



Cite this: Dalton Trans., 2024, 53, 1779

Received 7th October 2023,
Accepted 12th December 2023

DOI: 10.1039/d3dt03292b
rsc.li/dalton

EHDTA: a green approach to efficient Ln^{3+} -chelators†

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The rich coordination chemistry of lanthanoid ions (Ln^{3+}) is currently exploited in a vast and continuously expanding array of applications. Chelating agents are central in the development of Ln^{3+} -complexes and in tuning their physical and chemical properties. Most chelators for Ln^{3+} -complexation are derived from the macrocyclic DOTA or from linear DTPA platforms, both of which arise from fossil-based starting materials. Herein, we report a green and efficient approach to a chelating agent (EHDTA), derived from cheap and largely available furfurylamine. The oxygenated heterocycle of the latter is converted to a stereochemically defined and rigid heptadentate chelator, which shows good affinity towards Ln^{3+} ions. A combination of NMR, relaxometric, potentiometric and spectrophotometric techniques allows us to shed light on the interesting coordination chemistry of Ln^{3+} -EHDTA complexes, unveiling a promising ligand for the chelation of this important family of metal ions.

Introduction

The rich coordination chemistry of lanthanoid ions (Ln^{3+}) is the key to exploiting the unique properties of these fascinating elements for an increasing number of applications. Particularly relevant are the applications of Ln^{3+} -complexes in medicine and life sciences.^{1,2} Paramagnetic Gd^{3+} -complexes have been routinely employed for the last three decades as contrast agents for MRI scans.³ $^{177}\text{Lu}^{3+}$ -oxodotreotide was recently approved by the EMA and FDA for targeted radiotherapy of somatostatin receptor positive neuroendocrine tumours.⁴ The unique luminescence properties of Eu^{3+} - and Tb^{3+} -chelates are especially useful in optical imaging and bioanalysis.⁵

The properties of Ln^{3+} -complexes strongly depend on the employed chelating agents and on their match with the coordination sphere of the metal ion.⁶ All *in vivo* applications require highly stable complexes both in terms of thermodynamic stability and kinetic inertness, usually provided by macrocyclic chelating agents such as DOTA (Fig. 1).⁷ On the other hand, short-lived radioisotope ions need to be chelated in a reasonably short timeframe to avoid excessive loss of radioactivity, which is best achieved with flexible linear chelating agents such as DTPA (Fig. 1).⁸ Chelating agents for luminescent com-

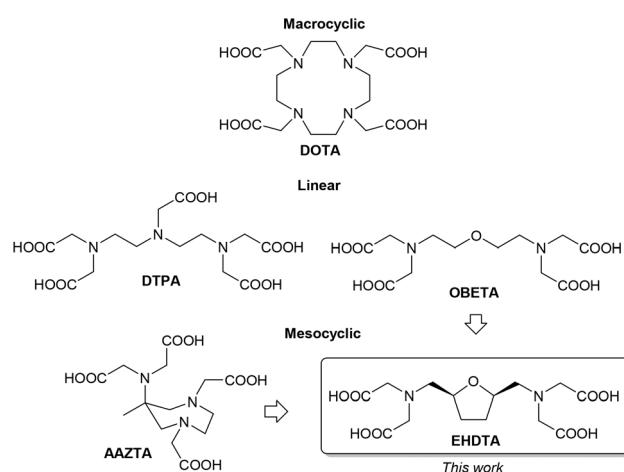


Fig. 1 Ln^{3+} -chelators.

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† Electronic supplementary information (ESI) available. CCDC 2235393. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d3dt03292b>



plexes should have active chromophores placed close to the metal ion, acting as an antenna for an efficient sensitization of lanthanide emission, and coordinated water molecules should be limited or absent to avoid fluorescence quenching.⁵ On the other hand, the number of coordinated water molecules ("q") is strictly related to the efficiency ("relaxivity") of Gd³⁺-based MRI contrast agents. In the octa- and nona-coordinated environment of Ln³⁺-ions, the denticity of the chelator and its steric hindrance define the *q* value, but the latter cannot be increased if not at the expense of the thermodynamic stability of the complex, implying a careful structural design devised to obtain a good match between stability and efficiency.³

Mesocyclic chelating agents such as AAZTA (Fig. 1) are able to coordinate Ln³⁺-ions more tightly than most linear congeners, combining a fast kinetics of formation and a slow kinetics of dissociation of the corresponding metal complexes with the possibility to modulate the number of coordinated water molecules.⁹

In this work, we report a green approach to the preparation of the mesocyclic chelating agent EHDTA[‡] (Fig. 1). EHDTA is formally related to OBETA, a linear heptadentate chelator that was recently shown to have an affinity for a metal ion stronger than the commercially available octadentate analogue EGTA.¹⁰ Moreover, a combination of NMR, relaxometric, potentiometric and spectrophotometric techniques is used to explore the coordination chemistry of Ln³⁺-EHDTA complexes.

Results and discussion

Design and synthesis

Our long-standing interest in the aqueous coordination chemistry of Ln³⁺-ions for biomedical applications led us to identify novel and efficient heptadentate chelators.⁹ Recently, a detailed investigation on Ln³⁺-OBETA complexes showed an unusual selectivity profile for this chelating agent in the Ln³⁺-series, which even overcomes the well-known higher-denticity homolog EGTA.¹⁰ We were then prompted to explore the possibility of improving the chelating ability of OBETA by acting on its molecular structure. Mesocyclic chelating agents, embodying a medium-sized ring in the backbone of the chelating agent, were shown to quickly form highly inert complexes. In this context, ring fusion of the linear OBETA molecule could rigidify the overall structure, providing even more stable metal complexes and eventually modulating the selectivity pattern through the Ln³⁺-series.³

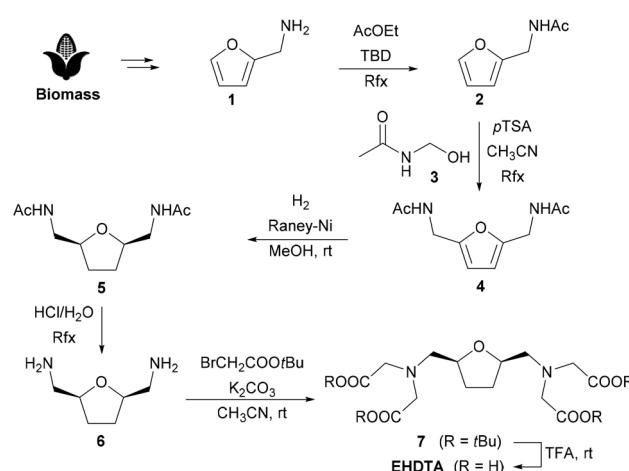
The tetrahydrofuran ring was chosen as a preferred scaffold for the development of improved chelating agents, as its oxygen atom encased in a strain-free five membered ring could easily play the role of the central oxygen atom of OBETA. Moreover, tetrahydrofuran derivatives are conveniently obtained by biomass processing, allowing us to shift the paradigm of current blockbuster chelating agents, obtained almost

completely by polyamines derived from petroleum-based chemicals.¹¹ As applications of Ln³⁺-complexes are constantly growing, sustainability issues must be taken into account in the design of the corresponding chelating agents. The formal ring fusion of the OBETA molecule with a tetrahydrofuran ring led to the chelating agent EHDTA, the subject of this work (Fig. 1). An early report on EHDTA dates back to 1977 with a preliminary study on the complexation of Pb²⁺.¹² Few details are given on the stereochemistry, inferred from the synthetic route to be *cis*.¹³ Moreover, the potential coordinating ability towards any other metal ion especially Ln³⁺ was overlooked, then and thereafter.

The original synthesis of EHDTA, even if amenable for a gram-scale preparation of these chelating agents, is far from being efficient and sustainable on a larger scale due to an inefficient atom economy and the use of toxic reagents, mainly related to the introduction of the amino groups through the Gabriel synthesis, performed on an intermediate di(*p*-toluenesulphonate).^{12,13} Therefore, the synthetic approach to EHDTA has been revised according to the principles of Green Chemistry.

The alternative preparation of EHDTA starts from furfurylamine **1**, a cheap biomass-derived chemical already endowed with a correctly placed amino group. *N*-Acetylation is accomplished by refluxing **1** in ethyl acetate, in the presence of TBD (1,5,7-triazabicyclo[4.4.0]dec-5-ene) as a transacylation catalyst, leading to *N*-furfurylacetamide **2**.

A Tscherniak-Einhorn amidomethylation¹⁴ with *N*-hydroxymethyl-acetamide¹⁵ **3** selectively implants the second (protected) amino group onto the furan ring, giving the symmetric heteroaromatic diamide **4**. Catalytic hydrogenation in methanol in the presence of RANEY®-Ni selectively provides the desired *cis*-disubstituted tetrahydrofuran **5**. Hydrolysis of the acetamide groups gives access to the diamine **6**, which is then exhaustively decorated with carboxymethyl groups in the classical alkylation-deprotection sequence (*t*-butyl bromoacetate/K₂CO₃, then TFA) to complete the synthesis of EHDTA (Scheme 1).



Scheme 1 Synthesis of EHDTA.

[‡] EHDTA = 2,5-EpoxyHexane-1,6-diamine-*N,N,N',N'*-TetraAcetic Acid.



Thermodynamic properties

The protonation constants of EHDTA were determined by pH potentiometry and ^1H NMR spectroscopy (Table S1 and Fig. S18†). The stability and protonation constants of some alkaline earth metals, transition metals and lanthanide complexes formed with EHDTA were also determined by pH-potentiometry (Tables S2 and S3†). Equations used for the evaluation of the equilibrium constants characterizing the protonation of the EHDTA ligand and of the stability and the protonation of the Mg^{2+} , Ca^{2+} , Sr^{2+} , Zn^{2+} , Cu^{2+} and Ln^{3+} -complexes are summarized in the ESI.†

The $\log K_{\text{LnL}}$ values of $\text{Ln}(\text{EHDTA})^-$ complexes increase from the La^{3+} - to Gd^{3+} -ion followed by a slight decrease for the latter members of the series, whereas the $\log K_{\text{LnL}}$ values of $\text{Ln}(\text{OBETA})^-$ decrease markedly at the end of the lanthanide series (Fig. 2). As a result, the $\Delta \log K_{\text{LnL}}$ values between the complexes formed by the two ligands with the same Ln^{3+} ion shows a minimum between La^{3+} and Gd^{3+} and a nearly monotonous increase up to Lu^{3+} (Fig. 2). The stability trend observed for $\text{Ln}(\text{OBETA})^-$ complexes is explained by the facts that the

flexible system embodying the ether oxygen and two nitrogen donor atoms and four carboxylate groups is well suited for the coordination of middle sized Ln^{3+} -ions, in particular the optimal coordination match with Gd^{3+} , resulting in a drop in the stability constants for the heavier Ln^{3+} ions.^{10,16} However, the stability trend of $\text{Ln}(\text{EHDTA})^-$ is very similar to that of $\text{Ln}(\text{DTPA})^{2-}$ complexes (Fig. 2), increasing from La^{3+} to Dy^{3+} and then being practically constant in the Ln -series with a slight decrease for the heavier Ln^{3+} ions, owing to the electrostatic repulsion between the negatively charged carboxylate groups in the complexes of the smaller Ln^{3+} ions.¹⁷ The presence of the five membered tetrahydrofuran ring results in an additional steric constraint of the donor atoms, which may explain the decrease of the $\log K_{\text{LnL}}$ values of $\text{Ln}(\text{EHDTA})^-$ complexes formed with the smaller Ln^{3+} -ions due to the increased electrostatic repulsion of the carboxylate groups.

