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# Two new red-emitting ternary europium(III) complexes with high photoluminescence quantum yields and exceptional performance in OLED devices†

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Two new organo-europium complexes (OEuCs) [Eu(btfa)<sub>3</sub>(Bathphen)] (**OEuC-1**) and [Eu(tta)<sub>3</sub>(Bathphen)] (**OEuC-2**) where btfa and tta are the anions of 4,4,4-trifluoro-1-phenyl-1,3-butanedione and 2-thenoyl-trifluoroacetone while Bathphen = Bathphenanthroline have been synthesized and characterized. Both complexes in the solid state exhibit strong red emissions with photoluminescence quantum yields (PLQYs) of 80% ± 10%. These complexes were tested as dopants in the emitting layer (EML) to fabricate red organic light emitting diodes (R-OLEDs). Through device engineering involving the amalgamation of appropriate host materials and complexes, we have achieved exceptional overall R-OLED performance for an **OEuC-2** based device, with maximum current efficiency (CE<sub>max</sub>) = 9.91 cd A<sup>-1</sup>, maximum power efficiency (PE<sub>max</sub>) = 9.15 lm W<sup>-1</sup>, maximum external quantum efficiency (EQE<sub>max</sub>) = 6.24%, brightness (B) of 545 cd m<sup>-2</sup>, and (CIE)<sub>x,y</sub> = 0.620, 0.323 at J = 10 mA cm<sup>-2</sup>.

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## 1 Introduction

Luminescent organo-lanthanide complexes (OLnCs), especially trivalent OEuCs, have been extensively applied in luminescence thermometry,<sup>1</sup> sensors<sup>2</sup> and OLED devices,<sup>3</sup> because of their innate optical properties.<sup>4</sup> In pursuit of OLED efficiency, it is vital that the complexes exhibit excellent photoluminescence quantum yields (PLQYs) *i.e.* >50% along with good electron-transport properties. A wide variety of bidentate chelating ligands have been tailored to develop OEuCs for this purpose.<sup>5</sup> Among them, mono-ionic β-diketone bidentate ligands in combination with neutral N<sup>∧</sup>N ligands dominate. A

search of the literature indicates that the PLQY of eight-coordinate [Eu(β-diket.)<sub>3</sub>(N<sup>∧</sup>N)] with a EuO<sub>6</sub>N<sub>2</sub> coordination sphere rarely achieves 70% target (Table S1 and Chart S1, ESI†). This is because Eu(III) luminescence in OEuCs can be quenched readily by high-energy X-H oscillators<sup>6</sup> present in the organic ligands. Two complexes that do show a greater than 70% PLQY are deuterated hexafluoroacetylacetone (hfaa) systems bearing derivatives of phosphine oxide (O<sup>∧</sup>O) as neutral ancillary ligands, *i.e.*, [Eu(hfa-D)<sub>3</sub>(TPPO)<sub>2</sub>] (PLQY = 90%)<sup>7</sup> and [Eu(hfa-D)<sub>3</sub>(BIPHEPO)] (PLQY = 87%)<sup>7</sup> (Table S2 and Chart S2, ESI†) with a EuO<sub>6</sub>O<sub>2</sub> coordination sphere [TPPO = triphenylphosphine oxide and BIPHEPO = 1,1'-biphenyl-2,2'-diyl bis(diphenylphosphineoxide)]. However, despite the very high PLQY, the potential application of these two complexes as EMLs in OLEDs has not been explored. Moreover, as observed by Kalyakina *et al.*,<sup>8</sup> despite having record PLQYs, ternary europium complexes of carboxylates bearing neutral phenanthroline (Phen) resulted in inferior electroluminescence (EL), due to the poor electron mobility of the Phen ligand. Even though the role of neutral ancillary ligand(s) in sensitizing the PL is almost two times lower (~35–25%) than that of primary anionic antenna ligand(s) (~65–75%),<sup>9</sup> their presence improves the physicochemical properties of OEuCs such as thermal stability, volatility, *etc.*<sup>10</sup> More importantly, they enhance the optoelectronic performance by serving as the main channel for charge and exciton harvesting.

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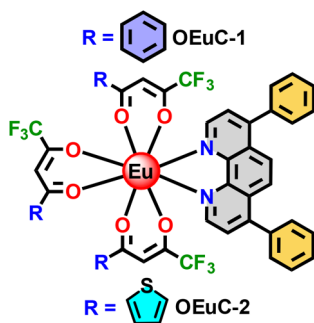


Chart 1 The molecular structure of the novel OEuCs.

