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Polyoxometalates**

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# New robust luminescent supramolecular assemblies based on $[\text{Ln}(\text{Mo}_8\text{O}_{26})_2]^{5-}$ ( $\text{Ln} = \text{Eu}$ and $\text{Sm}$ ) polyoxometalates

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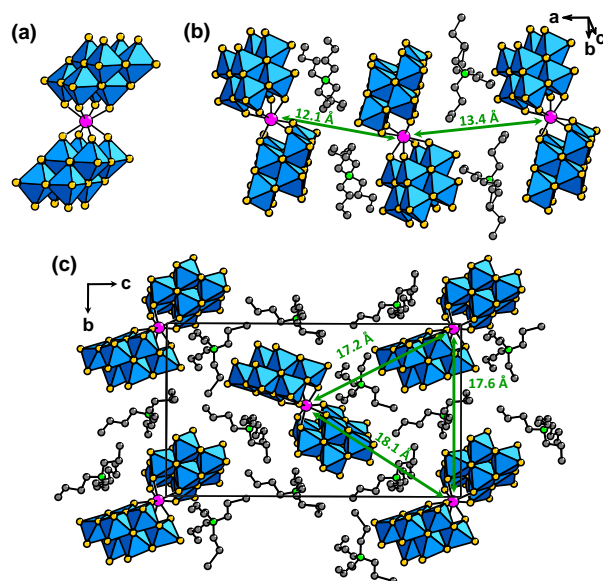
Supporting Information Placeholder

**ABSTRACT:** This work highlights for the first time the photoluminescence (PL) properties of two new  $[\text{Ln}(\text{Mo}_8\text{O}_{26})_2]^{5-}$  ( $\text{Ln} = \text{Eu}, \text{Sm}$ ) lanthanide-containing polyoxometalates. Stable crystals of the tetrabutylammonium salts  $(\text{TBA})_5[\text{Ln}(\text{Mo}_8\text{O}_{26})_2]$  were synthesized, and their structures were confirmed by single-crystal X-ray diffraction. The robustness of the  $[\text{Ln}(\text{Mo}_8\text{O}_{26})_2]^{5-}$  complexes in acetonitrile solution has been evidenced by FT-Raman and PL spectroscopies. Then, the tetraphenylphosphonium (TPP) $_5[\text{Ln}(\text{Mo}_8\text{O}_{26})_2]$  derivatives were obtained by a salt metathesis reaction. The two series exhibit a high thermal stability in air and are efficient solid-state phosphors at room temperature.

Polyoxometalates incorporating  $\text{Ln}^{3+}$  lanthanide ions (Ln-POMs) form a fascinating class of nanosized metal-oxide clusters of early transition metals ( $M = \text{Mo}, \text{W}, \text{V}$ ).<sup>1</sup> They are currently the focus of intense researches owing to their promising applications in the fields of single molecule magnets (SMMs),<sup>2</sup> catalysis,<sup>3</sup> and especially photoluminescence (PL).<sup>4</sup>  $\text{Na}_9[\text{EuW}_{10}\text{O}_{36}] \cdot x\text{H}_2\text{O}$  (**EuW<sub>10</sub>**) has been extensively investigated due to its remarkable PL properties<sup>5</sup> which were recently used to elaborate chemical probes for biology,<sup>6</sup> photocatalysts<sup>7</sup> and luminescent thermometers.<sup>8</sup> In the Weakley-type  $[\text{Eu}(\text{W}_5\text{O}_{18})_2]^{9-}$  complex, the POM ligands act as sensitizers to exalt the luminescence of the  $\text{Eu}^{3+}$  ion via an antenna effect.<sup>5</sup>  $[\text{Eu}(\text{W}_5\text{O}_{18})_2]^{9-}$  has been also ionically incorporated into organic polymers,<sup>9</sup> vesicles,<sup>10</sup> surfactants,<sup>11</sup> ionic liquids,<sup>12</sup> and metal-organic frameworks<sup>8</sup> to design hybrid organic-inorganic photoactive composites. However, because of its huge negative charge and the insolubility of **EuW<sub>10</sub>** in most of non-aqueous solutions, the assembly of  $[\text{Eu}(\text{W}_5\text{O}_{18})_2]^{9-}$  with organic cations into crystallized supramolecular frameworks is more critical and was very rarely reported.<sup>13</sup> Other Eu-POMs were since designed,<sup>4c,5,14</sup> but they are less efficient emitters than **EuW<sub>10</sub>** because i) they contain corner-shared  $\text{MO}_6$  octahedra that favor the electron-to-hole recombination into the POM ligands instead of POM-to- $\text{Eu}^{3+}$  energy transfers, ii) the  $\text{Ln}^{3+}$  ions are too close to each other leading to PL concentration quenching effects or iii) water molecules link the  $\text{Eu}^{3+}$  ions and promote non-radiative deactivation from its  $^5\text{D}_0$  emitting level.<sup>15</sup> In addition, most of these compounds are hydrated alkali salts which exhibit a limited thermal stability. Thus, the discovery of Ln-POMs with lower charges, and soluble in non-aqueous solvents is still highly challenging. In 1997, Kitamura *et al.* reported tetrabutylammonium (TBA) salts of the  $[\text{Ln}(\text{Mo}_8\text{O}_{26})_2]^{5-}$  complex ( $\text{Ln} = \text{La}, \text{Y}, \text{Ce}, \text{Pr}, \text{Nd}, \text{Gd}, \text{Yb}$ ) (Figure 1a).<sup>16</sup> Other Ln-derivatives (Tb, Dy, Ho, Er, Tm)

