

Novel Z-DNA binding domains in giant viruses

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Z-nucleic acid structures play vital roles in cellular processes and have implications in innate immunity due to their recognition by Z α domains containing proteins (Z-DNA/Z-RNA binding proteins, ZBPs). Although Z α domains have been identified in six proteins, including viral E3L, ORF112, and I73R, as well as, cellular ADAR1, ZBP1, and PKZ, their prevalence across living organisms remains largely unexplored. In this study, we introduce a computational approach to predict Z α domains, leading to the revelation of previously unidentified Z α domain-containing proteins in eukaryotic organisms, including non-metazoan species. Our findings encompass the discovery of new ZBPs in previously unexplored giant viruses, members of the *Nucleocytoviricota* phylum. Through experimental validation, we confirm the Z α functionality of select proteins, establishing their capability to induce the B-to-Z conversion. Additionally, we identify Z α -like domains within bacterial proteins. While these domains share certain features with Z α domains, they lack the ability to bind to Z-nucleic acids or facilitate the B-to-Z DNA conversion. Our findings significantly expand the ZBP family across a wide spectrum of organisms and raise intriguing questions about the evolutionary origins of Z α -containing proteins. Moreover, our study offers fresh perspectives on the functional significance of Z α domains in virus sensing and innate immunity and opens avenues for exploring hitherto undiscovered functions of ZBPs.

Nucleic acids adopt a variety of structures other than right-handed double helices. Both DNA and RNA can form triplexes, I-motifs, G-quadruplexes, and left-handed helices referred to as “Z-nucleic acids” (Z-DNA and Z-RNA) (1–3). These alternative structures are critical components of cellular functions, influencing a myriad of biological processes (4–6). In particular, it is now well established that the innate immune system exploits Z-nucleic acids as pathogen-

associated molecular patterns and damage-associated molecular patterns (7, 8).

Z-DNA/Z-RNA biology owes its now-recognized relevance to several proteins containing Z α domains, which are responsible for recognizing Z-nucleic acids (9). Z α domains are found in metazoan proteins involved in immunity and cancer, as well as in viral proteins. The first crystal structure revealed that the Z α domain of ADAR1 (adenosine deaminase acting on RNA 1) adopts a winged-helix-turn-helix (wHTH) fold. This structure is characterized by a compact α/β architecture encompassing a three-helix bundle (α 1 to α 3), juxtaposed to a twisted antiparallel β -sheet (β 1 to β 3) (10). Specific amino acids are important for the specific recognition of Z-nucleic acids, such as N173 and Y177 within helix α 3, and P192 and P193; W195 contributes to both protein stability and nucleic acid binding (10). This earlier work demonstrated that the commonly found wHTH domain across proteins only binds to Z-nucleic acids when these amino acids are present.

The only other proteins known to contain Z α domains are ZBP1 in mammals, amphibia, and reptilia (11), PKZ in salmoniform and cypriniform fish (12), E3L in poxviruses (13), ORF112 in cyprinid herpesviruses (CyHV-1-to-3) (14) and I73R in the asfarviridae African swine fever virus (ASFV) (15). Our current understanding of the biology of these proteins points to antagonistic relationships in innate immunity and a potential regulatory role in gene expression (16, 17). Briefly, the recognition of Z-nucleic acids by ADAR1 and ZBP1 is essential for the immune system's ability to detect and respond to viral threats effectively (18). Conversely, viruses have evolved Z α -containing proteins like E3L to subvert and antagonize host immune defenses, thereby enhancing their own virulence and survival within the host organism (19, 20). However, Z α domains have functional roles that need to be nuanced, as they cannot always be swapped, especially across different viruses (20, 21). This level of complexity implies an evolutionary balance between a virus and its host, where the unique characteristics of Z α domains are finely tuned to suit the particular dynamics of each virus-host relationship.

Because Z-nucleic acid biology is now viewed as a determinant for cell fate during infection and auto-immune diseases, one could ask the question of whether more proteins

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with Z α domains could be discovered, and in what species. Recent computational efforts based on structure similarity search have unveiled the existence of 14 potential Z α domains (22). Additionally, two other potential ZBPs were identified, namely RBP7910 in *Trypanosoma brucei* (23), and DprA in *Riemerella anatipestifer* (24). However, these studies did not address the absence of the complete set of required amino acids for binding Z-nucleic acids, and no experimental validations of these putative Z α domains were carried out.

Here, we introduce a computational approach for predicting Z α domains, which combines primary sequence analysis, three-dimensional modeling, and manual examination of the resulting hits through structural analysis. This method led to the discovery of 8 putative Z α domain-containing proteins in eukaryotic organisms, and 68 in giant viruses diverse and mainly unexplored giant viruses in the viral phylum *Nucleocytoviricota*. We also report Z α -like domains in bacterial proteins, which deviate somewhat from their eukaryotic and viral counterparts in primary amino acid sequence. Finally, we have experimentally validated our predictions by assessing the Z-DNA binding capacity and ability to shift the B-Z equilibrium for two Z α and two Z α -like candidates. This work offers fresh insights into the functional role of Z α domains in virus sensing and innate immunity. Furthermore, it may shed light on previously undiscovered functions of ZBPs, expanding our understanding of their roles in biological processes.

Results and discussion

A combinatory method to predict Z α -domain and Z α -like candidates

Initially, we employed BLAST search tools in conjunction with primary sequence analysis by domain prediction software SMART and PROSITE to identify the Z α -domain candidates. Subsequently, 3D structure analysis and prediction tools were used to refine our predictions. Leveraging our existing knowledge of the essential attributes of the Z α -domain, we further refined the results derived from our *in silico* analysis (Fig. 1A). We used multiple sequence alignments to compare the candidate sequences with some representative Z α domains in all 6 known ZBPs. We checked the presence of residues crucial for the functionality of the Z α -domain by primary sequence alignment. Among these, the NxxxY motif within α 3 is essential for Z-DNA binding, while the pPxW motif in the β -wing plays a significant role in both Z-DNA (10) binding and A-to-Z conversion (25). In the crystal structure of the HsZ α _{ADAR1}, tryptophan 195 has been demonstrated to be vital not only for protein stability but also for DNA binding. The double mutation involving asparagine 173 and tyrosine 177 within the α 3 helix completely abolishes Z α 's capacity to bind Z-DNA, rendering it functionally inoperative for both mammalian and viral ZBPs. Additionally, proline 192 and proline 193 contribute significantly through van der Waals interactions with DNA (10). Despite the limited overall sequence similarity typical to Z α domains, the aliphatic residues (marked in red) from the three helices, along with Tryptophan 195 in the β -wing, exhibit strong conservation in

all candidates. These residues are well-established for their role in supporting the hydrophobic core of the winged helix-turn-helix structure (Figs. 1B, and S1) (10). In line with this, the predicted 3D structures, generated by AlphaFold (26), reveal that all candidates adopt the characteristic winged-HTH conformation. The 3D fold comparisons underscore a remarkable degree of structural similarity between these candidates and the crystallized ADAR1 Z α domain, PDB ID 2GXB (27) (Fig. 1, C and D). Despite these similarities, some candidates found in bacteria lack conserved residues in the corresponding positions N173 and Y177 in HsZ α _{ADAR1} displaying instead a serine and an asparagine respectively. As such, we classify them as Z α -like candidates (Figs. 1B and S1).