In order to improve the EL performance of relatively cheap and easily accessible red-emitting OEuCs [crystal abundance 0.3 ppm]<sup>11</sup> that can be compared to better known but expensive red-emitting iridium (Ir)-based complexes [crystal abundance 0.000037 ppm],<sup>11</sup> we embarked on the design and development of two new luminescent OEuCs (Chart 1) with a  $\text{EuO}_6\text{N}_2$  coordination sphere. Compared to the preparation of many red-emitting Ir(III)-complexes which require specialized synthetic modifications and complicated purification processes,<sup>12</sup> Eu(III)-based complexes are relatively easy to prepare and exhibit highly efficient monochromatic red emissions (CIE  $\approx$  0.67, 0.33).<sup>13</sup> These Eu(III) complexes have a few basic prerequisites: strong light absorption properties of the ligand (s) and appropriate energy differences ( $\Delta E$ )<sup>14</sup> between the  $^3\pi\pi^*$  (triplet) and  $^1\pi\pi^*$  (singlet) states. We now report the synthesis, characterization, and photophysical and electrophysical properties of two strongly luminescent OEuCs (Chart 1) with a  $\text{EuO}_6\text{N}_2$  coordination sphere. This is achieved by employing readily available primary antenna ligands, Hbtfa and Htta ( $^3\pi\pi^* = 21\,400\text{ cm}^{-1}$  for btfa and  $20\,300\text{ cm}^{-1}$  for tta)<sup>15</sup> bearing an electron-donating phenyl/thienyl group at one terminus and an electron-withdrawing trifluoromethyl ( $-\text{CF}_3$ ) group at the other terminus, which also play a pivotal role in minimizing the vibrational quenching.<sup>6</sup> The rigid neutral Bathphen ligand with an extended  $\pi$ -system ( $^3\pi\pi^* = 21\,000\text{ cm}^{-1}$ )<sup>16</sup> is employed as the ancillary ligand. It has good electron mobility ( $5 \times 10^{-4}\text{ cm}^2\text{ V}^{-1}\text{ s}^{-1}$ )<sup>17</sup> and has been widely used as a hole or exciton-blocking layer.

## 2 Experimental

### 2.1 Chemical reagents and general instrumentation

All chemicals used in the synthesis and OLED fabrication were purchased from commercial sources and were used without further purification unless otherwise specified. Solvents used in the experiments were dried and distilled prior to use. Elemental analyses of OEuC-1 and OEuC-2 were performed on a Euro EA-CHN elemental analyser. The Fourier transform infrared (FT-IR) spectra of the solid complexes were obtained using a Cary 630 FT-IR spectrometer in the attenuated total reflectance (ATR) mode. Mass spectra were obtained using an

LCMS-8040, Shimadzu-Japan coupled to a triple quadrupole tandem mass spectrometer equipped with electrospray ionization (ESI). The thermal stability of the complexes was determined by thermogravimetric analysis (TGA) and differential thermogravimetric analysis (DTA) in the temperature range between 50 and 700 °C under a dinitrogen ( $\text{N}_2$ ) atmosphere and recorded on a TA Instruments model SDT Q600.

### 2.2 Synthesis

**[Eu(btfa)<sub>3</sub>(Bathphen)] (OEuC-1).** OEuC-1 was synthesized by reacting equimolar quantities of  $[\text{Eu}(\text{btfa})_3(\text{H}_2\text{O})_2]$ <sup>13</sup> (0.300 g, 0.360 mmol) and Bathphen (0.119 g, 0.360 mmol) in methanol (MeOH, 20 mL). The reaction mixture was stirred overnight at room temperature and left for slow solvent evaporation. The solid formed was washed with cold EtOH ( $5 \times 2\text{ mL}$ ) and toluene ( $5 \times 2\text{ mL}$ ). Yield: 80%. Microanalysis calculated for  $\text{C}_{54}\text{H}_{34}\text{EuF}_9\text{N}_2\text{O}_6$ , C, 57.41; H, 3.03; N, 2.48; observed C, 57.45; H, 3.01; N, 2.49%. FT-IR (solid,  $\text{cm}^{-1}$ ):  $\nu(\text{ar C-H st})$  3058  $\text{cm}^{-1}$ ;  $\nu(\text{C=O st})$  1609  $\text{cm}^{-1}$ ;  $\nu(\text{C=N st})$  1576  $\text{cm}^{-1}$ ;  $\nu(\text{C=C st})$  1519  $\text{cm}^{-1}$ ;  $\nu(\text{C-F st, CF}_3)$  1388, 1317  $\text{cm}^{-1}$ ; out-of plane asymmetric  $\nu(\text{C-F st})$  1180  $\text{cm}^{-1}$ ; in-plane  $\nu(\text{C-H bend})$  1129  $\text{cm}^{-1}$  (Fig. S1, ESI<sup>†</sup>); MS (positive mode):  $m/z$ : 1130.9  $[\text{M} + \text{H}]^+$ ; 1186.8  $[\text{M} + \text{Na} + \text{CH}_3\text{OH} + \text{H}]^+$  (Fig. S2, ESI<sup>†</sup>); decomposition temperature ( $T_d$ ) with 5% weight loss = 315 °C (Fig. S3, ESI<sup>†</sup>).

