

Circularly Polarized Luminescence from New Heteroleptic Eu(III) and Tb(III) Complexes

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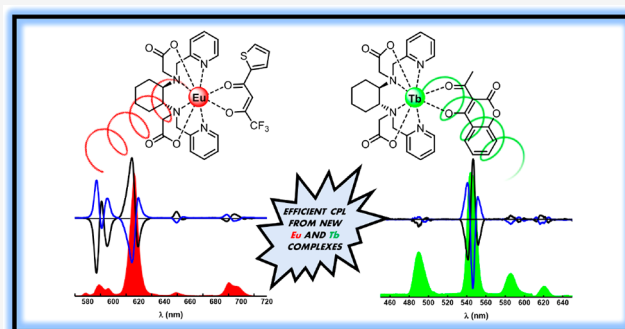


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ABSTRACT: The complexes [Eu(bpcd)(tta)], [Eu(bpcd)-(Coum)], and [Tb(bpcd)(Coum)] [tta = 2-thenoyltrifluoroacetyl-acetonate, Coum = 3-acetyl-4-hydroxy-coumarin, and bpcd = *N,N'*-bis(2-pyridylmethyl)-*trans*-1,2-diaminocyclohexane-*N,N'*-diacetate] have been synthesized and characterized from photophysical and thermodynamic points of view. The optical and chiroptical properties of these complexes, such as the total luminescence, decay curves of the Ln(III) luminescence, electronic circular dichroism, and circularly polarized luminescence, have been investigated. Interestingly, the number of coordinated solvent (methanol) molecules is sensitive to the nature of the metal ion. This number, estimated by spectroscopy, is >1 for Eu(III)-based complexes and <1 for Tb(III)-based complexes. A possible explanation for this behavior is provided via the study of the minimum energy structure obtained by density functional theory (DFT) calculations on the model complexes of the diamagnetic Y(III) and La(III) counterparts [Y(bpcd)(tta)], [Y(bpcd)(Coum)], and [La(bpcd)(Coum)]. By time-dependent DFT calculations, estimation of donor–acceptor (D–A) distances and of the energy position of the S_1 and T_1 ligand excited states involved in the *antenna* effect was possible. These data are useful for rationalizing the different sensitization efficiencies (η_{sens}) of the *antennae* toward Eu(III) and Tb(III). The tta ligand is an optimal *antenna* for sensitizing Eu(III) luminescence, while the Coum ligand sensitizes better Tb(III) luminescence ($\phi_{\text{ovl}} = 55\%$; $\eta_{\text{sens}} \geq 55\%$ for the [Tb(bpcd)(Coum)] complex}. Finally, for the [Eu(bpcd)(tta)] complex, a sizable value of g_{lum} (0.26) and a good quantum yield (26%) were measured.



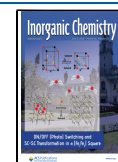
INTRODUCTION

Circularly polarized luminescence (CPL) is a chiroptical phenomenon that is attracting more interest in materials chemistry and physics thanks to the broad range of possible biological^{1–5} and technological applications,^{6–15} as in the case of the design of organic light-emitting diodes (OLEDs) emitting circularly polarized (CP) light.^{16–18} Focusing our attention on these latter devices, we can find a logical application in displays in which the emitted polarized light is exploited to prevent reflection of ambient light and obtain high-contrast three-dimensional images and true black.¹⁹ In addition, a CP screen reduces the perceived distortion found at some angles when the display is viewed through a linearly polarized (LP) filter, significantly improving the outdoor viewing of laptops or smartphones. In this scenario, as the emission of CP light by trivalent lanthanide ions is a quite efficient phenomenon, luminescent lanthanide complexes play a pivotal role.¹⁸ The new quantity recently proposed by some of us²⁰ and known as CPL brightness (B_{CPL}) can be considered as useful tool for the design of efficient CPL phosphors for

specific chiroptical applications, such as in CPL microscopy²¹ and possibly in CPL security inks.^{7,22} Moreover, a high B_{CPL} is correlated with a higher signal available for a CPL measurement.^{23,24} B_{CPL} takes into account the absorption extinction coefficient and quantum yield along with the g_{lum} factor and, for a selected lanthanide transition, is defined as $B_{\text{CPL}} = \beta_i \epsilon_\lambda \phi_{\text{ovl}} g_{\text{lum}} / 2 = \beta_i B g_{\text{lum}} / 2$, where β_i is the so-called branching ratio (β ; $0 \leq \beta \leq 1$), ϵ_λ is the molar extinction coefficient, ϕ_{ovl} is the overall quantum yield,²⁵ and g_{lum} ²⁶ is the dissymmetry factor associated with the considered transition $\beta_i = I_i / \sum_j I_j$, where I_i is the integrated intensity of the considered transition and $\sum_j I_j$ is the summation of the integrated intensities over all of the transitions. In the case of lanthanide complexes, optimal

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B_{CPL} values can be obtained in the presence of strong absorbing chromophores (high ϵ_{λ} values), good ligand-to-metal energy transfer (LMET or *antenna* effect), and luminescence efficiency from Ln(III) (both contributing to high ϕ_{ovl} values). Finally, the emitted light should be strongly polarized, with the difference in the intensity between the right- and left-handed components of the CP light being as large as possible. A promising strategy for obtaining a Ln(III)-based luminescent complex exhibiting sizable B_{CPL} values is to design heteroleptic complexes in which two different ligands are bound to the metal ion. One ligand takes care of efficiently sensitizing the Ln(III) luminescence, while the other one can trigger the required CPL activity, by virtue of the fact that it is chiral and nonracemic. As for the *antenna* that can sensitize Eu(III) and Tb(III) luminescence, the anion derived from 3-acetyl-4-hydroxycoumarin (Coum)²⁷ has been selected for both ions, while 2-thenoyltrifluoroacetyl-acetonate (tta) has been selected as the ligand for only Eu(III) (Figure 1). It is

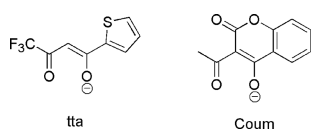


Figure 1. Ligands employed in this work.

well-known that the latter ligand efficiently sensitizes the luminescence of the Eu(III) ion by virtue of the small energy gap between the accepting levels of the europium ion and the triplet state of the ligand and the increased anisotropy around the metal ion.²⁸

As the chiral bpdc ligand can induce a significant chiroptical response stemming from Eu(III) and Tb(III) ions,²⁹ we decided to employ this ligand to produce CPL activity from our complexes. Therefore, in this work, we synthesized and spectroscopically characterized three different complexes (in both enantiomeric forms), namely, [Eu(bpdc)(tta)], [Eu(bpdc)(Coum)], and [Tb(bpdc)(Coum)]; the structures are shown in Figure 2.

Also, an ultraviolet–visible (UV–vis) titration study and density functional theory (DFT) calculations were carried out to characterize the stability and structural features of the complexes present in solution. Also, by means of time-dependent DFT (TD-DFT) calculations, the energies of the tta and Coum excited states involved in the energy transfer mechanism were determined, and a possible explanation for the different sensitization efficiencies of the different complexes has been provided.

EXPERIMENTAL SECTION

EuCl₃·6H₂O and TbCl₃·6H₂O (Aldrich, 98%) and 2-thenoyltrifluoroacetyl-acetone (Htta, Alfa Aesar, 98%) were stored under vacuum for several days at 80 °C and then transferred to a glovebox.

Both enantiomers of the *N,N'*-bis(2-pyridylmethyl)-*trans*-1,2-diaminocyclohexane-*N,N'*-diacetic acid (H₂bpdc) ligand, in the form of trifluoroacetate salt, were synthesized as previously reported in the literature.³⁰

Elemental Analysis. Elemental analyses were carried out by using an EACE 1110 CHNO analyzer.

ESI-MS. Electrospray ionization mass spectra (ESI-MS) were recorded with a Finnigan LXQ Linear Ion Trap (Thermo Scientific, San Jose, CA) operating in positive ion mode. The data were acquired under the control of Xcalibur software (Thermo Scientific). A methanol solution of the sample was properly diluted and infused into

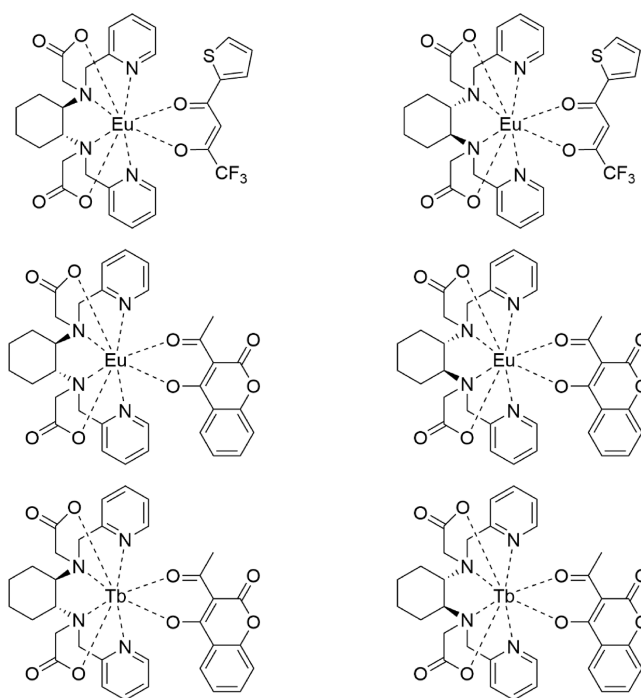


Figure 2. Complexes investigated in this work: (top) (*R,R*)-[Eu(bpdc)(tta)] (left) and (*S,S*)-[Eu(bpdc)(tta)] (right), (middle) (*R,R*)-[Eu(bpdc)(Coum)] (left) and (*S,S*)-[Eu(bpdc)(Coum)] (right), and (bottom) (*R,R*)-[Tb(bpdc)(Coum)] (left) and (*S,S*)-[Tb(bpdc)(Coum)] (right).

the ion source at a flow rate of 10 $\mu\text{L}/\text{min}$ with the aid of a syringe pump. The typical source conditions were as follows: transfer line capillary at 275 °C; ion spray voltage of 4.70 kV; and sheath, auxiliary, and sweep gas (N_2) flow rates of 10, 5, and 0 arbitrary units, respectively. Helium was used as the collision damping gas in the ion trap set at a pressure of 1 mTorr.