Kinetic inertness of $\text{Gd}(\text{EHDTA})^-$

The kinetic inertness of $\text{Gd}(\text{EHDTA})^-$ was determined by following either the transmetallation reactions with Cu^{2+} and Eu^{3+} by spectrophotometry or the transchelation reactions with TTHA by ^1H NMR relaxometry in the presence of Cu^{2+} , Eu^{3+} and TTHA excess to ensure pseudo-first-order conditions (H_6TTHA = triethylenetetra-aminehexaacetic acid). In order to determine the role of the ternary complexes formed with small endogenous ligands on the dissociation rate of $\text{Gd}(\text{EHDTA})^-$, the transchelation reactions between $\text{Gd}(\text{EHDTA})^-$ and TTHA were investigated in the presence of citrate, phosphate and carbonate ions. The rate (k_x) and equilibrium constants (K_x) characterizing the transmetallation and transchelation reactions of $\text{Gd}(\text{EHDTA})^-$ are summarized in Table 1. Definitions and equations used for the evaluation of the equilibrium and kinetic data are reported in the ESI.†

The transmetallation reactions of $\text{Gd}(\text{EHDTA})^-$ with Cu^{2+} and Eu^{3+} were monitored on the absorption band of the result-

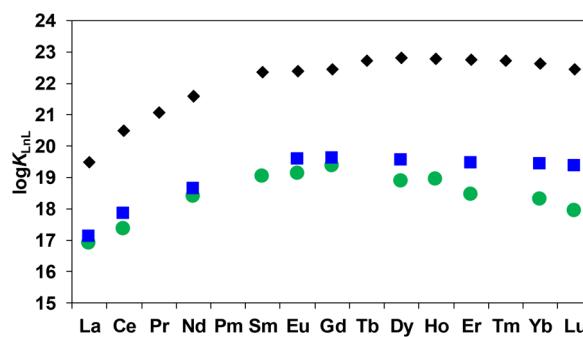
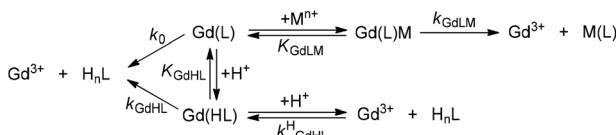


Fig. 2 Stability constants ($\log K_{\text{LnL}}$) of the $\text{Ln}(\text{EHDTA})^-$ (■), $\text{Ln}(\text{OBETA})^-$ (●) and $\text{Ln}(\text{DTPA})^{2-}$ (◆) complexes.

Table 1 Rate (k_x) and equilibrium constants (K_x) characterizing the dissociation reactions of $\text{Gd}(\text{EHDTA})^-$, $\text{Gd}(\text{OBETA})^-$ and $\text{Gd}(\text{DTPA})^{2-}$ complexes (25 °C)

Parameters	$\text{Gd}(\text{EHDTA})^-$ 0.1 M KCl	$\text{Gd}(\text{OBETA})^-$ ^{a,b} 0.1 M KCl	$\text{Gd}(\text{DTPA-BMA})^{2-}$ ^c 1.0 M KCl	$\text{Gd}(\text{DTPA})^e$ 1.0 M KCl
$k_1^f/\text{M}^{-1} \text{s}^{-1}$	3.3 ± 0.5	7.8	12.7	0.58
$k_2^g/\text{M}^{-2} \text{s}^{-1}$	$(3.6 \pm 0.4) \times 10^4$	2.1×10^5	—	9.7×10^4
$k_{\text{GdHL}}^h/\text{M}^{-1} \text{s}^{-1}$	0.37 ± 0.04	0.15	0.63	0.93
$k_3^i/\text{M}^{-1} \text{s}^{-1}$	$(2.8 \pm 0.3) \times 10^{-3}$	6.4×10^{-2}	—	4.9×10^{-4}
$k_{\text{cit}}/\text{M}^{-1} \text{s}^{-1}$	—	1.3×10^{-2}	5.6×10^{-3} ^d	4.4×10^{-5} ^d
$k_{\text{Hcit}}/\text{M}^{-1} \text{s}^{-1}$	0.33 ± 0.01	1.05	1.14 ^d	7.7×10^{-3} ^d
$k_{\text{CO}_3}/\text{M}^{-1} \text{s}^{-1}$	—	—	2.7×10^{-2} ^d	3.1×10^{-4} ^d
$k_{\text{HCO}_3}/\text{M}^{-1} \text{s}^{-1}$	$(7.0 \pm 0.2) \times 10^{-4}$	3.5×10^{-4}	—	2.2×10^{-5} ^d
$k_{\text{H}_2\text{PO}_4}/\text{M}^{-1} \text{s}^{-1}$	$(1.6 \pm 0.2) \times 10^{-3}$	5.5×10^{-2}	3.1×10^{-3} ^d	2.7×10^{-4} ^d
$K_{\text{GdHL}}/\text{M}^{-1}$	115 (pH-pot.)	158.5 (pH-pot.)	<2	100
$K_{\text{GdLCu}}/\text{M}^{-1}$	—	—	—	13
$K_{\text{GdLEu}}/\text{M}^{-1}$	—	23	—	19
$K_{\text{GdLCit}}/\text{M}^{-1}$	—	156	25 ^d	—
$K_{\text{GdLCO}_3}/\text{M}^{-1}$	942 ± 30	100	20 ^d	—
k_d/s^{-1}	1.7×10^{-5}	1.8×10^{-5}	3.8×10^{-6}	6.3×10^{-7}
$t_{1/2}/\text{h}$	12	11	50	305

^a Ref. 10. ^b Ref. 16. ^c Ref. 18. ^d Ref. 19 (0.15 M NaCl, 25 °C). ^e Ref. 20. ^f $k_1 = k_{\text{GdHL}} \times K_{\text{GdHL}}$, ^g $k_2 = k_{\text{H}_2\text{PO}_4}^{\text{H}} \times K_{\text{GdHL}}$, ^h $\text{h}_3\text{M}_3 = k_{\text{GdLM}} \times K_{\text{GdLM}}$.



Scheme 2 Spontaneous, proton- and metal-assisted dissociation of $\text{Gd}(\text{EHDTA})^-$.

ing Cu^{2+} - and $\text{Eu}(\text{EHDTA})^-$ complexes in the pH range 3.5–6.0 ($[\text{GdL}] = 0.1$ and 1.0 mM in the exchange reactions with Cu^{2+} and Eu^{3+} , 0.1 M KCl, 25 °C). The proposed mechanism for the transmetallation of $\text{Gd}(\text{EHDTA})^-$ is shown in Scheme 2.

The k_1 rate constant characterizing the proton-assisted dissociation of $\text{Gd}(\text{EHDTA})^-$ is about two times smaller than that of $\text{Gd}(\text{OBETA})^-$. Based on the mechanism of proton-assisted dissociation, the proton transfer from the carboxylic acid pendant of the protonated $\text{Gd}(\text{HL})$ intermediate to the N-atom of the ligand backbone takes place by the formation of a relatively labile protonated intermediate, which can dissociate into a free Gd^{3+} -ion and a protonated ligand. The structural preorganization of the EHDTA ligand results in less probable proton transfer and a slower rate of de-coordination of the Gd^{3+} ion from the protonated $\text{Gd}(\text{EHDTA})$ intermediate. However, the proton-assisted dissociation of both $\text{Gd}(\text{EHDTA})^-$ and $\text{Gd}(\text{OBETA})^-$ complexes is slower than for $\text{Gd}(\text{DTPA-BMA})$, which might be interpreted by a more rigid coordination environment in the Gd^{3+} -complexes of EHDTA and OBETA. This is further confirmed by the comparison of the rate constant characterizing the metal-assisted dissociation (k_3^{Cu}) of $\text{Gd}(\text{EHDTA})^-$, $\text{Gd}(\text{OBETA})^-$ and $\text{Gd}(\text{DTPA-BMA})$ (Table 1). The k_3^{Cu} rate constants of $\text{Gd}(\text{EHDTA})^-$ and $\text{Gd}(\text{OBETA})^-$ are comparable and about 2–4 times lower than that of $\text{Gd}(\text{DTPA-BMA})$,¹⁸ which indicates higher resistance of both $\text{Gd}(\text{EHDTA})^-$ and $\text{Gd}(\text{OBETA})^-$ complexes against Cu^{2+} -mediated dissociation due to the rigidity of both Gd^{3+} -complexes. Presumably, the stability of the $\text{Gd}(\text{L})\text{Cu}$ intermediate of $\text{Gd}(\text{EHDTA})^-$ and $\text{Gd}(\text{OBETA})^-$ is significantly lower than that of $\text{Gd}(\text{DTPA-BMA})$ resulting in a lower concentration of the kinetically active $\text{Gd}(\text{L})\text{Cu}$ intermediate and in the slower Cu^{2+} assisted dissociation of both Gd^{3+} -complexes.

