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1 Efficient Luminescence Control in Dithienylethene
2 Functionalized Cyclen Macrocyclic Lanthanide
3 Complexes

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23 TITLE RUNNING HEAD: Efficient Luminescence Control in a Dithienylethene Functionalized
24 Cyclen Macrocyclic Lanthanide Complex

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4 **Abstract.** We report the synthesis of an original ligand scaffold based on a dimethyl-cyclen
5 platform Medo2pa with two dithienylethene units attached to each picolinate arms and the
6 corresponding yttrium(III), europium(III) and ytterbium(III) complexes. All three compounds
7 show reversible photochromism with high photo-conversions. Photoluminescence experiments
8 demonstrate that this design is versatile and adapted for both europium and ytterbium emission
9 switching when measured in frozen organic glasses at 77 K. The OFF/ON luminescence ratio are
10 excellent in the case of europium (4 to 8 %) and still quite good in the case of ytterbium (around
11 13 %).

12

13 **Keywords:** photochromism, ytterbium(III), europium(III), dithienylethene, cyclen,
14 azamacrocycles, luminescence, molecular switch.

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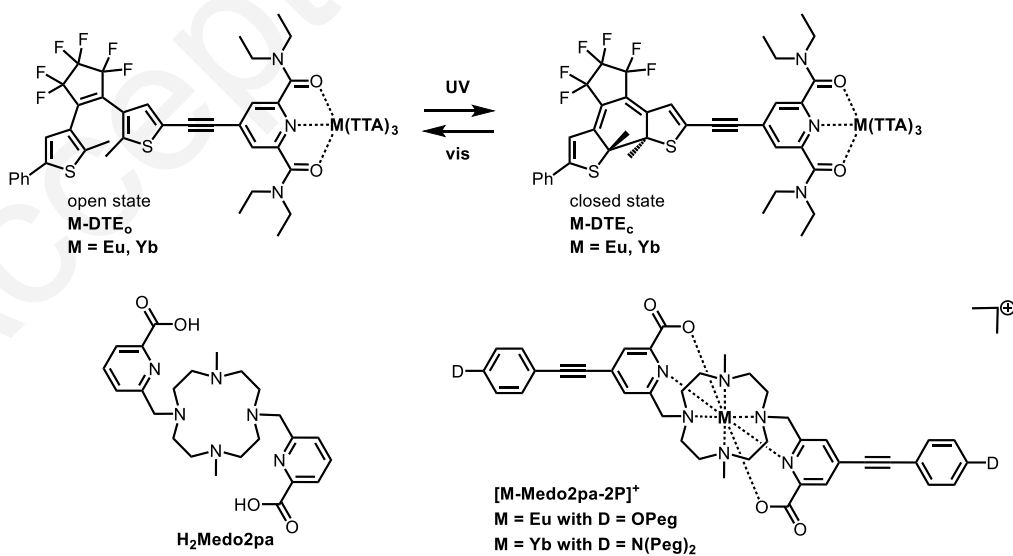
16 **Introduction**

17 Responsive materials in which a key property can be modulated by an external stimulus in a
18 controlled way are a great achievement in the field of molecular materials.¹⁻² Among them “all-
19 optical” systems, that are triggered by light to change their optical (absorption, emission)
20 properties, combine fast response, remote control and a low level of technical requirements for
21 their implementation in real life applications. Applications could be as diverse as labels for cell
22 imaging,³ super resolution imaging,⁴ anti-counterfeiting dyes,⁵⁻⁶ optical data-storage⁷ and many
23 others.

24 In this context, several research groups have explored the photo-modulation of lanthanide-based
25 luminescent systems,^{5, 8-16} mainly focusing on the association of photochromic compounds with
26 the red-emitting europium(III) ion.^{5, 10-16} The ubiquitous diarylethene (DAE) photochromic units,¹⁷
27 on top of their excellent photo-physical properties, fatigue resistance and thermal stability of both
28 open and closed isomers, is perfectly suited. Indeed, DAE scaffolds can be easily designed so that

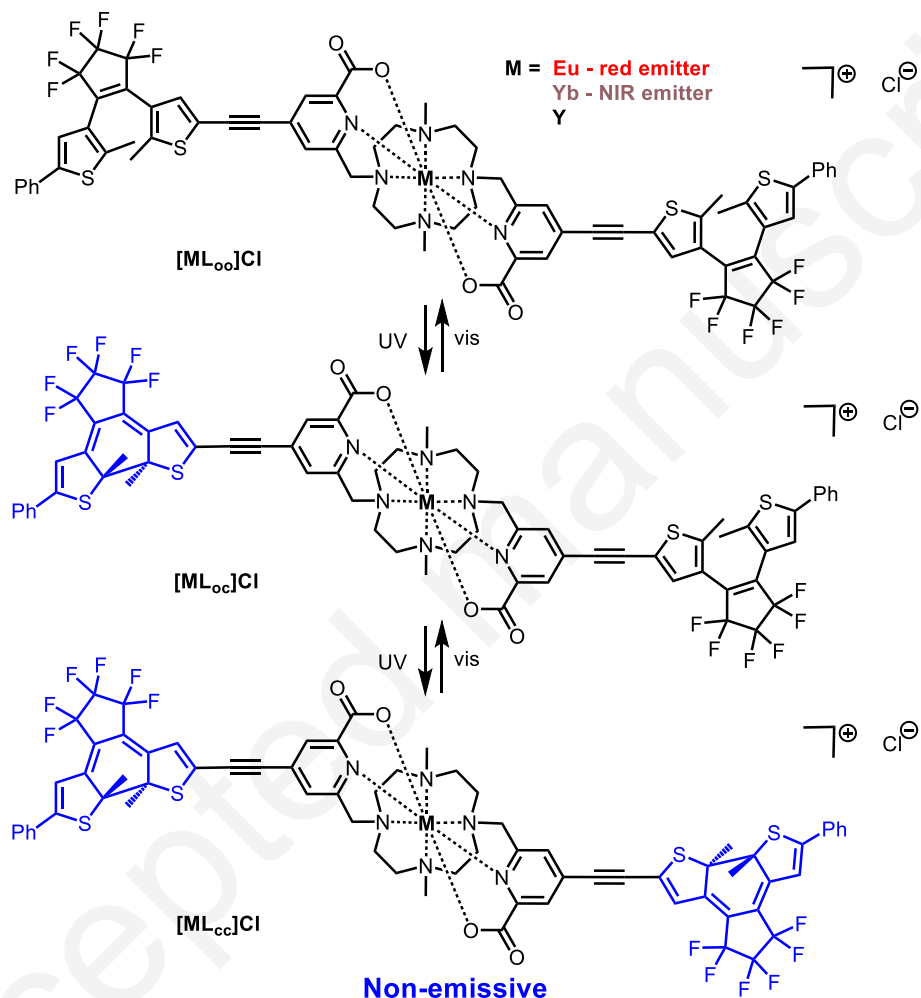
1 the closed isomers show strong absorptions around 610 nm, matching the narrow emission lines of
 2 europium(III) and then favoring emission quenching typically via an energy transfer. However,
 3 according to this strategy, a complete quenching of europium luminescence in the closed form has
 4 not been realized yet. The only total quenching of europium luminescence by a photochromic unit
 5 reported to date consist of a tris(dipicolinate)europium core decorated with three N[^]C chelate four
 6 coordinate organoboron T type (reversible upon heating) photoswitches.¹⁵ Therefore, it is highly
 7 desirable to achieve a complete optical control of ON/OFF switching of europium luminescence
 8 with the P (thermally stable) photochromic DAE. Recently, some of us reported an example of
 9 partial photo-modulation in a dithienylethene (DTE) appended dipicolinic amide europium
 10 complex (Chart 1),¹⁸ and we hypothesize that a partial lability of the metal complex could be a
 11 factor contributing to the moderate efficiency of the quenching in the closed form. At the same
 12 time, surprisingly, this previous paper showed that DTE photochromic units could actually be more
 13 versatile modulators of lanthanide luminescence than initially thought since ytterbium(III) NIR
 14 emission could be sensitized by the 580 nm absorption of the closed isomer. Based on this, two
 15 important goals remain to be achieved in this field: i) the improvement of the efficiency of
 16 europium(III) emission quenching by closed DTE system in order to reach real ON/OFF switching,
 17 and ii) the generalization and optimization of photo-modulation of ytterbium(III) ion by DTE units.
 18 These two goals thus require a better understanding of the underlying photo-physical mechanisms
 19 and the exploration of new systems combining DTE and lanthanide ions.

20



21

1 **Chart 1.** Previously studied systems. (top) **M-DTE** system shows versatile photo-modulation of
 2 emission of both ytterbium(III) and europium(III) ions upon switching between the open and closed
 3 state (TTA is 2-thenoyltrifluoroacetate).¹⁸ (middle) **Medo2pa** provides water soluble and stable
 4 lanthanide complexes and **M-Medo2pa-2P** chlorine salts enable cell imaging in the NIR range in
 5 the case of the ytterbium(III) complex.¹⁹⁻²⁰



6
7
8
9 **Chart 2.** Targeted luminescence switches.