The Coum ligand precursor was synthesized by modifying the procedure previously reported in the literature.²⁷ In a 50 mL round-bottom flask equipped with a condenser under a nitrogen atmosphere, 4-hydroxy coumarin (839 mg, 5.18 mmol), dry and freshly distilled pyridine (10 mL), and 4-(dimethylamino)pyridine (DMAP, 32 mg, 0.26 mmol) were added. To this stirred suspension was added acetic anhydride (0.57 mL, 5.29 mg, 5.18 mmol), and the mixture was stirred at 50 °C for 24 h. Pyridine was evaporated under reduced pressure; the residue was dissolved in ethyl acetate (AcOEt, 20 mL), and 1 M aqueous HCl was added until the pH reached 1. The water phase was extracted with AcOEt (3 \times 15 mL); the combined organic layers were dried over anhydrous Na_2SO_4 , and the solvent was removed under reduced pressure. The crude was subjected to flash column chromatography (SiO_2 ; 85:15 cyclohexane/AcOEt) to afford 3-acetyl-4-hydroxy-2*H*-chromen-2-one in 64% yield (673 mg, 3.3 mmol). Spectroscopic data (¹H NMR and ¹³C NMR in Figures S11 and S12) are in agreement with those reported in the literature.²⁷

[Eu(bpdc)(tta)] (yield of 94%) was synthesized as follows (Figure 3). At room temperature, KOH (60 mg, 1.07 mmol) was added to a methanol (8 mL) solution of the desired enantiomer (*S,S* or *R,R*) of H₂bpdc (200 mg, 0.31 mmol, as trifluoroacetate salt). After 15 min, EuCl₃·6H₂O (116 mg, 0.31 mmol) was added to form the first complex, [Eu(bpdc)]Cl. In another flask, KOH (21 mg, 0.37 mmol) was added to a methanol (4 mL) solution of Htta (2-thenoyltrifluoroacetyl-acetone, 68 mg, 0.31 mmol). This mixture was slowly added to the previously prepared solution containing the first complex, and the final mixture was stirred for 1 h. Then, the solvent was removed under reduced pressure, and the desired product was obtained as a yellow powder upon extraction in dichloromethane (8 \times 3 mL) followed by solvent removal *in vacuo*.

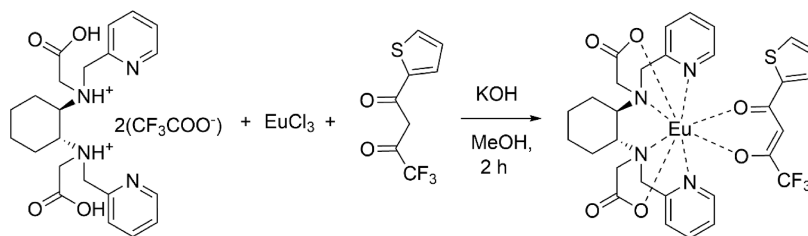


Figure 3. Synthesis of [Eu(bpdc)(tta)] complexes (the *R,R* enantiomer is chosen as a representative).

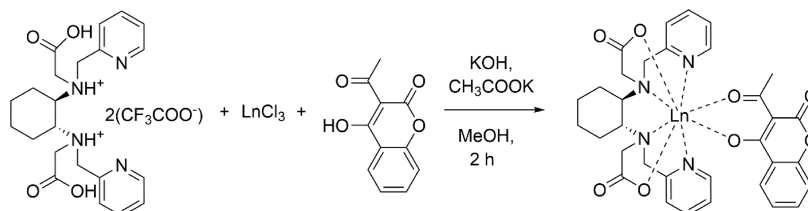


Figure 4. Synthesis of [Ln(bpdc)(Coum)] complexes (Ln = Eu or Tb). The *R,R* enantiomers are chosen as a representative.

Elemental analysis calcd for [Eu(bpdc)(tta)], $C_{30}H_{30}EuF_3N_4O_6S$ (MW 783.6): C, 45.98; H, 3.86; N, 7.15; O, 12.25. Found: C, 45.11; H, 3.80; N, 6.99; O, 12.37 (*R,R* isomer). Found: C, 45.28; H, 3.78; N, 7.07; O, 12.12 (*S,S* isomer). ESI-MS (scan ES+, m/z): 807.09 (100%), 805.09 (90%), 808.10 (35%) ([Eu(bpdc)(tta)] + Na)⁺; 784.09 (100%), 785.11 (30%), 783.11 (27%), 786.12 (10%) ([Eu(bpdc)(tta)] + H)⁺. ESI-MS (scan ES-, m/z): 220.99 (100%), 221.99 (10%) [tta]⁻.

[Eu(bpdc)(Coum)] and [Tb(bpdc)(Coum)] (yield of 95%) were synthesized as follows (Figure 4). At room temperature, $EuCl_3 \cdot 6H_2O$ (113 mg, 0.31 mmol) or $TbCl_3 \cdot 6H_2O$ (115.7 mg, 0.31 mmol) was added to a previously prepared methanol solution containing KOH (60 mg, 1.07 mmol) and H_2bpdc (*S,S* or *R,R*) (200 mg, 0.31 mmol, as trifluoroacetate salt). In another flask, 3-acetyl-4-hydroxy-2*H*-chromen-2-one (63.3 mg, 0.31 mmol) was solubilized in methanol and added to a solution of NaOMe (16.7 mg, 0.31 mmol), in the same solvent (10 mL), with some drops of acetonitrile to improve its solubility. The deprotonated Coum ligand was slowly added to the solution containing the Ln(III) complex. The final mixture was stirred for 2 h. Then, the solvent was removed under reduced pressure, and the desired product was obtained as a yellow powder upon extraction in dichloromethane (8 × 3 mL) followed by solvent removal in vacuo.

Elemental analysis calcd for [Eu(bpdc)(Coum)](CH_3OH), $C_{34}H_{37}EuN_4O_9$ (MW 797.64): C, 51.20; H, 4.68; N, 7.02; O, 18.05. Found: C, 51.22; H, 4.54; N, 7.11; O, 17.99 (*R,R* isomer). Found: C, 51.12; H, 4.60; N, 6.95; O, 17.91 (*S,S* isomer). ESI-MS (scan ES+, m/z): 789.14 (100%), 787.14 (90%), 790.14 (35%) ([Eu(bpdc)(Coum)] + Na)⁺.

Elemental analysis calcd for [Tb(bpdc)(Coum)], $C_{33}H_{33}N_4O_8Tb$ (MW 772.56): C, 51.30; H, 4.31; N, 7.25; O, 16.57. Found: C, 51.25; H, 4.22; N, 7.18; O, 16.12 (*R,R* isomer). Found: C, 51.19; H, 4.12; N, 7.19; O, 16.33 (*S,S* isomer). ESI-MS (scan ES+, m/z): 795.14 (100%), 796.15 (40%), 797.15 (8%) ([Tb(bpdc)(Coum)] + Na)⁺.

Luminescence and Decay Kinetics. Room-temperature luminescence was measured with a Fluorolog 3 (Horiba-Jobin Yvon) spectrofluorometer, equipped with a Xe lamp, a double-excitation monochromator, a single-emission monochromator (mod. HR320), and a photomultiplier in photon counting mode for the detection of the emitted signal. All of the spectra were corrected for the spectral distortions of the setup. The spectra in solution were recorded on methanol (50 μ M) solutions.

In decay kinetics measurements, a xenon microsecond flash lamp was used and the signal was recorded by means of a multichannel scaling method. True decay times were obtained using the convolution of the instrumental response function with an exponential function and the least-squares-sum-based fitting program (Spectra-Solve software package).

Circularly Polarized Luminescence. CPL spectra were recorded with the homemade spectrofluoropolarimeter described previously.³¹ The spectra were recorded on methanol (0.4 mM) solutions in a 1 cm cell. The samples were excited at 365 nm {for [Eu(bpdc)(tta)]} or 254 nm {for [Eu(bpdc)(Coum)] and [Tb(bpdc)(Coum)]}.

Electronic Circular Dichroism (ECD). ECD spectra were recorded with a Jasco J1500 spectropolarimeter on 1 mM CH_3OH solutions in a 0.01 cm cell.

Overall Quantum Yield Measurements. Overall quantum yields were measured by adopting the relative method. Fluorescein in 0.1 M NaOH (fluorescence quantum yield of 0.9) was used as the standard. Absorption spectra were recorded with a PerkinElmer Lambda 650 UV-vis spectrophotometer. Emission spectra were recorded with an Edinburgh FLS1000 fluorometer and corrected for excitation intensity and detector sensitivity. The samples were dissolved in methanol, while keeping their absorbance lower than 0.1. We obtained the same values of overall quantum yield for the two enantiomers.

Spectrophotometric Titrations. The UV-vis absorption spectra were recorded in the wavelength range of 220–500 nm on a Varian Cary 50 spectrophotometer using 1 cm optical path length quartz cells (Hellma Analytics) sealed with a Teflon stopper. The samples were prepared in dry methanol within a drybox with a N_2 atmosphere.

In our experiments, solutions of tta or Coum (initial concentrations of 5.0×10^{-5} mol L^{-1}) were titrated with a [Eu(bpdc)]Cl solution (1.0×10^{-3} mol L^{-1}) with a final ligand:complex molar ratio ~ 3.0 . The cell was stirred for ~ 10 min before each spectrum was recorded. The formation constants of the [Eu(bpdc)(tta)] and [Eu(bpdc)(Coum)] complexes were obtained by multiwavelength analysis of the absorption spectra using HypSpec.³²

DFT Calculations. All molecular structures of the complexes were obtained by means of DFT calculations performed in Gaussian 16 (version A.03).³³ In previous works,^{30,34} the paramagnetic Eu(III) ion was replaced by Y(III), which was a suitable substitute. This choice is also supported for the isostructural complexes found with analogous hexadentate ligands EDTA and CDTA. In the crystal structures with the latter two ligands,^{35–37} Y(III) and Eu(III) ions are nine-coordinated with EDTA (ligand and three water molecules bound to the metal) and eight-coordinated with CDTA (two bound water molecules). Because the ionic radius of Y(III) is smaller than those of Eu(III) and Tb(III), the calculations for the same complexes with the larger La(III) ion were also carried out. To identify the solvent molecules bound to the complexes, the geometries of [Ln(bpdc)(Coum)]·4 H_2O (L = Y or La), in which water molecules replaced methanol, were also considered.

