Visual Detection of Triethylamine and a Dual Input/Output Logic Gate Based on a Eu$^{3+}$-Complex

Bao-Ning Li, Yuan-Yuan Liu, Ya-Ping Wang and Mei Pan *

Abstract: A series of Ln$^{3+}$-metal centered complexes, Ln(TTA)$_3$(DPPI) (Ln = La, 1; Ln = Eu, 2; Ln = Tb, 3; or Ln = Gd, 4) [(DPPI = N-(4-(1H-imidazo[4,5-f][1,10]phenanthroline-2-yl)phenyl)-N-phenylbenzenamine) and (TTA = 2-Thenoyltrifluoroacetone)] have been synthesized and characterized. Among which, the Eu$^{3+}$-complex shows efficient purity red luminescence in dimethylsulfoxide (DMSO) solution, with a Commission International De L’Eclairage (CIE) coordinate at x = 0.638, y = 0.323 and $\Phi_{Eu}$ = 38.9%. Interestingly, increasing the amounts of triethylamine (TEA) in the solution regulates the energy transfer between the ligand and the Eu$^{3+}$-metal center, which further leads to the luminescence color changing from red to white, and then bluish-green depending on the different excitation wavelengths. Based on this, we have designed the IMPLICATION logic gate for TEA recognition by applying the amounts of TEA and the excitation wavelengths as the dual input signal, which makes this Eu$^{3+}$-complex a promising candidate for TEA-sensing optical sensors.

Keywords: Eu$^{3+}$-complex; TEA detection; logic gate

1. Introduction

Stimulus-response materials, which are new types of intelligent materials, have received widespread attention [1–4]. Molecular systems capable of exhibiting color-tunable luminescence in suitable substrates or under specific acid or base conditions are highly appreciated, since they favor sensors [5,6], molecular logic gates and fluorescence ratio meters, etc. [7]. Among which, lanthanide-based complexes are of particular interest owing to their specific luminescent properties, which can be applied in various areas, such as pure red organic light-emitting diodes [8], cellular recognition [5], fluorescence sensors [9–11], magnetic materials and catalysis [12]. As to the potential applications of stimulus-responsive luminescent complexes [13–15], there has been an unremitting pursuit of new organo-Eu$^{3+}$ complexes to achieve desirable fluorescence responses for micro-environment detection [16–18]. Owing to the unique f-f radiative transitions, lanthanide based emitters show sharp and narrow emission bands, which usually have stability towards the surrounding environments [19]. In suitable cases, by regulating the energy of the ligand and the outer stimulations, the energy transfer between the ligands and the Ln-metal centers can be controlled, leading the emission to be “Off-On”, or leading to a change in color and intensity [20], which leads to the emission becoming a favorable and promising candidate for visual detection [21,22]. Furthermore, the detection of toxic triethylamine (TEA) [23] possesses a direct relevance to the quality of the environment and our lives and is highly demanding [24]. However, the development of stimuli-responsive systems that show a wide range of luminescent color changes for sensitive and eye-catching observations upon TEA stimulations are still worth studying [17,25,26].

Herein, we developed a series of organo-Ln$^{3+}$ (Ln = La, Eu, Tb, Gd) complexes coordinated with the donor-acceptor structured DPPI ligand (Scheme 1), which possesses an ILCT (intra-ligand charge transfer) potential due to the combination of phenanthroline,
imidazole and triphenylamine groups, which can serve as sensitizers for the Eu$^{3+}$-centered emission and also stimulation sites for the detection of TEA. As a result, the molecular Eu$^{3+}$-complex showed good photophysical responses to both TEA concentrations and excitation wavelengths, and was applied well within the dual input/output logic gate design.

![Scheme 1. Reaction procedure for the synthesis of DPPI, and complexes 1–4.](image)

**2. Results and Discussion**

**2.1. Synthesis and Photophysical Properties of the Ln-Complexes**

The ligand DPPI was synthesized according to the typical reaction [27], as shown in Scheme 1. Further into its self-assembly with a deprotonated TTA ligand and LnCl$_3$·6H$_2$O (Ln = La, Eu, Tb, or Gd), a series of Ln(TTA)$_3$(DPPI) complexes (Ln = La, 1; Ln = Eu, 2; Ln = Tb, 3; or Ln = Gd, 4) were obtained and well-characterized by EA, FT-IR, $^1$H NMR and ESI-MS, respectively (Figures S1–S5). In particular, the $^1$H NMR spectra of complex 1 and the DPPI ligand reveals that the TTA$^-$ ligand and the rare earth ion [Ln$^{3+}$] are in a stipulated molar ratio of 3:1. In addition, the characteristic proton resonances ($\delta$ = 13.68 ppm) of the typical group ‘-NH’ relative to the DPPI ligand moved 0.11 ppm to the higher field of complex 1 ($\delta$ = 13.59 ppm), due to the coordination of the La$^{3+}$ ion by the metal-atomic effects [28,29]. Moreover, the ESI-MS spectra of these typically binary tris-$\beta$-diketone-Ln$^{3+}$-complexes show standard molecular ion peaks at 1288.97, 1302.03, 1308.99, and 1307.32, respectively (Ln = La, 1; Ln = Eu, 2; Ln = Tb, 3; or Ln = Gd, 4) [30–32].

The photophysical properties of the complexes were examined in dilute DMSO solution at RT or 77 K, and are summarized in Figure 1, Figures S6 and S7 and Table S1. In particular, the $^1$H NMR spectra of complex 1 and the DPPI ligand revealed that the TTA$^-$ ligand effects [28,29]. Moreover, the ESI-MS spectra of these typically binary tris-$\beta$-diketone-Ln$^{3+}$-complexes show standard molecular ion peaks at 1288.97, 1302.03, 1308.99, and 1307.32, respectively (Ln = La, 1; Ln = Eu, 2; Ln = Tb, 3; or Ln = Gd, 4) [30–32].