10 In parallel, macrocyclic lanthanide complexes have been widely studied as imaging bioprobes in
 11 general,^{3, 21-22} and as luminescent systems in particular.²³⁻²⁴ Among them, the cyclen based
 12 Medo2pa platform (Chart 1) has provided complexes of various lanthanide ions displaying high
 13 stability constants,²⁵ that are typically stable in water solutions.²⁶ This cyclen platform N-
 14 functionalized by two picolinate pendants and two methyl groups, when modified with two photon
 active conjugated antennas, provides bright luminescent europium(III) and ytterbium(III)

1 complexes that are spontaneously internalized into live cells,¹⁹⁻²⁰ the latter remaining highly
2 luminescent in biological media (Chart 1).²⁰ Based on these convincing results, and complementary
3 to another strategy on based DTE modified acetyl acetonate ligands that we are developing in
4 parallel,²⁷ we thought that the association of the Medo2pa platform with appropriate DTE units
5 could lead to “all optical” switches with improved stability and, therefore, better switching ratio
6 between the open and closed state, as well as to provide a new efficient ytterbium based switch in
7 the NIR range through the closed DTE unit sensitization. We therefore targeted the synthesis of a
8 new Medo2pa platform bearing two DTE units (on each picolinate arms) as shown in Chart 2. First
9 motivated by the ease of synthesis, the presence of two photochromic units within the same scaffold
10 could also be anticipated as an advantage to improve i) quenching efficiency in the case of the
11 europium(III) complex, and ii) sensitization through the closed DTE unit in the case of the
12 ytterbium(III) complex. In this paper, we report on the synthesis of this new ligand and of the
13 corresponding europium(III), ytterbium(III) and yttrium(III) complexes. We study in detail the
14 photo-switching of these three metal complexes by absorption and (¹H, ¹⁹F) NMR spectroscopies
15 to illustrate that a reversible and complete isomerization occurs, the two DTE units behaving
16 independently. Our strategy is proved effective in improving the quenching efficiency of europium
17 luminescence as shown by a residual intensity of 4-8 % of the initial one for the closed form as
18 compared to the open one when measured at 77 K. We also show that the ytterbium complex
19 luminescence can be modulated at 77 K although it does not exhibit any sensitization through the
20 closed DTE.

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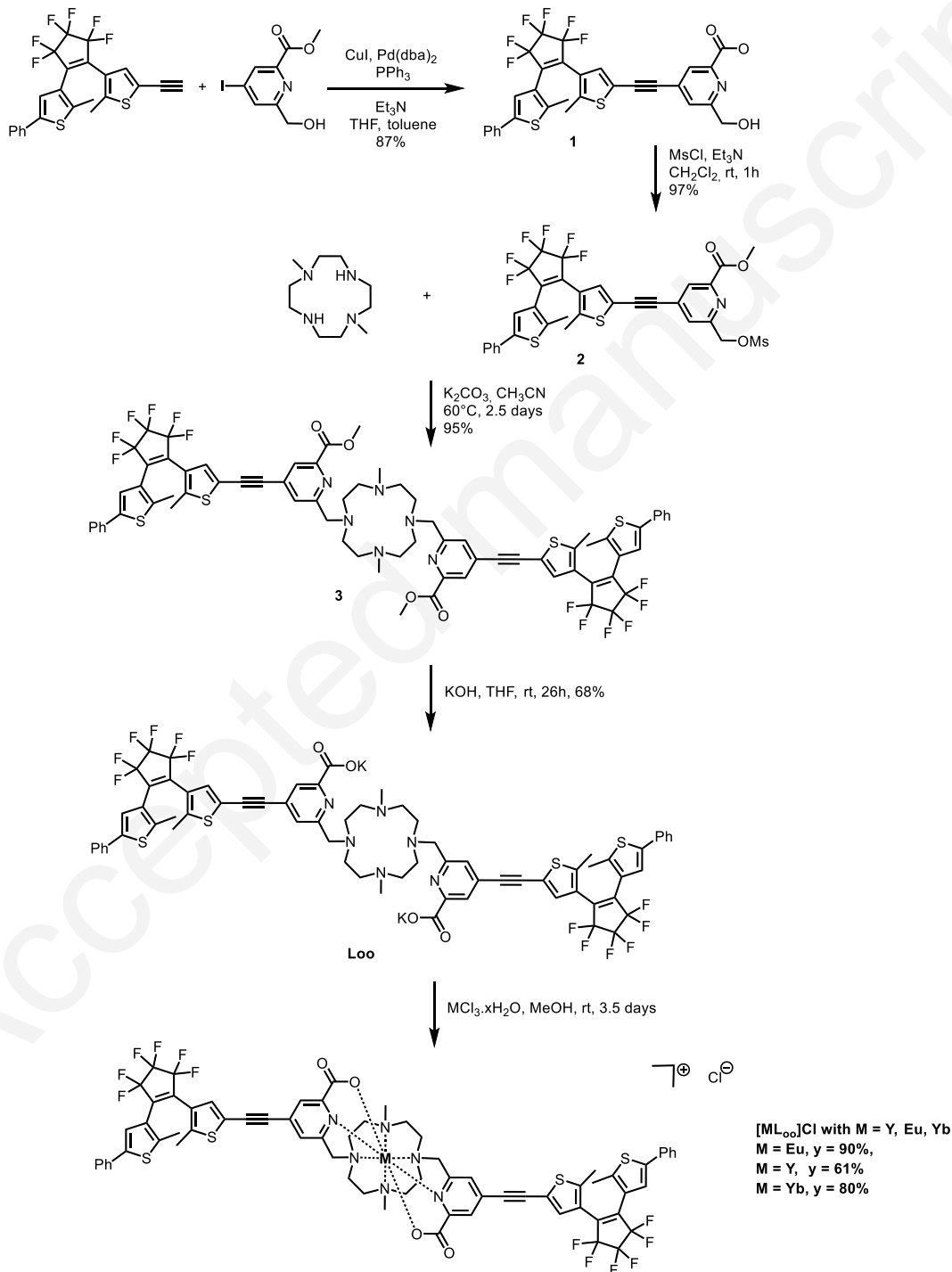
22 **Results and Discussion**

23 **Complex synthesis.** Synthesis of the target metal complexes [ML_{oo}]Cl (M = Y, Eu, Yb) is
24 described in Scheme 1. The DTE-photochromic-picolinate arm **1** was obtained by Sonogashira
25 coupling from the alkyne terminated DTE and methyl 6-(hydroxymethyl)-4-iodopicolinate²⁸ (see
26 SI). Mesylation of the latter was performed under usual conditions and *trans*-dialkylation of the
27 dimethyl-cyclen macrocycle with two equivalents of compound **2** in the presence of K₂CO₃ led to
28 the desired di-ester **3** with an excellent yield of 95%. Saponification of compound **3** in the presence
29 of KOH in THF led to L_{oo} as a potassium salt which was purified, thanks to a precipitation in an
30 EtOAc/hexane mixture. The synthesis of the compounds was further performed in MeOH at pH

1 around 7. Washings with water and precipitations in CH₂Cl₂/hexane gave the desired [ML_{oo}]Cl
2 with yields comprised between 61% and 90%. These new compounds were fully characterized (see
3 experimental section and SI). As characteristic features in its ¹H NMR spectrum, diamagnetic
4 [YL_{oo}]Cl exhibit shielded pyridine protons chemical shifts, similarly to other yttrium(III) dimethyl
5 cyclen complexes,²⁹ while the signals from the cyclen moiety become significantly broadened upon
6 coordination (Figure S12). In the case of [EuL_{oo}]Cl, additional paramagnetic shifts (pseudo contact
7 shifts) are observed. Typically, the photochromic moiety shows small paramagnetic shifts, of
8 around -0.1/-0.2 ppm as compared with the yttrium(III) complex, while the pyridine protons are
9 observed at $\delta = 38.4$ and 25.8 ppm and the cyclen protons give broad signals down to -16 ppm as
10 expected (Figure S21).¹⁹ For [YbL_{oo}]Cl, in line with the greater magnetic anisotropy tensor of
11 ytterbium(III) compared with europium(III),³⁰ shifts of the same sign but of greater magnitude are
12 observed, the pyridine protons being observed at $\delta = 83.8$ and 55.5 ppm and the cyclen ones down
13 to $\delta = -40.5$ ppm (Figure S18). The paramagnetic shifts observed for the photochromic moiety are
14 also larger with, for instance, the thiophene protons shielded to $\delta = 6.97$ and 6.37 ppm instead of δ
15 $= 7.47$ and 7.28 ppm in [YL_{oo}]Cl.

