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Quantitative determination of fluoride in pure water using luminescent europium complexes†

Stephen J. Butler

Two luminescent probes [Eu.L1–2]⁺ are reported for the rapid detection of fluoride in water. Probes [Eu.L1–2]⁺ exhibit exceptional enhancements in Eu emission in the presence of fluoride, permitting its selective determination within the environmentally relevant concentration range (20–210 µM).

Fluoride is considered essential for healthy teeth and bone growth; in several countries this has led to the artificial fluoridation of water supplies. Fluoride is also introduced into water supplies through the production of phosphate containing fertilizers and aluminium processing industries. Consumption of elevated levels of fluoride in drinking water can cause dental and skeletal fluorosis, as well as acute gastric problems and kidney failure. Therefore, controlling the level of fluoride in water supplies is a global governmental concern; the maximum recommended concentration of fluoride in drinking water defined by the World Health Organization guidelines is 4 mg L⁻¹, or 210 µM. Consequently, the development of convenient analytical methods for the rapid, quantitative detection of fluoride in water samples has become an active area of research.

Traditional methods for the accurate analysis of fluoride involve the use of ion-selective electrodes and ion chromatography. However, such approaches can be time-consuming, require expensive instrumentation and are not practical for measuring fluoride concentrations in vivo. In recent years, a number of alternative strategies have been devised, such as the development of colorimetric or luminescent molecular probes, and reaction-based (irreversible) chemodosimeters.¹–⁶ The creation of probes for fluoride recognition in water is particularly challenging because of the high free energy of hydration of fluoride (ΔGhydration = 465 kJ mol⁻¹) compared to other anions, together with its relatively small ionic radius (1.26 Å). Most probes reported to date have been limited in sensing applications, because they suffer from interference from other anions present in water samples (e.g. chloride, nitrate, sulfate and phosphate) that bind competitively. Furthermore, very few probes can actually detect fluoride ions in a pure aqueous medium;¹ the majority of reported systems are restricted for use in organic solvents such as acetonitrile,² DMSO,³ methanol⁴ or dichloromethane,⁵ or they require a solvent mixture of organic solvent and water.⁶ Reaction-based probes, such as those which rely on fluoride-mediated cleavage of a Si–O bond, are irreversible and typically suffer from delayed acquisition times (10 min to several hours).¹b,d

Over the last decade, several emissive lanthanide (Ln) complexes have been developed that can report changes in the concentrations of various anions, including citrate, lactate and bicarbonate, through modulation of emission spectral form or lifetime of the complex.⁷,⁸ However, Ln-based sensors for fluoride are scarce because the binding affinity between fluoride and Ln ions is generally too weak (log Kd = 1.5–3).⁹ In this case, the fluoride detection range was very low (0.2–50 µM), falling outside of the environmentally significant range (20–210 µM).

In this work, two water-soluble luminescent probes [Eu.L1–2]⁺ are reported (Fig. 1), each capable of binding and sensing fluoride in pure water samples, with minimal interference from other anions. Each probe is based on a C₂-symmetric mono-cationic europium complex, bearing two trans-related quinoline chromophores and a coordinated water molecule. Fluoride binds reversibly to each probe, displacing the coordinated water, resulting in a 9-fold enhancement in the overall emission intensity and

Fig. 1 Structure of luminescent europium complexes [Eu.L1–2]⁺.
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Dramatic changes in emission spectral form (Fig. 2). Complex $[\text{Eu.L}^2]^{+}$ represents one of the very few probes capable of signalling the presence of fluoride within the range relevant to water fluoridation (20–210 mM), and provides an instantaneous spectral readout signal.

Details of the synthesis of complexes $[\text{Eu.L}^{1-2}]^{+}$ are provided in the ESI. Selected photophysical data for $[\text{Eu.L}^{1-2}]^{+}$ are provided in Table 1. The UV-Vis absorption spectrum of $[\text{Eu.L}^{1-2}]^{+}$ is composed of a broad featureless band centred at 332 nm, whereas the absorption spectrum of $[\text{Eu.L}^1]^{+}$ features a narrow band with a maximum at 318 nm (Fig. S1, ESI†). The emission spectra of $[\text{Eu.L}^{1-2}]^{+}$ in water (25 mM HEPES, pH 7.4) were similar, each characterised by at least three components in the $\Delta J = 1$ transition (585–605 nm), indicating that both complexes adopt structures of low symmetry in water (Fig. 2a). In addition, the $\Delta J = 2$ transition (605–625 nm) is approximately equal in intensity to the $\Delta J = 4$ manifold (685–710 nm).