The functional B3LYP^{38,39} was used with the 6-31+G(d) basis set for all ligand atoms and MWB28 pseudopotential and valence

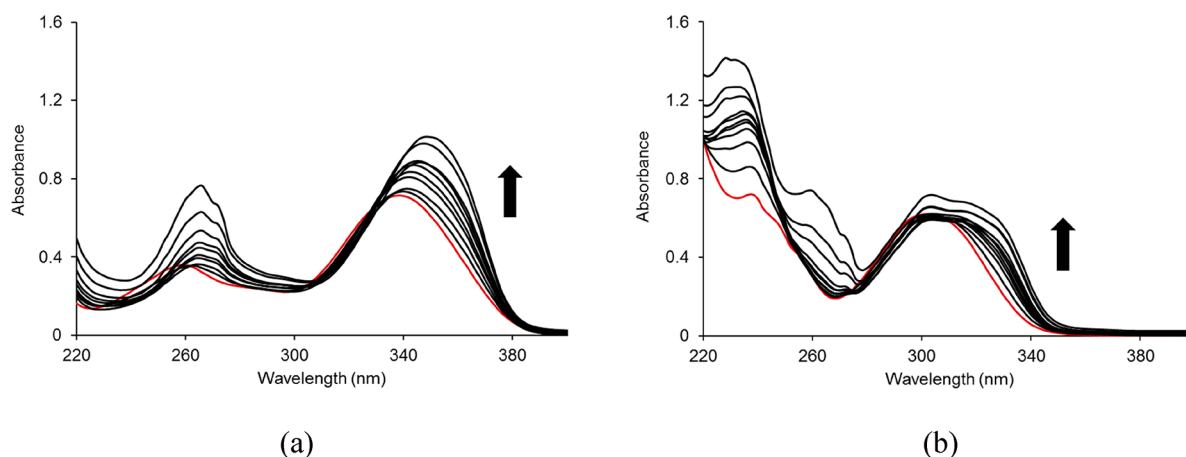


Figure 5. Changes in the UV–vis absorption spectra during the titration of (a) tta (50 μM) and (b) Coum (50 μM) upon addition of a solution of [Eu(bpcd)Cl] at 25 $^{\circ}\text{C}$. The spectra of the initial tta and Coum solutions are colored red.

electron basis set for the metal ions.^{40,41} Geometry optimizations were carried out at the DFT level with a polarizable continuum model (PCM) to simulate solvation.⁴²

As in a previous work,⁴³ the excited state (T_1 and S_1) energies were obtained by employing the time-dependent DFT approach (TD-DFT) on the [Y(bpcd)L] complexes using the same level of theory as in the geometry optimizations, as it was shown that the B3LYP functional accurately predicts the UV–vis spectra of coumarin derivatives.⁴⁴ Analyses were performed with Multiwfn version 3.8.⁴⁵

RESULTS AND DISCUSSION

Synthesis of the Complexes. Both enantiomers (*S,S* or *R,R*) of [Eu(bpcd)(tta)], [Eu(bpcd)(Coum)], and [Tb(bpcd)(Coum)] complexes have been obtained in very high chemical yields ($\sim 95\%$) and high degrees of purity, as confirmed by the elemental analysis and ESI-MS data. Additional details of the synthesis are reported in the [Experimental Section](#).

UV–Vis Absorption and ECD. The UV–vis electronic absorption spectra and ECD spectra of the [Eu(bpcd)(tta)], [Eu(bpcd)(Coum)], and [Tb(bpcd)(Coum)] complexes in methanol are shown in [Figure S1](#).

As it is well-known, the absorption band around 350 nm, in the case of [Eu(bpcd)(tta)], can be attributed to the diketonate-centered singlet–singlet π – π^* transition of the tta ligand⁴⁶ while the absorption band peaking around 270 nm is assigned to electronic transitions involving both pyridine ring (i.e., π – π^* and n – π^*) transitions of the bpcd²⁻ ligand.³⁰ The lower-energy ECD bands indicate that the chiral ligand can dictate a preferred sense of twist of the diketonate, as demonstrated by a dichroic signal around 350 nm, where the absorption of tta takes place.

The UV–vis electronic absorption spectra of [Eu(bpcd)(Coum)] and [Tb(bpcd)(Coum)] complexes are practically identical, demonstrating the ligand-centered nature of the electronic transitions, which is independent of the type of metal ion. The absorption peaks centered around 230, 300, and 320 nm, as previously demonstrated,²⁷ are related to Coum electronic transitions, while the shoulder at 270 nm is due to the absorption of the bpcd²⁻ ligand. Although both ECD spectra show an overall similar trend, the dichroic signals show some differences in intensity and shape especially in the range of 280–350 nm. Possibly, this could be due to the different size of the lanthanide cation (Eu^{3+} vs Tb^{3+}), which in turn impacts the position of the Coum moiety relative to the

bpcd ligand. Geometrical variations within a series of analogue lanthanide complexes are possible, especially across the so-called Gd break. For comparison, the ECD of the [Gd(bpcd)(Coum)] analogue was also measured. It displayed a trend similar to that shown by the Eu complex but with a somehow stronger band at 270 nm ([Figure S1](#)). Again, the CD signals in the range of 280–350 nm, which are associated with the achiral coumarin, indicate a stereochemically defined arrangement of the whole complex, including the coumarin.

UV–Vis Titrations. The absorption spectra of tta/Coum with an increasing concentration of [Eu(bpcd)]Cl were recorded in anhydrous methanol between 220 and 400 nm. The spectra of both systems present a maximum of absorption at 266 nm related to the pyridine group of the [Eu(bpcd)]⁺ complex.³⁰ As for tta ([Figure 5a](#)), the initial absorption maximum at 340 nm undergoes a marked red-shift upon formation of the [Eu(bpcd)(tta)] complex in solution. On the contrary, for Coum ([Figure 5b](#)), the intensity of the initial absorption at 310 nm increases with a slight red-shift and a shoulder at 320 nm. The data were analyzed by simultaneous least-squares fitting of the absorbance data in the wavelength range associated with the formation of the new species (300–380 and 290–340 nm for tta and Coum, respectively). As an example of the goodness of fit, calculated and experimental absorbances at selected wavelengths are reported in [Figure S2](#). The fitting procedure confirms the formation of quite stable 1:1 species ([Table 1](#)). Interestingly, we show that the complex

Table 1. Equilibrium Constants (log K) for the Formation of the [Eu(bpcd)]L Complexes in Anhydrous Methanol at 25 $^{\circ}\text{C}$ (L = tta or Coum) (charges omitted)

	log K	
	L = tta	L = Coum
[Eu(bpcd)] + L \rightleftharpoons [Eu(bpcd)L]	4.12 \pm 0.05	5.59 \pm 0.06

formed with Coum is more stable (~ 1.5 log units) than that with tta. In previous works, the formation of the [EuL₃]³⁻ species was studied in water for tta (log K_1 = 4.65, log β_2 = 9.67, and log β_3 = 12.0 at 25 $^{\circ}\text{C}$ in water)⁴⁷ and mixed solvents for Coum (50% water/dioxane; log K_1 = 3.92 and log β_2 = 6.89 at 35 $^{\circ}\text{C}$).⁴⁸ However, our results ([Table 1](#)) are not comparable with the literature data for the 1:1 species. The reaction reported in those works indeed occurs between the

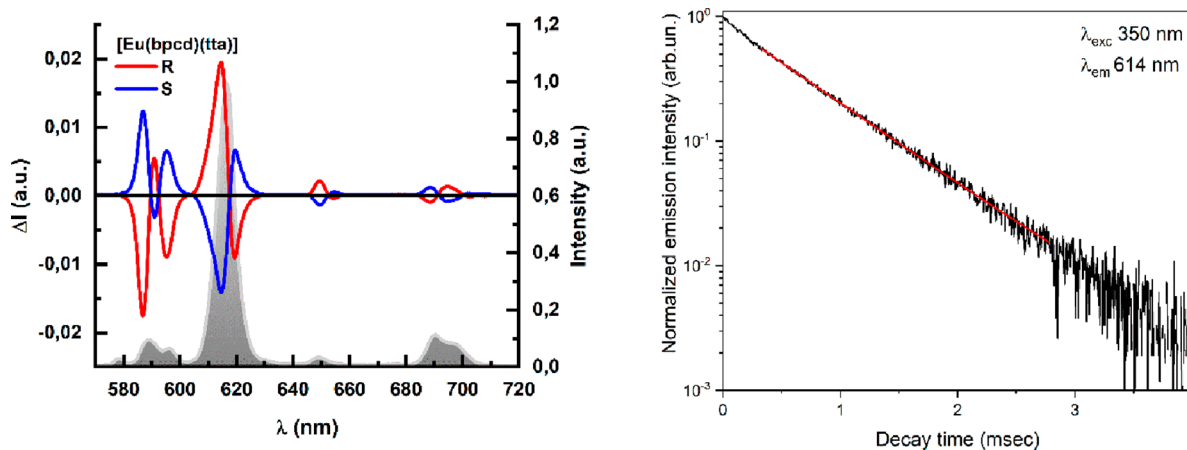


Figure 6. CPL spectra (left) of both enantiomers of [Eu(bpcd)(tta)], with the normalized total luminescence spectrum overlaid, upon excitation at 365 nm. Eu(III) 5D_0 luminescence decay curve (right) (the plot of the S,S isomer is shown and chosen as a representative). The equation of the fitting curve (red line) is $y = 0.903 \exp(-t/0.68) + 0.001$ [reduced $\chi^2 = 4.0 \times 10^{-5}$; R^2 (COD) = 0.99803].

