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Substituent effects on fluoride binding by lanthanide complexes of DOTA-tetraamides†

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Fluoride binding by a series of europium and ytterbium complexes of DOTA-tetraamide ligands derived from primary, secondary and tertiary amides has been studied by NMR and luminescence spectroscopies. In all the systems studied, fluoride binding results in a change in the nature of the magnetic anisotropy at the metal centre from an easy axis, to an easy plane anisotropy. This results in reversal of the peaks in the NMR spectra, and in changes to the fine structure of the luminescence spectra. Furthermore, changes to the periphery of the binding cavity are implicated in determining the affinity constant for fluoride. There are clear differences in the entropic contribution to the free energy of activation between systems with benzylic amides and those with methylamides.

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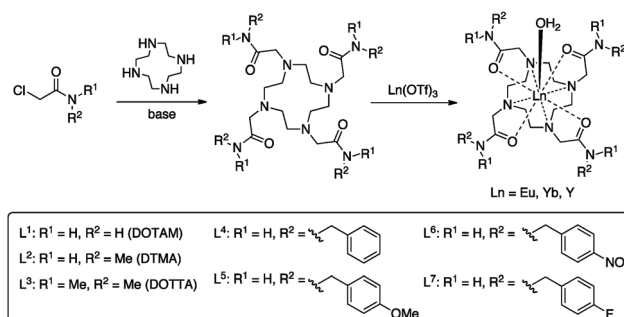
The spectroscopic and magnetic properties of lanthanide complexes have been widely exploited over the course of many years, particularly in magnetic resonance imaging contrast agents and in time-resolved bioassays, and more recently in the field of molecular magnetism. In all of these areas, optimisation of the properties of these lanthanide containing systems is contingent on the fundamental understanding of their electronic structure, and the interaction of open shell f-electrons with their surroundings.

Bleaney's approach to understanding the effect of lanthanide magnetism on surrounding nuclear spins has underpinned the development of the field for almost fifty years.¹ Defining these theories required a number of approximations to be made; and, while many have pointed out discrepancies between theoretical prediction and experimental observations (particularly in low symmetry systems), it is only recently that supporting theoretical methods have become available to study and interpret more unusual aspects of lanthanide behaviour in molecular systems.

We recently investigated the binding of fluoride ions to lanthanide complexes of the tetraamide ligand DTMA and determined that the replacement of water with fluoride induces a change in the nature of the magnetic anisotropy and

in the sign of the crystal field parameter $B_0^{2,2,3}$. The change resulted in dramatic changes to NMR, EPR and luminescence spectra, and the phenomenon was further understood using *ab initio* calculations. These results, taken together with a number of recent studies on single molecule magnets⁴ and luminescent complexes,⁵ provide a rapidly growing body of data that reveal the subtleties inherent to the magnetic behaviour of lanthanide complexes.

In this manuscript we seek to determine how the nature of the ligand affects the fluoride-binding event in symmetric DOTA tetraamide complexes. To this end we have synthesised and studied the fluoride binding by a number of complexes related to DTMA (L^1 , L^3 – L^7), shown in Scheme 1. This array of related complexes has allowed us to explore the effect of variations in amide structure and hydrophobicity on fluoride binding and its spectroscopic consequences.



Scheme 1 Synthesis of the complexes studied.

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† Electronic supplementary information (ESI) available: Synthetic procedures, Bleaney plots, $\chi_{||}$ and χ_{\perp} values, EXSY spectra, $\text{Eu}^{\text{X}}\text{-}^1\text{H}$ NMR spectra, emission spectra, binding isotherms, $\text{Lu}^{\text{X}}\text{-}^1\text{H}$ NMR spectra, inversion recovery plots, exchange rate data. See DOI: 10.1039/c5dt04394b

Results and discussion

Synthesis

A series of tetraamide ligands derived from a variety of different amines, were synthesised by the well-established procedure outlined in Scheme 1. The chloroacetamides were synthesised from the appropriate primary or secondary amine and reacted with cyclen to give the 8-coordinate ligands. The complexes were formed by reaction of the ligands with the appro-

priate lanthanide triflate salts and characterised by NMR, mass spectrometry and CHN analysis (see ESI†).

