

RESEARCH ARTICLE

Functional Significance of Calcium Binding to Tissue-Nonspecific Alkaline Phosphatase

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Abstract

The conserved active site of alkaline phosphatases (AP) contains catalytically important Zn²⁺ (M1 and M2) and Mg²⁺-sites (M3) and a fourth peripheral Ca²⁺ site (M4) of unknown significance. We have studied Ca²⁺ binding to M1-4 of tissue-nonspecific AP (TNAP), an enzyme crucial for skeletal mineralization, using recombinant TNAP and a series of M4 mutants. Ca²⁺ could substitute for Mg²⁺ at M3, with maximal activity for Ca²⁺/Zn²⁺-TNAP around 40% that of Mg²⁺/Zn²⁺-TNAP at pH 9.8 and 7.4. At pH 7.4, allosteric TNAP-activation at M3 by Ca²⁺ occurred faster than by Mg²⁺. Several TNAP M4 mutations eradicated TNAP activity, while others mildly influenced the affinity of Ca²⁺ and Mg²⁺ for M3 similarly, excluding a catalytic role for Ca²⁺ in the TNAP M4 site. At pH 9.8, Ca²⁺ competed with soluble Zn²⁺ for binding to M1 and M2 up to 1 mM and at higher concentrations, it even displaced M1- and M2-bound Zn²⁺, forming Ca²⁺/Ca²⁺-TNAP with a catalytic activity only 4–6% that of Mg²⁺/Zn²⁺-TNAP. At pH 7.4, competition with Zn²⁺ and its displacement from M1 and M2 required >10-fold higher Ca²⁺ concentrations, to generate weakly active Ca²⁺/Ca²⁺-TNAP. Thus, in a Ca²⁺-rich environment, such as during skeletal mineralization at pH 7.4, Ca²⁺ adequately activates Zn²⁺-TNAP at M3, but very high Ca²⁺ concentrations compete with available Zn²⁺ for binding to M1 and M2 and ultimately displace Zn²⁺ from the active site, virtually inactivating TNAP. Those ALPL mutations that substitute critical TNAP amino acids involved in coordinating Ca²⁺ to M4 cause hypophosphatasia because of their 3D-structural impact, but M4-bound Ca²⁺ is catalytically inactive. In conclusion, during skeletal mineralization, the building Ca²⁺ gradient first activates TNAP, but gradually inactivates it at high Ca²⁺ concentrations, toward completion of mineralization.

Introduction

Alkaline phosphatases (APs) occur widely in nature, and are found in many organisms from bacteria to man [1, 2]. *In vitro*, APs are quite promiscuous in their substrate specificity, being able to catalyze both the hydrolysis of monoesters of phosphoric acid and a transphosphorylation reaction in the presence of large concentrations of phosphate acceptors [1]; however their

in vivo functions are quite specific [2]. Four isozymes, with differential tissue expression and encoded by distinct genes, are found in humans: tissue-nonspecific AP (TNAP, also known as liver-bone-kidney type), placental AP (PLAP), germ cell AP and intestinal AP (IAP). Mammalian APs in general, the human isozymes in particular, are homodimeric enzymes and each catalytic site contains three metal ions, two Zn²⁺ (M1 and M2) and one Mg²⁺ (M3), which are perfectly conserved throughout speciation and required for enzymatic activity [3]. An additional metal-binding site M4, that appears to be occupied by Ca²⁺ and is not present in the bacterial enzymes, was revealed upon solving the PLAP 3D structure [4, 5]. This fourth metal site is conserved in all human and mouse APs [6] and presumably represents a novel feature common to many if not all mammalian APs. However, the structural and functional significance of this new M4 metal site remains to be established. Here we have investigated the functional role of this M4 site for TNAP catalysis, an enzyme crucial for skeletal and dental mineralization.

Hypomorphic mutations in *ALPL*, the gene encoding human TNAP (*Alpl* in mice) lead to hypophosphatasia, a heritable form of rickets or osteomalacia. Hypophosphatasia is caused by accumulation of inorganic pyrophosphate (PP_i), the physiological substrate of TNAP and a potent mineralization inhibitor, in the cartilage and bone extracellular matrix [7, 8]. Thus, a crucial function of TNAP is to hydrolyze PP_i in skeletal and dental tissues, restricting the extracellular pool of this mineralization inhibitor [9] and allowing calcification to proceed. In addition, TNAP can also produce phosphate (P_i) from ATP, which helps drive mineralization in the presence of Ca²⁺ [10]. The current model of the initiation of skeletal and dental mineralization involves crystal formation inside the chondrocyte- and osteoblast-derived matrix vesicles (MVs) favored by P_i accumulation resulting from both PHOSPHO1-mediated intravesicular production and transporter-mediated influx of P_i produced primarily by the ATPase activity of TNAP. Next, extravesicular calcification is mainly supported by the pyrophosphatase activity of TNAP, and is driven by the availability of P_i and Ca²⁺ ions and the presence of a collagenous fibrillar scaffold and guided by other ECM mineral-binding proteins [11]. Early studies indicated that TNAP in cartilage is a Ca²⁺ binding glycoprotein [12], but whether Ca²⁺ binding occurs at M4 or any other site and whether Ca²⁺-binding functionally modulates TNAP activity remains unknown. The overall structure of the M4 site comprises 76 residues (209–285) folded into two β-strands flanked by two α-helices. In PLAP, this region includes a glycosylation site at N249, stabilized by a stacking interaction with W248, and a metal ion coordinated by carboxylates from residues E216, E270 and D285, the carbonyl of F269 and a water molecule, all of which suggest that M4 is occupied by a Ca²⁺ ion [4]. Interestingly, hypomorphic mutations of the corresponding residues in TNAP (W253, E218, E274, and D289) cause hypophosphatasia (http://www.sesep.uvsq.fr/03_hypo_mutations.php).

In this report, we have investigated the functional significance of Ca²⁺ binding to all four metal ion-binding sites in TNAP to better understand how the activity of TNAP is regulated during skeletal mineralization in an environment with high local Ca²⁺ gradients, further aiming to understand the pathophysiological basis for hypophosphatasia.

Materials and Methods

Mutagenesis and expression of TNAP-FLAG and PLAP-FLAG enzymes

Site-directed mutagenesis was performed to generate a series of PLAP and TNAP Ca²⁺-binding site (M4) and peripheral site mutants, using pcDNA3/PLAP-FLAG or pcDNA3.1/TNAP-FLAG vectors as templates, respectively. Site-directed mutagenesis was performed with a Quickchange Site-Directed Mutagenesis kit (Stratagene, San Diego, CA, USA), according to the manufacturer's instructions, using the oligonucleotide primers listed in Table A in [S1 File](#). COS-1 (ATCC

CRL-1650) cells were transfected with plasmids and FLAG-tagged enzymes were collected from the culture supernatant, as described previously [13]. A

Western Blotting

Culture supernatants of each FLAG-tagged mutant enzyme and the respective native enzymes were purified using an anti-FLAG M2 monoclonal antibody (AbM2) column (Sigma, St Louis, MO, USA) according to the manufacturer's instructions (the antibody M2 will be referred to as AbM2, to avoid confusion with the metal ion-binding site M2). The protein concentration of each purified sample was determined with a Pierce BCA Protein Assay (Thermo Scientific, Rockford, IL, USA). For Western blots, electrophoresed proteins were transferred to reinforced nitrocellulose membrane (Whatman, Dassel, Germany) followed by blocking in SuperBlock Blocking Buffer in Tris-buffered saline (Thermo Scientific, Rockford, IL, USA). Subsequently, the membranes were incubated with 1 µg/ml AbM2, followed by detection as described [14].

Kinetic measurements

PLAP and TNAP activity were measured as a function of the concentration of the reference substrate p-nitrophenyl phosphate (pNPP; Sigma, St Louis, MO), at the enzyme's pH optimum in 1 M diethanolamine buffer, pH 9.8, containing 20 µM ZnCl₂ (Merck, Darmstadt, Germany) and 1 mM MgCl₂, (Merck) and Lineweaver-Burk plots were constructed to calculate K_m and V_{max}. From the V_{max} values, k_{cat} was calculated by comparison with V_{max} for a known concentration of native TNAP and historical k_{cat} values [13]. Molar concentrations of p-nitrophenol were calculated, using a molar extinction coefficient $\epsilon = 18,000 \text{ M}^{-1}\text{cm}^{-1}$, at pH 9.8 (no conversion was made at pH 7.4).

Functional analysis on the M1-M4 metal sites

Buffers and pNPP substrate were Chelex-treated prior to addition of ZnCl₂ and/or CaCl₂ and or MgCl₂, to minimize contamination with unknown divalent metal ions. Microtiter plates were coated with AbM2 (0.2–0.4 µg/ml) overnight at 4°C, after which plates were blocked with 1% human serum albumin (hSA) for 1 h in Tris-buffered saline (TBS: 50 mM Tris, 137 mM NaCl, 2.6 mM KCl) pH 8.0. TNAP and its mutants were then incubated in TBS, 0.1% hSA, for 3 h at room temperature at various dilutions, taking into account the specific activity for each mutant and the pH at which subsequent analyses would be carried out; dilution factors ranged from 10–200 for the native TNAP solution (stock concentration 20 nM). After washing, plates were subjected to one of the following treatments: 1. Incubation with 1 mM EDTA in TBS, 0.1% hSA, for 2 h; 2. Incubation with CaCl₂ (0–20 mM) in TBS, 0.1% hSA, up to 16 h; 3. Incubation with 20 µM ZnCl₂ + 1 mM MgCl₂ for up to 16 h. After washing, EDTA pre-treated plates were incubated with increasing [ZnCl₂] (0–40 µM, but mostly 20 µM) to upload bound enzyme, for 2 h. Microtiter plates were then incubated for 60–90 min with the substrate pNPP (1 or 10 mM), dissolved in 1 M Tris-HCL buffer, pH 7.4 [13], containing ZnCl₂, CaCl₂ (Merck) and/or MgCl₂, as specified. Alternatively, pNPP was dissolved in 1 M DEA-buffer, pH 9.8 [13], containing ZnCl₂, CaCl₂ and/or MgCl₂, as specified. The formation of p-nitrophenol was then followed kinetically, via repetitive measurements of A_{405nm}, at 1 or 2 min intervals, up to 90 or 120 min, after which plots of A_{405nm} vs. time were constructed. For the indicated reference interval (mostly 60–90 min, where 405nm increased linearly with time), the mean rate of hydrolysis was calculated as $\Delta m_{405nm}/min$. Acceleration and deceleration of TNAP activity was measured from calculation of $\Delta m_{405nm}/min$ at a given time point, and these slopes were plotted as a function of time, or vs. metal ion concentration. Slopes were derived using the GraphPad Prism (San Diego, CA) and represent a measure for the activity of TNAP for the

chosen interval. This enzyme kinetics representation was chosen to allow the direct comparison of enzyme activities in conditions where specific activities fluctuated over time. When specified, TNAP bound to microtiter plates was incubated overnight at room temperature for 16h with 250 μM EDTA in TBS, 0.1% hSA, to prepare holo-TNAP.