Table 2. Most Relevant Photophysical Parameters of the Eu(III) and Tb(III) Complexes in Methanol Investigated in This Work^a

complex	τ_{obs} (ms)	τ_{obs} (ms), CD ₃ OD	τ_{rad} (ms)	ϕ_{int} (%)	ϕ_{ovl} (%)	η_{sens} (%)	m^b
[Eu(bpcd)(tta)]	0.68(1)	1.18(1)	2.38(1)	26.4(1)	26	100	1.3(5) ^c
[Eu(bpcd)(Coum)]	0.80(1)	1.40(1)	2.42(1)	33.1(1)	7	21.1	1.1(5) ^c
[Tb(bpcd)(Coum)]	1.42(1)	1.59(1)	—	—	55	$\geq 55^a$	0.6(5) ^d

^aThe reported values are the same for both enantiomers. ^bNumber of methanol molecules in the inner coordination sphere. ^cCalculated as described in ref 50. Following the Judd–Ofelt theory,⁵⁶ one can calculate τ_{rad} from the emission spectra, only in the case of the Eu(III) ion. ^dCalculated by a modification for Tb(III) of the equation discussed for the Eu(III)⁵⁰ ion.

Table 3. Photophysical Parameters and B_{CPL} Values of CPL-Active Eu(III) and Tb(III) Complexes Investigated in This Work

complex	ϵ (M ⁻¹ cm ⁻¹) [λ_{abs} (nm)]	ϕ_{ovl} (%)	$ g_{\text{lum}} $ [λ (nm)]	B_{CPL} (M ⁻¹ cm ⁻¹) ^a
[Eu(bpcd)(tta)]	17000 (350)	26	0.26 (586)	13.2
			0.13 (595)	7.8
			0.02 (615)	22.3
[Eu(bpcd)(Coum)]	12000 (318)	7	0.06 (596)	1.4
			0.01 (614)	2.0
			0.01 (624)	0.8
			0.04 (555)	6.9
[Tb(bpcd)(Coum)]	12000 (313)	55	0.05 (537)	11.7
			0.02 (547)	12.0
			0.04 (555)	6.9

^aCalculated according to a modified formula (see the Supporting Information for more details).

solvated Eu(III) ion and the ligands, with the stepwise formation constants for the 1:3 [$\log K_3 = 2.33$ for the reaction $\text{Eu}(\text{tta})_2 + \text{tta} \rightleftharpoons \text{Eu}(\text{tta})_3$]⁴⁷ and 1:2 [$\log K_2 = 2.97$ for the reaction $\text{Eu}(\text{Coum}) + \text{Coum} \rightleftharpoons \text{Eu}(\text{Coum})_2$] ratios,⁴⁸ clearly higher, due to the weaker solvation of the metal complex and the ligands in methanol than in water.⁴⁹

Total Luminescence (TL), CPL, and Luminescence Decay Kinetics. [Eu(bpcd)(tta)]. The luminescence excitation and emission spectra of a methanol solution (50 μM) of the complex are shown in Figure S3. The excitation spectrum is very similar to the absorption spectrum, and the presence of two excitation peaks at 270 and 350 nm is related to an efficient ligand-to-metal energy transfer involving both ligands. The emission spectrum is dominated by the presence of the hypersensitive $^5D_0 \rightarrow ^7F_2$ transition of Eu(III), which is usually observed when the emitting Eu(III) is located in a site, whose point symmetry lacks the inversion center. The high value (9.54) of the asymmetry ratio

$$R = \frac{I(^5D_0 \rightarrow ^7F_2)}{I(^5D_0 \rightarrow ^7F_1)} \quad (1)$$

is compatible with a highly distorted geometric environment around the metal ion.

The luminescence decay curve of the 5D_0 excited state of Eu(III) in methanol can be properly fitted by a single-exponential function (Figure 6, right). The calculated observed lifetime [0.68(1) ms] is reported in Table 2.

To estimate the number of methanol molecules in the inner coordination sphere of the Eu(III) ion (m), the observed lifetime was also measured in CD₃OD (1.18 ms). The value obtained here for m is 1.3(5) and has been determined by using the equation $m = 2.1(1/\tau_{\text{MeOH}} - 1/\tau_{\text{CD}_3\text{OD}})$ (Table 2).⁵⁰ This number has a significant impact on the luminescence efficiency of the emitting lanthanide ion. In fact, the OH group of methanol possesses high-energy vibrations (3300–3400 cm^{-1}), which are particularly effective in the nonradiative

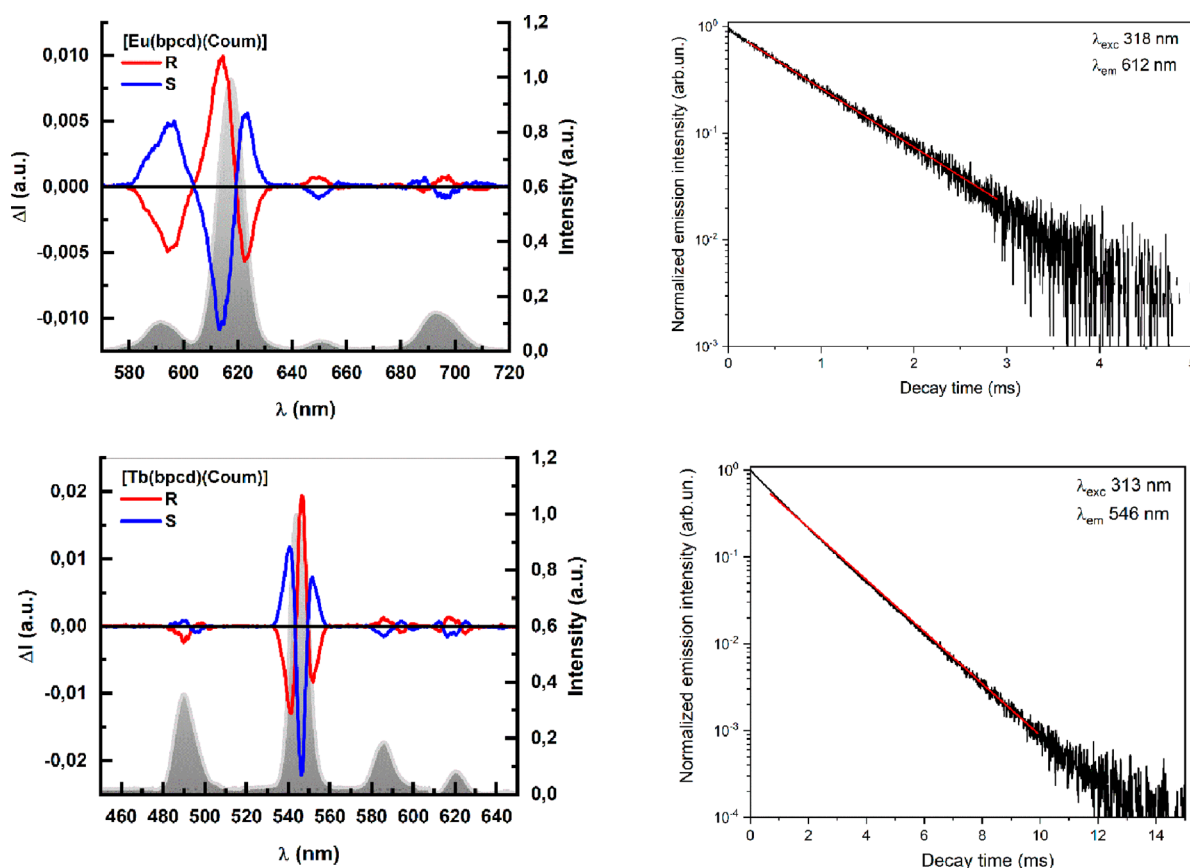


Figure 7. Overlap of the CPL and normalized total luminescence spectra (left) and luminescence decay curves (right) of [Eu(bpcd)(Coum)] (top) and [Tb(bpcd)(Coum)] (bottom) complexes. The luminescence spectra were recorded upon excitation at 254 nm. The decay curves of the S₁S isomers are shown and chosen as a representative. The equations of the fitting curve (red line) are $y = 0.93577 \exp(-t/0.80) + 0.0004$ [reduced $\chi^2 = 1.6 \times 10^{-4}$; R^2 (COD) = 0.99531] for [Eu(bpcd)(Coum)] and $y = 0.965 \exp(-t/1.42) + 0.004$ [reduced $\chi^2 = 3.2 \times 10^{-6}$; R^2 (COD) = 0.99981] for [Tb(bpcd)(Coum)].

quenching of the lanthanide emitting level, by means of the multiphonon relaxation process (MPR).⁵¹ A higher m value would lead to stronger quenching of the Ln(III) luminescence. By using the equation reported by Werts,⁵² which is known to be applicable to only the emission spectra of Eu(III), we calculated the radiative lifetime (τ_{rad}). We also determined the intrinsic quantum yield ($\phi_{\text{int}} = \tau_{\text{obs}}/\tau_{\text{rad}}$; i.e., the ratio of emitting/absorbed photons upon direct excitation into a luminescent level of the lanthanide ion) and the overall quantum yield upon excitation of the tta ligand (ϕ_{ovl} ; i.e., the ratio of emitting/absorbed photons upon excitation of the ligand) by using the secondary methods described in the literature⁵³ and η_{sens} which is the overall energy transfer efficiency ($\eta_{\text{sens}} = \phi_{\text{ovl}}/\phi_{\text{int}}$). It is worth noting that the obtained value of η_{sens} is $\sim 100\%$, which underlines the high efficiency of the antenna effect of tta molecules in sensitizing the Eu(III) luminescence in the [Eu(bpcd)(tta)] complex.

The CPL spectra of [Eu(bpcd)(tta)] (Figure 6) show intense bands with g factors on the order of 10^{-1} and 10^{-2} for the $^5D_0 \rightarrow ^7F_1$ and 7F_2 transitions, respectively (see Table 3 and Figure S4). Notably, three and two bands are clearly resolved for the $^5D_0 \rightarrow ^7F_1$ and 7F_2 transitions, respectively, corresponding to the M_J splitting due to the crystal field.