Identification of novel Z α -domain candidates in eukaryotic organisms

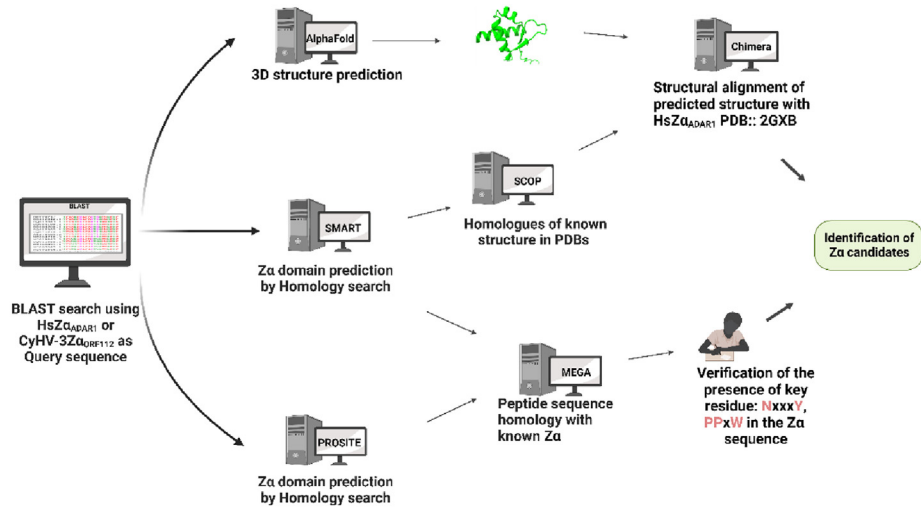
All predicted Z α candidates maintain the conserved residues N173 and Y177 in HsZ α _{ADAR1} crucial for Z-DNA binding (10). In Anthozoa, a class of marine invertebrates that includes sea anemones, stony corals, and soft corals, we predicted a new Z α containing protein. Interestingly, in some families such as the Poritidae or Actinidea, the novel Z α displays all important residues found in the known Z α . However, some families such as Acroporidae or Caryophylliidae display a Z α domain in which the Y177 is substituted by phenylalanine and W195 is replaced by a tyrosine (Fig. S1). We predicted other potential Z α domain-containing proteins in other eukaryotic organisms out of metazoan and DNA viruses, such as a protein annotated as transcription activator p15 in phytoplankton, and another annotated as acetate-CoA ligase in unicellular microalgae of the Symbiodiniaceae family (Figs. S1 and S3).

Identification of novel Z α -domain candidates in giant viruses

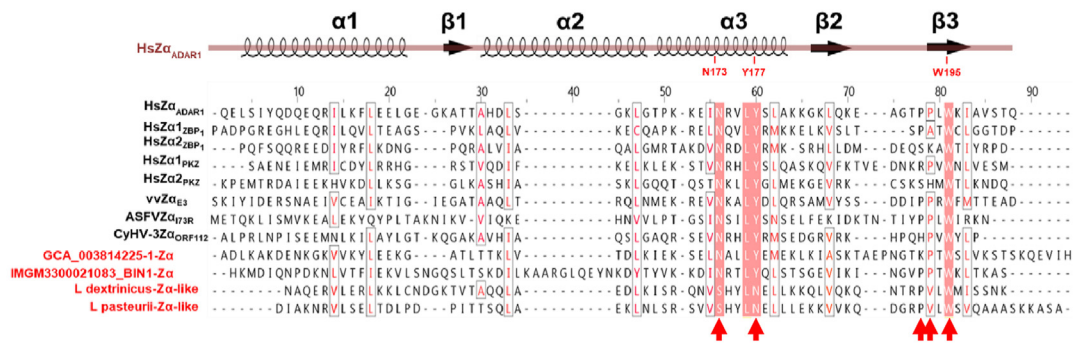
The most substantial contingent of novel ZBP candidates emerged within giant virus metagenome-assembled genomes (GVMAGs), revealing the presence of potential ZBPs in members of the orders pimascovirales, pandoravirales, and imitervirales. These proteins are distinct from previously documented ZBPs in fish-herpesviruses, poxviruses, and African swine fever virus (ASFV) (Fig. 2). These giant viruses were recovered in our previous metagenomic survey of samples from marine and freshwater, soil, and thermal environments (Fig. S2) (28), but also included additional sequences from deep-sea sediments (29) and permafrost (from 49 to 53,000 years old) (30). Our analysis shows that the candidate ZBPs in these giant viruses share similar domain architectures as in previously identified viral-containing proteins (13, 15, 20). The only exception is the viral ZBP candidate in which Z α is associated with tRNA metabolism domains (IMGM3300021083) (Fig. S3).

In our phylogenetic analysis, the clade topology in the ZBPs protein tree does not align with the topology found in the *Nucleocytoviricota* species tree (Fig. 2). This finding suggests that proteins with Z α domains have been independently acquired, likely multiple times, originating from a specific host *via* gene transfer. This possibility, as proposed by Kuś *et al.*,

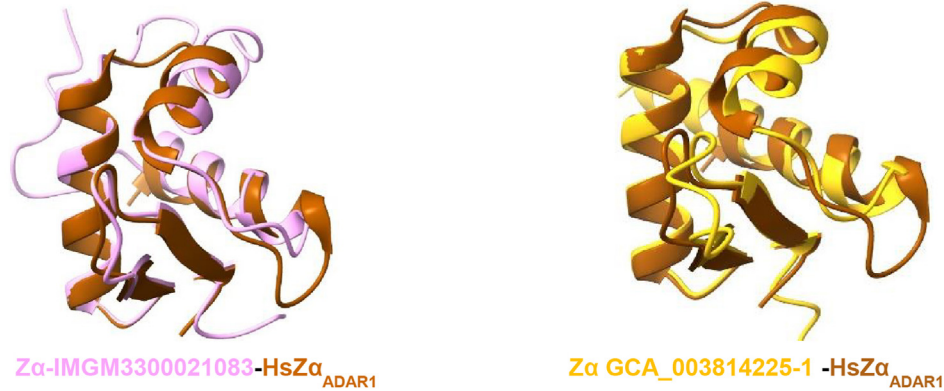
A



B



C



D



Figure 1. Prediction of new Zα candidates. A, the summary of the workflow for identifying Zα candidates. B, alignment of two predicted Zα and two Zα-like with known Zα domains from human and mouse ADAR1 (HsZα_{ADAR1}), human ZBP1 (HsZα_{ZBP1}), and viral from vaccinia virus and cyprinid-herpesvirus-3 (vvZα_{E3} ASFV Zα_{73R} CyHV-3Zα₁₁₂). The consensus structure Zα domain is shown at the top. The key DNA-interacting residues in ADAR1 (corresponding to Asn174, Tyr177, P192, P193, and Trp195 in HsZα_{ADAR1}) are pointed with red arrows. Hydrophobic residues in the hydrophobic core are highlighted in red front. C, structural alignment of AlphaFold predicted structure of Zα candidates with HsZα_{ADAR1} (brown, PDB ID 2GXB). D, structural alignment of AlphaFold predicted structure of Zα-like candidate with HsZα_{ADAR1} (brown, PDB ID 2GXB).