were recently obtained which exhibit interesting SMM properties.<sup>17</sup> However, crystals of these efflorescent acetonitrile solvates degrade rapidly in air, leading to structurally uncharacterized powders. In addition, the photophysical properties of these Ln-POMs have never been investigated, and their Eu and Sm counterparts have not been designed yet.

We report herein the synthesis and structural characterization of two new thermally robust  $(\text{TBA})_5[\text{Ln}(\text{Mo}_8\text{O}_{26})_2]$  compounds (**TBA-EuMo<sub>16</sub>** and **TBA-SmMo<sub>16</sub>**), and their  $\text{Y}^{3+}$ -counterpart (**TBA-YMo<sub>16</sub>**). Their tetraphenylphosphonium (TPP) derivatives  $(\text{TPP})_5[\text{Ln}(\text{Mo}_8\text{O}_{26})_2]$  (**TPP-LnMo<sub>16</sub>**) were also prepared in acetonitrile by a salt metathesis reaction. The photophysical properties of these phosphors were thoroughly investigated and discussed.



**Figure 1.** a) Structure of the  $[\text{Eu}(\text{Mo}_8\text{O}_{26})_2]^{5-}$  complex. b) Supramolecular  $\{(\text{TBA})_5[\text{Eu}(\text{Mo}_8\text{O}_{26})_2]\}^{3+}$  ribbon. c) Crystal packing in **TBA-EuMo<sub>16</sub>**. Green arrows indicate the Eu...Eu distances (blue octahedra =  $\text{MoO}_6$ , gold sphere: oxygen, pink sphere: europium, green sphere: nitrogen, grey sphere: carbon. H-atoms are omitted).

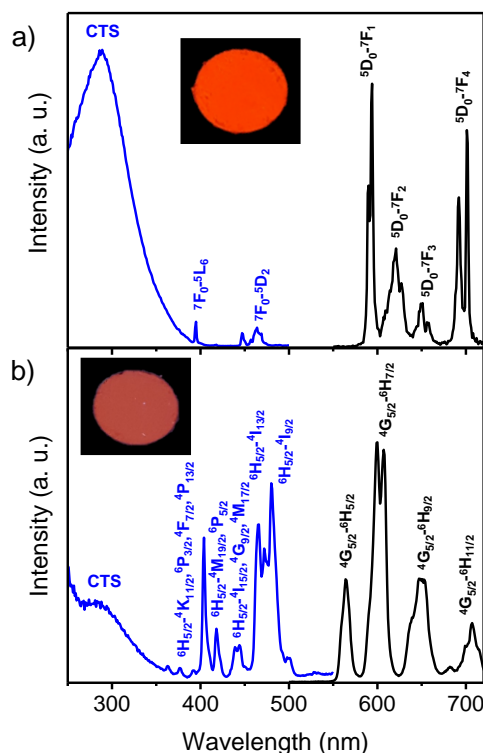
The syntheses of the six compounds are described in the Supporting Information. The TBA-LnMo<sub>16</sub> systems were obtained as colorless crystals by slightly modifying the synthesis route developed for other Ln-derivatives.<sup>17</sup> Our synthetic procedure also differs from that previously reported for **TBA-YMo<sub>16</sub>**.<sup>16</sup> FT-IR spectroscopy (Figure S1), TGA measurements (Figure S2) along with elemental analyses confirm the chemical composition

