Structural and functional characterization of a cold-adapted stand-alone TPM domain reveals a relationship between dynamics and phosphatase activity

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The TPM domain constitutes a family of recently characterized protein domains that are present in most living organisms. Although some progress has been made in understanding the cellular role of TPM-containing proteins, the relationship between structure and function is not clear yet. We have recently solved the solution and crystal structure of one TPM domain (BA42) from the Antarctic bacterium Bizonia argentinensis. In this work, we demonstrate that BA42 has phosphoric monoester hydrolase activity. The activity of BA42 is strictly dependent on the binding of divalent metals and retains nearly 70% of the maximum at 4 °C, a typical characteristic of cold-adapted enzymes. From HSQC, 15N relaxation measurements, and molecular dynamics studies, we determine that the flexibility of the crossing loops was associated to the protein activity. Thermal unfolding experiments showed that the local increment in flexibility of Mg2+-bound BA42, when compared with Ca2+-bound BA42, is associated to a decrease in global protein stability. Finally, through mutagenesis experiments, we unambiguously demonstrate that the region comprising the metal-binding site participates in the catalytic mechanism. The results shown here contribute to the understanding of the relationship between structure and function of this new family of TPM domains providing important cues on the regulatory role of Mg2+ and Ca2+ and the molecular mechanism underlying enzyme activity at low temperatures.

Introduction

The TPM domains (Pfam PF04536) represent a superfamily of poorly characterized protein domains widely distributed in eukarya and eubacteria that have been implicated in diverse cellular processes such as photosynthesis and synaptic transmission [1,2]. The TPM domain family, previously referred to as ‘DUF477’
and ‘Repair_PSII’, has received its name from the first three proteins studied, TLP18.3, Psb32, and MOLO-1 [2,3].

In recent years, the structure of four TPM domains present in several proteins has been determined (CG2496 from Corynebacterium glutamicum [PDB code: 2KPT], PG0361 from Porphyromonas gingivalis [PDB code: 2KW7], BA42 from Bizonia argentinensis [PDB codes: 2MPB and 4OA3] [4], and AtTLP18.3 from Arabidopsis thaliana [PDB code: 3PTJ] [2]). These structures, despite sharing a very low sequence identity (<15%), present a similar αβα ‘sandwich’ conformation, not previously found in other protein domains. The function of CG2496, PG0361, and BA42 remains unknown, whereas AtTLP18.3 was classified as a thyrosynthetic protein that participates in the repair of photodamaged photosystem II from photodamage and accelerates its repair. Very recently, a functional analysis of the three proteins studied, TLP18.3, Psb32, and MOLO-1, has received its name from the first three proteins studied, TLP18.3, Psb32, and MOLO-1 [2,3].

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In this context, we have recently solved the solution and crystal structure of BA42, a protein from the Antarctic bacteria B. argentinensis [4]. BA42 is the first structure of a member of the PF04536 family comprised of a stand-alone TPM domain. The structure reveals a new topological variant of the four β-strands constituting the central β-sheet of the αβα architecture and a double metal-binding site stabilizing a pair of crossing loops, not observed in previous structures of proteins belonging to this family. The metal-bound form of BA42 is characterized by a notable rigidity, with the exception of the first seven residues. Without the metal, BA42 undergoes a transition to a more flexible structure, presenting a long C-terminal tail of 17 residues largely disordered [4]. Furthermore, BA42 lacks both an N-terminal signal sequence and a transmembrane region; therefore, in contrast to the other characterized TPM family members, it is most probably localized in the cytoplasm.

In order to characterize the functional role of BA42 and based on the previously characterized TPM domains [2,7], we tested the possible phosphatase activity of this protein. Here we demonstrate that BA42 has in vitro phosphoric-monoster hydrolyase activity (EC 3.1.3), similar to other TPM members. Further characterization indicated that the activity of BA42 is strictly dependent on the presence of divalent cations, being Mg\(^{2+}\) the most efficient to increase the enzyme activity. The protein retained nearly 70% of maximum activity at 4 °C, a typical characteristic of cold-adapted enzymes. From HSQC, 15N relaxation measurements and molecular dynamics (MD) simulations, we determined that the binding of Mg\(^{2+}\) to BA42, when compared with Ca\(^{2+}\), induces local changes in the metal-binding site and on residues of the crossing loops. Although the global structure of Mg\(^{2+}\)-bound BA42 seemed to be similar to the Ca\(^{2+}\)-bound BA42, an increased flexibility was observed in the region of the crossing loops indicating that changes in local flexibility are related to enzymatic activity, as observed for other cold-adapted enzymes [9,10]. Thermal unfolding experiments showed that the increased flexibility of Mg\(^{2+}\)-bound BA42 correlates with a reduction of protein stability. Furthermore, the results obtained from mutagenesis experiments were consistent with our previous results and allowed the identification of a region encompassing the metal-binding site that participates in the catalytic reaction. These results characterize, for the first time at atomic level, the function of a TPM-containing protein and strongly support that this domain family has phosphatase activity. Moreover, the activity of BA42 is regulated by Mg\(^{2+}\) and Ca\(^{2+}\) and the increased flexibility of the crossing loops induced by Mg\(^{2+}\) emerged as an important feature that may explain the catalytic mechanism underlying enzymatic activity at low temperatures.

