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Cite this: *Inorg. Chem. Front.*, 2023, 10, 5772Guest exchange in a biomimetic Zn^{II} cavity-complex: kinetic control by a catalytic water, through pore selection, 2nd sphere assistance, and induced-fit processes†N. Nyssen,^{a,b} N. Giraud,^a J. Wouters,^c I. Jabin,^b L. Leherte^{*c} and O. Reinaud^{id} ^{*a}

The kinetics of ligand exchange at a Zn^{II} center confined in a biomimetic environment is studied. The metal ion is bound to a TMPA unit covalently capping the calix[6]arene core. As such, the system is rigidified in a cone conformation open to the solvent at the large rim. At the small rim, the methyl groups of the anisole units lead to steric crowding, thus defining a small pore leading to the metal center. The exchange of an organic guest ligand (MeCN) for another (EtCN) is studied by ¹H NMR spectroscopy. It is shown that it is very slow under dry conditions (several hours at mM concentrations), but much faster in the presence of appreciable amounts of water. Kinetic analyses indicate that the exchange process involves the transient formation of an aqua complex that was independently synthesized and well characterized. Computer studies suggest that transient water coordination first occurs through the small rim pore, in *exo*-position. The water molecule then flips in *endo*-position, as the MeCN guest decoordinates and exits from the cavity through the large rim. This allows another organic guest ligand to get in, with release of the catalytic water to the solvent. Steered molecular dynamics simulations also show the assistance of the second coordination sphere of the metal ion for the *exo/endo* flipping process of the water ligand through H-bonding to the oxygenated small rim of the calixarene macrocycle. Importantly, it appeared that only water can play such a catalytic role, due to size selection by the pores defined by the calixarene small rim. From a biological point of view, it further substantiates the importance of water molecules and micro-environment on metal ion lability.

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Introduction

Zn^{II} coordination in biology is important, displaying structural (e.g. Zn fingers) or functional roles (Zn enzymes).^{1–3} In these biological structures, Zn^{II} is coordinated to 3 or 4 amino-acid residues. In enzymes, an extra site is generally occupied by a water molecule that is the reactant (in hydrolases) or that is

displaced by the reactant (e.g. a phosphate derivative as in alkaline phosphatases⁴ or an alcohol in alcohol dehydrogenase⁵). For all these enzymes, ligand exchange at the metal center is a key step in the catalysis. Ligand exchange is also a key factor that determines the stability of a structural site (inertness), or the Zn homeostasis. These exchange processes are difficult to study within the biological systems. Therefore, small model complexes are currently elaborated and studied to get specific insights into mechanisms at the molecular level.

Poly-aza tripods are classically used to mimic the poly-histidine rich biological sites found in metallo-enzymes. One of them, namely TMPA (Tris(2-PyridylMethyl)Amine), has been long studied for Fe and Cu biomimetic complexes. Less has been reported with Zn. While few examples of TMPA Zn complexes exist in literature, the main area of focus has been the utilization of Zn^{II} complexes in the cleavage of phosphodiester bonds. In those catalytic transformations, the kinetically active form was determined to be the corresponding Zn-aqua complex.⁶ The introduction of hydrogen bond donors has been shown to decrease the pK_a of the bound water molecule and enhance the activity of such complexes, while also favor-

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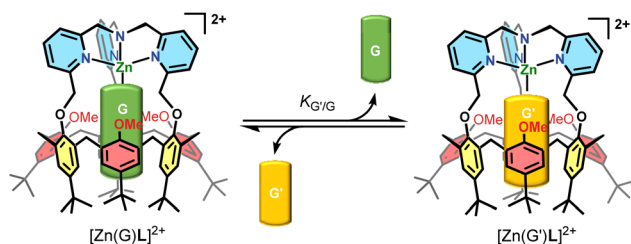
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^bLaboratoire de Chimie Organique, Université Libre de Bruxelles (ULB), Avenue F. D. Roosevelt 50, CP160/06, B-1050 Brussels, Belgium^cLaboratory of Structural Biological Chemistry, Unit of Theoretical and Structural Physical Chemistry, Department of Chemistry, NAMUR Research Institute for Life Sciences (NARILIS), NAMUR Institute of Structured Matter (NISM), NAMUR Medicine & Drug Innovation Center (NAMEDIC), University of Namur, Rue de Bruxelles 61, B-5000 Namur, Belgium†Electronic supplementary information (ESI) available: Synthetic and procedure and characterization of [Zn(H₂O)L](ClO₄)₂, thermodynamic and kinetic measurements of guest exchanges through NMR spectroscopy, computational details. See DOI: <https://doi.org/10.1039/d3qi01271a>

ing the binding of phosphates ions.^{7,8} Indeed, it has long been recognized that the microenvironment around metal ions is a major factor that controls the reactivity/stability of Zn centers. One way to explore the supramolecular aspects relative to this microenvironment is to construct a well-defined and tunable microstructure around the metal ion labile site and evaluate the impact on its properties.⁹ Such a strategy allows to control the second coordination sphere of the metal ion, but also cavity effects related to ligand exchange. Being open to the solvent, the cavity acts not only as a receptor pocket, but also as a pore that will allow or not exogenous molecules to reach the metal center, thereby controlling its reactivity.

Ligands based on the calix[6]arene macrocycle connected to a tripodal nitrogenous core at the small rim are good mimics of the histidine-rich protein cavity.^{10,11} One of them, namely calix[6]TMPA (**L**, Scheme 1), presents a structure that is highly congested although open to the solvent. The calix[6]arene macrocycle is tris-methylated on alternate position of the phenol units, and covalently capped by a TMPA unit through short methylene linkers to the three other phenol units. As such, access to the metal ion bound in the TMPA cap through the small rim pores is highly restricted by steric congestion between the methoxy substituents and the cap methylene linkers.

In a recent study,¹² we reported the synthesis of the corresponding Zn^{II} acetonitrile complex that is 5-coordinate with a guest acetonitrile molecule bound in the calixarene cavity. A thermodynamic study of ligand exchanges monitored by NMR spectroscopy showed that the macrocyclic structure is rigidified in a cone conformation. However, it keeps some flexibility as the three anisole “walls” can rotate along the calix methylene groups for a few degrees, as a function of the size of the included guest ligand. Along our studies, we noticed some surprisingly slow processes: first, coordination of the Zn^{II} dication required heating for hours; second, the exchange of the MeCN guest for EtCN took several hours to reach equilibrium under some experimental conditions. These observations stand in strong contrast with the behavior of simple TMPA metal complexes, devoted of cavity, for which metalation and ligand exchange proceed almost instantaneously. Wanting to know more about these phenomena, we undertook a detailed study relative to the kinetics of exchange between two representative organic guest ligands (**G** and **G'**, Scheme 1), namely MeCN and EtCN, which is the subject of the present article.



Scheme 1 Guest exchange G/G' equilibrium inside the cavity of [Zn(G)L]²⁺.