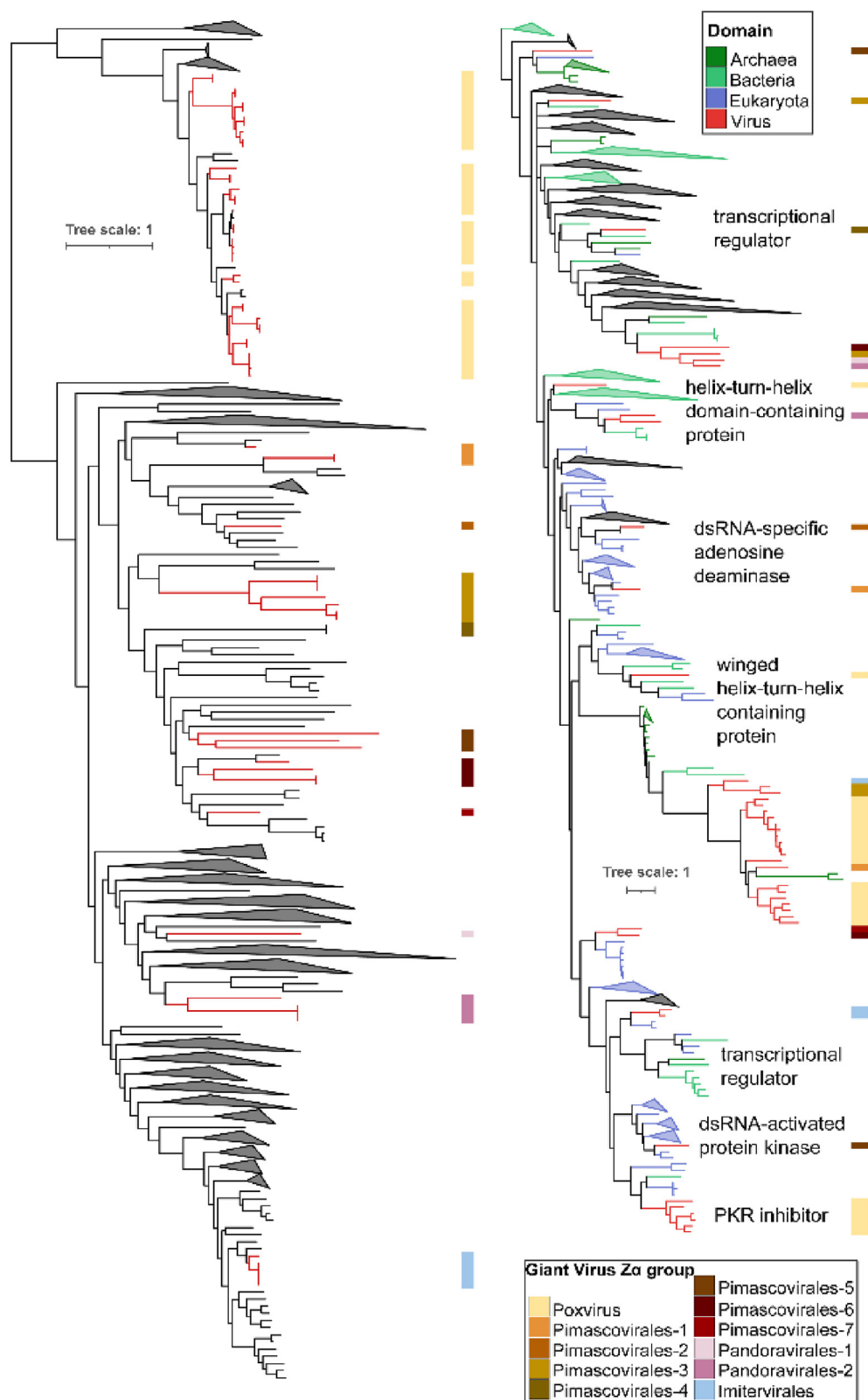


Figure 2. Phylogenetic tree of proteins that contain Z α and Z α -like domains. Proteins that were included in the analysis are giant virus ZBPs and homologs identified and extracted from the NCBI nr database.

suggests that CyHV-3 ORF112 might have been co-opted independently from the host by the common ancestor of the cyprinid herpesviruses, rather than through horizontal

transfer from a poxvirus (31). Such scenarios are further supported by the fact that in our tree, proteins from different giant virus lineages group together and are simultaneously

intertwined in monophyletic clades with eukaryotic and archaeal homologs.

Our results imply that giant viruses with the Z α domain likely infect a wide range of organisms ranging from dinoflagellates to marine vertebrates and arthropods as well as early diverging metazoans such as cnidarians (32). The presence of ZBP in diverse *Nucleocytoviricota* lineages raises the question of the role of Z α in the interactions of these viruses with their hosts, and how this fits into our current understanding of the role of ZBPs in animal innate immunity. Furthermore, the manifold presence of ZBPs in divergent giant virus lineages, known to infect protists and algae, sparks curiosity. This observation is particularly intriguing given that virus-protist interactions entail mechanisms distinctly different from the innate immunity found in animals (33). Could the presence of ZBP indicate that these giant viruses infect metazoan hosts? In line with this, the ASFV infects two phylogenetically distant hosts, functioning as an arthropod-borne DNA virus (34), significantly impacting pig farming due to its high virulence (19).

Experimental validation of Z α and Z α -like candidates

To validate our prediction, we experimentally studied some of these Z α and Z α -like domains by circular dichroism, an established method to probe their ability to induce B(A)-to-Z conversion (35). We found that two predicted Z α domains from two distinct members of the viral order pimascovirales are able to efficiently convert B-DNA duplex of different lengths d(CpG)₃ and d(CpG)₆ to Z-form (Fig. 3, A and B). Interestingly, we found that these two Z α domains are not able to induce A-to-Z conversion of RNA duplex as seen for the positive control HsZ α _{ADAR1} or in the high salt concentration condition (Fig. 3C). Consequently, our study identifies new cases of Z α capable of converting B-DNA to Z-form but not A-RNA. Indeed, the conversion of A-RNA to Z-RNA is a less favorable and energetically costly process compared with B-DNA to Z-DNA conversion (35). To test the binding capacity to sequence contexts that have a lower B-Z energy barrier or preformed Z-form nucleic acids, we used a singly methylated RNA duplex 8mG4 r(CpG)₃ which stabilized 50% of the population in Z-form (36, 37) (Fig. 3D). In this case, the Z α domains are capable of flipping the RNA to Z-form. This indicates that Z α domains bind to the pre-stabilized Z-RNA since they cannot flip A-RNA to the Z-form.

It is interesting to speculate on the distinction between competent A/B-to-Z flipping Z α domains, Z α domains that can only flip B-DNA, and the nonfunctional Z α -like domains, and their evolutionary/functional relevance. Numerous studies have tried to identify key residues/regions within the Z α domain that are important for binding and flipping nucleic acids to the Z-conformation. Kim *et al.* were able to restore the partial function of ADAR1's Z β domain by mutating Ile335 back to a tyrosine as is typical for all other ZBDs (38). Additionally, residues within the β -wing play a large role in not only the stabilization of the Z-conformation but also the rate at which it converts nucleic acids to the Z-conformation (39, 40).

Furthermore, distantly related WTH domains with no ability to convert nucleic acids to the Z-conformation can be converted into good Z-binders and converters through directed mutagenesis (41). It is likely that numerous factors contribute to the ability of any given Z α domain's ability to bind and convert nucleic acids to the Z-conformation, and the loss of one stabilizing interaction may result in the inability to convert A-RNA while still retaining the ability to convert B-DNA.

As expected, we found that for the Z α -like domain, there was no conversion of B-DNA to Z-DNA or A-RNA to Z-RNA, in contrast to our positive control HsZ α _{ADAR1} (Fig. 3, E and F). To test the binding capacity to sequence contexts that have a lower B-Z energy barrier or preformed Z-form nucleic acids, we used a methylated DNA duplex d(m⁵CpG)₃ and the singly methylated RNA duplex 8mG4 d(CpG)₃ respectively (36, 37). The Z α -like domains show no ability to stabilize or shift the B-Z transition equilibrium of the Z-nucleic acid analogs (Fig. 3, G and H). The Z α -like domain is found in the bacterial protein annotated as sigma 54 (σ^{54})-interacting transcriptional regulator. σ^{54} also encompasses an ATPase AAA region, which overlaps with the Holliday junction DNA helicase RuvB. RuvB is a component of the RuvABC revolvase, crucial for resolving Holliday junctions (42). Remarkably, both Holliday junctions and Z-DNA are found in bacterial biofilms (43, 44). Studies have shown that RuvA protein or DNABII stabilizes Holliday junctions, inducing supercoiling that reduces the energy needed for B-Z transition and promotes Z-DNA formation in the extracellular DNA biofilm matrix (43). Therefore, it would be interesting to test the B-to-Z conversion of the predicted association between the Z α -like domain and the helicase RuvB in σ^{54} . Moreover, knowing that bacteria of the genus of *Lactobacillus* are biofilm-forming bacteria (45), we hypothesize that σ^{54} could be involved in the biofilm matrix formation.