The photophysical properties of the complexes were examined in dilute DMSO solution at RT or 77 K, and are summarized in Figure 1, Figures S6 and S7 and Table S1. In particular, the $^1$H NMR spectra of complex 1 and the DPPI ligand showed good photophysical responses to both TEA concentrations and excitation wavelengths, and was applied well within the dual input/output logic gate design.

For complex 2, the luminescent emission (Figure 1) shows the Eu$^{3+}$-centered characteristic peaks ($^5D_{0} \rightarrow ^7F_j$, $j = 0–4$) with the maximum at 612 nm [26], giving the pure red-light CIE coordinate of (x = 0.638, y = 0.323) and a quantum yield of ($\Phi_{\text{Eu}}^\text{L} = 38.9\%$). For complex 3, photoexcitation ($\lambda_{\text{exc}} = 406$ nm) gave a weak Tb$^{3+}$-centered characteristic emission at 545 nm, while a strong and wide residual emission ($\lambda_{\text{em}} = 506$ nm) assigned to the DPPI ligand was observed [35]. However, in contrast to complexes 2 and 3 at RT, complex 4 shows that the 0-0 transition phosphorescence ($\lambda_{\text{em}} = 519$ nm, $\tau = 11.3 \mu$s) was under 77 K (Figure S7), assigned to the ($^5\pi-\pi^*$) triplet energy level (20198 cm$^{-1}$) of the ligand. Meanwhile, the ($^\pi-\pi^*$) singlet energy level (25313 cm$^{-1}$) was obtained by the lowest wavelength of the UV-visible edge. As a result, an energy gap of $\Delta E$ ($^1\pi-\pi^* \rightarrow ^3\pi-\pi^* = 5115$ cm$^{-1}$) greater than 5000 cm$^{-1}$ was obtained [36], as shown in Figure S8, which enabled an effective ISC (intersystem crossing) according to Reinholdt’s empirical rule [37]. Furthermore, through the check of the energy level match between the ligand and the first excited state level of $^3D_0$ (17286 cm$^{-1}$) of the Eu$^{3+}$ ion, the suitable energy gap
\(\Delta E \) of 2912 cm\(^{-1}\) within a 2500–4500 cm\(^{-1}\) range from Latva’s empirical rule confirmed the effective sensitization of Eu\(^{3+}\). However, the triplet energy level of the ligand was lower than \(5D_4\) (20545 cm\(^{-1}\)) of Tb\(^{3+}\), and the excessive nonradiative transitions resulted in the quenching of the Tb\(^{3+}\)-centered effective emission [20]. The Eu\(^{3+}\)-based (\(\lambda_{em} = 612\) nm) lifetime of 2 was found to be \(\tau_{obs} = 217\) \(\mu\)s, and a high overall quantum yield (\(\Phi_{Eu} = 38.9\)%) was obtained, further manifesting the efficient energy transfer from the ligand. Noticeably, the Gd-complex 4 also gave an attractive quantum yield (\(\Phi_{em} = 8.9\)%) of blue light emission (\(x = 0.211\) and \(y = 0.219\)) from the organic ligands TTA and DPPI. Moreover, the TG analysis of the Ln-complex revealed good thermal stability to be about 300 °C (Figure S9).

![Figure 1](https://example.com/image1.png)

Figure 1. (a) Emission and excitation spectra of complexes 2–4 and DPPI and (b) CIE coordinates of complex 2 in DMSO solution (10\(^{-5}\) M) at room temperature (at around \(\lambda_{ex} = 400\) nm, and \(\lambda_{em} = 612\) nm for complex 2, \(\lambda_{em} = 545\) nm for complex 3, \(\lambda_{em} = 495\) nm for complex 4 and DPPI).

2.2. Adjustable Color Emission of Complex 2 with TEA Stimulate

As shown in Figure 2a,b, the Eu\(^{3+}\)-centered red-luminescence intensity of complex 2 in a DMSO solution gradually decreased when TEA is added dropwisely. Noticeably, a new broad blue-emission peak (\(\lambda_{em} = 500\) nm) appeared that was dependent on the different excitations and ranged from 315–420 nm. Near white light emission (\(x = 0.381, y = 0.382\)) was successfully constructed through the two emission color components (red and bluish green) at the excitation of 385 nm, and the white light emission index parameters are summarized in Table S2. In addition, the fluorescence photographs record the process of the emitting color change upon the addition of different amounts of TEA under ultraviolet light (365 nm), which is shown in Figure S10. As the amounts of TEA increased, the emission intensity of complex 2 at 613 nm gradually weakened and disappeared (Figure 2c–f); meanwhile, the blueish green light emission intensity at around 500 nm gradually increased. When the content of TEA reached 10 \(\mu\)L, it is recorded that the intensity of the residual ligand fluorescence at around 495 nm became the main emission in the whole spectra, and the Eu\(^{3+}\)-centered emission was very weak and even non-observable.