Results

Characterization of the aqua complex

Complex [Zn(MeCN)L](ClO₄)₂ was synthesized as previously reported, by reacting the calixarene ligand **L** with a stoichiometric amount of the Zn^{II} perchlorate salt in MeCN. Noticeably, the complexation process required heating the solution for several hours. Indeed, protonation of **L** occurs immediately in solution in a first step and complexation necessitates the displacement of the proton by the metal ion in a second slow, but thermodynamically favored, step.¹² The dicationic complex was isolated as the acetonitrile complex and characterized by ¹H NMR spectroscopy. The overall signature attested to a C_{3v} symmetry with the Zn^{II} center bound to the TMPA cap and one molecule of acetonitrile sitting inside the calixarene cavity as attested by its high field δ shift (−0.62 ppm). We noticed however that, when dissolved in wet acetone, a second species was present. The latter did not correspond to the protonated ligand and vanished upon addition of small amounts of MeCN. Suspecting that it could correspond to the aqua complex, we repeated the complexation process with a 1 : 1 mixture of ligand **L** and Zn^{II} salt in a non (/poorly)-coordinating solvent, namely acetone. The complexation also required heating the solution for several hours. The isolated complex was characterized by various spectroscopies. The IR spectrum confirmed the presence of two perchlorates as counterions (see the ESI†). The ¹H NMR spectrum (Fig. 1, bottom-blue) is typical of a C_{3v} symmetrical structure with a cone conformation for the calix[6]arene macrocycle. As for the nitrile complex, the pyridyl protons are low field shifted compared to the free ligand, thus attesting to the coordination of the Lewis acidic metal ion Zn^{II}. The two sets of aromatic H_{Ar} and *t*Bu protons display closer δ shifts for the aqua complex ($\Delta\delta \sim 0.2$ ppm) than for the MeCN complex (~ 0.7 ppm). This suggests an average straighter conformation for the calixarene host, consistently with the smaller size of the guest water molecule. The coordinated water molecule could be identified under relatively dry conditions, through two signals at 8.28 and 8.46 ppm, each one integrating for one proton. These two signals vanished upon D₂O addition. Saturation transfer experiments with free water and NOESY measurements further confirmed their attribution (see the ESI†). These low-field shifted δ values indicate that the water molecule is bound to the Lewis acidic Zn^{II} center and sits at the level of the small rim of the calixarene structure, away from the anisotropic cone of the aromatic walls. It is noteworthy that each proton of the coordinated water molecule is differentiated from the other. This reflexes strong electrostatic interaction with the oxygen atoms belonging to the calixarene small rim that slows down the rotation of the water molecule inside the structure. The differentiation of the two protons can be due to a stronger interaction of one of the water protons with a calixarene oxygen atom. Addition of an aliquot of MeCN to this aqua complex led to the appearance of a high-field peak at −0.62 ppm corresponding to the MeCN ligand included in the cavity (Fig. 1, top-purple), thus attesting to the formation of the MeCN complex. The exchange was slow at the ¹H NMR shift time scale, but fast at the experi-



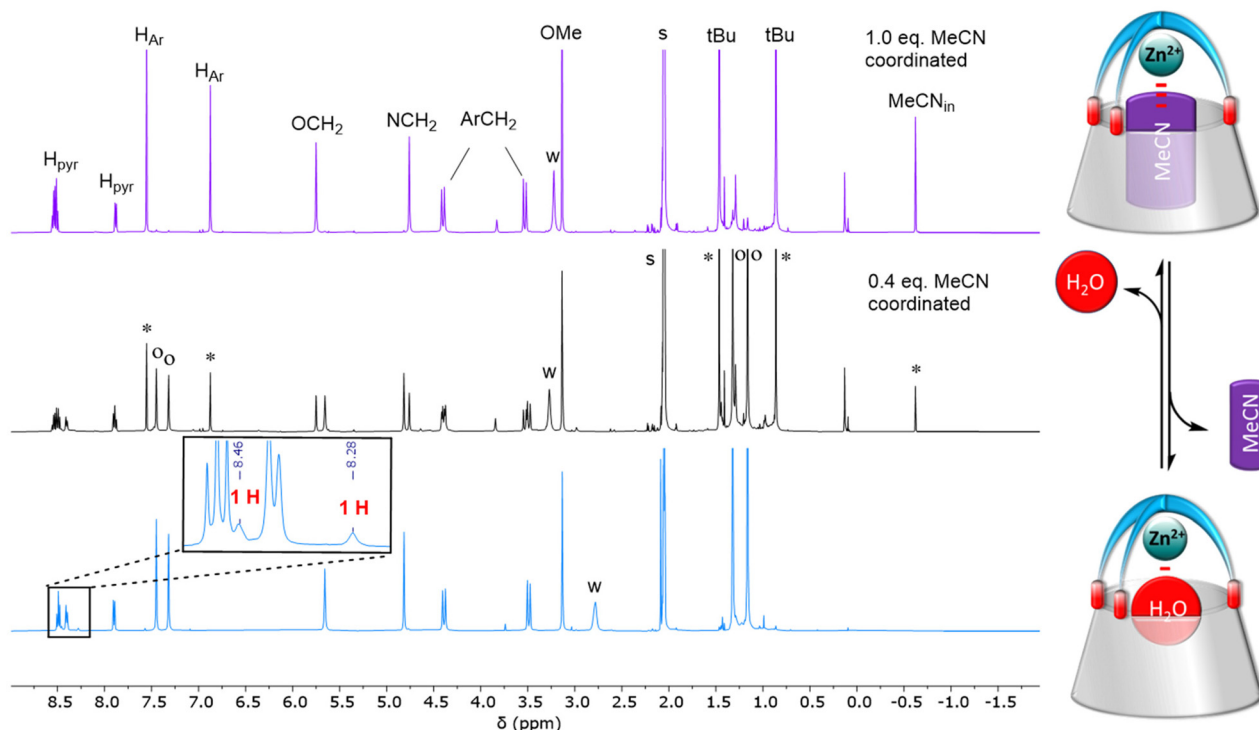


Fig. 1 ^1H NMR spectra (300 K, 500 MHz) of $[\text{Zn}(\text{H}_2\text{O})\text{L}](\text{ClO}_4)_2$ dissolved in acetone- d_6 before (bottom-blue) and after the addition of MeCN (ca. 0.5 and 1 equiv. middle-black and top-purple, respectively). Inset: Zoom on the low-field region to show the peaks associated to the water ligand. The spectra where a nitrile was added were recorded immediately after the addition. o: aqua complex, * MeCN complex.

mental time scale. A careful titration of the aqua complex by acetonitrile (see the ESI†) confirmed that one water molecule is coordinated to the Zn^{II} ion, and no extra water guest is present, either in the cavity or in *exo*-position.