Experimental procedures

In silico prediction of Z α domain

To identify new Z α domain candidates, we developed a combinatorial approach. Using the Basic Local Alignment Search Tool, blastp algorithm of NCBI (46), we performed taxid-exclusion blast and taxid-targeted blast using the Z α peptides from CyHV-3 and HsADAR1 as the query sequences. The candidates with a minimum of three crucial residues conserved or featuring less drastic substitutions (such as in A *digitifera*-Z α , where phenylalanine replaces the tyrosine in helix- α 3 and a tyrosine replaces the tryptophane in the β -wing Fig. S1), were considered as the best hits and were selected. Their primary sequences were subjected to Z α domain prediction using SMART (47) and PROSITE (48, 49) software enabling us to determine the peptide length covered by the Z α domain. Leveraging our knowledge of the essential attributes of the Z α -domain, we further refined the results derived from our *in silico* prediction. We check the presence of certain residues crucial for the functionality of the Z α -domain by primary sequence alignment using MEGA-11 (50) and UGENE (51) tools. The presence or absence of these critical motifs served as

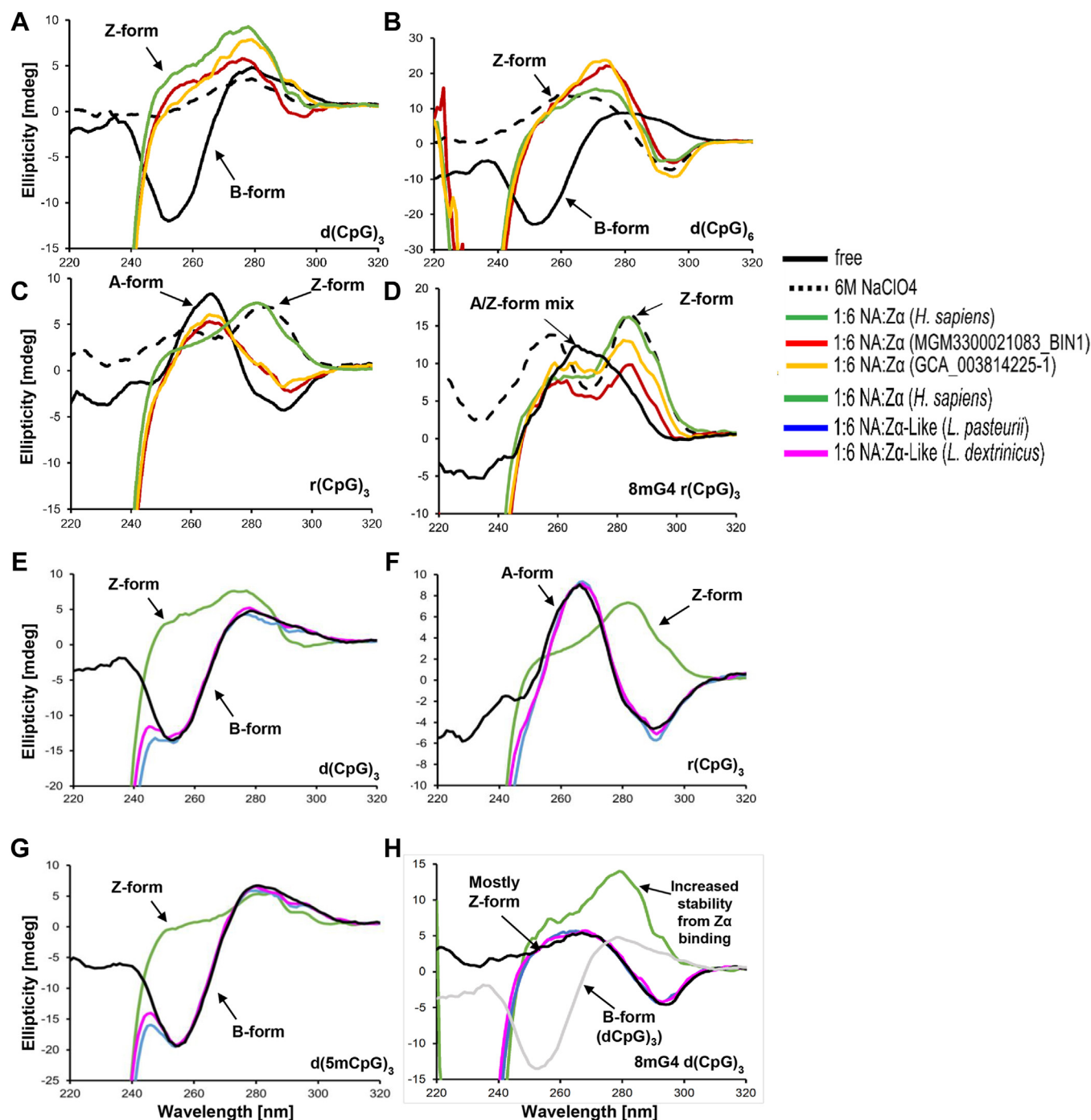


Figure 3. Experimental validation by circular dichroism. A and B, the predicted viral Z α domains induce B-to-Z conversion of DNA. C, viral Z α domains do not convert A-RNA to Z-RNA. D, viral Z α domains do convert an RNA with prestabilized Z-conformation (A:Z form ~50%/50%) to full Z-RNA. E and F, Z α -like domains are not able to induce B(A)-to-Z conversion. G and H, Z α -like domains do not bind to Z-nucleic acids. For comparison, the profile of a B-form DNA is shown in grey in (H), demonstrating the presence of a large portion of Z-form in 8mG4 d(CpG)₃. The profiles overlaid with profiles from second independent measurements are provided in Fig. S4.

the final determinant in our analysis, allowing us to confidently identify Z α candidates. Any candidates lacking tyrosine and asparagine were categorized as Z α -like, given their alignment with other defining characteristics. SMART software also outputs structure primary sequence homologs of known in the protein data bank (PDB) or blast with known SCOPs (structural classification of proteins) of the Z α , for example, d1qbj (52).

Finally, to strengthen the Z α prediction, a structure prediction using AlphaFold (26, 53) confirms the winged helix-turn-helix fold superposable with the crystallized Z α available with the PDBs using UCSF ChimeraX (54). With the predicted 3D structures, we performed a 3D-BLAST search (55) to confirm that the best hits are entries corresponding to the Z α domain. Figures 1A and S3 were created with BioRender.com.

Sequence source and annotation

Most of the giant viruses used in our analysis were identified in our metagenomic analysis performed in samples from different biotopes (S2) (28). We also included other giant viruses identified in other studies such as Pithovirus (29) and Marseillesvirus (30). The sequencing of the full viral genome or the viral genome fragment covering the coding sequence of the ZBP allowed the complete identification of the open reading frame corresponding to the ZBP. For the other organisms, the ZBPs are annotated at the PDB genomic level and (protein and mRNA level for *Micromonas pusilla* (56). Z α domain sequence and the ID numbers of ZBPs are listed in Table S6.

Phylogenetic analysis

Novel Z α and Z α -like domain sequences were clustered at 40% identity using cd-hit (57) and the resulting cluster representative sequences were aligned using mafft-linsi (version 7.505) (58). We used the hmmbuild (version 3.3.2, hmmer.org) to create a profile hidden Markov model from the representative sequence alignment and then used it as a query for the hmmsearch program (version 3.3.2, hmmer.org) on the NCBI nr database (59). Protein sequences that had an overall e-value <0.005 and contained at least one domain with length $\geq$ 60 amino acids were considered as significant hits to the model. Then we de-replicated the protein sequences with significant hits with cd-hit at 90% identity and an alignment length covering at least 60% the length of the longer sequence (-aL 0.6). The representative protein sequences were aligned together with Z α containing sequences from the viral phylum Nucleocytoviricota using the mafft FFT-NS-2 method (58). The alignment was trimmed with trimal with the -gt 0.1 option (60). A protein tree was built using IQ-Tree (version 1.6.12) (61) with 1000 ultra fast bootstrap replicates (62) and the JTT+R10 model, which was selected with the ModelFinder feature in IQ-Tree (63). The tree was visualised using iTol (64).

To build the giant virus species tree the nsgtree pipeline was used (<https://github.com/NeLLi-team/nsgtree>) on representative genome of the viral phylum Nucleocytoviricota (65) and additional GVMAGs that contained proteins with a Z α domain. In brief, 7 giant virus orthologous groups (GVOGs) were identified using hmmsearch (version 3.1/b2, hmmer.org), extracted GVOGs were aligned with mafft (version 7.31) (58), trimmed with trimal (-gt 0.1, v1.4) and concatenated. A species tree was built from the supermatrix alignment using IQ-Tree (version 2.03) (66) with LG+G4 + I + G4 and visualized in iTol (64).