**[Eu(tta)<sub>3</sub>(Bathphen)] (OEuC-2).** OEuC-2 was synthesized by a similar method with  $[\text{Eu}(\text{tta})_3(\text{H}_2\text{O})_2]$ <sup>9b</sup> (0.300 g, 0.305 mmol) and Bathphen (0.101 g, 0.305 mmol) in methanol (MeOH; 20 mL). Yield: 75%. Microanalysis calculated for  $\text{C}_{48}\text{H}_{28}\text{EuF}_9\text{N}_2\text{O}_6\text{S}_3$ , C, 50.23; H, 2.46; N, 2.44; observed C, 50.29; H, 2.41; N, 2.41%. FTIR (solid,  $\text{cm}^{-1}$ )  $\nu(\text{ar C-H st})$  3096  $\text{cm}^{-1}$ ;  $\nu(\text{C=O st})$  1625  $\text{cm}^{-1}$ ;  $\nu(\text{C=N st})$  1594  $\text{cm}^{-1}$ ;  $\nu(\text{C=C st})$  1507  $\text{cm}^{-1}$ ;  $\nu(\text{C-F st, CF}_3)$  1353, 1302  $\text{cm}^{-1}$ ; out-of plane asymmetric  $\nu(\text{C-F st})$  1181  $\text{cm}^{-1}$ ; in-plane  $\nu(\text{C-H bend})$  1127  $\text{cm}^{-1}$  (Fig. S4, ESI<sup>†</sup>); MS (positive mode):  $m/z$ : 1200.9  $[\text{M} + \text{Na} + \text{CH}_3\text{OH} - \text{H}]^+$  (Fig. S5, ESI<sup>†</sup>);  $T_d$  with 5% weight loss = 330 °C (Fig. S6, ESI<sup>†</sup>).

### 2.3 Spectroscopic measurements, determination of photophysical parameters and the OLED fabrication process

Spectroscopic measurements of the complexes, which include UV-visible absorption, excitation, emission spectra, decay profiles and absolute PLQY values were obtained at room temperature; details of the measurements have been reported previously.<sup>13</sup> Optical absorption spectra were obtained using a Varian Cary 5000 UV-visible-NIR spectrophotometer while excitation, emission spectra and decay profiles were recorded on an Edinburgh FS5 fluorimeter. The absolute PLQYs of OEuCs in the solid state were determined using a Hamamatsu, C-9920-02 spectrometer equipped with a calibrated integrating sphere of 3.3 inches in radius and measured within the range of 300–750 nm. Excitation was provided with a 150 W xenon lamp after passing through a monochromator. The powder was pressed directly into the sample holder provided by the manufacturer. Photophysical parameters such as the  $J$ - $O$  parameters ( $\Omega_2$  and  $\Omega_4$ ),  $A_{\text{Rad}}$ ,  $A_{\text{NRad}}$  decay rates, radiative lifetime



( $\tau_{\text{rad}}$ ),  $Q_{\text{Eu}}^{\text{Eu}}$  and  $\eta_{\text{sen}}$  were calculated by applying the following set of equations and details are reported elsewhere.<sup>18</sup>

$$\Omega_{\lambda}^{\text{exp}} = \frac{3\hbar A_{\text{Rad}}[{}^5\text{D}_0 \rightarrow {}^7\text{F}_J]}{32e^2\pi^3\chi\nu[{}^5\text{D}_0 \rightarrow {}^7\text{F}_J]^3 |\langle {}^5\text{D}_0 || U^{(\lambda)} || {}^7\text{F}_J \rangle|^2} \quad (1)$$

$$A_{\text{Rad}} = \sum_{J=0}^4 A_{\text{Rad}}[{}^5\text{D}_0 \rightarrow {}^7\text{F}_J] \quad (2)$$

$$A_{\text{Rad}}[{}^5\text{D}_0 \rightarrow {}^7\text{F}_J] = \frac{\nu[{}^5\text{D}_0 \rightarrow {}^7\text{F}_1]}{\nu[{}^5\text{D}_0 \rightarrow {}^7\text{F}_J]} \times \frac{A[{}^5\text{D}_0 \rightarrow {}^7\text{F}_1]}{A[{}^5\text{D}_0 \rightarrow {}^7\text{F}_J]} A_{\text{Rad}}[{}^5\text{D}_0 \rightarrow {}^7\text{F}_1] \quad (3)$$

$$A_{\text{tot}} = \frac{1}{\tau_{\text{obs}}} = A_{\text{Rad}} + A_{\text{NRad}} \quad (4)$$

$$\tau_{\text{R}} = \frac{1}{A_{\text{Rad}}} \quad (5)$$

$$Q_{\text{Eu}}^{\text{Eu}} = \frac{\tau_{\text{obs}}}{\tau_{\text{Rad}}} = \frac{A_{\text{Rad}}}{A_{\text{Rad}} + A_{\text{NRad}}} \quad (6)$$

$$\eta_{\text{sen}} = \frac{Q_{\text{Eu}}^{\text{L}}}{Q_{\text{Eu}}^{\text{Eu}}} \quad (7)$$