Two different commercial sources of CaCl₂ were used in these studies: Calcium chloride dihydrate pro analysi (Merck KGaA, Darmstadt, Germany) with Sr (≤ 0.05%) and Mg (≤ 0.005%) as the most relevant major divalent ion contaminants; and Calcium chloride solution BioUltra, for molecular biology (≈ 1M, Sigma-Aldrich, Saint-Louis, MO), also with Sr (≤ 20 mg/kg) and Mg (≤ 5 mg/kg) as the most relevant major divalent ion contaminants, also containing other divalent ions (≤ 5 mg/kg).

TNAP protein structure modeling

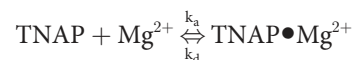
The primary sequence of human TNAP was submitted to the SWISS-MODEL server [15] to model their tertiary structures, based on homology to human placental alkaline phosphatase (1ZED). The resulting molecular structures for TNAP and its M4 mutants were visualized and analyzed using Chimera v1.7 [16] and Swiss-PdbViewer [17].

TNAP (mutant) structural analysis

The structural impact of TNAP mutations surrounding M4 was analyzed by antibody mapping and heat inactivation. In the first approach, a TNAP-epitope mapped antibody panel was coated onto microtiter plates, after which a standard concentration of TNAP was added, as previously described [13]. Bound TNAP or TNAP mutant was then detected, using AbM2 [13]. Heat inactivation studies of PLAP and its mutants were performed by incubation at different temperatures, after which remaining PLAP (mutant) activity was measured with pNPP as a substrate [18]. Heat inactivation of the more heat-labile TNAP (mutants) was analyzed by measuring residual activity, after incubation of TNAP (mutant) at 56°C in TBS, as a function of time [13].

Mathematical model

Binding of Mg²⁺ to the M3 site in TNAP was represented by the following general model:



and $K_d = K_d/K_a = [\text{TNAP}] \cdot [\text{Mg}^{2+}]/[\text{TNAP} \bullet \text{Mg}^{2+}]$,

In which k_a and k_d represent the association and dissociation rate constant of this reaction, respectively and K_d is the dissociation constant.

The apparent first order rate constant k_{app} for the binding was calculated from the time to reach 50% of the maximal TNAP activity, $t_{1/2}$ ($k_{app} = \ln 2/t_{1/2}$) and was then fitted to the equation

$$k_{app} = k_a \cdot [\text{Mg}^{2+}] + k_d.$$

Plots of k_{app} vs. $[\text{Mg}^{2+}]$ were therefore constructed, to derive k_d and k_a , enabling calculation of K_d and comparison to directly determine the K_d from dose-response studies. These dose-response curves for TNAP activation vs. metal ion concentration were calculated after a steady-state was reached, i.e. from the “reference” interval from 60–90 min, by fitting the data to a one-site binding model (GraphPad Prism), from which plots the K_d , maximal activity (plateau) and Hill coefficient were derived.

Statistical analysis

All experiments were carried out at least three times and were confirmed at different enzyme dilutions. When executed in identical conditions, data were averaged and represented as the mean values \pm SD; when repeated with different concentrations, one representative example is shown. Dissociation constants are expressed with their SD, calculated from the fitted lines (GraphPad Prism). Groups and dissociation constants were compared using unpaired Student *t* tests, calculating two-tailed *p*-values, defined in the text or figure legends, as required.

Results

Allosterism for Ca²⁺ binding to Zn²⁺-TNAP

Various lengths of TNAP demetalation and remetation were tested, prior to selection of a standardized approach. [S1 Fig.](#) illustrates that the demetalation strategy selected in the majority of cases was a compromise resulting in a low residual baseline TNAP activity, but guaranteeing full enzyme recovery upon remetation with predefined metal ions. Therefore, in these cases, TNAP activity profiles shown were constructed after EDTA-treatment for 2 h, followed by a loading step with 20 μ M ZnCl₂ (yielding Zn²⁺-TNAP with Zn²⁺ in M1 and M2 but free M3). Prior to investigating TNAP activation as a result of binding of Ca²⁺ to M3, we verified whether human TNAP activity complies with the model described for bovine kidney AP [19, 20]. [Fig. 1A](#) shows that at high TNAP-concentration (AbM2-bound at 1 nM), increasing concentrations of MgCl₂ strongly enhanced Zn²⁺-TNAP activity up to 12-fold between 0–1 mM, from baseline (35 mA405nm/min) to a maximum of 417 mA405nm/min, which is similar to the activity observed for native TNAP (non-EDTA treated), measured with 1 mM MgCl₂ in the pNPP substrate, at pH 9.8 (not shown). Repetition at 15-fold lower TNAP concentration ([Fig. 1A](#), right panel, max. activity 28 mA405nm/min) confirmed strong allosteric TNAP activation by Mg²⁺, but also illustrated the low rate of Mg²⁺-binding to Zn²⁺-TNAP, from the progressive acceleration (i.e. increasing slope) of TNAP activity with time. At high [Mg²⁺] (1 mM), binding was almost immediate (constant slope for Δ A405nm/time). The first derivative of these curves (i.e. the plot of the slopes vs time) describing the formation of Mg²⁺/Zn²⁺-TNAP (plotted and fitted in [Fig. 1B](#), left panel) confirmed this formation to result from a bimolecular reaction, requiring over 90 min to complete for the lowest [Mg²⁺] tested. Calculation of *t*_{1/2} and corresponding apparent first order rate constants for this reaction, followed by plotting *k*_{app} vs. [Mg²⁺] ([Fig. 1B](#), right panel, *r*² = 0.977) allowed estimating the kinetic constants of Mg²⁺ binding (*k*_d = 4.4 \pm 0.2 \times 10⁻⁴ s⁻¹ and *k*_a = 23 M⁻¹s⁻¹), resulting in *K*_d = *k*_d/*k*_a = 17 μ M (95% confidence interval: 9.3–39.8), consistent with previously determined values for bovine kidney AP at pH 8 (*k*_a = 7 M⁻¹s⁻¹ and *k*_d = 4 \times 10⁻⁴ s⁻¹). In other words, the allosteric effect of [Mg²⁺] on human TNAP is consistent with that reported for bovine kidney AP [19], with a very slow *k*_a but high affinity (activity measured with 10 mM pNPP, see below). These experiments confirmed that binding of Mg²⁺ stimulated TNAP activity slowly but potently.

Similar incubations as in [Fig. 1A](#) (left panel) with increasing [CaCl₂] also dose-dependently stimulated Zn²⁺-TNAP ([Fig. 2A](#)) to a maximal activity of 158 mA405nm/min at 625 μ M CaCl₂, i.e. only 2.6-fold weaker than the maximum for Mg²⁺/Zn²⁺-TNAP in [Fig. 1A](#). However, the binding kinetics (exponential in some cases, although not further analyzed) occurred detectably faster than those for Mg²⁺ binding, at the higher CaCl₂ concentrations needed to achieve full activation. [Fig. 2A](#), left panel shows functional binding of Ca²⁺ to M3, but high [Ca²⁺] (1.25–10 mM) inhibited Ca²⁺/Zn²⁺-TNAP activity to a level equating 13 mA405nm/min (at 10 mM), i.e. below that of the baseline (22 mA405nm/min). This activity, representing 8% of that of Ca²⁺/Zn²⁺-TNAP (and 3% of that of Mg²⁺/Zn²⁺-TNAP, analyzed at the same [TNAP]),

