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Concentration-Independent pH Detection with a Luminescent imetallic Eu(III)-Based Probe

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Abstract

A pH-responsive, luminescent, dimetallic Eu(III)-containing complex has been synthesized and exhibits a unique mechanism of response. The luminescence-decay rate of the complex is slow, due to a lack of water molecules coordinated to the Eu(III) ions. However, the luminescence-decay rate decreases with increasing pH over a biologically relevant range of 4 to 8. Physical characterization and computational analysis suggest that the pH response is due to protonation of a bridging alkoxide at lower pH values. Modulation of the luminescence-decay rate is independent from the concentration of Eu(III), which we expect to be useful in the non-invasive imaging of *in vivo* pH.

Molecular imaging with fluorescence microscopy is a powerful tool in biochemical and biomedical research. Organic fluorescent probes that respond to the presence of specific molecular analytes are limited by photobleaching, broad emission bands (~100 nm), small Stokes shifts (<20 nm), and short-lived emissions (<100 ns). Lanthanide-based probes overcome many of the limitations associated with organic fluorophores because they offer atomic-based emissions that do not photobleach, emit line-like bands (<20 nm bandwidth), have large Stokes shifts, and exhibit long luminescence lifetimes (ms). Due to these advantages, many examples of lanthanide-based probes have been reported for the detection of biologically important cations,¹ anions,² neutral species,³ proteins and peptides,⁴ and DNA.⁵ Many of these probes rely on the ratio of two emission peaks to determine analyte concentration, but the luminescence-decay rate of Eu(III)-containing complexes should enable the determination of analyte concentration without knowledge of probe concentration or extinction coefficient. Reports describing time-resolved microscopy,⁶ luminescence-lifetime imaging,⁷ and phosphorescence-quenching microscopy⁸ demonstrate that luminescence-decay rates can be determined from microscopy and are, therefore, a useful parameter for sensing applications. Lanthanide-based probes often exhibit low sensitivity that can be addressed by the use of antennae or through the use of multimetallic complexes that increase sensitivity additively.⁹ In targeting the multimetallic strategy, we synthesized a new dimetallic Eu(III)-containing complex, **1**, that responds to pH over a physiologically relevant range of 4 to 8 by a new mechanism, based on luminescence-decay rates, that is independent of the probe concentration and selective for only proton concentration. To our knowledge, no system of this type has been reported to respond to pH by luminescence-decay rate.

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ASSOCIATED CONTENT

Supporting Information

Synthetic details, characterization, table with optimized distances, and the complete Gaussian citation. This material is available free of charge via the Internet at <http://pubs.acs.org>.

We hypothesized that bridging two Eu(III) ions with an alcohol would decrease electron density on the bridging oxygen and increase the acidity of the alcohol compared to an uncoordinated alcohol. We anticipated that the acid-base equilibrium of the bridging oxygen would result in a pH-dependent change in the luminescence-decay rate because the luminescence-decay rate of Eu(III) increases with the number of hydroxyl oscillators coordinated to the metal ion.¹⁰ We expected this change to be independent of the concentration of **1** because luminescence-decay rate is not dependent on metal concentration. Further, we expected that this mechanism would be selective for protons because hydroxyl oscillators have the correct vibrational energy levels to efficiently quench Eu(III) luminescence. Here, we present the synthesis, characterization, and spectroscopic and computational analyses of dimetallic Eu(III)-containing complex **1**. Additionally, we propose a mechanism for the pH-dependent luminescence-decay rate supported by our experimental and computational data.

Our ligand design incorporates two Eu(III)-containing 2,2',2''-(1,4,7,10-tetraazacyclododecane-1,4,7-triyl)triacetate complexes bridged by an *iso*-propanol linker that is positioned to form two *ortho*-fused five-membered rings containing the two metal ions. The synthesis of **1** was completed in four steps (Scheme 1). Briefly, 1,3-dibromo-2-propanol was converted to triethylsilyl ether **2**, which was used to alkylate two equivalents of 1,4,7,10-tetraazacyclododecane-1,4,7-tris(*t*-butyl acetate) yielding the protected ligand **3**. We found it imperative that the alcohol be protected for the success of this alkylation. Ligand **4** was obtained by global deprotection of **3** with trifluoroacetic acid (TFA), and Eu(III) complexation was achieved by addition of an aqueous solution of Eu(III) trifluoromethanesulfonate (OTf) to ligand **4** at pH 6.5. Dialysis yielded dimetallic Eu(III)-containing complex **1**. A xylene orange test indicated that there was no unchelated Eu(III) ions present at pH 5.8 even after storing in solution at ambient temperature for 24 h.¹¹