NMR studies

Proton NMR studies focussed on the Yb^{3+} complexes of ligands L^{1-7} due to the dominance of the pseudocontact shift for this ion. The ^1H NMR spectra in D_2O , and the effect of addition of fluoride on the spectra are shown in Fig. 1. All of the Yb^{3+} complexes apart from YbL^3 appear to only be present as the SAP



Fig. 1 ^1H NMR of complexes YbL^{1-7} in the absence (black) and presence (red) of an excess of sodium fluoride (D_2O , 400 MHz, 298 K).



isomer. YbL³ contains approximately 7% TSAP in D₂O at 298 K. Upon addition of fluoride, all Yb complexes display a new set of peaks in slow exchange with the original set, as originally observed with YbL² (YbDTMA).² In all cases, EXSY spectra show a reversal of the peak order between hydrated and fluoride-bound forms, implying a change of sign of B_0^2 (see ESI†). In our earlier studies, we rationalised these observations by a change in the order of the m_J states arising from ligand field splitting of the $^2F_{7/2}$ ground state giving rise to a change in the nature of the magnetic anisotropy at the lanthanide centre, and it is reasonable to infer that this phenomenon is general across this series of complexes. The observation of such dramatic changes to the NMR spectra upon fluoride coordination must be explained in terms of changes to the Boltzmann populations within the m_J manifold as a result of changing the relative energies of the various m_J states, and it is clear that the overall anisotropy of the metal ion must reflect all of these states.

Inspection of the spectra in Fig. 1 reveals that while the range of chemical shifts observed for the hydrated complexes are fairly similar, those for the fluoride-bound species show significant differences between complexes. Differences in the spread of observed ^1H chemical shifts between complexes reflect the relative magnitudes of the axial magnetic anisotropy resulting from the effects of different ligand fields. For systems with axial symmetry and negligible contact shift contribution, the effect of the crystal field on the observed shifts is conveniently expressed using the Bleaney equation¹ and the crystal field parameter, B_0^2 :

$$\delta_{\text{PC}} = \frac{2C_J\beta^2(3\cos^2\theta - 1)}{(kT)^2} \frac{B_0^2}{r^3} \quad (1)$$

The crystal field coefficient, C_J , and the parameters B_0^2 , β , k and T are often grouped together into one parameter labelled D_1 to give:

$$\delta_{\text{PC}} = D_1 \frac{(3\cos^2\theta - 1)}{r^3} \quad (2)$$

Plots of δ_{PC} vs. $(3\cos^2\theta - 1)/r^3$, yield lines with gradients corresponding to D_1 (eqn (2)), which are proportional to B_0^2 . δ_{PC} are approximated from δ_{obs} by subtraction of 2.9 ppm (the average shift for the protons on the ligand) to take account of the diamagnetic contribution and $(3\cos^2\theta - 1)/r^3$ values are taken from closely related crystal structures.^{6,7} These experimentally derived D_1 values are given in Table 1 for each

complex in both hydrated and fluoride-bound forms (plots are given in ESI†) along with the difference between the two. The reported values are uncorrected for changes in the bulk magnetic anisotropy with the lanthanide: the same method was used for both the fluoride-free and fluoride bound systems, and the relative change between the two can be assumed to be correct.

For the hydrated species, the magnitudes of D_1 fall within a relatively narrow range. Observed differences correlate with variations in electron demand, although solvation is also expected to play a role. The nature of the *para*-benzyl substituent has a small but significant influence on the crystal field, with the NO₂ group giving the smallest B_0^2 of the benzyl-substituted complexes.

It is possible to use the data in Table 1 to estimate the degree of anisotropy in each of the complexes. Since:

$$\delta_{\text{PC}} = \frac{1}{2N_A} \left[\frac{(3\cos^2\theta - 1)}{r^3} (\chi_{\parallel} - \chi_{\text{av}}) \right] \quad (3)$$

and

$$\chi_{\text{av}} = (\chi_{\parallel} + 2\chi_{\perp})/3 \quad (4)$$

It is possible to define both χ_{\parallel} and χ_{\perp} from D_1 provided we know χ_{av} . In previous studies, we modelled χ_{av} for Yb-L²-OH₂ and for Yb-L²-F⁻ obtaining values for $\chi_{\text{av}}T$ at 300 K of 2.49 and 2.51 cm³ mol⁻¹ K respectively, while the free ion χ_{av} would be expected to be 2.57 cm³ mol⁻¹ K.² In our previous study, theoretical models were also used to support the hypothesis that axial ligand exchange (*i.e.* replacing water with fluoride) did not alter the structure of the macrocyclic ligand significantly. In the case of this work, it is reasonable to assume that axial ligand exchange occurs- since substitution of a fluoride ligand for one of the other donor atoms to Ln would break the observed symmetry inferred from the NMR spectra.

Thus the crystal field of the ligands undoubtedly has an effect upon the value of χ_{av} , albeit a small one. For the purposes of this study, a qualitative map of the anisotropy can be obtained by estimating χ_{av} as 2.5 cm³ mol⁻¹ K for the systems studied in this manuscript. The results of applying this approach can be seen in Fig. 2, which represents the magnetic susceptibilities as a series of spheroids in which the z -axis defines the molecular axis. Values for χ_{\parallel} and χ_{\perp} are tabulated in the ESI† to this paper. From Fig. 2, it should be clear that the anisotropy takes very different forms in the water-bound and fluoride-bound complexes, while much smaller (though still significant) differences in anisotropy are observed between complexes with different macrocyclic ligands. It should be noted that, at the ambient temperatures studied in this work, free rotation about the Ln-O bond in the aquated complex results in averaging of χ_X and χ_Y on the timescale of the NMR experiment (meaning that the anisotropy can be treated as a spheroid rather than an ellipsoid on these timescales).