Kinetic study of the acetonitrile/propionitrile exchange

The exchange between two nitriles was followed by ^1H NMR spectroscopy by recording multiple spectra over the course of several hours of a solution of one nitrile complex in presence of a known amount of the other nitrile in deuterated acetone. A representative example is shown in Fig. 2. The amount of each complex was calculated by integrating the two signatures in slow exchange on the NMR shift time scale (ESI†). However, we noticed that the rate of exchange varied not only with the amount of MeCN or EtCN, but also with the amount of water: under “dry” conditions, it was slow, and under “wet” conditions it was much faster. In light of the ability of water to be complexed to the metal center, we hypothesized that the exchange mechanism involves its transient coordination (pre-equilibrium displayed in Scheme 2). Hence, starting from the MeCN complex dissolved in acetone- d_6 , we followed by ^1H NMR spectroscopy, as a function of time, the exchange process of the MeCN guest for EtCN, varying three factors: $[\text{MeCN}]$, $[\text{EtCN}]$, and $[\text{H}_2\text{O}]$. Table 1 summarizes the conditions of 5 different experiments A–E (the detailed quantifications are reported in the ESI†). Qualitatively, these results show that the rate of exchange increases when the concentration of MeCN

decreases (experiments A–C) and when the concentration of EtCN and water increases (experiments C–E).

Consistently, the more the pre-equilibrium is displaced to the right, towards the formation of the aqua complex, the faster the formation of the EtCN complex (Scheme 2). The rate of exchange of the acetonitrile guest for the propionitrile guest was expressed in eqn (1) according to the hypothesis made in Scheme 2. Importantly, in none of the experiments reported in Table 1, the aqua complex was detected in the ^1H NMR spectra. This is obviously related to the large relative affinities of nitrile guests that occupies the calixarene cavity compared to water that is too small and lies above. Indeed, from titration experiments, we found that $K_{\text{MeCN}/\text{H}_2\text{O}} = 200 \pm 12$ (see the ESI†), and knowing that $K_{\text{EtCN}/\text{MeCN}} = 30 \pm 1$,¹² we can deduce that $K_{\text{EtCN}/\text{H}_2\text{O}} = 6000 \pm 540$. The kinetic data measured in experiments A, B, C (Table 1 and Fig. 3) were then analyzed according to eqn (1)–(4) obtained by applying the steady state approximation for aqua intermediate concentration:

$$\nu = k_2[\text{w}_{\text{in}}][\text{EtCN}_{\text{out}}] = \frac{k_2K_1[\text{w}_0][\text{EtCN}_{\text{out}}][\text{MeCN}_{\text{in}}]}{[\text{MeCN}_{\text{out}}]}. \quad (1)$$

Setting that $x = [\text{MeCN}_{\text{in}}]$ and $c = [\text{total calixarene}]$, we obtain:

$$-\frac{dx}{dt} = k_2K_1[\text{w}_0] \frac{([\text{EtCN}_0] - c + x)x}{[\text{MeCN}_0] - x} \quad (2)$$



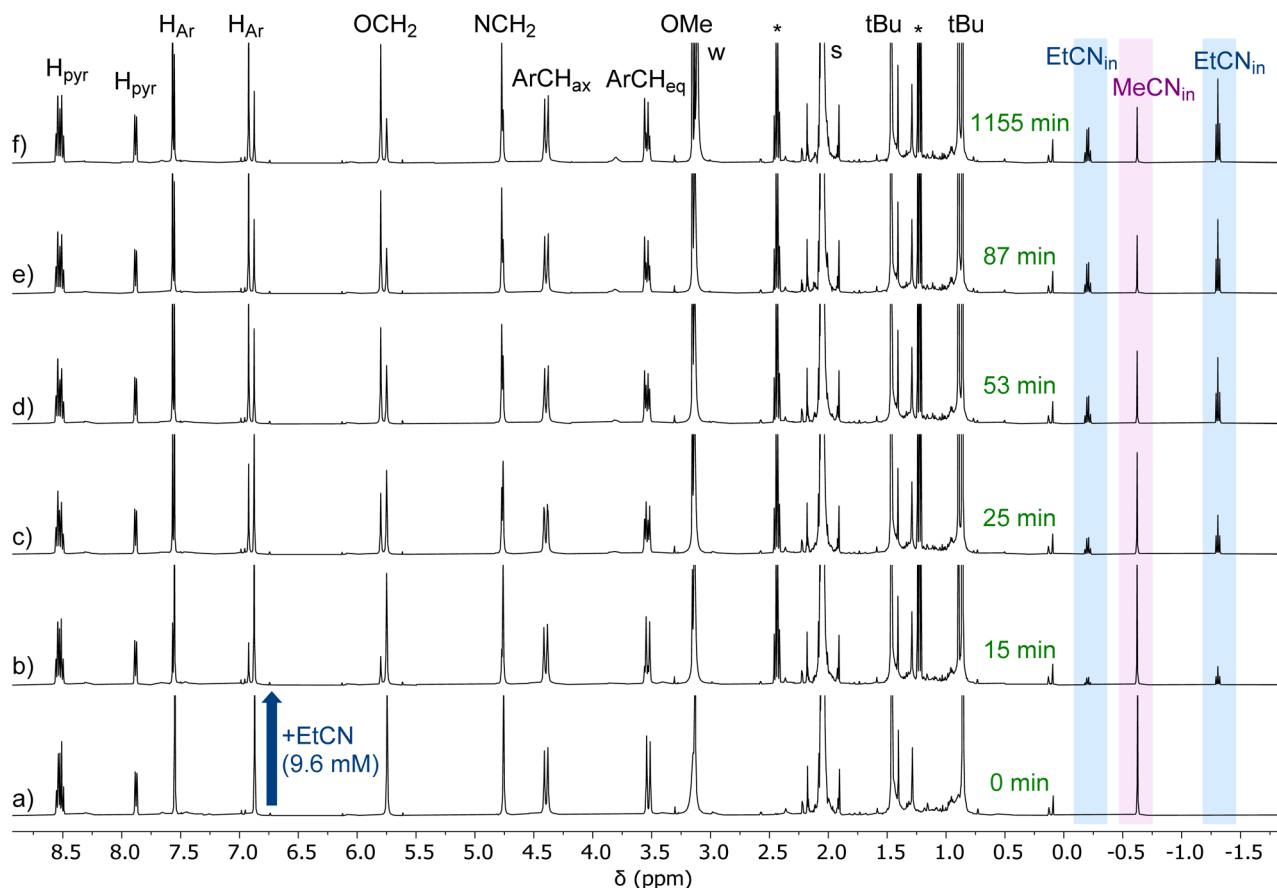
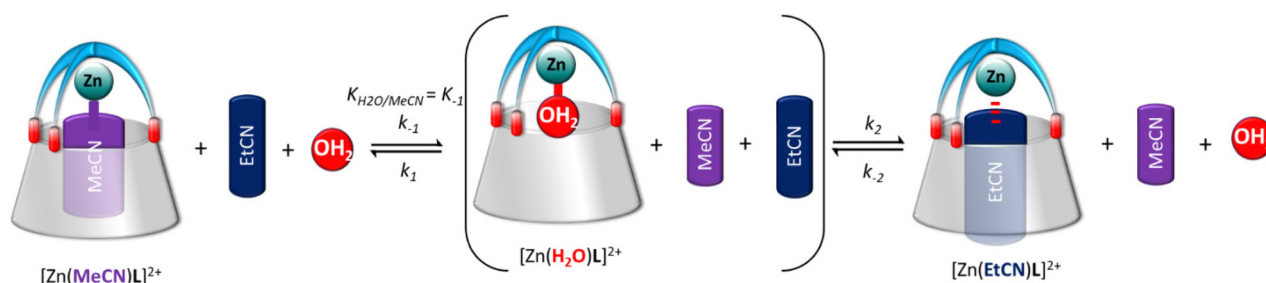