It is known that bis-hydrated Gd^{3+} -complexes might form ternary species with endogenous ligands *via* the substitution of the inner-sphere water molecules.^{16,21–23} In order to investigate the role of endogenous ligands in determining the kinetic inertness of $\text{Gd}(\text{EHDTA})^-$ the transchelation reactions with TTHA were monitored by ^1H NMR relaxometry in the presence of excess citrate, phosphate and carbonate. Previous studies indicate that the transchelation rate of the Gd^{3+} -complexes increases with [TTHA] in the pH range of 6.5–11.0.²⁴ The contribution of TTHA to the transchelation rate of $\text{Gd}(\text{EHDTA})^-$ should be minimized in order to obtain reliable kinetic data for the catalytic effect of citrate, phosphate and carbonate ions. Based on our model calculations, the transchelation between $\text{Gd}(\text{EHDTA})^-$ and TTHA takes place with 100% conversion even in the presence of a twofold TTHA excess

(Fig. S21†) due to the large difference between the stability constant of $\text{Gd}(\text{EHDTA})^-$ and $\text{Gd}(\text{TTHA})^{3-}$ ($\log K_{\text{Gd}(\text{EHDTA})} = 19.62(3)$, $\log K_{\text{Gd}(\text{TTHA})} = 23.53(1)$, 0.1 M KCl, 25 °C). Therefore, the kinetic inertness of $\text{Gd}(\text{EHDTA})^-$ had been studied by following the transchelation reactions with TTHA in the pH range 6.0–10.0 in the presence of 2 eq. of TTHA, 0–8 eq. of citrate, 0–9 eq. of phosphate and 0–30 eq. of carbonate ($[\text{GdL}] = 1.0$ mM, $[\text{TTHA}]_t = 2.0$ mM, $[\text{Cit}^{3-}]_t = 0$ –8 mM, $[\text{PO}_4^{3-}]_t = 0$ –9 mM, $[\text{CO}_3^{2-}]_t = 0$ –30 mM, 0.1 M KCl, 25 °C). The rate (k_X) and equilibrium constants (K_X) characterizing the contribution of citrate, phosphate and carbonate ions to the rate of the transchelation reaction between $\text{Gd}(\text{EHDTA})^-$ and TTHA are shown in Table 1. By taking into account the rate and equilibrium constants of Table 1, the k_d rate constants and the half-life ($t_{1/2} = \ln 2/k_d$) values for the dissociation reactions of $\text{Gd}(\text{EHDTA})^-$, $\text{Gd}(\text{OBETA})^-$ and $\text{Gd}(\text{DTPA-BMA})$ were calculated under physiological conditions ($[\text{H}^+] = 3.98 \times 10^{-8}$ M (pH = 7.4), $[\text{Cit}]_{\text{tot}} = 1.1 \times 10^{-4}$ M, $[\text{CO}_3]_{\text{tot}} = 2.5 \times 10^{-2}$ M, $[\text{PO}_4]_{\text{tot}} = 1 \times 10^{-3}$ M).

The transchelation reaction of $\text{Gd}(\text{EHDTA})^-$ with TTHA can take place with the assistance of Cit^{3-} , HCit^{2-} , H_2PO_4^- , HCO_3^- and CO_3^{2-} -ions *via* the formation of ternary $[\text{Gd}(\text{EHDTA})\text{X}]$ species. In the ternary $[\text{Gd}(\text{EHDTA})\text{X}]$ species, the electrostatic repulsion between the negatively charged donor atoms results in a fast rearrangement and rapid dissociation of the Gd^{3+} -complex into the free EHDTA ligand and Gd^{3+} ion, the latter quickly reacting with the exchanging TTHA ligand. Additionally, the proton transfer in the ternary $[\text{Gd}(\text{EHDTA})\text{X}]$ species formed with HCit^{2-} , H_2Cit^- , H_2PO_4^- and HCO_3^- -ions can also occur *via* the formation of a protonated $\text{Gd}(\text{EHDTA})$ complex, characterized by a significantly lower kinetic inertness than $\text{Gd}(\text{EHDTA})^-$. Comparison of the rate constants characterizing the citrate and phosphate assisted dissociation of Gd^{3+} -complexes reveals that the HCit^{2-} and H_2PO_4^- catalysed dissociation of $\text{Gd}(\text{EHDTA})^-$ is about 3 and 34 times slower than those of $\text{Gd}(\text{OBETA})^-$, respectively. Moreover, the HCO_3^- assisted dissociation of $\text{Gd}(\text{EHDTA})^-$ (k_{HCO_3}) is about 2 times faster than that of $\text{Gd}(\text{OBETA})^-$. Interestingly, the stability constant of the ternary $[\text{Gd}(\text{EHDTA})\text{CO}_3]^{3-}$ species is about 10 times higher than that of $[\text{Gd}(\text{OBETA})\text{CO}_3]^{3-}$. Such a difference between the kinetic properties of $\text{Gd}(\text{OBETA})^-$ and $\text{Gd}(\text{EHDTA})^-$ might be attributed to the preorganized rigid structure of the EHDTA ligand and to the less efficient proton transfer in the ternary $[\text{Gd}(\text{EHDTA})\text{X}]$ complex formed with HCit^{2-} and H_2PO_4^- ions. In order to examine the possible contribution of the proton transfer processes to the rate of HCit^{2-} , H_2PO_4^- and HCO_3^- assisted dissociation of $\text{Gd}(\text{EHDTA})^-$, the k_X rate constants ($X = \text{HCit}^{2-}$, H_2PO_4^- and HCO_3^-) of $\text{Gd}(\text{EHDTA})^-$ and $\text{Gd}(\text{OBETA})^-$ were plotted as a function of the protonation constants ($\log K_X^{\text{H}}$, Table S5†) of the related ions (Fig. 3).

In Fig. 3, the k_X rate constants characterizing the protonated-ion assisted dissociation of $\text{Gd}(\text{EHDTA})^-$, $\text{Gd}(\text{OBETA})^-$ and $\text{Gd}(\text{DTPA})^{2-}$ decrease monotonously with the increase in the protonation constants of the H_2Cit^- , HCit^{2-} , HCO_3^- and H_2PO_4^- anions, which highlights the role of the protonated

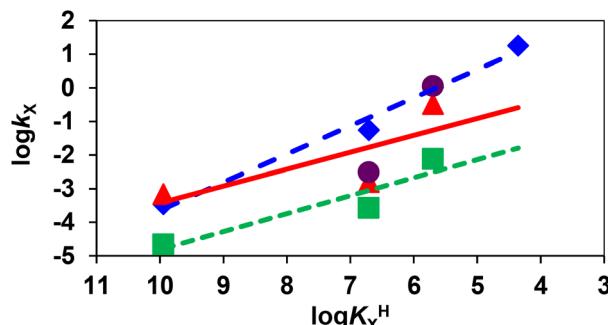


Fig. 3 Brønsted plot for the dissociation reaction of $\text{Gd}(\text{EHDTA})^-$ (▲), $\text{Gd}(\text{OBETA})^-$ (◆) and $\text{Gd}(\text{DTPA})^{2-}$ (■) and $\text{Gd}(\text{DTPA-BMA})$ (●) assisted by H_2Cit^- , HCit^{2-} , HCO_3^- and H_2PO_4^- ions (25 °C).

citrate, phosphate and carbonate ions in the dissociation of the Gd^{3+} -complexes. The rate of the proton catalysed reactions can take place by general acids, and the k_x rate constants are directly proportional to the acid strength of the general acid expressed by $\log K_x^{\text{H}}$ ($\log k_x = \alpha \times \log K_x^{\text{H}} + C$).²⁵ In the Brønsted plot, the reactions proceeding through an ideal proton transfer process are characterized by straight lines with slope $\alpha = 1.0$.²⁵ In the Brønsted plot (Fig. 3) the slope for the HCit^{2-} , HCO_3^- and H_2PO_4^- assisted dissociation reactions of $\text{Gd}(\text{EHDTA})^-$, $\text{Gd}(\text{OBETA})^-$ and $\text{Gd}(\text{DTPA})^{2-}$ was found to be $\alpha = 0.50$, 0.84 and 0.54, respectively. Based on the α values it can be assumed that the dissociation reactions of $\text{Gd}(\text{EHDTA})^-$ and $\text{Gd}(\text{DTPA})^{2-}$ take place with similar mechanisms *via* the fast rearrangement and the proton transfer processes in the ternary $[\text{Gd}(\text{EHDTA})\text{X}]$ intermediate. On the other hand the dissociation of $\text{Gd}(\text{OBETA})^-$ occurs mainly by the general acid catalysed pathway through proton transfer from the protonated HCit^- , HCO_3^- and H_2PO_4^- anions to OBETA in the ternary $[\text{Gd}(\text{OBETA})\text{X}]$ adducts.¹⁶ A somewhat lower contribution of the general acid catalysed pathway to the overall dissociation rate of $\text{Gd}(\text{EHDTA})^-$ can be interpreted by the preorganized structure of the EHDTA ligand and the less probable proton transfer from the protonated carboxylate group to the nitrogen donor atoms of the backbone.

Although, the Gd^{3+} -ion in $\text{Gd}(\text{DTPA-BMA})$ is coordinated by eight donor atoms of the ligand, the HCit^{2-} and H_2PO_4^- assisted dissociation rate of $\text{Gd}(\text{DTPA-BMA})$ is somewhat faster than that of the Gd^{3+} -complex formed with the heptadentate EHDTA ligand. The comparison of the k_d and $t_{1/2}$ values characterizes the dissociation rate of the Gd^{3+} -complexes close to physiological conditions and indicates that the kinetic inertness of $\text{Gd}(\text{EHDTA})^-$ and $\text{Gd}(\text{OBETA})^-$ is very similar and about 5 times lower than that of $\text{Gd}(\text{DTPA-BMA})$ due to the relatively fast HCO_3^- assisted dissociation of both Gd^{3+} -complexes.

¹H and ¹⁷O NMR relaxometric data

A detailed and complete relaxometric investigation of paramagnetic complexes in aqueous solution involves the measurement of the longitudinal ($R_1 = 1/T_1$) and transverse ($R_2 = 1/T_2$)

relaxation rates of the solvent nuclei as a function of pH, temperature and applied magnetic field. The global analysis of these data allows us to obtain valuable information on the stability of the complexes in aqueous solution, on the hydration state of the metal ion, on the exchange regime of the coordinated water molecules and an accurate estimation of the molecular parameters that control their effectiveness as relaxants.²⁶ This effectiveness is expressed by the relaxivity parameter. Relaxivity represents the increase in the longitudinal (r_1) or transverse (r_2) nuclear magnetic relaxation rates of the water protons normalized to a 1 mM aqueous solution of the Gd^{3+} complex.²⁷ At neutral pH and 298 K, for $\text{Gd}(\text{EHDTA})^-$ r_1 assumes the values of 7.1 and $6.9 \text{ mM}^{-1} \text{ s}^{-1}$ at 0.5 and 1.5 T, respectively. These values are fully comparable to those reported for $\text{Gd}(\text{OBETA})^-$, despite the lower molecular mass of the latter.^{10,16} Thus, also in this case two water molecules are presumably present in the inner coordination sphere of the complex ($q = 2$). Relaxivity does not show a pH dependence in the range 4–10 (Fig. S27†), confirming the good stability of the complex and the low propensity to form ternary complexes with dissolved carbonate, as previously observed for $\text{Gd}(\text{OBETA})^-$. In addition, this behaviour indicates that over this range of pH values, the complex does not hydrolyse and does not vary the hydration number q . One can safely conclude that in this pH range, all the seven donor atoms are coordinated to the metal, *i.e.* the structure of the complex is stable over a change of six orders of magnitude of proton concentration.