YbL¹ and YbL² behave similarly on addition of fluoride. YbL³-F has anomalous chemical shift ranges and a much less

Table 1 D_1 values (ppm Å³) from linear fits to the Bleaney plots of YbL¹⁻⁷ for the hydrated and fluoride bound complexes using ligand ^1H resonances, and ΔD_1 for each complex

	YbL ^x -OH ₂	YbL ^x -F	ΔD_1
YbL ¹	3027	-791	3818
YbL ²	3216	-890	4106
YbL ³	3613	-356	3969
YbL ⁴	3382	-1350	4732
YbL ⁵	3329	-1379	4708
YbL ⁶	2976	-1666	4642
YbL ⁷	3234	-1404	4638



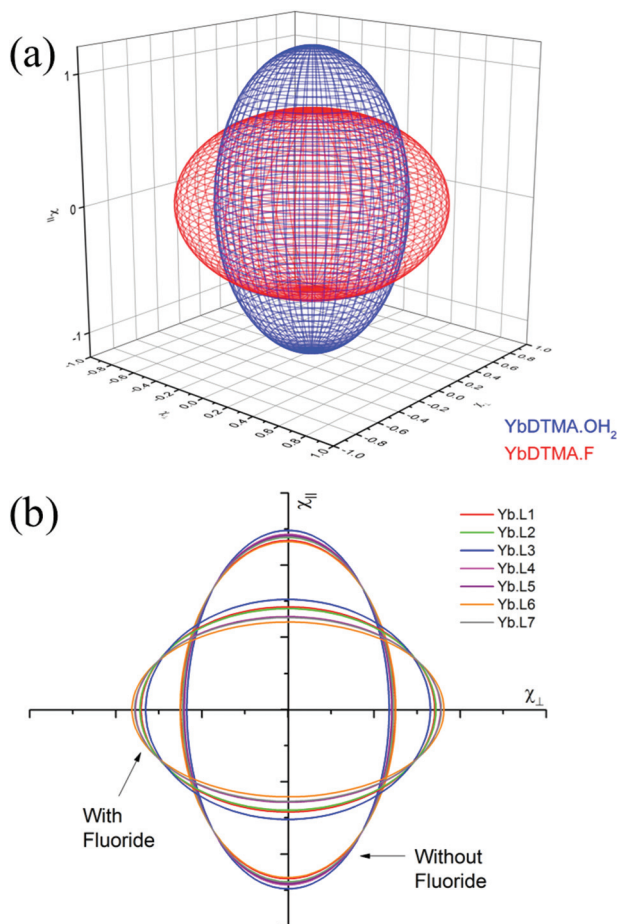


Fig. 2 Representations of the anisotropy in $\chi_{||}$ and χ_{\perp} . (a) Representing the spheroidal anisotropy with $\chi_{||}$ on the z-axis- and χ_{\perp} on the x- and y-axes for $\text{YbL}^2\text{-OH}_2$ and $\text{YbL}^2\text{-F}$; (b) Two dimensional representations for all the complexes studied, showing slices through the spheroids in the xz plane at $y = 0$.

negative D_1 , which are likely to be a consequence of YbL_3 being the only tertiary tetraamide studied. The change in slope of the Bleaney plot on binding fluoride is however similar to $\text{YbL}^{1,2}$, being in the range 3800–4100 ppm \AA^3 , so although the magnetic anisotropy in $\text{YbL}^3\text{-F}$ is smaller than the other complexes, this is due to the axial fluoride offsetting a stronger equatorial ligand field. The fluoride bound benzylamide derivatives, YbL^{4-7} , have more negative D_1 than YbL^{1-3} while all show changes in D_1 in the range 4600–4700 ppm \AA^3 . This would argue that exchange of water for fluoride has a greater impact on the crystal field for YbL^{4-7} compared with YbL^{1-3} . This is likely to be a consequence of reduced solvation of the fluoride in the binding cavity making the fluoride a better donor. The observed value of D_1 does, however, vary from complex to complex as a consequence of the electronic influences of the peripheral substituents on the benzyl groups. $\text{YbL}^6\text{-F}$ has a more negative gradient than the other benzylamides, however this is a consequence of a weaker equatorial ligand field (as indicated by the trend in the series of hydrated benzylamide complexes).