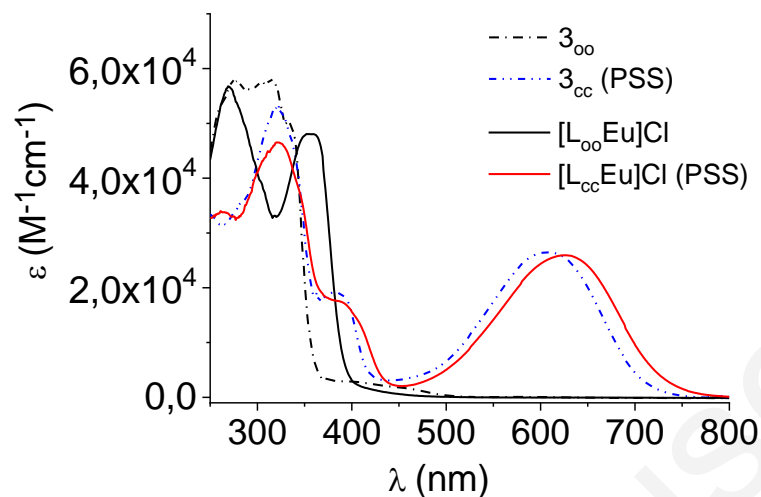
16 **Electronic absorption spectra and photochromism of 3_{oo} and [ML_{oo}]Cl.** The absorption
17 spectrum of 3_{oo} in DCM shows several intense bands in the UV range (Figure 1) that can be
18 assigned to local $\pi-\pi^*$ transition of the picolyl unit (275 nm) overlapping with one of the DTE
19 open form (315 nm). Upon irradiation at 330 nm, a decrease of absorption is observed at $\lambda_{\max} = 272$
20 nm while two new bands appear at $\lambda_{\max} = 382$ and 607 nm (Figure 1). The initial spectrum can be
21 recovered by 580 nm irradiation. This is in line with the usual photochromic behavior of DTE
22 units¹⁸ and consistent with the above mentioned assignment of the bands. Photo-cyclisation is
23 evidenced by the characteristic lower energy band ($\lambda_{\max} = 607$ nm) ascribed to an intra-ligand (IL)
24 transition centered on the closed DTE moiety.¹⁷ In this system with two DTE units, isomerization
25 proceeds through the intermediate 3_{oc} compound with one closed ring. However, at intermediate
26 photo-conversions, no shifting of the lower energy transition was observed, suggesting that the two
27 DTE units are electronically decoupled and behave independently in that case (Figure S24).³¹ The
28 isomerization was also studied by ¹H NMR spectroscopy that proved that a high photo-
29 isomerization conversion (up to 94 % of 3_{cc} and 6 % of 3_{oc}) can be reached in the photo-stationary
30 state (PSS) (Figure S29). Typically, the thienyl protons chemical shifts change from $\delta = 7.25$ and

1 7.31 ppm in **3_{oo}** to 6.72 and 6.43 ppm in **3_{cc}**. In the NMR conditions ($[c] = 1.2 \times 10^{-3}$ M), the
 2 cycloreversion process is almost quantitative with the recovery of **3_{oo}** in 94 % yield accompanied
 3 by unknown species, probably coming from partial degradation upon prolonged exposure to light.
 4 This behavior is in contrast to the more diluted UV-vis experiment that displays quantitative
 5 recovering.



1

Scheme 2. Synthetic pathway yielding the target compounds.



2

3 **Figure 1.** Electronic absorption spectra of **3_{oo}** and **[EuL_{oo}]Cl** in CH₂Cl₂ solutions. Upon UV light irradiation
4 to PSS, the spectra of **3_{cc}** and **[EuL_{cc}]Cl** are obtained. The other compounds (Y, Yb) show similar behavior
5 (see Table 1 and Figures S27 and S28). The initial spectra were recovered after bleaching at 580 nm.

6

7 Once the cyclen group is coordinated, clean photochromic behavior, without photo-degradation
8 was observed as evidenced by the presence of isobestic points. The absorption spectra of the three
9 metal complexes are very similar with two main transitions at $\lambda_{\text{max}} = 269$ nm and 350-360 nm and
10 Figure 1 shows the representative behavior of the europium complex (the cases of Y and Yb
11 complexes are depicted in Figures S27 and S28 respectively). Both bands are strongly modified
12 upon UV irradiations and subsequent ring closure, and new transitions appear with λ_{max} values of
13 330 and 627 nm. The lower energy transition is slightly red shifted upon coordination as compared
14 with **3_{cc}**. Under visible light irradiation ($\lambda_{\text{max}} = 580$ nm), the cycloreversion process is triggered as
15 attested by the quantitative recovery of the initial spectra. Further ¹H NMR spectroscopy
16 monitoring of the process unambiguously shows that the photochromic process upon UV
17 irradiation is almost complete with the reaching of a photo-stationary state composed of ca. 95 %
18 of closed DTE units and a recovery of the initial spectra upon 580 nm irradiation, in contrast to the
19 organic precursor. Details of the changes in the NMR spectra are highlighted in Figures 2 and S31.

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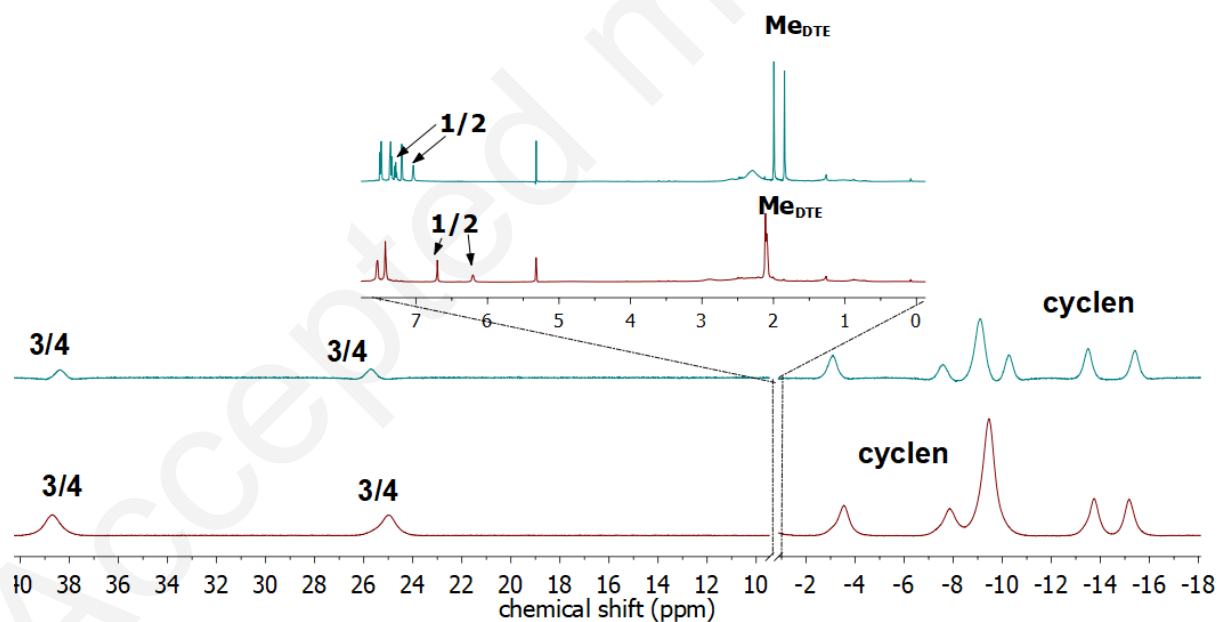
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compound	$\lambda_{\text{max}}[\text{nm}]$, corresponding energy [cm^{-1}]	$\epsilon [\text{M}^{-1} \cdot \text{cm}^{-1}]$
3_{oo}	340, 29400	40000
3_{cc}^(a)	382, 26200 607, 16500	19200 28000
[YL_{oo}]Cl	269, 37200 362, 27600	55600 48300
[YL_{cc}]Cl^(b)	330, 30300 627, 15900	42900 27800
[EuL_{oo}]Cl	269, 37200 352, 28400	53400 45300
[EuL_{cc}]Cl^(b)	339, 29500 627, 16000	39400 24700
[YbL_{oo}]Cl	269, 37200 347, 28800	57400 45300
[YbL_{cc}]Cl^(b)	330, 30300 627, 15900	46000 25900

2 a) Photo-stationary state after 330 nm irradiation, b) photo-stationary state after 364 nm irradiation

3 **Table 1.** Electronic absorption data for all compounds measured in DCM solutions.

4

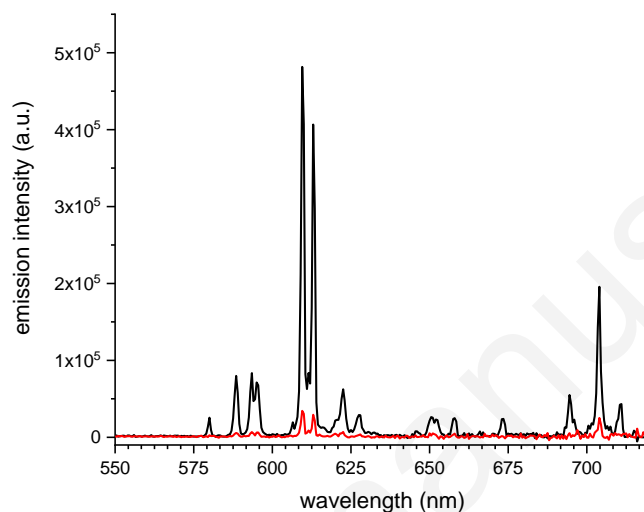


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6 **Figure 2.** ^1H NMR spectra (500 MHz, CD_2Cl_2 , 298 K) measured during the course of $[\text{L}_{\text{oo}}\text{Eu}]\text{Cl}$
7 (blue line) isomerization to the PSS containing $[\text{L}_{\text{cc}}\text{Eu}]\text{Cl}$ and $[\text{L}_{\text{oc}}\text{Eu}]\text{Cl}$ with 95 % of overall
8 closed DTE units (red line) upon 364 nm excitation in CD_2Cl_2 . See Scheme 3 (experimental
9 section) for numbering.