[Eu(bpcd)(Coum)] and [Tb(bpcd)(Coum)]. The luminescence excitation and emission spectra of a methanol solution (50 μM) of the complexes are shown in Figure S3. Both excitation spectra are very similar to the corresponding

absorption spectra. The presence of the excitation bands at 270 and 320 nm indicates, also for these Coum-based complexes, the presence of an efficient ligand-to-metal energy transfer involving both ligands. As in the case of the [Eu(bpcd)(tta)] complex, the emission spectrum of [Eu(bpcd)(Coum)] is dominated by the presence of the hypersensitive $^5D_0 \rightarrow ^7F_2$ transition of Eu(III) (Figure S3), although the geometric environment of the metal ion is less distorted than in the case of the tta-based complex, as suggested by the lower value of the asymmetry ratio (7.13). In the case of the [Tb(bpcd)(Coum)] complex upon excitation in the ligand absorption bands, the typical Tb(III) emission stemming from the $f-f$ transition is detected (Figure S3).

The luminescence decay curves of the excited states of Eu(III) and Tb(III) (5D_0 and 5D_4 levels, respectively) are fitted by a single-exponential function (Figure 7), and the calculated observed lifetimes (reported in Table 2) are 0.80(1) and 1.42(1) ms for the Eu(III) and Tb(III) complexes, respectively. As discussed above, the number of methanol molecules bound to the metal ion (m) can be obtained by measuring the observed lifetimes also in deuterated methanol. This number is slightly greater than 1 in the case of the [Eu(bpcd)(Coum)] complex and slightly less than 1 for [Tb(bpcd)(Coum)] (Table 2). A possible explanation for this behavior is given in DFT Calculations. A similar trend was observed by Arauzo et al.⁵⁴ in heteroleptic Eu and Tb complexes containing the Coum and phenanthroline ligands.

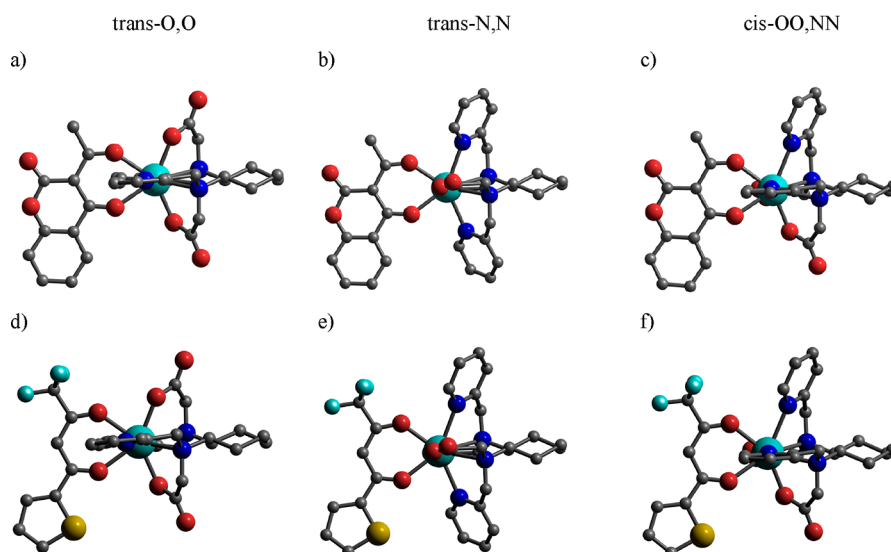


Figure 8. Minimum energy structures of the (a–c) [Y(bpcd)(Coum)] and (d–f) [Y(bpcd)(tta)] complexes obtained from DFT calculations. Hydrogen atoms bound to carbons have been omitted for the sake of clarity.

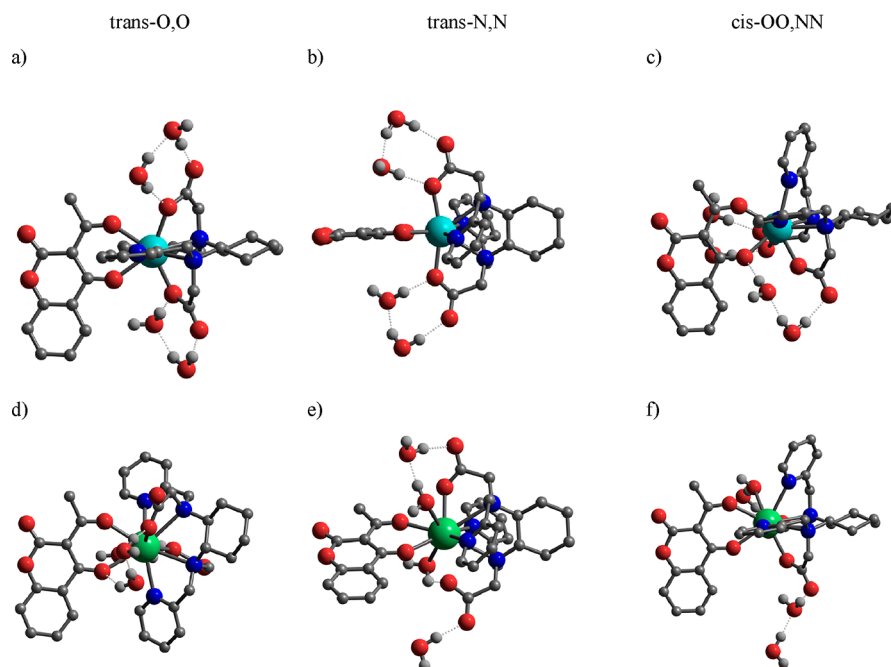


Figure 9. Minimum energy structures of the (a–c) [Y(bpcd)(Coum)]·4H₂O and (d–f) [La(bpcd)(Coum)]·4H₂O complexes obtained from DFT calculations. Hydrogen atoms bound to carbons have been omitted for the sake of clarity.

They observed the presence of one solvent molecule (H₂O) in the inner coordination sphere for only the Eu(III) derivative, while the Tb(III) complex lacks a coordinated solvent. Interestingly, we can also conclude that the Coum ligand is not a good sensitizer for Eu(III) luminescence (as η_{sens} is only ~21%) but can better sensitize Tb(III) luminescence [$\phi_{\text{ovl}} = 55\%$; $\eta_{\text{sens}} \geq 55$ (Table 2)]. This statement is in agreement with previous works on Tb(III) and Eu(III) tris chelates of Coum,²⁷ where the values of ϕ_{ovl} are 29% and 12%, respectively, and coumarin dipicolinate Eu(III) complexes ($\phi_{\text{ovl}} < 2\%$).⁵⁵ When heteroleptic complexes of the Coum ligand contain phenanthroline or bathophenanthroline ligands, the overall quantum yields increase for both Eu(III) (40–45%) and Tb(III) (58–76%).⁵⁴ This feature is related to the

involvement of the phenanthroline-based ligands in the sensitization of Ln(III) luminescence.

As expected, both CPL spectra of [Eu(bpcd)(Coum)] and [Tb(bpcd)(Coum)] (Figure 7) show mirror image spectra. Unlike the pattern of bands shown by [Eu(bpcd)(tta)] for the $^5D_0 \rightarrow ^7F_1$ transition, the coumarin complex has a single monosignate band (Figure 7). [Tb(bpcd)(Coum)] shows transitions associated with $^5D_4 \rightarrow ^7F_{6,5,4,3}$, with the most prominent feature being the manifold $^5D_4 \rightarrow ^7F_5$ transition around 547 nm. Despite the relatively low g_{lum} factors (Figure S6), because of its high quantum yield, [Tb(bpcd)(Coum)] shows high values of B_{CPL} , comparable to those obtained for [Eu(bpcd)(tta)] (Table 3).

A survey of the literature data reveals that the values of ϕ_{ovl} and g_{lum} of [Eu(bpcd)(tta)] and of B_{CPL} for both [Tb(bpcd)-(Coum)] and [Eu(bpcd)(tta)] complexes are in line with the average values reported for Eu(III) and Tb(III) luminescent complexes.⁵⁷

DFT Calculations. DFT calculations were carried out to obtain structural information about the [Ln(bpcd)L] complexes by studying the diamagnetic analogues of the Eu and Tb complexes. As previously described,^{30,58} several isomeric forms can be present, depending on the arrangement of the pyridine and acetate groups in the bpcd ligand, namely, *trans*-OO, *trans*-NN, and *cis*-OO,NN. These isomers can host either the Coum (Figure 8a–c) or tta (Figure 8d–f) ligand, which replaces the coordinated solvent molecules. The energy of the isomers does not differ significantly ($\Delta E < 0.3$ and 1.0 kcal mol⁻¹ for the adduct with Coum and tta, respectively), so that a mixture should be present in solution.

In our previous work,³⁰ the number of water molecules coordinated to the metal ion in the structures of [Y(bpcd)-H₂O_{*n*}]⁺ (*n* = 2–5) complexes was always 2. The additional waters remained dissociated and formed hydrogen bonds with the ligand or the coordinated ones. Nevertheless, a similar calculation for the [La(bpcd)H₂O_{*n*}]⁺ counterpart showed that more than two water molecules could interact directly with the metal ion.

The calculated number of coordinated methanol molecules in this study is 1.3 and 1.1 for Eu(III) and 0.6 for Tb(III). To confirm the solvent coordination, we considered the [Y(bpcd)(Coum)]·4H₂O complexes, in which water was employed in place of methanol to reduce the number of degrees of freedom of the system. The first result is that water cannot interact with the metal in all [Y(bpcd)(Coum)]·4H₂O isomers (Figure 9a–c). Second, the differences in energy [$\Delta E = E(\text{isomer}) - E(\text{a})$] are 1.2 and 4.6 kcal mol⁻¹ for structures b and c, respectively, in agreement with our previous observation that the *cis*-OO,NN isomer was the less stable one.

In the obtained minimum energy structures, the shortest metal–O_{water} distances fall in the range of 4.2–4.5 Å. Even if it has been proposed that closely diffusing solvent molecules (like those shown in Figure 9a–c) can contribute to decrease the luminescence decay lifetime, the number of methanol molecules calculated from the experimental measurements is definitely greater than 1 for Eu(III).