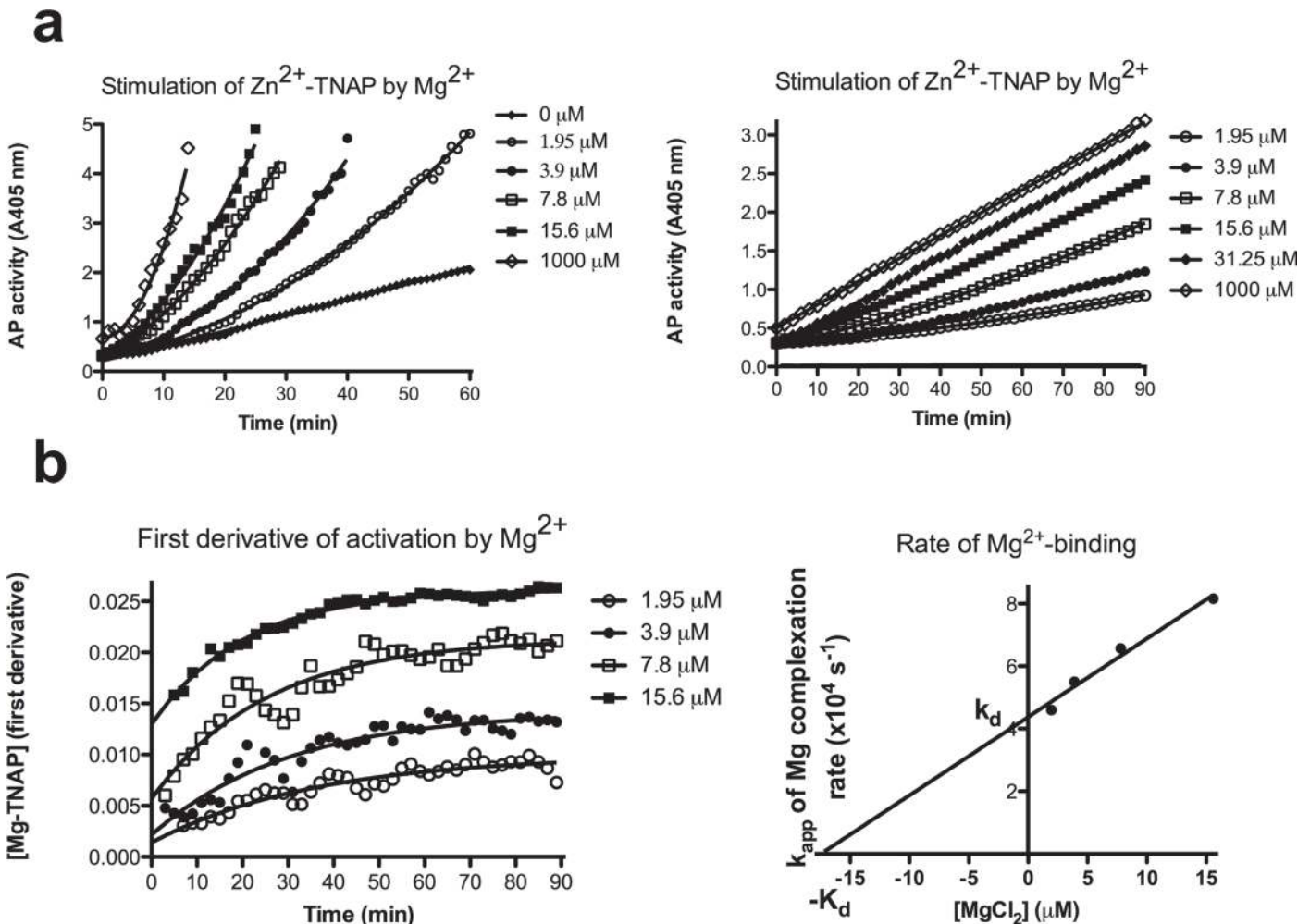


Fig 1. Allosteric activation of Zn²⁺-TNAP by MgCl₂. a. Progressive AbM2-bound Zn²⁺-TNAP activation, visualized as increasing slopes in plots of A405 nm vs. time, for the indicated [Mg²⁺], added to Chelex-pretreated pNPP (10 mM) at pH 9.8 (left panel); repeat of the experiment in (a) at 15-fold lower [TNAP] over a time-interval 0–90 min, for the indicated [Mg²⁺] (right panel); b. Slopes (first derivatives) to the lines in Fig. 1A, right panel vs. time, for the indicated [Mg²⁺], describing formation of Mg²⁺/Zn²⁺-TNAP as a function of time (left panel); plots of k_{app} (calculated from Fig. 1B left panel) vs. [Mg²⁺] and determination of k_a and k_d for binding of Mg²⁺ to Zn²⁺-TNAP (right panel). Experiments representative of at least three replicates with variable enzyme and MgCl₂ concentrations.

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thus abrogated the role of Zn²⁺ in the baseline activity of Zn²⁺-TNAP (Fig. 2A, right panel). Further analysis (S2 Fig.) confirmed that this drop was the result of Ca²⁺ binding to M1 and M2, resulting in the formation of poorly active Ca²⁺/Ca²⁺-TNAP. Analysis at 15-fold less TNAP (as in Fig. 1A, right panel) and plotting the apparent TNAP activity (calculated for the interval range 60–90 min, see S2 and S1 Figs.) vs. [MgCl₂] or [CaCl₂] (Fig. 2B) revealed a one-site saturation profile for the binding of Mg²⁺ to M3, with an apparent K_d = 4.4 ± 0.23 μM and an activity plateau at 31 mA405nm/min, at pH 9.8, using a standard [pNPP] = 10 mM. In the same conditions, the binding of Ca²⁺ followed a biphasic pattern (from baseline activity of Zn²⁺-TNAP), with the ascending limb describing a one-site saturation, with an estimated K_d = 220 ± 26 μM and a pseudo-plateau at 8.3 mA405nm/min, i.e. 3.7-fold lower than the Mg²⁺ plateau. The descending limb revealed potent TNAP inactivation with an IC₅₀ = 5.4 mM CaCl₂ and a Hill coefficient = -2.2, compatible with binding to M1 and M2 (see S2 Fig.). The presence of 20 μM ZnCl₂ in the substrate buffer only displaced the dose-response curves weakly (Fig. 2B), compatible with negligible

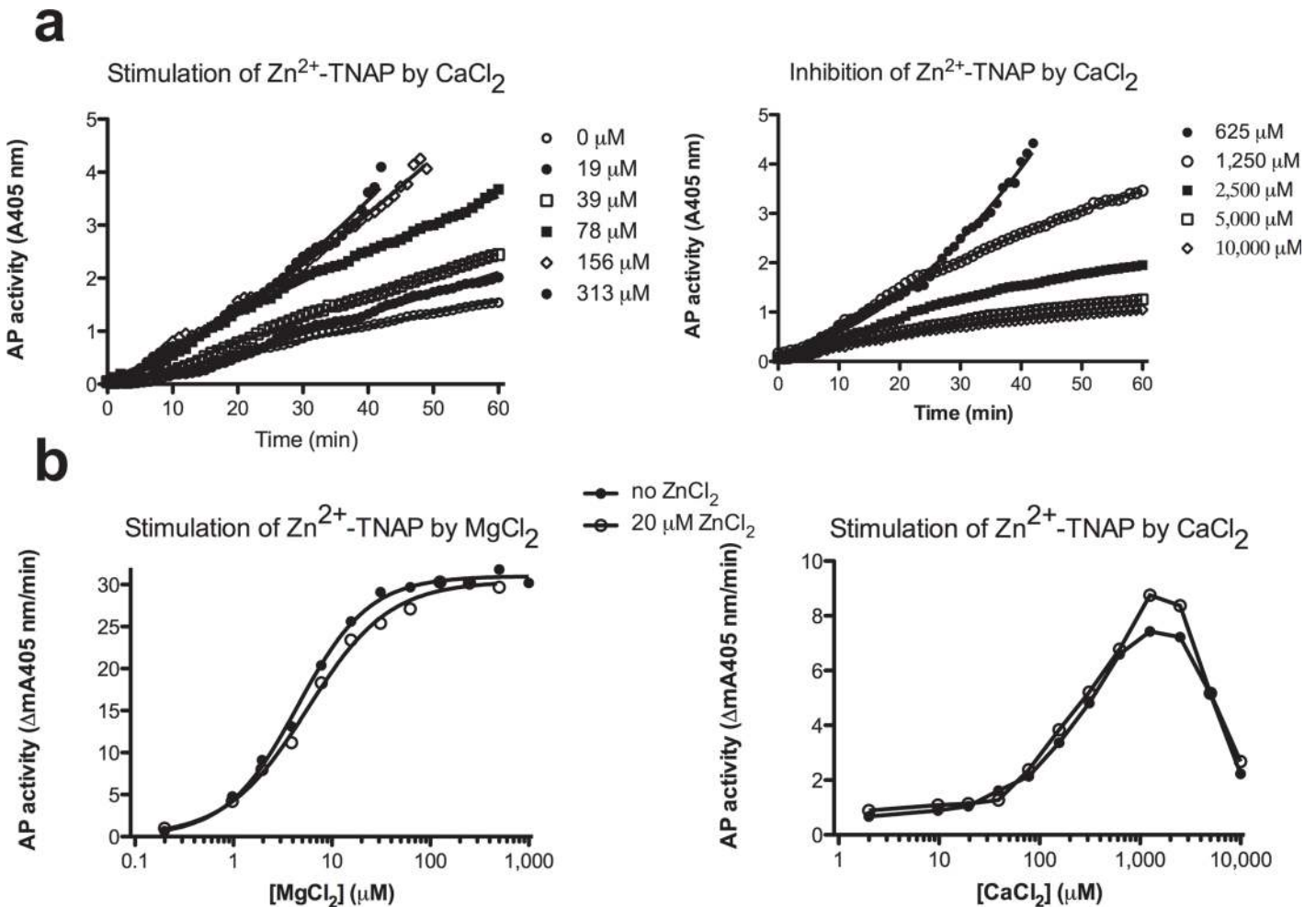


Fig 2. Allosteric activation of Zn²⁺-TNAP by CaCl₂. a. Progressive AbM2-bound Zn²⁺-TNAP activation, measured as A405 nm vs. time, for the indicated [Ca²⁺], added to Chelex-pretreated pNPP (10 mM) at pH 9.8, showing dose-dependent activation (left panel) and inhibition at high concentrations (right panel); b. Dose-response of generated AP activity (mean mA405nm/min) in steady-state (i.e. the slope measured between 60–90 min in Figs. 1A and 2A) for increasing [MgCl₂] and [CaCl₂] at identical AbM2-bound [Zn²⁺-TNAP], reflecting the plateau and pseudo-plateau at high [MgCl₂] and [CaCl₂] respectively, followed by a steep drop of the TNAP activity in the case of CaCl₂ (right panel). Activities were measured in Chelex-treated pNPP (10 mM) at pH 9.8, supplemented with MgCl₂ and CaCl₂, as indicated. Experiments representative of at least three replicates with variable enzyme and MgCl₂ concentrations.

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functional competition between Mg²⁺ or Ca²⁺ and low [Zn²⁺] at M3, at pH 9.8, and essentially confirming that the loading of TNAP with 20 μM ZnCl₂ allowed the measurement of allosterism in conditions where Zn²⁺-TNAP had been preformed.

To account for complexation between Ca²⁺ (and Mg²⁺) and dissociated phosphate ions in pNPP at pH 9.8 [21], dose-response studies were also repeated at 1 mM pNPP, to reduce the loss of metal ion as a result of its inactivating complexation to pNP-PO₄³⁻, while taking care to choose sufficiently low enzyme concentrations to not disrupt pseudo-first order conditions (Fig. 3A). M3 saturation by MgCl₂ occurred with an identical plateau (33.7 mA405nm/min), i.e. did not affect TNAP activity when M3 was saturated with Mg²⁺, but the saturation curve underwent a considerable leftward-shift with K_d = 0.52 ± 0.03 μM (8-fold lower as value at 10 mM pNPP, two tailed p < 0.0001). Also, M3 saturation by CaCl₂ (K_d = 66 ± 4 μM, with a more accurately determined plateau at 14.2 mA405nm/min) also shifted to lower concentrations (3-fold lower as value at 10 mM pNPP, two tailed p < 0.0001), now also reflecting a

