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Primary amine recognition in water by a calix[6]aza-cryptand incorporated in dodecylphosphocholine micelles†

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Water is a unique solvent and the design of selective artificial hosts that can efficiently work in an aqueous medium is a challenging task. It is known that the calix[6]tren zinc complex can recognize neutral guests in organic solvents. This complex was incorporated into dodecylphosphocholine micelles (DPC) and studied by NMR. The incorporated complex is able to extract selectively primary amines from the aqueous environment driven by an important hydrophobic effect which also affects the selectivity of the complex for these amines. This work shows how the incorporation of organo-soluble receptors in micelles can be an elegant and very efficient strategy to obtain water compatible nanosized supramolecular recognition devices which can be prepared via a straightforward self-assembly process.

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Introduction

The design of artificial receptors that can selectively bind, with a high affinity, charged or neutral species is one of the major objectives of supramolecular chemistry.¹ Indeed, such receptors have potential applications in the sensing of chemical species in the fields of biological and environmental analysis.² The development of such receptors is nevertheless challenging because water is a very competitive solvent.³ Exploiting the hydrophobic effect which can drive apolar guests into apolar cavities is an interesting strategy. In this context, many examples of neutral guest complexation have been reported with water-soluble cavitands having a hydrophobic inner space such as cyclodextrins⁴ and cucurbiturils.⁵ These macrocycles

however remain difficult to modify selectively⁶ which limits the design of sophisticated receptors. In the case of the other families of cavitands such as calixarenes, fastidious syntheses are often required in order to make them water-soluble.

Micelles are extensively used in a wide variety of industrial and recovery processes to incorporate lipophilic compounds⁷ and are thus a possible solution to overcome the solubility problem of organic receptors in water. This strategy is interesting, as no synthesis needs to be undertaken in order to introduce hydrophilic groups on the receptor. A few examples of receptors and sensors incorporated in micelles have been reported in the literature⁸ and in some cases their binding properties have been investigated and shown to be enhanced by the micellar environment. Rebek and co-workers have, for instance, observed that a conformational reorganization of a resorcinarene cavitand incorporated into dodecylphosphocholine (DPC) micelles increases significantly its binding capacity towards different guests.⁹ It has also been reported that uranyl-salophen receptors incorporated into cetyltrimethylammonium bromide (CTABr) micelles have an affinity for fluoride which is two order of magnitude larger than the affinity of the related water-soluble receptor.^{10,11}

Calix[6]arenes are a well-known class of concave macrocyclic compounds presenting a cavity well adapted for the inclusion of organic guests and which can be selectively modified.^{12,13} Depending on their functionalization, they have revealed to be efficient receptors for a variety of neutral or charged species. In this context, different generations of biomimetic receptors based on calix[6]arene ligands bearing nitrogenous coordinating arms on the narrow rim have been developed.¹⁴ In particular,

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† Electronic supplementary information (ESI) available: 2D DOSY experiment of 1-Zn²⁺ incorporated in DPC. NMR PRE experiment with 1-Zn²⁺ in DPC. ¹H NMR spectrum of 1-nH⁺ in DPC. Determination of the *pseudo* pK_a for PrNH₂ with 1-Zn²⁺ in DPC. ¹H NMR spectra of 1-Zn²⁺ in the absence and presence of EtNH₂ in DPC-d₃₈. ¹H NMR spectra of 1-Zn²⁺ in the absence and presence of HeptylNH₂ in DPC. Experimental conditions for titrations of 1-Zn²⁺ with amines, alcohols and aminoalcohols in DPC. See DOI: 10.1039/c4ob02495h



