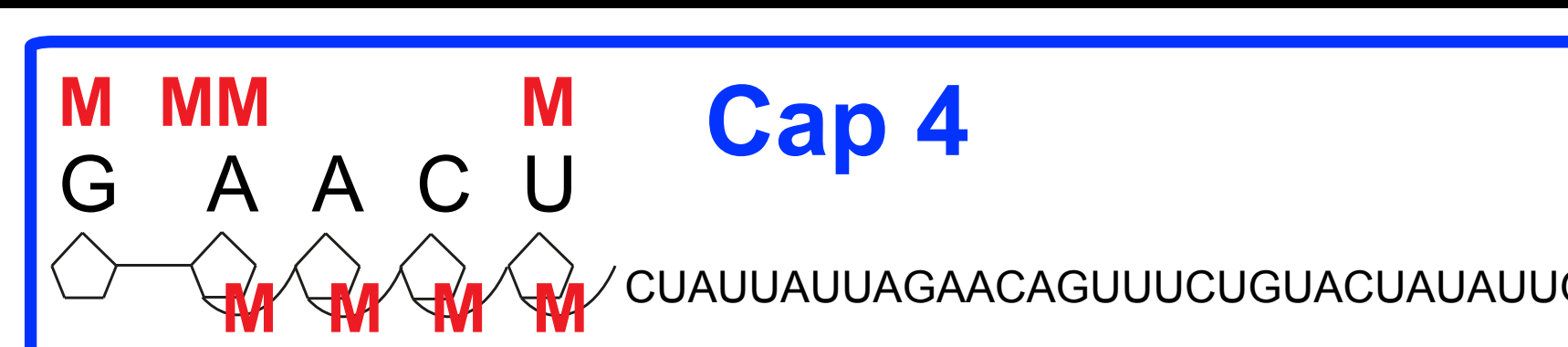
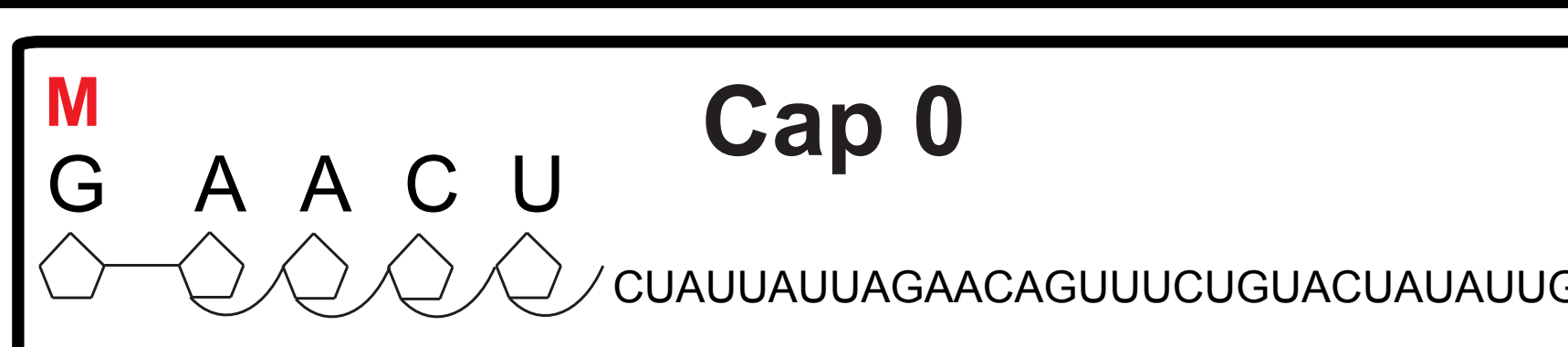
ApaH like phosphatases are widespread throughout all eukaryotes in a patchy way.



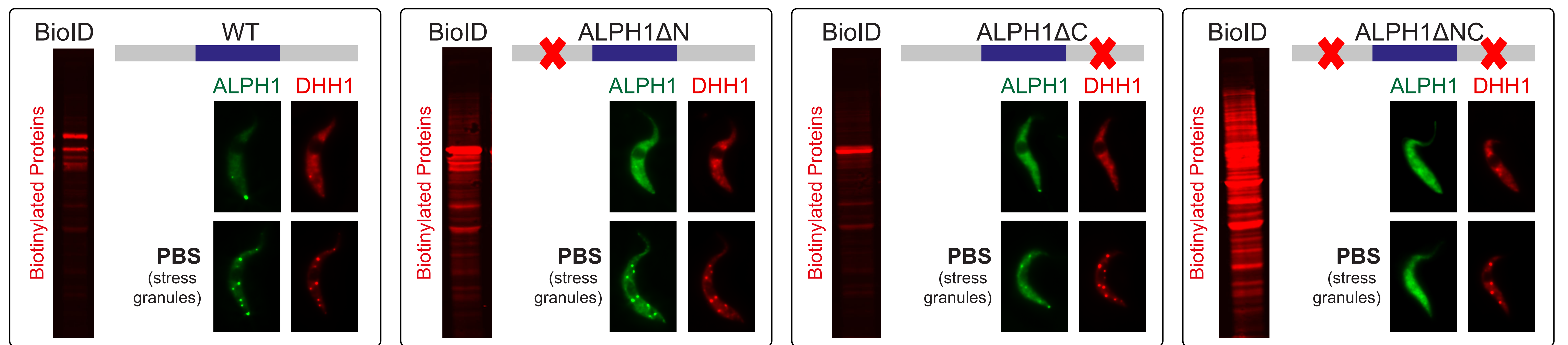
Most ApaH like phosphatases consist of the catalytic domain only, and are likely not cytoplasmic.

Kinetoplastida have two ALPH1 isoforms, but the ones without N- and C-terminal extension are likely not in the cytoplasm.

## The catalytic domain of ALPH1 is sufficient for decapping *in vitro*.



## The N and C-terminal extensions are important for ALPH1 localisation and protein interactions.



ALPH1 ΔN/- is viable, but BSF cells grow slower.