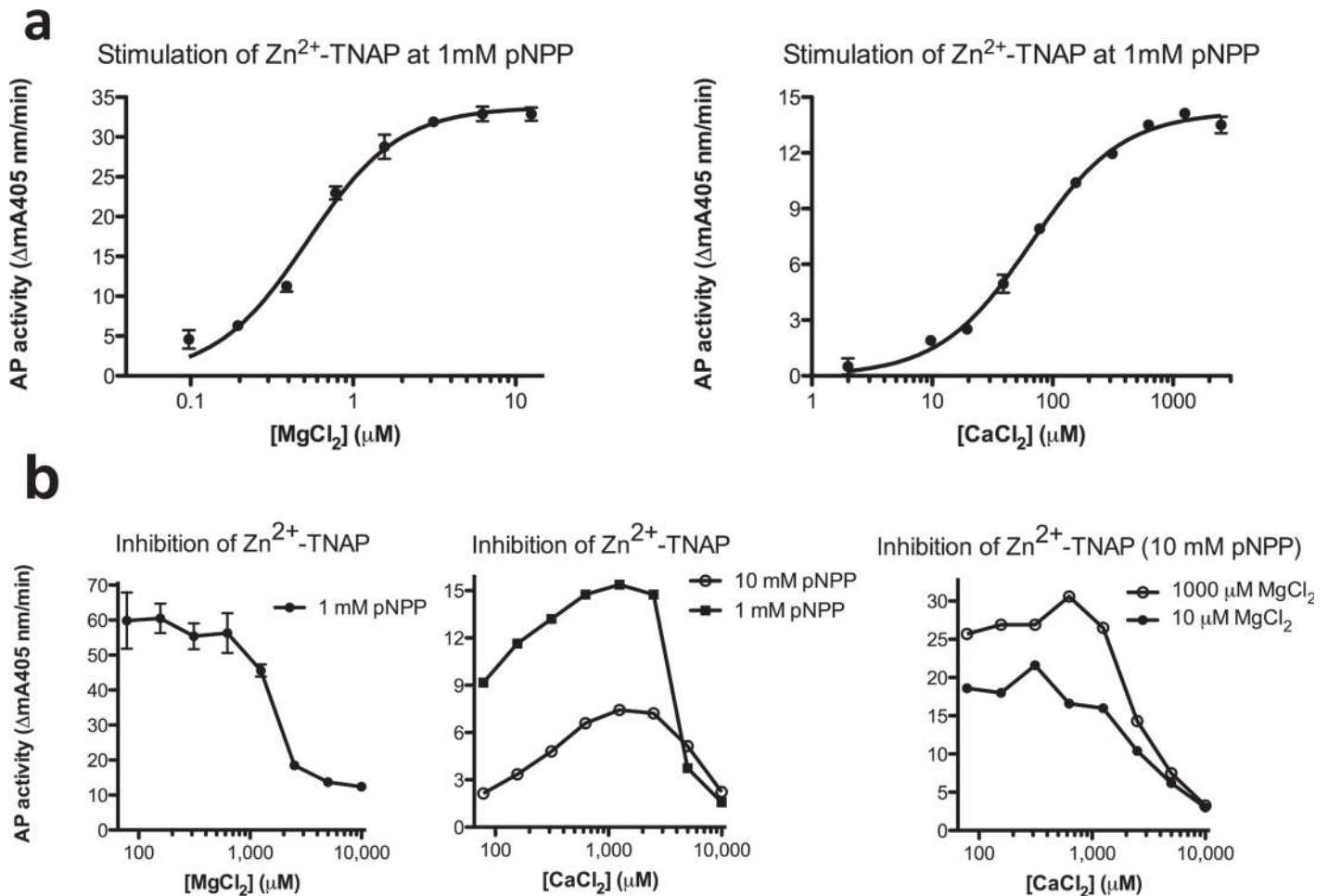


Fig 3. Activation vs. inhibition of Zn^{2+} -TNAP by MgCl_2 and CaCl_2 . **a.** Dose-response of generated AP activity (mean mA405nm/min) in steady-state (i.e. the slope measured between 60–90 min) for increasing $[\text{MgCl}_2]$ and $[\text{CaCl}_2]$ at identical AbM2-bound $[\text{Zn}^{2+}$ -TNAP], measured in Chelex-treated pNPP (1 mM) at pH 9.8; **b.** TNAP inhibition at high $[\text{MgCl}_2]$ and $[\text{CaCl}_2]$, measured in Chelex-treated pNPP (1 or 10 mM as indicated) at pH 9.8, in the presence of the indicated concentrations of MgCl_2 . Results represent mean \pm SD for 3 identical experiments or are representative examples of experiments, performed in 3-fold (b, middle and right panel).

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more precise ratio corresponding to a 2.4-fold lower maximal activity for $\text{Ca}^{2+}/\text{Zn}^{2+}$ -TNAP than for $\text{Mg}^{2+}/\text{Zn}^{2+}$ -TNAP. High $[\text{MgCl}_2]$ inhibited TNAP activity between 1–10 mM (Fig. 3B, left panel), consistent with the known inhibitory role of Mg^{2+} at M1 and M2 [19] at pH 9.8. Likewise, the descending limbs of the CaCl_2 curves in Fig. 2B, middle and right panel hardly differed, as a function of the concentration of pNPP or MgCl_2 in the test tube, excluding the possibility that this drop in activity was determined by the degree of Ca^{2+} -pNP- PO_4^{3-} complexation or the degree of M3 saturation by Mg^{2+} (Fig. 3B, left panel). The activity of $\text{Ca}^{2+}/\text{Ca}^{2+}$ -TNAP at 10 mM CaCl_2 was again about 10% of that of $\text{Ca}^{2+}/\text{Zn}^{2+}$ -TNAP and 4.2% of that of $\text{Mg}^{2+}/\text{Zn}^{2+}$ -TNAP (Fig. 3B, middle and right panel).

Functional relevance of the M4 site

The crystal structure of PLAP had uncovered a putative Ca^{2+} -binding site (M4) in a peripheral location [22], but its significance for AP function remains unknown. To investigate whether binding of Ca^{2+} to M4 also contributes to AP activity, in addition to its binding to M3, a series

of site-directed mutants were produced, in which residues potentially coordinating Ca²⁺ in M4 were mutated to alanine in PLAP (where the site was documented) and at the homologous residues in TNAP. Fig. 4A-C displays how those residues (W248, R250, E216, F269, E270, D285 in PLAP; E218, W253, R255, E273, E274, D289 in TNAP) are positioned around the fixed Ca²⁺ ion in a structural homology model of TNAP. Part a in S3 Fig. shows that all mutants were secreted as FLAG-tagged enzymes. Classical kinetic activity measurements showed that one PLAP mutant and three TNAP mutants were inactive. The remainder of the mutants showed mildly affected kinetic parameters when analyzed via Michaelis-Menten kinetics, using pNPP as substrate without added CaCl₂ (Table 1). The overall structural impact of most mutations was very limited for the PLAP mutants, with most mutants showing heat inactivation curves comparable to that of reference PLAP (Part b in S3 Fig.). Although TNAP is structurally less stable than PLAP [13], little structural influence on heat stability was noted for the active mutants (Part c in S3 Fig.); in these cases the analysis was done as a function of time at a constant temperature to more gently inactivate the more labile reference TNAP and its mutants.

Structural effects were further investigated both for the active and inactive TNAP mutants, applying an anti-TNAP monoclonal antibody mapping approach using a panel of 19 epitope-mapped antibodies [13], to measure relative affinities in the presence or absence of 1 mM CaCl₂. Fig. 5 shows that the inactive mutants reacted poorly with the four most discriminating antibodies, indicating that these mutants were not folded properly to maintain a functionally active site. However, this approach could not identify any effect of CaCl₂ on the affinity of the antibody panel for TNAP or any mutant. Hence, this structural probing, essentially targeting the entire TNAP surface [13] revealed that some mutations perturbed the 3D structure of the resulting TNAP mutants, but that these structural changes occurred independently of the presence of CaCl₂, as sensed by the anti-TNAP antibody panel.

To further investigate whether Ca²⁺ binding to M4 contributes to the allosteric activation of TNAP by CaCl₂, activation of each mutant was analyzed as a function of the concentration of Mg²⁺ and Ca²⁺. The rationale was that M4 mutations would not differentially alter the functional consequences of Mg²⁺ or Ca²⁺ binding to M3, and would impact the overall stimulation by Ca²⁺ only if the M4 site would significantly contribute to TNAP activity, a regulation expected to be defective in at least some mutants. Fig. 6 shows the biphasic Ca²⁺-saturation curves for the three active TNAP mutants. Since the K_m for pNPP varies slightly at pH 9.8 for the various mutants, these experiments were conducted at 10 mM pNPP, i.e. yielding rather apparent than true K_ds for the binding of Mg²⁺ and Ca²⁺. Mutations in the active mutants affected the affinity of Mg²⁺ and Ca²⁺ for M3 to some extent, but also that of Zn²⁺ for M3, as evident from the different relative inhibition of TNAP mutants in the presence of 20 μM ZnCl₂ (Fig. 6). However, the apparent K_ds measured for Mg²⁺ and Ca²⁺ (in the absence of added ZnCl₂) correlated strongly (r² = 0.97), showing that Mg²⁺ and Ca²⁺ regulate Zn²⁺-TNAP at the same functionally relevant site, i.e. M3, which was affected by mutations to a comparable degree for both Mg²⁺ and Ca²⁺. These findings identify the TNAP M4 site as a structural determinant, indirectly determining TNAP activity, rather than as a Ca-site directly implicated in the control of enzyme catalysis. As observed above, all M4 mutants were partly inactivated at 10 mM CaCl₂, an inhibition independent of the presence of 20 μM ZnCl₂ (Fig. 6).

Ca²⁺ in TNAP regulation at physiological pH

TNAP activity is routinely analyzed at its alkaline pH optimum, but to properly understand the impact of Ca-homeostasis on the physiological TNAP activity, which is to hydrolyze ATP and pyrophosphate primarily during mineralization [23], we also studied ionized Ca²⁺-interactions with TNAP at pH 7.4. At this pH, coordinating active site residues are more protonated

