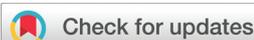


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Air stable and efficient rare earth Eu(II) hydro-tris(pyrazolyl)borate complexes with tunable emission colors†

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Luminescent rare earth Eu(II) complexes with a characteristic 5d–4f transition have shown potential in various applications. However, the exploration and further application of typical Eu(II) complexes are strongly impeded by their instability arising from the easily oxidation of Eu²⁺ ions. In this work, three new Eu(II) complexes bis(hydro-tris(3-phenyl-5-methylpyrazolyl)borate)europium(II) (Eu(Tp^{Ph,Me})₂), bis(hydro-tris(3-phenylpyrazolyl)borate)europium(II) (Eu(Tp^{Ph})₂), and bis(hydro-tris(3,5-diphenylpyrazolyl)borate)europium(II) (Eu(Tp^{Ph₂})₂) were designed and synthesized. Their crystal structures, photophysical properties and air stabilities were characterized and discussed. It is found that the introduction of the bulky phenyl group not only increases the air stability by sterically hindering the central Eu²⁺ ions, but also tunes the maximum emission wavelength by influencing the ligand field. This work provides rules for regulating the luminescence properties of Eu(II) complexes as well as design basis for achieving stable and efficient Eu(II) complexes.

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Introduction

Rare earth elements are widely used in many luminescent materials due to their abundant orbital energy levels and excellent luminescence characteristics.^{1,2} From the perspective of the luminescence mechanism, rare earth elements can be mainly divided into f–f transition and d–f transition ones. For traditional f–f transition rare earth compounds, the f orbitals are in the inner shell and shielded by the filled 5s²5p⁶ electron orbitals. This gives rise to special photophysical properties: relatively fixed absorption and emission wavelengths, narrow-line emission, long excited state lifetime and not easily quenched luminescence by concentration or temperature.¹ Based on these characteristics, f–f transition rare earth compounds are widely used in organic electroluminescence,^{3,4} biological imaging^{5–7} and silicon multicrystalline solar cells.⁸

Different from the f–f transition, the parity-allowed d–f transition gives rare earth compounds a relatively short excited life-

time and broadband emission. Since the 5d electron is not protected by the shielding effect, its energy levels can easily be varied with the external environment. Thus, the emission color of d–f transition rare earth compounds can be tuned in a very broad range.⁹ These unique properties make d–f transition compounds a good prospect in many fields such as near-infrared-emitting phosphors,¹⁰ visible-light-promoted photoredox catalysis,¹¹ phosphor-converted white light-emitting diodes (pc-WLEDs)¹² and organic electroluminescence.¹³ Despite the great progress, d–f transition rare earth complexes, most of them are divalent rare earth complexes, still have a long way to go. The high instability of these complexes to oxygen is a great challenge for the application of such materials.¹⁴ Among them, the Eu(II) ion has an accessible d–f transition state because of its half-filled 4f⁷ electronic configuration and a considerably higher stabilization from exchange energy with a standard electrode potential of –0.35 V.¹⁵ Currently, the research on Eu(II) complexes mainly focuses on macrocyclic complexes,^{16,17} cyclopentadienyl complexes,¹⁸ *etc.* However, the complexes can remain stable in air for a long time are very rare. Solving the stability problem of Eu(II) complexes is the first step to solve the stability problem of divalent rare earth complexes.

Large steric hindrance is an effective way to solve the instability of Eu(II) complexes by protecting the Eu(II) core to minimize its interaction with the environment.¹⁹ Among the reported Eu(II) complexes, Eu(II)-hydro-tris(pyrazolyl)borates

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have demonstrated the highest photoluminescence quantum yield (PLQY).^{13,20,21} Hence, we chose hydro-tris(pyrazolyl) borates as the skeleton ligand, and introduced the big phenyl group at the 3-position and systematically adjusted the substituent at the 5-position of pyrazole. By investigating the structure, photoluminescence properties, and air stability of the resulting three complexes, it is found that they all showed good stability in air with typical and tunable d–f transitions.