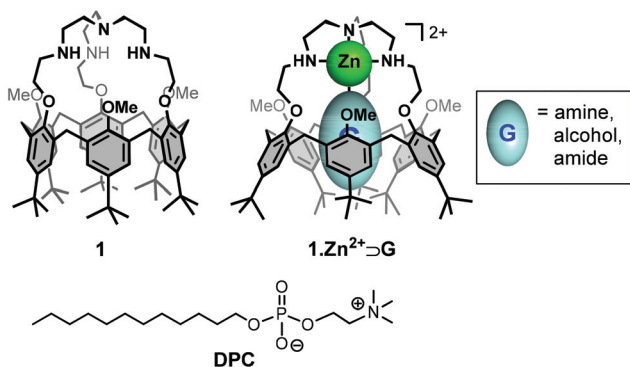


Fig. 1 Structures of ligand **1**, complex **1·Zn²⁺ ⊃ G** and DPC surfactant.

the calix[6]tren receptor **1**,¹⁵ belonging to the third generation has been shown to display unique and versatile binding properties, notably toward metal ions and neutral guests (Fig. 1).¹⁶ Indeed, the tren cap offers a strong binding site for a metal ion (*e.g.* Zn²⁺, Cu²⁺). Once coordinated, the metal ion presents a single coordination site accessible for an exogenous neutral guest, which is fully controlled by the hydrophobic funnel. Hence, complex **1·Zn²⁺** (Fig. 1) can bind with a high selectivity various neutral organic molecules (*e.g.* amines, alcohols, amides, *etc.*) inside the calix[6]arene core in different organic solvents.¹⁷

With the aim of developing selective molecular receptors that exhibit high affinity for specific guests in water, we were interested in evaluating the binding properties of complex

1·Zn²⁺ in an aqueous medium. In order to transfer this lipophilic metal complex into water, we thus envisioned its incorporation into micelles. Herein, we report the incorporation of calix[6]tren **1** and of its corresponding metal complex **1·Zn²⁺** into DPC micelles (Fig. 1). Characterization of the calix[6]-arene/micelle systems and evaluation of the binding properties of the incorporated receptors were achieved by NMR spectroscopy.

Results and discussion

Incorporation of complex 1·Zn²⁺ and ligand 1 into DPC micelles. The receptor **1·Zn²⁺** was successfully incorporated into DPC micelles by simple mixing of the receptor and the surfactant (20 mM) in D₂O at neutral pH (~7.6) and concentrations of up to 0.5 mM of **1·Zn²⁺** could be obtained.¹⁸ The ¹H NMR spectrum of the incorporated receptor **1·Zn²⁺** in DPC-*d*38 micelles shows a NMR pattern which is very similar to the one recorded in CDCl₃ (Fig. 2a vs. 2b). No significant changes were observed in the ¹H NMR spectrum between 283 K and 330 K. As shown previously, the complexity of the NMR profile arises from the heterochirality of the three nitrogen stereocenters which leads to a nonsymmetrical C₁ calixtren-Zn complex.¹⁷ Similarly to **1·Zn²⁺** in CDCl₃, the downfield shift of the OMe signals of the incorporated complex ($\delta_{\text{OMe}} > 3.9$ ppm in CDCl₃ and in D₂O)¹⁹ suggests that these groups have been expelled