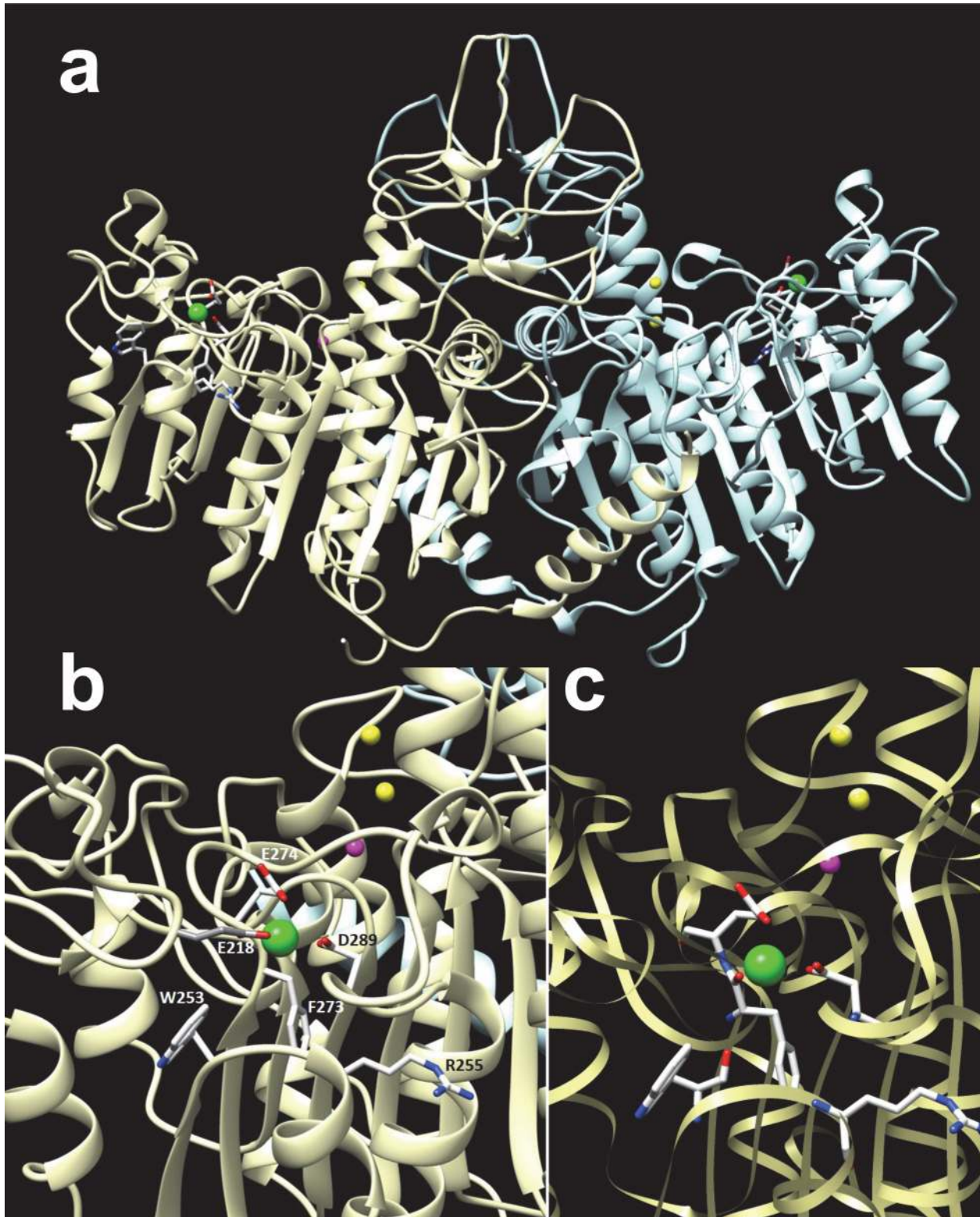


Fig 4. Representation of M4 ligands in the modeled structure of TNAP. Rendering of the 3D structure of the entire TNAP dimer in front view (a) and coordinating residues, detailed in a lateral zoom of the shoulder region harboring M4, in larger detail (b) and in ribbon representation (c).

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Table 1. Kinetic Parameters of PLAP, TNAP and the M4-site mutants, measured in 1M DEA buffer, pH 9.8, with pNPP as a substrate.

enzyme	K _m (mM)	k _{cat} (s ⁻¹)
Placental Alkaline Phosphatase (PLAP)		
WT	1.8	460*
E216A	0.9	236
W248A	1.5	165
R250A	1.8	424
F269A	inactive	inactive
E270A	1.1	200
D285A	0.7	18
Tissue-Nonspecific Alkaline Phosphatase (TNAP)		
WT	0.6	1102*
E218A	inactive	inactive
W253A	0.6	212
R255A	2.3	307
F273A	inactive	inactive
E274A	1.1	384
D289A	inactive	inactive

* based on historical values [13]

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and the pKs describing phosphate dissociation favor preponderance of HPO₄²⁻, an ion with a higher solubility product for Ca²⁺ than PO₄³⁻, the predominant phosphate ion at pH 9.8 [21]. Since, moreover, the K_m for pNPP is very low at this pH, [pNPP] was kept at 1 mM in all cases. Also at pH 7.4, Mg²⁺ binding to M3 is slow and exponential (Fig. 7A, left panel), i.e. Mg²⁺/Zn²⁺-TNAP complex formation is not complete at 90 min, even at [MgCl₂] = 100 μM). In contrast, binding at M3 is relatively fast for Ca²⁺, reaching steady-state after 10–20 min. In a physiological environment where Ca²⁺ and Mg²⁺ are both present, they display additive effects on M3 during activation of Zn²⁺-TNAP. As a consequence of its faster binding at pH 7.4, Ca²⁺ has a relative competitive advantage during binding, 1 mM CaCl₂ capable of enhancing the activity of forming Mg²⁺/Zn²⁺-TNAP, due to faster Ca²⁺-binding to M3 (Fig. 7A, left panel), Fig. 7A, right panel further illustrates the additive interplay of both metal ions on Zn²⁺-TNAP activation at pH 7.4, illustrating their near-equivalence.

At pH 7.4, TNAP was dose-dependently activated by CaCl₂ (Fig. 7B, left panel). From the ascending limb a K_d = 217 ± 50 μM was calculated for the binding of Ca²⁺, with a plateau activity of 11.1 mA405nm/min. This value differed from K_d = 66 ± 4 μM, determined at pH 9.8 by a factor 3 only (two tailed p < 0.0001). Similar plots of TNAP saturation by Mg²⁺ (Part a in S4 Fig., left panel) yielded a K_d = 82 ± 27 μM for the binding of Mg²⁺, i.e. considerably higher than K_d = 0.52 ± 0.03 μM determined at pH 9.8 (two-tailed p < 0.0001) with a plateau activity of 27.4 mA405nm/min. In conclusion, at M3 Ca²⁺ is almost equipotent to Mg²⁺ at pH 7.4, with plateau activities again differing 2.5-fold. Yet, some TNAP inactivation was noted at 10 and 20 mM CaCl₂ (Fig. 7B, left panel), milder than at pH 9.8, but not absent.

Ca²⁺-mediated loss of activity at pH 7.4

Fig. 7B, right panel shows that the binding of TNAP to AbM2 in the presence of 5–20 mM CaCl₂ for 3 h, resulted in a dose-dependent 2.5-fold reduction of TNAP activity at 20 mM,

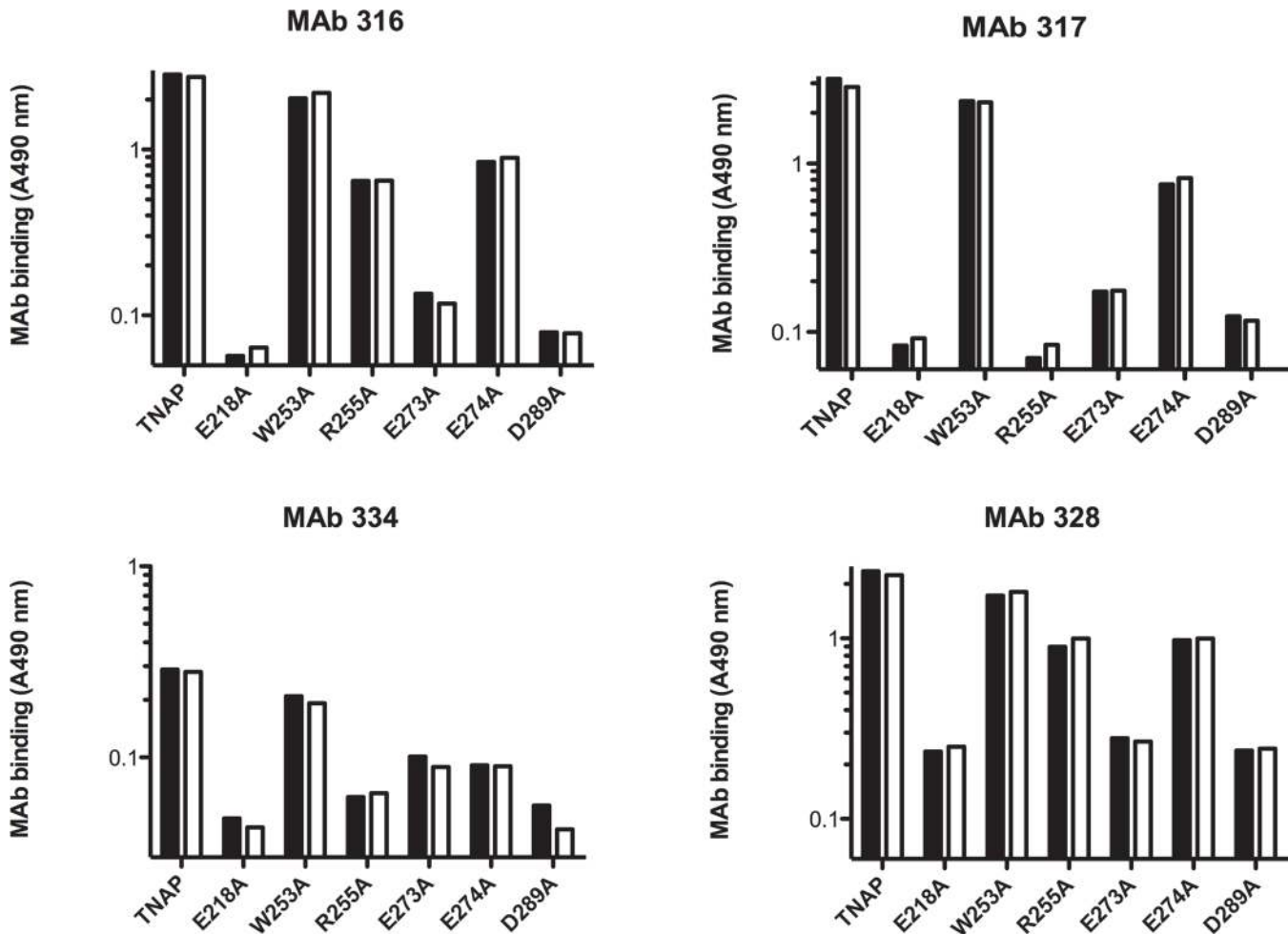


Fig 5. Structural mapping of TNAP M4 mutants. Degree of binding of TNAP and the indicated TNAP mutants to four epitope-mapped monoclonal anti-TNAP antibodies (of 19 studied). Binding was analyzed in the absence (black bars) and presence (white bars) of 1 mM CaCl₂. TNAP was bound to microtiter plates coated with AbM2 and bound TNAP (mutant) was detected with AbM2, recognizing the FLAG tag.