Results and discussion

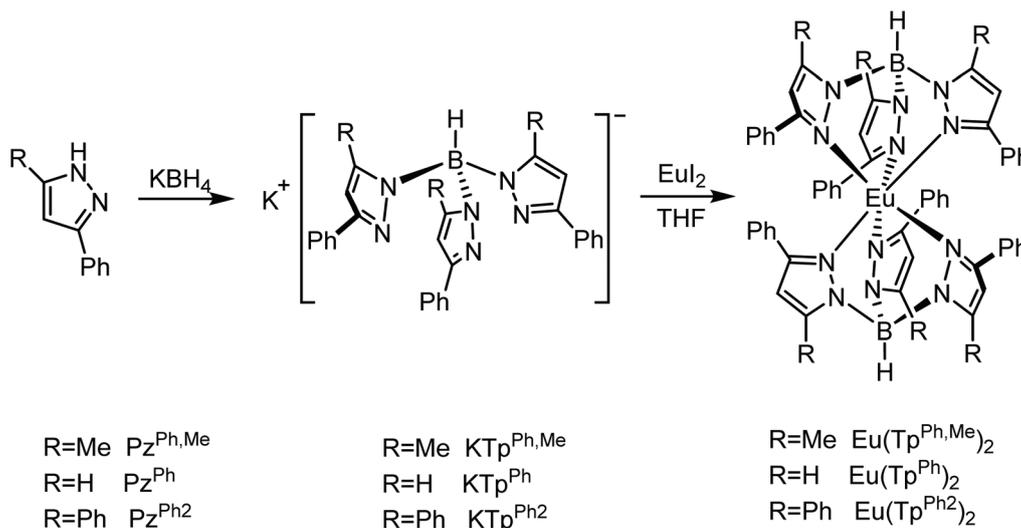
Synthesis and structures

The three Eu(II) complexes $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$, $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ and $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ were synthesized according to the routes shown in Scheme 1. Single crystals of $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$, $\text{Eu}(\text{Tp}^{\text{Ph}})_2$, and $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ were obtained by the slow evaporation of dichloromethane/hexane solution, toluene solution, and sublimation, and characterized by X-ray diffraction analysis (SC-XRD, Table S1†), respectively. The crystal structures of these three complexes are shown in Fig. 1. As shown in the figure, the complexes are all hexa-coordinated, with six nitrogen atoms belonging to pyrazole. $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ is similar in structure to the reported complexes of $\text{Sm}(\text{Tp}^{\text{Ph}})_2$ and $\text{Yb}(\text{Tp}^{\text{Ph}})_2$, which have the same ligands.²² From $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$, $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ to $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$, the average Eu–N bonds increase gradually from 2.652 Å, 2.662 Å to 2.714 Å, respectively (Table 1, the small difference between $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$ and $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ is the result of averaging the difference in bond lengths, as the maximum difference is ~0.03 Å). Besides, there are six phenyl groups around the Eu^{2+} core protecting it from the environment. With the change of substituents, the molecular symmetry of the three complexes increases from $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$ to $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ and $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$. Compared with $\text{Eu}(\text{Tp}^{\text{Ph}})_2$, $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ has a higher symmetry of C_3 , while the “bent sandwich-like” structure of $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$ decreases its symmetry. Moreover,