The measurement of the variation of r_1 as a function of the magnetic field strength allows us to obtain the so-called Nuclear Magnetic Resonance Dispersion profile (¹H NMRD). These contain key information on the physicochemical parameters correlating r_1 with the structural and dynamic properties of the complexes.²⁶ The ¹H NMRD spectra of $\text{Gd}(\text{EHDTA})^-$ were recorded at 283, 298 and 310 K in the proton Larmor frequency range 0.01–120 MHz, corresponding to the magnetic field strengths varying between 2.34×10^{-4} and 3.0 T (Fig. 4).

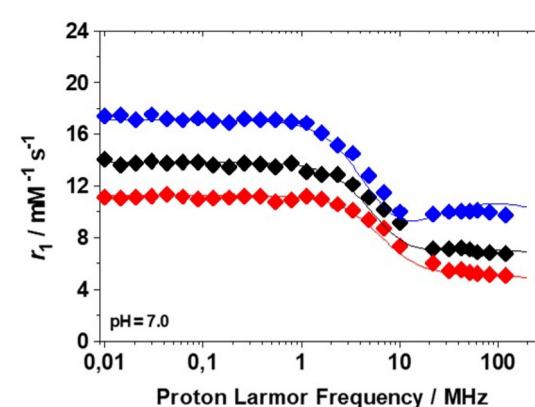


Fig. 4 $1/T_1$ ¹H NMRD profiles of $\text{Gd}(\text{EHDTA})^-$ at 283 (▲), 298 (◆) and 310 K (◆). The solid lines represent the results of the best fitting procedure to the experimental data (see Table 2).

The profiles are quite typical of low molecular weight Gd^{3+} complexes, whose salient features are the presence of a constant relaxivity in a wide range of low frequencies (up to *ca.* 1 MHz), the occurrence of a dispersion around at 3–6 MHz and another plateau with lower and almost constant relaxivity values at higher frequencies (>20 MHz). When these conditions occur, r_1 is largely dominated by the rotational dynamics, described by the rotational correlation time τ_R . Furthermore, the amplitude of the profiles decreases with increasing temperature (τ_R gets shorter) indicating that the system is in the fast exchange regime.²⁷ This implies that the rate of coordinated water exchange does not influence or limit relaxivity. The profiles were analysed using the Solomon–Bloembergen–Morgan^{28–30} and Freed³¹ equations that describe the contributions of the water molecules in the inner- and outer-sphere to the relaxation, respectively (see the ESI† for more details on the equations used).

Detailed and quantitative information on the exchange process of Gd -bound water molecules is derived from the analysis of data on the temperature dependence of the reduced ^{17}O NMR solvent transverse relaxation rates, R_{2r} ($= 1/T_{2r}$), and shift, $\Delta\omega_r$.²⁷ These experiments were carried out at 11.75 T on a 16.8 mM solution of the complex at neutral pH and the results are reported in Fig. 5. The reduced transverse relaxation rate increases with decreasing temperature until reaching a maximum of around 290 K, and then decreasing at lower temperatures. This is indicative of a rate of water exchange slightly slower than that of Gd(OBETA)^- but faster than those measured for $[\text{Gd(DTPA)}]^{2-}$ and $[\text{Gd(DOTA)}]^-$ for which the maximum is shifted at higher temperatures.³² As expected, $\Delta\omega_r$ exhibits an inflection point at a temperature quite similar to that in which $1/T_{2r}$ reaches the maximum value.

The data are conveniently analysed using the Swift–Connick equations that depend on a number of parameters, among which the most relevant are:³³ (i) the parameters associated with the electronic relaxation times $T_{1,2e}$, *i.e.* the trace of the square of the zero-field splitting, Δ^2 ; the correlation time describing the modulation of the zero-field splitting, τ_V , and its activation energy, E_V ; (ii) the enthalpy of activation for the water exchange process, ΔH_M ; (iii) the hyperfine Gd^{3+} – $^{17}\text{O}_{\text{water}}$ coupling constant, A/\hbar . Many of these parameters also affect the temperature dependence of the ^{17}O NMR shift. Due to the large number of parameters, for the global data fitting it is necessary to fix some of them to well-established or reasonably estimated values. Initially, the hydration number q was fixed to 2; the distance between the metal ion and the protons of the bound water molecules, r , was fixed to 3.0 Å; a and the D , the relative diffusion coefficient between solute and solvent molecules, were set to 4.0 Å and $2.3 \times 10^{-5} \text{ cm}^2 \text{ s}^{-1}$ (at 298 K), respectively; E_V was fixed to 1.0 kJ mol⁻¹.

Unexpectedly, the results of the global analysis of ^1H and ^{17}O NMR data were rather unsatisfactory. In particular, the calculated NMRD profiles consistently showed slightly lower values than the experimental ones. Excellent results could be obtained only by assuming an unreasonably long distance between the Gd^{3+} ion and the coordinated water molecules.

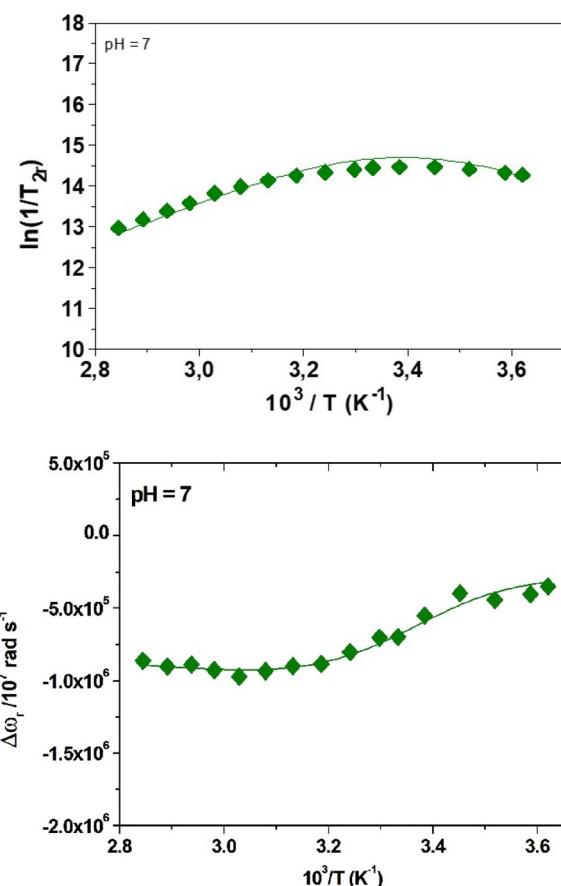


Fig. 5 Temperature dependence of the reduced water ^{17}O NMR transverse relaxation rates (top) and chemical shifts (bottom) at 11.75 T and pH = 7 for a 16.8 mM solution of $\text{Gd}(\text{EHDTA})^-$.

Since no evidence supports this hypothesis, we preferred to proceed with the fit by treating q as a variable parameter. An excellent result was obtained with the parameters reported in Table 2, where the hydration number q assumes a value equal to 1.8. A fractional q value implies the presence in solution of a hydration equilibrium between species differing in their hydration state ($q = 1$ and $q = 2$). This result is not at all unusual and has been previously reported in the case of

Table 2 Parameters obtained from the simultaneous analysis of ^1H NMRD profiles and ^{17}O NMR data (11.75 T) for the Gd^{3+} complexes of EHDTA and OBETA¹⁶

Parameters	$\text{Gd}(\text{EHDTA})^-$	$\text{Gd}(\text{OBETA})^-$
$^{22}r_{1(298 \text{ K})} (\text{mM}^{-1} \text{ s}^{-1})$	7.25	7.20
$\Delta^2/10^{19} \text{ s}^{-2}$	5.8	4.3
τ_V/ps	13.0	17.0
τ_M/ns	125	77
$\Delta H_M/\text{kJ mol}^{-1}$	41.0	40.1
τ_R/ps	79	65
$E_R/\text{kJ mol}^{-1}$	17	16
$A_O/\hbar/10^6 \text{ rad s}^{-1}$	-2.9	-2.9
q	1.8	2



Gd^{3+} ³⁴ and Mn^{2+} -complexes.³⁵ Furthermore, a similar conclusion is confirmed by photophysical measurements conducted on the corresponding Eu^{3+} and Tb^{3+} complexes (see *infra*).

The other best-fit parameters confirm the previous, qualitative conclusions. The residence lifetime of the inner sphere water molecule(s) is less than a factor of two longer than that of Gd(OBETA)^- , while it is about half that reported for $[\text{Gd(DTPA)}]^{2-}$ and $[\text{Gd(DOTA)}]^-$.³² The parameters of the electron relaxation times, Δ^2 and τ_v , are quite comparable to those found for the parent complex Gd(OBETA)^- , whereas the slightly longer rotational correlation time reflects the larger molecular weight of Gd(EHDTA)^- .

It is well known that complexes with $q > 1$ are able to form ternary compounds with bidentate oxyanions of biological relevance (e.g., lactate, citrate, oxalate, and carbonate) through the displacement of one or both coordinated water molecules.²¹ When this process takes place, we can easily observe a decrease in relaxivity that reflects the decrease in q . We monitored the variation of relaxivity of Gd(EHDTA)^- as a function of increasing concentration of bicarbonate, phosphate, oxalate, lactate and citrate up to 115 equivalents, at 32 MHz and 298 K (Fig. 6). The relaxometric titration data thus obtained can be analysed using the proton relaxation enhancement (PRE) method which allows the assessment of the apparent affinity constant K_A and the relaxivity of the ternary complex, r_1^b .²⁷ The curves of Fig. 6 and the analysis of the data (Table 3) clearly indicate that only oxalate has a weak but evident interaction with the complex ($K_A = 179 \text{ M}^{-1}$) and that the resulting ternary complex exhibits $q = 0$, as suggested by the very low value of relaxivity that is typical of pure outer sphere Gd-chelates. In all other cases the interaction is extremely weak and the variations in relaxivity are barely detectable. The values of the association constants are all in the range of *ca.* 40–50 M^{-1} and the calculated relaxivity of the ternary complexes ($6.5\text{--}7.0 \text{ mM}^{-1} \text{ s}^{-1}$) typical of systems with $q = 1$. These data confirm for Gd

Table 3 Parameters calculated by analysis of the titration curves of Fig. 6

	Phosphate	Lactate	Oxalate	Citrate	Carbonate
K_A/M^{-1}	39	51	179	46	43
$r_1^b/\text{mM}^{-1} \text{ s}^{-1}$	6.8	5.1	2.0	7.0	6.4

(EHDTA)[–] what was previously found in the case of Gd(OBETA)^- , *i.e.* a very weak affinity to interact with bidentate oxyanions to form ternary complexes under physiological conditions.