**Results**

**Phosphoric hydrolase activity of BA42**

Previous studies showed that TPM domains of Rv2345 and AtTLP18.3 are capable of removing phosphate groups from different substrates [2,7]. Although the structure of the TPM domain of Rv2345 is currently unknown, a pairwise sequence alignment between this
domain and the TPM domain of AtTLP18.3 [7] showed conserved regions of key residues proposed to be involved in substrate binding. Consistent with this, a pairwise sequence alignment between BA42 and AtTLP18.3 (15% of sequence identity) showed conserved amino acid residues reported to be essential for substrate binding in the acid phosphatase of *A. thaliana* (Fig. 1A). Val101, Asp102, and Lys112 have been shown to interact with the substrate in the crystalline structure of the AtTLP18.3-Ser complex [2]. The alignment showed the conservation of Val17 and Asp18 in the primary sequence of BA42. However, we

![Fig. 1.](http://www.ebi.ac.uk/clustalW)

Yellow box indicates conservation of the Val101 and Asp102 proposed to be involved in substrate binding in AtTLP18.3. Residues shaded in blue are strictly conserved. (B) Sequence alignment of BA42 and five homologs belonging to the phylum Bacteroidetes. Red dots indicate residues involved in metal coordination in the crystal structure of BA42. The conservation of the down loop and β-strand 1 residues is shown boxed in red. Residues shaded in blue are strictly conserved and those shaded in light blue are partially conserved. (C) Schematic superposition of crystal structures of BA42 (blue) and AtTLP18.3 (brown). The solid spheres represent the Ca^{2+} ions in both structures. The residues of the ligand-binding site of AtTLP18.3 and the conserved amino acids in BA42 are highlighted in red.
were unable to find a conserved residue in BA42 corresponding to Lys112 in AtTLP18.3. In this respect, it is worth mentioning that Lys112 has been shown to be exclusively conserved in land plants [2].

These previous observations led us to the hypothesis that BA42 might also have a role as phosphoric monoester hydrolase. However, despite this sequence conservation, a structural alignment between BA42 and AtTLP18.3 clearly showed that conserved residues are located in different secondary structure elements of the TPM fold (Fig. 1C). Although the substrate-binding residues in AtTLP18.3 are positioned at the β-sheet I, the conserved hypothetical substrate-binding residues in BA42 are placed in α-helix I. In this sense, the spatial arrangement of these residues in the structure of BA42 is significantly different to the spatial configuration observed in the substrate-binding site of AtTLP18.3. In consequence, this feature hinders the possibility that both domains could share the same substrate-binding site. Moreover, as mentioned before, BA42 has a distinctive folding topology that introduces a remarkable conformational difference in the region encompassing the substrate and the metal-binding sites in the structure of AtTLP18.3 (Fig. 1C) [4]. In this regard, from a detailed inspection of the proteins classified in the PF04536 family, we noted that most members (polypeptides with a size ranging between 270 and 290 amino acids) have a similar topology characterized by the presence of an N-terminal signal sequence followed by the TPM domain, a transmembrane domain, and a C-terminal low complexity region with variable length. This group includes AtTLP18.3 and other TPM-containing proteins characterized so far. In contrast, as mentioned above, BA42 is a stand-alone TPM domain with a distinctive topology and predicted to be cytoplasmic. Relevant to these observations, from a sequence similarity search using the BLASTP program, we detected that proteins homologous to BA42 (i.e., stand-alone TPM domains) are almost exclusively conserved within the phylum Bacteroidetes. Interestingly, a multiple alignment of homologous BA42 sequences showed no strict conservation of the VDK triad corresponding to the substrate-binding site in AtTLP18.3 (Fig. 1B). However, a remarkable conservation was observed for residues involved in the metal-binding site, the down crossing loop, and the β-sheet I in BA42 (Fig. 1B).

Although BA42 may lack an active site homologous to AtTLP18.3, the hypothesis that BA42 might acts as a phosphoric hydrolase could not be ruled out. First, similar to AtTLP18.3, BA42 contains metals in its structure, being a characteristic feature of other phosphoric-monoester hydrolases [11,12]. Second, BA42 belongs to a psychrophilic organism and therefore may have evolved an alternative active site with dynamics and structural characteristics suitable for cold catalysis. Finally, through a structural comparison with DALI [13] and MATRAS [14] programs, we found that BA42 shared high structural similarity with hydrolases, as previously observed for AtTLP18.3 [2].