the interligand C–H...N bonds (2.696 Å and 2.671 Å, shown in Fig. 1) between the H on the phenyl and the N on the pyrazolyl become another driving force for forming the bent structure.^{23,24} The “bent sandwich-like” structure is not usual,^{22,25,26} but the same as the divalent Eu, Yb, Sm, and Tm complexes of $\text{Ln}(\text{Tp}^{\text{iPr}_2})_2$.^{23,27} The B–Eu–B angle of $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$ is 164.28°, greater than 151.07° and 153.88° of $\text{Eu}(\text{Tp}^{\text{iPr}_2})_2$ in the two independent molecules.²⁷ A probable reason is that methyl is less sterically hindered than isopropyl. Without the steric effect of the 5-methyl group, $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ adopts a trigonal antiprismatic molecular structure comprising a linear B–Eu–B arrangement. The phenyl groups around the Eu^{2+} core have C_2 symmetry. For $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$, the three 5-phenyl groups at each vertex form a C_3 symmetry axis in an outward-sloping configuration (the face angles between each of the two phenyl groups are all 62.92°), which greatly reduces the repulsion. So $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ does not adopt a “bent sandwich-like” configuration like $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$, but uses a linear configuration. The 3-phenyl groups around the $\text{Eu}(\text{II})$ core adopt a C_6 symmetry.

Photophysical properties

It is found that $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ is almost insoluble in common solvents such as dimethyl sulfoxide, dichloromethane, and toluene, while $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$ and $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ showed very weak emissions when dissolved in these solvents; hence the photoluminescence properties of the three complexes were only studied in crystalline powder. The maximum emission peaks (λ_{em}) of $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$, $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ and $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ are located at 617 nm, 578 nm and 538 nm (Table 2 and Fig. 2a), demonstrating emission colors as orange-red, orange, and green (Fig. 2b), respectively. As the emission energy distribution of an $\text{Eu}(\text{II})$ complex is generally related to the average bond lengths,²⁸ this blue shift can be explained in the view of comparing the Eu–N bond lengths in these complexes. Among the



Scheme 1 Synthetic routes of $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$, $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ and $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$.

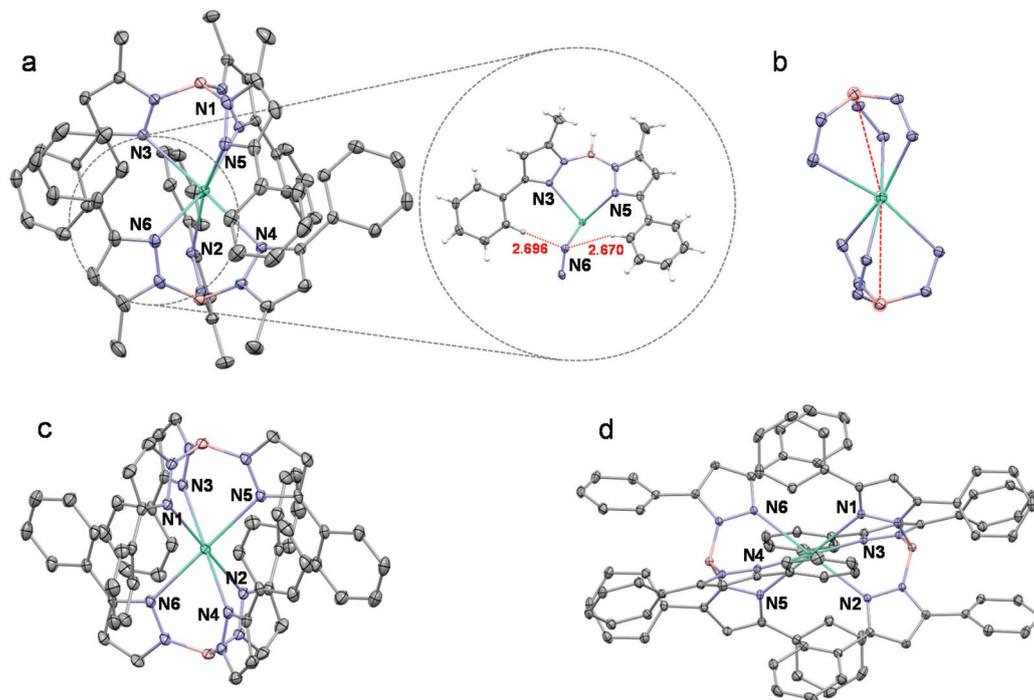


Fig. 1 Molecular structures of $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$ (a, left), $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ (c) and $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ (d). The illustration of interligand C–H...N bonds (a, right) and the description of the “bent sandwich-like” structure of $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$ (b). The atoms are drawn with 30% probability ellipsoids. H atoms are omitted for clarity. The blue-green sphere represents Eu, the pink one represents B, the purple blue one represents N and the gray one represents C.