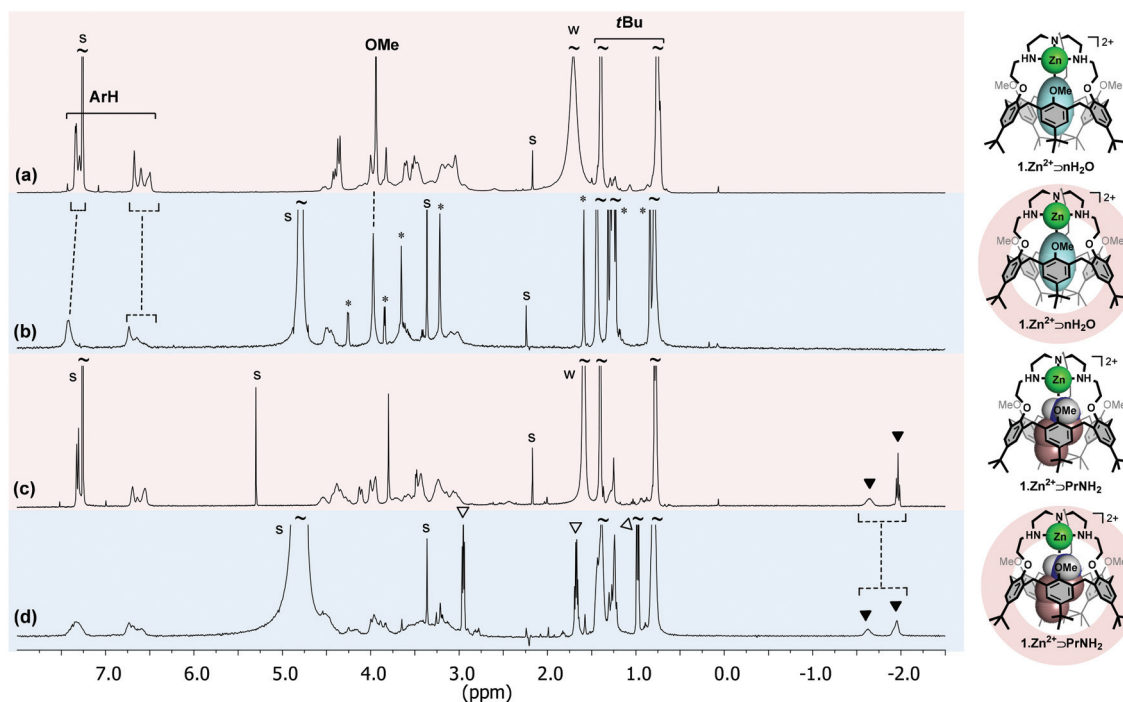


Fig. 2 ¹H NMR spectra (298 K) of (a) **1·Zn²⁺** in CDCl₃ (600 MHz); (b) **1·Zn²⁺** in DPC-*d*38 (20 mM in D₂O at pH ~7.6, 600 MHz); (c) **1·Zn²⁺** in CDCl₃ after the addition of ~1 equiv. of PrNH₂ (400 MHz); (d) **1·Zn²⁺** in DPC-*d*38 (20 mM in D₂O at pH ~8.6) after the addition of 5 equiv. of PrNH₂ (600 MHz). ▼: PrNH₂ in; ▽: PrNH₂ out; *: residual DPC; s: solvent; w: water.



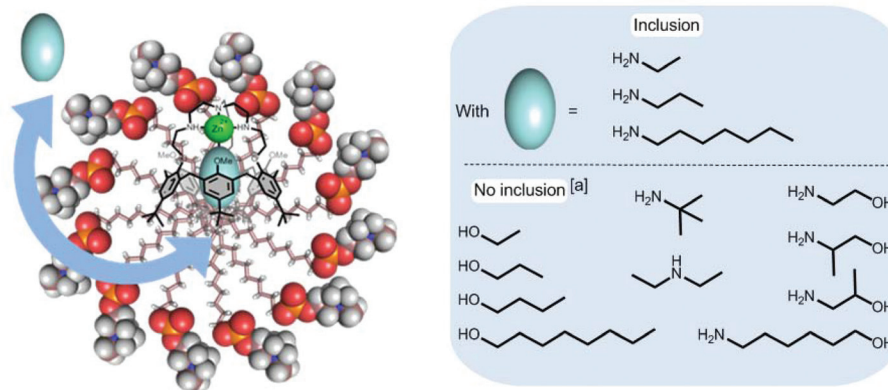


Fig. 3 Summary, using a schematic representation, of the host–guest properties of the receptor $1\cdot\text{Zn}^{2+}$ incorporated in DPC micelles. [a]: see ESI† for experimental conditions.²¹

from the cavity by a coordinated solvent water molecule.²⁰ Two very distinct sets of signals can be observed for the *t*Bu as well as for the ArH protons, indicating that the calixarene adopts a major flattened cone conformation. The incorporation of $1\cdot\text{Zn}^{2+}$ inside the DPC micelles was unambiguously confirmed by DOSY NMR experiments as both receptor and micelle exhibit the same diffusion coefficient, corresponding to a hydrodynamic radius of ~ 2 nm.²¹ Moreover, knowing that the micellar aggregation number is around 50,²² it is reasonable to estimate that there is 1 molecule of complex $1\cdot\text{Zn}^{2+}$ per micelle.