Details of the structure determination by single crystal X-ray diffraction, semi-empirical Sparkle/PM7 modelling, device fabrication and instrumentation involved are described in sections 1.1–1.3 of the ESI†

## 3 Results and discussion

### 3.1 Synthesis, characterization and structure determination

The complexes **OEuC-1** and **OEuC-2** were synthesized by employing a two-step method<sup>9b,13</sup> reported earlier and were characterized by elemental analysis, FT-IR spectroscopy and mass spectrometry (Fig. S1–S6, ESI†), which supports the formulation of the complexes shown in Chart 1. The structure of **OEuC-1** (Fig. 1) was established using single-crystal X-ray diffraction while the structure of **OEuC-2** was elucidated using the computational Sparkle/PM7 method (Fig. S7, ESI†). The data obtained are listed in Tables S3–S7, ESI† Both complexes have similar average Eu–O bond distances  $\approx 2.38$  Å, which are well comparable to those of reported analogues.<sup>19</sup> The symmetry of the octa-coordinate  $\text{EuO}_6\text{N}_2$  geometry around the Eu(III) centre

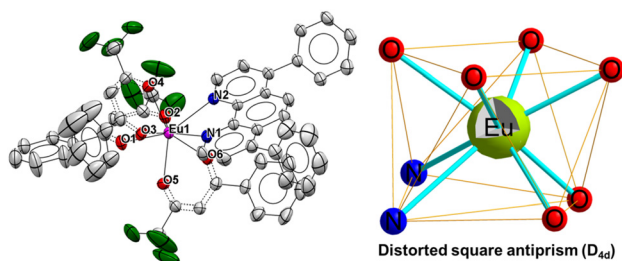


Fig. 1 Single crystal X-ray structure of **OEuC-1** with distorted square antiprism geometry ( $D_{4d}$ ).

was determined using the SHAPE software package.<sup>20</sup> **OEuC-1** adopts a distorted square antiprismatic geometry ( $D_{4d}$ ; with a deviation of 0.715 from the idealized square antiprism, Fig. 1 and Table S8†) while **OEuC-2** exhibits a distorted triangular dodecahedral geometry ( $D_{2d}$ ; with a deviation of 1.687 from the idealized triangular dodecahedron, Fig. S7b and Table S8, ESI†).

### 3.2 Photophysical studies

Optical absorption spectroscopy was employed to gain information on the light absorbing capability of the complexes. The electronic absorption spectra are similar (Fig. S8, ESI†) except for the change in the molar absorptivity value. The complexes displayed broad spectra in the region between 250 and 400 nm with  $\lambda_{\text{max}}^{\text{abs}} = 277$  nm ( $5933 \text{ M}^{-1} \text{ cm}^{-1}$ ) and 320 nm ( $5320 \text{ M}^{-1} \text{ cm}^{-1}$ ) for **OEuC-1** and  $\lambda_{\text{max}}^{\text{abs}} = 277$  nm ( $8424 \text{ M}^{-1} \text{ cm}^{-1}$ ) and 341 nm ( $6038 \text{ M}^{-1} \text{ cm}^{-1}$ ) for **OEuC-2**, respectively. Moreover, the spectrum of **OEuC-2** exhibited a red shift of 21 nm compared to that of **OEuC-1** presumably due to the enhanced quinoidal character of the electron-donating thienyl ring.<sup>21</sup> Both complexes in the solid state and in DCM solution exhibited bright red emission (Fig. S9–S12, ESI†) under UV excitation (Fig. S13–S16, ESI†) with five innate well-resolved transitions from  ${}^5\text{D}_0$  to  ${}^7\text{F}_J$  states ( $J = 0-4$ ) (Fig. 2 and Fig. S15 and S16†). The photophysical data obtained are presented in Table 1. The complexes displayed similar emission spectra, with the induced electric dipole (ED)  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  transition contributing  $\approx 85\%$  of the total integrated intensity with a high asymmetric ratio ( $R_{21}$ ) value, consistent with an asymmetric coordination environment.<sup>18</sup> This is further supported by the large values of  $\Omega_2 = 25.64 \times 10^{-20} \text{ cm}^2$  and  $26.83 \times 10^{-20} \text{ cm}^2$  for **OEuC-1** and **OEuC-2**, respectively, generated by the asymmetric structures of the complexes. Furthermore, the prevalence of the  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  transition advocates that the forced ED and the dynamic coupling mechanism dominate over the magnetic dipoles (MDs).<sup>22</sup> The excited state lifetime ( $\tau_{\text{obs}}$ ) of the  ${}^5\text{D}_0$  emitting state for the complexes calculated by the fitting of the decay