In this context, we analyzed the phosphoric-monoester hydrolase activity (hereafter referred to as phosphatase) of BA42 by use of the general substrates p-nitrophenyl phosphate (pNPP) and phosphoserine (pSer) at 25 °C. Initial results indicated that BA42 contained phosphatase activity (Fig. 2A). In these assays, we used the TEV protease as a control [15], a protein lacking phosphatase activity. BA42 showed enzymatic activity with both substrates, while being

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**Fig. 2.** Phosphatase activity of BA42. (A) The activity was assayed with the general substrates p-nitrophenyl phosphate (pNPP) and phosphoserine (pSer) by incubating samples of BA42 in 20 mM Tris/HCl (pH 7.5) at 25 °C in the presence or absence of EDTA. TEV protease was used as a control. (B) Effect of divalent metal ions on the phosphatase activity. BA42 was treated with EDTA, extensively dialyzed to remove the EDTA, and finally tested its phosphatase activity in 25 mM HEPES (pH 7.0) at 22 °C and in the presence of several divalent metals. All experiments were performed in triplicate and results are expressed as mean ± SEM.
more efficient to hydrolyze the pSer. Interestingly, the enzymatic activity of BA42 was dramatically inhibited after pretreatment with EDTA. Consequently, to characterize the effect of bivalent metals on the phosphatase activity, we performed the assay with the metal-free BA42 in the presence of several divalent ions (Mg$^{2+}$, Ca$^{2+}$, Mn$^{2+}$, and Zn$^{2+}$) using pSer as substrate (Fig. 2B). When incubated with Mg$^{2+}$, the activity of BA42 was significantly increased (up to fivefold) in comparison to the other divalent metal ions assayed.

We determined the optimal enzymatic conditions of Mg$^{2+}$-bound BA42 for pSer. The highest activity was observed at pH 7.0 and around 20–22 °C (Fig. 3A,B). Of note, as observed for psychrophilic enzymes [16], the protein retained nearly 70% of maximum activity at 4 °C. Kinetics assay of BA42 were performed for pSer and the results were best fitted to an allosteric sigmoidal (Fig. 3C). The apparent substrate-binding activity affinity ($K_m$) was 5.63 ± 1.82 mM, the catalytic velocity ($V_{max}$) was 3.95 ± 1.26 nmol·min$^{-1}$·mg$^{-1}$, and the $k_{cat}$ was 0.067 ± 0.002 min$^{-1}$ at the optimal condition of 20–22 °C and pH 7.0.

The binding of Mg$^{2+}$ to BA42 is associated with an increase in enzyme activity

We have previously shown that it is possible to follow conformational changes associated to Ca$^{2+}$ binding of metal-free BA42 by inspection of resonances in the range of 0.5 to −1.5 p.p.m in the 1D $^1$H NMR spectrum [4]. Accordingly, we used the same procedure to monitor conformational changes of metal-free BA42 upon binding of other bivalent metals (Fig. 4A). All different metals tested were able to bind BA42 and displayed a similar behavior of the 1D $^1$H NMR spectrum as previously observed for Ca$^{2+}$. Hence, we hypothesized that binding of divalent metals results in a global conformational change similar to that reported for Ca$^{2+}$. Surprisingly, unlike the other cations tested, Mg$^{2+}$ presented a differential behavior. The dissociation constant for this bivalent ion was 87 μM, whereas the other metals showed dissociation constants of 48 μM in average. But most important, the maximum activity in the presence of Mg$^{2+}$ was on average fivefold greater than with Ca$^{2+}$, Mn$^{2+}$, or Zn$^{2+}$ (Fig. 4B). The observed behavior when the enzyme was titrated with Mg$^{2+}$ was better appreciated by plotting the activity and the conformational change together in the same graph (Fig. 4C). It was very clear that the binding of Mg$^{2+}$, followed by monitoring the conformational change from metal-free to Mg$^{2+}$-bound BA42, was associated with an increase in enzyme activity. In this regard, it is interesting to note that even though all bivalent metals tested were able to induce the conformational change of the metal-free
BA42, Mg\textsuperscript{2+} was the most efficient in increasing the enzymatic activity. This suggests a strict specificity in the type of metal in the catalytic mechanism.

Mg\textsuperscript{2+}-bound BA42 exhibits local conformational changes with respect to Ca\textsuperscript{2+}-bound BA42

We recently proved that Ca\textsuperscript{2+} binding is crucial for stabilization of the BA42 structure, involving the closure of the C-terminal segment anchored to the $\beta\alpha$ region, generating a rigid conformation [4]. To investigate whether Mg\textsuperscript{2+} induced the same conformational change, we analyzed the \textsuperscript{1}H-\textsuperscript{15}N HSQC spectra of metal-bound BA42 with Ca\textsuperscript{2+} and Mg\textsuperscript{2+} (Fig. 5).