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when subsequently measured in Chelex-treated pNPP, without further metal ions. ZnCl₂, added to the pNPP substrate hardly affected the activity measured, but added MgCl₂ recovered activity fully after pre-incubation with 5 mM CaCl₂ and for about 80% after pre-incubation with 10 and 20 mM CaCl₂ ($p < 0.005$ for 10 and 20 mM combined vs. 0 mM). These experiments illustrated that CaCl₂ readily displaced M3-bound Mg²⁺, reducing TNAP activity 2.5-fold, as expected, a loss easily recovered by Mg²⁺ added to the pNPP substrate.

However, the irreversible loss of TNAP (20%) at higher [Ca²⁺] (10–20 mM) was compatible with some Ca²⁺-induced TNAP inactivation. To enable proper study of the interaction of Ca²⁺ and M1 and M2, we have prepared holo-TNAP, by overnight incubating AbM2-bound TNAP with 250 μM EDTA, at room temperature. This procedure fully stripped TNAP from its bound metal ions (Fig. 8A, left panel), resulting in complete TNAP inactivation, as measured with chelex-treated pNPP, and showing minor enzyme activity upon inclusion of 20 μM ZnCl₂ in the substrate. Whereas 1 mM MgCl₂ did not cause activation, combined, ZnCl₂ + MgCl₂ reconstituted TNAP over 1 h to over 80% of its initial activity. Fig. 8A, right panel shows the calculated enzyme activities for reconstituted TNAP, measured in different conditions. Overnight

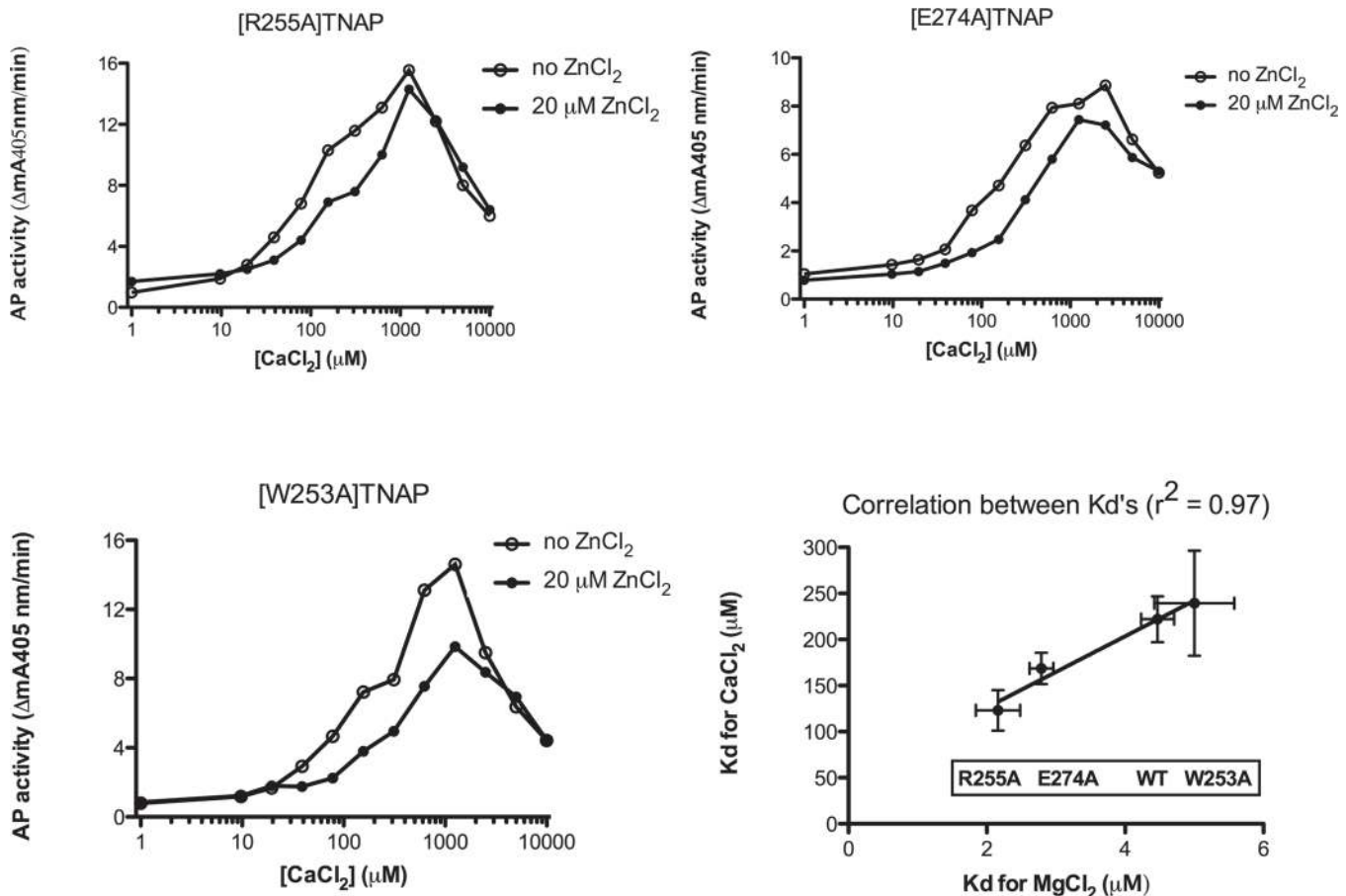


Fig 6. Role of Ca²⁺ binding to M4 in TNAP activity. Dose-response of stimulation of AP activity (mean mA405nm/min) in steady-state (i.e. the slope measured between 60–90 min) at pH 9.8 (10 mM pNPP) for increasing [CaCl₂], added to AbM2-bound Zn²⁺-TNAP and the indicated mutants, after pre-treatment with EDTA (2 h) and loading with 20 μM Zn²⁺; correlation between calculated apparent K_ds for the functionally relevant Mg²⁺ and Ca²⁺ binding to TNAP and the three indicated TNAP mutants. Lines constructed in the absence and presence of 20 μM ZnCl₂ are as indicated; Apparent K_ds are represented with their respective SD.

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incubations with chelex-treated TBS, lacking EDTA were less efficient, confirming the presence of low residual TNAP-bound Zn²⁺, acting in concert with MgCl₂ in the substrate.

Fig. 8B, left panel shows that [CaCl₂] unexpectedly triggered holo-TNAP activation dose-dependently, with an apparent K_d = 509 ± 49 μM. To quench this non-specific TNAP activation, triggered by divalent metal ion contaminants [24], we performed further dose-response studies in the presence of known concentrations of Zn²⁺ (0.2–20 μM). This procedure shifted Ca-saturation curves to lower [CaCl₂] with K_d = 144 ± 49 μM for 0.2 μM and K_d = 137 ± 24 μM for 2 μM, i.e. in fair agreement with K_d = 217 ± 50 μM, for the binding of CaCl₂ to Zn²⁺-TNAP at M3 (Fig. 7B, left panel).

Therefore, to analyze the competition between Ca²⁺ and Zn²⁺ vs. Ca²⁺-induced Zn-displacement from M1 and M2, first AbM2-bound holo-TNAP was incubated with high [CaCl₂] in the presence of 2 or 20 μM ZnCl₂ (thus quenching non-specific TNAP activation by divalent metal ions in the CaCl₂ solutions). Fig. 8B, right panel confirms that CaCl₂ competes with ZnCl₂ for binding to M1 and M2 at [CaCl₂] > 10 mM, more efficiently when [ZnCl₂] is low (2 μM), with an apparent IC₅₀ around 250 mM. On the contrary, when AbM2-bound TNAP was fully