In the [La(bpcd)(Coum)]·4H₂O structures (Figure 9d–f), on the contrary, two water molecules can coordinate to the metal ion in the *trans*-OO and *trans*-NN isomers (La–O_{water} bond lengths of 2.64–2.97 Å), while one is found in the *cis*-OO,NN form (La–O_{water} distance of 2.77 Å). The energy difference between the different structures is <2.8 kcal mol⁻¹, which suggests the presence of a mixture in solution. Because La(III) and Y(III) have ionic radii, longer and shorter than that of Eu(III),⁶ respectively, it is likely that the small differences allow the coordination of an additional solvent molecule in solution. Interestingly, the experimental *m* (Table 2) clearly decreases from Eu (1.3 or 1.1) to Tb (0.6), in agreement with the hypothesis that small changes in ionic radius here determine a different number of coordinated solvent molecules in the complex. Therefore, in this case, Y(III) models are more representative of the [Tb(bpcd)-(Coum)] coordination in solution.

The energies of the S₁ and T₁ excited states for both tta and Coum ligands were determined by TD-DFT calculations for the Y- and La-based complexes (Table 4). The HOMO and

Table 4. Energies of the S₁ and T₁ States (cm⁻¹) Obtained from TD-DFT Calculations and Estimated Donor–Acceptor Distances, R_L (Å), Corresponding to the Unoccupied Molecular Orbitals Centroid to the Metal Ion

complex	S ₁	T ₁	difference	R _L
[Y(bpcd)(Coum)]				
<i>trans</i> -O,O	31514	25636	5878	3.37
<i>trans</i> -N,N	31865	25606	6259	3.47
<i>cis</i> -O,O-N,N	31769	25612	6157	3.28
[Y(bpcd)(tta)]				
<i>trans</i> -O,O	29008	19640	9368	3.51
<i>trans</i> -N,N	29138	19582	9556	3.54
<i>cis</i> -O,O-N,N	29122	19631	9491	3.50
[La(bpcd)(Coum)(H ₂ O) _{<i>n</i>}]				
<i>trans</i> -O,O (<i>n</i> = 1)	32614	26033	6581	3.98
<i>trans</i> -N,N (<i>n</i> = 2)	32552	25882	6670	4.08
<i>cis</i> -O,O-N,N (<i>n</i> = 1)	32045	25632	6413	3.83
[La(bpcd)(tta)(H ₂ O) _{<i>n</i>}]				
<i>trans</i> -O,O (<i>n</i> = 1)	29674	20007	9667	3.89
<i>trans</i> -N,N (<i>n</i> = 2)	29671	20026	9645	3.96
<i>cis</i> -O,O-N,N (<i>n</i> = 1)	29179	19512	9667	3.76

LUMO molecular orbitals, mainly localized on these two ligands, are mostly involved in the S₀ → S₁ and S₀ → T₁ transitions (Figures S7–S10), and the two most intense peaks at lower energies [around 315 nm for Coum-based complexes and around 350 nm for tta-based complexes (Figures 5 and 6)] are related to these S₀ → S₁ transitions. As previously demonstrated by some of us, at shorter wavelengths (at 270 nm), an additional electronic transition involving the molecular orbitals localized on the bpcd ligand can be exploited to sensitize Eu(III)⁴³ and Tb(III)³⁰ luminescence (not discussed here).

Assuming the participation of the excited triplet state in the *antenna* process, the sensitization of Tb(III) luminescence by tta can be ruled out. In fact, the lowest excited state of Tb(III) (⁵D₄) is located around 20500 cm⁻¹; that being above the energy of the triplet state of the ligand (19618 cm⁻¹ averaged on the possible isomers) could be responsible for a back energy transfer mechanism (from the metal to the ligand) that makes the *antenna* process ineffective. On the contrary, the energy position of the T₁ state of tta seems to be optimal to transfer energy to the ⁵D₀ emitting level of Eu(III) (located at 17300 cm⁻¹), in agreement with the calculated η_{sens} of ~100% (Table 2). As for the energy transfer mechanism in the case of the Coum ligand, both the emitting ⁵D₄ level of Tb(III) located at 20500 cm⁻¹⁵⁹ and the ⁵D₂ level of Eu(III) (around 21500 cm⁻¹) can be considered suitable acceptor states of the T₁ donor state of the Coum molecule [around 25600/26000 cm⁻¹ (Table 4)]. However, in our complexes, the sensitization efficiency of Eu(III) luminescence by the Coum ligand is lower than that of Tb(III) (η_{sens} values around 20% and ≥55% for the [Eu(bpcd)(Coum)] and [Tb(bpcd)(Coum)] complexes, respectively). Although a detailed study of the dynamics of the energy transfer process would be necessary to clearly understand the reasons underlying this behavior, a possible explanation for the lower sensitization efficiency in the case of the Eu(III) complex can be found in the D–A (donor–acceptor) distance (R_L) and the S₁–T₁ energy gap (fifth and fourth columns, respectively, in Table 4). In fact, it is well-known that the R_L distance strongly affects the probability of energy transfer from D to A: the shorter the R_L, the higher the

probability.⁶⁰ As it can be evinced from the inspection of Table 4, with an increase in the ionic radius of the lanthanide ion (passing from Y to La), the D–A distance increases. Therefore, we expect a sensitization efficiency that improves as the lanthanide ion becomes smaller. The decrease ($\Delta r_l = 0.03 \text{ \AA}$) in the ionic radius (and in turn of the R_L distance) by passing from the Eu(III) to Tb(III) ion can justify the higher sensitization efficiency observed in the case of the Tb(III)-based complex. Analogously, it seems that an increase in the S_1 – T_1 energy gap occurs with an increase in the size of the metal ion [from Y to La (Table 4) and presumably from Tb to Eu]. The increase in this energy gap is often related to a decrease of the intersystem crossing and sensitization efficiencies. Finally, as usual for lanthanide-based coordination compounds, in the energy transfer process we can assume the exchange mechanism as dominant, being the most sensitive to the D–A distance.⁶⁰ Therefore, its involvement could account for the significant decrease in sensitization efficiency, even if only a very small increase in the ionic radius (and in turn the R_L distance) occurs by passing from the Tb(III) to Eu(III) ion.

CONCLUSIONS

Upon excitation of the antenna ligands, a moderate to good overall quantum yield is measured for the [Eu(bpcd)(tta)] (26%) and [Tb(bpcd)(Coum)] (55%) complexes. This is mainly due to the very good sensitization efficiency of the Eu(III) and Tb(III) luminescence by tta and Coum antenna ligands, respectively. A shorter D–A distance and a better intersystem crossing efficiency, in the case of [Tb(bpcd)(Coum)], could account for the observed higher sensitization efficiency of the Coum ligand toward Tb(III) ($\geq 55\%$), with respect to Eu(III) (21%). In addition, the access to the first coordination sphere of the metal ion by the methanol molecules, which decrease the luminescence efficiency, is more limited in the case of the Tb(III) complex ($m = 0.6$) than in the case of the Eu(III) complex ($m = 1.1$). This is probably due to an ionic size effect, Eu(III) being slightly larger than Tb(III) ($\Delta r = 0.03 \text{ \AA}$; r is the ionic radius in water). As suggested by the D–A distances (R_L), the Coum ligand is more tightly bound to the metal ion than tta, and this reflects the fact that the stability of [Eu(bpcd)(Coum)] ($\log K = 5.59$) is higher than that of [Eu(bpcd)(tta)] ($\log K = 4.12$). The ECD experiments suggested that the chiral bpcd ligand can dictate a preferred sense of twist of both tta and Coum ligands and the CPL activity of the magnetic dipole transition around 586 nm of Eu(III) in [Eu(bpcd)(tta)] complex is remarkable ($g_{\text{lum}} = 0.26$). The reported values of B_{CPL} for these complexes are in line with the average values reported in the literature for Eu(III) and Tb(III) luminescent complexes.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.inorgchem.3c00196>.

Excitation and emission spectra and plots of g_{lum} versus wavelength of the Eu(III) and Tb(III) complexes under investigation; Kohn–Sham molecular orbital composition of the S_1 and T_1 states for the Y(III) and La(III) counterparts; additional experimental details, materials, and methods of the chiroptical instrumentation and definition of the B_{CPL} formula employed in this work; and ^1H and ^{13}C NMR data of the Coum ligand (PDF)

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Notes

The authors declare no competing financial interest.

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ADDITIONAL NOTES

^aBecause (i) ϕ_{int} , ϕ_{ovl} , and η_{sens} can assume values between 0 and 1, (ii) ϕ_{ovl} must be less than or equal to ϕ_{int} , and (iii) ϕ_{ovl} is 55% for the [Tb(bpcd)(Coum)] complex, we can conclude that η_{sens} must be $\geq 55\%$.

^bThe recently revised⁶¹ ionic radii in a water solution are 1.250 (1.216), 1.12 (1.066), and 1.090 (1.04) Å for La(III), Eu(III), and Tb(III), respectively (Shannon radii⁶² in parentheses). For Y(III), the ionic radius calculated from the Y–O(water) distance obtained experimentally⁶³ using the water radius proposed in ref 61 is 1.020 (1.019) Å.