In order to gain information about the position of $1\cdot\text{Zn}^{2+}$ in the micelle, paramagnetic relaxation enhancement (PRE) experiments were undertaken.²³ It is indeed possible to obtain reliable information on the localization of receptors in micelles by comparing the PRE data obtained for the receptor protons with those for the surfactant protons.¹⁰ When plotting the relaxation rate enhancement as a function of paramagnetic species concentration, a linear relationship was obtained and the slope, known as the relaxivity, was extracted. The normalized relaxivities for the aromatic and *t*Bu protons of $1\cdot\text{Zn}^{2+}$ have values similar to those of the tail of the surfactant but very different from those of the hydrophilic part.²¹ It was thus possible to conclude that the calixarene positions itself in the hydrophobic core. Interestingly, the spectrum of $1\cdot\text{Zn}^{2+}$ in DPC micelles was not significantly modified by the addition of DCl or NaOD in the pH range from 6 to 9, highlighting the stability of the system in this pH window.²⁴

The calix[6]tren ligand **1** was also successfully incorporated into DPC micelles, by simple mixing in D_2O of the receptor and the surfactant, as attested by the NMR spectrum.²¹ It was however necessary to lower the pH to ~ 3 , showing that the ligand must be protonated at the level of its basic tren cap ($1\cdot n\text{H}^+$) in order to be incorporated.²⁵ This result suggests that the favourable electrostatic interaction of the positively charged $1\cdot n\text{H}^+$ and $1\cdot\text{Zn}^{2+}$ with the phosphate group of the zwitterionic surfactant is a driving force for their incorporation (Fig. 3). It is noteworthy to mention that the OMe groups of the incorporated receptor $1\cdot n\text{H}^+$ are pointing inside the cavity

($\delta_{\text{OMe}} = 2.41$ ppm), clearly highlighting the absence of a guest molecule inside the cavity. The efficient incorporation of $1\cdot n\text{H}^+$ and $1\cdot\text{Zn}^{2+}$ into DPC micelles encouraged us to undertake host–guest complexation studies with these systems.

Guest complexation by $1\cdot\text{Zn}^{2+}$ in DPC micelles

Propylamine (PrNH_2) is known to display a high affinity for $1\cdot\text{Zn}^{2+}$ in CDCl_3 .¹⁷ A titration with PrNH_2 of the receptor in DPC micelles was monitored by ^1H NMR spectroscopy. As the DPC solution was not buffered, the addition of the amine increased the pH somewhat during the titration (from 7.6 to 8.6). The following observations can be made from the spectrum recorded with the addition of 5 equiv. of PrNH_2 (pH = 8.6) (Fig. 2d):

(i) High-field signals for the protons of the alkyl chain of PrNH_2 (*i.e.* -1.62 and -1.95 ppm) attest of its inclusion inside the cavity and thus of the formation of the complex $1\cdot\text{Zn}^{2+} \supset \text{PrNH}_2$. These chemical shifts are similar to those obtained with the same system in CDCl_3 suggesting a similar positioning of the guest in the cavity (Fig. 2c vs. 2d). The exchange process is slow on the NMR chemical shift timescale.

(ii) There are only small changes in the chemical shifts of the aromatic and *t*Bu calixarene protons, which shows that the conformation of the receptor essentially does not change upon inclusion of a small amine such as propylamine. The receptor maintains a flattened cone conformation clearly evidenced by the large difference between the resonances of the *t*Bu or ArH protons pointing inside and pointing outside the cavity ($\Delta\delta_{\text{tBu}} \sim 0.65$ ppm and $\Delta\delta_{\text{ArH}} \sim 0.67$ ppm).