Table 1 The Eu–N bond distances (Å) in $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$, $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ and $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$

| | $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$ | $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ | $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ |
|----------------|---|--------------------------------------|--|
| Eu–N1 | 2.632 | 2.634 | 2.712 |
| Eu–N2 | 2.632 | 2.634 | 2.712 |
| Eu–N3 | 2.655 | 2.657 | 2.713 |
| Eu–N4 | 2.655 | 2.657 | 2.714 |
| Eu–N5 | 2.666 | 2.695 | 2.715 |
| Eu–N6 | 2.666 | 2.695 | 2.715 |
| Eu–N (average) | 2.652 | 2.662 | 2.714 |

three complexes, the average Eu–N bond lengths for $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$, $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ and $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ increased gradually. Since a shorter bond length makes a stronger ligand field splitting of the d orbitals, the energy levels of the lowest energy 5d orbital increase in the order of $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$, $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ and $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ (Fig. 2c). So, the transition energy from the lowest energy d orbitals to f orbitals increases, assuming that the energy of f orbitals is unchanged in these complexes, which

results in the observed blue shift in emission spectra. In addition, all the three complexes showed broadband emissions, with the full widths at half maximum (FWHMs) gradually decreasing from $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$ (125 nm) to $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ (101 nm) and $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ (70 nm). Such a broad emission band is typical of the 5d–4f transition of the Eu(II) core, and the tendency from $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$ to $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ might be caused by the increasing rigidity in turn. By comparing the excitation spectra of the three Eu(II) complexes and their corresponding ligands (Fig. S1†), an obvious broadening of the spectra was observed, indicating that both excitation from the ligand and the central Eu^{2+} ion are accessible to achieve an efficient 5d–4f transition in the three Eu(II) complexes.

The excited state lifetimes (τ) of these three complexes were also measured (Fig. 2d and Table 2). Coincidentally, an increased lifetime from $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$ (34 ns) to $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ (118 ns) and $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ (471 ns) was observed. These lifetimes are comparable to those of the reported Eu(II) complexes,¹⁶ but the huge differences in these three complexes are worthy of further study. Therefore, PLQYs were measured to deduce the

Table 2 Photophysical data of $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$, $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ and $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ in crystalline powder

| Complex | λ_{ex} (nm) | λ_{em} (nm) | τ (ns) | PLQY (%) | FWHMs (nm) | k_r (10^6 s^{-1}) | k_{nr} (10^6 s^{-1}) |
|---|----------------------------|----------------------------|-------------|----------|------------|---------------------------------|---|
| $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$ | 465 | 617 | 34 | 3 | 125 | 0.88 | 28.5 |
| $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ | 440 | 578 | 118 | 20 | 101 | 1.69 | 6.78 |
| $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ | 414 | 538 | 471 | 70 | 70 | 1.49 | 0.63 |