Fig. 2 ^1H NMR spectra (300 K, 500 MHz, acetone- d_6) recorded during the titration of $G = \text{MeCN}$ by $G' = \text{EtCN}$. Initial conditions in (a) $[\text{Zn}(\text{MeCN})\text{L}]^{2+} = 3.7 \text{ mM}$, $[\text{free MeCN}] = 92 \text{ mM}$, $[\text{EtCN}] = 0 \text{ mM}$, $[\text{H}_2\text{O}] = 20 \text{ mM}$. Percentage of EtCN complex in solution: (a) 0%, (b) 17%, (c) 38%, (d) 41%, (e) 59%, (f) 71%. This experiment is reported as entry A in Table 1. Full conversion corresponds to the amount of EtCN complex formed at equilibrium under these experimental (71%). The “ t (50% conversion)” is then the time at which we have 35.5% of EtCN complex in solution. *: free EtCN.



Scheme 2 Proposed pathway for the guest exchange involving the transient formation of the aqua complex.

and thus:

$$-\frac{1}{K_1 \frac{([\text{EtCN}]_0 - c + x)x}{[\text{MeCN}]_0 - x}} dx = k_2 [w_0] dt \quad (3)$$

which gives, after integration:

$$F(x) = \frac{[\text{MeCN}]_0 \ln(x) - ([\text{EtCN}]_0 + [\text{MeCN}]_0 - c) \ln([\text{EtCN}]_0 - c + x)}{K_1([\text{EtCN}]_0 - c)} = -k_2 [w_0] t \quad (4)$$

(the subscript “0” defines “total amount of”; $[w_{\text{out}}] \approx [w_0]$, “in” and “out” refer to the calixarene cavity).

Fig. 3 shows full agreement with the working hypothesis as the data recorded for the 3 different experiments show a linear relationship in agreement with eqn (4). The exchange kinetic constant is thus: $k_2 = 0.090 \pm 0.004 \text{ mM}^{-2} \text{ min}^{-1}$. From this value, we can deduce that $k_{-2} = 0.015 \pm 0.002 \times 10^{-3} \text{ mM}^{-2} \text{ min}^{-1}$.

NOESY measurement

As the MeCN/ H_2O exchange process is much faster than that of MeCN/EtCN, we have chosen to monitor it by NMR using



Table 1 Variation of the guest exchange rate monitored by ^1H NMR analyses of a solution initially containing complex $[\text{Zn}(\text{MeCN})\text{L}]^{2+}$, as a function of the initial content in complex, MeCN, EtCN and water (5 independent experiments referred as A–E and used for detailed kinetic analyses, see below)

Experiment #	$[\text{Zn}(\text{MeCN})\text{L}]^{2+}$ (mM)	$[\text{EtCN}]$ (mM)	$[\text{MeCN}]$ (mM)	$[\text{Water}]$ (mM)	t (50% conversion) (min)
A	3.7	10	96	20	23
B	2.0	10	36	16	19
C	3.1	9	15	15	7
D	2.2	66	14	18	$<5^a$
E	2.8	70	14	129	$\ll 5^b$

^a 86% conversion was already achieved at the first measurement after 5 min. ^b The first measurement after 5 min already showed full conversion. Full conversion refers to the quantity of $[\text{Zn}(\text{EtCN})\text{L}]^{2+}$ at equilibrium.

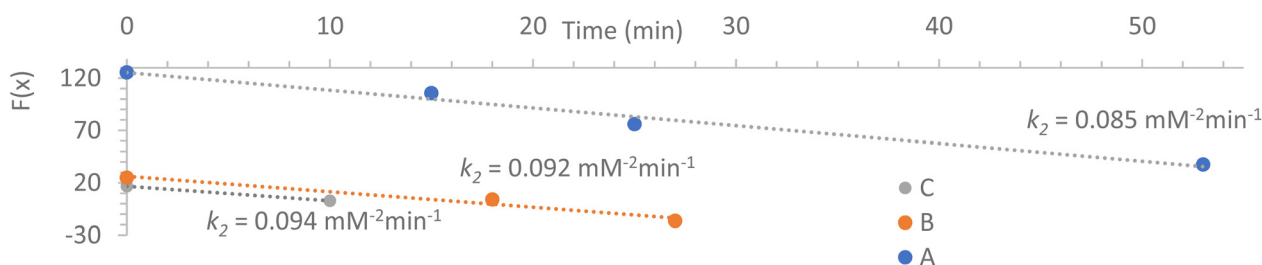
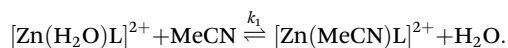


Fig. 3 Kinetic fit of the ^1H NMR analyses. For the expression of $F(x)$, see (eqn (4)). A, B, C refers to experiments of Table 1.

EXchange Spectroscopy (EXSY). For this purpose, a solution containing a mixture of the MeCN and aqua complexes was prepared, and we have recorded a ^1H – ^1H 2D NMR NOESY spectrum on this sample. On the resulting data, we clearly observe several sets of correlations for the complex, revealing a slow exchange on the NMR shifts time scale between the MeCN and the aqua form. Fig. 4 shows the region of the resulting 2D spectrum where this correlation pattern is observed. We have thus recorded a series of EXSY spectra, varying the mixing time to monitor this slow exchange process (a representative 2D map acquired at 600 MHz is described in the ESI†). The volume of the resolved correlation patterns was measured for each mixing time value. The resulting build-up curves that were obtained for the selected diagonal and cross peaks arising from the exchange process between the aqua and MeCN complexes can be modelled using modified Bloch equations accounting for proton longitudinal relaxation and the formal equilibrium:



The analysis of the NOESY correlations involving (i) an aromatic proton from the calixarene ligand, and (ii) the methyl protons from MeCN, both undergoing a chemical exchange between the aqua and MeCN complexes, yielded k_1 values from 2.2 to 11 $\text{min}^{-1} \text{mM}^{-2}$ and $k_{-1} = 0.02 \pm 0.02 \text{ min}^{-1} \text{mM}^{-2}$. This result yields a calculated value for $K_{\text{MeCN}/\text{H}_2\text{O}} = k_1/k_{-1} = 110$ to 500, in good agreement with the value of 200 measured by titration (see above and Scheme 2). These kinetic data, when compared to those measured for the EtCN/ H_2O exchange ($k_2 = 0.090 \text{ mM}^{-2} \text{ min}^{-1}$ and $k_{-2} = 0.015 \times 10^{-3} \text{ mM}^{-2} \text{ min}^{-1}$),