(TBA)<sub>5</sub>[Ln(Mo<sub>8</sub>O<sub>26</sub>)<sub>2</sub>]. Noticeably, crystals are stable **in air**, and no efflorescence appears even after several months. Powder X-Ray diffraction (PXRD) analysis shows that the three compounds are isostructural (Figure S3). The structures of **TBA-EuMo<sub>16</sub>** and **TBA-SmMo<sub>16</sub>** were resolved by single-crystal X-ray diffraction (Table S1), and that of **TBA-EuMo<sub>16</sub>** is described below. In [Eu(Mo<sub>8</sub>O<sub>26</sub>)<sub>2</sub>]<sup>5-</sup>, two β-[Mo<sub>8</sub>O<sub>26</sub>]<sup>4-</sup> units connect a central Eu<sup>3+</sup> ion (Figure 1a) which is located into a distorted square antiprismatic site. Supramolecular {(TBA)<sub>2</sub>[Eu(Mo<sub>8</sub>O<sub>26</sub>)<sub>2</sub>]}<sup>3-</sup> ribbons run along the *a* axis (Figure 1b) with long intermolecular Eu⋯Eu distances of 12.1 Å and 13.4 Å. Ribbons are separated by TBA units (Figure 1c) with the shortest inter-ribbon Eu⋯Eu distance of 17.2 Å. As suspected, the structures do not show any crystallized acetonitrile molecules. This is consistent with the absence in the Raman spectra of the intense ν(C≡N) stretching mode around 2260 cm<sup>-1</sup> (Figure S4).<sup>18</sup> In addition, TGA measurements (Figure S2) **confirm the stability of compounds in air up to 275°C**.

FT-Raman spectroscopy is a powerful tool to characterize POMs in solution and in the solid state.<sup>19</sup> Figure S5 displays the Raman spectra of **TBA-EuMo<sub>16</sub>** in acetonitrile solutions at three concentrations i.e., 2 · 10<sup>-2</sup> M, 10<sup>-2</sup> M, and 5 · 10<sup>-3</sup> M. In the 150-1000 cm<sup>-1</sup> range, the Raman peaks characteristic of [Eu(Mo<sub>8</sub>O<sub>26</sub>)<sub>2</sub>]<sup>5-</sup> match with those recorded for the **TBA-EuMo<sub>16</sub>** solid phase. In this concentration range, no additional signals that could indicate its partial degradation are observed. Moreover, the intensity of the main line at 977 cm<sup>-1</sup> **attributed** to the terminal Mo=O bonds increases linearly with the Eu-POM concentration (Figure S6). In addition, a statistical analysis by “separation curve method” reveals that the spectra recorded in solution are perfectly similar to the solid one. **This quantitatively confirms** the integrity of the Ln-POM complex in solution. Then, by exchanging TBA cations by TPP ones, **TPP-LnMo<sub>16</sub>** (Ln = Eu, Sm) and their Y<sup>3+</sup> counterpart were prepared as microcrystalline powders. Attempts to isolate single-crystals were unsuccessful. However, PXRD analyses show that they are isostructural (Figure S3). Finally, FT-IR (Figure S1) and FT-Raman (Figure S7) spectroscopies, elemental analyses, and TGA measurements (Figure S2) indicate a TPP:Ln-POM ratio of 5 : 1, and show that the compounds are thermally stable in air up to 355°C.