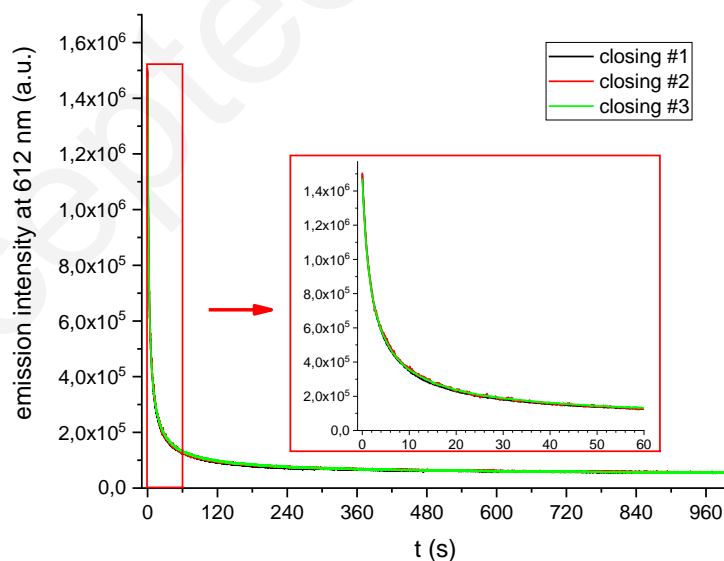
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1 **Photoluminescence of [ML_{oo}]Cl.** We further studied the photoluminescence of all three metal
2 complexes (M = Y, Eu, Yb). The yttrium complex serves as a reference to understand the photo-
3 physics of the ligand since no metal-based emission is expected for this compound. Thus, upon
4 excitation at $\lambda_{\text{ex}} = 350$ nm of [YL_{oo}]Cl in an ethanol:methanol glass (77 K), a ligand-based
5 fluorescence centered at $\lambda_{\text{em}} = 395$ nm was observed (Figure S32) with the presence of additional
6 peaks in its tail. A time-gated measurement performed with a 1 ms delay allows us to assign
7 unambiguously these features to a simultaneous structured phosphorescence with maximum at λ_{em}
8 = 517 nm and a corresponding lifetime of 14 ms at 77 K (Figure S33). This phosphorescence
9 process corresponds to a ligand-centered triplet at around 19 000 cm⁻¹. Upon continuous irradiation
10 at 350 nm and closing of the DTE units, both fluorescence and phosphorescence progressively
11 disappeared, and at the PSS the closed yttrium(III) complex was almost non-emissive (Figure S32).
12 Concerning the spectroscopy of the europium(III) complex, [EuL_{oo}]Cl was studied at room
13 temperature and at 77 K. At room temperature, excitation at 350 nm induces both emission and
14 competitive closing of the DTE units. The spectrum is actually dominated by an intense ligand-
15 centered emission at $\lambda_{\text{em}} = 395$ nm accompanied by a weak europium emission at 616 nm (Figure
16 S34). In contrast, at 77 K in a methanol/ethanol organic glass, ligand centered emission is
17 drastically decreased as compared with the sharp f-f transitions. The difference in the response of
18 the system with temperature could be ascribed to the occurrence of thermally activated back energy
19 transfer that is hampered at 77 K. We also observed a drastic slowing down of the closing reaction
20 by this lowering of temperature and immobilization in an organic glass that allows one to measure
21 the emission spectrum of pure [EuL_{oo}]Cl with an excellent resolution. Therefore, the characteristic
22 europium(III) emission profile assigned to the ⁵D₀ → ⁷F_J (J = 0-4) transitions were detected at λ_{em}
23 = 580 (J = 0), 588, 593, 595 (J = 1), 610, 613, 622, 627 (J = 2), 646, 650, 652, 658, 673 (J =
24 3), and 694, 704, 711 nm (J = 4) (Figure 4) and overall, the spectrum and particularly the crystal
25 field splitting, is very similar to the one of a previously published europium(III) complexes with a
26 similar Medo₂pa ligand for which a C₂ symmetry was calculated by DFT.¹⁹ The same measurement
27 at 77 K was performed on [EuL_{cc}]Cl (PSS state) and showed that an impressive quenching of
28 europium luminescence occurs after closing of the DTE since only very weak emission (about 8 %
29 of the original intensity determined by integration of the open state more intense band (J = 2), see
30 Figure 3) was detected. It is also possible to follow the emission quenching in the glass at 77 K
31 upon successive scans, highlighting the progressive closing of the DTE during each luminescence

1 measurement (Figure S35). An attempt to reach the PSS was performed upon irradiation of the
2 glass during 1000 s. A 90 % quenching was achieved after only 40 s but the complete closing was
3 not reached at the end of the experiment where less than 4% of the initial emission was still
4 observed. A perfect reproducibility of the behavior was observed after several re-opening were
5 performed with white light irradiation (Figure 4).



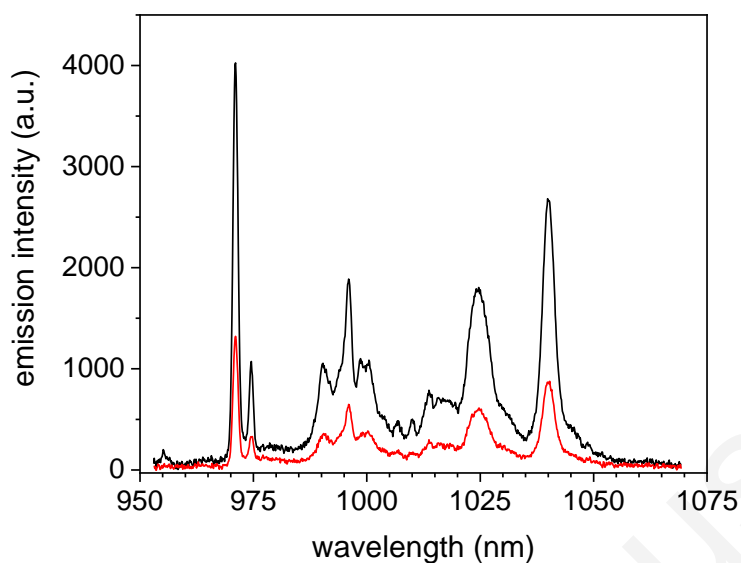
17 **Figure 3.** Emission spectra ($\lambda_{\text{ex}} = 350 \text{ nm}$) of $[\text{L}_{\text{oo}}\text{Eu}]\text{Cl}$ (black line) and $[\text{L}_{\text{cc}}\text{Eu}]\text{Cl}$ (red line) in
18 ethanol: methanol (4:1) organic glass at 77K.



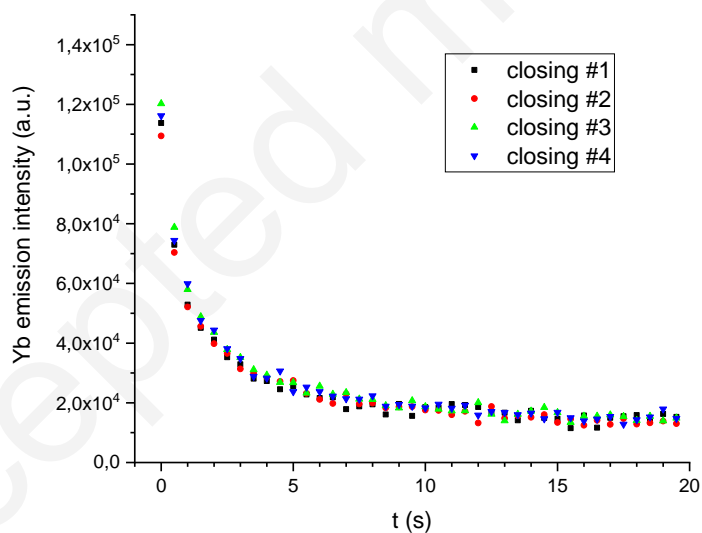
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1 **Figure 4.** Emission at 612 nm ($\lambda_{\text{ex}} = 350$ nm) of **[L_{oo}Eu]Cl** while isomerizing to **[L_{cc}Eu]Cl** in
2 ethanol : methanol (4:1) organic glass at 77K. Two other successive measurements are performed
3 after discoloration with white light (at RT).