To probe the origins of the above emission responses, we speculate that as the TEA ratio in the solution increased, the electronic distribution and intra-ligand charge transfer state between the donor-acceptor groups on the DPPI ligand was changing gradually, accompanying the changes of energy transfer from the DPPI to the Eu\(^{3+}\) metal centers [38]. Basically, due to the stimulation of TEA, the energy level of DPPI was elevated (Figures S10–S12, as confirmed by the UV-visible absorption titration tests discussed below), and this caused an incomplete energy transfer to the Eu\(^{3+}\) center. Furthermore, the energy transfer processes are sensitive to the excitations. As discussed before, the DPPI ligand absorbs the UV light by either \(\pi-\pi^*\) and/or ILCT singlet states. After transitioning to the triplet states via intersystem crossing (ISC), the energy can be further transferred to the metal centers and can generate the red Eu\(^{3+}\)-emissions. However, the bluish green emission of the DPPI
ligand can only be excited by the ILCT spectral region (see Figure 1a) [39,40]. As shown in the emission spectra of complex 2 at different excitations without TEA (Figure S13), basically only the Eu\(^{3+}\)-centered emissions were observed, manifesting that the emission of the DPPI ligand itself in complex 2 was hampered due to the total energy transfer to the metal center. After the addition of TEA, both the ligand and the Eu\(^{3+}\)-centered emissions appeared due to the changes in the ligand electronic states and energy transfer processes to the Eu\(^{3+}\) center. However, according to Figure S14, it is clear that the Eu\(^{3+}\)-centered emission (recorded at 613 nm) can be excited by both the π-π* and ILCT states (showing two excitation peaks), while the ligand-centered emission can only be excited by the ILCT states (basically only one excitation peak is observed). As a result, the alternative intensity between the Eu\(^{3+}\)-centered and residual fluorescence of the DPPI ligand can be tuned by applying different excitation wavelengths in complex 2 after the addition of TEA, accompanying with various color emissions as a whole (Figure 3).

![Figure 2](image-url)

*Figure 2.* Emission spectra of complex 2 in DMSO solution (1 \(\times\) 10\(^{-5}\) M) stimulated by 2 µL (a), 5 µL (c) and 10 µL TEA (0.1 mM) (e) dependent on the excitations (\(\lambda_{\text{ex}}\) = 315–425 nm, each interval of 10 nm) at RT. (b,d,f) corresponding CIE graph dependent on the excitations.
Basically, due to the stimulation of TEA, the energy level of DPPI ligand can be tuned by applying different excitation wavelengths in complex 1. Such phenomena confirm that the basic-attributed TEA can cause electronic redistribution in the ligand DPPI, increase its energy levels and regulate the energy transfer from the ligand to the 

As shown in Figures S11–S14, in order to further verify the mechanism of the fluorescence changes, the UV-visible absorption titration of complex 2 upon TEA addition was taken. It can be seen that the absorbance intensity at 230, 270, 285, and 340 nm was obviously enhanced. At the same time, comparing the spectra before and after titration, there was a significant blue shift for the long wavelength absorption peak of nearly 20 nm after the addition of TEA, which shifted from 369 to 349 nm. For comparison, a red-shift tendency was observed upon the addition of acetic acid into the DMSO solution of complex 2. Such phenomena further confirms that the basic-attributed TEA can cause electronic redistribution in the ligand DPPI, increase its energy levels and regulate the energy transfer from the ligand to the 

To further strengthen the reason for the fluorescence alternating processes, dynamic 

1H NMR spectra were performed for complex 1 (Figure S15). The results recorded that with the addition of TEA, the proton resonances of the typical group ‘-NH’ (δ = 13.59 ppm) were gradually widened or weakened. Meanwhile, the whole chemical shift peaks shifted slightly, due to the intermolecular forces between the Ln-complexes and TEA [42,43]. These observations can be attributed to the unique structure of DPPI with imidazole and triphenylamine groups, which are sensitive to the micro-environments changed by the addition of TEA [44–46]. On one hand, TEA provides active hydrogen protons that result in acid-base interactions; on the other hand, it causes the electronic distribution and energy state changes in the ILCT-structured DPPI ligand due to molecular interactions [47], and leads to the color emission variations and the TEA detecting capabilities of the Eu³⁺-metal centers [41].

2.3. Potential Application in IMPLICATION Logic Gate

As discussed above, by increasing the amounts of TEA into the solution of complex 2 at the same excitation wavelength (λex = 365 nm), the color tunable emission can be obtained, which can range from a red emission with a CIE chromatic coordinate of (x = 0.638, y = 0.323), to a white light emission (x = 0.366, y = 0.372), then to a green emission (x = 0.280, y = 0.420) (Figures 3 and 4). The amounts of TEA may affect the rate and degree of the molecular interaction between the TEA molecule and the DPPI ligand, further leading the emission color to change, combined with the results in Figures 2 and 3 and Figures S11 and S12. This is a key factor in determining the fold of the ratio metric enhancement.
between the two emission peaks at 613 nm and 500 nm of complex 2. The strongest fluorescence intensity for Eu\(^{3+}\) at 613 nm of 2 was achieved without the addition of TEA under \(\lambda_{\text{ex}} = 365\) nm. With the addition of TEA, the emission intensity at 613 nm decreased, while the fluorescence intensity at 500 nm assigned to the ligand was gradually enhanced. The corresponding histograms of ratiometric signals vs. various excitations and TEA values are displayed in Figure 4c,d. To this end, the intensity ratio (R = \(I_{613}/I_{500}\)) between the Eu\(^{3+}\) emission at 613 nm and the metal disturbed ligand emission at 500 nm is set as the standard, with a threshold of 2.5. If the luminescence ratio value is higher than the default threshold, the ratiometric signals show turn-on. When the ratio is smaller than the threshold, it can be described as the turn-off state.