Photoluminescence

In consideration of the fractional q value emerging from the relaxometric data, we decided to carry out photoluminescence (PL) measurements on the Eu(EHDTA)^- complex to retrieve further evidence of the coordination environment around the lanthanide ion. Since the intrinsic quantum yield of Eu^{3+} emission is highly sensitive to the presence of water molecules in the first or second coordination sphere, we investigated the photophysical properties of the complex in H_2O and D_2O solutions (10^{-3} M) as well as on a solid state sample obtained by lyophilisation from an aqueous solution. Steady-state PL spectra, reported in Fig. 7, display the main distinctive Eu^{3+} f-f peaks in the visible range, associated with the $^5\text{D}_0 \rightarrow ^7\text{F}_J (J = 0, \dots, 4)$ transitions. The most intense (hypersensitive) line, corresponding to the $^5\text{D}_0 \rightarrow ^7\text{F}_2$ transition, is centred at 615 nm in aqueous solution and it is slightly shifted (617 nm) in the solid. A weak broad band, likely attributed to residual ligand emission, is observed in the region 400–650 nm, as con-

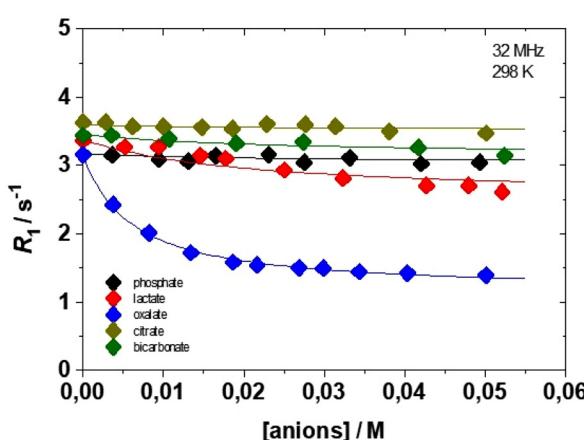


Fig. 6 Plot of the observed relaxation rate ($\text{pH} = 7.0$, 32 MHz, 298 K) of Gd(EHDTA)^- , as a function of $[\text{PO}_4^{3-}]$ ($[\text{Gd}^{3+}] = 0.39 \text{ mM}$), $[\text{lactate}]$ ($[\text{Gd}^{3+}] = 0.42 \text{ mM}$), $[\text{oxalate}]$ ($[\text{Gd}^{3+}] = 0.39 \text{ mM}$), $[\text{citrate}]$ ($[\text{Gd}^{3+}] = 0.45 \text{ mM}$) and $[\text{HCO}_3^-]$ ($[\text{Gd}^{3+}] = 0.43 \text{ mM}$).

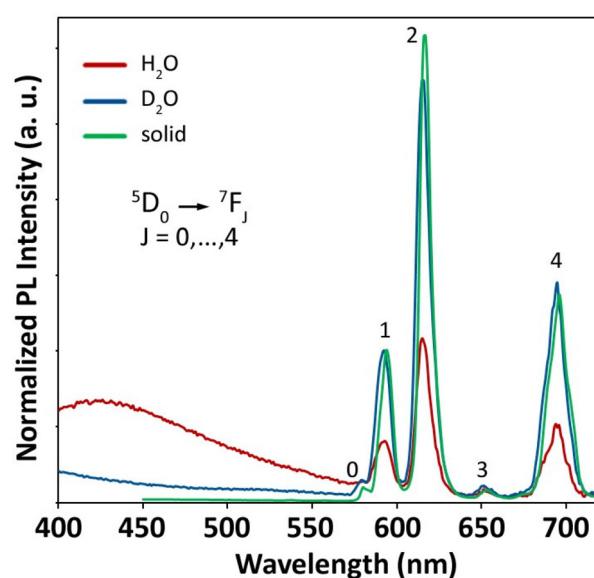


Fig. 7 Photoluminescence spectra of Eu(EHDTA)^- in the solid state (grey), and in H_2O (red) and D_2O (blue) solutions under excitation at 280 nm. Solution spectra are normalized to the $^5\text{D}_0 \rightarrow ^7\text{F}_3$ emission line of Eu^{3+} at 652 nm. The spectrum of the solid sample is normalized to the $^5\text{D}_0 \rightarrow ^7\text{F}_1$ emission line of the D_2O solution sample at 593 nm.



firmed by the PL spectrum of the optically silent Gd^{3+} analog (Fig. S28a†). The role of the ligand in the emission photocycle is evidenced by the excitation (PLE) spectra monitored at the Eu-centred emission peak at 615 nm and reported in Fig. S28b,† which displays a broad band centred at ~ 280 nm, alongside the characteristic sharp Eu^{3+} peaks (Table S6†). The appearance of such a broad band is the typical signature of a ligand-to-metal sensitization process.

The spectral shape and the relative intensity of the Eu^{3+} emission peaks are highly sensitive to the crystal/ligand field environment of the lanthanide ion, particularly for the hypersensitive $^5\text{D}_0 \rightarrow ^7\text{F}_2$ transition, and except for the purely magnetic $^5\text{D}_0 \rightarrow ^7\text{F}_1$ line (593 nm). Generally, a lowering of the symmetry results in a lifting of the selection rules and in a higher oscillator strength, which also yields an increase of the emission integrated intensity.³⁶ To achieve a quantitative estimation of the subtle coordination symmetry changes on going from the solid state to aqueous solution, we assessed the Eu^{3+} radiative rate constants, κ_{RAD} , in the three media. This parameter directly relates to the emitter's oscillator strength in the specific environment and corresponds to the reciprocal of the radiative lifetime τ_{RAD} ($\kappa_{\text{RAD}} = 1/\tau_{\text{RAD}}$), that is, the radiative emission lifetime in the absence of any quenching phenomena. κ_{RAD} values were obtained by applying a simplified Einstein's equation for spontaneous emission³⁷ to the integrated spectral data (ESI†). To minimize spurious effects related to the overlap of the ligand-centred emission band, solution spectra have been deconvoluted and subtracted for ligand contribution prior to the analysis (Fig. S29†). The obtained parameters are summarized in Table 4.

As expected, the results of H_2O and D_2O solutions are equivalent within experimental error. On the other hand, the radiative rate constant is remarkably higher in the solid state, an effect that is not only attributable to the difference in the refractive indexes ($\kappa_{\text{RAD}}(\text{solid})/\kappa_{\text{RAD}}(\text{H}_2\text{O}) = 1.57$ and $n(\text{solid})^3/n(\text{H}_2\text{O})^3 = 1.43$). To discriminate the contribution of the refractive index n of the different media from the local symmetry effects, we also removed the offset by calculating κ_{RAD} for the solid state sample using the same n value of water (last column in Table 4). The fact that the retrieved value is still significantly higher than that found in solution is a clear indication of a lowering of the symmetry of the complex upon sample drying.

In this context, time-resolved PL spectroscopy provides a powerful tool to achieve a more in-depth picture of the coordination environment in such compounds. Fig. 8 shows the

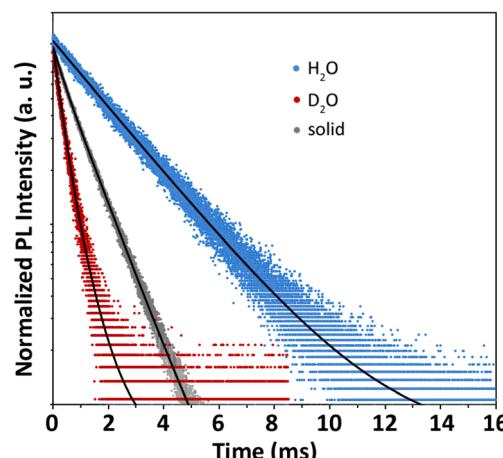


Fig. 8 Normalized time-resolved PL decay of $\text{Eu}(\text{EHDTA})^-$ in H_2O (red dots), D_2O (blue dots) and in the solid state (grey dots) upon excitation at 280 nm. Black solid curves represent the best fit to data (see the text). The vertical axis is reported on a logarithmic scale.

decay curves of $\text{Eu}^{3+} \text{ }^5\text{D}_0 \rightarrow ^7\text{F}_2$ transition for the $\text{Eu}(\text{EHDTA})^-$ complex in the different media upon excitation at 280 nm. Photophysical parameters including observed decay time constants, τ , rate constants $\kappa = 1/\tau$ and associated signal contributions (%) extrapolated from curve fitting are reported in Table 5 together with calculated emission intrinsic quantum yields ($\phi_{\text{Eu}}^{\text{Eu}} = \tau/\tau_{\text{RAD}}$).

As expected, the overall temporal dynamics in the deuterated solvent is remarkably slower than in water as a consequence of the replacement of the high-energy H_2O oscillators with D_2O molecules of lower phonon energy (Fig. S30†). In the solid state, where second-sphere solvent effects are removed, the decay dynamics remains significantly faster than in D_2O solution, indicating the occurrence of water molecules directly coordinated to the metal. Solid state data are best fitted with a biexponential function, where the contribution of the slow decay component (τ_{slow}) is dominant (%slow = 85%). In H_2O , the decay also follows a biexponential trend, but the population distribution is reversed and 72% of the signal is associated with the fast component (τ_{fast}). The observed double-exponential deactivation behaviour indicates the presence of two populations of Eu^{3+} emitters which undergo quenching to a different extent, suggesting the existence of molecular species bearing one of two H_2O quenchers in the first coordination sphere. This is consistent with the inferred lowering of the symmetry in the solid state, which likely stems from the predominance of less symmetric and less quenched mono-hydrated molecular species with respect to bis-hydrated complexes, which instead represent the majority in water solution. In D_2O , signal decay is monoexponential, as expected, since water-quenching effects are suppressed. This also means that the complex has a stable single configuration only subtly differing for the number of coordinated H_2O molecules. Combining the steady-state with time-resolved data it can be retrieved that $\tau_{\text{RAD}} = 4.37$ ms for mono-hydrated and $\tau_{\text{RAD}} =$

Table 4 Radiative rate constants and lifetimes of $\text{Eu}(\text{EHDTA})^-$ in different media

	Solid	H_2O	D_2O	Solid (corrected)
Refractive index, n	1.5	1.33	1.33	1.33
κ_{RAD}^a ($\times 10^{-3} \mu\text{s}^{-1}$)	320.0	203.7	204.7	223.1
τ_{RAD} (μs)	3125	4908	4885	4483

^aThe error was estimated to be 10%.