Interestingly, although both spectra showed a similar pattern of cross-peaks with minor variations, a group of cross-peaks in the Mg\textsuperscript{2+}-bound BA42 spectrum could not be detected (Fig. 5). This was the case of amide signals of residues 21, 26, 28, 31, adjacent to the metal-binding site, and residues 131–137, comprising the up crossing loop in the Ca\textsuperscript{2+}-bound BA42 (Fig. 5). The disappearance of these signals could be attributed to: i) an increased flexibility associated to slow or intermediate conformational motions, ii) a higher solvent exposition or iii) large chemical shift differences in the Ca\textsuperscript{2+} and Mg\textsuperscript{2+}-bound states that make it difficult to match the corresponding resonances, hence the apparent disappearance. However, no resonances in the Mg\textsuperscript{2+}-bound state could be assigned to the missing resonances, ruling out this option. Furthermore, we were unable to detect the appearance of these amide signals in HSQC spectra performed at low pH (up to the limit of precipitation of the protein), thus discarding the disappearance of the signals due to a greater exchange with the solvent. Therefore, we hypothesized that binding of Mg\textsuperscript{2+} destabilizes the conformation of the crossing loops.

To further analyze the differences between Mg\textsuperscript{2+}-bound and Ca\textsuperscript{2+}-bound BA42 HSQCs, we performed a chemical shift perturbation (CSP) study (Fig. 6A). We found major changes in the amide signals of residues that directly interact with the metals and others close to the binding site, specifically in the down crossing loop and the loop that connects the $\beta$-sheet B with the $\alpha$-helix II (Fig. 6E,F). Residues 22, 24, 25, 27, 30, 59, 60, 61, and 140 showed CSP values above the threshold line.

Taken together, these results clearly indicated that the binding of Mg\textsuperscript{2+} to BA42 (in place of Ca\textsuperscript{2+}) induced local changes in the metal-binding site and residues of the crossing loops. Although the global structure of Mg\textsuperscript{2+}-bound BA42 seemed to be similar to the Ca\textsuperscript{2+}-bound BA42, the increased flexibility of the crossing loops induced by Mg\textsuperscript{2+} emerged as an important feature that may probably explain the enhancement of the phosphatase activity.
BA42 with Ca$^{2+}$ and Mg$^{2+}$. As reported previously [4], Ca$^{2+}$-bound BA42 exhibits rather uniform values of relaxation rates along the protein sequence, a sign of a compact globular monomeric protein. Indeed, the average $T_1/T_2$ value is $8.0 \pm 0.2$ ns at 28 °C. The Mg$^{2+}$-bound BA42 $T_1/T_2$ ratios were very similar to
those Ca\(^{2+}\)-bound BA42, including anisotropic tumbling effects as a result of the shape of the molecule [4] (Fig. 6B). However, some residues in Mg\(^{2+}\)-bound BA42 displayed relative high \(T_1/\tau_2\) ratios and a shorter \(\tau_2\). They were those in the D22-E30 region, comprising the down crossing loop, and N140-K144, belonging to the C-terminal region, suggesting the presence of low-frequency motions (\(\mu\)-ms), timescales often associated with conformational exchange. In addition, the loop that connects the \(\beta\)-sheet B with the \(\alpha\)-helix II that displayed significant variations in the CSP experiment, showed minor changes in dynamics between Mg\(^{2+}\)-bound and Ca\(^{2+}\)-bound BA42.

On the other hand, the \(\text{hnNOE}\) values obtained for Mg\(^{2+}\)-bound BA42 were almost identical to the values obtained for Ca\(^{2+}\)-bound BA42, thus indicating that no new fast internal motions (ps-ns) appeared in the protein bound to Mg\(^{2+}\) (Fig. 6C). Therefore, in line with our previous results, the backbone dynamics experiments clearly indicated that Mg\(^{2+}\)-bound BA42, when compared with Ca\(^{2+}\)-bound BA42, presented increased flexibility (in the slow timescale, \(\mu\)-ms) in the region comprising the down crossing loop and the C-terminal region.

Additionally, to further studying differences in the dynamics of BA42 with Ca\(^{2+}\) and Mg\(^{2+}\), we performed all-atom MD in explicit solvent and in the presence of one of the two ions in both coordination sites. MD simulations were converged, as shown by RMSD vs time plots (Fig. 6G). When analyzing fluctuations by residue, we observed a significant increase in the C-terminal region of the Mg\(^{2+}\)-bound BA42, as can be seen in the RMSF plot (Fig. 6D), and in agreement with our previous NMR experiments.

**Mg\(^{2+}\)**-bound BA42 displays changes in thermal stability with respect to Ca\(^{2+}\)-bound and metal-free BA42

We have previously shown the strong difference in the conformational stability between metal-free and Ca\(^{2+}\)-bound BA42 [4]. In order to assess whether the conformational change induced by the binding of Mg\(^{2+}\) affected the stability of the protein, we carried out thermal denaturation experiments followed by circular dichroism (CD). In a first experiment, we found that the far-UV spectra of Ca\(^{2+}\)-bound and Mg\(^{2+}\)-bound BA42 were almost identical to each other, but slightly different to the spectrum of EDTA-treated (metal-free) BA42. The three spectra were dominated by an \(\alpha\)-helix content with two minima at 209 and 223 nm (Fig. 7A). We next investigated the thermal unfolding of metal-free, Ca\(^{2+}\)-bound and Mg\(^{2+}\)-bound BA42 by recording the molar ellipticity at 222 nm at different temperatures (Fig. 7B). All the three variants tested showed a reversible unfolding–folding process and displayed a cooperative behavior in the thermal denaturation process. However, Ca\(^{2+}\)-bound BA42 turned out to be more stable than Mg\(^{2+}\)-bound BA42 and the latter more stable than metal-free BA42, being the calculated Tm 61.3 ± 0.2 °C, 57.0 ± 0.2 °C, and 47.9 ± 0.3 °C, respectively (Fig. 7B).