3. Experimental Section

3.1. Synthesis of 1,10-Phenanthroline-5,6-Dione (PHD)

The general procedure for the synthesis of PHD followed the methods in the literature [50]. An ice-cooled mixture solution of 20 mL H\(_2\)SO\(_4\) and 10 mL HNO\(_3\) was added by (11.0 mmol, 2.0 g) 1,10-phenanthroline and (16.8 mmol, 2.0 g) KBr, stirred at room temperature for 20 min. After this procedure, the mixture was stirred at 130 °C for another 3 h. Then the hot yellow solution was poured into 500 mL of ice water and neutralized by a saturated Na\(_2\)CO\(_3\) aqueous solution until pH = 7–8. The solution was then extracted with CHCl\(_3\), dried with anhydrous Mg\(_2\)SO\(_4\) and the solvent was removed. There was a yield of

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**Figure 4.** Visible emission spectra (a), and corresponding CIE graph (b) of complex 2 in DMSO solution stimulated by different amounts of TEA at 1 × 10\(^{-5}\) M at RT with the excitation 365 nm. (c) corresponding histograms of the fluorescence intensity ratio (R) of 2 at 613 and 500 nm (R = \(I_{613}/I_{500}\)) with 2 \(\mu\)L TEA at different excitation. (d) corresponding histograms of the fluorescence intensity ratio (R') of 2 at 613 and 500 nm (R' = \(I_{500}/I_{613}\)) with different amounts of TEA at \(\lambda_{\text{ex}} = 365\) nm. (e) The IMPLICATION logic gate dependent on the excitation and amounts of TEA as input signals, and the color of the emission and the ratio intensity as dual output signals.
It is noted that such color tunable changes of fluorescence intensity are dependent both on the excitation wavelength and amounts of TEA, which are conducive to the construction of the logic gate as illustrated in Figure 4e. The following interpretation of two input modes and two output modes can be applied: (a) viewing the overall emission color, and (b) detecting the readout ratios for comparing the relative intensities at 613 and 500 nm. As we can see, with different input signals, the output 1 signals (overall emission color) can be red, yellowish pink, white, or blueish green, while the output 2 signals will change from ON ($I_{613}/I_{500} > 2.5$) to OFF ($I_{613}/I_{500} < 2.5$). Therefore, we can design a dual-input/dual-output color IMPLICATION logic gate.

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$^1$H NMR (400 MHz, DMSO-$d_6$): $\delta$ 9.05 (d, 2H, -pyridyl), 8.56 (d, 2H, -pyridyl), 7.84 (t, 2H, -pyridyl). FT-IR (KBr, cm$^{-1}$): 1684 (s), 1560 (m), 1459 (w), 1414 (s), 1313 (w), 1293 (s), 1205 (w), 1115 (w), 1010 (w), 925 (w), 807 (m), 736 (vs), 668 (w), 613 (w), 540 (w).

The elemental analysis calculated (%) for C$_{12}$H$_6$N$_2$O$_2$ was C, 68.57; H, 2.88; N, 13.33. C, 68.49; H, 2.77; N, 13.21. MS (ESI$^+$): m/z: 211.04(100%), [M − H]$^+$ was found.

3.2. Synthesis of DPPI

The DPPI was synthesized by refluxing a mixture of 1,10-phenanthroline-5,6-dione (0.21 g, 1 mmol), 4-(diphenylamino)-benzaldehyde (0.27 g, 1 mmol), NaH$_2$PO$_4$ (0.012 g, 1 mmol) and ammonium acetate (1.54 g, 20 mmol) in glacial acid (40 mL) for 6 h under a N$_2$ atmosphere, according to the literature [51]. After being stirred overnight, the mixture was cooled to room temperature, and then the aiming compound was obtained by the addition of a K$_2$CO$_3$ solution to the mixture filtration and from washing with enough water. The crude product was further extracted by CHCl$_3$ and vacuum dried. There was a yield of 46%.

$^1$H NMR (400 MHz, DMSO-$d_6$): $\delta$ 13.68 (s, 1H, -NH), 9.04 (d, 2H, -pyridyl), 8.91 (d, 2H, -pyridyl), 8.17 (t, 2H, -pyridyl), 7.83 (m, 2H, -phenyl), 7.45–7.36 (m, 4H, -phenyl), 7.25–7.07 (m, 8H, -phenyl). FT-IR (KBr, cm$^{-1}$): 3060 (w), 1590(m), 1521 (w), 1480 (s), 1397 (w), 1317 (w), 1280 (m), 1190 (w), 1126 (w), 1069 (w), 1029 (w), 952 (w), 840 (w), 804 (w), 740 (s), 694 (vs), 636 (w), 619 (w), 530 (w).

The elemental analysis calculated (%) for C$_{25}$H$_{23}$N$_3$ was C, 85.87; H, 5.02; N, 9.10. C, 86.06; H, 5.18; N, 9.37. MS (ESI$^+$): m/z: 486.17(100%), [M − H]$^+$ was found.