4
5 Concerning **[YbL_{oo}]Cl**, no ytterbium emission was detected upon 350 nm excitation at room
6 temperature. In contrast, in an ethanol/methanol organic glass at 77 K, the typical emission of
7 ytterbium(III) was detected in the NIR. In order to avoid distortion of the signal due to concomitant
8 closing, the emission was detected with a CCD camera. First, a resolved spectrum can be obtained,
9 clearly showing the different lines expected for the $^2F_{5/2} \rightarrow ^2F_{7/2}$ transition and again very similar
10 to previously reported ytterbium(III) complexes with C_2 symmetry,²⁰ with the main crystal field
11 splitting lines at 971, 996, 1025 and 1040 nm (Figure 5). In order to follow the effect of photo-
12 isomerization on ytterbium emission, fast-acquired successive spectra were obtained, clearly
13 showing a 10 fold quenching of luminescence due to the closing reaction (Figure S36). Note that
14 the quenching ratio is not rendered by Figure 5 because the initial intensity actually corresponds to
15 a system already undergoing a significant amount of closing. Rather, the ratio between the initial
16 and final states can be obtained from integration of the fast acquired data (Figure 6), giving a 13 %
17 ratio. Finally, we have addressed the possibility of sensitization by excitation at 600 nm, and unlike
18 **Yb-DTE_c** (Scheme 1) no ytterbium emission was detected in such case.¹⁸ For both europium and
19 ytterbium complexes, it is unclear whether the remaining emission after closing arises from the
20 closed species or whether a PSS composition different from the one in DCM solutions at room
21 temperature (95 % of closed units, no remaining oo isomer) is reached due to immobilization in a
22 frozen organic glass.



1
 2 **Figure 5.** Emission spectra ($\lambda_{\text{ex}} = 350 \text{ nm}$) of $[\text{L}_{\text{oo}}\text{Yb}]\text{Cl}$ (black line) and $[\text{L}_{\text{cc}}\text{Yb}]\text{Cl}$ (red line) in
 3 ethanol : methanol (4:1) organic glass at 77K.



4
 5 **Figure 6.** Emission (integrated intensity from Figure S36) at 1000 nm ($\lambda_{\text{ex}} = 350 \text{ nm}$) of $[\text{L}_{\text{oo}}\text{Yb}]\text{Cl}$
 6 while isomerizing to $[\text{L}_{\text{cc}}\text{Yb}]\text{Cl}$ in ethanol : methanol (4:1) organic glass at 77 K. Two other
 7 successive measurements are performed after discoloration with white light (at RT).

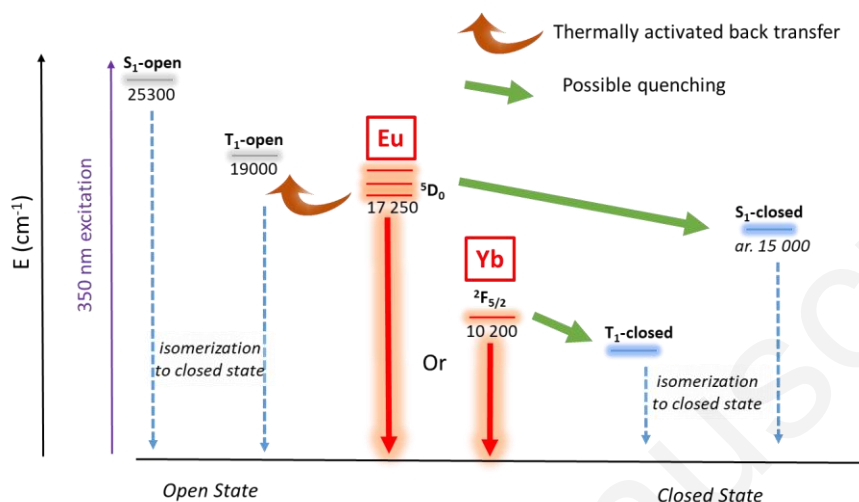
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 9

Compounds	Conditions	Ratio of Europium based luminescence (OFF state/ON state)	reference
Eu-DTE_o	RT, DCM	60 %	18
DAE-tpy-Eu(TTA) ₃	RT, CHCl ₃	20 %	14
Host guest complex of (terpyridinyldibenzo-24-crown-8)Eu(TTA) ₃ and DAE	RT, 1:1 CH ₃ CN/CHCl ₃	20 %	16
Triangle terthiazole complex Eu(hfac) ₃	RT, dioxane	36 %	10-11
	80 K, EtOH/MeOH glass matrix	ND ^a	10-11
Sulfone-DTE complex Eu(hfac) ₃	RT, ethylacetate	60 %	32
(5,6-dithienyl-1,10-phenanthroline)Eu(acac) ₃	RT, ethanol	30 %	5
(DTE-β-diketonate) ₃ Eu(phenanthroline)	RT, DCM	1 %	33
	RT, Film	5 %	
[L_{oo}Eu]Cl	77 K, ethanol : methanol (4:1)	4 – 8 %	This work
(DTE-β-diketonate) ₃ Yb(bpy-Ru-alkynyl)	RT, DCM	1.4 %	27
(DTE-β-diketonate) ₃ Yb(phenanthroline)	RT, DCM	2 %	33
	RT, film	3 %	
[L_{oo}Yb]Cl	77 K, ethanol : methanol (4:1)	13 %	This work

1 **Table 2.** Previously studied complexes showing reversible luminescence switching with DAE
2 photochromic units compared with this work. TTA =2-thenoyltrifluoroacetone, hfac = 1,1,1,5,5,5-
3 hexafluoroacetylacetone, aca = acetylacetone, tpy = terpyridine.

1 a. emission QY are 10 % for the open state and 7.6 % for the closed one and no photoreaction
2 occurs.

3



4

5 **Figure 7.** Tentative Jablonski diagram showing the different photophysical processes.

6

7 **Discussion.** Altogether, and in light with the objectives mentioned in the introduction, the results
8 of the photoluminescence experiments deserve a few comments. First, temperature/medium
9 dependence of the response is very spectacular for both systems and in both cases, no lanthanide
10 based emission can be detected at room temperature, in contrast to previously published systems
11 gathered in Table 2. In the case of europium, this is probably because of thermally activated back-
12 transfer, hence causing ligand-centered emission as suggested by the presence of the open form
13 ligand triplet state at 19000 cm^{-1} (see Figure 7). In the case of ytterbium, it is more likely that
14 luminescence is inherently weak due to efficient non-radiative processes and therefore difficult to
15 detect without causing the closing of the DTE. At 77 K in an organic glass, the non-radiative
16 processes are drastically slowed down as well as the closing reaction and both factors favor the
17 observation of ytterbium emission. Second, when measured in appropriate conditions, the contrast
18 between the responses of the two states for our europium complex is much higher than in previous
19 photoswitchable systems based on europium and diarylethene combinations (Table 2) and only one
20 example relying on N⁺C chelate four coordinate organoboron photoswitches of T type previously
21 showed better quenching ratio.¹⁵ Provided that back transfer and non-radiative processes are
22 reduced by further chemical engineering, our design with a macrocycle bearing two DTE units
23 could lead to very efficient RT europium luminescence switches, more suitable for applications.³³

1 Nonetheless, this design leads to the third example of efficient ytterbium luminescence photo-
2 control reported so far. In that case, the mechanism for emission quenching does not rely on spectral
3 overlap between the closed DTE and the lanthanide emission lines and we are currently
4 investigating the possibility of a low lying triplet state quenching the emission in the closed state
5 (Figure 7). We also postulate that the position of this state is not favorable to sensitization of
6 ytterbium emission through the visible transition of the closed DTE unit unlike in **Yb-DTEc**. This
7 leaves room for improvement of ligand design in order to obtain optimized positioning of this state
8 depending on the targeted behavior *ie* UV sensitization with quenching by a low lying state or
9 controllable visible light sensitization.

11 **Conclusion.**

12 With this work, we report the synthesis of an original ligand scaffold with two DTE units attached
13 to a cyclen based macrocycle designed for luminescence switching and the corresponding
14 complexes of yttrium(III), europium(III) and ytterbium(III). All three metal complexes show
15 reversible photochromism with high photo-conversions. Our design proved to be versatile and
16 adapted for both europium and ytterbium emission switching, when measured in frozen organic
17 glasses. The OFF/ON luminescence ratio are excellent in the case of europium compared to all
18 previously published compounds and still quite good in the case of ytterbium, that represents the
19 second example of such behavior. More important, our study, combined with on-going in depth
20 photo-physical studies, will contribute to the understanding of important factors for the design of
21 further improved molecular switches with custom switching, excitation and emission wavelengths.

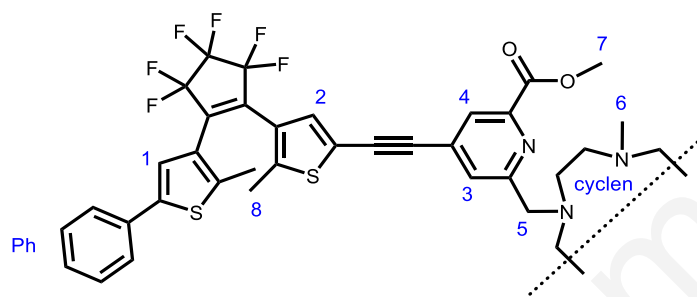
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24 A. S. thanks GDR MCM-2 for funding a short stay at the Laboratoire de Chimie in Lyon.