To assess the coordination environment of the Eu(III) ions, the number of water molecules coordinated to the Eu(III) ion, q , was determined using the method developed by Horrocks and coworkers.¹⁰ Complex **1** was found to have a q value of 0 in the absence of buffer (pH 5.4), suggesting that no water was bound to the Eu(III) ions. When we considered possible structures to satisfy this data, we were unable to construct a structure lacking a bridge between Eu(III) ions that contained no water molecules bound to the Eu(III) ions. This q data indicated that the Eu(III) ions are bridged by an alcohol donor from the ligand. Because coordinated OH oscillators influence the luminescence-decay rate of Eu(III) ions, we hypothesized that there would be a decrease in the luminescence-decay rate of **1** with increasing pH above 5.4 due to deprotonation of the bridging alcohol.

To test this hypothesis, we prepared 0.5 mM solutions of **1** in a series of buffers from pH 4.0 to 8.0 in increments of 0.2 pH units. The luminescence-decay rate of each solution was determined and plotted against pH values (Figure 1). We confirmed that the pH response is independent of the concentration of **1** by repeating the luminescence-decay rate measurements with 1 mM solutions of **1**. These experiments resulted in identical pH-dependent changes in luminescence-decay rates. The resulting plot supports our hypothesis that the luminescence-decay rate depends on pH and indicates that the pH response extends over a biologically relevant range of 4 to 8. The second derivative of this plot was used to estimate the pK_a of the protic source at 5.8, placing the maximum sensitivity of the probe very close to the pH of diseased tissue.¹²

Structures of **H1** and **1**⁻ were optimized with Gaussian 09 at the PBE0/SDDall level (small core for all atoms) using the default convergence criteria.¹³ The optimized structures suggest that the alcohol bridges the Eu(III) ions and that two carboxylate oxygens are also bridging (Figure 2). This tri-bridged structure is consistent with a 9-coordinate Eu(III) environment

that is saturated by the ligand and corroborates the luminescence-decay data that indicate the absence of water molecules bound to Eu(III). Interestingly, to obtain a suitable initial guess for the optimization, it was necessary to include the two bridging carboxylates. To achieve this, the orientation of the carboxylate arms on the two chelates had to be in opposite conformations from each other (Δ vs Λ) to minimize the bridged structure. Further, the steric constraints of the ligand prevent a bridge as small as alkoxide without two bridging carboxylates. These structures corroborate both the exceptionally long luminescence lifetime at higher pH values and the luminescence quenching that results from lowering pH.

To further elucidate the mechanism of the pH response, we compared the emission spectra of **1** at different pH values (Figure 3 and Figure S3). At pH 4.4, we observed that the multiplicity of the $\Delta J = 1$ manifold (centered at 592 nm) is at the maximum of three peaks. This high multiplicity indicates that the symmetry around the Eu(III) ions is low. However, the spectrum would indicate that the symmetry of the left and right chelates is similar because we do not observe greater than $2J+1$ peaks for any manifold in the spectrum. Surprisingly, the spectrum of **1** at pH 7.4 contains four peaks in the $\Delta J = 1$ manifold and two major peaks in the $\Delta J = 2$ manifold. The source of these additional peaks at high pH is not immediately obvious. One explanation that is consistent with all other data is that the loss of the intramolecular hydrogen bond at high pH values increases the flexibility of the ligand, resulting in a change in conformation about one or both of the Eu(III) ions. Previous studies of similar monometallic Eu(III) complexes by Horrocks and coworkers demonstrated that inter-converting isomers can be observed by luminescence spectroscopy.¹⁴ NMR analyses of interconverting lanthanide macrocyclic complexes is also well established and these methods are currently under investigation.

The spectroscopic and computational data suggest a novel mechanism of pH response wherein protonation of the bridging alkoxide under acidic conditions introduces a hydroxyl oscillator in the Eu(III) coordination sphere (Figure 4). The inner-sphere hydroxyl oscillator quenches the Eu(III) excited state resulting in a faster luminescence-decay rate. Further, because coordination of other cations to the alkoxide will not produce oscillators with the proper energy to quench the Eu(III) emission, we expect that this system will specifically respond to proton concentration. Moreover, this process is reversible such that continuous monitoring of pH is possible. To our knowledge, no other system of this kind has been reported to respond to pH by luminescence-decay rate.