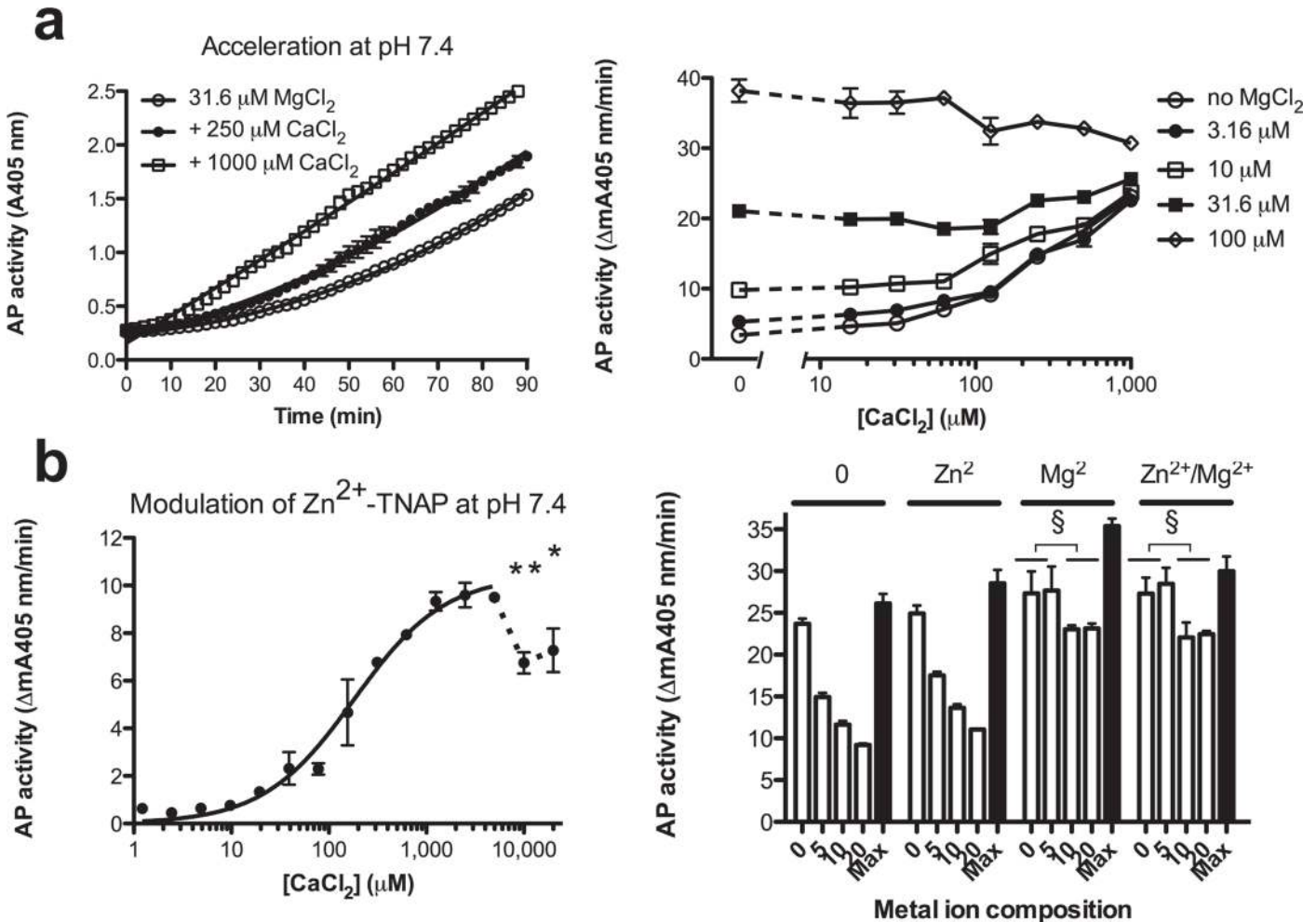


Fig 7. Consequences of Ca²⁺ binding to TNAP at pH 7.4. **a.** Kinetics of TNAP activation by low [Mg²⁺] and acceleration of activation during formation of Mg²⁺/Zn²⁺-TNAP by increasing [CaCl₂] as indicated, measured as A405 nm (left panel); additive effects for various combinations of Mg²⁺ and Ca²⁺ during TNAP activation (measured in steady-state, right panel); **b.** Dose-response of activation and partial inhibition of TNAP by CaCl₂, at the indicated concentrations (left panel); TNAP activity recovery in chelex-treated pNPP, containing the indicated metal ion composition (0: no metal ion; [Zn²⁺] = 20 μM; [Mg²⁺] = 1 mM; Zn²⁺/Mg²⁺ = 20 μM Zn²⁺ + 1 mM Mg²⁺) after 3 h of TNAP binding to AbM2 in the presence of co-incubated CaCl₂ (0–20 mM, as indicated) (right panel); black bars: corresponding activity for maximally active TNAP, (pre-incubation for 3 h in TBS, containing 20 μM ZnCl₂ + 1 mM CaCl₂). Results represent mean ± SD for 3 identical experiments. (*p<0.05 and **p<0.01 vs. plateau, §p<0.005 vs. [CaCl₂] = 0).

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loaded overnight with 20 μM ZnCl₂, CaCl₂ only displaced bound Zn²⁺ at concentrations as high as 100 mM, independently of the presence of soluble ZnCl₂ (0–20 μM).

Discussion

APs are zinc metalloenzymes, with Zn²⁺ binding to M1 and M2, and allosterically activated by Mg²⁺ binding to M3 [4, 6]. Whereas substitution of Zn²⁺ by most metal ions, except Co²⁺ inactivates enzyme activity, allosteric activation of APs can also be provided at M3 by other ions like Mn²⁺, Co²⁺, Ni²⁺, including Ca²⁺ [25]. Because matrix vesicle-induced mineralization is a process that requires the generation of reaction products by TNAP, the present work was undertaken to study TNAP functionality during exposure to an increasing Ca²⁺ gradient, including the mechanism of TNAP inhibition, at high [CaCl₂] [26, 27]. Inspired by the existence of a fourth metal ion site, occupied by Ca²⁺ [4], we have presently investigated contributions by all

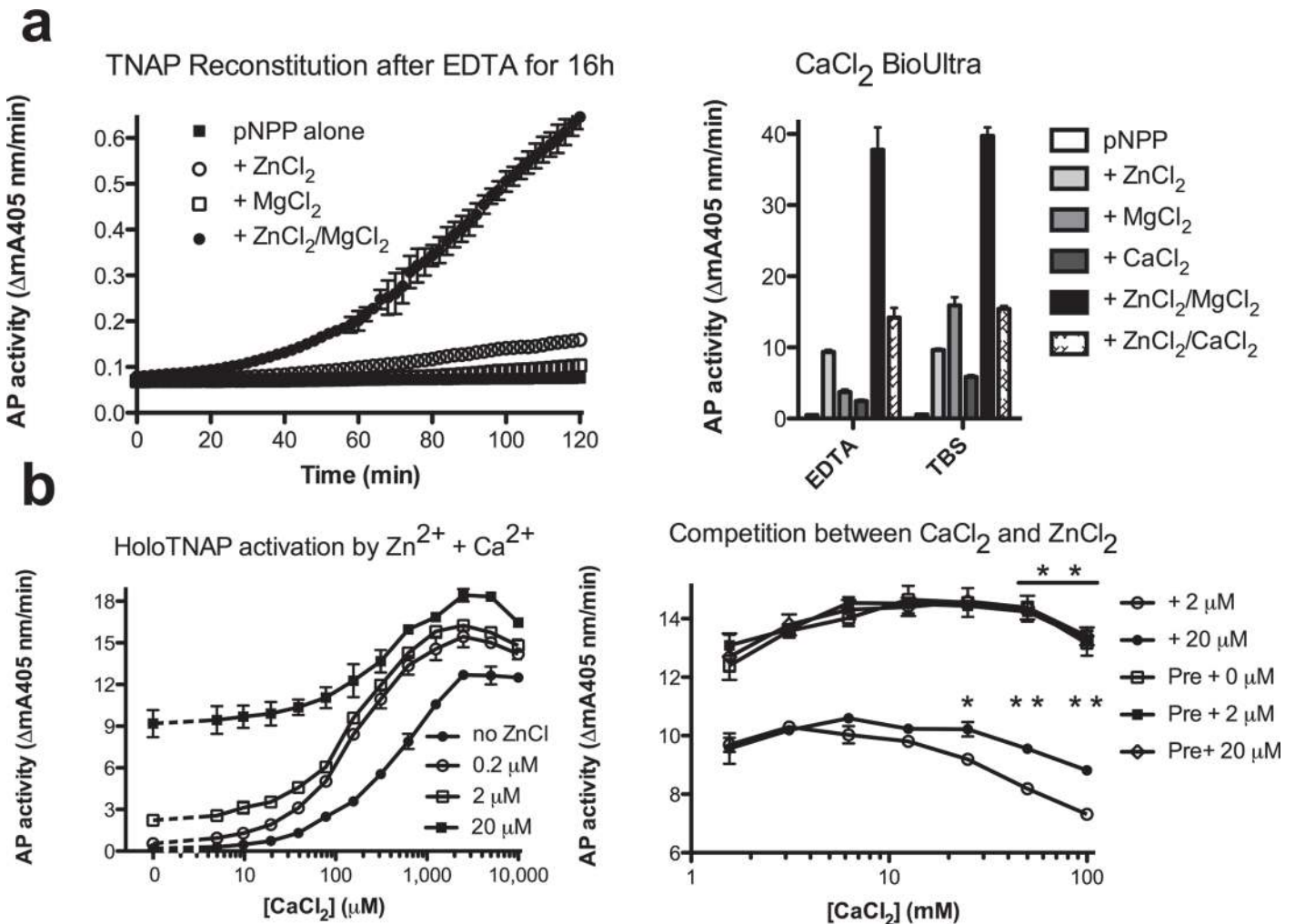


Fig 8. TNAP inactivation by high [CaCl₂] at pH 7.4. a. Reconstitution of TNAP activity by 20 μ M Zn²⁺ + 1 mM Mg²⁺, but not by the individual metal ions, added to fully demetalated holo-TNAP, starting after 60 min (left panel); comparison of specific TNAP activity of (holo)-TNAP, after reconstitution with the indicated metal ion composition, after overnight treatment with TBS, with or without added EDTA (250 μ M); b. Dose-dependency of holo-TNAP reconstitution by CaCl₂ (0–10 mM), in the absence or presence of the indicated [ZnCl₂] (0–20 μ M, left panel); competition between the indicated concentrations of Zn²⁺ (2 μ M and 20 μ M) and increasing concentrations of CaCl₂; minor displacement of TNAP-bound Zn²⁺ (overnight Zn²⁺ preloading indicated as “Pre”) by increasing concentrations of CaCl₂, independently of the presence of free Zn²⁺. (0–20 μ M). Results represent mean \pm SD for 3 identical experiments (*p<0.003, **p<0.0001.).

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four metal sites. We found that Ca²⁺, by binding to M3 is a fairly good allosteric activator of TNAP when bound to M3, but that binding at M4 hardly influences the catalytic activity of TNAP. A strong determinant of TNAP activity is the availability of Zn²⁺, free Zn²⁺ being competed out by high Ca²⁺ concentrations and TNAP-bound Zn²⁺ being displaced from M1 and M2 at still higher Ca²⁺ concentrations, both resulting in virtual TNAP inactivation. Elegant zinc mapping studies in osteons [28] have revealed co-distribution of alkaline phosphatase with zinc at the calcification front, providing an explanation for the long-lasting presence of Zn²⁺-TNAP in bone matrix.

In humans, baseline plasma Zn²⁺ concentrations average around 12 μ M [29] and free plasma Mg²⁺ averages 0.4–0.6 mM, with free Ca²⁺ averaging 1.1–1.3 mM [30]. The present affinity determinations at physiological pH therefore predict circulating TNAP is properly charged at M1 and M2 with Zn²⁺ and is saturated at M3 primarily with Mg²⁺. However, in an

environment where matrix vesicles generate a gradient of Ca²⁺ during early mineralization and TNAP generates P_i from ATP, PP_i and other physiological substrates, the relative balance between divalent metal ions as found in plasma will gradually be disturbed by gradients of P_i and PP_i, inducing formation of poorly soluble hydroxyapatite. Our present findings confirm at pH 7.4 that Ca²⁺ and Mg²⁺ are quite complementary in the allosteric activation of TNAP. Thus, a relative drop of [Mg²⁺] is not a matter of concern, since transported Ca²⁺ is capable of adequately substituting for Mg²⁺. Indeed, at physiological pH, the affinities of Ca²⁺ and Mg²⁺ for M3 only differ 2–3 fold and the maximal activity is only 2.5 fold weaker for Ca²⁺/Zn²⁺-TNAP than for Mg²⁺/Zn²⁺-TNAP.

Mg²⁺ can also easily be replaced at M3 by Mn²⁺, Co²⁺ and Ni²⁺ [19, 25], but our present findings confirm that Mg²⁺ does not generate activity when incubated with the apoenzyme, as a result of binding to M1 and M2 [19]. At physiological pH, TNAP has a low K_m for common substrates [13, 18] or relative catalytic efficiency comparisons between different activity states are dictated by the catalytic rate constants primarily. TNAP has a 50-fold lower k_{cat} for pNPP at pH 7.4 than at pH 9.8. Correspondingly, also the affinities of catalytically active metal ions differ at both pHs and physiologically relevant comparisons for metal ion substitutions in the TNAP active site can only be made representatively at pH 7.4. Correspondingly, we presently found that Ca²⁺ binds to M1 and M2, rapidly at pH 9.8, but more slowly at pH 7.4, a process completing dissociation of bound Zn²⁺, a slow process, because of the high affinity of Zn²⁺ [20]. Hence, the main conclusion of our present work is that TNAP is extremely robust in a Ca²⁺-rich (patho)-physiological environment. The rapid substitution of Mg²⁺ for Ca²⁺ in M3 hardly results in any loss-of-function. The slow substitution at pH 7.4 of Zn²⁺ for Ca²⁺ as a result of competition or Zn-displacement at M1 and M2 generates an enzyme (Ca²⁺/Ca²⁺-TNAP) 20-fold less active as the parent Mg²⁺/Zn²⁺-TNAP and still 10-fold less active as Ca²⁺/Zn²⁺-TNAP.