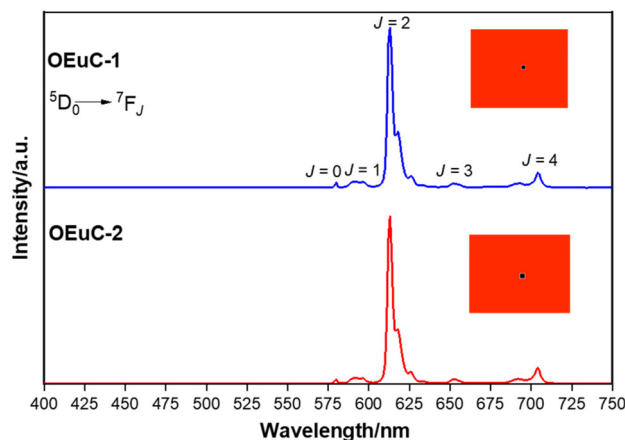


Fig. 2 Solid-state PL emission spectra of the OEuCs. The inset shows a magnified view of the 1931 CIE chromaticity diagram.



**Table 1** Photophysical properties of **OEuC-1** and **OEuC-2** in the solid state

Photophysical parameters	<b>OEuC-1</b>	<b>OEuC-2</b>
${}^5D_0 \rightarrow {}^7F_0$	17253.97 cm <sup>-1</sup> [0.74%]	17257.41 cm <sup>-1</sup> [0.71%]
${}^5D_0 \rightarrow {}^7F_1$	16866.49 cm <sup>-1</sup>	16869.98 cm <sup>-1</sup>
${}^5D_0 \rightarrow {}^7F_2$	16246.87 cm <sup>-1</sup> [85.51%]	16247.97 cm <sup>-1</sup> [84.31%]
${}^5D_0 \rightarrow {}^7F_3$	15308.77 cm <sup>-1</sup> [2.78%]	15312.50 cm <sup>-1</sup> [2.91%]
${}^5D_0 \rightarrow {}^7F_4$	14301.46 cm <sup>-1</sup> [10.96%]	14310.08 cm <sup>-1</sup> [12.01%]
FWHM of ${}^5D_0 \rightarrow {}^7F_2$ (nm)	3.76	3.36
Asymmetric ratio ( $R_{21}$ ) <sup>a</sup>	14.71	15.38
(CIE) <sub>x,y</sub> colour coordinates	0.666; 0.330	0.668, 0.330
$\tau_{obs}$	586.00 ± 2.16 μs ( $\chi^2 = 1.04$ )	623 ± 0.76 μs ( $\chi^2 = 1.02$ )
$\Omega_2^b$ ( $\times 10^{-20}$ cm <sup>2</sup> )	25.64	26.83
$\Omega_4^b$ ( $\times 10^{-20}$ cm <sup>2</sup> )	7.67	8.84
$A_{Rad}^c$ (s <sup>-1</sup> )	962.04	1018.32
$A_{NRad}^c$ (s <sup>-1</sup> )	744.44	586.81
$\tau_R^d$ (μs)	1039.50	982
$Q_{Eu}^{Eu}$ (%)	56.38	63.44
$Q_{Eu}^L$ (%)	80.00 ± 10	80.09 ± 10
$\eta_{Sen}^f$ (%)	70.45	79.21
$\Delta E$ ( ${}^3\pi\pi^*$ (β-diketone) - ${}^5D_0$ ) [cm <sup>-1</sup> ]	4100	3000
$\Delta E$ ( ${}^3\pi\pi^*$ (Bphen) - ${}^5D_0$ ) [cm <sup>-1</sup> ]	3700	3700

Values in the square brackets are % contribution in relation to  ${}^5D_0 \rightarrow {}^7F_1$  MD transition. <sup>a</sup> Ratio of the integrated intensity of the induced electric dipole (ED)  ${}^5D_0 \rightarrow {}^7F_2$  to the  ${}^5D_0 \rightarrow {}^7F_1$  MD transition. <sup>b</sup>  $\Omega_2$  and  $\Omega_4$  were calculated by applying eqn (1) and (2). <sup>c</sup>  $A_{Rad}$  and  $A_{NRad}$  were calculated by applying eqn (2)–(4),  $ESI_{\dagger}^d$   $\tau_R$  was calculated by applying eqn (5),  $ESI_{\dagger}^e$   $Q_{Eu}^{Eu}$  was calculated by applying eqn (6),  $ESI_{\dagger}^f$   $\eta_{Sen}$  was calculated by applying eqn (7),  $ESI_{\dagger}$

curves (Fig. S17–S20, ESI†) lies on the microsecond (μs) time-scale (Table 1). Both the **OEuCs** exhibited high absolute PLQYs in the solid state reaching 80.00 ± 10%, which could be due to the optimum  $\Delta E$  values (Table 1). To the best of our knowledge, these values are the highest to date for systems with a  $EuO_6N_2$  coordination environment. Interestingly,  $\eta_{Sen}$  of **OEuC-2** is ≈10% higher than that of **OEuC-1**, despite both complexes having similar PLQYs. A plausible explanation is that there is a ≈ 21% reduction in  $A_{NRad}$  of **OEuC-2**, which has six fewer C–H oscillators than **OEuC-1**, a factor that is directly related to eqn (7). It may be noted that there are a few nona-coordinated homoleptic  $Eu(III)$  complexes<sup>23</sup> bearing anionic tridentate ligands exhibiting PLQY values >80%, reaching up to 94%. However, they possess ultra-long  $\tau_{obs}$  (1.65 ms–2.74 ms)<sup>23</sup> and thus would have a greater probability of hindering EL performance.