These NMR data show that the host–guest properties of $1\cdot\text{Zn}^{2+}$ toward amines are maintained in the micellar environment (Fig. 3). An apparent binding constant for propylamine (K_{app}) of $\sim 5 \times 10^3 \text{ M}^{-1}$ for a 1 : 1 binding was determined by signal integration at pH ~ 8 .²⁶ At pH ~ 10 , this constant was estimated to be $>10^4 \text{ M}^{-1}$. The complex $1\cdot\text{Zn}^{2+} \supset \text{PrNH}_2$ was observed on a larger pH window (pH from ~ 3 to ~ 11) in the micelle than complex $1\cdot\text{Zn}^{2+}$ devoid of organic guest coordinated to the Zn atom (*vide supra*). From these data, a *pseudo* pK_a of ~ 4.3 for complexed propylamine could be esti-



mated.²¹ This value can be compared to the pK_a value of free propylamine of 10.6. This remarkable result shows the extraordinary stabilization of the amine due to its complexation to the metal cation inside the calixarene cavity. This pK_a shift is comparable to the one estimated for heptylamine with a related water-soluble Zn-calixarene receptor bearing imidazole moieties.²⁷ It is interesting to point out that this related receptor does not complex $PrNH_2$ in water, which highlights the role played by the micellar environment on the complexation properties of the calix[6]tren-based system.

Complex $1\cdot Zn^{2+}$ in DPC micelles also binds ethylamine ($EtNH_2$) and heptylamine ($HeptylNH_2$) as evidenced by the presence of high field signals for their alkyl chain protons in the 1H NMR spectra.²¹ The affinity constants for these amines relative to $PrNH_2$ were evaluated by competition experiments (Table 1)²⁸ and the following sequence of affinities was observed at $pH \sim 11.1$ (± 0.1): $EtNH_2 > PrNH_2 > HeptylNH_2$. They are all of the same order of magnitude, with, however, a significant advantage to the smallest amine, in spite of its lower lipophilicity. This may be explained by a better fit of the guest considering the volume of the calix[6]arene cone as defined by the *t*Bu substituents at the large rim. Indeed, with other related *p*-*t*Bu-substituted calix[6]arene-based metal complexes studied in chloroform, it has been shown that 3- and 4-atom length guests that are fully encapsulated in the cavity (as shown here by the high-field shifted 1H -NMR signals), display the highest affinities. In contrast, the alkyl chain of the larger amino guest has to protrude outside the cavity and, in order to avoid a steric clash with the *t*Bu groups, the calix[6]-arene skeleton is forced to adopt an energetically less favourable straight conformation.²⁹ Finally, it is interesting to note that the relative affinity for $HeptylNH_2$ (vs. $PrNH_2$) in the micellar solution is spectacularly much higher than the one measured in $CDCl_3$ (i.e. 0.5 vs. 0.00025). Such a difference may well reflect the benefit of the hydrophobic effect for lipophilic amine binding in water.

Very interestingly, the binding of $tBuNH_2$ as well as the secondary amine $(Et)_2NH$ was not observed (from $pH \sim 8$ to $pH \sim 11$) despite the fact that these amines are less hydrophilic than $EtNH_2$ or $PrNH_2$. This result shows that steric hindrance at the level of amino group is a major factor of selectivity and thus highlights the cavity-based selectivity