We have noted before that specific amino acid substitutions affecting catalysis at pH 9.8 did not have a similar effect at physiological pH [18, 31]. Yet, the relative residual activity, measured for Ca²⁺/Ca²⁺-TNAP at pH 9.8 and pH 7.4 are comparable. On a relative scale, on which Mg²⁺/Zn²⁺-TNAP is 100% active at pH 9.8, Ca²⁺/Zn²⁺-TNAP is 40% active and Ca²⁺/Ca²⁺-TNAP is 5% active. On that same scale, at pH 7.4, Mg²⁺/Zn²⁺-TNAP is 2% active and Ca²⁺/Zn²⁺-TNAP is 0.8% active, Ca²⁺/Ca²⁺-TNAP extrapolated to be virtually inactive. From a physiological perspective, calcium incorporation in TNAP does not destroy TNAP, but the relative balance between ionic calcium, P_i, PP_i, other divalent ions and the relative availability of Zn²⁺ during synthesis of TNAP are all crucial factors, determining proper charging at M1 and M2 during long exposure to high Ca²⁺ concentrations.

We found that the impact of Ca²⁺ on TNAP activity could be explained entirely by its interactions with M1–3. In contrast, contributions by M4 were structural. Mutations of several TNAP ligands coordinating the M4 binding site did affect the conformation of the resulting mutants to a variable degree, from minor effects for some mutants to complete loss of the 3D-structure for others, as concluded from combined epitope analysis by an antibody panel of 10 antibodies, heat inactivation studies and classical kinetic analysis. Compared to native TNAP, some mutants manifested a mildly influenced affinity for Mg²⁺ binding to M3, indicative both of gain-of-function, as well as loss-of-function. The relative effects on the affinity for Ca²⁺ were very similar, i.e. the respective K_ds for Mg²⁺ and Ca²⁺ correlated well for the various mutants and native TNAP. These TNAP M4 mutants manifested slight changes in their allosteric properties, which could fully be explained by the allosteric properties of M3, i.e. we did not find any evidence for a role of M4 in catalysis. Instead, our structural analyses identified some ligands of the M4 site to be critical structural elements of TNAP which, despite their distant location from the active site have a dominant role on the active site integrity when mutated to residues such as encountered in some hypophosphatasia patients (http://www.esep.uvsq.fr/03_hypo_mutations.php). Yet, we did not

observe any structural change for the anti-TNAP monoclonal antibody panel, when affinities were measured in the absence or presence of 1 mM CaCl_2 , despite detection of primary changes in 3D-structure in the TNAP mutants. These findings also rule out that Ca^{2+} binding to M4 will participate in structural folding of the M4 ligand area.

We have previously described APs as allosteric enzymes in which asymmetry between monomers generates activity patterns which differ from the expected weighed properties of the two monomers [32]. In particular, negative cooperativity can be generated between both monomers, when they are differently metalated. It is to be expected that a gradient of CaCl_2 will not cause parallel substitutions in both monomers, i.e. generate asymmetry. Such may generate mixed enzymatic properties as complex as those presently found during the simultaneous reconstitution of partially demetalated TNAP with mixtures of Zn^{2+} and Ca^{2+} , respectively, or Zn^{2+} , Ca^{2+} and Mg^{2+} . In case of active site asymmetry, cross-occupation of binding sites by the “wrong” metal may generate response profiles, hardly predicted by more straightforward approaches, based on preloaded TNAP in equilibrium conditions. Since each TNAP dimer needs to accommodate 4 metal ions in M1 and M2, before reaching symmetry, some activity measurements in intermediate stages may reflect such more complex behavior. Presently, such conditions were met during the interaction of Zn^{2+} -TNAP with 10 and 20 mM CaCl_2 , modulating TNAP-activity in a time-dependent manner. Even highly pure CaCl_2 contains metal ion contaminants, the most abundant one Sr^{2+} , a potent TNAP activator [24]. Inclusion of standardized $[\text{ZnCl}_2]$ could overcome this limitation, allowing proper competition and displacement studies between ZnCl_2 and high $[\text{CaCl}_2]$ at pH 7.4.

Medial vascular calcification is associated with chondrocyte transdifferentiation and expression of TNAP [33]. It is to be expected that also in this environment, where a gradient of calcium builds up, TNAP will be gradually substituted with Ca^{2+} at all metal ion sites. Our work also suggests that TNAP in other Ca^{2+} -rich environments can act as $\text{Ca}^{2+}/\text{Zn}^{2+}$ -TNAP, e.g. regulating the hydrolysis of phospholamban in the sarcoplasmic reticulum of cardiomyocytes and in skeletal muscle [34]. The injection of Zn acetate into the tail vein of mice enhanced TNAP activity in the sarcoplasmic reticulum of the cardiac sarcomere, leading to increased dephosphorylation of phospholamban. This finding supports the interpretation that also in the cardiac sarcomere TNAP activity is tempered by high prevailing Ca^{2+} levels [34]. The strong correlation between the loss of TNAP activity and the accumulation of calcium during MV-mediated mineralization, observed by Genge et al. [26] can also be explained by our present findings. In these studies TNAP activity present at the site of the MV-dependent mineralization process was found to be profoundly reduced by the mineralization process, a finding that can be explained by $[\text{CaCl}_2]$ -dependent conversion of $\text{Ca}^{2+}/\text{Zn}^{2+}$ -TNAP into $\text{Ca}^{2+}/\text{Ca}^{2+}$ -TNAP.

In conclusion, our work has identified that ionized calcium supports TNAP activity in Ca^{2+} -rich milieus, until very high concentrations of Ca^{2+} occupy M1 and M2 leading to greatly reduced enzymatic activity.

Supporting Information

S1 Fig. Short-term reconstitution of Zn^{2+} -TNAP activity at pH 7.4. a. Progressive Zn^{2+} -TNAP formation, measured from the increase of A405 nm vs. time during pNPP hydrolysis, after incubation of AbM2-bound TNAP with 1 mM EDTA (2 h) followed by addition of the indicated $[\text{Zn}^{2+}]$ (2 h), dissolved in Chelex-treated TBS, followed by addition of Chelex-treated pNPP (10 mM) at pH 9.8; b. Slope (first derivative) to the line for 40 μM ZnCl_2 in Fig. 1A; c. Dose-response of Zn^{2+} -TNAP formation, from plots of initial AP activity (ΔA405 nm/20 min) vs. the indicated $[\text{Zn}^{2+}]$; the red line represents the corresponding AP activity for native non-EDTA treated

AbM2-bound TNAP. Results are representative of 3 independent experiments. (TIF)

S2 Fig. Activation and inhibition of EDTA-treated TNAP by CaCl₂, at pH 9.8. a. Progressive activation and inhibition of AbM2-bound EDTA-treated TNAP in the absence (left panel) and presence (right panel) of 2 μM ZnCl₂, measured from its activity at A405 nm vs. time in Chelex-treated pNPP (10 mM) at pH 9.8; Note the curvi-linearity at low [CaCl₂] and linearity at 5 and 10 mM CaCl₂ respectively (left panel); the maximal activity (“max”) represents activity of fully metalated TNAP in pNPP, pH 9.8, measured in the presence of 20 μM ZnCl₂ and 1 mM MgCl₂; b. Dose-response of TNAP activation and inhibition, from plots of AP activity (ΔmA405nm/min calculated between 60–90 min) vs. the indicated [CaCl₂] (left panel) or [MgCl₂] (right panel); ●: absence of ZnCl₂; ■: 2 μM ZnCl₂ mixed with CaCl₂; μ: 2 μM ZnCl₂ mixed with MgCl₂. *Insert*: one-site binding model fit for the ascending limb, without added Zn²⁺. Results are representative of 3 independent experiments. (TIF)

S3 Fig. Production and stability of TNAP M4 mutants. a. Western blots of TNAP, PLAP and their mutants, after purification from COS-1 cellular medium, via AbM2 detection; b. Heat inactivation curves of PLAP and the indicated mutants, plotted as residual activity after 10 min incubation at the indicated temperature; c. Heat inactivation curves of TNAP and the indicated mutants, plotted as residual activity after incubation for the indicated time interval at 56°C in TBS. Results are representative of 3 independent experiments. (TIF)

S4 Fig. Inhibition kinetics of TNAP by MgCl₂ and CaCl₂. a. Dose-response of Zn²⁺-TNAP inhibition by high [MgCl₂] (0–20 mM) at pH 7.4; AP activity was measured as mean mA405nm/min in steady-state (between 60–90 min) (left panel); tracings of TNAP activity vs. time, in the presence of the indicated medium to high [MgCl₂] at pH 7.4, illustrating a slight deflection in hydrolysis rate after 60 min (solid line vs. dotted line) (right panel); b. Similar tracings, in the presence of the indicated medium to high [CaCl₂], illustrating clear deflection in hydrolysis rate after 50 min (left panel); kinetics of TNAP inactivation by high [CaCl₂], at pH 9.8, reaching steady-state after 10 min (right panel). Activities were measured in Chelex-treated pNPP (1 mM); results represent mean ± SD for 3 identical experiments. (TIF)

S5 Fig. Impact of pNPP concentration on affinity assessment at pH 7.4. Dose-response of generated AP activity (mean mA405nm/min) in steady-state (between 60–90 min) for increasing [MgCl₂] (a) and [CaCl₂] (b) at identical AbM2-bound [Zn²⁺-TNAP]; activities were measured in Chelex-treated pNPP (1 mM or 10 mM as indicated) at pH 7.4. Results represent mean ± SD for 3 identical experiments. (TIF)

S1 File. Data Supplement. Sequence of the primers used for site-directed mutagenesis and presentation of the results and discussion of the data reported in the 5 supplemental figures. (DOCX)

Author Contributions

Conceived and designed the experiments: MFH JLM. Performed the experiments: MFH SVk TKM CS SN. Analyzed the data: MFH JLM. Contributed reagents/materials/analysis tools: TKM CS SN. Wrote the paper: MFH JLM.

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