REFERENCES

- (1) Frawley, A. T.; Pal, R.; Parker, D. Very Bright, Enantiopure Europium(III) Complexes Allow Time-Gated Chiral Contrast Imaging. *Chem. Commun.* **2016**, 52 (91), 13349–13352.
- (2) Çoruh, N.; Riehl, J. P. Circularly Polarized Luminescence from Terbium(III) as a Probe of Metal Ion Binding in Calcium-Binding Proteins. *Biochemistry* **1992**, 31 (34), 7970–7976.
- (3) Abdollahi, S.; Harris, W. R.; Riehl, J. P. Application of Circularly Polarized Luminescence Spectroscopy to Tb(III) and Eu(III) Complexes of Transferrins. *J. Phys. Chem.* **1996**, 100 (5), 1950–1956.
- (4) Yuasa, J.; Ohno, T.; Tsumatori, H.; Shiba, R.; Kamikubo, H.; Kataoka, M.; Hasegawa, Y.; Kawai, T. Fingerprint Signatures of Lanthanide Circularly Polarized Luminescence from Proteins Covalently Labeled with a β -Diketonate Europium(III) Chelate. *Chem. Commun.* **2013**, 49 (41), 4604–4606.
- (5) Leonzio, M.; Melchior, A.; Faura, G.; Tolazzi, M.; Bettinelli, M.; Zinna, F.; Arrico, L.; Di Bari, L.; Piccinelli, F. A Chiral Lactate Reporter Based on Total and Circularly Polarized Tb(III) Luminescence. *New J. Chem.* **2018**, 42 (10), 7931–7939.
- (6) Wong, H.-Y.; Lo, W.-S.; Yim, K.-H.; Law, G.-L. Chirality and Chiroptics of Lanthanide Molecular and Supramolecular Assemblies. *Chem.* **2019**, 5 (12), 3058–3095.
- (7) Kitagawa, Y.; Wada, S.; Islam, M. D. J.; Saita, K.; Gon, M.; Fushimi, K.; Tanaka, K.; Maeda, S.; Hasegawa, Y. Chiral Lanthanide Lumino-Glass for a Circularly Polarized Light Security Device. *Commun. Chem.* **2020**, 3 (1), 119.
- (8) Willis, O. G.; Zinna, F.; Di Bari, L. NIR-Circularly Polarized Luminescence from Chiral Complexes of Lanthanides and d-Metals. *Angew. Chem., Int. Ed.* **2023**, No. e202302358.
- (9) Willis, O. G.; Petri, F.; Pescitelli, G.; Pucci, A.; Cavalli, E.; Mandoli, A.; Zinna, F.; Di Bari, L. Efficient 1400–1600 Nm Circularly Polarized Luminescence from a Tuned Chiral Erbium Complex. *Angew. Chem., Int. Ed.* **2022**, 61 (34), No. e202208326.
- (10) Mukhtar, N. F. M.; Schley, N. D.; Ung, G. Strong Circularly Polarized Luminescence at 1550 Nm from Enantiopure Molecular Erbium Complexes. *J. Am. Chem. Soc.* **2022**, 144 (14), 6148–6153.
- (11) Willis, B.-A. N.; Schnable, D.; Schley, N. D.; Ung, G. Spinolate Lanthanide Complexes for High Circularly Polarized Luminescence Metrics in the Visible and Near-Infrared. *J. Am. Chem. Soc.* **2022**, 144 (49), 22421–22425.
- (12) Willis, O. G.; Pucci, A.; Cavalli, E.; Zinna, F.; Di Bari, L. Intense 1400–1600 Nm Circularly Polarized Luminescence from Homo- and Heteroleptic Chiral Erbium Complexes. *J. Mater. Chem. C* **2023**, 11 (16), 5290–5296.
- (13) Dhbaibi, K.; Grasser, M.; Douib, H.; Dorcet, V.; Cador, O.; Vanthuyne, N.; Riobé, F.; Maury, O.; Guy, S.; Bensalah-Ledoux, A.; Baguenard, B.; Rikken, G. L. J. A.; Train, C.; Le Guennic, B.; Atzori, M.; Pointillart, F.; Crassous, J. Multifunctional Helicene-Based Ytterbium Coordination Polymer Displaying Circularly Polarized Luminescence, Slow Magnetic Relaxation and Room Temperature Magneto-Chiral Dichroism. *Angew. Chem., Int. Ed.* **2023**, 62 (5), No. e202215558.
- (14) Lefeuvre, B.; Mattei, C. A.; Gonzalez, J. F.; Gendron, F.; Dorcet, V.; Riobé, F.; Lalli, C.; Le Guennic, B.; Cador, O.; Maury, O.; Guy, S.; Bensalah-Ledoux, A.; Baguenard, B.; Pointillart, F. Solid-State Near-Infrared Circularly Polarized Luminescence from Chiral Yb(III)-Single-Molecule Magnet. *Chem. - Eur. J.* **2021**, 27 (26), 7362–7366.
- (15) Adewuyi, J. A.; Schley, N. D.; Ung, G. Vanol-Supported Lanthanide Complexes for Strong Circularly Polarized Luminescence at 1550 Nm. *Chem. - Eur. J.* **2023**, No. e202300800.
- (16) Geng, Y.; Trajkovska, A.; Culligan, S. W.; Ou, J. J.; Chen, H. M. P.; Katsis, D.; Chen, S. H. Origin of Strong Chiroptical Activities in Films of Nonfluorenes with a Varying Extent of Pendant Chirality. *J. Am. Chem. Soc.* **2003**, 125 (46), 14032–14038.
- (17) Yang, Y.; Da Costa, R. C.; Smilgies, D. M.; Campbell, A. J.; Fuchter, M. J. Induction of Circularly Polarized Electroluminescence from an Achiral Light-Emitting Polymer via a Chiral Small-Molecule Dopant. *Adv. Mater.* **2013**, 25 (18), 2624–2628.
- (18) Zinna, F.; Pasini, M.; Galeotti, F.; Botta, C.; Di Bari, L.; Giovannella, U. Design of Lanthanide-Based OLEDs with Remarkable Circularly Polarized Electroluminescence. *Adv. Funct. Mater.* **2017**, 27 (1), 1603719.
- (19) Brandt, J. R.; Wang, X.; Yang, Y.; Campbell, A. J.; Fuchter, M. J. Circularly Polarized Phosphorescent Electroluminescence with a High Dissymmetry Factor from PHOLEDs Based on a Platinahelicene. *J. Am. Chem. Soc.* **2016**, 138 (31), 9743–9746.
- (20) Arrico, L.; Di Bari, L.; Zinna, F. Quantifying the Overall Efficiency of Circularly Polarized Emitters. *Chem. - Eur. J.* **2021**, 27 (9), 2920–2934.
- (21) Stachelek, P.; MacKenzie, L.; Parker, D.; Pal, R. Circularly Polarized Luminescence Laser Scanning Confocal Microscopy to Study Live Cell Chiral Molecular Interactions. *Nat. Commun.* **2022**, 13 (1), 553.
- (22) MacKenzie, L. E.; Pal, R. Circularly Polarized Lanthanide Luminescence for Advanced Security Inks. *Nat. Rev. Chem.* **2021**, 5 (2), 109–124.
- (23) De Rosa, D. F.; Stachelek, P.; Black, D. J.; Pal, R. Rapid Handheld Time-Resolved Circularly Polarized Luminescence Photography Camera for Life and Material Sciences. *Nat. Commun.* **2023**, 14 (1), 1537.
- (24) Baguenard, B.; Bensalah-Ledoux, A.; Guy, L.; Riobé, F.; Maury, O.; Guy, S. Theoretical and Experimental Analysis of Circularly Polarized Luminescence Spectrophotometers for Artifact-Free Measurements Using a Single CCD Camera. *Nat. Commun.* **2023**, 14 (1), 1065.
- (25) Wong, K.-L.; Bünzli, J.-C. G.; Tanner, P. A. Quantum Yield and Brightness. *J. Lumin.* **2020**, 224, 117256.
- (26) Zinna, F.; Di Bari, L. Lanthanide Circularly Polarized Luminescence: Bases and Applications. *Chirality* **2015**, 27 (1), 1–13.
- (27) Guzmán-Méndez, O.; González, F.; Bernès, S.; Flores-Alamo, M.; Ordóñez-Hernández, J.; García-Ortega, H.; Guerrero, J.; Qian, W.; Aliaga-Alcalde, N.; Gasque, L. Coumarin Derivative Directly Coordinated to Lanthanides Acts as an Excellent Antenna for UV–Vis and Near-IR Emission. *Inorg. Chem.* **2018**, 57 (3), 908–911.
- (28) Binnemans, K. *Chapter 225 - Rare-Earth Beta-Diketonates*; Gschneidner, K. A., Bünzli, J.-C. G., Pecharsky, V. K. B. T.-H., Eds.; Elsevier, 2005; Vol. 35, pp 107–272.
- (29) Piccinelli, F.; Nardon, C.; Bettinelli, M.; Melchior, A.; Tolazzi, M.; Zinna, F.; Di Bari, L. Lanthanide-Based Complexes Containing a Chiral Trans-1,2-Diaminocyclohexane (DACH) Backbone: Spectro-

scopic Properties and Potential Applications. *ChemPhotoChem* **2022**, *6* (2), No. e202100143.

(30) Leonzio, M.; Melchior, A.; Faura, G.; Tolazzi, M.; Zinna, F.; Di Bari, L.; Piccinelli, F. Strongly Circularly Polarized Emission from Water-Soluble Eu(III)- and Tb(III)-Based Complexes: A Structural and Spectroscopic Study. *Inorg. Chem.* **2017**, *56* (8), 4413–4422.

(31) Zinna, F.; Bruhn, T.; Guido, C. A.; Ahrens, J.; Bröring, M.; Di Bari, L.; Pescitelli, G. Circularly Polarized Luminescence from Axially Chiral BODIPY DYEmers: An Experimental and Computational Study. *Chem. - Eur. J.* **2016**, *22* (45), 16089–16098.

(32) Gans, P.; Sabatini, A.; Vacca, A. Investigation of Equilibria in Solution. Determination of Equilibrium Constants with the HYPERQUAD Suite of Programs. *Talanta* **1996**, *43* (10), 1739–1753.