Protein expression and purification

The Z α domains from *Homo sapiens* ADAR1, *L. pasteurii*, *L. dextrinicus*, giant virus (IMG3300021083 and GCA_003814225-1) were cloned into the pet-28a(+) plasmid (N-terminal 6x His-tag and thrombin cleavage site between His tag and the Z α sequence). The plasmids were transformed and expressed in BL21(DE3) *E. coli* cells. The cell cultures were grown in Luria Broth (LB) to an OD₆₀₀ of $\sim$ 0.4 and induced with IPTG to a final concentration of 1 mM and allowed to express Z α for 4 h at 37 °C, then centrifuged to

collect the cell pellets. Cell pellets were resuspended in lysis buffer (50 mM Tris-HCl (pH 8.0), 300 mM NaCl, 10 mM imidazole, 1 mM BME) and sonicated. Lysate was centrifuged and the supernatant was applied to a His-trap column, (50 mM Tris-HCl (pH 8.0), 1 M NaCl, 10 mM imidazole, 1 mM BME), and eluted in 20 ml of elution buffer washed with 40 ml of lysis buffer, 80 ml of wash buffer (50 mM Tris-HCl (pH 8.0), 300 mM NaCl, 500 mM imidazole, 1 mM BME). The eluents were concentrated to $\sim$ 2 ml and applied to a HiLoad 16/600 Superdex 200 prep grade Gel Filtration Column (GE Healthcare) and the peak corresponding to pure protein (which elutes at $\sim$ 80 ml in 20 mM potassium phosphate (pH 6.4), 300 mM NaCl) was collected and concentrated using an Amicon 3 kDa cutoff centrifugal filter (Millipore-Sigma, Burlington, MA). Z α was dialyzed and concentrated into 20 mM potassium phosphate (pH 6.4), 25 mM, 0.5 mM EDTA and concentrated to $\sim$ 2 mM using an Amicon 3 kDa cutoff centrifugal filter (Millipore-Sigma). Subsequent dilutions were made in 20 mM potassium phosphate (pH 6.4), 25 mM, 0.5 mM EDTA as needed for the different experiments. We confirmed that all recombinant proteins were free of contaminating nucleic acid (Fig. S5).

DNA and RNA constructs and preparation

The d(CpG)₃, d(CpG)₆, and d(5mCpG)₃ constructs were synthesized by Integrated DNA Technologies. The r(CpG)₃ construct was synthesized by Dharmacon (a part of Horizon Discovery). The 8mG4 d(CpG)₃ construct was synthesized by the Yale School of Medicine oligo synthesis resource. All nucleic acid constructs in this study were heat annealed prior to use at 95 °C for 5 minutes followed by slow cooling at room temperature for 30 min. For circular dichroism (CD) measurements, all of the constructs were 50 μ M in 20 mM potassium phosphate, 25 mM NaCl (pH 6.4), 0.5 mM EDTA, or the same buffer but with 6 M NaClO₄ or a 1:6 M ratio of nucleic acid:Z α .

Circular dichroism

All CD measurements were collected using a JASCO J-815 CD spectrometer (run using Spectra Manager version 2 (JASCO)) in a 0.1 cm quartz cuvette. Z-form adoption by high-salt and Z α binding was carried out by incubating the constructs in 6M NaClO₄ or with saturating amounts of Z α at a 1:6 M ratio of nucleic acid:Z α for 1 h at 42 °C before cooling to 25 °C and measuring. Spectra were collected in 1-nm steps from 320 to 220 nm with an average of two scans. All measurements with unmodified nucleic acids as well as d(5mCpG)₃ and 8mG4 r(CpG)₃ were conducted twice, and the profiles shown in Figure 3 overlaid with profiles from second independent measurements are provided in Fig. S4. All duplicate profiles confirmed the high reproducibility of the measurements.

Conclusion

In summary, our study employed a computational combined with an experimental approach that enabled the

identification of potential Z α -containing proteins across various organisms. These ZBPs open up new avenues in studies of Z α domains and Z-nucleic acids in virus sensing and innate immunity. The domain architectures within predicted Z-DNA binding proteins imply that they potentially serve functions distinct from those currently attributed to the known ZBPs (Fig. S3). These functions could be conserved in mammals but in the form of synergy between a ZBP and another protein like the cooperative function between ADAR1_{p150} and DHX9 (67). Furthermore, our experimental validation of two Z α -containing proteins confirms not only their presence in currently uncultivated giant viruses but also their functionality. Our findings open new questions regarding the role of Z α domains in these giant viruses and suggest that the interplay between ZBPs/Z-nucleic acid is an evolutionarily ancient feature as shown for the cGAS/STING pathway (68). It is tempting to speculate that the ZBPs we predicted in giant viruses and early diverging metazoan could be involved in an ancient pathogen detection mechanism that could provide functional and evolutionary insight into innate immune signaling pathways in animals. Our validation results underscore the need for heightened caution in Z α domain prediction. We demonstrate that the mere detection of a Z α domain by prediction tools or the observation of structural similarities between predicted wHTH and crystallized Z α domains fall short of substantiating experimental B(A)-to-Z conversion or Z-nucleic acid-binding. These findings emphasize that the ability to bind to Z-nucleic acid should be the definitive criterion for establishing a wHTH as a true Z α domain.

In our study, we focused on validating the Z α and Z α -like domains found mainly in multi-domain proteins. Although we demonstrate that wHTH domains we classify as Z α -like cannot bind Z-nucleic acid nor convert B-to-Z, we cannot exclude the possibility that the proteins with Z α -like domains may still be involved in Z-nucleic biology. This is because other proteins lacking Z α domains have been found to bind Z-DNA (69) or play a role in regulating Z-RNA sensing (70).

Data availability

The data that support the findings of this study are available from the corresponding authors upon request. The primary sequence and the id-numbers of the new ZBPs and Z α -like containing proteins are summarized in Fig. S3 and Table S6. Other sequences used in this work are available from Frederik Schulz and Mamadou Amadou Diallo upon reasonable request.

Supporting information—This article contains supporting information.

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Conflicts of interest—The authors declare that they have no conflicts of interest with the contents of this article.