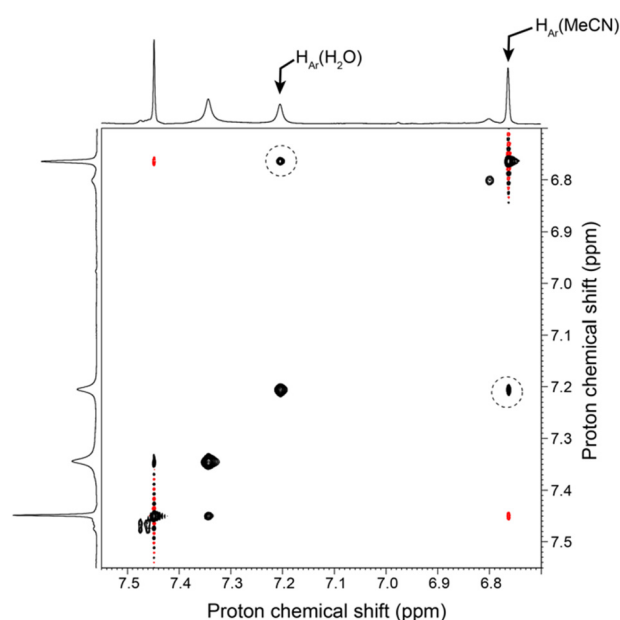


Fig. 4 Selected region of a 2D ^1H – ^1H NOESY spectrum (500 MHz) of a solution containing a mixture of the aqua and MeCN Zn complex in acetone- d_6 . This spectrum was recorded with a mixing time of 1.5 s. The sample was prepared with 0.9 eq. of MeCN and 200 eq. of H_2O . Correlations reflecting the exchange process between MeCN and H_2O in the cavity of the Zn complex are highlighted with a dotted circle. Other regions of interest of this 2D map are shown together with the 1D ^1H NMR spectrum in ESI.†

indicate that both, entrance and exit of the nitrilo guest are three orders of magnitude faster for MeCN than for EtCN. Hence, the rates are obviously related to the size of the guest.



This corresponds to the energy cost to open the small rim door constituted by the 3 *t*Bu substituents of the most rigid aromatic units, namely those connected to the TMPA cap. Such a kinetic differentiation is reminiscent of a previous study reported for a Cu^I complex obtained with a calix[6]arene functionalized by three α -methylpyridyl groups. Indeed, it was shown that part of the activation energy for the exit of the acetonitrile guest stems from the *t*Bu door into which the guest “bump” during the dissociative process.¹³

In order to clarify the role of the water molecule and the reason for which the aqua-complex is involved as a necessary intermediate for the exchange between two nitrile guests, we have carried out molecular computing studies. The idea was to start with the acetonitrile complex and identify the pathway through which the water molecule becomes in interaction with the Zn^{II} center, whereas the MeCN guest leaves the cavity.

Steered molecular dynamics simulation of the guest-water exchange

To favor the guest-water exchange in the calixarene structure, Molecular Dynamics (MD) and Steered MD (SMD) simulations of a calixarene–MeCN complex interacting with a water molecule were carried out in vacuum at 300 K using the program GROMACS2020,¹⁴ as detailed in the ESI†. Water and the small molecules were modeled using the TIP3P and CHARMM27 force fields, respectively. The simulations were carried out with four atom position restraints placed on the N atoms of the calixarene cap, while restraining the four Zn–N_{cap} distances to avoid exaggerated displacement of the Zn²⁺ ion during the approach and the insertion of water inside the calixarene cavity. The system was first optimized using a steepest descent algorithm (Fig. 5, structure A). Then, the MD total equilibration stage duration was set to 2100 ps. At the end of this stage, the water molecule more closely interacts with Zn^{II} (Fig. 5, structure B). Afterwards, the SMD stage was carried out for a period of 60 ps. A pulling force was applied to the center of mass of MeCN to progressively evacuate MeCN from the calixarene cavity. As illustrated in the ESI†, the time-dependence of the water interaction energy with its neighborhood allows to model the substitution of MeCN as a stepwise process.

Step 1: coordination of water. At the end of the preliminary optimization stage, the total intermolecular interaction energy (IIE) value of the molecule of water (defined as the sum of IIEs with the calixarene, Zn, and MeCN) is equal to -40 kJ mol^{-1} . Starting from the optimized structure (Fig. 5, structure A), the water molecule evolves around the calixarene until it reaches a stabilized position facing the Zn²⁺ ion, at 1701.0 ps (Fig. 5, structure B). It is hydrogen-bonded to an O_{methoxy} atom of the calixarene structure (see the ESI†) and a deformation of the calixarene cap is observed. The interaction energy value drastically decreases from a value of -54 kJ mol^{-1} at $t = 1700.0 \text{ ps}$ to a value of -142 kJ mol^{-1} at $t = 1701.0 \text{ ps}$, and the O_{water}–Zn distance is reduced to a value of about 2 Å (see the ESI†). This gives rise to a stable 6-coordinate intermediate presenting a pseudo-octahedral environment due to the simultaneous

coordination of water and MeCN. Such an intermediate is observed until the end of the equilibration stage, *i.e.*, $t = 2100.0 \text{ ps}$.

Step 2: substitution of MeCN by water. Snapshots at various times during the insertion procedure (60 ps) are displayed in Fig. 5 (structures I1 to I4). When the pulling force is applied to the MeCN ligand, the substitution of MeCN by H₂O does not immediately take place. Indeed, the water molecule position and orientation remain stable during 29.7 ps. H₂O closely approaches the calixarene cavity after 29.8 ps of the pulling stage (Fig. 5, structure I1). At that moment, the water molecule is hydrogen-bonded to a methoxy group of the calixarene structure (Fig. 5 inset, and ESI†) and its second H atom is oriented towards the N_{MeCN} atom at a distance of 1.91 Å. Also, the water interaction energy value has decreased to a value of -173 kJ mol^{-1} . The O_{water}–Zn distance remains slightly larger than 2 Å (ESI†), while the N_{MeCN}–Zn distance begins to increase (ESI†). Shortly after, at $t = 29.9 \text{ ps}$ when MeCN is farther from Zn, the water molecule begins its insertion into the calixarene (Fig. 5, structure I2). Its interaction energy value is -192 kJ mol^{-1} . At $t = 30.0 \text{ ps}$ (Fig. 5, structure I3), the water interaction energy reaches a value of -231 kJ mol^{-1} .