Analogous effects of fluoride binding are also observed in the NMR spectra of the europium complexes, although chemical shifts are significantly affected by the contact shift and so eqn (1) cannot be employed. Furthermore, EuL^{1-3} are present as mixtures of SAP and TSAP isomers in solution.⁷ The exchange between isomers in these cases causes broadening of the ^1H NMR spectra and cooling is required to distinguish the peaks. The benzyl-substituted $\text{Eu}(\text{III})$ complexes however display much sharper NMR spectra since only the SAP isomer appears to be present (see ESI†), as has been previously noted for larger secondary substituents of this nature.⁸ EuL^3 is a mixture of SAP and TSAP in *ca.* 1 : 2 ratio and two new sets of ^1H signals appear on addition of fluoride, in a ratio of *ca.* 1 : 3 (Fig. 3), presumably corresponding to fluoride bound SAP and TSAP isomers.

During our previous study we observed the ^{19}F signals corresponding to bound fluoride for several different lanthanide complexes of ligand $\text{L}^{2,3}$. In the case of the pseudo-lanthanide yttrium the signal is a doublet due to coupling between ^{19}F and ^{89}Y nuclei, as confirmed by HMQC.³ Bound fluoride resonances are reported here for Y^{3+} , Eu^{3+} and Yb^{3+} complexes (Table 2). In the case of EuL^3 , two bound fluoride signals are observed and integration indicates that the peak at –500 ppm (298 K) corresponds to the major isomer of the fluoride-bound form.

The yttrium complexes allow us to assess the diamagnetic contributions to the ^{19}F chemical shift. Complexes of L^{1-3} have bound fluoride shifts of a similar magnitude to one another, while complexes of L^{4-7} give less negative bound fluoride



Fig. 3 NMR spectra of EuL^3 (D_2O , 500 MHz, 278 K). Top: ^1H NMR in the absence (black) and presence (red) of an excess of sodium fluoride; Bottom: ^{19}F NMR in the presence of an excess of sodium fluoride.



Table 2 Chemical shifts of bound fluoride (ppm) in D₂O at 298 K, coupling constants are given in brackets for Y complexes (Hz)

	Y	Eu	Yb
L ¹	−62 (59)	−487	−833
L ²	−59 (61)	−481	−859
L ³	−60 (57)	−414, −500	−721
L ⁴	−54 (65)	−483	−993
L ⁵	−54 (65)	−479	−1016
L ⁶	−52 (66)	−477	−1098
L ⁷	−54 (66)	−483	−1023

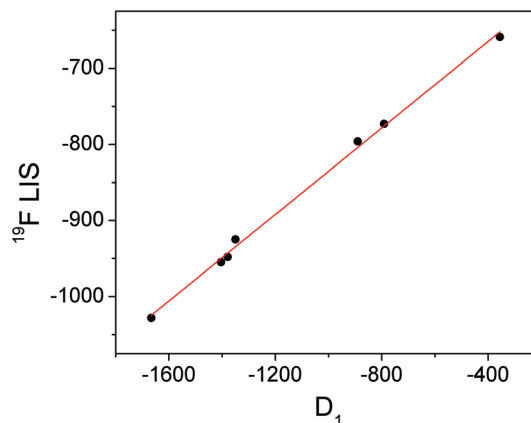
shifts. Y–F coupling constants are also separated into these two groups. The larger coupling constants and less negative chemical shifts (a larger shift from free fluoride) would imply a significant difference to the nature of the interaction between fluoride and the benzylamide-based complexes.

In the paramagnetic complexes, the diamagnetic contribution to the shift is small relative to the effect of the paramagnetic metal centres. The europium complexes have bound fluoride chemical shifts that are broadly similar, with the exception of EuL³. Here the lanthanide induced shift (LIS) is dominated by the contact shift, especially considering that the fluoride is directly coordinated to the metal. The phenomenon of spin–spin coupling between fluorine and spatially adjacent nuclei by virtue of orbital overlap (“through-space couplings”) is well known and has been extensively studied.⁹ Note that this does not imply any “formal” bond between the fluoride ion and the metal, nor does it necessarily tell us anything about the contribution of the contact shift for other nuclei (such as protons) in the complex. The similarity of the lanthanide induced fluoride shifts therefore indicates that the contact shift felt by the fluoride ion is relatively constant across the series of complexes studied and implies that the nature of the contact interaction is insensitive to the substituents.

Conversely, the bound fluoride shifts of the ytterbium complexes are dominated by the pseudocontact shift,³ and therefore vary with the size of the crystal field coefficient. A plot of D_1 for the fluoride-bound Yb³⁺ complexes taken from Table 1 vs. LIS of the ytterbium bound fluoride gives a straight line (Fig. 4). The shifts of the yttrium analogues are used to subtract the diamagnetic contributions. The linear correlation would imply that the geometric factor, $(3 \cos^2 \theta - 1)/r^3$, remains constant across the series of complexes and since we can assume that θ is also constant, we can infer that the distance between fluoride and lanthanide does not vary with the identity of the ligand.