The solid-state absorption spectra of the TBA-LnMo<sub>16</sub> series (Figure S8) show two broad bands at 270 and 305 nm, which are assigned to mixed O→Mo and O→Eu ligand-to-metal charge-transfer (LMCT) transitions. Similar bands are observed for the TPP-LnMo<sub>16</sub> series, with a more intense transition at 270 nm, because the cation absorbs at this wavelength. The six compounds have an optical band gap of 370 nm. In addition, room-temperature time-resolved PL spectroscopy has been performed to characterize the intrinsic luminescence of the POM ligands. Upon photoexcitation into the O→Mo LMCT band, the Ln-free reference **TBA-YMo<sub>16</sub>** shows a broad emission band centered at 385 nm (Figure S9) with a short mean luminescence lifetime of τ = 0.382 μs (Table S2). This band can be decomposed into two contributions at 364 (27473 cm<sup>-1</sup>) and 418 nm (23923 cm<sup>-1</sup>) (Table S2). Comparable emissions are also observed for the TBA-LnMo<sub>16</sub> (Ln = Eu, Sm) compounds and their TPP counterparts (Figure S9), with similar decay times (Table S2). Based on earlier studies,<sup>10,4c</sup> these bands have been assigned

to the <sup>3</sup>T<sub>1u</sub>→<sup>1</sup>A<sub>1g</sub> transitions originating from the two O→Mo LMCT emitting triplet states of the POM ligands.



**Figure 2.** a) Excitation spectrum (blue line) monitored at 594 nm and emission spectrum (black line) monitored at 286 nm of **TBA-EuMo<sub>16</sub>**. Inset: Photograph of the powder of **TBA-EuMo<sub>16</sub>** upon irradiation at 254 nm. b) Excitation spectrum (blue line) monitored at 286 nm and emission spectrum (black line) monitored at 286 nm of **TBA-SmMo<sub>16</sub>**. Inset: Photograph of the powder of **TBA-SmMo<sub>16</sub>** upon irradiation at 254 nm.

The room-temperature steady-state photoluminescence excitation (PLE) and emission spectra of **TBA-EuMo<sub>16</sub>** and **TBA-SmMo<sub>16</sub>** are depicted in Figure 2. Upon irradiation at 254 nm with a standard UV lamp, **TBA-EuMo<sub>16</sub>** exhibits a strong orange-red emission (Figure 2a). Its PLE spectrum monitored at 594 nm shows a broad intense band at 286 nm (34965 cm<sup>-1</sup>) which corresponds to the charge-transfer states (CTS), mainly dominated by the <sup>1</sup>A<sub>1g</sub>→<sup>1</sup>T<sub>1u</sub> transition of the O→Mo LMCT photoexcitation.<sup>4c</sup> Several sharp lines are also observed which are characteristic of the intra-configurational f-f transitions of Eu<sup>3+</sup> i.e., <sup>7</sup>F<sub>0</sub>→<sup>5</sup>L<sub>6</sub> (395 nm) and <sup>7</sup>F<sub>0</sub>→<sup>5</sup>D<sub>2</sub> (464 nm). Noticeably, the f-f transitions got a very weak intensity which indicates that the most efficient excitation process corresponds to the CTS. Thus, the β-[Mo<sub>8</sub>O<sub>26</sub>]<sup>4-</sup> ligands are effective light-harvesting antennae to sensitize the Eu<sup>3+</sup> ion through an intramolecular energy transfer from the <sup>3</sup>T<sub>1u</sub> excited levels to the emitting <sup>5</sup>D<sub>0</sub> level, **lying** at a lower energy (Figure S10). Upon photoexcitation into the CTS band, the PL spectrum of **TBA-EuMo<sub>16</sub>** consists in a multi-band emission characteristic of the <sup>5</sup>D<sub>0</sub>→<sup>7</sup>F<sub>J</sub> (J = 0-4) transitions of an Eu<sup>3+</sup> ion occupying an asymmetric site with approximate D<sub>4d</sub> symmetry.<sup>20</sup> Indeed, the symmetry-forbidden <sup>5</sup>D<sub>0</sub>→<sup>7</sup>F<sub>0</sub> electric-dipole transition is absent. The strongest <sup>5</sup>D<sub>0</sub>→<sup>7</sup>F<sub>1</sub> magnetic-dipole transition is split into two Stark components at