3.3. Synthesis and Characterization of Ln(TTA)$_3$(DPPI) (Ln = La, 1; Ln = Eu, 2; Ln = Tb, 3; Ln = Gd, 4)

A solution of 2-Thenoyltrifuoroacetone (HTTA) (0.20 g, 0.9 mmol), and sodium hydroxide (0.36 g, 0.9 mmol) was added in 5 mL of mixed solvent (MeOH/chloroform = 3: 5) and stirred for 30 min. Then DPPI (0.13 g, 0.3 mmol) was added, and then heated at 50 °C for 30 min with stirring. Following this, the solution of LnCl$_3$−6H$_2$O (0.3 mmol; Ln = La, 0.10 g; Ln = Eu, 0.11 g; Ln = Tb, 0.12 g or Ln = Gd, 0.11 g) was dissolved in 2 mL of MeOH and added dropwise and stirred at 50 °C for 12 h. The light yellow product was isolated by precipitation in ice-cold n-hexane and dried for 24 h in a vacuum oven at 30 °C to obtain a milk-white powder. The detailed process is shown in Scheme 1. The series of complexes Ln(TTA)$_3$(DPPI) (Ln = La, 1; Ln = Eu, 2; Ln = Tb, 3; Ln = Gd, 4) were obtained and well-characterized by EA, FT-IR, $^1$H NMR and ESI-MS, respectively.
For 1 (C$_{35}$H$_{33}$F$_{6}$O$_{6}$N$_{5}$S$_{2}$La), $^1$H NMR (400 MHz, DMSO-$d_6$): δ 13.59 (s, 1H, -NH), 9.05–9.81 (t, 4H, -phenyl, and -pyridyl), 8.17 (d, 2H, -phenyl), 7.83 (m, 8H, -phenyl, -thienyl and -pyridyl), 7.38 (m, 4H, -phenyl, -thienyl and -pyridyl), 7.15–7.12 (m, 11H, -phenyl, -thienyl and -pyridyl), 6.17 (s, 3H, -CH=C-). FT-IR (KBr, cm$^{-1}$): 1599 (s), 1536 (s), 1481 (m), 1454 (m), 1411 (s), 1355 (w), 1298 (vs), 1246 (w), 1229 (w), 1186 (s), 1137 (s), 1078 (w), 1061 (w), 933 (w), 859 (w), 842 (w), 786 (m), 767 (w), 718 (w), 696 (m), 680 (w), 640 (m), 604 (w), 579 (m), 520 (w). The elemental analysis (%) calculated for C$_{55}$H$_{33}$F$_{6}$O$_{6}$N$_{5}$S$_{2}$La was C, 52.18; H, 2.63; N, 5.53. C, 52.21; H, 2.49; N, 5.60. MS (ESI$^+$): m/z: 1288.97(100%), [M + Na]$^+$ was found. The yield was 60%.

For 2 (C$_{55}$H$_{33}$F$_{6}$O$_{6}$N$_{5}$S$_{2}$Eu), FT-IR (KBr, cm$^{-1}$): 1594 (s), 1537 (s), 1505 (m), 1454 (m), 1410 (s), 1354 (w), 1306 (vs), 1247 (w), 1229 (w), 1181 (s), 1135 (s), 1076 (w), 1062 (w), 933 (w), 860 (w), 842 (w), 788 (m), 768 (w), 719 (w), 696 (m), 683 (w), 640 (m), 605 (w), 579 (m), 521 (w). The elemental analysis (%) calculated for C$_{55}$H$_{33}$F$_{6}$O$_{6}$N$_{5}$S$_{2}$Eu was C, 51.65; H, 2.60; N, 5.48. C, 51.62; H, 2.59; N, 5.46. MS (ESI$^+$): m/z: 1302.05(100%), [M + Na]$^+$ was found. The yield was 63%.

For 3 (C$_{55}$H$_{33}$F$_{6}$O$_{6}$N$_{5}$S$_{2}$Tb), FT-IR (KBr, cm$^{-1}$): 1594 (s), 1537 (s), 1505 (m), 1454 (m), 1410 (s), 1354 (w), 1306 (vs), 1247 (w), 1229 (w), 1181 (s), 1135 (s), 1076 (w), 1062 (w), 933 (w), 860 (w), 842 (w), 788 (m), 768 (w), 719 (w), 696 (m), 683 (w), 640 (m), 605 (w), 579 (m), 521 (w). The elemental analysis (%) calculated for C$_{55}$H$_{33}$F$_{6}$O$_{6}$N$_{5}$S$_{2}$Tb was C, 51.37; H, 2.59; N, 5.45. C, 51.29; H, 2.52; N, 5.39. MS (ESI$^+$): m/z: 1308.99(100%), [M + Na]$^+$ was found. The yield was 65%.

For 4 (C$_{55}$H$_{33}$F$_{6}$O$_{6}$N$_{5}$S$_{2}$Gd), FT-IR (KBr, cm$^{-1}$): 1598 (s), 1538 (s), 1480 (m), 1454 (m), 1411 (s), 1355 (w), 1304 (vs), 1247 (w), 1230 (w), 1188 (s), 1137 (s), 1078 (w), 1062 (w), 934 (w), 860 (w), 842 (w), 787 (m), 768 (w), 718 (w), 695 (m), 681 (w), 641 (m), 605 (w), 581 (m), 521 (w). The elemental analysis (%) calculated for C$_{55}$H$_{33}$F$_{6}$O$_{6}$N$_{5}$S$_{2}$Gd was C, 51.44; H, 2.59; N, 5.45. C, 51.42; H, 2.53; N, 5.39. MS (ESI$^+$): m/z: 1307.32 (100%), [M + Na]$^+$ was found. The yield was 61%.

4. Conclusions

In summary, a purity-red emitter (x = 0.638, y = 0.323; $\Phi_{Eu}^L$ = 38.9%) of Eu$^{3+}$-complex was synthesized, which showed color tunable emissions that ranged from red, white, to bluish green under the different excitation wavelengths and various amounts of TEA stimulations. Relying on the changeable color emission, we successfully designed the dual-input/dual-output logic gate for the visual detection of toxic TEA in a solution, which expands the research applications of lanthanide Eu$^{3+}$ luminescent materials.

Supplementary Materials: The following are available online. Figure S1 Partial $^1$H NMR spectra of PHD, DPPI and complex 1. Figure S2 FT-IR spectrum of DPPI. Figure S3 FT-IR spectra of complexes 1–4. Figure S4 ESI mass spectrum of DPPI. Figure S5 ESI mass spectrum of complex 2. Figure S6 UV-visible absorption spectra of complexes 2–4, the ligand HTTA and DPPI. Figure S7 Emission and excitation spectra of complex 1 with the addition of TEA. Figure S8 Energy transfer process from DPPI ligand to Eu$^{3+}$ or Tb$^{3+}$ in complexes 2–3. Figure S9 TG curves of 2 and DPPI. Figure S10 Fluorescent photographs of complex 2 under Sunlight and UV light. Figure S11 UV-visible absorption spectra changes of complex 2. Figure S12 UV-visible absorption spectra of complex 2 by different excitation wavelengths ($\lambda_{ex}$ = 290 to 410 nm) in DMSO solution without TEA. Figure S13 Emission spectra of complex 2 monitored at 495 nm and 613 nm with the simulation of TEA in solution at room temperature. Figure S14 Excitation spectra of complex 2 monitored at 495 nm and 613 nm with the simulation of TEA in solution at room temperature. Figure S15 $^1$H NMR (400 MHz, DMSO-$d_6$) of complex 1 with the addition of TEA (0.1 mM). Table S1 Photophysical properties of complex monomers 2–4, HTTA and DPPI. Table S2 White light emission index parameters.