Luminescence studies

We also studied the effects of fluoride binding on the luminescence properties of the europium complexes since information regarding the ligand field can readily be extracted from fine structure.¹⁰ Changes to the shape of the spectrum are observed in all bands (see ESI†) upon addition of fluoride. The $\Delta J = 1$ band is the most appropriate to analyse since in axial

**Fig. 4** Graph of D_1 (derived from Bleaney plots of ^1H shifts of YbL^{1–7}) vs. the observed LIS of bound fluoride for YbL^{1–7} with linear fit in red.

symmetry we would expect the band to consist of two peaks as a result of the $2J + 1$ degeneracy being lifted by the ligand field. The parameter B_0^2 can be extracted directly from the splitting of this band. The fine structure of the $\Delta J = 1$ bands in the presence and absence of fluoride are shown for EuL^{1–7} in Fig. 5.

In the absence of fluoride, the $\Delta J = 1$ regions of EuL^{1–3} appear to consist of more than two transitions, especially for EuL³ which displays a particularly broad band. For EuL^{4–7}, $\Delta J = 1$ is split into two easily distinguishable peaks. This correlates with the presence of two conformational isomers with significantly different ligand field splitting, SAP and TSAP, for EuL^{1–3} and the predominance of SAP in EuL^{4–7}. Therefore, while the shape of the $\Delta J = 1$ band for EuL^{1–3} is complicated by the presence of multiple isomers, the shape of this band for EuL^{4–7} is determined only by B_0^2 of the SAP isomer. It is instantly apparent from the splitting of $\Delta J = 1$ that L⁶ invokes a smaller crystal field splitting than L^{4,5,7}, which correlates with the ^1H NMR studies of the Yb³⁺ complexes (Table 1).

Upon addition of fluoride, the emission spectra of the Eu complexes change shape in accordance with a change in the ligand field (see ESI†). The change is similar for all complexes with a decrease in the magnitude of splitting within the $\Delta J = 1$ band (Fig. 5). For EuL^{1–3} the $\Delta J = 1$ peaks merge into one, corresponding to the reduction in the magnitude of B_0^2 in the fluoride bound complexes. EuL³ displays a particularly narrow band, which correlates with the small magnitude of D_1 of YbL³ compared with YbL^{1–2} (Table 1). The magnitude of the splitting is unresolvable in the EuL^{1–3} emission spectra.

For the benzyl substituted complexes in the presence of fluoride, the splitting of $\Delta J = 1$ is resolvable and is small compared with the original complex, in line with the observations from ^1H NMR of the Yb complexes. The fact that the splitting is resolvable for EuL^{4–7} but not EuL^{1–3} tallies with the larger magnitude of D_1 observed for the former (Table 1). The peak separation is larger for EuL⁶ than EuL^{4–5} in line with the more negative D_1 for the former. Furthermore, while it is not possible to clearly identify the A and E components of the $\Delta J = 1$



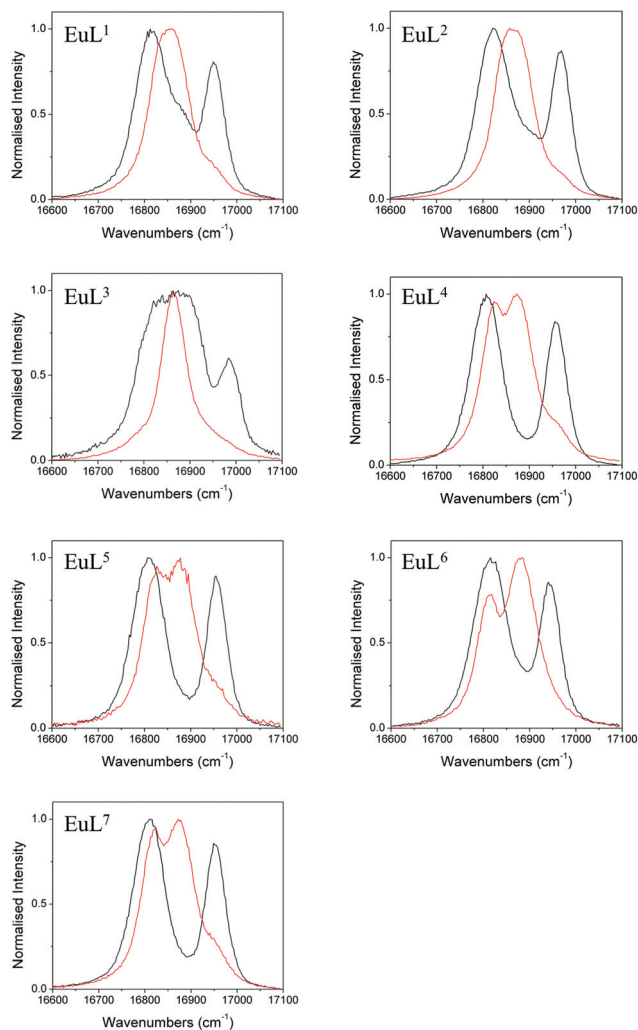


Fig. 5 Normalised emission bands for the 5D_0 – 7F_1 transitions of EuL^{1-7} in the absence (black) and presence (red) of an excess of sodium fluoride (exit slit = 0.2 nm).