(33) Frisch, M. J.; Trucks, G. W.; Schlegel, H. B.; Scuseria, G. E.; Robb, M. A.; Cheeseman, J. R.; Scalmani, G.; Barone, V.; Petersson, G. A.; Nakatsuji, H.; Li, X.; Caricato, M.; Marenich, A. V.; Bloino, J.; Janesko, B. G.; Gomperts, R.; Mennucci, B.; Hratchian, H. P.; Ortiz, J. V.; Izmaylov, A. F.; Sonnenberg, J. L.; Williams-Young, D.; Ding, F.; Lipparini, F.; Egidi, F.; Goings, J.; Peng, B.; Petrone, A.; Henderson, T.; Ranasinghe, D.; Zakrzewski, V. G.; Gao, J.; Rega, N.; Zheng, G.; Liang, W.; Hada, M.; Ehara, M.; Toyota, K.; Fukuda, R.; Hasegawa, J.; Ishida, M.; Nakajima, T.; Honda, Y.; Kitao, O.; Nakai, H.; Vreven, T.; Throssell, K.; Montgomer, J. A., Jr.; Peralta, J. E.; Ogliaro, F.; Bearpark, M. J.; Heyd, J. J.; Brothers, E. N.; Kudin, K. N.; Staroverov, V. N.; Keith, T. A.; Kobayashi, R.; Normand, J.; Raghavachari, K.; Rendell, A. P.; Burant, J. C.; Iyengar, S. S.; Tomasi, J.; Cossi, M.; Millam, J. M.; Klene, M.; Adamo, C.; Cammi, R.; Ochterski, J. W.; Martin, R. L.; Morokuma, K.; Farkas, O.; Foresman, J. B.; Fox, D. J. *Gaussian 16*, rev. A03; Gaussian, Inc.: Wallingford, CT, 2016.

(34) Arrico, L.; De Rosa, C.; Di Bari, L.; Melchior, A.; Piccinelli, F. Effect of the Counterion on Circularly Polarized Luminescence of Europium(III) and Samarium(III) Complexes. *Inorg. Chem.* **2020**, *59* (7), 5050–5062.

(35) Wang, J.; Wang, Y.; Zhang, Z. H.; Zhang, X. D.; Tong, J.; Liu, X. Z.; Liu, X. Y.; Zhang, Y.; Pan, Z. J. Syntheses, Characterization, and Structure Determination of Nine-Coordinate $\text{Na}[\text{Y}^{\text{III}}(\text{Edta})(\text{H}_2\text{O})_3]\cdot 5\text{H}_2\text{O}$ and Eight-Coordinate $\text{Na}[\text{Y}^{\text{III}}(\text{Cydta})(\text{H}_2\text{O})_2]\cdot 5\text{H}_2\text{O}$ Complexes. *J. Struct. Chem.* **2005**, *46* (5), 895–905.

(36) Mondry, A.; Janicki, R. From Structural Properties of the EuIII Complex with Ethylenediaminetetra(Methylenephosphonic Acid) (H_8EDTMP) towards Biomedical Applications. *Dalton Trans.* **2006**, *39*, 4702–4710.

(37) Wang, J.; Hu, P.; Liu, B.; Xu, R.; Wang, X.; Wang, D.; Zhang, L. Q.; Zhang, X. D. $\text{NH}_4[\text{Eu}^{\text{III}}(\text{Cydta})(\text{H}_2\text{O})_2]\cdot 4.5\text{H}_2\text{O}$ and $\text{K}_2[\text{Eu}_2^{\text{III}}(\text{Pdta})_2(\text{H}_2\text{O})_2]\cdot 6\text{H}_2\text{O}$. *J. Struct. Chem.* **2011**, *52* (3), 568.

(38) Lee, C. T.; Yang, W. T.; Parr, R. G. Development of the Colle-Salvetti Correlation-Energy Formula Into A Functional of the Electron-Density. *Phys.Rev.B* **1988**, *37* (2), 785–789.

(39) Becke, A. D. A New Mixing of Hartree-Fock and Local Density-Functional Theories. *J. Chem. Phys.* **1993**, *98* (2), 1372–1377.

(40) Cao, X.; Dolg, M. Segmented Contraction Scheme for Small-Core Lanthanide Pseudopotential Basis Sets. *J. Mol. Struct. THEOCHEM* **2002**, *581* (1), 139–147.

(41) Andrae, D.; Häussermann, U.; Dolg, M.; Stoll, H.; Preuss, H. Energy-Adjusted Ab Initio Pseudopotentials for the Second and Third Row Transition Elements. *Theor. Chim. Acta* **1990**, *77* (2), 123–141.

(42) Tomasi, J.; Mennucci, B.; Cammi, R. Quantum Mechanical Continuum Solvation Models. *Chem. Rev.* **2005**, *105* (8), 2999–3094.

(43) Carneiro Neto, A. N.; Moura, R. T. J.; Carlos, L. D.; Malta, O. L.; Sanadar, M.; Melchior, A.; Kraka, E.; Ruggieri, S.; Bettinelli, M.; Piccinelli, F. Dynamics of the Energy Transfer Process in Eu(III) Complexes Containing Polydentate Ligands Based on Pyridine, Quinoline, and Isoquinoline as Chromophoric Antennae. *Inorg. Chem.* **2022**, *61* (41), 16333–16346.

(44) Van Bay, M.; Hien, N. K.; Tran, P. T. D.; Tuyen, N. T. K.; Oanh, D. T. Y.; Nam, P. C.; Quang, D. T. TD-DFT Benchmark for UV-Vis Spectra of Coumarin Derivatives. *Vietnam J. Chem.* **2021**, *59* (2), 203–210.

(45) Lu, T.; Chen, F. Multiwfn: A Multifunctional Wavefunction Analyzer. *J. Comput. Chem.* **2012**, *33* (5), 580–592.

(46) Andreiadis, E. S.; Gauthier, N.; Imbert, D.; Demadrille, R.; Pécaut, J.; Mazzanti, M. Lanthanide Complexes Based on β -Diketonates and a Tetradentate Chromophore Highly Luminescent as Powders and in Polymers. *Inorg. Chem.* **2013**, *52* (24), 14382–14390.

(47) Ikeda, N.; Kimura, K.; Asai, H.; Oshima, N. Extraction Behavior of Europium with Thenoyltrifluoroacetone (TTA). *Radioisotopes* **1970**, *19* (1), 1–6.

(48) Manku, G. C. S. The Formation Constants of Some Metal Complexes with 3-Acetyl-4-Hydroxycoumarin, Dehydroacetic Acid, and Their Oximes. *Aust. J. Chem.* **1971**, *24* (5), 925–934.

(49) Di Bernardo, P.; Melchior, A.; Tolazzi, M.; Zanonato, P. L. Thermodynamics of Lanthanide(III) Complexation in Non-Aqueous Solvents. *Coord. Chem. Rev.* **2012**, *256* (1–2), 328–351.

(50) Holz, R. C.; Chang, C. A.; Horrocks, W. D. Spectroscopic Characterization of the Europium(III) Complexes of a Series of $\text{N,N}'$ -Bis(Carboxymethyl) Macrocylic Ether Bis(Lactones). *Inorg. Chem.* **1991**, *30* (17), 3270–3275.

(51) Weber, M. J. Radiative and Multiphonon Relaxation of Rare-Earth Ions in Y_2O_3 . *Phys. Rev.* **1968**, *171* (2), 283–291.

(52) Werts, M. H. V.; Jukes, R. T. F.; Verhoeven, J. W. The Emission Spectrum and the Radiative Lifetime of Eu^{3+} in Luminescent Lanthanide Complexes. *Phys. Chem. Chem. Phys.* **2002**, *4* (9), 1542–1548.

(53) Eaton, D. F. Reference Materials for Fluorescence Measurement. *Pure Appl. Chem.* **1988**, *60* (7), 1107–1114.

(54) Arauzo, A.; Gasque, L.; Fuertes, S.; Tenorio, C.; Bernès, S.; Bartolomé, E. Coumarin-Lanthanide Based Compounds with SMM Behavior and High Quantum Yield Luminescence. *Dalton Trans.* **2020**, *49* (39), 13671–13684.

(55) Di Pietro, S.; Iacopini, D.; Moscardini, A.; Bizzarri, R.; Pineschi, M.; Di Bussolo, V.; Signore, G. New Coumarin Dipicolinate Europium Complexes with a Rich Chemical Speciation and Tunable Luminescence. *Molecules* **2021**, *26* (5), 1265.

(56) Peacock, R. D. *The Intensities of Lanthanide $f \leftrightarrow f$ Transitions BT - Rare Earths*; Nieboer, E., Jorgensen, C. K., Peacock, R. D., Reisfeld, R., Eds.; Structure and Bonding; Springer: Berlin, 1975; Vol. 22.

(57) Islam, M. J.; Kitagawa, Y.; Tsurui, M.; Hasegawa, Y. Strong Circularly Polarized Luminescence of Mixed Lanthanide Coordination Polymers with Control of 4f Electronic Structures. *Dalton Trans.* **2021**, *50* (16), 5433–5436.

(58) McLauchlan, C. C.; Florián, J.; Kissel, D. S.; Herlinger, A. W. Metal Ion Complexes of $\text{N,N}'$ -Bis(2-Pyridylmethyl)-*Trans*-1,2-Diaminocyclohexane- $\text{N,N}'$ -Diacetic Acid, H_2 Bpcd: Lanthanide-(III)– Bpcd^{2-} Cationic Complexes. *Inorg. Chem.* **2017**, *56* (6), 3556–3567.

(59) Carnall, W. T.; Fields, P. R.; Rajnak, K. Electronic Energy Levels of the Trivalent Lanthanide Aquo Ions. II. Gd^{3+} . *J. Chem. Phys.* **1968**, *49* (10), 4443–4446.

(60) Dexter, D. L. A Theory of Sensitized Luminescence in Solids. *J. Chem. Phys.* **1953**, *21* (5), 836–850.

(61) D'Angelo, P.; Zitolo, A.; Migliorati, V.; Chillemi, G.; Duvail, M.; Vitorge, P.; Abadie, S.; Spezia, R. Revised Ionic Radii of Lanthanoid(III) Ions in Aqueous Solution. *Inorg. Chem.* **2011**, *50* (10), 4572–4579.

(62) Shannon, R. D. Revised Effective Ionic Radii and Systematic Studies of Interatomic Distances in Halides and Chalcogenides. *Acta Crystallogr., Sect. A* **1976**, *32* (5), 751–767.

(63) Lindqvist-Reis, P.; Lambie, K.; Pattanaik, S.; Persson, I.; Sandström, M. Hydration of the Yttrium(III) Ion in Aqueous Solution. An X-Ray Diffraction and XAFS Structural Study. *J. Phys. Chem. B* **2000**, *104* (2), 402–408.