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**References**

1. Lee, J.; Farha, O.K.; Roberts, J.; Scheidt, K.A.; Nguyen, S.T.; Hupp, J.T. Metal-organic framework materials as catalysts. *Chem. Soc. Rev.* 2009, 38, 1450–1459. [CrossRef]

2. Ramaswamy, P.; Wong, N.E.; Shimizu, G.K. MOFs as proton conductors-challenges and opportunities. *Chem. Soc. Rev. 2014, 43, 5913–5932. [CrossRef]

3. Guo, M.Y.; Li, P.; Yang, S.L.; Bu, R.; Piao, X.Q.; Gao, E.Q. Distinct and Selective Amine- and Anion-Responsive Behaviors of bis(diimine)dicarboxylic acid europium(III) complexes. *J. Mater. Chem. C 2015, 3, 5775–5782. [CrossRef] [PubMed]

4. Li, P.; Li, Z.; Yao, D.; Li, H. Colorimetric sensor arrays for amines based on responsive lanthanide complex entrapment. *J. Mater. Chem. C 2017, 5, 6805–6811. [CrossRef]

5. Chu, B.; Song, B.; Ji, X.; Su, Y.; Wang, H.; He, Y. Fluorescent Silicon Nanorods-Based Ratiometric Sensors for Long-Term and Real-Time Measurements of Intracellular pH in Live Cells. *Anal. Chem. 2017, 89, 12152–12159. [CrossRef] [PubMed]

6. Fu, P.; Mo, J.; Sun, S.; Yin, S.; Wang, H.; Pan, M. Acid-base Vapor Sensing Enabled by ESITP-attributed Cd(II) Coordination Polymer with Switchable Luminescence. *Chem. Res. Chin. Univ. 2020, 36, 755–759. [CrossRef]

7. Du, B.B.; Zhu, Y.X.; Pan, M.; Yue, M.Q.; Hou, Y.J.; Wu, K.; Zhang, L.Y.; Chen, L.; Yin, S.Y.; Fan, Y.N.; et al. Direct white-light and a dual-channel barcode module from Pr(III)-MOF crystals. *Chem. Commun. 2015, 51, 12533–12536. [CrossRef] [PubMed]

8. Biju, S.; Xu, L.-J.; Sun, C.-Z.; Chen, Z.-N. White OLEDs based on a novel EuIII-tetrakis-β-diketonate doped into 4,4′-N,N′-dicarbazolebiphenyl as emitting material. *J. Mater. Chem. C 2015, 3, 5775–5782. [CrossRef]

9. Yin, S.-Y.; Fu, P.-Y.; Pan, M.; Guo, J.; Fan, Y.-N.; Su, C.-Y. Reverse photoluminescence responses of Ln(III) complexes to methanol vapor clarify the differentiated energy transfer pathway and potential for methanol detection and encryption. *J. Mater. Chem. C 2020, 8, 16907–16914. [CrossRef]

10. Okutani, K.; Nozaki, K.; Iwamura, M. Specific chiral sensing of amino acids using induced circularly polarized luminescence of bis(diimine)dicarboxylic acid europium(III) complexes. *Inorg. Chem. 2014, 53, 5527–5537. [CrossRef]

11. Wang, G.-D.; Li, Y.-Z.; Shi, W.-J.; Zhang, B.; Hou, L.; Wang, Y.-Y. A robust cluster-based Eu-MOF as multi-functional fluorescence sensor for detecting antibiotics and pesticides in water. *Sens. Actuators B 2021, 331, 129377. [CrossRef]

12. Ma, Y.-J.; Hu, J.-X.; Han, S.-D.; Pan, J.; Li, J.-H.; Wang, G.-M. Manipulating on/off single-molecule magnet behavior in a Dy(III)-based photochromic complex. *J. Am. Chem. Soc. 2020, 142, 2682–2689. [CrossRef]

13. Jeon, H.-G.; Kim, H.; Byeon, S.-H. Flexibly transparent luminescent organic-inorganic-polymer composite films: Intense full-color emissions at a single excitation wavelength. *Chem. Eng. J. 2021, 405, 126675. [CrossRef]

14. Li, P.; Zhang, Y.; Wang, Y.; Wang, Y.; Li, H. Luminescent europium(III)-beta-diketonate complexes hosted in nanozelite L as turn-on sensors for detecting basic molecules. *Chem. Commun. 2015, 50, 13680–13682. [CrossRef] [PubMed]

15. Lucas, F.; Quinton, C.; Fall, S.; Heiser, T.; Tondelier, D.; Geffroy, B.; Leclerc, N.; Rault-Berthelot, J.; Poriel, C. Universal host materials for red, green and blue high-efficiency single-layer phosphorescent organic light-emitting diodes. *J. Mater. Chem. C 2020, 8, 16354–16367. [CrossRef]

16. Wang, Y.; Zhao, G.; Chi, H.; Yang, S.; Niu, Q.; Wu, D.; Cao, W.; Li, T.; Ma, H.; Wei, Q. Self-Luminescent Lanthanide Metal–Organic Frameworks as Signal Probes in Electrochemiluminescence Immunoassay. *J. Am. Chem. Soc. 2020, 143, 504–512. [CrossRef]