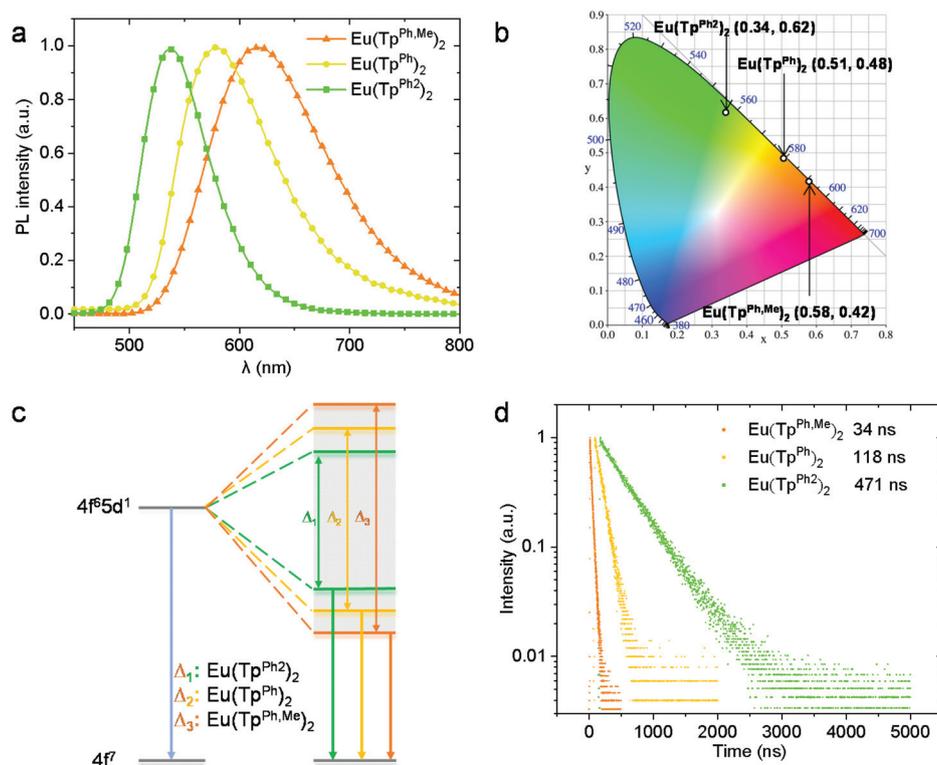


Fig. 2 (a) Solid steady-state PL spectra (photoexcitation wavelength: 365 nm) of $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$, $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ and $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$. (b) The CIE 1931 chromaticity diagram of $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$, $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ and $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$. (c) Orbital-energy diagram for the ligand field splitting of the d orbitals (double arrow) and the energy transition (single arrow) of $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$, $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ and $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$. (d) Transient PL spectra of $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$, $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ and $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$.

radiative rate constant (k_r) and non-radiative rate constant (k_{nr}). Among these complexes, $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ showed a high PLQY of 70%, which is the highest value for an green emission Eu(II) complex up to now, while $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$ and $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ showed a relatively lower PLQY of 3% and 20%, respectively. So far, the maximum PLQY of Eu(II) complexes is 96%,²¹ but the PLQYs of most Eu(II) complexes are relatively low, below 50% (Table S2†). The radiative rate constants, calculated from the aforementioned data, are shown in Table 2. In order to better understand the regularities, we also introduced a published complex bis(hydro-tris(3,5-dimethylpyrazolyl)borate)europium(II) ($\text{Eu}(\text{Tp}^{\text{Me}_2})_2$) with good photoluminescence properties for comparison¹³ ($\tau = 987$ ns, PLQY = 85%, $k_r = 0.86 \times 10^6$ s⁻¹, $k_{nr} = 0.15 \times 10^6$ s⁻¹). It can be concluded that their radiative rate constants are similar, while the non-radiative rate constants vary in an order of magnitude. We propose that increased rigidity decreases the non-radiative transition rate. 5-Phenyl in $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ and 5-methyl in $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$ would increase the molecular rigidity. But the forming “bent sandwich-like” structure can be more randomly arranged in space, which allows us to ignore the rigidity of the methyl group. So the rigidity increases from $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$, $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ to $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$. As for $\text{Eu}(\text{Tp}^{\text{Me}_2})_2$, the decreased non-radiative transition rate may be due to a stronger combination between Eu(II) and the coordination N atoms in the ligands (the average length of Eu–N

bonds is 2.585 Å, much shorter than those for $\text{Eu}(\text{Tp}^{\text{Ph,Me}})_2$, $\text{Eu}(\text{Tp}^{\text{Ph}})_2$ and $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$).¹⁶