**BA42 mutants unable to bind Mg\(^{2+}\)** reveal the location of the active site

As mentioned above, the crystalline structure of the AtTLP18.3-Ser complex [2] revealed a VDK triad shown to constitute the substrate-binding site. Moreover, a calcium ion-binding site stabilized by six hydrogen bonds was also suggested to be involved in AtTLP18.3 phosphatase activity. In contrast, the VDK triad is clearly not present in the structure of BA42 (Fig. 1B,C) and two calcium ions are bound on the other side of the molecule (i.e., shifted 20 Å of the corresponding metal-binding site in AtTLP18.3). At this point, based on our experiments, we could not exclude the coexistence of two metal-binding sites in BA42: one structural identified in the BA42 crystal structure and one catalytic as proposed in AtTLP18.3. If structural sites had higher affinity than the catalytic site, it would be difficult to identify the two sites from titrations. Therefore, in order to evaluate these possibilities, we conducted mutagenesis experiments. First, we generated a BA42 variant comprised of residues 1–130 (BA42-ΔCT). BA42-ΔCT lacks the C-terminal region where most of the residues constituting the metal-binding site are located (Fig. 5A,B). Interestingly, no activity in the presence of Mg\(^{2+}\) was detected for BA42-ΔCT as compared with the wild-type protein (Fig. 8D), thus ruling out the presence of a catalytic site in BA42 in a region different from that constituted by the metal-binding site. In addition, in contrast to the wild-type protein where the structural change in the presence of Mg\(^{2+}\) was clearly evident, 1D \(^{1}H\) NMR spectra of BA42-ΔCT were virtually unchanged in the presence of the divalent ion (Fig. 8C).

However, although the results obtained with BA42-ΔCT clearly demonstrate the absence of a catalytic site outside the metal-binding site in BA42, the generation of this mutant involves the removal of a large number of amino acids which can lead to a subtle conformational change that may indirectly affect the potential active site. Therefore, we considered that to better evaluate the enzyme activity of BA42, it would be more appropriate to produce a protein variant with...
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fewer mutated residues which, similar to BA42-ΔCT, had the characteristic of not binding Mg²⁺. In this respect, we generated BA42 variants carrying single-point mutations in residues involved in the metal-binding site. We found that the single-point mutation of Glumatic acid 30 to Alanine (BA42-E30A) generated a protein which lacked the ability to bind Mg²⁺ (Fig. 8C). It is interesting to note that Glutamic acid 30 has the particularity of interacting with the two metals ions in the structure of BA42 (Fig. 8B) and is highly conserved among stand-alone TPM proteins (Fig. 1B). Consistent with the behavior observed for BA42-ΔCT, the mutant BA42-E30A showed no significant differences between the 1D ¹H NMR spectra of the metal-free protein and in the presence of Mg²⁺ (Fig. 8C). But most importantly, BA42-E30A did not show phosphatase activity (Fig. 8D). This observation was congruent with the previous results with BA42-ΔCT, but in this case the loss of activity was associated to a single-point mutation of a residue specifically involved in the metal-binding site of BA42, discarding any artifact due to the removal of a large number of amino acids.

As a control, CD spectra of the BA42-E30A and BA42-ΔCT were found to be practically identical to the spectrum of the metal-free wild-type protein (Fig. 8E), thus indicating the integrity of the secondary structure and the absence of large conformational changes of mutant proteins with respect to the metal-free wild-type BA42.

**Discussion**

BA42 is the first member of the PF04536 family comprised of a stand-alone TPM domain structurally and biochemically characterized to date. The structure of BA42 uncovers a new topological variant of the 4βα architecture and a double metal-binding site stabilizing a pair of crossing loops, not observed in previous structures of proteins belonging to this family. Furthermore, in contrast to other characterized TPM family members, BA42 has predicted cytoplasmic localization.

Previously, another structural characterized TPM-containing protein, AtTPL18.3, was reported to have phosphatase activity [2]. Here we show that beside the fact that BA42 has the conserved residues involved in the catalytic mechanism, they do not have a spatial arrangement compatible with the active site reported for AtTPL18.3 [2]. Despite this discrepancy in the structure of the potential active site, in this work we demonstrate that BA42 has phosphatase activity and displays characteristics compatible with psychrophilic enzymes [16] presenting a maximum activity at 20–22 °C. In this sense, it is worth to mention that the most favorable growth condition of *B. argentinensis* is around 22–25 °C, which is surprisingly similar to the optimal catalytic temperature reported here for BA42. Although the absolute values of activity are low compared with other hydrolases enzymes [17], they are consistent with those reported previously for other members of the TPM family [2,7]. Furthermore, as the physiological substrate of BA42 is not known, the low activity observed *in vitro* may also be attributed to suboptimal reaction conditions.