17. Zhao, Y.S.; Wu, J.; Huang, J. Vertical Organic Nanowire Arrays: Controlled Synthesis and Chemical Sensors. *J. Am. Chem. Soc. 2009, 131, 3158–3159. [CrossRef]

18. Li, X.; Zhang, D.; Li, J. Emission “Off-On” effect from europium complexes triggered by AcO anion: Synthesis, characterization and sensing performance. *Spectrochim. Acta Part A 2014, 127, 1–9. [CrossRef] [PubMed]

19. Zhang, D.; Shi, M.; Liu, Z.; Li, F.; Yi, T.; Huang, C. Luminescence Modulation of a Terbium Complex with Anions and Its Application as a Reagent. *Eur. J. Inorg. Chem. 2006, 2006, 2277–2284. [CrossRef]

20. Xu, J.; Corneille, T.M.; Moore, E.G.; Law, G.-L.; Butlin, N.G.; Raymond, K.N. Octadentate Cages of Tb(III) 2-Hydroxyisophthalamides: A New Standard for Luminescent Lanthanide Labels. *J. Am. Chem. Soc. 2011, 133, 19900–19910. [CrossRef]

21. Sato, T.; Higuchi, M. A vapoluminescent Eu-based metallo-supramolecular polymer. *Chem. Commun. 2012, 48, 4947–4949. [CrossRef]
22. Chen, P.; Li, Q.; Grindley, S.; Holten-Andersen, N. White-Light-Emitting Lanthanide Metallogels with Tunable Luminescence and Reversible Stimuli-Responsive Properties. *J. Am. Chem. Soc.* 2015, 137, 11590–11593. [CrossRef] [PubMed]

23. Liang, M.H.; Xue, L.L.; Jiang, J.G. Two-Stage Cultivation of Dunaliella tertiolecta with Glycerol and Triethylamine for Lipid Accumulation: A Viable Way To Alleviate the Inhibitory Effect of Triethylamine on Biomass. *Appl. Environ. Microbiol.* 2019, 85, 02614–02618. [CrossRef] [PubMed]

24. Meng, X.; Yao, M.; Mu, S.; Wang, Y. Oxygen Vacancies Enhance Triethylamine Sensing Properties of SnO2 Nanoparticles. *ChemistrySelect* 2019, 4, 11268–11274. [CrossRef]

25. Lu, Y.; Yan, B. Lanthanide-organic-inorganic hybrids based on functionalized metal-organic frameworks (MOFs) for a near-UV white LED. *Chem. Commun.* 2014, 50, 15443–15446. [CrossRef] [PubMed]

26. Li, B.; Dong, J.-P.; Zhou, Z.; Wang, R.; Wang, L.-Y.; Zang, S.-Q. Robust lanthanide metal–organic frameworks with “all-in-one” multifunction: Efficient gas adsorption and separation, tunable light emission and luminescence sensing. *J. Mater. Chem. C* 2021, 9, 3429–3439. [CrossRef]

27. Zheng, H.-Q.; Guo, Y.-P.; Yin, M.-C.; Fan, Y.-T. Synthesis, characterization of a new photosensitive compound [Ru(bpy)2(TPAD)(PF6)2] and its application for photocatalytic hydrogen production. *Chem. Phys. Lett.* 2016, 653, 17–23. [CrossRef]

28. Zheng, H.; Deng, Y.-K.; Ye, M.-Y.; Xu, Q.-F.; Kong, X.-J.; Long, L.-S.; Zheng, L.-S. Lanthanide-Titanium Oxo Clusters as the Luminescence Sensor for Nitrobenzene Detection. *Inorg. Chem.* 2020, 59, 12404–12409. [CrossRef] [PubMed]

29. Liu, L.; Fang, M.; Chen, H.; Fu, G.; Li, B.; Lü, X.; Wang, L. Efficient and high colour-purity green-light polymer light-emitting diodes (PLEDs) based on a PVK-supported Tb3+-containing metallopolymers. *J. Mater. Chem. C* 2017, 5, 9021–9027. [CrossRef]

30. Ren, Y.; Feng, J. Poly(MMA-co-FMA) as a platform for tuning emission by clicking with luminescent lanthanide complexes. *J. Mater. Chem. C* 2018, 6, 10202–10206. [CrossRef]

31. Wang, Z.; Meng, Q.; Wang, C.; Fan, D.; Wang, Y. Full color-emitting (Y,Tb,Eu)NbO4 nanophosphors: Calcination-assisted hydrothermal synthesis, energy interaction, and application in deep UV chip-based WLEDS. *J. Mater. Chem. C* 2020, 8, 14548–14558. [CrossRef]

32. Liao, W.-M.; Li, C.-J.; Wu, X.; Zhang, J.-H.; Wang, Z.; Wang, H.-P.; Fan, Y.-N.; Pan, M.; Su, C.-Y. Homometallic Ln(III)-complexes from an ILCT ligand with sensitized vis-NIR emission, excitation-dependent PL color tuning and white-light emission. *J. Mater. Chem. C* 2018, 6, 3254–3259. [CrossRef]

33. Sun, S.-S.; Wang, Z.; Wu, X.W.; Zhang, J.-H.; Li, C.-J.; Yin, S.-Y.; Chen, L.; Pan, M.; Su, C.-Y. ESIPt-Modulated Emission of Lanthanide Complexes: Different Energy-Transfer Pathways and Multiple Responses. *Chem. Eur. J.* 2018, 24, 10091–10098. [CrossRef] [PubMed]