band in $\text{EuL}^{4,5,7}$, for EuL^6 the transition from 5D_0 to the doubly degenerate E level of 7F_1 (which appears broader in the spectrum) is now at higher energy than the A level. The energetic ordering of the A and E levels is thus reversed with respect to the hydrated complex, presenting further direct evidence of a change in sign of B_0^2 .

The luminescence lifetimes of the Eu-based emission were measured in the presence and absence of fluoride in both H_2O and D_2O and are given along with q values in Table 3.¹¹ Lifetimes are lengthened on addition of fluoride corresponding to a change in the number of bound water molecules from one to zero, confirming the displacement of water by fluoride in each case.

Association constants

The association constants for fluoride with EuL^{1-7} were determined both by tracking changes in the emission spectra and by following changes in ^1H and/or ^{19}F NMR intensities as a

Table 3 Luminescence lifetimes ($\lambda = 616$ nm) with and without excess fluoride in H_2O and D_2O with associated q values

	$\text{EuL}^x\text{-OH}_2$			$\text{EuL}^x\text{-F}$		
	$\tau_{\text{D}_2\text{O}}$ (ms)	$\tau_{\text{H}_2\text{O}}$ (ms)	q	$\tau_{\text{D}_2\text{O}}$ (ms)	$\tau_{\text{H}_2\text{O}}$ (ms)	q
EuL^1	2.22	0.51	1.0	2.50	0.96	0.0
EuL^{2a}	1.72	0.54	0.9	2.41	1.07	0.0
EuL^3	1.88	0.61	1.0	2.29	1.34	0.1
EuL^4	2.27	0.59	0.8	2.60	1.01	0.1
EuL^5	2.31	0.57	0.9	2.61	1.10	0.0
EuL^6	2.17	0.54	1.0	2.37	1.07	0.0
EuL^7	1.92	0.54	0.9	2.56	1.13	0.0

^a Previously published.³

Table 4 Association constants (M^{-1}) for EuL^{1-7} with sodium fluoride in D_2O at 298 K (unless otherwise stated) measured by luminescence and ^1H and/or ^{19}F NMR spectroscopies. 95% confidence intervals are given in parentheses

	Luminescence	NMR
EuL^1	90.9 [82.5–100.4]	82.8 [72.7–94.7] ^a
EuL^2	59.9 [57.1–62.7]	57.5 [47.7–69.5] ^b
EuL^3	33.3 [31.4–35.3]	17.7 [16.0–19.5] ^c
EuL^4	21.0 [19.5–22.6]	21.7 [19.3–24.4] ^a
EuL^5	20.8 [18.2–23.8]	19.9 [16.8–23.5] ^a
EuL^6	83.4 [77.4–89.9]	85.2 [73.5–99.6] ^a
EuL^7	40.7 [37.9–43.9]	43.7 [37.8–50.7] ^d

^a Using ^1H NMR data. ^b Previously published.³ ^c At 278 K using ^1H and ^{19}F data from the same titration. ^d Following the ^{19}F resonance of the ligand.

function of fluoride concentration, which provide a direct measure of the concentrations of each species. The data was fitted using a one to one binding model in Dynafit¹² and the values are given in Table 4 with associated confidence intervals. NMR titrations provided K values within error of those obtained from the luminescence titrations.

The titration data shows that there are significant differences in K with varying the ligand substituents, although they are of a similar magnitude. As we incorporate methyl groups from L^1 to L^3 , the K value decreases. Similarly, as the electron withdrawing nature of the benzyl substituent is increased, the K value increases. This would indicate that residual charge on the metal plays a major role in determining the association constant. Correlation of $\ln K$ with the D_1 values (for the hydrated Yb^{3+} complexes) from Table 1 reveals a broadly negative correlation (Fig. S17†), although a lack of linearity would imply that there are additional factors contributing. The introduction of hydrophobic benzyl substituents does not appear to have a significant effect of the magnitude of K .