Air stability and its rationalization

It is well known that the air stability of typical Eu(II) complexes is poor because the standard electrode potential of Eu^{2+} is as low as -0.35 V. So we introduced phenyl as a large steric hindrance group around Eu(II) to improve its air stability. In view of this, the complex $\text{Eu}(\text{Tp}^{\text{Me}_2})_2$ was adopted as a reference. The luminescence stability of the complexes was described by the change of the PLQY in air and which is further used to understand the stability of the complexes. As shown in Fig. 3a, the PLQY of $\text{Eu}(\text{Tp}^{\text{Me}_2})_2$ decreased to 7% of its original value after one day and to only 3% after 4 days. In contrast, the PLQY of $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ decreased to 94% after one day and to 87% even after 10 days. After the tenth day of measurement, $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ was placed in a glove box and taken out the next day to measure its PLQY. To our delight, the PLQY returned to the original value. This indicates that the decrease of the PLQY of $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ is not caused by the oxidation of Eu(II) but probably due to the quenching effect of oxygen or water in air. To investigate further, two atmospheres, one with pure oxygen and another with water-containing nitrogen, were created. Then $\text{Eu}(\text{Tp}^{\text{Ph}_2})_2$ was loaded into the two atmospheres for 3 hours. It was found that the PLQYs of the samples in a pure oxygen

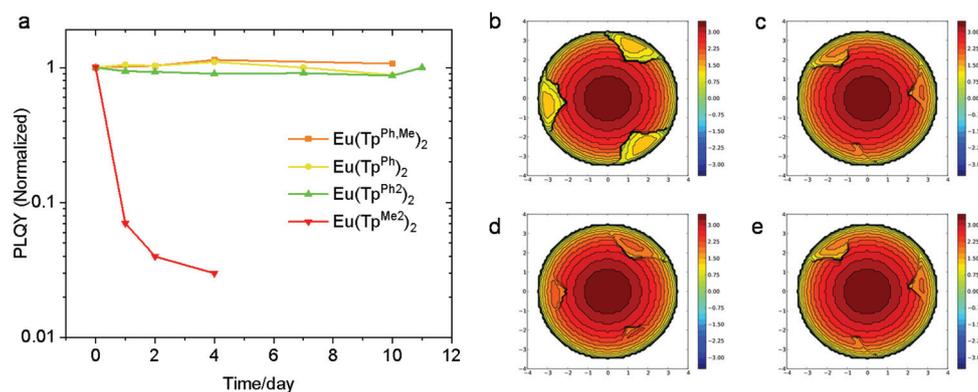


Fig. 3 (a) The normalized PLQY changes of crystalline Eu(Tp^{Me2})₂, Eu(Tp^{Ph,Me})₂, Eu(Tp^{Ph})₂ and Eu(Tp^{Ph2})₂ in air. Topographic steric maps of Eu(Tp^{Me2})₂ (b), Eu(Tp^{Ph,Me})₂ (c), Eu(Tp^{Ph})₂ (d) and Eu(Tp^{Ph2})₂ (e). Red and blue indicate the more- and less-hindered zones, respectively.²⁹

