

# Temperature influence on the emission intensity of a $\text{Eu}^{3+}/\text{Tb}^{3+}$ bimetallic complex with bipyrimidinic and beta-diketone ligands

Ariane Carolina Ferreira Beltrame<sup>1</sup>, Rodolpho Alessandro Nesta Silva<sup>2</sup>, Fernando Eduardo Maturi<sup>3</sup>, Luis D. Carlos<sup>3</sup>, Airton G. Bispo-Jr<sup>4</sup>, Flavia Artizzu<sup>2</sup>, Sergio Antonio Marques de Lima<sup>5</sup>, Ana Maria Pires<sup>5</sup>

<sup>1</sup>São Paulo State University, <sup>2</sup>Università degli Studi del Piemonte Orientale, <sup>3</sup>Universidade de Aveiro, <sup>4</sup>Universidade de São Paulo (*Institute of Chemistry*), <sup>5</sup>UNESP (*Chemistry and Biochemistry*)

*e-mail: ariane.beltrame@unesp.br*

Real-time, non-contact thermal monitoring is crucial in industrial applications where temperature directly affects device functionality. Developing efficient remote sensing technologies can overcome limitations of conventional contact-based methods, such as low spatial resolution [1]. In this sense, this study investigated the temperature-dependent emission behavior of a bimetallic Eu-Tb complex,  $[\text{TbEu}(\text{acac})_6\text{-bpm}]$ , using acetylacetonate (acac) as a sensitizer and 2,2'-bipyrimidine (bpm) as a bridging ligand. The complex was synthesized via a one-pot method ( $\text{Tb}^{3+}:\text{Eu}^{3+} = 1:1$ ;  $\text{acac}:\text{bpm} = 6:1$ )[2] and characterized by mass spectrometry (MS), diffuse reflectance (DR), and photoluminescence (PL). MS data confirmed the proposed structure, while the DR spectrum showed a maximum absorption at 263 nm, characteristic of the acac ligand. Emission spectra and time-resolved decay curves were collected over a temperature range of 12 K to 272 K, with intervals of 26 K and a fixed excitation wavelength at the ligand (330 nm). The emission spectra display characteristic  $\text{Eu}^{3+}$  ( ${}^5\text{D}_0 \rightarrow {}^7\text{F}_{0,5}$ ) and  $\text{Tb}^{3+}$  ( ${}^5\text{D}_4 \rightarrow {}^7\text{F}_6$ ,  ${}^5\text{D}_4 \rightarrow {}^7\text{F}_5$ ) transitions, with intensities decreasing as temperature rises due to enhanced non-radiative energy losses, reducing ligand-to-metal sensitization. Notably, the  $\text{Eu}^{3+}$  spectrum reveals the  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_5$  transition, which is typically weak or absent in conventional  $\text{Eu}^{3+}$  emission profiles, suggesting an altered electronic environment in the complex. Emission decay measurements ( $\lambda_{\text{ex}} = 330$  nm;  $\lambda_{\text{em}} = 702$  nm for  $\text{Eu}^{3+}$ , 540 nm for  $\text{Tb}^{3+}$ ) show a sigmoidal curve profile, with non-linear lifetime reductions of the  $\text{Eu}^{3+}$   ${}^5\text{D}_0$  and  $\text{Tb}^{3+}$   ${}^5\text{D}_4$  states, supporting emission trends. These findings highlight the system's potential for thermometric applications, as both emission intensity and lifetime respond to thermal stimuli.

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