Table 5 Photophysical parameters for Eu(EHDTA)⁻ in different media

	τ_{fast} (μs)	% _{fast}	τ_{slow} (μs)	% _{slow}	$\tau_{\text{amplitude}}^a$ (μs)	$\tau_{\text{Intensity}}$ (μs)	κ^b (×10 ⁻³ μs ⁻¹)	$\Phi_{\text{Eu}}^{\text{Eu}}$ (%)
Solid	344(8)	15%	1079(3)	85%	969	1040	0.962	31
H ₂ O	312(6)	72%	697(3)	28%	419	489	2.04	10
D ₂ O	—	—	2357(2)	100%	—	—	0.424	48

^a cf. Experimental section. ^b Calculated on average intensity time constants for double-exponential decays.

5.10 ms for bis-hydrated complexes, in agreement with literature reports.³⁸ To further confirm this interpretation, we applied the empirical model developed by Horrocks to retrieve the number of inner sphere water molecules (q), through the simple equation $q = A(\kappa_{\text{H}_2\text{O}} - \kappa_{\text{D}_2\text{O}})$, where A is a proportionality constant, whose value was taken as $A = 1.05$ ms while $\kappa_{\text{H}_2\text{O}}$ and $\kappa_{\text{D}_2\text{O}}$ represent the decay rate constants ($=1/\tau$) of Eu³⁺ emission in H₂O and D₂O, respectively.³⁹ For the double-exponential observed decay in water, the intensity average time constant (Table 5) was used (see the ESI†), and the model equation yielded $q = 1.7$. This non-integer value of the number of coordinated water molecules is in perfect agreement with a population distribution of bis-hydrated and mono-hydrated molecular species of 72% and 28%, which was inferred from time-resolved data. Following this convincing result, it can be assumed that the average number of water molecules in the solid state is 1.15. This observation points out that the second coordinated water molecule is rather labile and can be easily removed under vacuum (after solution lyophilisation) yielding preferentially mono-hydrated octa-coordinated species. This is consistent with the observed fractional q value in water, where, despite the presence of excess solvent molecules, octa-coordinated complexes still coexist with bis-hydrated nine-coordinated species.

The Eu³⁺ intrinsic quantum yield $\Phi_{\text{Eu}}^{\text{Eu}} = \tau/\tau_{\text{RAD}}$ in D₂O, where both directly coordinated H₂O quenchers as well as second-sphere effects are lifted, indicates that water accounts for more than 50% of emission quenching, and in H₂O $\Phi_{\text{Eu}}^{\text{Eu}}$ reduces to 10%. In the solid state, the combined effect of solvent removal and oscillator strength increase accounts for the observed intermediate value (31%) of the intrinsic quantum yield.

X-Ray diffractometric studies of the ternary [Gd(EHDTA)CO₃]³⁻ complex

Single crystals of the composition [C(NH₂)₃]₃[Gd(EHDTA)CO₃]³⁻·2H₂O suitable for X-ray diffraction studies were grown from an aqueous solution of the Gd(EHDTA)⁻ complex in the presence of 3.0 molar equivalents of guanidinium carbonate ([C(NH₂)₃]₂CO₃). The complex crystallizes in space group $P2_1/n$ ($Z = 4$). The uncoordinated water molecules and guanidinium counter ions are involved in an intricate hydrogen-bonds network with oxygen atoms of carboxylate groups of the ligand and the carbonate anion (Fig. S31 and Table S8 in the ESI†). The same arrangement and hydrogen bonding network were observed for [C(NH₂)₃]₃[Lu(OBETA)CO₃]³⁻·2H₂O¹⁶ which is in fact isostructural (Fig. S33 and S34†).

The similarities between both complexes thus extend also to the central ion: the Gd³⁺ ion is coordinated by seven donor atoms of the EHDTA ligand, whereas an eighth and ninth coordination site is occupied by an η^2 -carbonate ligand, coordinating in a bidentate fashion (Fig. 9). As for [Lu(OBETA)CO₃]³⁻, the carbonate ion coordinates in an almost perfect symmetrical binding mode. Likewise, the coordination polyhedron around the Gd³⁺ ion in [Gd(EHDTA)(CO₃)]³⁻ can be best described as a slightly distorted monocapped square antiprism defined by two nearly parallel pseudo planes (Fig. S31†) with the upper one going through O4, O9, O2 and O11 (mean atomic distance 0.163 Å) as well as the lower one going through N1, O7, O1, O5 (mean atomic distance 0.188 Å). The N2 amine nitrogen occupies the capping position above the upper plane.

Compared to [Lu(OBETA)CO₃]³⁻ the atomic distances to the best planes are smaller even though the angle between them is significantly larger (9° compared to 0.8°) probably owing to the higher rigidity and the preorganized structure of the EHDTA ligand. The average bond distances in the ternary [Gd(EHDTA)CO₃]³⁻ complex were found to be 2.453 Å (Gd–O) and 2.729 Å (Gd–N) respectively. This is significantly longer than for [Lu(OBETA)(CO₃)]³⁻ with 2.352 Å (Lu–O) and 2.619 Å (Lu–N),¹⁶ which is expected for metals in lanthanide series⁴⁰ due to

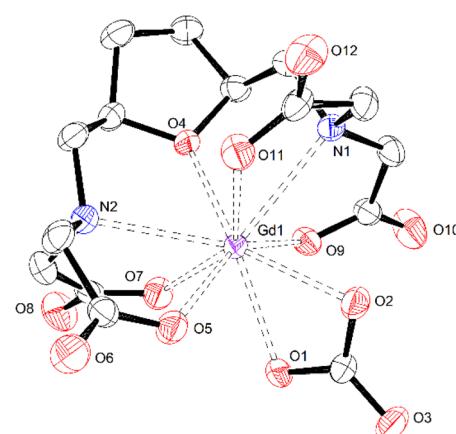


Fig. 9 Molecular structure of the [Gd(EHDTA)(CO₃)]³⁻ complex ion present in the crystal structure of [C(NH₂)₃]₃[Gd(EHDTA)(CO₃)]³⁻·2H₂O. Anisotropic displacement parameters are shown at the 50% probability level. Selected bond distances (Å): Gd1–N1 2.724(2); Gd1–N2 2.734(2); Gd1–O1 2.4429(18); Gd1–O2 2.4511(17); Gd1–O4 2.4793(17); Gd1–O5 2.4505(18); Gd1–O7 2.4234(18); Gd1–O9 2.4245(17); Gd1–O11 2.5002(18).



the smaller size of the Lu³⁺-ion (0.977 Å) compared to the Gd³⁺-ion (1.053 Å).

The change of the ligand framework from OBETA to EHDTA also leads to a slight change of the molecular conformation of both complexes. In [Gd(EHDTA)CO₃]³⁻ the five-membered chelate rings of the 2,2'-bis(aminomethyl)furan at O4 have opposite helicity *i.e.* (λδ) or (δλ) (both enantiomers present as space group is enantiogenic) while they are identical in [Lu(OBETA)(CO₃)]³⁻.⁴⁰ This is most likely caused by the higher rigidity of the EHDTA ligand. The remaining chelate rings in both complexes have the same relationship of helicity thus leading to an overall conformation of (λδ)(λδδδ) or (δλ)(δλλλ).

Experimental

General

Solvents and starting materials were purchased from Merck or TCI and used without further purification. All aqueous solutions were prepared from ultrapure laboratory grade water (18 MΩ cm) obtained from a Millipore/Milli-Q purification system. ¹H and ¹³C NMR spectra were recorded on a Bruker Avance Neo 400 spectrometer operating at 9.4 T. Chemical shifts are reported in ppm with the protic impurities of the deuterated solvent as the internal reference. Mass spectra were obtained with a Thermo Finnigan LCQ-Deca XP-PLUS ion trap spectrometer equipped with an electrospray source. High resolution mass spectra were registered on a Thermo Scientific Q-Exactive Plus spectrometer. TLC was performed with silica gel (MN Kieselgel 60 F254) and visualized by UV light or sprayed with the Dragendorff reagent or alkaline KMnO₄. Column chromatography was carried out on Macherey-Nagel Silica gel 60 (0.063–0.200 mm).

N-Furfurylacetamide (2). Furfurylamine (1, 25.0 g, 257 mmol) was dissolved in 200 mL of ethyl acetate and 1,5,7-triazabicyclo[4.3.0]oct-5-ene (TBD) (0.400 g, 2.87 mmol) was added. The mixture was refluxed for 30 h and checked by TLC (ethyl acetate or dichloromethane/methanol 9 : 1). The mixture was then cooled to room temperature, washed with 2 M aqueous HCl solution, NaHCO₃ saturated solution in water and brine and the organic phase was dried (Na₂SO₄) and filtered and the solvent was removed under vacuum to give compound 1 as a yellow oil (35.8 g, quant.). ¹H NMR (400 MHz, CDCl₃, 298 K) δ 7.32 (dd, *J* = 1.9, 0.9 Hz, 1H), 6.29 (dd, *J* = 3.2, 1.8 Hz, 1H), 6.20–6.19 (m, 2H), 4.38 (d, *J* = 5.5 Hz, 2H), 1.97 (s, 3H) ppm. ¹³C NMR (101 MHz, CDCl₃) δ 170.1 (C), 151.4 (C), 142.2 (CH), 110.5 (CH), 107.5 (CH), 36.6 (CH₂), 23.1 (CH₃) ppm. MS (ESI⁺): *m/z* = 140.07 (100%, [M + H]⁺). Calc. for C₇H₉NO₂ + H⁺: 140.07.