We have demonstrated a concentration-independent luminescence response to pH achieved through the protic equilibrium of an alkoxide in **1**. Our characterization indicates that the Eu(III) ions have no coordinated water molecules resulting in relatively slow luminescence-decay rates, especially at pH values above 7. The absence of coordinated water molecules makes this system less sensitive to changes in the composition of the surrounding media relative to systems that contain coordinated water molecules. Spectral changes with pH are indicative of a change in the coordination sphere in agreement with our proposed mechanism. Computationally optimized structures also are consistent with our experimental observations. Complex **1** represents a unique system using a new mechanism by which pH can be detected and offers a useful method for concentration-independent, non-invasive, continuous pH imaging. We expect that this research is a step toward powerful tools for biomedical and biochemical research because pH is often an indicator of disease.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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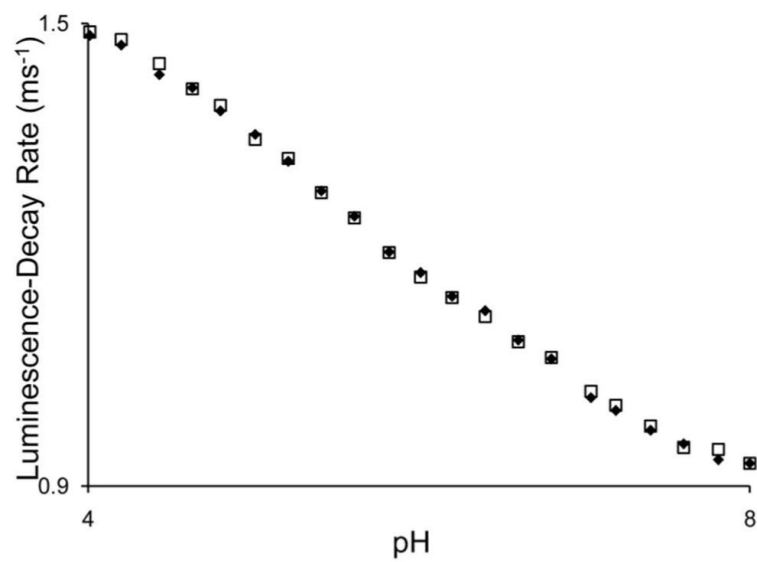


Figure 1. Luminescence-decay rate of 0.5 (◆) and 1.0 mM (□) **1** as a function of pH (50 mM citrate, 50 mM phosphate, $\lambda_{\text{ex}} = 395$ nm, $\lambda_{\text{em}} = 595$ nm).

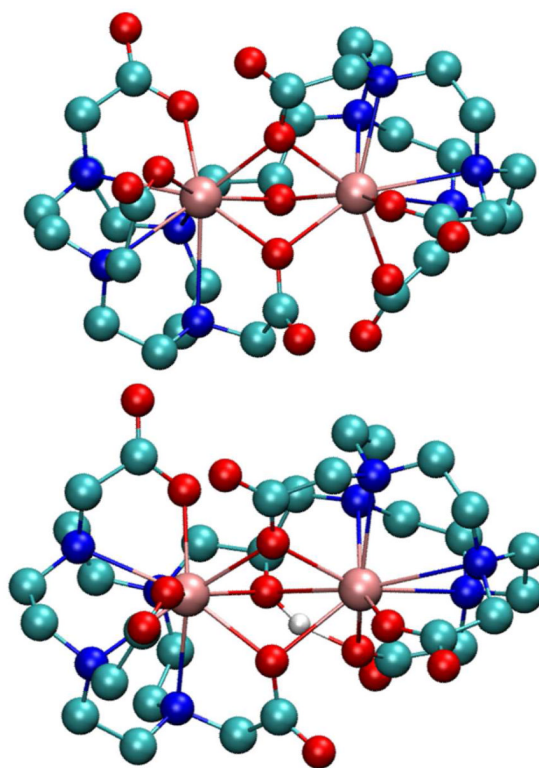


Figure 2. Optimized structures of $\mathbf{1}^-$ (top) and $\mathbf{H1}$ (bottom): Eu = pink, C = cyan, N = blue, O = red, and H = white. Non-exchangeable hydrogen atoms are omitted for clarity.