ensured by the incorporated receptor. The complexation of other families of guests such as alcohols (ethanol, propanol, butanol, octanol) and aminoalcohols (ethanolamine, (\pm)-1-amino-2-propanol, (\pm)-2-amino-1-propanol, 6-amino-1-hexanol) was also studied. Again, no sign of inclusion was evidenced for any of these guests (Fig. 3). The fact that octanol, which should like heptylamine benefit from the hydrophobic effect, is not complexed by the calixarene suggests that the amino group plays a key role for complexation. It has indeed been shown that the affinity of $1\cdot Zn^{2+}$ for ethanol in $CDCl_3$ is four orders of magnitude lower than for propylamine ($K_{EtOH/PrNH_2} \sim 2.5 \times 10^{-4}$).¹⁷ In order to understand why the aminoalcohols, and in particular the small ethanolamine, are not recognized by $1\cdot Zn^{2+}$ incorporated into DPC micelles, the affinity for ethanolamine was measured in $CDCl_3$. The relative affinity compared to $PrNH_2$ was determined to be close to 1 ($K_{PrNH_2/Ethanolamine} \sim 0.8$, see Table 1). Such a discrepancy between the results obtained in water/micelle conditions and in an organic solvent is also assignable to the hydrophobic effect that obviously plays a major role in the host-guest recognition processes and in the observed selectivities. In this context, it is worth pointing out that DOSY experiments undertaken with DPC solutions showed that $PrNH_2$ does not incorporate into the DPC micelles but that $HeptylNH_2$ co-micellizes. The diffusion coefficient of $PrNH_2$ in the presence of DPC micelles is indeed identical to the diffusion coefficient of the amine in D_2O and this is not the case for $HeptylNH_2$ which clearly partitions itself between the aqueous and micellar environments. This difference can be rationalized in terms of the hydrophobicity of the amines which are more hydrophobic with increasing chain length.

Guest complexation by $1\cdot nH^+$ into DPC micelles

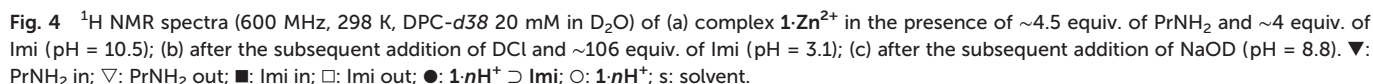
We have previously shown that calix[6]arene-based receptors bearing either a polyammonium,³¹ polyamido³² or polyureido cap³³ can bind with a high affinity the neutral guest 2-imidazolidinone (Imi) in their cavity. As shown on the XRD structure of one of these host-guest systems,³⁴ the high affinity for Imi stems from a complementary DAAD-ADDA quadruple H-bonding array between the urea guest and the calixarene host. Imi is also recognized by the protonated calixtren derivative $1\cdot nH^+$ in $CDCl_3$ and, to date, behaves as the best "key" for this receptor.³⁵ The binding of this neutral guest was thus evaluated in the micellar environment by NMR spectroscopy. It was observed that Imi (~ 230 equiv.) is not recognized by $1\cdot Zn^{2+}$ incorporated in DPC micelles (from $pH \sim 8$ to ~ 11). However, $1\cdot nH^+$ incorporated in DPC micelles at low pH complexes Imi under a slow exchange regime on the NMR chemical shift timescale. A singlet at 0.22 ppm is observed for the two methylene groups of the included Imi and an impressive downfield shift of the signal corresponding to the OMe groups ($\Delta\delta_{OMe} = 1.35$ ppm) attests to the ejection of these groups from the cavity upon guest inclusion. A large excess of Imi (40 equiv.), which is very soluble in water, was however required to observe the binding. Unfortunately, the addition of Imi also leads to a pH increase and ultimately to a partial

Table 1 Affinities, relative to $PrNH_2$ ($pK_a = 10.58$), of various guests G for host $1\cdot Zn^{2+}$

G (pK_a) ^b	Relative affinities ^a	
	DPC (20 mM in D_2O)	$CDCl_3$
$EtNH_2$ (10.67)	1.3 ($pH \sim 11.1$)	—
$HeptylNH_2$ (10.65)	0.5 ($pH \sim 11.2$)	~ 0.00025
$HO(CH_2)_2NH_2$ (9.50)	n.d. ^c ($pH \sim 9.4$)	0.8

^a Defined as $[G_{in}]/[PrNH_{2in}] \times [PrNH_{2Free}]/[G_{Free}]$ and measured at 298 K. The subscript "in" stands for "included". Errors estimated $\pm 20\%$. ^b For the pK_a values, see ref. 30. ^c Not detected with up to 50 equiv. of $HO(CH_2)_2NH_2$.