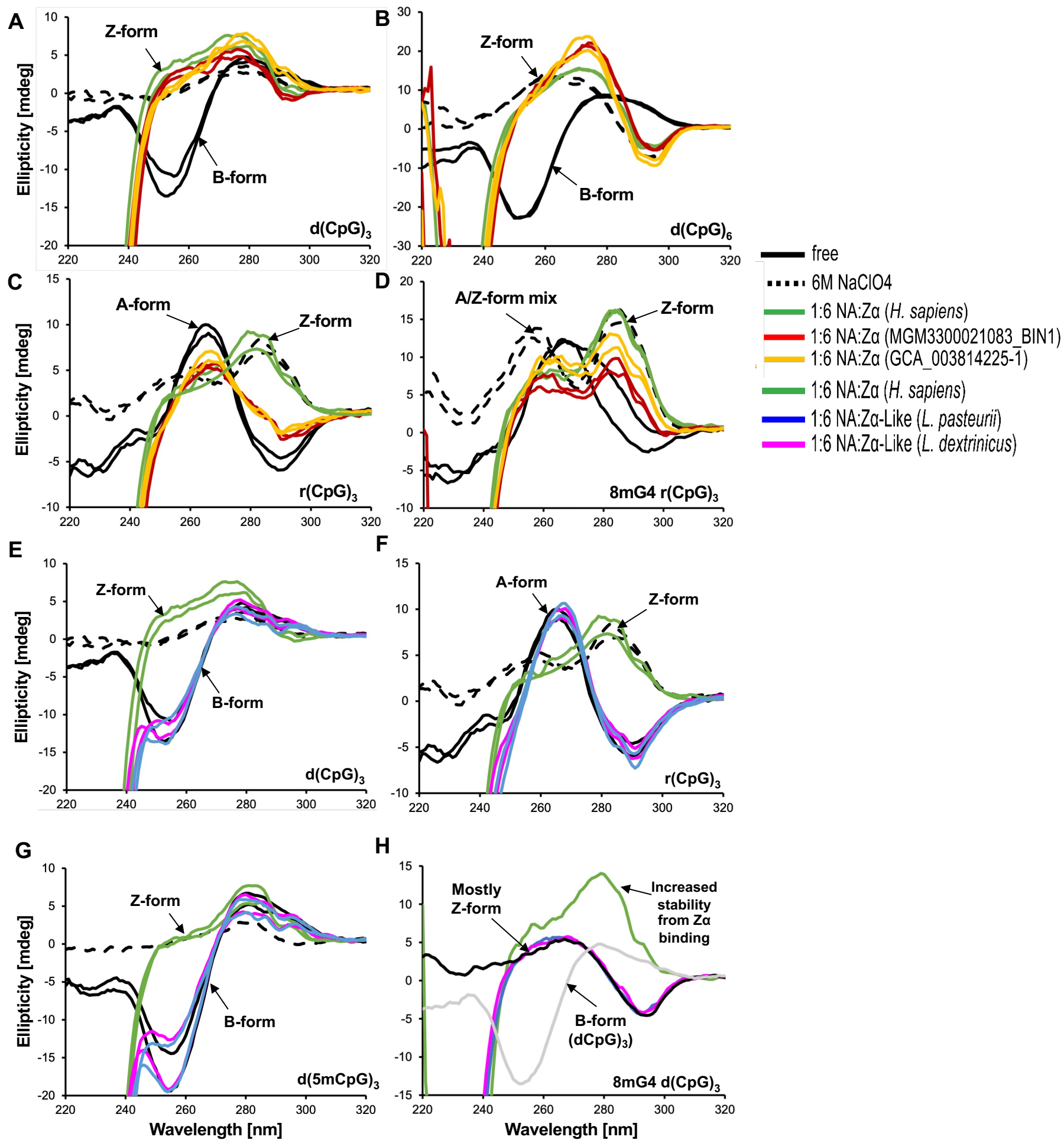
Abbreviations—The abbreviations used are: wHTH, winged-helix-turn-helix; ZBPs, Z-RNA binding proteins.

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Supplemental Figure 1: Alignment of new Z α with known eukaryotic Z α (HsZ α _{ADAR1} and Mm Z α _{ADAR1}) and viral Z α (ASFVZ α _{I73R} vVZ α _{E3} and CyHV-3Z α _{ORF112}). The consensus structure Z α domain is shown at the top. The key DNA-interacting residues in ADAR1 (corresponding to Asn174, Tyr177, P192, P193 and Trp195 in HsZ α _{ADAR1}) are pointed with red arrows. Hydrophobic residues in the hydrophobic core are highlighted in red front. Validated Z α in this study are indicated in green front and those that are validated as Z α -like are in red front.

Supplemental Figure 2: Giant viruses' environmental distribution. Each nucleocytoplasmic large DNA virus (NCLDV) is facing its sampling environment.

Supplemental Figure 3: Domains architecture of several representative cases of new ZBPs. Domains are predicted by SMART[®].

Supplemental Figure 4: Overlay of CD profiles of two independent experiments.

Supplemental Figure 5: Purity of expressed Z α domains. **(A):** Purified recombinant proteins were tested for protein contamination by SDS-PAGE using Novex[™] 4-20% Tris-Glycine Plus WedgeWell[™] Gels. Each lane shows one band corresponding to the correct expected molecular weights shown in the table below. **(B):** Purified recombinant proteins were tested for RNA contamination by spectral analysis using a NanoDrop 2000. Each recombinant protein was free of contaminating nucleic acid as shown by the 260/280 ratios (~0.60 is expected for pure proteins whereas values of ~2.00 is expected for RNA). Full spectral traces for each measurement are provided.

Supplemental table S6: Summary of information on the available sequences of Z α and Z α -like used in this study