N,N'-Furan-2,5-diylbis(methylene))diacetamide (4). Acetamide (10.0 g, 169 mmol), paraformaldehyde (6.00 g, 200 mmol) and potassium carbonate (0.100 g, 0.723 mmol) were mixed and melted at 100 °C with stirring. After 1 h at this temperature the mixture was cooled to 60 °C and potassium hydroxide (0.0100 g, 0.178 mmol) was added with stirring. The mixture was cooled to RT and the *N*-hydroxymethylacetamide 3 was used without further purification. *N*-Furfurylacetamide 2

(17.8 g, 128 mmol, 1.0 eq.) and *p*-toluenesulfonic acid (PTSA) (1.21 g, 6.38 mmol, 0.05 eq.) were dissolved in 250 mL of acetonitrile and added to crude 3. The resulting mixture was refluxed for 15 h, replacing the distilled acetonitrile with an equal amount of fresh solvent by means of Dean–Stark apparatus. The product starts to separate during the reaction and its crystallization is completed by standing overnight at 4 °C. The product is collected by vacuum filtration and washed with acetone. The resulting brownish crystals were recrystallized from methanol to give compound 4 as off-white crystals (15.7 g, 59%). M. p.: 159.4–161.0 °C. ¹H NMR (400 MHz, DMSO-d₆, 298 K) δ 8.28 (br t, *J* = 5.7 Hz, 2H), 6.14 (s, 2H), 4.19 (d, *J* = 5.6 Hz, 4H), 1.83 (s, 6H) ppm. ¹³C NMR (100.6 MHz, DMSO-d₆, 298 K) δ 169.0 (C), 151.6 (C), 107.5 (CH), 35.5 (CH₂), 22.5 (CH₃) ppm. MS (ESI⁺): *m/z* = 211.11 (100%, [M + H]⁺). Calc. for C₁₀H₁₄N₂O₃ + H⁺: 211.11. HRMS (ESI⁺): *m/z* = 233.08952 (100%, [M + Na]⁺), 152.07063 (11%). Calc. for C₁₀H₁₄N₂O₃ + Na⁺: 233.08966.

cis-N,N'-((Tetrahydrofuran-2,5-diyl)bis(methylene))diacetamide (5). Compound 4 (11.3 g, 53.8 mmol) was dissolved in 100 mL of methanol. RANEY®-nickel suspension (12.0 g, 50% p/p in water) was washed with methanol and centrifuged 3 times to completely remove water. The catalyst was then added to the solution of 4 and the suspension was placed in a Parr hydrogenator and stirred under a hydrogen atmosphere (10 bar), checking the progress by TLC (ethyl acetate/methanol 8 : 2). After 48 h the mixture was filtered through Celite®, and the filtrate was evaporated under vacuum. The product was crystallized from methanol, collected by vacuum filtration and washed with methanol to give compound 5 as white crystals (10.7 g, 93%). M. p.: 120.0–121.9 °C. ¹H NMR (400 MHz, DMSO-d₆, 298 K) δ 7.88 (t, *J* = 5.9 Hz, 2H), 3.83 (p, *J* = 5.0 Hz, 2H), 3.19 (dt, *J* = 13.6, 6.2 Hz, 2H), 3.04 (dt, *J* = 13.6, 5.1 Hz, 2H), 1.85–1.80 (m, 8H), 1.56–1.45 (m, 2H) ppm. ¹³C NMR (101 MHz, DMSO-d₆, 298 K) δ 169.4 (C), 77.8 (CH), 42.7 (CH₂), 27.9 (CH₂), 22.6 (CH₃) ppm. HRMS (ESI⁺): *m/z* = 237.12086 (100%, [M + Na]⁺), 215.13906 (35%, [M + H]⁺). Calc. for C₁₀H₁₈O₃N₂ + H⁺: 215.13902.

cis-(Tetrahydrofuran-2,5-diyl)dimethanamine dihydrobromide (6). The diamide 5 (1.00 g, 4.67 mmol) was dissolved in 10 mL of 1.2 M aqueous hydrochloric acid and refluxed for 20 h. Then, the mixture was cooled to 0 °C, in an ice bath, and basified with sodium hydroxide. The mixture was extracted three times with dichloromethane. The pooled organic phases were dried over Na₂SO₄ and Na₂CO₃ and filtered and the solvent was removed by evaporation under reduced pressure. The residue was dissolved in 10 mL of isopropanol and 1.3 mL of 48% aqueous HBr were slowly added at 0 °C. This mixture was refluxed for 5 minutes until a solution was obtained. White crystals were formed after standing overnight at 4 °C and were collected by vacuum filtration, washed with isopropanol and diethyl ether, and dried under vacuum to give compound 6 (1.35 g, 98%). ¹H NMR (400 MHz, D₂O, 298 K) δ 4.37–4.23 (m, 2H), 3.28 (dd, *J* = 13.2, 3.2 Hz, 2H), 3.05 (dd, *J* = 13.2, 9.2 Hz, 2H), 2.30–2.17 (m, 2H), 1.85–1.73 (m, 2H) ppm. ¹³C NMR (101 MHz, D₂O, 298 K) δ 76.1 (CH), 43.6 (CH₂), 28.1



(CH₂) ppm. HRMS (ESI⁺): *m/z* = 131.11804 (100%, [M + H]⁺), 114.09148 (32%). Calc. for C₆H₁₄N₂O + H⁺: 130.11061.

cis-(Tetrahydrofuran-2,5-diyl)dimethanamine-*N,N,N',N'*-tetra-acetic acid, tetra-*t*-butyl ester (7). The diamine dihydribromide 6 (1.04 g, 3.54 mmol) was suspended in 10 mL of acetonitrile and potassium carbonate (2.94 g, 21.3 mmol) was added. Then, *t*-butyl bromoacetate (2.7 mL, 17.7 mmol) was added very slowly, while the mixture was cooled to 0 °C with an ice bath. Then, the reaction mixture was stirred at room temperature and checked by TLC (petroleum ether/ethyl acetate 8 : 2). After 18 h, the reaction mixture was filtered, and the filtrate was evaporated under reduced pressure. The crude product was purified by gravity-flow chromatography on a silica-gel column (petroleum ether/ethyl acetate 8 : 2) to give compound 7 as a yellow oil (1.42 g, 68%). ¹H NMR (400 MHz, CDCl₃, 298 K) δ 4.00–3.93 (m, 2H), 3.51 (d, *J* = 17.3 Hz, 4H), 3.42 (d, *J* = 17.4 Hz, 4H), 2.87 (dd, *J* = 13.5, 6.3 Hz, 2H), 2.71 (dd, *J* = 13.6, 4.8 Hz, 2H), 1.93–1.85 (m, *J* = 4.3 Hz, 2H), 1.65–1.56 (m, 2H), 1.43 (s, 36H) ppm. ¹³C NMR (101 MHz, CDCl₃, 298 K) δ 171.0 (C), 80.8 (C), 79.3 (CH), 59.0 (CH₂), 57.0 (CH₂), 29.3 (CH₂), 28.3 (CH₃) ppm. HRMS (ESI⁺): *m/z* = 609.37145 (100%, [M + Na]⁺), 587.38981 (63%, [M + H]⁺), 473.32211 (10%), 214.08949 (13%). Calc. for C₃₀H₅₄N₂O₉ + H⁺: 587.39021.

cis-(Tetrahydrofuran-2,5-diyl)dimethanamine-*N,N,N',N'*-tetra-acetic acid (EHDTA). The diaminotetraester 7 (1.42 g, 6.25 mmol) was dissolved in 15 mL of trifluoroacetic acid and refluxed for 24 h. Then, the solvent was removed by evaporation under reduced pressure, the residue was digested three times in diethyl ether and the solvent was discarded. The product was dried under vacuum to give EHDTA as an off-white solid (0.600 g, 68%). ¹H NMR (400 MHz, D₂O, 298 K) δ 4.48–4.42 (m, 2H), 4.24 (d, *J* = 17.3 Hz, 4H), 4.18 (d, *J* = 17.2 Hz, 4H), 3.66 (d, *J* = 13.4 Hz, 2H), 3.45 (dd, *J* = 13.6, 10.0 Hz, 2H), 2.23–2.16 (m, 2H), 1.74–1.66 (m, 2H) ppm. ¹³C NMR (101 MHz, D₂O, 298 K) δ 168.6 (C), 74.8 (CH), 59.6 (CH₂), 55.6 (CH₂), 28.2 (CH₂) ppm. HRMS (ESI[−]): *m/z* = 361.12547 (100%, [M – H][−]). Calc. for C₁₄H₂₂N₂O₉ – H⁺: 361.12525.

Equilibrium studies

Materials. The chemicals used for the experiments were of the highest analytical grade. The concentrations of the MgCl₂, CaCl₂, SrCl₂, ZnCl₂, CuCl₂ and LnCl₃ solutions (Fluka) were determined by complexometric titration with standardized Na₂H₂EDTA and xylanol orange (ZnCl₂ and LnCl₃), murexide (CuCl₂), Patton & Reeder's (CaCl₂), methylthymol blue (SrCl₂) and Eriochrome Black T (MgCl₂) as indicators. The concentrations of the stock solutions of H₄EHDTA and H₆TTHA were determined by pH-potentiometric titration in the presence and absence of a large (40-fold) excess of CaCl₂. The pH-potentiometric titrations were carried out with standardized 0.2 M KOH.

pH-potentiometry. The protonation constants of EHDTA and TTHA, the stability and protonation constants of Mg²⁺, Ca²⁺, Sr²⁺, Zn²⁺, Cu²⁺ and Ln³⁺ complexes formed by the EHDTA ligand were determined by pH-potentiometric titration. The metal-to-ligand concentration ratio was 1 : 1, but for the Cu²⁺

and Zn²⁺ complexes titrations were also made at a metal-to-ligand ratio of 2 : 1 (the concentration of the ligand was generally 0.002 M). For the pH measurements and titrations, a Metrohm 785 DMP Titrino titration workstation and a Metrohm-6.0233.100 combined electrode were used. Equilibrium measurements were carried out at a constant ionic strength (0.1 M KCl) in 6 mL samples at 25 °C. The solutions were stirred under a N₂ atmosphere. The titrations were made in the pH range of 1.7–11.7. KH-phthalate (pH = 4.005) and borax (pH = 9.177) buffers were used to calibrate the pH meter. For the calculation of [H⁺] from the measured pH values, the method proposed by Irving *et al.* was used.⁴¹

A 0.01 M HCl solution was titrated with the standardized KOH solution in the presence of 0.1 M KCl ionic strength. The differences between the measured (pH_{read}) and calculated pH (–log[H⁺]) values were used to obtain the equilibrium H⁺ concentration from the pH values, measured in titration experiments. The ionic product of water (pK_w) was also determined in the presence of 0.1 M KCl. The protonation and stability constants were calculated with the PSEQUAD program.⁴²

Kinetic measurements

Transmetallation reactions. The kinetic inertness of Gd(EHDTA)[−] was characterized by the rates of the exchange reactions taking place between Gd(EHDTA) and Eu³⁺ or Cu²⁺. The exchange reactions with Eu³⁺ or Cu²⁺ were studied by spectrophotometry, following the formation of the Eu³⁺ or Cu²⁺ complexes at 250 or 330 nm with a Cary 1E spectrophotometer. The concentration of the Gd(EHDTA)[−] complex was 0.1 (Cu²⁺ exch.) and 1.0 mM (Eu³⁺ exch.), while the concentration of the Cu²⁺ and Eu³⁺ was 10 to 40 times higher, in order to guarantee pseudo-first-order conditions. The temperature was maintained at 25 °C and the ionic strength of the solutions was kept constant, 0.1 M for KCl. The exchange rates were studied in the pH range of about 3.5–6.0. For keeping the pH values constant, 1,4-dimethylpiperazine (pH range 3.1–4.1), *N*-methylpiperazine (pH range 4.1–5.2) and piperazine (pH range 5.2–6.0) buffers (0.02 M) were used. The pseudo-first-order rate constants (*k*_d) were calculated by fitting the absorbance data to eqn (1):

$$I_t = (I_0 - I_p)e^{-k_d t} + I_p \quad (1)$$

where *I*_t, *I*₀ and *I*_p are the absorbance values at time *t*, the start of the reaction and at equilibrium, respectively.