Step 3: relaxation of the calixarene 3D structure. Starting from the coordination step (step #1), H₂O remains hydrogen-bonded to the calixarene structure (Fig. 5 inset, and ESI†). At $t = 30.0 \text{ ps}$, the deformation of the calixarene cap observed at step #1 is still visible, but the C₃ symmetry is recovered at around 42.0 ps (see the ESI†). Then, due to the folding of a *t*Bu group inside the calixarene, the C₃ symmetry is preserved only at the level of calixarene cap, as illustrated at $t = 60.0 \text{ ps}$ of the SMD trajectory (Fig. 5, structure I4 and ESI†) where the two shortest H_{water}–O_{methoxy} distances are equal to 2.64 and 2.94 Å. A slight relaxation is observed in the mean Zn–N_{calix} distance profile with an overall decrease of 0.003 nm (see the ESI†) and no large energy changes are observed. At that moment, the water interaction energy value is equal to -236 kJ mol^{-1} . Hence, after substitution of MeCN by H₂O around 43 ps of the SMD trajectory, the calixarene structure undergoes a folding of one *t*Bu group towards the calixarene cavity, which coincides with a decrease in the H₂O and *t*Bu interaction energy values (ESI† and Fig. 5 inset, respectively). To confirm the likelihood of such a fold, the structure I4 was submitted to an optimization stage at the MP7 semi-empirical level using the program MOPAC¹⁵ which corroborates the SMD results, in agreement with the NMR observations (Fig. 5, structure C). Particularly, in the final optimized structure, the water molecule reorients to form a H-bond with one of the calixarene methoxy groups. Such a self-inclusion process of a *t*Bu substituent in the cavity was previously observed by ¹H NMR spectroscopy at low *T* with a closely related calix[6]arene-based Cu^I funnel complex binding carbon monoxide.¹⁶ With this small guest ligand, a “3-step valve” was evidenced between three *t*Bu substituents going alternatively in “in” and “out” positions. When fast at the NMR shift time scale (*i.e.* at RT), this led to a pseudo C_{3v} signature with an apparent straight conformation, exactly as observed here for the aqua-complex.



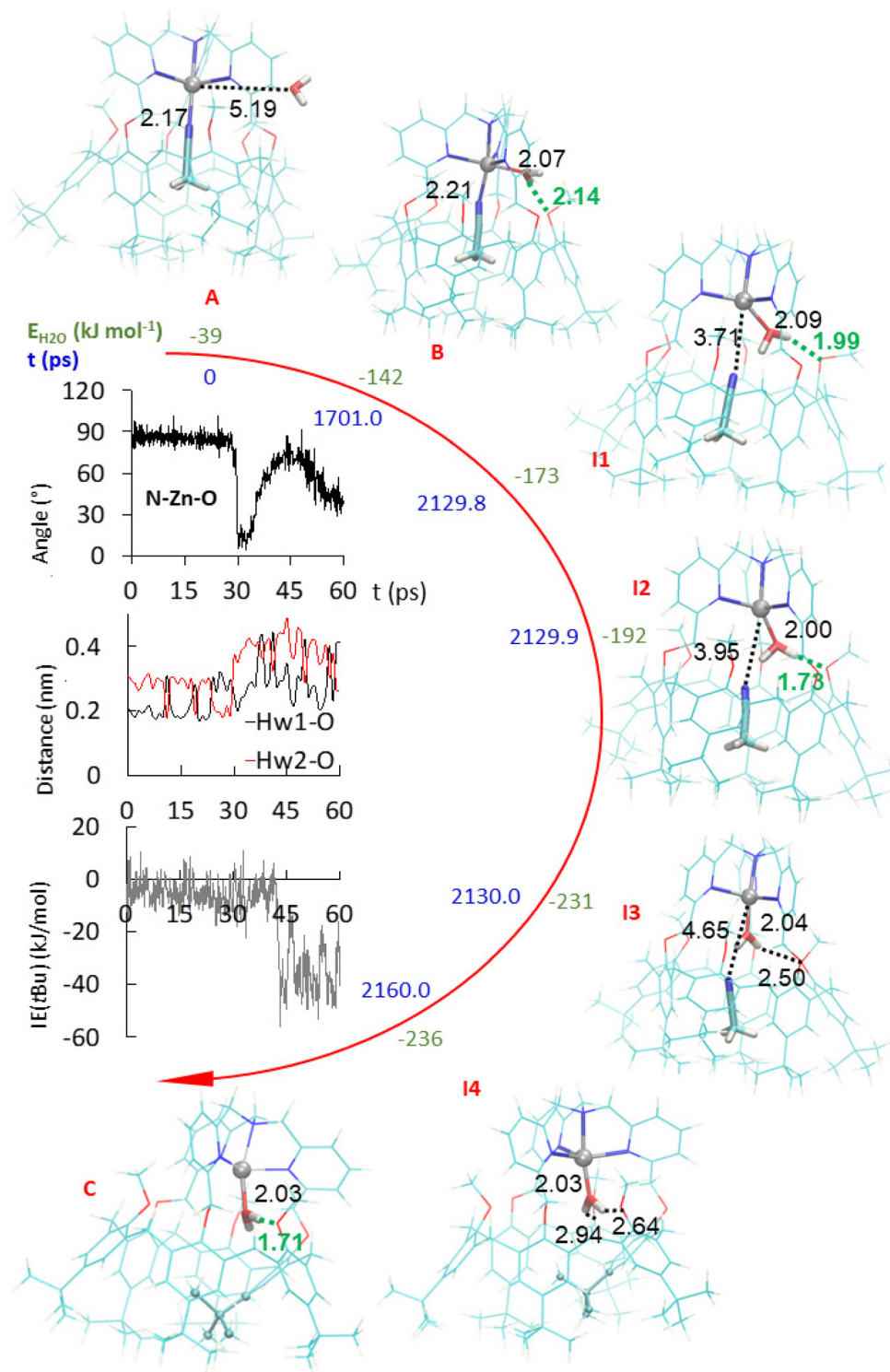


Fig. 5 Stepwise substitution of MeCN guest by a water molecule following an associative pathway, as obtained from the 2160 ps MD and SMD trajectories in vacuum at 300 K. Initial optimized structure – A, snapshot at 1701 ps of the MD trajectory (coordination of H₂O – B), at $t = 29.8$ of the SMD trajectory (insertion stage – I1), 29.9 ps (insertion stage – I2), 30.0 ps (insertion stage – I3), and 60.0 ps (final structure – I4). Semi-empirical MP7 optimized structure (C). Dotted lines refer to selected interatomic distances (in Å). Green dotted lines depict hydrogen bonds. The folded tBu moiety is displayed using ball and stick representations in I4 and C. Inset from top to bottom (obtained from the 60 ps long SMD trajectory): variation of the NMeCN-Zn^{II}-O_{water} angle and H_{water}...OMe distances; tBu-calixarene short-range interaction energy profile.



To support the observation, the 60 ps long SMD simulation was continued by a 4000 ps conventional MD run using initial random atom velocities. Unfolding-refolding events were observed with exchange between the *t*Bu moieties until $t = 1576.3$ ps (see the ESI†). Then, *t*Bu exchange stops and the last folded conformation remains present until the end of the simulation. In this structure, a strong H-bond connects regularly one of the hydrogen atoms of the water guest to an anisole unit during 4% of the time, a percentage equally shared by both H atoms of the water molecule.