34. Cui, Y.; Yue, Y.; Qian, G.; Chen, B. Luminescent functional metal-organic frameworks. *Chem. Rev.* 2012, 112, 1126–1162. [CrossRef] [PubMed]

35. Chen, X.-J.; Hu, W.-J.; Qin, Y.; Zhao, G.-L. Synthesis, Crystal Structure and Fluorescence Spectrum of a Europium Compound [Eu(NO3)3(H2O)2]·(C10H16N2)·(NO3)2. *Asian J. Chem.* 2013, 25, 8418–8422. [CrossRef]

36. Vishwakarma, A.; Sengupta, S.K.; Pandey, O.P. Synthesis, Characterization and Photo-Physical Properties of Europium(III) and Terbium(III) Complexes with Thiosemicarbazones. *Asian J. Chem.* 2020, 32, 952–958. [CrossRef]

37. Xu, H.; Zhu, R.; Zhao, P.; Huang, W. Monochromatic Red-Emitting Nonconjugated Copolymers Containing Double-Carrier-Trapping Phosphine Oxide Eu3+ Segments: Toward Bright and Efficient Electroluminescence. *J. Phys. Chem. C* 2011, 115, 15627–15638. [CrossRef]

38. Dou, C.; Han, L.; Zhao, S.; Zhang, H.; Wang, Y. Multi-Stimuli-Responsive Fluorescence Switching of a Donor-Acceptor π-Conjugated Compound. *J. Phys. Chem. B* 2012, 116, 666–670. [CrossRef]

39. Yang, Q.Y.; Wu, K.; Jiang, J.J.; Hsu, C.W.; Pan, M.; Lehn, J.M.; Su, C.Y. Pure white-light and yellow-to-blue emission tuning in single crystals of Dy(III) metal framework. *Chem. Commun.* 2014, 50, 7702–7704. [CrossRef]

40. Yang, Q.Y.; Wu, K.; Jiang, J.J.; Hsu, C.W.; Pan, M.; Lehn, J.M.; Su, C.Y. Pure white-light and yellow-to-blue emission tuning in single crystals of Dy(III) metal framework. *Chem. Commun.* 2014, 50, 7702–7704. [CrossRef]

41. Francke, R.; Little, R.D. Optimizing Electron Transfer Mediators Based on Arylimidazoles by Ring Fusion: Synthesis, Electrochemistry, and Computational Analysis of 2-Aryl-1-methylphenanthro[9,10-d]imidazoles. *J. Am. Chem. Soc.* 2013, 136, 427–435. [CrossRef]

42. Yang, P.; Zhao, J.; Wu, W.; Yu, X.; Liu, Y. Accessing the long-lived triplet excited states in bodipy-conjugated 2-(2-hydroxyphenyl) benzothiazole/benzoxazoles and applications as organic triplet photosensitizers for photooxidations. *J. Org. Chem.* 2012, 77, 6166–6178. [CrossRef]

43. Chuang, W.T.; Hsieh, C.C.; Lai, C.H.; Lai, C.H.; Shih, C.W.; Chen, K.Y.; Hung, W.Y.; Hsu, Y.H.; Chou, P.T. Excited-state intramolecular proton transfer molecules bearing o-hydroxy analogues of green fluorescent protein chromophore. *J. Org. Chem.* 2011, 76, 8189–8202. [CrossRef]

44. Goudappagouda; Asokan, K.; Nayak, R.; Krishnan, R.; Babu, S.S. Tuning phosphorescence features of triphenylamines by varying functional groups and intermolecular interactions. *Dyes Pigments* 2020, 173, 107931. [CrossRef]

45. Hosseinzadeh, B.; Beni, A.S.; Azari, M.; Zarandi, M.; Karami, M. Novel D–π–A type triphenylamine based chromogens for DSSC: Design, synthesis and performance studies. *New J. Chem.* 2016, 40, 8371–8381. [CrossRef]
46. Perrier, A.; Jacquemin, D. Theoretical investigation of the photochromic properties of [2.2]paracyclophane-bridged imidazole dimers and bisimidazole) dimers. *Tetrahedron* 2017, 73, 4936–4949. [CrossRef]

47. Boxi, S.; Jana, D.; Parui, P.P.; Ghorai, B.K. Dibenzo[a,c]phenazine-Based Donor-Acceptor (D-A) Tetra Branched Molecules: Fine Tuning of Optical Properties. *ChemistrySelect* 2018, 3, 6953–6959. [CrossRef]

48. Wang, K.; Zheng, C.J.; Liu, W.; Liang, K.; Shi, Y.Z.; Tao, S.L.; Lee, C.S.; Ou, X.M.; Zhang, X.H. Avoiding Energy Loss on TADF Emitters: Controlling the Dual Conformations of D-A Structure Molecules Based on the Pseudoplanar Segments. *Adv. Mater.* 2017, 29, 170146. [CrossRef] [PubMed]

49. Li, M.H.; Lv, S.L.; You, M.H.; Lin, M.J. Three-component D-A hybrid heterostructures with enhanced photochromic, photomodulated luminescence and selective anion-sensing properties. *Dalton Trans.* 2020, 49, 13083–13089. [CrossRef]

50. Zhao, B.; Fang, Y.; Xu, Y.; Deng, Q.; Liu, T.; Kan, W.; Wang, L. Dual-responsive pH sensor based on a phenanthro[9,10-d]imidazole fluorophore modified by amino diacetate. *Tetrahedron Lett.* 2016, 57, 1825–1830. [CrossRef]

51. Jang, H.; Shin, C.-H.; Jung, B.-J.; Kim, D.-h.; Shim, H.-K.; Do, Y. Synthesis and Characterization of Dinuclear Europium Complexes Showing Pure Red Electroluminescence. *Eur. J. Inorg. Chem.* 2006, 2006, 718–725. [CrossRef]