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Experimental section

Chemicals

DPC and DPC-*d38* were purchased from Avanti Lipids and used with no further purification. D₂O and CDCl₃ were purchased from VWR BDH PROLABO. Compounds **1** and **1**·Zn²⁺ were synthesized according to the literature.^{15,16}

General procedures

¹H NMR spectra were recorded on a 600 MHz, 400 MHz or a 300 MHz spectrometer. For the 1D ¹H spectra, parameters (acquisition time, recycling times, and signal accumulation) were chosen so as to ensure that quantitative data could be obtained from signal integration. Traces of residual solvents were used as internal chemical shift references. Chemical shifts are quoted on the δ scale. The NMR spectra were recorded at 298 K unless otherwise stated.

NMR solutions preparation

All solutions for NMR studies were prepared following a similar protocol: **1**·Zn²⁺ (~4 mg) and DPC (~35 mg, deuterated or not depending on the experiments) were added to 5 mL of D₂O in order to have, respectively, concentrations of ~0.5 mM and ~20 mM. Solutions were stirred until a clear solution was obtained.

¹H NMR characterization of various complexes **1**·Zn²⁺ \supset *n*H₂O

¹H NMR (DPC 20 mM in D₂O, 298 K, 600 MHz): δ (ppm) 0.79 (bs, 27H, *t*Bu), 1.44 (bs, 27H, *t*Bu), 3.97 (bs, 9H, OMe), 2.87–4.61 (m, 24H, ArCH₂^{eq}/ArCH₂^{ax}/CH₂N/CH₂O), 6.49–6.81 (m, 6H, ArH^{OMe}), 7.40 (bs, 6H, ArH^{cap}). **1**·Zn²⁺ \supset PrNH₂: ¹H NMR (DPC 20 mM in D₂O, 298 K, 600 MHz): * δ (ppm) –1.95 (bs, 3H, CH₃^{PrNH₂,in}), –1.63 (bs, 2H, CH₂^{PrNH₂,in}), 0.74–0.87 (m, 27H, *t*Bu), 1.36–1.57 (m, 27H, *t*Bu), 2.99–4.68 (m, 33H, ArCH₂^{eq}/ArCH₂^{ax}/CH₂N/CH₂O/OMe), 6.49–6.99 (m, 6H, ArH^{OMe}), 7.22–7.51 (m, 6H, ArH^{cap}). * The chemical shift of the CH₂N of PrNH₂ (in) was not observed due to overlapped signals.

Determination of the apparent affinity of PrNH₂ toward complex **1**·Zn²⁺ through ¹H NMR titration in DPC (20 mM in D₂O)

Different aliquots of a solution of amine (PrNH₂) were progressively added to 500 μ L of a DPC solution (20 mM in D₂O) containing the complex **1**·Zn²⁺. Non-deuterated DPC was used and the integration of the CH_{2 β} and CH_{2 γ} signals were used as internal reference to determine the concentration of **1**·Zn²⁺. The integrations of the free guests and of the included guests were used to calculate the apparent affinity defined as $[\text{PrNH}_{2\text{in}}]/(1 - [\text{PrNH}_{2\text{in}}] [\text{PrNH}_{2\text{Free}}][\text{1}\cdot\text{Zn}^{2+}])$ (errors estimated $\pm 20\%$).

Determination of the relative affinities of heptyl- and ethylamine compared to propylamine through ¹H NMR competitive binding studies in DPC (20 mM in D₂O)

At room temperature, ~7 equivalents of heptylamine (HeptylNH₂) or ~3 equivalents of ethylamine (EthylNH₂) were added to 500 μ L of a DPC solution (20 mM in D₂O) containing the complex **1**·Zn²⁺. Propylamine (PrNH₂) was then added (~5 equivalent). The ¹H NMR spectrum showed the guest resonances of both complexes **1**·Zn²⁺ \supset PrNH₂ and **1**·Zn²⁺ \supset HeptylNH₂ or **1**·Zn²⁺ \supset EthylNH₂ and signals corresponding to the free amines. The integrations of the signals corresponding to the free guests and included guests were used to calculate relative affinity defined as $[\text{G}_{\text{in}}]/[\text{PrNH}_{2\text{in}}] \times [\text{PrNH}_{2\text{Free}}]/[\text{G}_{\text{Free}}]$ (errors estimated $\pm 20\%$).