In the present study, we show the strict dependence of the activity of BA42 on the presence of divalent ions. Although the protein presents no activity when pretreated with EDTA, activity is observed after treatment with different metallic cationic ions. Interestingly, an increase of about fivefold is observed in the presence of Mg²⁺ in comparison with other metals tested. This increase in the enzyme activity is associated to a conformational change of BA42 from a structure with an unfolded C-terminal region to a closed structure with the C-terminal tail bound to the protein core. However, all metals tested induce a similar conformational change in BA42 and even with higher affinity than Mg²⁺. The major differences in the conformation of Mg²⁺-bound BA42 reside in the dynamics of the crossing loops and the metal-binding site. Two independent experiments denote an increase in the...
flexibility of these regions. First, the HSQC spectra of the Mg$^{2+}$-bound protein reveal the disappearance of amide signals of residues adjacent to the cation-binding site and residues 131–137, comprising the up crossing loop in the Ca$^{2+}$-bound BA42, which presented a defined conformation. As neither a large chemical shift variation nor an increase in solvent exposure was detected in this region, we conclude that this is due to a gain in flexibility (in the scale of slow or intermediate movements, μs-ms). Second, $^{15}$N relaxation experiments clearly show an increase in slow movements (μs-ms) of residues D22-E30, comprising the down crossing loop, and N140-K144, belonging to the C-terminal portion. Furthermore, the lower conformational stability of Mg$^{2+}$-bound BA42 with respect to Ca$^{2+}$-bound BA42, detected in thermal denaturation experiments, could be attributed to an increase in the flexibility of this region. Consistent with this, MD simulations provide further evidence of increased flexibility in residues belonging to the C-terminal portion.

Taken together, these results indicate that binding of Mg$^{2+}$ to BA42 induces local flexibility in the region comprising the crossing loops and residues involved in metal binding. This characteristic is associated with a decrease in the global conformational stability and an improvement of the catalytic activity, compared with Ca$^{2+}$-bound BA42.

In order to further evaluate the relationship between the flexibility and the catalytic activity, we compared the enzyme activity between Ca$^{2+}$-bound and Mg$^{2+}$-bound BA42 as a function of temperature (Fig. 7C). Although the activity of Ca$^{2+}$-bound BA42 was lower than Mg$^{2+}$-bound BA42 over the whole temperature range studied, major differences were detected below 26 °C. Interestingly, the maximum activity of Ca$^{2+}$-bound BA42 was found at 37 °C, shifted about 17 °C relative to that observed for the Mg$^{2+}$-bound BA42 (around 20–22 °C). In consequence, these results were compatible with the idea that the presence of a flexible active site in BA42 generates a more efficient enzyme to catalyze at low temperatures.

In response to low temperatures, psychrophiles use diverse strategies for adaptation [18,19]. In most organism, the flexibility of the structural elements that are involved in the catalytic cycle of enzymes is commonly adjusted, resulting in a loose structure characterized by a low local conformational stability and/or by destabilization of the whole molecule [9,20]. This involves a reduction in the number and strength of all types of weak interactions or the disappearance of stability factors, resulting in improved dynamics of active site residues in the cold. In this scenario, based on our results, we can hypothesize that the region comprising the crossing loops and residues involved in metal binding in BA42 is directly related to the catalytic mechanism and may contain the active site. Furthermore, it

Fig. 7. (A) Circular dichroism spectra of BA42. The CD spectra of metal-free (black line), Mg$^{2+}$-bound (red line), and Ca$^{2+}$-bound BA4 (blue line) were recorded at 25 °C. (B) Thermal unfolding process of BA42. Metal-free (black line), Mg$^{2+}$-bound (red line), and Ca$^{2+}$-bound BA4 (blue line) were incubated at different temperatures from 25 to 75 °C and the molar ellipticity at 222 nm was recorded at intervals of 1 °C. (C) Effect of temperature on enzyme activity of BA42. The phosphatase activity was measured at different temperatures in 25 mM HEPES (pH 7.0) and in the presence of 1 mM CaCl$_2$ (blue dots) or 1 mM MgCl$_2$ (red dots). The activity assays were performed in triplicate and the results are expressed as mean ± SEM.
has been observed in several enzymes that ‘crossing loops’ structures are part of the active site [21,22]. In this respect, the results obtained with BA42-ΔCT and BA42-E30A were crucial to circumscribe the active site of BA42 in a region constituted by the metals binding site and to rule out the existence of an active site.
similar to that reported for AtTLP18.3 [2]. At this point, given the differences observed between BA42 and AtTLP18.3 regarding the folding topology, the metal-binding site and the location of the active site, we propose that to study TPM-containing proteins more appropriately, they might be classified in at least two groups: membrane-associated TPMs and cytoplasmic stand-alone TPMs. Cytoplasmic stand-alone TPMs seem to be almost exclusively conserved within the phylum Bacteroidetes, while membrane-associated TPMs appear to be conserved over a wide range of organisms (including plants, fungi, and bacteria). In particular, as both groups of TPM-containing proteins are found in Bacteroidetes, we hypothesize that these domains have evolved independently. In addition, cytoplasmic stand-alone TPMs are mainly found in marine organisms adapted to low temperatures and, therefore, we assume that the members of this group conserve structural features compatible with the activity at these environmental conditions, similar to those described for BA42.