26 **Supporting Information Available:** Synthetic procedures for compounds **1** and **2**. NMR spectra
27 and HRMS of all metal complexes. Photo-isomerization studies (absorption and NMR
28 spectroscopy). Additional photoluminescence details.

30 **Experimental Section.**

1
2 **General comments:** Reagents were purchased from ACROS Organics and from ALDRICH
3 Chemical Co. Cyclen was purchased from Chematech (Dijon, France). Dimethyl-Cyclen was
4 synthesized as previously described.²⁶ The solvents were freshly distilled prior to use and according
5 to the standard methods. NMR spectra (¹H, ¹⁹F and ¹³C) were recorded at the core facilities of the
6 University of Brest, with Bruker Advanced 500 (500 MHz) or Bruker AMX-3 300 (300 MHz)
7 spectrometers or at the Centre Régional de Mesures Physiques de l'Ouest on a Bruker Avance III
8 HD 500 (500 MHz). The HR-MS analyses were performed at the Institute of Analytic and Organic
9 Chemistry, ICOA in Orléans. Elemental Analysis were performed at Centre Régional de Mesures
10 Physiques de l'Ouest. All experiments were performed in the dark.

11



12

13

14 Scheme 3. Numbering used for NMR assignment.

15

16 **Compound 3.** A solution of dimethyl-cyclen (60 mg, 0.3 mmol) and K₂CO₃ (165 mg, 1.2 mmol,
17 4 eq) in CH₃CN (7.5 mL) was stirred at room temperature for 30 min. To this solution was added
18 dropwise a solution of compound 2 (447 mg, 0.63 mmol, 2.1 eq) in CH₃CN (11.4 mL). The reaction
19 mixture was stirred at 60 °C for 2.5 days before evaporation of solvents to dryness. The residue
20 was taken up in CHCl₃ and the residual salts were filtered off. The filtrate was evaporated to
21 dryness to give a brown oil which was dissolved in the minimum amount of ethyl acetate. A large
22 amount of hexane was added and the formation of a dark precipitate was observed. The precipitate
23 was filtered and the filtrate was evaporated and dried under vacuum to give compound 3 (410 mg,
24 0.29 mmol, 95 %) as an orange oil. R_f (CH₂Cl₂/MeOH, neutral alumina) = 0.25. ¹H NMR (400
25 MHz, CDCl₃) δ 8.48 (s, 2H, H₄), 7.96 (s, 2H, H₃), 7.53 (d, *J* = 7.6 Hz, 5H), 7.38 (t, *J* = 7.6 Hz,
26 5H), 7.30 (d, *J* = 7.2 Hz, 2H), 7.26 (s, 2H), 3.95 (s, 6H, H₇), 3.87 (s, 4H, H₅), 2.66 (br s, 8H, CH₂
27 cyclen), 2.63 (brs, 8H, CH₂ cyclen), 2.13 (s, 6H, 2* H₆), 1.92 (s, 6H, 2* H₈), 1.90 (s, 6H, 2* H₈).

1 ^{19}F NMR (282 MHz, CDCl_3) δ -107.52, -107.69, -129.29. ^{13}C NMR (125 MHz, CDCl_3) δ 165.4
2 (C=O), 162.3, 147.2, 144.9, 142.6, 141.3 (Cq), 137.1 (t, $J = 23$ Hz, $\underline{\text{C}}\text{CF}_2$), 134.9 (t, $J = 23$ Hz, $\underline{\text{C}}\text{q}$ -
3 CF_2), 133.2 (Cq), 132.8, 132.2 (Cq), 129.1, 128.1, 127.9, 125.6 (CH), 125.5, 125.4 (Cq), 124.7,
4 122.1 (CH), 120.2 (Cq), 116.0 (tt, $^1J_{\text{C-F}} = 255$ Hz, $^2J_{\text{C-F}} = 23$ Hz, external CF_2), 110.9 (tquint, $^1J_{\text{C-F}}$
5 $= 270$ Hz, $^2J_{\text{C-F}} = 25$ Hz, central CF_2), 91.7, 86.5 (alkyne), 61.0 (CH_2 -pico), 56.1, 53.9 (CH_2 cyclen),
6 52.9 (COOCH_3), 44.3 (CH_3 cyclen), 14.54, 14.49 (C_8). ESI-HR-MS (positive, MeOH) m/z calcd.
7 for $[\text{C}_{72}\text{H}_{63}\text{F}_{12}\text{N}_6\text{O}_4\text{S}_4]^+$: 1431.3597, found: 1431.3598, $[\text{M}+\text{H}]^+$; calcd. for $[\text{C}_{72}\text{H}_{62}\text{F}_{12}\text{N}_6\text{NaO}_4\text{S}_4]^+$:
8 1453.3416, found: 1453.3420, $[\text{M}+\text{Na}]^+$; calcd. for $[\text{C}_{72}\text{H}_{64}\text{F}_{12}\text{N}_6\text{O}_4\text{S}_4]^{2+}$: 716.1835, found:
9 716.1842, $[\text{M}+2\text{H}]^{2+}$.

10 **L_{oo}**. A solution of compound **3** (100 mg, 69.8 μmol) and KOH (1M, 4.37mL) in THF (8.73 mL)
11 was stirred at room temperature for 26 h. Phases were separated and the aqueous phase was
12 extracted with CH_2Cl_2 (2×10 mL). The combined organic phases were dried over MgSO_4 and
13 solvents were evaporated to dryness. The residue was dissolved in the minimum of ethyl acetate
14 and a large amount of hexane was added. The precipitate was filtered and taken up in CHCl_3 to
15 give **L_{oo}** (70 mg, 47.3 μmol , 68 %) as an orange oil. ^1H NMR (500 MHz, CDCl_3) δ 7.84 (s, 2H),
16 7.52 (d, $J = 7.5$ Hz, 4H), 7.37 (t, $J = 7.5$ Hz, 4H), 7.33 (s, 2H), 7.31 – 7.23 (m, 4H), 7.09 (s, 2H),
17 4.12 - 2.16 (H of the cyclen), 1.94 (s, 6H, 2* H_8), 1.92 (s, 6H, 2* H_8), 1.81 (s, 6H, 2* H_6). ^{13}C NMR
18 (125 MHz, CDCl_3) δ 170.3, 157.3, 144.4, 142.6, 141.4 (Cq), 137.0 (t, $J = 23$ Hz, $\underline{\text{C}}\text{CF}_2$), 135.2 (t,
19 $J = 23$ Hz, $\underline{\text{C}}\text{CF}_2$), 133.2 (Cq), 132.5 (CH), 131.1 (Cq), 129.1, 128.0, 125.6 (CH), 125.5, 125.3
20 (Cq), 125.1, 124.5, 122.2 (CH), 120.9 (Cq), 116.0 (tt, $^1J_{\text{C-F}} = 255$ Hz, $^2J_{\text{C-F}} = 25$ Hz, external CF_2),
21 110.9 (tquint, $^1J_{\text{C-F}} = 270$ Hz, $^2J_{\text{C-F}} = 25$ Hz, central CF_2), 92.2, 85.3 (alkyne), 62.1 (CH_2 pico), 51.1
22 (CH_2 cyclen), 43.2 (CH_3 cyclen), 14.7 (C_8), 14.5 (C_8). ESI-HR-MS (positive, MeOH) m/z calcd.
23 for $[\text{C}_{70}\text{H}_{59}\text{F}_{12}\text{N}_6\text{O}_4\text{S}_4]^+$: 1403.3284, found: 1403.3296, $[\text{M}+\text{H}]^+$; calcd. for $[\text{C}_{70}\text{H}_{60}\text{F}_{12}\text{N}_6\text{O}_4\text{S}_4]^{2+}$:
24 702.1678, found: 702.1688, $[\text{M}+2\text{H}]^{2+}$ with M referring to the conjugated diacid of **L_{oo}**.