Determination of the relative affinity of ethylamine compared to propylamine in CDCl₃ via ¹H NMR competitive binding studies

At room temperature, ~30 equivalents of propylamine (PrNH₂) and ~15 equivalents of ethylamine (EthylNH₂) were added in 500 μ L of a CDCl₃ solution of complex **1**·Zn²⁺. A ¹H NMR spectrum showed the guest resonances of both complexes **1**·Zn²⁺ \supset PrNH₂ and **1**·Zn²⁺ \supset EthylNH₂ and signals corresponding to the free amines. The integrations of the signals corresponding to free guests and the included guests were used to calculate relative affinity defined as $[\text{EtNH}_{2\text{in}}]/[\text{PrNH}_{2\text{in}}] \times [\text{PrNH}_{2\text{Free}}]/[\text{EtNH}_{2\text{Free}}]$ (errors estimated $\pm 20\%$).

Determination of the relative affinity of heptylamine compared to DMF in CDCl₃ via ¹H NMR competitive binding studies

At room temperature, ~40 equivalents of dimethylformamide (DMF) and ~100 equivalents of heptylamine (HeptylNH₂) were added in 500 μ L of a CDCl₃ solution of complex **1**·Zn²⁺. A ¹H NMR spectrum showed the guest resonances of both complexes **1**·Zn²⁺ \supset DMF and **1**·Zn²⁺ \supset HeptylNH₂ and the signals corresponding to free DMF and HeptylNH₂. The integrations of the signals corresponding to the free and included guests were used to calculate relative affinity defined as $[\text{DMF}_{\text{in}}]/[\text{HeptylNH}_{2\text{in}}] \times [\text{HeptylNH}_{2\text{Free}}]/[\text{DMF}_{\text{Free}}]$ (errors estimated $\pm 20\%$). In order to obtain the value corresponding to $K_{\text{PrNH}_2/\text{HeptylNH}_2}$, we used the value for $K_{\text{PrNH}_2/\text{DMF}}$ from ref. 17 (1610).

Reversible interconversion of host-guest adducts

1·Zn²⁺ \supset PrNH₂ and **1**·nH⁺ \supset Imi

PrNH₂ (4.5 equiv.) and Imi (4 equiv.) were added to a ~0.5 mM solution of **1**·Zn²⁺ in DPC-*d38* micelles (pH ~10.5). In a second stage, DCl and a large excess of Imi (~106 equiv.) were added to the system (pH ~3.1). In the final stage, NaOD was added to reach pH = 8.8.

DOSY experimental parameters

15 values for the magnitude of the gradient pulses (ranging from 2 G cm^{–1} to 50 G cm^{–1}), diffusion delay 50 ms, time-length of the gradient 5 ms, acquisition time 2 s, relaxation delay 7 s, 32 transients.



PRE experimental parameters

For the PRE experiments, aliquots of 5 μl of a 5 mM solution of $\text{K}_3[\text{Cr}(\text{CN})_6]$ in D_2O were added to the 600 μl solution under study. T1 measurements were undertaken using the classical inversion recovery ($180 - \tau - 90 - \text{acquisition}$) sequence, with 15 points and the delay varying between 0 and 7 s. For each signal monitored, the increase in the longitudinal relaxation rate induced by the paramagnetic species ($\Delta 1/T_1$; the difference between the longitudinal relaxation rate measured in the presence and in the absence of the paramagnetic species) was plotted as a function of the concentration of paramagnetic species. A linear regression was undertaken with the experimental data points from which the relaxivity value was derived (slope of the line).

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