On this basis, we hypothesize that BA42 might have evolved a catalytic site where activity at low temperature is regulated, at least in part, by varying the flexibility of the active site. In the presence of Ca\(^{2+}\), the active site becomes rigid and the activity is depressed; whereas the binding of Mg\(^{2+}\) enhances the flexibility in a way compatible with catalysis at low temperatures (Fig. 9). In this sense, Mg\(^{2+}\) is the most abundant divalent metal in living cells. It has been reported that the intracellular levels of Mg\(^{2+}\) in marine bacteria may vary from 130 to 710 mM in normal growth conditions [23] and even up to 910 mM under carbon limitation or dormancy [24]. In addition, the intracellular Ca\(^{2+}\) concentration in these organisms ranges from 10 to 80 mM [23]. Therefore, given the high concentration of these two metals, it can be predicted that BA42 may be found in the cytoplasm as metal-bound protein and the relative levels of Ca\(^{2+}\) and Mg\(^{2+}\) could be an important factor to modulate the phosphatase activity.

In conclusion, the results shown here contribute to the understanding of the relationship between structure and function of this recently characterized family of TPM domains. Our work, in conjunction with other previously described to date, suggest that the phosphoric hydrolase activity of the TPM scaffold has been conserved throughout evolution. However, more research is needed to uncover the catalytic and regulatory mechanisms relevant to its cellular function.

Materials and methods

Construction of BA42 mutants

BA42-ΔCT gene was amplified by PCR using *B. argentinensis* genomic DNA as template, 5′-ATGTCTAAAAATAGAAA-GAGTTTTTAAC-3′ (primer A) as 5′-primer and 5′-CATGGAAAATATTTTGCCAAAGC-3′ as 3′-primer. BA42-E30A site-directed mutant was generated by the megaprimer method using *B. argentinensis* genomic DNA as template. The megaprimer was synthesized in a first PCR reaction using primer A as 5′-primer and 5′-GAACACG-TATGGCGCCAGACGTG-3′ as mutagenic primer. The

![Fig. 9. A schematic representation of the metal-dependent behavior of the catalytic region of BA42. The figure illustrates the postulated mechanism of phosphatase activity modulation by varying the flexibility of the active site. In the absence of divalent metal ions, the C-terminal region of BA42 is extremely disordered and the active site is not formed. In the presence of Ca\(^{2+}\), the active site acquires a rigid configuration and the activity is low. The binding of Mg\(^{2+}\) induces an increment of the flexibility of the active site, in comparison with the protein bound to Ca\(^{2+}\), and the protein activity is highly increased.](image)
second PCR reaction was performed using the megaprimer as 5' primer and 5'-TCCCTTGTAAAAATTGTTTGG-3' as 3' primer. The final PCR products were purified and cloned between NdeI and BamHI restriction sites of the pET-28a(+) vector (Qiagen). Recombinant proteins were transformed into a TEV protease cleavage site followed by 6×His Histidine tag for affinity purification. The quality of the preparations was analyzed by DNA sequencing. Finally, the recombinant expression plasmids were transformed in *Escherichia coli* BL21 (DE3) competent cells (Stratagene, La Jolla, CA).

**Protein expression and purification**

BA42 recombinant proteins were expressed and purified as previously described [4].

**Phosphorico-monoester hydrolase activity**

The phosphorico-monoester hydrolase activity of BA42 was assayed by using p-nitrophenolphosphate (pNPP) (New England Biolabs) and O-phosphoserine (pSer) (Sigma-Aldrich) as substrates. The reaction solution contained 15 μg of purified BA42, 25 mM HEPES (pH 7), and the substrates in a total volume of 0.1 mL. The reactions were initiated by the addition of 20 μM pNPP or pSer, and quenched by the addition of 10 μL of 1 M NaOH. When pNPP was used as a substrate, the activity was measured by monitoring the change in the absorbance at 405 nm (ε405 nm = 18 300 M⁻¹ cm⁻¹) as pNPP (colorless) is converted to p-nitrophenol (pNP, yellow). For pSer, the method of Eibl and Lands [25] was used to detect the Pi released. All the assays were performed in triplicate. Enzymatic kinetics was analyzed by fitting the data to the allogenic sigmoidal equation using the nonlinear regression equation

\[
V = \frac{V_{\text{max}} [S]^h}{K_m + [S]^h},
\]

where \(V\) is the initial velocity, [S] is the concentration of the substrate, and \(h\) is the Hill slope.