590 and 594 nm. The  $^5D_0 \rightarrow ^7F_2$  electric-dipole transition (611–630 nm), which is highly sensitive to the chemical surrounding at the vicinity of  $\text{Eu}^{3+}$  and the  $^5D_0 \rightarrow ^7F_3$  transition (648–661 nm) got weak intensities. In contrast, the  $^5D_0 \rightarrow ^7F_4$  split-band (692 and 701 nm) is intense. Thus, the emission of **TBA-EuMo<sub>16</sub>** is characterized by CIE chromaticity coordinates of (0.6255, 0.3681) (Figure S11). The  $^5D_0 \rightarrow ^7F_1$  transition exhibits a long luminescence lifetime of  $5.24 \pm 0.03$  ms (Figure S12). Noticeably, the emission spectrum of **TBA-EuMo<sub>16</sub>** shows the same feature than that of **EuW<sub>10</sub>** and their PL performances are close.<sup>5</sup> This can be explained considering that  $[\text{Eu}(\text{Mo}_8\text{O}_{26})_2]^{5-}$  and  $[\text{Eu}(\text{W}_5\text{O}_{18})_2]^{9-}$  have some structural similarities. Indeed, they contain POM ligands only built upon edge-shared  $\text{MO}_6$  octahedra that discriminates against thermally activated hopping of d<sup>1</sup> electrons and limits the M→O non-radiative deactivation processes.<sup>22,4c</sup> They also exhibit low average M-O-Eu bond angle values ( $131.3^\circ$  and  $130.17^\circ$  for **TBA-EuMo<sub>16</sub>** and **EuW<sub>10</sub>**,<sup>21</sup> respectively). This contributes to prevent the PL quenching of  $\text{Eu}^{3+}$  due to d<sup>1</sup> electron hopping through  $f\pi$ - $\pi$ - $d\pi$  orbital mixing, which is rather favored for higher bond angle values.<sup>4b,c</sup> Moreover, the  $\text{Eu}^{3+}$  ion occupies a similar distorted square antiprismatic site without coordinated water molecules likely to quench its luminescence.<sup>22</sup> Finally, the long  $\text{Eu}\cdots\text{Eu}$  distances, higher than 12 Å, may suggest that side PL concentration quenching effects should be strongly minimized. The orange emission of **TBA-SmMo<sub>16</sub>** at room temperature is less intense than that of **TBA-EuMo<sub>16</sub>** (Figure 2b). The PLE spectrum contains the CTS band (286 nm) and sharp lines associated to the f-f transitions of  $\text{Sm}^{3+}$ , i.e.  $^6H_{5/2} \rightarrow ^4K_{11/2}$ ,  $^6P_{3/2}$ ,  $^4F_{7/2}$ ,  $^4P_{13/2}$  (404 nm),  $^6H_{5/2} \rightarrow ^4M_{19/2}$ ,  $^6P_{5/2}$  (418 nm),  $^6H_{5/2} \rightarrow ^4I_{15/2}$ ,  $^4G_{9/2}$ ,  $^4M_{17/2}$  (440–452 nm),  $^6H_{5/2} \rightarrow ^4I_{13/2}$  (466 nm),  $^6H_{5/2} \rightarrow ^4I_{9/2}$  (480 nm).<sup>23</sup> They are more intense than the CTS band, which indicates that the emission process can be stimulated upon photoexcitation into the CTS band or directly into the f-f transitions of  $\text{Sm}^{3+}$ . The emission spectrum of **TBA-SmMo<sub>16</sub>** ( $\lambda_{\text{ex}} = 286$  nm) consists in characteristic emission peaks of the  $\text{Sm}^{3+}$  ion at 564 nm ( $^4G_{5/2} \rightarrow ^6H_{5/2}$ ), 599 and 607 nm ( $^4G_{5/2} \rightarrow ^6H_{7/2}$ ), 650 nm ( $^4G_{5/2} \rightarrow ^6H_{9/2}$ ), and 707 nm ( $^4G_{5/2} \rightarrow ^6H_{11/2}$ ).<sup>23</sup> Its emission color is more orange than for **TBA-EuMo<sub>16</sub>** with CIE chromaticity coordinates of (0.5841, 0.4065) (Figure S11). The decay time of the  $^4G_{5/2} \rightarrow ^5H_{7/2}$  emission ( $\lambda_{\text{exc}} = 286$  nm) is  $0.28 \pm 0.01$  ms (Figure S12).