Rates of exchange

We previously determined that the rate of exchange of fluoride at the metal centre is relatively slow compared with, for



example, water exchange. Qualitative information about the relative rates of exchange can be gleaned by observing the change in the proton NMR spectra of the Lu^{3+} complexes as the fluoride concentration is increased. Here the differences in chemical shifts of the aliphatic protons are small between complexes unlike in the paramagnetic complexes. Furthermore, exchange is not complicated by the presence of multiple isomers. The exchange rate of fluoride at the lanthanide appears fast in LuL^{1-3} (see ESI† and ref. 3†) since all ^1H peaks are seen to shift upfield with increasing fluoride concentration. Whereas in LuL^4 the exchange rate is slower, with the spectra exhibiting coalescence and line shape changes upon heating in the “intermediate” exchange regime (see ESI†).

A quantitative assessment of the exchange rate is achieved by using a selective inversion NMR technique appropriate for a system in slow exchange. In a sample containing a mixture of the fluoride-bound and hydrated complexes, the effects of exchange on a pair of resonances is monitored following selective inversion of one of the signals. The evolution of the magnetisations is governed by both the exchange rate (k) and the spin-lattice relaxation rates ($R_1 = 1/T_1$). A second experiment monitors the magnetisation following a non-selective inversion pulse and fitting of all data is performed by varying k and R_1 using the CIFT2 program.¹³ In order to extract k , an appropriate pair of resonances is required and in this case the main difficulty is that R_1 is very fast for nuclei near the paramagnetic centres. The ^{19}F nuclei of the L^7 ligand are ideal for this study since they are far enough from the lanthanide to have reasonable R_1 s and other ^{19}F resonances do not overlap. The methyl protons of L^2 have much faster R_1 s than the ^{19}F of L^7 , but fortunately the rate of exchange is large enough to compensate in the case of YbL^2 .

The rate of fluoride exchange measured with EuL^7 is significantly slower than with YbL^7 at 298 K (Table 5). This tallies with our previous observation that the process is faster for LuL^2 compared with $\text{YL}^{2,3}$ and indicates that the difference in rates is linked with the ionic size of the lanthanide ion. The ligand is able to envelop the smaller metal ions (Yb^{3+} , Lu^{3+}) more extensively, accelerating the dissociation of the axial ligand, as suggested by Aime *et al.* with DOTA complexes.¹⁴

There is also a significant difference in k measured between Yb^{3+} complexes of ligands L^2 and L^7 , with the aromatic substituents conferring a slower rate of exchange between species. This corroborates our qualitative observations with Lu complexes of L^2 and L^4 above. A similar observation is made for the exchange rate of water at lanthanide centres in related



Fig. 6 Eyring plots for YbL^2 and YbL^7 in D_2O for the exchange of fluoride-bound and hydrated complexes.

complexes— hydrophobic groups tend to slow the rate of water exchange.¹⁵ In order to explore the origins of the ligand effect on the rate, k was measured for YbL^2 and YbL^7 at a range of temperatures (see ESI†). Eyring plots display good linear correlations and are shown in Fig. 6.

The thermodynamic parameters extracted from the slopes and intercepts of the plots are given in Table 5. It is clear from the data that the ΔH^\ddagger values for the two complexes are the same within error and thus that the electrostatic interaction between Yb^{3+} and F^- is independent of the ligand framework in this instance. However, the ΔS^\ddagger values extracted from the plots are significantly different, with the benzyl-appended complex having a more negative value. This implies that the interaction of fluoride with lanthanide tetraamide complexes is appreciably affected by the nature of the ligand substituents and that the rearrangement of solvent in the vicinity of the metal centre plays a key role. The change in solvation around the complex decorated with hydrophobic groups incurs a greater entropic cost during fluoride binding, resulting in a slower rate of exchange. Calculated ΔG^\ddagger values at 298 K are within error of one another.

Conclusions

From these results, several things become clear. Firstly, the effect of fluoride on the magnetic anisotropy of lanthanide tetraamide complexes appears to be a general phenomenon: in all the systems studied fluoride binding results in a change in the anisotropy from a prolate to an oblate electron distribution as a consequence of the effect of the axial fluoride donor atom. Further, Eyring analysis of the data shows that entropy plays a large part in defining the free energy of activation. The results highlight the complexity of the influences on the fluoride binding event, even within complexes of similar chemical

Table 5 Rates and kinetic parameters derived from Eyring plots for the exchange between fluoride-bound and hydrated complexes

	k (298 K) (s^{-1})	ΔH^\ddagger (kJ mol^{-1})	ΔS^\ddagger ($\text{J K}^{-1} \text{mol}^{-1}$)	ΔG^\ddagger (298 K) (kJ mol^{-1})
YbL^2	215.7 ± 3.0	47.2 ± 1.5	-41.8 ± 5.1	59.7 ± 3.0
YbL^7	61.2 ± 3.6	45.5 ± 0.8	-58.1 ± 2.9	62.8 ± 1.7
EuL^7	0.41 ± 0.01	—	—	—



structure, with electrostatics, sterics and solvation clearly playing intricate roles in the nature of the interaction.