In order to analyze the effect of divalent metal ions on the phosphorico-monoester hydrolase activity of BA42, the protein was treated with 1 mM EDTA for 10 min and then extensively dialyzed against 25 mM HEPES (pH 7.0). After this treatment and before starting the enzymatic reaction, the enzyme was incubated with 1 mM CaCl₂, 1 mM MgCl₂, 1 mM MnCl₂ or 1 mM ZnCl₂ in the reaction buffer at 22 °C for 10 min, as indicated. The optimal reaction conditions for BA42 were determined using pSer as substrate. The optimal pH was determined by using 25 mM acetate-acetate buffer (pH 5–7), 25 mM Tris/HCl buffer (pH 7–9), and 25 mM glycine-NaOH buffer (pH 9–12).

**CD measurements**

Circular dichroism measurements were made on a Jasco J-815 spectropolarimeter (Jasco Inc., Easton, MD, USA). Circular dichroism spectra of BA42 were obtained in the range of 185–250 nm using a 0.2 mm path length and expressed in molar ellipticity (θ) (deg·cm²·dmol⁻¹). All measurements were carried out at a protein concentration of 2 μM in 20 mM HEPES pH 7.0.

Unfolding transitions as a function of temperature were monitored at 222 nm. Temperature was varied from 25 to 75 °C at a constant rate of 1 °C·min⁻¹ and sampling at intervals of 1 °C.

**NMR spectroscopy**

**NMR samples, data acquisition, and processing**

Samples for NMR experiments contained 0.3–0.8 mM 15N-labeled BA42 or 0.05–0.1 mM unlabeled BA42 dissolved in a buffer containing 25 mM NaCl, 20 mM HEPES, and 5% D₂O, pH 7. NMR experiments were performed at 298 K in a Bruker 600 MHz Avance III spectrometer (Bruker Instruments, Inc., Bellerica, MA, USA) equipped with a 5-mm triple resonance cryoprobe incorporating shielded z-axis gradient coils. The NMR data were processed using NMRPipe [26] and analyzed using NMRVIEWJ [27].

**¹H NMR measurements**

The 1-1 Jump-and-return water suppression pulse sequence was used for ¹H 1D NMR experiments. In order to follow the conformational change of BA42 in the presence of divalent metals, we measured the peak integral of methyl groups of residues 1100 and V88, as reported previously [4].

**¹⁵N spin relaxation measurements**

Measurements of ¹⁵N T1, T2, and steady-state ¹H-¹⁵N NOE (hnNOE) of BA42 were performed at a nitrogen frequency of 60.82 MHz and 298 K, using standard pulse schemes in an interleaved manner. Both T1 and T2 data were acquired with six relaxation delays. The delays used were 7, 300, 600, 900, 1200, and 1600 ms for the T1 calculations and 15.7, 31.4, 62.8, 125.5, 156.9, and 219.7 ms for the T2 calculations. The rate analysis routine of NMRVIEWJ was used to analyze the data [27]. The hnNOE values were determined by the ratio of peak volumes of spectra recorded with and without ¹H saturation, employing a net relaxation delay of 4 s for each scan in both experiments. Typically, errors were ~1% for T1 and T2, and 2–5% for hnNOE measurements.

**Molecular dynamics**

The initial structure files were obtained from the structure of BA42 from the Protein Data Bank (PDB code: 4OA3) [4]. The AMBER99SB force field was used [28] and TLEAP program was used to create the topology and...
The SHAKE algorithm was used for constraining the bonds kept constant at 1 atm by the Berendsen barostat [31]. 300 K with a 0.1 ps coupling constant and pressure was constant by the Berendsen thermostat algorithm set at 200 ns long production runs, the temperature was kept for keeping pressure constant at 1 atm [31]. During temperature with a coupling constant of 0.1 ps for keeping processes of 50 ps with a 1 fs time step using Berendsen thermalization. This was followed by an equilibration process of 50 ps with a 1 fs time step using Berendsen temperature with a coupling constant of 0.1 ps for keeping temperature constant at 300 K and Berendsen barostat for keeping pressure constant at 1 atm [31]. During 200 ns long production runs, the temperature was kept constant by the Berendsen thermostat algorithm set at 300 K with a 0.1 ps coupling constant and pressure was kept constant at 1 atm by the Berendsen barostat [31]. The SHAKE algorithm was used for constraining the bonds that contained an H atom [32] and electrostatic interactions were calculated by the Particle Mesh Ewald decomposition algorithm [33]. A 2 fs time step was used. vmd program was used for visualization and RMSD and RMSF calculations [34].

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Author contributions
LAP and CS equally contributed to this work. LAP, CS, REI, and MA planned experiments, performed experiments, analyzed data, and wrote the paper. AGT and DOC analyzed data and wrote the paper.

Conflicts of interest
The authors declare that there is no conflict of interest regarding the publication of this paper.

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