Transchelation reactions. To investigate the role of endogenous ligands in the kinetic inertness of Gd(EHDTA)[−] its transchelation reactions with TTHA were monitored by ¹H NMR relaxometry in the presence of citrate, phosphate and carbonate excess. Since the relaxivity of Gd(EHDTA)[−] (*r*₁ = 7.2 mM^{−1} s^{−1}) differs considerably from that of Gd(TTHA)^{3−} (*r*₁ = 2.4 mM^{−1} s^{−1}) at 20 MHz and 298 K, the progress of the transchelation with TTHA was followed by measuring the water proton relaxation rates (1/*T*₁) of the samples with a Bruker MQ-20 spectrometer at 20 MHz and 298 K. The longitudinal relaxation time was measured by the “inversion recovery”



method ($180^\circ - \tau - 90^\circ$) with the use of ten different τ values. The measurements were made with 1.0 mM $\text{Gd}(\text{EHDTA})^-$ solution in the pH range 6.0–10.0 in the presence of 2 fold TTHA, 0–8 fold citrate, 0–9 fold phosphate and 0–30 fold carbonate excess ($[\text{GdL}]_t = 1.0 \text{ mM}$, $[\text{TTHA}]_t = 2.0 \text{ mM}$, $[\text{Cit}^{3-}]_t = 0\text{--}8 \text{ mM}$, $[\text{PO}_4^{3-}]_t = 0\text{--}9 \text{ mM}$, $[\text{CO}_3^{2-}]_t = 0\text{--}30 \text{ mM}$). Since the protonation/deprotonations of TTHA take place in the pH range of 6.0–10.0, no buffer was used to keep the pH values constant in these experiments. The ionic strength of the solutions was kept constant with KCl (0.1 M). The pseudo-first-order rate constants (k_d) were calculated by fitting the relaxivity data to eqn (1) where I_t , I_0 and I_p are the relaxivity values at time t , the start of the reaction and at equilibrium, respectively.

Relaxometric measurements

The ^1H $1/T_1$ NMRD profiles were obtained with a fast-field cycling Stelar SmartTracer relaxometer (Mede, Pavia, Italy) varying the magnetic-field strength from 0.00024 to 0.25 T (0.01–10 MHz range). The $1/T_1$ values are measured with an absolute uncertainty of $\pm 1\%$. Temperature was controlled with a Stelar VTC-91 airflow heater equipped with a calibrated copper–constantan thermocouple (uncertainty of $\pm 0.1 \text{ K}$). Data at high fields (0.5–3 T, corresponding to 20–120 MHz proton Larmor frequency) were collected with a High Field Relaxometer (Stelar) equipped with the HTS-110 3T Metrology Cryogen-free Superconducting Magnet. The measurements were performed with a standard inversion recovery sequence (20 experiments, 2 scans) with a typical 90° pulse width of 3.5 μs , and the reproducibility of the data was within $\pm 0.5\%$. The exact concentration of Gd^{III} was determined by measurement of bulk magnetic susceptibility shifts of a $t\text{BuOH}$ signal.⁴³

The ^{17}O NMR data were acquired on a Bruker Avance III spectrometer (11.7 T) using a 5 mm probe under temperature control. An aqueous solution of the complex was enriched to achieve 2.0% of the ^{17}O isotope (Cambridge Isotope). The transverse relaxation rates were measured from the signal width at half-height as a function of temperature in the 278–350 K range.

PL measurements

PL measurements were performed with a Horiba Jobin-Yvon Fluorolog-3 spectrofluorimeter. Steady state spectra were acquired using a xenon DC lamp source and time-resolved data were collected with a SpectraLED S-280 pulsed light source. Appropriate optical filters were used. The solid state powder samples used for PL measurements were obtained by lyophilisation of a solution of $\text{Eu}(\text{EHDTA})^-$ in water, after the removal of excess Eu^{3+} by precipitation as hydroxide. Average decay time constants of biexponential curves were calculated as amplitude–average lifetimes and intensity–average lifetimes $\tau_{\text{average}} = \sum_i A_i \tau_i^2 / \sum_i A_i \tau_i$ where A_i is the amplitude associated with the i -th time constant τ_i (see the ESI† for details).

XRD measurements

Single crystal X-ray diffraction (SC-XRD) data were collected with a Smart APEXII CCD area-detector diffractometer (BRUKER). The radiation source was a molybdenum anode ($\text{Mo-K}\alpha$, $\lambda = 0.71073 \text{ \AA}$) with the generator working at 50 kV and 30 mA. The data reduction was carried out with CrysAlis Pro⁴⁴ version 1.171.42.60a using an empirical absorption correction with spherical harmonics (SCALE3 ABSPACK). The structure was solved by dual space methods with SHELXT-2015⁴⁵ and refined with SHELXL-2018⁴⁶ using the WinGX program suite.⁴⁷ Structure refinement was done using full-matrix least-square routines against F^2 . All hydrogen atoms, besides the ones in the ligand, were refined using soft restraints. The pictures were generated with the programs ORTEP-3⁴⁷ (representation with anisotropic displacement parameters), and MERCURY⁴⁸ (hydrogen bonding network). Hydrogen atoms were omitted for clarity in these representations. CCDC 2235393 contains the supplementary crystallographic data for the structure of $[\text{C}(\text{NH}_2)_3]_5[\text{Gd}(\text{EHDTA})(\text{CO}_3)] \cdot 2\text{H}_2\text{O}$.†

Conclusions

In this work, we have considered a new OBETA-derived ligand featuring the incorporation of a tetrahydrofuran ring in the backbone and analysed various physicochemical properties of the corresponding Ln^{3+} complexes. The main objectives of this study were two: (i) to explore a green approach to the synthesis of multidentate ligands for Ln^{3+} ions for biomedical applications; (ii) to understand the impact on the properties of the complexes derived from the introduction of a stereochemical rigidity element in the chelator.

The synthetic procedure based on the use of a biomass-derived chemical has proved effective and convenient. This initial example should stimulate new and multiple approaches along this direction, with the aim of substantially reducing the environmental impact of this class of diagnostic probes that is widely and increasingly used.

The presence of the tetrahydrofuran ring in the ligand backbone has a marginal influence on the thermodynamic stability of Ln^{3+} complexes with the Ln^{3+} ion from the first half of the series. However, the $\log K_{\text{LnL}}$ values of the $\text{Ln}(\text{EHDTA})^-$ complexes formed with the heavier Ln^{3+} ions are higher than that of the corresponding $\text{Ln}(\text{OBETA})^-$ complexes. The tetrahydrofuran ring in the skeleton of the EHDTA ligand slows down the proton assisted dissociation of $\text{Gd}(\text{EHDTA})^-$ due to the more rigid coordination environment of the Gd^{3+} ion. The endogenous ligand (citrate, phosphate and carbonate) can accelerate the dissociation of $\text{Gd}(\text{EHDTA})^-$ via the formation $[\text{Gd}(\text{EHDTA})X]$ intermediates ($X = \text{HCit}^-, \text{HCO}_3^-$ and H_2PO_4^- anions). The dissociation reactions of $[\text{Gd}(\text{EHDTA})X]$ intermediates take place with the fast rearrangement and the proton transfer processes, whereas the dissociation of $\text{Gd}(\text{OBETA})^-$ occurs mainly by the general acid catalysed pathway through proton transfer from the protonated $\text{HCit}^-, \text{HCO}_3^-$ and H_2PO_4^- anions to OBETA in the ternary $[\text{Gd}(\text{OBETA})X]$ adducts.



On the other hand, unlike the case of $[\text{Gd(OBETA)}]^-$, the multifrequency and multinuclear NMR relaxometric study showed that $[\text{Gd(EHDTA)}]^-$ exists in solution as a mixture of two species that differ in the hydration state, *i.e.* $q = 1$ and $q = 2$. While the bis-hydrated species is prevalent, the presence of a detectable population of the mono-hydrated species signals a certain degree of distortion in the molecular structure which is confirmed by the X-ray diffractometric study of the Gd^{3+} complex. This effect, associated with the higher degree of stereochemical rigidity in EHDTA, should have an increasing impact along the series following the contraction of the ionic radius of Ln^{3+} ions.

In conclusion, EHDTA appears to be a ligand with good affinity towards trivalent lanthanoids ions, giving rise to complexes with good thermodynamic stability and acceptable kinetic inertness. These features make the $[\text{Ln(EHDTA)}]^-$ complexes good candidates in preclinical studies of molecular bioimaging.

Author contributions

Conceptualization, project administration and supervision: M. B., G. B. G. and Z. B.; funding acquisition: M. B., G. B. G., and J. B.; preparation of the chelating agent and complexes: F. T. and M. L. M.; relaxometric studies: M. B. and M. L. M.; luminescence studies: F. A.; complexometric studies: Z. B. and J. B.; diffractometric studies: T. G.; writing and finalization of the manuscript: all authors.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This publication is part of the project NODES which has received funding from the MUR – M4C2 1.5 of PNRR with grant agreement no. ECS00000036 (M. B., G. B. G., F. T.). J. B. acknowledges the financial support of the János Bolyai Research Scholarship of the Hungarian Academy of Sciences and of the Hungarian National Research, Development and Innovation Office (FK-134551) project, the New National Excellence Program ÚNKP-22-5.

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