### 3.3 Electroluminescence and electrophysical properties of **OEuC** based OLEDs

Good thermal stability of the newly synthesized materials (here **OEuC-1** and **OEuC-2**) is an important pre-requisite for the fabrication of EL devices, since poor thermal stability hampers the device performance along with the operational lifetime at the peak of its operation owing to the Joule heating effect

when current flows through the organic layer.<sup>3b,24</sup> The thermal stability of the **OEuCs** was evaluated in the temperature range of 50 to 700 °C. The thermograms of **OEuCs** (Fig. S3 & S5, ESI†) revealed that both the **OEuCs** possess good thermal stability with a  $T_d$  (with 5% weight loss) of 315 °C for **OEuC-1** and 330 °C for **OEuC-2**, respectively, implying that they can be readily employed to fabricate EL devices by the vacuum thermal evaporation method. Moreover, the thermal stability of the complexes is well comparable to that of analogous efficient red-emitting complexes *i.e.*, [Eu(btfa)<sub>3</sub>DPEPO] ( $T_d = 320$  °C),<sup>25</sup> [Eu(nta)<sub>3</sub>DPEPO] ( $T_d = 318$  °C),<sup>25</sup> and [Eu(btfa)<sub>3</sub>(Ph-TerPyr)] ( $T_d = 313$  °C),<sup>3a</sup> and higher than that of the complexes [Eu(btfa)<sub>3</sub>]<sub>2</sub>bpm ( $T_d = 302$  °C),<sup>26</sup> [Eu(tfac)<sub>3</sub>DPEPO] ( $T_d = 287$  °C),<sup>27</sup> [Eu(btfa)<sub>3</sub>Py-Im] ( $T_d = 286$  °C),<sup>28</sup> [Eu(tfac)<sub>3</sub>(TB-Im)] ( $T_d = 241$  °C)<sup>3b</sup> and [Eu(hfac)<sub>3</sub>(TB-Im)] ( $T_d = 239$  °C)<sup>3b</sup> (DPEPO = bis(2-(diphenylphosphino)phenyl)ether oxide; nta = 4,4,4-trifluoro-1-(2-naphthyl)-1,3-butanedione; Ph-TerPyr = 4'-phenyl-2,2':6',2''-terpyridine; bpm = 2,2'-bipyrimidine; Py-Im = 2-(2-pyridyl)benzimidazole; tfac = trifluoroacetylacetone; TB-Im = 2-(4-thiazolyl)benzimidazole; hfac = hexafluoroacetylacetone). Encouraged by the excellent photophysical properties and good thermal stability of **OEuC-1** and **OEuC-2** and to demonstrate their potential application as the emitting material in red OLEDs, we have fabricated two single-EML devices (**Devices A** and **E**) with a hole-transporting host CzSi and a fixed doping concentration of **OEuCs** (2 wt%). The EL spectra, CIE colour diagrams and current density ( $J$ )–brightness ( $B$ )–voltage ( $V$ ) curves are shown in the ESI (Fig. S21–S35, ESI†), while the device performance is summarized in Table 2. Both the single-EML devices displayed typical  $Eu(III)$  emissions due to the long-range Förster energy transfer (ET) from the host to the complexes and carrier trapping by the **OEuCs** as observed by one of us.<sup>29</sup> This gains support from the partial overlap of the CzSi emission with the **OEuC** absorption spectra and the PLQY of **OEuC-1** (2 wt%):CzSi (PLQY = 40%, Table S9, ESI†). Importantly, both devices start illuminating at the same low  $V_{turn-on} = 4.4$  V implying that the carrier injection and transport abilities of both **OEuCs** are similar. **Device A** displayed pure red EL with  $B = 156$  cd m<sup>-2</sup>,  $CE_{max} = 1.94$  cd A<sup>-1</sup>,  $PE_{max} = 1.39$  lm W<sup>-1</sup> and  $EQE_{max} = 1.10\%$ . Moreover, the EL performance of **Device E** is similar; however, the emitted colour is magenta due to the presence of the CzSi emission. A plausible explanation could be because of the greater electron donating capability of thiophene which may have generated extra electron(s) during the electrical excitation resulting in an imbalance between the electron–hole pair.<sup>30</sup>

