Circularly polarized luminescence of chiral Eu(III) complexes with phenanthrene unit

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Chiral Eu(III) complexes exhibit strong circularly polarized luminescence (CPL) *via* photosensitized energy transfer from the ligand triplet (T₁) state.^{1,2} The T₁ spin-polarization is induced by sublevel-selective intersystem crossing.^{3,4} In this study, the effect of spin-polarization on the CPL properties of chiral Eu(III) complex containing two photosensitizers, 2,7bis(diphenylphosphoryl) phenanthrene (dpph) and (+)-3-(heptafluorobutyryl) camphorate ligands (+hfc), is reported for the first time.

The Eu(III) complex (**Fig. 1**, **Eudpph**) was synthesized by reacting $Eu(+hfc)_3(H_2O)_n$ with dpph in methanol. The compound was identified by ESI-MS and FT-IR. The

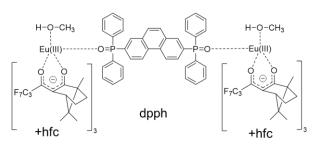


Fig. 1 Chemical structure of Eu-dpph.

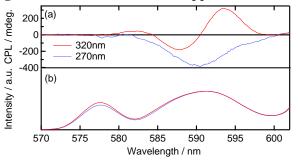


Fig. 2 (a) CPL and (b) emission spectra of **Eu-dpph** at 250 K. Red line: $\lambda_{ex} = 320$ nm, Blue line: $\lambda_{ex} = 270$ nm.

photophysical properties were characterized by absorption, emission, and CPL spectra.

Strong absorption bands were observed at 320 and 270 nm, which were assigned to the π - π^* transitions of +hfc and dpph, respectively. The CPL and emission spectra for the ${}^5D_0 \rightarrow {}^7F_{0,1}$ transitions are shown in **Fig. 2**. The positive and negative CPL signals at 594 and 588 nm were observed by +hfc excitation ($\lambda_{ex} = 320$ nm), while the dpph excitation ($\lambda_{ex} = 270$ nm) gave rise to one negative CPL signal at 590 nm. This is the first example of CPL depending on excitation wavelength for Eu(III) complexes, suggesting effective T₁ spin-polarization on the 4f-4f CPL properties.

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