To assess the ability of $[\text{Eu.L}^{1-2}]^{+}$ to signal the presence of different anions, the emission spectrum of $[\text{Eu.L}^1]^{+}$ (20 μM) was recorded in water (25 mM HEPES, pH 7.4) in the presence of a 100-fold excess of a range of anions (Fig. 2). The addition of fluoride resulted in a 9-fold enhancement in the overall emission intensity of $[\text{Eu.L}^1]^{+}$, as well as substantial perturbations in the emission spectral form (Fig. 2b). The limiting spectrum indicated the formation of one major fluoride-bound species, as defined by a single $\Delta J = 0$ transition at 579 nm and two intense components within the $\Delta J = 1$ transition, centred at 586 and 596 nm respectively. In addition, the relative intensity of the hypersensitive $\Delta J = 2$ transition centred at 617 nm increased by a factor of 15. The only other anion that induced a significant spectral response was bicarbonate. In the presence of $\text{HCO}_3^−$, the limiting spectrum of $[\text{Eu.L}^1]^{+}$ (Fig. 2c) was distinctly different from that observed with fluoride. Most notably, a larger ratio between the transitions $\Delta J = 2/\Delta J = 1$ was observed. In contrast, all other anions typically present in water samples, including chloride, phosphate, sulfate and nitrate induced essentially no change (<10%) in emission intensity or spectral form (Fig. 2d).

**Table 1** Selected photophysical data for Eu complexes $[\text{Eu.L}^{1-2}]^{+}$ (H$_2$O, or as stated)

<table>
<thead>
<tr>
<th>Complex</th>
<th>$\lambda_{\text{max}}$/nm</th>
<th>$\varepsilon$/mM$^{-1}$ cm$^{-1}$</th>
<th>$\Phi_{\text{em}}$/%$^a$</th>
<th>$\tau_{(0,0)}$/ms</th>
<th>$\tau_{(1,0)}$/ms</th>
<th>$q^b$</th>
</tr>
</thead>
<tbody>
<tr>
<td>$[\text{Eu.L}^2]^{+}$</td>
<td>332</td>
<td>12.5</td>
<td>7</td>
<td>0.49</td>
<td>1.38</td>
<td>1.2</td>
</tr>
<tr>
<td>$[\text{Eu.L}^1]^{+}$</td>
<td>318</td>
<td>11.8</td>
<td>23</td>
<td>0.51</td>
<td>1.37</td>
<td>1.1</td>
</tr>
</tbody>
</table>

$^a$ Errors on quantum yields and lifetimes are ±15%. $^b$ Values of hydration state, $q$ (±20%) are derived using methods in ref. 10.

An affinity constant was determined for fluoride binding by making incremental additions of NaF to a solution of $[\text{Eu.L}^1]^{+}$ (20 μM) at pH 7.4 (25 mM HEPES). The change in the emission intensity ratio, 596/615 nm, was measured as a function of anion concentration, and the data were analysed using a non-linear least squares curve-fitting procedure based on a 1:1 binding model (Fig. S4 and S5, ESI†). $[\text{Eu.L}^1]^{+}$ was found to bind to fluoride with log $K_a = 3.5$ (±0.1). Under the same conditions, $[\text{Eu.L}^1]^{+}$ showed a slightly weaker affinity for bicarbonate [log $K_a = 3.0$ (±0.1)]. To eliminate competitive binding to $\text{HCO}_3^−$ entirely, a fluoride titration experiment was conducted at pH 6 using 25 mM MES buffer (at pH 6, residual HCO$_3^−$ is readily removed as CO$_2$) (Fig. 3). Under these conditions, an affinity...
constant for the binding of fluoride to $[\text{Eu.L}^1]^+$ was log $K_a = 4.1$ (±0.1). This is 5 times stronger than that determined at pH 7.4, reflecting the absence of HCO$_3^-$ at lower pH. Complex $[\text{Eu.L}^2]^+$, which lacks acetamide groups, showed slightly weaker binding to fluoride (log $K_a = 3.5$ ± 0.1) under the same conditions.

The affinity of $[\text{Eu.L}^{1-2}]^+$ for fluoride is substantially higher than that of previously reported Ln–F systems. This can be ascribed to a synergistic effect between Eu(n) coordination and additional hydrogen bonding interactions to the electron deficient quinoline units. A molecular model of $[\text{Eu.L}^1]^+$ with a coordinated fluoride ion [optimised at B3LYP/3-21G* with the Gaussian 09 package, using water as solvent$^1$] indicated that the bound fluoride ion can form two short C–H···F$^-$ contacts with each of the quinoline units (Fig. 4). The average C–H···F$^-$ distance is 2.81 Å, with an average H···F$^-$ distance of 1.88 Å. Both C–H···F$^-$ angles are acceptably linear (avg. angle is 141°).$^1$ It is hypothesised that these bifurcated C–H···F···H–C contacts significantly stabilise the metal-bound fluoride. Such interactions would also restrict rotation of the quinoline N–Eu bonds, thus stabilising a single diastereomer, consistent with that observed in the Eu emission spectrum.