This study adds further weight to the increasing body of evidence that the ligand field is important in lanthanide coordination chemistry, and that the relative populations (and indeed ordering) of the Stark sub-levels of the ground state are critical to defining the spectroscopic properties of the complexes.

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Notes and references

- (a) B. Bleaney, *J. Magn. Reson.*, 1972, **8**, 91–100; (b) B. Bleaney, C. M. Dobson, B. A. Levine, R. B. Martin, R. J. P. Williams and A. V. Xavier, *J. Chem. Soc., Chem. Commun.*, 1972, 791b–7793.
- O. A. Blackburn, N. F. Chilton, K. Keller, C. E. Tait, W. K. Myers, E. J. L. McInnes, A. M. Kenwright, P. D. Beer, C. R. Timmel and S. Faulkner, *Angew. Chem., Int. Ed.*, 2015, **54**, 10783–10786.
- O. A. Blackburn, A. M. Kenwright, P. D. Beer and S. Faulkner, *Dalton Trans.*, 2015, **44**, 19509–19517.
- (a) M.-E. Boulon, G. Cucinotta, J. Luzon, C. Degl'Innocenti, M. Perfetti, K. Bernot, G. Calvez, A. Caneschi and R. Sessoli, *Angew. Chem., Int. Ed.*, 2012, **52**, 350–354; (b) N. F. Chilton, D. Collison, E. J. L. McInnes, R. E. P. Winpenny and A. Soncini, *Nat. Commun.*, 2013, **4**, 1–7, 1AD; (c) J. D. Rinehart and J. R. Long, *Chem. Sci.*, 2011, **2**, 2078.
- (a) A. M. Funk, P. Harvey, K.-L. N. A. Finney, M. A. Fox, A. M. Kenwright, N. J. Rogers, P. K. Senanayake and D. Parker, *Phys. Chem. Chem. Phys.*, 2015, **17**, 16507–16511; (b) A. M. Funk, K.-L. N. A. Finney, P. Harvey, A. M. Kenwright, E. R. Neil, N. J. Rogers, P. Kanthi Senanayake and D. Parker, *Chem. Sci.*, 2015, **6**, 1655–1662; (c) L. Di Bari, G. Pintacuda, P. Salvadori, R. S. Dickins and D. Parker, *J. Am. Chem. Soc.*, 2000, **122**, 9257; (d) L. Di Bari and P. Salvadori, *Coord. Chem. Rev.*, 2005, **249**, 2854; (e) S. Di Pietro, S. Lo Piano and L. Di Bari, *Coord. Chem. Rev.*, 2011, **255**, 2810; (f) R. Berardozzi and L. Di Bari, *Inorg. Chem.*, 2013, **52**, 11514.
- R. S. Dickins, J. A. K. Howard, C. W. Lehmann, J. Moloney, D. Parker and R. D. Peacock, *Angew. Chem., Int. Ed., Engl.*, 1997, **36**, 521–523.
- S. Aime, A. Barge, J. I. Bruce, M. Botta, J. A. K. Howard, J. M. Moloney, D. Parker, A. S. de Sousa and M. Woods, *J. Am. Chem. Soc.*, 1999, **121**, 5762–5771.
- (a) G. Zucchi, R. Scopelliti, P.-A. Pittet, J.-C. G. Bünzli and R. D. Rogers, *J. Chem. Soc., Dalton Trans.*, 1999, 931–938; (b) M. Woods, S. Zhang, V. H. Ebron and A. D. Sherry, *Chem. – Eur. J.*, 2003, **9**, 4634–4640.
- J. Hilton and L. H. Sutcliffe, *Progr. NMR Spectrosc.*, 1975, **10**, 27.
- K. Binnemans and C. Görller-Walrand, *Chem. Phys. Lett.*, 1995, **245**, 75.
- A. Beeby, I. M. Clarkson, R. S. Dickins, S. Faulkner, D. Parker, L. Royle, A. S. de Sousa, J. A. G. Williams and M. Woods, *J. Chem. Soc., Perkin Trans. 2*, 1999, 493–504.
- P. Kuzmic, *Anal. Biochem.*, 1996, **237**, 260–273.
- A. D. Bain and J. Cramer, *J. Magn. Reson., Ser. A*, 1996, **118**, 21–27.
- S. Aime, M. Botta, M. Fasano, M. P. M. Marques, C. F. Gherdes, D. Pubanz and A. E. Merbach, *Inorg. Chem.*, 1997, **36**, 2059–2068.
- S. Aime, A. Barge, A. S. Batsanov, M. Botta, D. D. Castelli, F. Fedeli, A. Mortillaro, D. Parker and H. Puschmann, *Chem. Commun.*, 2002, 1120–1121.