The computer simulations were also conducted with acetone as a solvent instead of vacuum and similar results were obtained (see the ESI†). The water molecule first gets close to the calixarene structure and deforms it while it coordinates Zn^{II} . In the solvated case however, H_2O is first hydrogen bonded to two acetone molecules. Then, similarly to the vacuum case, it forms a hydrogen bond with a methoxy group of the calixarene. The simulations were also performed with EtCN as an initial guest instead of MeCN, and same results were obtained for the substitution process. Finally, when the water molecule initially out of the cavity is replaced by MeCN, close interaction with the Zn center in *exo*-position was observed but at a distance that is much too long to correspond to a coordination bond. Consistently, the interaction energy of the *exo*-MeCN molecule is not significantly reduced compared to other positions. It remains very mobile and, after a while, tends to be trapped at the level of the cap of the calixarene. Similar results were obtained with MeOH replacing H_2O (see the ESI†). Coordination of the MeCN and MeOH molecules by Zn^{II} are then triggered by removing the *endo*-ligand MeCN during the whole MD simulations. It likely favors the strong deformation of the calixarene structure, but it does not initiate the insertion of the molecule that stays outside the calixarene cavity. Nevertheless, the insertion of MeOH can be artificially produced by the application of a steering force to pull the ligand inside the calixarene through the small rim. Such a SMD simulation was carried out using the same methodology as used for pulling the *endo*-ligand outside the cavity. Interestingly, MeCN does not enter the calixarene using that approach. Hence, water is the only ligand to spontaneously enter the calixarene cavity through the small rim: it is small enough to flip from the *exo*-position to the *endo*-position through the small pore existing at the calixarene small rim, whereas MeCN and MeOH appear to be either too large or not energetically stabilized at the level of the small pore. This suggests that the only (or most probable) pathway for an organic ligand to coordinate the metal center in *endo*-position, inside the cavity, is to go through the large rim of the calix-funnel. This inclusion process is, likely for a steric reason, impossible when the cavity is already occupied by an organic guest, whereas a water guest can get in from the small rim.

In summary, the simulations show that the preliminary step for a ligand to enter the cavity through the calixarene small rim is to be immobilized strongly enough, through a coordination bond to the metal center to the calixarene structure. Without such binding, the ligand remains free to move

around the calixarene structure. Then, if the *exo*-ligand is small enough, it can flip into the cavity when the guest ligand leaves, thereby avoiding the formation of an “empty cavity” intermediate during guest exchange. Hence, water seems very unique as it can bind to Zn^{II} in *exo*-position and flip into the cavity in *endo*-position, thus allowing the guest-Zn bond cleavage and the exit of the guest, and conversely. Quite interestingly, and subtly, the key flipping process involves transient assistance of an oxygen atom present at the small rim of the calixarene acting as H-bond acceptor in the second coordination sphere of the metal ion.

Discussion

It may appear surprising that a simple dissociative mechanism for guest-exchange is not at work in this system. Indeed, direct displacement of the guest would leave a 4-coordinate intermediate, which should not be so unstable. One explanation could rely on the rigidity of the system. Indeed, the capping TMPA unit of ligand **L** constrains the calixarene macrocycle in a cone conformation with aromatic units lying alternatively in in/out position relative to the cavity. This alternate cone conformation is the opposite to the one adopted by the more flexible ligands: three imidazole arms¹⁷ or a TREN (tris(2-aminoethyl)amine) unit.¹⁸ The relatively straight conformation adopted by calixarene **L** is explained by the fact that the aromatic units connected to the cap are obliged to sit in a position that projects their oxygen atom toward the outside (Fig. 6). Consequently, the three other aromatic units, bearing methoxy groups, are constrained to orient their oxygen atom toward the inside. This creates a high steric hindrance that defines the pore for *exo*-binding and allows 2nd sphere assistance for water *exo*-binding and *endo*-flipping. Importantly also, the rigidified calix core of $\text{Zn}^{\text{II}}\text{L}$ restricts the induced-fit process observed with the other systems that can adopt a flattened conformation, thus shrinking the cavity space in the absence of guest. As a result, the energy cost for leaving a fully empty cavity is probably much higher with **L**; it explains the fact that the water ligand flips into the cavity whereas the guest leaves. The resulting new 5-coordinate intermediate then undergoes partial insertion of one of its *t*Bu substituent present at the large rim in order to optimize the cavity filling. This leads to a fully stable aqua complex that could be independently synthesized and fully characterized.

Finally, it is worth noting that the *t*Bu group that is partially included in the cavity space of the aqua complex constitutes a gate that needs to be open for guest leaving and guest entering. These in/out phenomena may explain the impressive difference of rate exchange for the MeCN and EtCN guests in both directions. The corresponding difference in activation energies (*ca.* 3 orders of magnitude $\sim 17 \text{ kJ mol}^{-1}$) may correspond to the interaction energy difference between the *t*Bu gate and the ethyl *vs.* methyl group of the guest during the leaving and entering processes. Interestingly, such an inertness under dry conditions was previously observed¹⁹ with the Cu com-



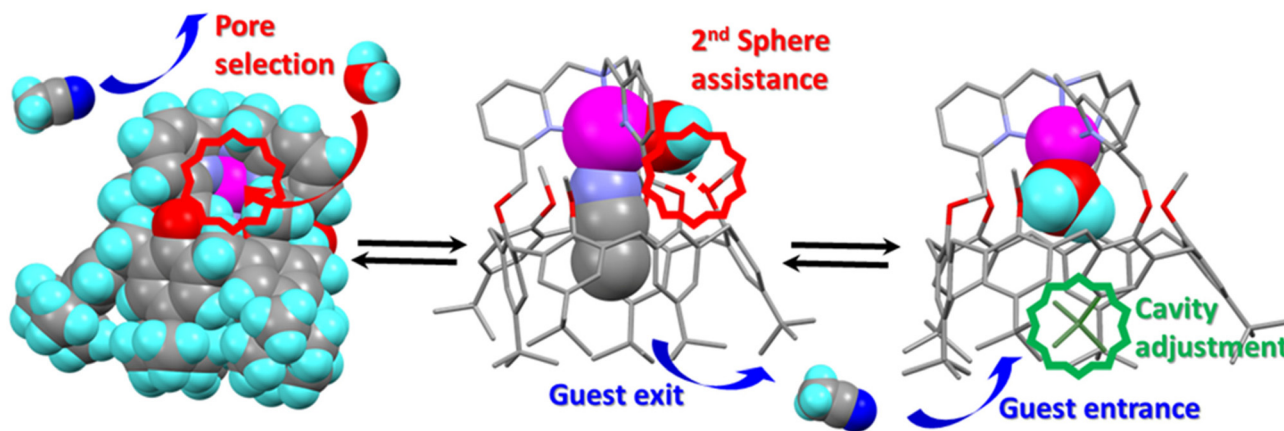


Fig. 6 Key steps for the guest exchange catalyzed by a water molecule evidencing: (left) pore-size selection for the *exo*-access to the metal center; (center) H-bonding assistance for the water ligand flipping from an *exo* to an *endo* position; (right) *t*Bu partial insertion for filling the cavity space.

plexes of the very same ligand **L** and it was shown that water guests regulate its redox activity.²⁰ In the present study, we evidence the role of water for ligand exchange, a role that was not elucidated before.