atmosphere and water-containing nitrogen atmosphere were reduced to 52% and 85% of the initial value, respectively. After the test, the samples were transferred into a glove box (under a nitrogen atmosphere and at room temperature, with the oxygen concentration below 1.0 ppm and the water concentration below 0.1 ppm), and their PLQYs were measured again a day later. The PLQYs of both the samples returned to the initial values. Based on this phenomenon, we propose that the complex can absorb oxygen and water and both oxygen and water can quench the luminescence of the complex to a certain extent, in which oxygen is the main quencher. When the sample is returned to the glove box, oxygen and water are desorbed from the surface of the complex, allowing the PLQY of Eu(Tp^{Ph2})₂ to recover. The PLQYs of Eu(Tp^{Ph,Me})₂ and Eu(Tp^{Ph})₂ were kept almost the same to the original data (the accurate PLQY values of the four complexes are shown in Table S3†). To further understand the air stability of these complexes, time dependent XRD spectra were obtained as shown in Fig. S2.† During the four-day test, the XRD peak position did not change, and no new peak was observed. In addition, mass change monitoring upon isothermal exposition of the complexes under an air flow has also been conducted (Fig. S3†). The maximum weight change is 0.0060 mg, which is within the allowable range of measurement error of the instrument. These findings further demonstrate the good air stability of the three complexes.

In order to understand the excellent air stability of the three complexes, we investigated the extent to which the central metal is coated in the three complexes and Eu(Tp^{Me2})₂. The spacefill views along the largest exposed area of the four complexes are shown in Fig. S4.† At a glance, the exposed area of Eu(Tp^{Me2})₂ is much larger than those of the other three complexes. To further understand the protection of the ligands to the central Eu²⁺ ions, buried volume descriptor (% *V*_{Bur}) was introduced to quantify the steric hindrance.³⁰ With the help of SambVca 2.1,³¹ the topographic steric maps of the four complexes are drawn in Fig. 3b–e, and the calculated % *V*_{Bur} values of Eu(Tp^{Me2})₂, Eu(Tp^{Ph,Me})₂, Eu(Tp^{Ph})₂ and Eu(Tp^{Ph2})₂ are 84.1%, 91.0%, 91.7% and 92.9%, respectively.³⁰

Based on these results, it can be seen that the contribution of the methyl and phenyl groups to steric hindrance is very different. By substituting the methyl group with the phenyl group at the 3-position of pyrazole, Eu(Tp^{Ph,Me})₂ showed an obviously increased %*V*_{Bur} as compared with Eu(Tp^{Me2})₂. However, the further substitution of the methyl group with the phenyl group at the 5-position shows only a little difference in %*V*_{Bur}. Thus, the steric hindrance is mainly caused by introducing the phenyl group at the 3-position of pyrazole.²⁹

Conclusions

In summary, three new Eu(II) complexes Eu(Tp^{Ph,Me})₂, Eu(Tp^{Ph})₂, and Eu(Tp^{Ph2})₂ were designed and successfully synthesized. Due to different steric hindrances of the ligand, these complexes adopt a “bent sandwich-like” structure for Eu(Tp^{Ph,Me})₂, and a linear configuration for Eu(Tp^{Ph})₂ and Eu(Tp^{Ph2})₂. All the complexes exhibited broad peak emission with the excited state lifetime in dozens to hundreds of nanoseconds, which is a typical 5d–4f transition characteristic. From Eu(Tp^{Ph,Me})₂, Eu(Tp^{Ph})₂, to Eu(Tp^{Ph2})₂, the average Eu–N length increases gradually, leading to decreased splitting energy of the d orbital, hence a blue shifted emission spectrum. However, the rigidity increases gradually in the order of Eu(Tp^{Ph,Me})₂, Eu(Tp^{Ph})₂, and Eu(Tp^{Ph2})₂, resulting in a significant increase in the excited state lifetime and PLQY. Consequently, Eu(Tp^{Ph2})₂ showed the highest PLQY up to 70%. To our delight, all the three complexes showed excellent stability in air, as demonstrated by measuring their time dependent PLQYs, XRD spectra and mass change in the presence of air. By comparing the spacefill views and buried volume descriptors of the three complexes with a reference Eu(II) complex Eu(Tp^{Me2})₂, it is found that the introduction of a phenyl group at the 3-position of pyrazole accounts for the improved stability. This work systematically reports the relationships between the photoluminescence/air stability and the structure of molecular Eu(II) complexes, which gives the feasibility to discover

highly efficient and air stable luminescent Eu(II) complexes with specific emission colors.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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