To further improve the OLED performance, two double-EML devices (**Devices B** and **F**) with CzSi and another hole-transport and hole-injection TcTa material were fabricated. As expected, the double-EML device of **OEuC-1** (**Device B**) exhibited almost 70% improvement in the CE and EQE with a marginal increase in  $B = 163$  cd m<sup>-2</sup> compared to **Device A**. The **OEuC-2** based device (**Device F**) also exhibited a three-fold enhancement in the overall performance; however, its colour remained in the light magenta region due to the marginal decrease in host emission *i.e.*, improved ET. To further tune



Table 2 Key EL performance of Devices A–H of OEuCs

Device	$V_{\text{turn-on}}^a$	$B_{\text{max}}^b$	$\text{EQE}_{\text{max}}^c$ (%)	$\text{CE}_{\text{max}}^d$	$\text{PE}_{\text{max}}^e$	$\text{FWHM}^f$	$(\text{CIE})_{x,y}$	$\text{EQE} (\%)@100 \text{ cd m}^{-2}$
<b>OEuC-1-based devices</b>								
Device A	4.4	156	1.10	1.94	1.39	6.52	0.608, 0.327	0.09
Device B	3.8	163	1.57	2.78	2.30	6.65	0.611, 0.323	0.11
Device C	3.3	425	4.83	7.89	7.51	6.34	0.630, 0.328	0.67
Device D	3.5	557	5.91	8.58	7.70	6.31	0.636, 0.326	1.18
<b>OEuC-2-based devices</b>								
Device E	4.4	228	1.08	1.38	0.99	6.00	0.500, 0.300	0.14
Device F	3.5	179	3.63	5.27	4.73	5.97	0.520, 0.300	0.18
Device G	3.2	433	5.01	8.22	8.07	6.00	0.603, 0.320	0.69
Device H	3.4	545	6.24	9.91	9.15	6.02	0.620, 0.323	1.31

<sup>a</sup> Turn-on voltage in volt. <sup>b</sup> Maximum brightness in  $\text{cd m}^{-2}$ . <sup>c</sup> External quantum efficiency in %. <sup>d</sup> Maximum current efficiency in  $\text{cd A}^{-1}$ . <sup>e</sup> Maximum power efficiency in  $\text{lm W}^{-1}$ . <sup>f</sup> Full width at half maxima for  $^5\text{D}_0 \rightarrow ^7\text{F}_2$  transition in nm under electrical excitation; Commission Internationale de l'éclairage at  $J = 10 \text{ mA cm}^{-2}$ .

the OLED performances, two single-EML and two double-EML OLEDs [Devices C and D (OEuC-1) and Devices G and H (OEuC-2), Fig. S36–S46, ESI†] were fabricated by replacing the unipolar host CzSi with a bipolar host 26DCzPPy (Fig. 3). As can be seen from Table 2, all the devices exhibited red EL (Fig. 4) with low  $V_{\text{turn-on}} = 3.2\text{--}3.5 \text{ V}$ . The EL performance of the OEuC-1 based devices *i.e.*, Devices C ( $\text{EQE}_{\text{max}} = 4.83\%$ ) and D ( $\text{EQE}_{\text{max}} = 5.91\%$ ) is more than three times higher than that of Devices A ( $\text{EQE}_{\text{max}} = 1.10\%$ ) and B ( $\text{EQE}_{\text{max}} = 1.57\%$ ). Moreover, the OEuC-2 based Devices G and H also displayed a similar improvement, which is approximately two times higher than that of the CzSi-based devices (Devices E and F). It is important to remember that the emission of 26DCzPPy partially overlaps with the absorption spectra of OEuCs (Fig. S47,

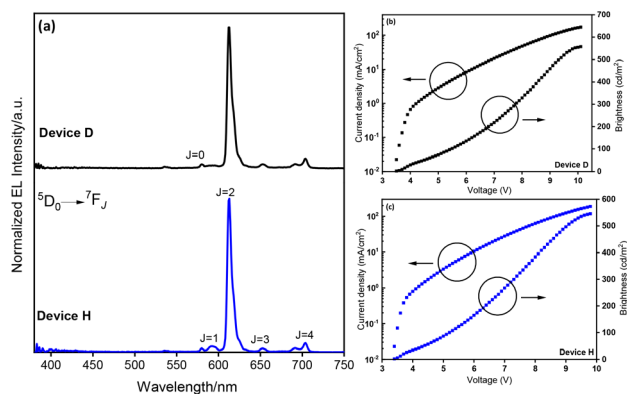


Fig. 4 (a) Normalized EL spectra of Devices D and H ( $J = 10 \text{ mA cm}^{-2}$ ).  $J$ - $V$ - $B$  curves of Devices (b) D and (c) H.