Conclusion

In conclusion, we have described a very special system where the Zn^{II} cation is embedded in a nitrogenous cap connected to a cavity open to the solvent, for which guest ligand exchange is under the control of a single water molecule. The exchange process involves a first associative step where a water molecule binds the Zn center in *exo*-position, thus leading to an octahedral intermediate. The second step is the intramolecular substitution of the *endo*-bound guest (**G**) by the *exo*-bound water to give rise to a 5-coordinate aqua Zn complex with concomitant expulsion of the initial guest (**G**). In virtue of the micro-reversibility principle, the substitution of the *endo*-bound aqua ligand by a new organic guest (**G'**) follows the same pathway, with entrance of the new guest (**G'**) at the large rim, and expulsion of the water molecule by the side, at the small rim. Hence, the water molecule acts as a catalyst and is very unique in its catalytic role: it first binds to the metal center in an *exo*-position, and flips into the pore at the calixarene small rim with the assistance of an oxygen atom of the host structure acting as H-bond acceptor. Other studied molecules are too large or do not, under the selected calculation conditions, spontaneously initiate the insertion step through coordination to Zn^{II} , which is required before insertion under the current calculation conditions. As a spectacular result, **G** exchanges for **G'** is fast only in the presence of water, and water plays the role of a very unique catalyst that controls the kinetics of guest exchange at the metal center thanks to a pore selection and 2nd coordination sphere assistance.

All in all, this study represents the second case ever reported in the literature of well-characterized molecular metal complex for which ligand exchange is under the unique

control of a single water molecule.¹⁷ Here, the control relies on pore selection, 2nd sphere assistance through transient H-bonding, and host-guest induced fit. From a biological point of view, it further substantiates the importance of water molecules on metal ion lability, a property that is key for the stability of structural Zn sites in proteins, for Zn-enzyme activity, as well as for homeostasis.

Conflicts of interest

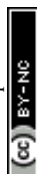
There are no conflicts to declare.

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Notes and references

- W. N. Lipscomb and N. Sträter, Recent Advances in Zinc Enzymology, *Chem. Rev.*, 1996, **96**, 2375–2434.
- W. Maret and Y. Li, Coordination Dynamics of Zinc in Proteins, *Chem. Rev.*, 2009, **109**, 4682–4707.
- A. Krężel and W. Maret, The biological inorganic chemistry of zinc ions, *Arch. Biochem. Biophys.*, 2016, **611**, 3–19.



- 4 B. Stec, K. M. Holtz and E. R. Kantrowitz, A revised mechanism for the alkaline phosphatase reaction involving three metal ions, *J. Mol. Biol.*, 2000, **299**, 1303–1311.
- 5 B. V. Plapp, B. R. Savarimuthu, D. J. Ferraro, J. K. Rubach, E. N. Brown and S. Ramaswamy, Horse Liver Alcohol Dehydrogenase: Zinc Coordination and Catalysis, *Biochemistry*, 2017, **56**, 3632–3646.
- 6 G. Feng, J. C. Mareque-Rivas, R. T. M. de Rosales and N. H. Williams, A Highly Reactive Mononuclear Zn(II) Complex for Phosphodiester Cleavage, *J. Am. Chem. Soc.*, 2005, **127**, 13470–13471.
- 7 J. C. Mareque-Rivas, R. Prabakaran and R. T. M. de Rosales, Relative importance of hydrogen bonding and coordinating groups in modulating the zinc–water acidity, *Chem. Commun.*, 2004, 76–77.
- 8 J. C. Mareque-Rivas, R. T. M. de Rosales and S. Parsons, The affinity of phosphates to zinc(II) complexes can be increased with hydrogen bond donors, *Chem. Commun.*, 2004, 610–611.
- 9 R. Gramage-Doria, D. Armspach and D. Matt, Metallated cavitands (calixarenes, resorcinarenes, cyclodextrins) with internal coordination sites, *Coord. Chem. Rev.*, 2013, **257**, 776–816.
- 10 J.-N. Rebilly, B. Colasson, O. Bistri, D. Over and O. Reinaud, Biomimetic cavity-based metal complexes, *Chem. Soc. Rev.*, 2015, **44**, 467–489.
- 11 D. Coquière, S. Le Gac, U. Darbost, O. Sénèque, I. Jabin and O. Reinaud, Biomimetic and self-assembled calix[6]arene-based receptors for neutral molecules, *Org. Biomol. Chem.*, 2009, **7**, 2485–2500.
- 12 P. Aoun, N. Nyssen, S. Richard, F. Zhurkin, I. Jabin, B. Colasson and O. Reinaud, Selective Metal-ion Complexation of a Biomimetic Calix[6]arene Funnel Cavity Functionalized with Phenol or Quinone, *Chem. – Eur. J.*, 2023, **29**, e202202934.
- 13 Y. Rondelez, A. Duprat and O. Reinaud, Calix[6]arene-Based Cuprous “Funnel Complexes”: A Mimic for the Substrate Access Channel to Metalloenzyme Active Sites, *J. Am. Chem. Soc.*, 2002, **124**, 1334–1340.
- 14 M. J. Abraham, T. Murtola, R. Schulz, S. Páll, J. C. Smith, B. Hess and E. Lindahl, GROMACS: High performance molecular simulations through multi-level parallelism from laptops to supercomputers, *SoftwareX*, 2015, **1–2**, 19–25.
- 15 J. J. P. Stewart, *Stewart Computational Chemistry*, Colorado Springs, CO, USA, 2016, <https://OpenMOPAC.net>, last accessed 20 Dec. 2022.
- 16 Y. Rondelez, O. Sénèque, M.-N. Rager, A. F. Duprat and O. Reinaud, Biomimetic Copper(I)–CO Complexes: A Structural and Dynamic Study of a Calix[6]arene-Based Supramolecular System, *Chem. – Eur. J.*, 2000, **6**, 4218–4226.
- 17 E. Brunetti, L. Marcellis, F. E. Zhurkin, M. Luhmer, I. Jabin, O. Reinaud and K. Bartik, A Water Molecule Triggers Guest Exchange at a Mono-Zinc Centre Confined in a Biomimetic Calixarene Pocket: a Model for Understanding Ligand Stability in Zn Proteins, *Chem. – Eur. J.*, 2021, **27**, 13730–13738.
- 18 U. Darbost, X. Zeng, M.-N. Rager, M. Giorgi, I. Jabin and O. Reinaud, X-ray and Solution Structures of the First Zn Funnel Complex Based on a Calix[6]aza-cryptand, *Eur. J. Inorg. Chem.*, 2004, 4371–4374.
- 19 N. Le Poul, B. Douziech, J. Zeitouny, G. Thiabaud, H. Colas, F. Conan, N. Cosquer, I. Jabin, C. Lagrost, P. Hapiot, O. Reinaud and Y. Le Mest, Mimicking the Protein Access Channel to a Metal Center: Effect of a Funnel Complex on Dissociative versus Associative Copper Redox Chemistry, *J. Am. Chem. Soc.*, 2009, **131**, 17800–17807.
- 20 N. Le Poul, B. Colasson, G. Thiabaud, D. J. Dit Fouque, C. Iacobucci, A. Memboeuf, B. Douziech, J. Řezáč, T. Prangé, A. de la Lande, O. Reinaud and Y. Le Mest, Gating the electron transfer at a monocopper centre through the supramolecular coordination of water molecules within a protein chamber mimic, *Chem. Sci.*, 2018, **9**, 8282–8290.