The dramatic changes in Eu emission spectral form of $[\text{Eu.L}^{1-2}]^+$ observed in the presence of fluoride are indicative of alteration of the Eu(n) metal coordination environment. To probe this change further, the effect of fluoride on the $^1$H NMR spectrum of $[\text{Eu.L}^1]^+$ was investigated in D$_2$O (pD 6.4). In the absence of added fluoride, two sets of proton resonances were clearly discernible in the $^1$H NMR spectrum of $[\text{Eu.L}^1]^+$, consistent with the presence of two diastereomers in solution (Fig. 5). The addition of 1 eq. of NaF resulted in the appearance of a new set of resonances corresponding to the fluoride-bound species, in slow exchange with the original hydrated complex on the NMR timescale. In the presence of excess NaF (5 eq.), the original signals disappeared and only signals for the fluoride-bound complex remained. The number of apparent signals has halved compared to the hydrated complex, indicating that a single fluoride-bound species has formed. Coordination of fluoride was also observed in the $^19$F NMR spectra, which showed a resonance at $-123$ ppm for unbound fluoride, and the emergence of a second signal at $-474$ ppm, corresponding to fluoride bound at the Eu metal centre (Fig. S11, ESIF$^\dagger$).

The remarkable sensitivity of the probe was demonstrated by plotting the emission intensity ratio, 596/601 nm, as a function of added NaF in the concentration range 0–260 μM (Fig. 6). The plot showed very good linearity ($R^2 = 0.9994$) within this range. Notably, a 200% enhancement in the intensity ratio was obtained after the addition of 260 μM NaF. Competitive binding studies revealed that $[\text{Eu.L}^{1-2}]^+$ can detect low micromolar levels of fluoride with essentially no interference from other anions typically found in drinking water (e.g. Cl$^-$, HPO$_4^{2-}$, NO$_3^-$ and SO$_4^{2-}$), even when present in high concentrations (5 mM) (Fig. S9, ESIF$^\dagger$). Thus, $[\text{Eu.L}^1]^+$ represents one of the most sensitive probes for the selective determination of fluoride levels in drinking water.

To validate the practicable utility of probe $[\text{Eu.L}^1]^+$, tap water samples buffered at pH 6 (25 mM MES) were spiked with known concentrations of NaF. The amount of fluoride was measured by comparing the recorded emission spectra with the calibration curve in Fig. 6 (Table 2). The results obtained using $[\text{Eu.L}^1]^+$

![Fig. 4](image-url) An optimised structure of $[\text{Eu.L}^3]^+$ with a coordinated fluoride ion, stabilised by two short C–H···F$^-$ contacts (avg. C–H···F$^-$ distance is 2.81 Å, avg. C–H···F$^-$ angle is 141°). The model geometry was optimised at B3LYP/3-21G* (Gaussian 09 package) using the crystal structure of an Eu complex reported in ref. 14 as a starting point.

![Fig. 5](image-url) $^1$H NMR spectra of $[\text{Eu.L}^1]^+$ (2.4 mM) upon addition of NaF (0–5 eq.). Measured in D$_2$O (pD 6.4, 298 K).

**Table 2** Measurement of fluoride ion concentration in spiked tap water samples, using $[\text{Eu.L}^1]^+$ and a fluoride-selective electrode (FSE)

<table>
<thead>
<tr>
<th>Spiked [F$^-$] μM</th>
<th>Using $[\text{Eu.L}^1]^+$</th>
<th>% Variation</th>
<th>Using FSE$\dagger$</th>
</tr>
</thead>
<tbody>
<tr>
<td>30</td>
<td>36 (±6)</td>
<td>3.0</td>
<td>35 (±3)</td>
</tr>
<tr>
<td>65</td>
<td>65 (±4)</td>
<td>7.0</td>
<td>68 (±3)</td>
</tr>
<tr>
<td>115</td>
<td>113 (±4)</td>
<td>6.0</td>
<td>111 (±4)</td>
</tr>
<tr>
<td>160</td>
<td>154 (±6)</td>
<td>6.5</td>
<td>165 (±4)</td>
</tr>
<tr>
<td>210</td>
<td>204 (±8)</td>
<td>5.0</td>
<td>212 (±4)</td>
</tr>
</tbody>
</table>

$^a$ Values are the mean average of two independent measurements. $^b$ Calculated based on an initial F$^-$ ion concentration of 5 μM.
were in close agreement with the expected fluoride levels, with a maximal percentage variation of 7%. The results were also compared to those obtained independently using a fluoride-selective electrode (FSE); both analytical methods showed very good agreement.

In summary, two luminescent europium complexes $[\text{Eu.L}^{1-2}]^+$ have been synthesised, each capable of selective binding to fluoride in water in the 20–260 µM range. Such strong binding to fluoride can be attributed to a synergistic effect between Eu metal coordination and two stabilising C–H...F$^-$ interactions. $[\text{Eu.L}^1]^+$ was used for the quantitative determination of fluoride ions in drinking water samples, within the range relevant to water modelling studies.

In the context of drinking water treatment, the ligands were used for the quantitative determination of fluoride in drinking water samples, within the range relevant to water modelling studies.

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References
11 The concentration of HCO$_3^-$ in water rises with increased pH. A pH titration revealed that the emission spectral profile of $[\text{Eu.L}^1]^+$ did not change at all within pH range 3.5–7.0 (where the level of HCO$_3^-$ is negligible), but was sensitive to the increased amount of HCO$_3^-$ present in the pH range 7.1–8.5 (Fig. S3, ESf).
12 Energy minimised structures were determined using the polarised continuum solvent model (PCM), using water as solvent.