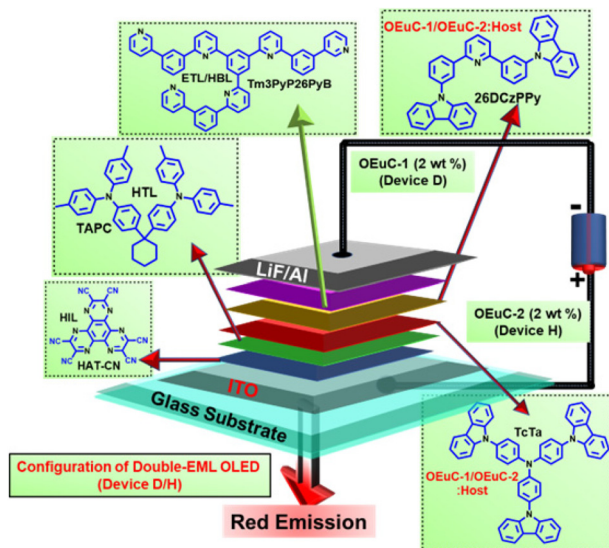


Fig. 3 General device configuration of double-EML OLEDs (HIL: Hole Injection Layer; HTL: Hole Transport Layer; ETL: Electron Transport Layer; HBL: Hole Blocking Layer; HAT-CN: 1,4,5,8,9,11-hexaazatriphenylene hexacarbonitrile; TAPC: di-[4-(*N,N*-ditolyl-amino)-phenyl]cyclohexane; Tm3PyP26PyB: 1,3,5-tris(6-(3-(pyridin-3-yl)phenyl)pyridin-2-yl)benzene; host: 26DCzPPy: 2,6-bis(3-(9*H*-carbazol-9-yl)phenyl)pyridine; TcTa: 4,4',4''-tris(carbazole-9-yl)triphenylamine).

ESI†), which implies that the carrier trapping process could be the dominant EL process in these devices. To confirm this, we have determined the photophysical properties of the OEuCs: host films (Tables S9 and S10†), which revealed that carrier trapping is the dominant process in the cases of Devices C, D and H, while for Device G, both Förster ET and carrier trapping play a crucial role (PLQY = 52.8%). It was also apparent that regardless of the large or small PLQYs of complex: host films (Tables S9 and S10, ESI†), their brightness remains dependent on the excited state lifetime, *e.g.*, Device A (PLQY = 40.4%,  $\tau_{\text{obs}} = 564.27 \mu\text{s}$ ) and Device B (PLQY = 10.3%,  $\tau_{\text{obs}} = 566.79 \mu\text{s}$ ) showed similar brightness (Table 2). Importantly, the overall OLED performance of Device H is the best among the  $\text{O}_6\text{N}_2/\text{O}_6\text{O}_2$  coordinated OEuCs (Chart S3 and Table S11†) with  $\text{CE}_{\text{max}} = 9.91 \text{ cd A}^{-1}$ ,  $\text{PE}_{\text{max}} = 9.15 \text{ lm W}^{-1}$ ,  $\text{EQE}_{\text{max}} = 6.24\%$ ,  $B$  of  $545 \text{ cd m}^{-2}$ , and  $(\text{CIE})_{x,y} = 0.620, 0.323$  at  $J = 10 \text{ mA cm}^{-2}$ . Despite the impressive OLED performance of OEuC-2-based Device H among OEuC-based devices, its EL performance falls behind that of the red-emitting Ir(III) complexes, which usually exhibit  $\text{EQE} > 10\%$  at a luminance of  $1000 \text{ cd m}^{-2}$ .<sup>31</sup> This could be due to the very short excited state lifetime of 0.1–5  $\mu\text{s}$  of Ir(III) complexes<sup>31b</sup> compared to that of the OEuCs (100–800  $\mu\text{s}$ ).



## 4 Conclusions

We have synthesized two highly luminescent red-emitting OEuCs displaying PLQY = 80% ± 10 and successfully applied them as EMLs to fabricate four single-EML and four double-EML devices. Through the combination of appropriate host materials, the device based on OEuC-2 (**Device H**) exhibited a remarkable EL performance with  $CE_{\max} = 9.91 \text{ cd A}^{-1}$ ,  $PE_{\max} = 9.15 \text{ lm W}^{-1}$ ,  $EQE_{\max} = 6.24\%$ ,  $B$  of  $545 \text{ cd m}^{-2}$ , and  $(CIE)_{x,y} = 0.620, 0.323$  at  $J = 10 \text{ mA cm}^{-2}$ , which, we believe, is the best among the  $O_6N_2/O_6O_2$  coordinated OEuCs (Chart S3 and Table S11†). There are few important observations that we made from the present study: the brightness of the device is independent of the PLQY of the complex: host film but depends on the excited state lifetime of the complex: host film e.g., **Device A** ( $B = 156 \text{ cd m}^{-2}$ , PLQY = 40.4% and  $\tau_{\text{obs}} = 564.27 \mu\text{s}$ ) and **Device B** ( $B = 163 \text{ cd m}^{-2}$ , PLQY = 10.3% and  $\tau_{\text{obs}} = 566.79 \mu\text{s}$ ) and hole-injection TcTa in conjunction with the bipolar 26DCzPPy material could be an excellent choice of host for OEuCs to develop more efficient R-OLEDs.

## Conflicts of interest

There are no conflicts to declare.

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