25 **[YL_{oo}]Cl**. To a solution of compound **L_{oo}** (60 mg, 41 μmol) in MeOH (HPLC grad, 10 mL) was
26 added $\text{YCl}_3 \cdot 6\text{H}_2\text{O}$ (37 mg, 122 μmol , 3 eq). The pH was controlled at 7 and the reaction mixture
27 was stirred at room temperature for 3.5 days. Solvents were evaporated to dryness and water was
28 added to the residue. Water was then filtered on cotton and the solid kept on the cotton was
29 dissolved with CH_3CN (HPLC grad). CH_3CN was evaporated to dryness and the residue was
30 dissolved in the minimum of CH_2Cl_2 . A large amount of hexane was added and the precipitated

1 was filtered, washed with hexane and dried under vacuum to yield [**YL_{oo}**]Cl (38 mg, 25 μmol,
2 61%) as a pale yellow solid. ¹H NMR (500 MHz, CDCl₃, 253K) δ : 8.10 (s, 2H), 7.61 (s, 2H), 7.55
3 (d, *J* = 7.5 Hz, 4H), 7.46 (s, 2H), 7.40 (t, *J* = 7.2 Hz, 4H), 7.32 (d, *J* = 7.7 Hz, 2H), 7.29 (s, 2H),
4 4.68 (d, *J* = 13.3 Hz, 4H), 4.49 (br s, 2H), 4.05 (d, *J* = 15.2 Hz, 2H), 3.17 (br s, 2H), 3.05 (d, *J* =
5 10.7 Hz, 2H), 2.89 (d, *J* = 11.8 Hz, 2H), 2.80 (s, 2H), 2.70 (d, *J* = 10.5 Hz, 2H), 2.24 – 2.12 (m,
6 2H), 2.04 (s, 6H, 2* H₆), 1.92 (s, 6H, 2* H₈), 1.89 (s, 6H, 2* H₈). ¹³C NMR (125 MHz, CDCl₃) δ
7 168.2 (C=O), 152.6, 146.5, 142.7, 141.3 (Cq), 137.6 (m, Cq-CF₂), 137.3 (Cq), 134.8 (m, Cq-CF₂),
8 134.4 (C2), 133.1 (Cq), 129.0, 128.1, 125.9 (Ar), 125.8 (Cq), 125.6 (Ar), 125.3 (Cq), 122.0 (C1),
9 119.1 (Cq), 115.9 (tt, ¹J_{C-F} = 255 Hz, ²J_{C-F} = 25 Hz, external CF₂), 110.9 (tquint, ¹J_{C-F} = 270 Hz,
10 ²J_{C-F} = 25 Hz, central CF₂), 91.3, 90.2 (Cq alkyne), 54.2 (CH₂ cyclen), 46.1 (CH₃ cyclen), 14.64
11 (C₈), 14.58 (C₈). ¹⁹F NMR (376 MHz, CDCl₃) δ -110.55, -110.78, -132.34. ⁸⁹Y NMR (HMBC
12 ¹H/⁸⁹Y, CDCl₃) δ 250.9. ESI-HR-MS (positive, MeOH) *m/z* calcd. for [C₇₀H₅₆F₁₂N₆O₄S₄Y]⁺:
13 1489.2107, found: 1489.2114, [M+H]⁺; calcd. for [C₇₀H₅₇F₁₂N₆O₄S₄Y]²⁺: 745.1090, found:
14 745.1102, [M+2H]²⁺.

15 [**YbL_{oo}**]Cl. Compound **L_{oo}** (60 mg, 0.041 mmol) was dissolved in 13 mL of methanol and
16 YbCl₃·6H₂O (24 mg, 0.061 mmol) was then added. After stirring at room temperature for 3.5 days,
17 the solvent was evaporated under vacuum. Water was added and the resulting suspension was
18 filtered. The solid was then extracted with acetonitrile. After evaporation of the solvents, the crude
19 solid was dissolved in the minimum amount of dichloromethane and precipitated in hexane
20 yielding [**YbL_{oo}**]Cl as a white bluish solid (51 mg, 80 %). ¹H NMR: (500 MHz, CD₂Cl₂) δ 83.88
21 (2H, H₃/H₄), 55.60 (2H, H₃/H₄), 27.35 (2H, H_{cyclen}), 20.29 (2H, H_{cyclen}), 15.43 (2H, H_{cyclen}), 10.47
22 (2H, H_{cyclen}), 7.35-7.22 (10H, H_{ph}), 6.97 (2H, H₂), 6.37(2H, H₁), 1.60-1.50 (12H, CH₃), -2.53 (2H,
23 H_{cyclen}), -7.12 (2H, H_{cyclen}), -15.92 (2H, H_{cyclen}), -30.94 (6H, H₆), -40.64(2H, H_{cyclen}). EA C 49.07;
24 H 3.78; S 6.52; N 4.41, Calcd for C₇₀H₅₆ClF₁₂N₆O₄S₄Yb·5H₂O C 49.46; H 3.91; S 7.54; N 4.94
25 ESI-HR-MS (positive, MeOH) *m/z* calcd. for [C₇₀H₅₆YbF₁₂N₆O₄S₄]⁺: 1574.2456, found:
26 1574.2437, [M+H]⁺; calcd. for [C₇₀H₅₇YbF₁₂N₆O₄S₄]²⁺: 787.6277, found: 787.6295, [M+2H]²⁺.

27 [**YbL_{cc}**]Cl was obtained upon UV irradiation (see SI). ¹H NMR (500 MHz, CD₂Cl₂) δ 84.99(2H,
28 H₃/H₄), 53.37 (2H, H₃/H₄), 27.44 (2H, H_{cyclen}), 21.74 (2H, H_{cyclen}), 14.92 (2H, H_{cyclen}), 11.24-9.50
29 (2H, H_{cyclen}), 7.42-7.33 (10H, H_{ph}), 6.51 (2H, H₂), 5.65 (2H, H₁), 1.76-1.70 (12H, CH₃), -2.24 (2H,
30 H_{cyclen}), -6.44 (2H, H_{cyclen}), -16.34(2H, H_{cyclen}), -32.25(6H, H₆), -38.40(2H, H_{cyclen}).

1 **[EuL_{oo}]Cl**. Compound **L_{oo}** (46 mg, 0.031 mmol) was dissolved in 10 mL of methanol (HPLC grad)
2 and EuCl₃·6H₂O (28 mg, 0.076 mmol) was then added. After stirring at room temperature for 3.5
3 days, the solvent was evaporated under vacuum. Water was added and the resulting suspension was
4 filtered. The solid was then extracted with acetonitrile (HPLC grad). After evaporation of the
5 solvents, the crude solid was dissolved in the minimum amount of dichloromethane and
6 precipitated in hexane yielding **[EuL_{oo}]Cl** as a pale yellow solid (44 mg, 90 %). ¹H NMR (500
7 MHz, CD₂Cl₂) δ 38.43(2H, H₃/H₄), 25.75(2H, H₃/H₄), 7.49 (d, 4H, H_{ph}), 7.36 (t, 4H, H_{ph}), 7.28 (t,
8 4H, H_{ph}), 7.20 (2H, H₂), 7.04 (2H, H₁), 2.59-2.34 (10H, H₅ + 6H_{cyclen}) 1.99 (6H, 2* H₈), 1.85 (6H,
9 2* H₈), -3.09 (2H, H_{cyclen}), -7.56 (2H, H_{cyclen}), -9.13 (6H, H₆), -10.28 (2H, H_{cyclen}), -13.52 (2H,
10 H_{cyclen}), -15.45 (2H, H_{cyclen}). EA C 50.94; H 3.81; S 7.39; N 4.76, Calcd for C₇₀H₅₆ClF₁₂N₆O₄S₄Eu·4H₂O
11 C 50.62; H 3.88; S 7.72; N 5.06. ESI-HR-MS (positive, MeOH) *m/z* calcd. for [C₇₀H₅₆EuF₁₂N₆O₄S₄]⁺:
12 1553.2261, found: 1553.2264, [M+H]⁺; calcd. for [C₇₀H₅₇EuF₁₂N₆O₄S₄]²⁺: 777.1167, found:
13 777.1178, [M+2H]²⁺.

14 **[EuL_{cc}]Cl** was obtained upon UV irradiation (see SI), ¹H NMR (500 MHz, CD₂Cl₂) δ 38.68 (2H,
15 H₃/H₄), 24.96 (2H, H₃/H₄), 7.55-7.43 (10H, H_{ph}), 6.70 (2H, H₂), 6.20 (2H, H₁), 2.89 (10H, H₅ +
16 6H_{cyclen}), 2.11-2.09 (12H, CH₃), -3.52 (2H, H_{cyclen}), -7.91 (2H, H_{cyclen}), -9.50 (8H, 6H₆ and 2H_{cyclen}),
17 -13.79 (2H, H_{cyclen}), -15.11 (2H, H_{cyclen}).