S1

Anthozoa

Phytoplankton

Symbiodiniaceae

Viral

Bacteria

HsZa_{ADAR1}HsZa_{ADAR1}MmZa_{ADAR1}

A tenebrosa-Za

A digitifera-Za

M pusilla-Za

M commoda-Za

O tauri-Za

P provasolii-Za

C goreau-Za1

C goreau-Za2

S microadriaticum-Za

ASFV-173R-Za

VvZa_{E3}CyHV3Za₁₁₂

LCPAC304-Za

LCPAC406-Za

GVMAG-S-1025707-429-Za1

GVMAG-S-1025707-429-Za2

IMGM3300010938 BIN245-Za1

IMGM3300010938 BIN245-Za2

IMGM3300008255 BIN298-Za

CAH6421103.1-Za

IMGM3300010938 BIN801-Za

GCA_003814225-1-Za

IMGM3300006467 BIN4-Za

IMGM3300014493 BIN257-Za

GVMAG-M-3300005987-15-Za1

GVMAG-M-3300005987-15-Za2

IMGM3300021083 BIN1-Za

GVMAG-M-3300020182-33-Za

IMGM3300019458 BIN240-Za

IMGM3300042450 BIN843-Za

IMGM3300042541 BIN490-Za

IMGM3300037467 BIN618-Za

IMGM3300014838 BIN820-Za

IMGM3300023174 BIN273-Za

IMGM3300009183 BIN399-Za

IMGM3300021079 BIN283-Za

L dextrinicus-Za-like

L pasteurii-Za-like

L johnsonii-Za-like

L plantarum-Za-like

L rhamnosus-Za-like

S terrae-Za-like

 $\alpha 1$ $\beta 1$ $\alpha 2$ $\alpha 3$ $\beta 2$ $\beta 3$

	10	20	30	40	50	60	70	80	90	100	110	120
	QELSIYQDQEQ-R	I L K F L E E		L G E G K A T T	A H D L S G K L G T P			K K E I N R V L Y	S L A K K G K L Q K E	-- AGT - P	P L W K I A V S T Q	
	ELSIQSPEQ-KV	L N R L E E		L G E G K A T T	A H V L A R E L R I P			K R D I N R I L Y	S L E K K G K L H R G	-- RGK - P	P L W S L V P L S	
	SNLELNXLQK-S	I V E V M T Q		Y N R P L T T L E	I S K A I G L T T			K R D V N S A L Y	A M E R A G M I Q R V	-- KYQ - P	P V W S L K S R K	
	SDDVPLDRGEK-R	I F E L L R N		N G P L V A G D I	A T L G V D T			E D R V N H C L Y	T L Y K R G L V E I S	-- RSS - P	P L Y A V S D S F	
	AAPGAPVPEDK	L L A A L Q L		K G D D G - A T	A K E L A G F L K I G			K S E V N K A L Y	R L Q S S G R V R S D	- G A G G G - A	P T W R V G S S S	
	VPEDK	L L A A L T L		K G D D G - A T	A K E L A G F L K M G			K S E V N K A L Y	R L Q S S G R V R S D	- G A G G G - A	P T W R V G S S S	
	ATPPSPAVAESR	V L A A L R A		L G E D G - G C	A K T I A E F L K R D			K G A V N K I L Y	A A K G R G L A R C A E S E R G A	- P R W F I A D G C		
	SSSTTPEET-IV	L S A I S D		G M P - R K	A Q E I A K A T K L E			K K V V N S I L Y	K L A S R G L A V K H	- E D F T -	-- W T A K G A P	
	GTSVEDQSLKQ-R	I L E Y L K A		T G D W V P L S T	I V K V S K K W G A T K			K K D V N R C L Y	A L K D Q G L V E L I T T E N A N	- P Q W K A L A P G		
	KGGVDVDPTE-K	I M D Y F R R		T A G T E S T S L A	I A K A V I G K D G T			A S N V N R F L Y	S L K N K G Q I E L K	-- K K D G K	P Y W K L A G V Q	
	YNAVNMDYET-S	I L C Y V R E		K L V V T T K Q	I A D H F E V P			K K A V N R I L Y	E M E E F G A V H S I	- I Q S - P	P T W S T D E S	
	METQKLIS-M	V K E A L E K		Y Q Y P - L T	A K N I K V V I Q K E H N V V L P			T G S I N S I L Y	S N S E L F E K I D K T	-- N T I Y P	P L W I R K N	
	SKIYIDERSNAEI	V C E A I K T		I G I E G - A T	A A Q L T R Q L N M E			K R E V N K A L Y	D L Q R S A M V Y S S	-- D D I - P	P R W F M T T E A D	
	ALPRLNPISEEM	N L K I L A Y L G T		K Q G - A K	A V H I A Q S L G A Q			R S E V N R H L Y	R M S E D G R V R K H P Q	-- H P V W Y L P		
	MESLLN-L	I V T Y L S D		K T E P - V P T L Q	I A K Y V L G S K A T			K K M V N P S L Y	T L Q K L G K V R K I A E E N G G	- N P R W I L T K		
	KLNEKDENVHQ-W	I L N Y L E D		K E A I D E Y S A I S P L S	I A K H V F G K K A T			K K T V N S Y L Y	T M K G D G K V N K I T E E H M K	- R P R W Y L S G		
	MASTLRD-E	L N V L Y S		V - D Q - I P K R E	L F S K F P T L T			S K E I N S E L Y	T L K Q G K V K K I	-- Q D V - P	P I W A L N A T P	
	NDDFADQI	Y E F I L E		Q G A V E L K K I	A Q H F K A D			K T A I N K I L Y	N T L A K Q G K V K K I	-- Q D G - P	P V W T A V E D E	
	MAELEQ-R	I L T L S S V V D P I T R Q P K A D G P I R P	M K T I D I A K E	L F G T N G E				S K M V N P T L Y	G L E K K G L L V K I A D E G G K	- N P R W T V A K P G		
	KLSDAEAKARI	L E V L R N		S S D P - V P T L Q	I S R A L Y G P K A T			R K M V N S Q L Y	S L S K N L V S K S S N P D G T	- N P H W S L V Q A P		
	MDEVQN-K	I L E Y L L E		C D E E - V P T L Q	I S K H V F G D K A T			K K M V N K L Y	A L E K E G S V K S S E E N G T	- K P K W R I V K N D		
	MQANAEDAEIER	- A V L A W F P G		A S G G A S A L E	L T R S V A G C R V H			K T D M N R V L Y	A L E K R G V L H R D	-- G K T - P	P R W T L A T T A	
	ASDADLET-K	I Q H A L Q T		A V H P - L P A K D I	C K L L G S S V H			R S D V N R V L Y	S L L K R S Q V V K N	- E G T - P	P T W G V N L A A S	
	ADLKAKDENKGV	V K Y L E E		K G A T L T T K L V T D	L K I E			K S E L N A L Y	E M E K L K I A S K T A E P N G T	- K P T W S L V K S T S K Q E V I H		
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	SEELTA-K	I V A A L K T		K P - Q T	A R N L V S S L K E A			K S A I N S I L Y	A L Q K K G I L Q K D	-- D S T - A	P V W S I V D Q D G	
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	HKMDIQNPDKNL	V T - F I E K V L S N		G - Q S - L T S K D I	L K A A R G L Q E Y N K D Y T V			K K D I N R T L Y	Q L S T S G E V I K I	-- N G V - P	P T W K L T K A S	
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	MASPWSKVEARE	- K V E E V F R S		H D Y P - I S A V W I	A N K T E M T			K K D V N G M L Y	Q M L K E G A V E K L	-- E E M S P	V L W R L T D T	
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	EEPL	T E Y L S K		S P A T A R E L V K H	F S E N K N L L N P K F F S H G T I L T R H V			V N S C L Y	Y L L H K K R V N K S	-- D T T - P	P V W T V N T D	
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	NAQER	- V L R L K K		L C N D G K T - V T A Q Q	L A E D L K I S			R Q N V S H Y L N	E L L K K Q L V Q K Q	-- N T R - P	V L W M I S S N K	
	DIAKNRV	L S E L T D		L P D P - I T T S Q L A E K	L N L S			R S V V S H Y L N	E L L L E K K V V K Q	-- D G R - P	V L W S V Q A A S K K A S A	
	DKGKN-NV	L T Y L N M		L S H D H A L T T A Q L A D S	L N L S			R S V V S H Y L N	E L F R E E K I Q K I	-- A G R - P	V K W T K K G S M	
	MVKAKY-RV	L D V L K Q		L V A D N E T N I S T E M V A T A	A K I T			R G V T S A Y L S	Q L H K E G L V R K Y	-- G T K - P	V Y W K L Q R R T	
	MNAIDKLY	V A I V R Q		F P N E R I T T N Q A A E A	V G L T			R G V T S S Y L S	K L E R A G K L L K T	-- G S R - P	V Y W Q V K H E H	
	KAQD-RV	Y E A L V H		V G N D G - V T A A Q	L A K K L E L S			R P V V S H Y L N	G L L K E E R A A K K	-- G T K - P	V Y W T A I A E R	

NxxxY

PPxW

Color code

Experimentally confirmed Za

New Za candidates

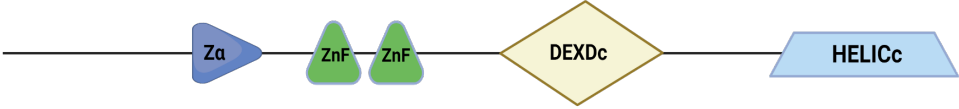
Experimentally confirmed Za-like

Za-like candidates

NCLDV-ID	Environmental distribution
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3300005573	Thermal springs
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3300023174	Freshwater
3300014838	Soil
3300009183	Freshwater
3300021079	Freshwater
3300014493	Soil
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3300010938	Thermal springs
3300042541	Freshwater
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003814225	Soil

Anthozoa

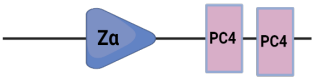
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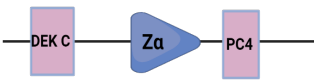
endoribonuclease Dicer

Phytoplankton

M commoda



M pusilla

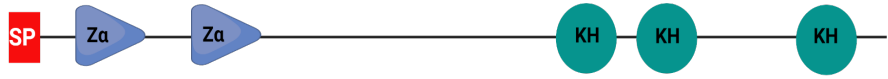


Z-binding domain-containing protein

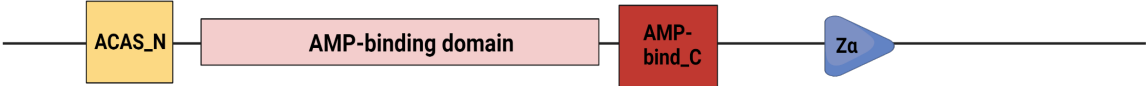
transcriptional coactivator p15

Symbiodiniaceae

Cladocopium goreau



S microadriaticum



acetate--CoA ligase

acetate--CoA ligase

NCLDV

GVMAG-M-3300014838-63



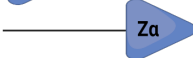
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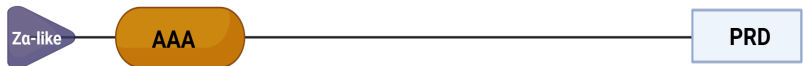