**TBA-EuMo<sub>16</sub>** and **TBA-SmMo<sub>16</sub>** dissolved in acetonitrile exhibit an orange luminescence at room temperature (Figure S13). The PL spectrum of  $[\text{Sm}(\text{Mo}_8\text{O}_{26})_2]^{5-}$  is similar to that of **TBA-SmMo<sub>16</sub>**, confirming the robustness of the Sm-POM. The PL spectrum of  $[\text{Eu}(\text{Mo}_8\text{O}_{26})_2]^{5-}$  in solution shows slight changes compared to that monitored in the solid state. In particular, the weakly intense  $^5D_0 \rightarrow ^7F_0$  transition (576 nm) is distinguishable. In addition, the  $^5D_0 \rightarrow ^7F_2/{}^5D_0 \rightarrow ^7F_1$  intensity ratio is increased (0.64 vs 0.38 for **TBA-EuMo<sub>16</sub>**). This indicates that the site symmetry of the  $\text{Eu}^{3+}$  ion decreases.<sup>12</sup> Nevertheless, as no change is detectable in the Raman spectrum (Figure S5), this symmetry reduction should be limited, and anyway much less pronounced than for **EuW<sub>10</sub>** which shows a strong inversion of the  $^5D_0 \rightarrow ^7F_2/{}^5D_0 \rightarrow ^7F_1$  intensity ratio in water.<sup>8</sup>

**TPP-EuMo<sub>16</sub>** has an intense orange-red solid-state emission at room temperature (Figure S14), with CIE chromaticity coordinates of (0.6281, 0.3666) very close to those of **TBA-EuMo<sub>16</sub>**. Their PLE and PL spectra (Figure S15) are compara-

ble excepted for the  $^5D_0 \rightarrow ^7F_2/{}^5D_0 \rightarrow ^7F_1$  intensity ratio which is increased to 0.5 for **TPP-EuMo<sub>16</sub>**. Upon photoexcitation at 286 nm, the decay time of the  $^5D_0 \rightarrow ^7F_1$  transition is  $5.30 \pm 0.03$  ms (Figure S12), similarly as observed for its TBA-counterpart. This evidences that the substitution of flexible TBA cations by more rigid TPP ones does not significantly impact the high PL performances of  $[\text{Eu}(\text{Mo}_8\text{O}_{26})_2]^{5-}$  in the solid state. In contrast, the orange emission of **TTP-SmMo<sub>16</sub>**, which is characterized by CIE chromaticity coordinates of (0.5757, 0.4054), is more intense than that of **TBA-SmMo<sub>16</sub>** (Figure S14), and the decay time of the  $^4G_{5/2} \rightarrow ^5H_{7/2}$  transition ( $\lambda_{\text{exc}} = 286$  nm) reaches  $0.66 \pm 0.01$  ms. This is consistent with the change in the PLE spectrum profile in which the CTS band has the same intensity than the f-f transitions of  $\text{Sm}^{3+}$  (Figure S16), suggesting a better POM-to- $\text{Sm}^{3+}$  energy transfer.

In summary, the  $[\text{Ln}(\text{Mo}_8\text{O}_{26})_2]^{5-}$  (Ln = Eu, Sm) complexes are new luminescent Ln-POMs both in solution and in the solid state. The nonsolvated TBA-LnMo<sub>16</sub> compounds have been well structurally characterized. They show a high thermal stability in air and an efficient emission at room temperature. Noticeably, the PL performances of **TBA-EuMo<sub>16</sub>** are close to those of the **EuW<sub>10</sub>** reference, owing to structural similarities between  $[\text{Eu}(\text{Mo}_8\text{O}_{26})_2]^{5-}$  and  $[\text{Eu}(\text{W}_5\text{O}_{18})_2]^{9-}$ . This makes **TBA-EuMo<sub>16</sub>** a promising alternative Ln-POM emitter which is furthermore easily soluble in polar non-aqueous solvents. Both FT-Raman and PL spectroscopies confirmed the integrity of the  $[\text{Ln}(\text{Mo}_8\text{O}_{26})_2]^{5-}$  complexes in acetonitrile. The TPP-derivatives were obtained by a cation metathesis reaction. They are also thermally robust and efficient phosphors. To go further, this work evidences the possibility to design crystallized supramolecular assemblies by combining the luminescent  $[\text{Ln}(\text{Mo}_8\text{O}_{26})_2]^{5-}$  units with photoactive organic cations, paving the way toward novel multifunctional optical hybrid systems.

## ASSOCIATED CONTENT

### Supporting Information

Experimental section, crystallographic data, IR and Raman spectra, ATG measurements, PXRD patterns, Absorption, PL and PLE spectra, CIE diagrams, luminescence decay curves and extracted lifetimes. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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### NOTES

The authors declare no competing financial interests.

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