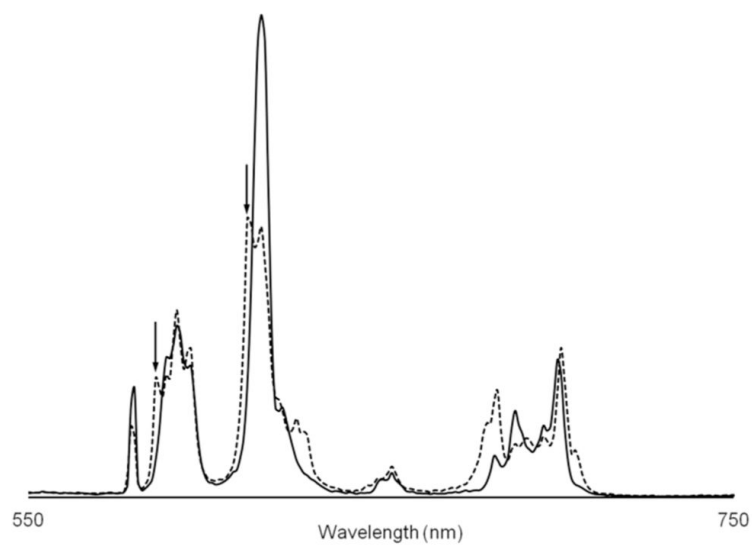
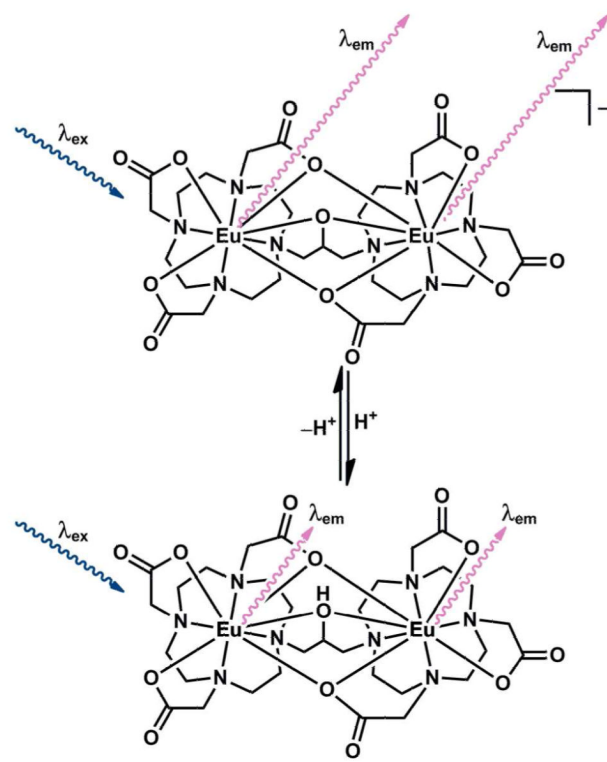
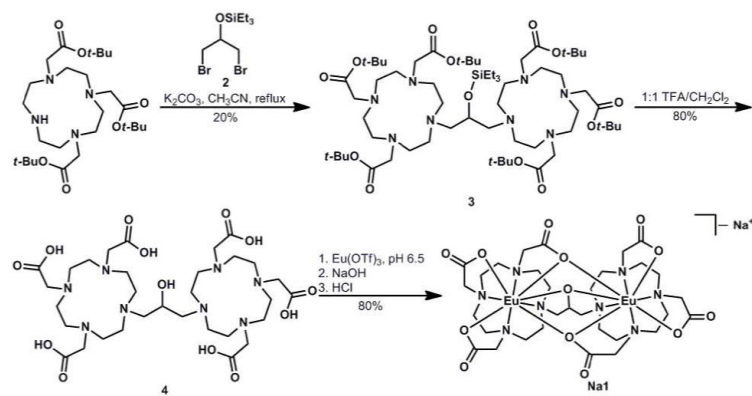


Figure 3. Comparison of emission spectra of **1** at pH 4.4 (solid line) and pH 7.4 (dashed line) (0.5 mM **1**, 50 mM citrate, 50 mM phosphate, $\lambda_{\text{ex}} = 395$ nm). Arrows point to peaks that may result from a second Eu(III) environment.

**Figure 4.**

Proposed mechanism for the response of **1** to changes in pH. At high pH (top), the lack of OH oscillators leads to long-lived luminescence (long wavy arrows, λ_{em}). At low pH (bottom), the bridging OH oscillator quenches the Eu(III) excited state leading to rapid luminescence decay (short wavy arrows, λ_{em}).



Scheme 1. Synthetic route to 1