1 References

- 2 (1) Andreasson, J.; Pischel, U., Smart molecules at work-mimicking advanced logic operations. *Chem.*
3 *Soc. Rev.* **2010**, *39* (1), 174-188.
- 4 (2) Browne, W. R.; Feringa, B. L., Light Switching of Molecules on Surfaces. In *Annu. Rev. Phys. Chem.*,
5 2009; Vol. 60, pp 407-428.
- 6 (3) Heffern, M. C.; Matosziuk, L. M.; Meade, T. J., Lanthanide Probes for Bioresponsive Imaging. *Chem.*
7 *Rev.* **2014**, *114* (8), 4496-4539.
- 8 (4) Tian, Z. Y.; Li, A. D. Q., Photoswitching-Enabled Novel Optical Imaging: Innovative Solutions for Real-
9 World Challenges in Fluorescence Detections. *Acc. Chem. Res.* **2013**, *46* (2), 269-279.
- 10 (5) Mei, J. F.; Lv, Z. P.; Lai, J. C.; Jia, X. Y.; Li, C. H.; Zuo, J. L.; You, X. Z., A novel photo-responsive
11 europium(III) complex for advanced anti-counterfeiting and encryption. *Dalton Trans.* **2016**, *45* (13),
12 5451-5454.
- 13 (6) Qi, Q.; Li, C.; Liu, X.; Jiang, S.; Xu, Z.; Lee, R.; Zhu, M.; Xu, B.; Tian, W., Solid-State Photoinduced
14 Luminescence Switch for Advanced Anticounterfeiting and Super-Resolution Imaging Applications. *J. Am.*
15 *Chem. Soc.* **2017**.
- 16 (7) Zhang, J. J.; Zou, Q.; Tian, H., Photochromic Materials: More Than Meets The Eye. *Adv. Mater.* **2013**,
17 *25* (3), 378-399.
- 18 (8) Zhou, W.; Chen, Y.; Yu, Q.; Li, P.; Chen, X.; Liu, Y., Photo-responsive cyclodextrin/anthracene/Eu³⁺-
19 supramolecular assembly for a tunable photochromic multicolor cell label and fluorescent ink. *Chem. Sci.*
20 **2019**, *10* (11), 3346-3352.
- 21 (9) Yuasa, J.; Nakagawa, T.; Kita, Y.; Kaito, A.; Kawai, T., Photoactivatable europium luminescence turn-
22 on by photo-oxygenation of β -diketone having pyrrole rings. *Chem. Commun.* **2017**, *53* (50), 6748-6751.
- 23 (10) Nakagawa, T.; Hasegawa, Y.; Kawai, T., Photoresponsive europium(III) complex based on
24 photochromic reaction. *J. Phys. Chem. A* **2008**, *112* (23), 5096-5103.
- 25 (11) Nakagawa, T.; Atsumi, K.; Nakashima, T.; Hasegawa, Y.; Kawai, T., Reversible luminescence
26 modulation in photochromic europium(III) complex having triangle terthiazole ligands. *Chem. Lett.* **2007**,
27 *36* (3), 372-373.
- 28 (12) Koji, M.; Yoshio, N.; Keiichi, K., Photochemical Modulation of Europium Ion Fluorescence Using a
29 Tetraazamacrocyclic Derivative Bearing a Spirobenzopyran and Three Carboxymethyl Moieties. *Bull.*
30 *Chem. Soc. Jpn.* **2009**, *82* (4), 472-474.
- 31 (13) Hasegawa, Y.; Nakagawa, T.; Kawai, T., Recent progress of luminescent metal complexes with
32 photochromic units. *Coord. Chem. Rev.* **2010**, *254* (21-22), 2643-2651.
- 33 (14) Cheng, H. B.; Hu, G. F.; Zhang, Z. H.; Gao, L.; Gao, X.; Wu, H. C., Photocontrolled Reversible
34 Luminescent Lanthanide Molecular Switch Based on a Diarylethene-Europium Dyad. *Inorg. Chem.* **2016**,
35 *55* (16), 7962-8.
- 36 (15) Wang, N.; Wang, J.; Zhao, D.; Møllerup, S. K.; Peng, T.; Wang, H.; Wang, S., Lanthanide Complexes
37 with Photochromic Organoboron Ligand: Synthesis and Luminescence Study. *Inorg. Chem.* **2018**, *57* (16),
38 10040-10049.
- 39 (16) Cheng, H.-B.; Zhang, H.-Y.; Liu, Y., Dual-Stimulus Luminescent Lanthanide Molecular Switch Based
40 on an Unsymmetrical Diarylperfluorocyclopentene. *J. Am. Chem. Soc.* **2013**, *135* (28), 10190-10193.
- 41 (17) Irie, M.; Fulciniti, T.; Matsuda, K.; Kobatake, S., Photochromism of Diarylethene Molecules and
42 Crystals: Memories, Switches, and Actuators. *Chem. Rev.* **2014**, *114* (24), 12174-12277.
- 43 (18) He, X.; Norel, L.; Hervault, Y.-M.; Métivier, R.; D'Aléo, A.; Maury, O.; Rigaut, S., Modulation of Eu(III)
44 and Yb(III) Luminescence Using a DTE Photochromic Ligand. *Inorg. Chem.* **2016**, *55* (24), 12635-12643.
- 45 (19) Bui, A. T.; Beyler, M.; Liao, Y.-Y.; Grichine, A.; Duperray, A.; Mulatier, J.-C.; Guennic, B. L.; Andraud,
46 C.; Maury, O.; Tripier, R., Cationic Two-Photon Lanthanide Bioprobes Able to Accumulate in Live Cells.
47 *Inorg. Chem.* **2016**, *55* (14), 7020-7025.

- 1 (20) Bui, A. T.; Beyler, M.; Grichine, A.; Duperray, A.; Mulatier, J.-C.; Guyot, Y.; Andraud, C.; Tripier, R.;
2 Brasselet, S.; Maury, O., Near infrared two photon imaging using a bright cationic Yb(III) bioprobes
3 spontaneously internalized into live cells. *Chem. Commun.* **2017**, *53* (44), 6005-6008.
- 4 (21) Stasiuk, G. J.; Long, N. J., The ubiquitous DOTA and its derivatives: the impact of 1,4,7,10-
5 tetraazacyclododecane-1,4,7,10-tetraacetic acid on biomedical imaging. *Chem. Commun.* **2013**, *49* (27),
6 2732-2746.
- 7 (22) Clough, T. J.; Jiang, L.; Wong, K.-L.; Long, N. J., Ligand design strategies to increase stability of
8 gadolinium-based magnetic resonance imaging contrast agents. *Nat. Commun.* **2019**, *10* (1), 1420.
- 9 (23) Moore, E. G.; Samuel, A. P. S.; Raymond, K. N., From Antenna to Assay: Lessons Learned in
10 Lanthanide Luminescence. *Acc. Chem. Res.* **2009**, *42* (4), 542-552.
- 11 (24) Shuvaev, S.; Starck, M.; Parker, D., Responsive, Water-Soluble Europium(III) Luminescent Probes.
12 *Chem. Eur. J.* **2017**, *23* (42), 9974-9989.
- 13 (25) Rodríguez-Rodríguez, A.; Garda, Z.; Ruscsák, E.; Esteban-Gómez, D.; de Blas, A.; Rodríguez-Blas, T.;
14 Lima, L. M. P.; Beyler, M.; Tripier, R.; Tircsó, G.; Platas-Iglesias, C., Stable Mn²⁺, Cu²⁺ and Ln³⁺
15 complexes with cyclen-based ligands functionalized with picolinate pendant arms. *Dalton Trans.* **2015**, *44*
16 (11), 5017-5031.
- 17 (26) Rodríguez-Rodríguez, A.; Esteban-Gómez, D.; de Blas, A.; Rodríguez-Blas, T.; Fekete, M.; Botta, M.;
18 Tripier, R.; Platas-Iglesias, C., Lanthanide(III) Complexes with Ligands Derived from a Cyclen Framework
19 Containing Pyridinecarboxylate Pendants. The Effect of Steric Hindrance on the Hydration Number.
20 *Inorg. Chem.* **2012**, *51* (4), 2509-2521.
- 21 (27) Al Sabea, H.; Norel, L.; Galangau, O.; Hijazi, H.; Metivier, R.; Roisnel, T.; Maury, O.; Bucher, C.; Riobe,
22 F.; Rigaut, S., Dual Light and Redox Control of NIR Luminescence with Complementary Photochromic and
23 Organometallic Antennae. *J. Am. Chem. Soc.* **2019**, *141* (51), 20026-20030.
- 24 (28) Bourdolle, A.; Allali, M.; Mulatier, J.-C.; Le Guennic, B.; Zwier, J. M.; Baldeck, P. L.; Buezli, J.-C. G.;
25 Andraud, C.; Lamarque, L.; Maury, O., Modulating the Photophysical Properties of Azamacrocyclic
26 Europium Complexes with Charge-Transfer Antenna Chromophores. *Inorg. Chem.* **2011**, *50* (11), 4987-
27 4999.
- 28 (29) Xing, Y.; Jindal, A. K.; Regueiro-Figueroa, M.; Le Fur, M.; Kervarec, N.; Zhao, P.; Kovacs, Z.; Valencia,
29 L.; Pérez-Lourido, P.; Tripier, R.; Esteban-Gómez, D.; Platas-Iglesias, C.; Sherry, A. D., The Relationship
30 between NMR Chemical Shifts of Thermally Polarized and Hyperpolarized 89Y Complexes and Their
31 Solution Structures. *Chem. Eur. J.* **2016**, *22* (46), 16657-16667.
- 32 (30) Pintacuda, G.; John, M.; Su, X.-C.; Otting, G., NMR structure determination of protein-ligand
33 complexes by lanthanide labeling. *Acc. Chem. Res.* **2007**, *40* (3), 206-212.
- 34 (31) Fihey, A.; Perrier, A.; Browne, W. R.; Jacquemin, D., Multiphotochromic molecular systems. *Chem.*
35 *Soc. Rev.* **2015**, *44* (11), 3719-3759.
- 36 (32) Nakagawa, T.; Hasegawa, Y.; Kawai, T., Nondestructive luminescence intensity readout of a
37 photochromic lanthanide(III) complex. *Chem. Commun.* **2009**, (37), 5630-5632.
- 38 (33) Al Sabea, H.; Norel, L.; Galangau, O.; Roisnel, T.; Maury, O.; Riobé, F.; Rigaut, S., Efficient photo-
39 modulation of visible Eu(III) and invisible Yb(III) luminescences using DTE photochromic ligands for
40 optical encryption *Adv. Funct. Mater.* **2020**, 2002943

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