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Bacteria

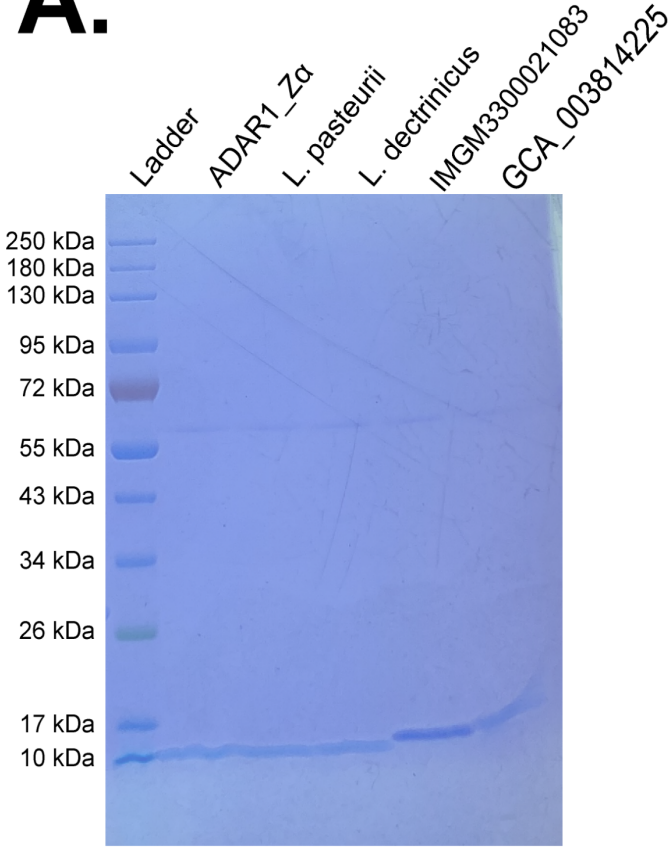
L dextrinicus



DNA translocase FtsK

sigma 54-interacting transcriptional regulator

A.

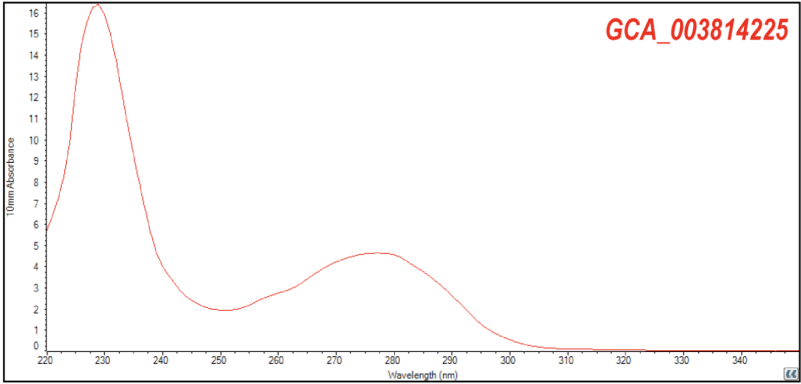
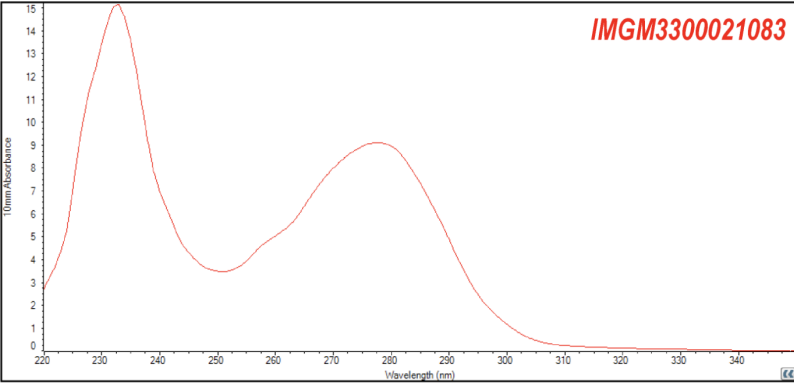
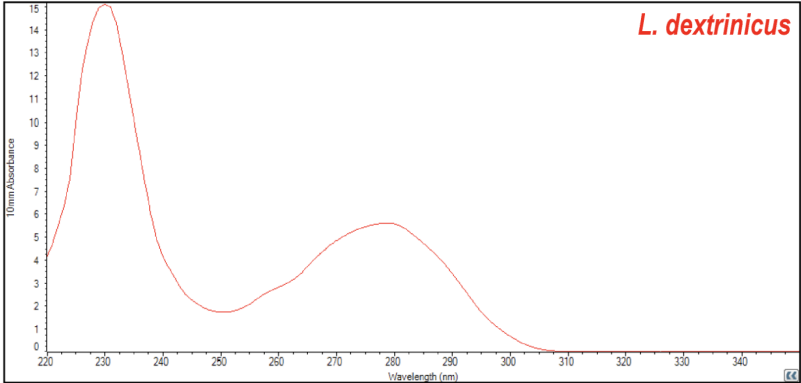
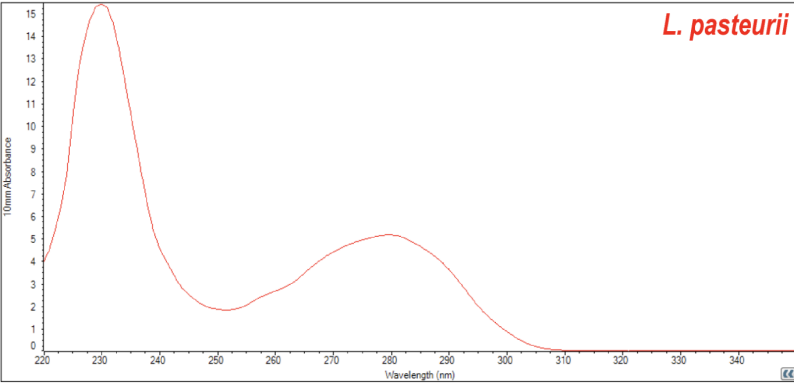
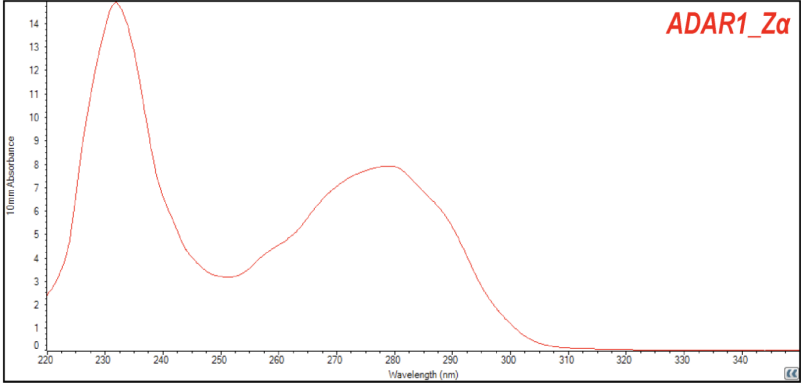


Expected Molecular Weights (kDa):

ADAR1_Zα	9.25
L. pasteurii_Zα	9.76
L. dextrinicus_Zα	10.01
IMG3300021083_Zα	11.49
GCA_003814225_Zα	11.00

B.

#	Sample ID	Protein	Unit	A280 (Abs)	260/280	Sample Type
1	ADAR1_Zα	7.893	mg/ml	7.893	0,57	1Abs = 1 mg/ml
2	L. pasteurii	5.163	mg/ml	5.163	0,51	1Abs = 1 mg/ml
3	L. dextrinicus	5.565	mg/ml	5.565	0,50	1Abs = 1 mg/ml
4	IMG3300021083	8957	mg/ml	8957	0,56	1Abs = 1 mg/ml
5	GCA_003814225	4547	mg/ml	4547	0,60	1Abs = 1 mg/ml



Name

>Z α -Like-Lapidilactobacillus dextrinicus
>Z α -Like-Lactobacillus pasteurii
>Z α -Like-Lactobacillus johnsonii
>Z α -Like-Lactiplantibacillus plantarum
>Z α -Like-Lacticaseibacillus rhamnosus
>Z α -Like-Sporolactobacillus terrae
>Z α -Actinia tenebrosa
>Z α -Acropora digitifera
>Z α -Symbiodinium microadriaticum
>Z α 1-Cladocopium goreau
>Z α 2-Cladocopium goreau
>Z α -Micromonas pusilla
>Z α -Micromonas commoda
>Z α -Ostreococcus tauri
>Z α -Pycnococcus provasolii
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