

## PAPER

View Article Online  
View Journal | View IssueCite this: *J. Mater. Chem. C*, 2023,  
11, 8263Color tuning of multi-resonant thermally activated  
delayed fluorescence emitters based on fully fused  
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Two novel  $\pi$ -extended amine/carbonyl-based multi-resonance thermally activated delayed fluorescence (MR-TADF) emitters have been designed and synthesized. The two emitters are isomeric, composed of nine fused rings and show green-yellow emission. **Sym-DiDiKta** and **Asym-DiDiKta** possess *tert*-butyl groups distributed in a symmetrical and asymmetrical fashion, respectively, which significantly impact the single-crystal packing structure. The two compounds possess similar singlet-triplet energy gaps,  $\Delta E_{ST}$ , of around 0.23 eV, narrowband emission characterized by a full-width at half-maximum, FWHM, of 29 nm and a photoluminescence quantum yield,  $\Phi_{PL}$ , of 70% and 53% for the symmetric and asymmetric counterparts, respectively, in toluene. Investigation in OLEDs demonstrated that the devices with **Sym-DiDiKta** and **Asym-DiDiKta** displayed electroluminescence maxima of 543 and 544 nm, and maximum external quantum efficiencies (EQE<sub>max</sub>) of 9.8% and 10.5%, respectively. The maximum EQE was further improved to 19.9% by employing a hyperfluorescence strategy. We further present the first example of a neutral MR-TADF emitter incorporated in a LEC device where **Sym-DiDiKta** acts as the emitter. The LEC shows a  $\lambda_{EL}$  at 551 nm and FWHM of 60 nm with luminance of 300 cd m<sup>-2</sup> and a fast turn-on time of less than 2 s to 100 cd m<sup>-2</sup>.

Received 20th February 2023,  
Accepted 1st May 2023

DOI: 10.1039/d3tc00641g

rsc.li/materials-c

10<sup>th</sup> Anniversary statement

The journal of Materials Chemistry C has been one of our favourite journals for electroluminescent materials and device research. The quality and breadth of the science covered in the journal particularly in the area of emitter development for electroluminescent devices has been excellent. It is always a pleasure to peruse each week's table of contents and to then read exciting and new science in optoelectronic materials. We look forward to the next 10 years and beyond and will continue to support JMCC.

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† The research data supporting this publication can be accessed at <https://doi.org/10.17630/f844ce22-cf6b-413d-ae79-70e2e38a1f74>

‡ Electronic supplementary information (ESI) available: A summary of prior examples of carbonyl-containing MR-TADF emitters, experimental details, synthesis procedures and characterization data, NMR spectra, HRMS, HPLC, X-ray crystallographic data, supplemental photophysical data, electrochemical data, xyz coordinate file for computations. CCDC 2225501–2225503. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d3tc00641g>

## Introduction

Organic light-emitting diodes (OLEDs) have now matured as a technology and are now integrated in efficient, flexible and ultra-high contrast next-generation displays.<sup>1,2</sup> Organic thermally activated delayed fluorescence (TADF) materials are increasingly viewed as an attractive alternative emitter class to phosphorescent complexes that contain noble metals as they do not contain scarce elements yet can likewise harvest both singlet and triplet excitons in the device to generate light at comparable efficiencies.<sup>3–7</sup> Unlike phosphorescent OLEDs that funnel the emitting excitons *via* the lowest-lying triplet excited state, TADF compounds convert triplet excitons into singlets *via* reverse intersystem crossing (RISC),<sup>8,9</sup> which is possible at ambient temperatures due to the small  $S_1$ – $T_1$  energy gap,  $\Delta E_{ST}$ .<sup>10</sup> Compounds that possess a small

$\Delta E_{ST}$  show a small overlap of the electron density between the highest occupied molecular orbital (HOMO) and lowest unoccupied molecular orbital (LUMO). To adhere to this requirement, the great majority of organic TADF emitters are based on a highly twisted donor-acceptor architecture and emit from a long-range charge transfer (LRCT) state.<sup>7</sup> This strategy, however, leads to molecules for which the conformational space is large in the excited state<sup>11,12</sup> that manifests in broadband emission with full width half maximum (FWHM) values of around 70–100 nm.<sup>13</sup> This is unattractive to the display industry as the color purity of these devices is poor and color filters must therefore be used to meet the industry standards for red, green and blue pixels. Hence, significant effort has been devoted in recent years to suppress vibrational and conformational relaxation within the emitter through the development of narrowband emitters.<sup>14–20</sup>

An exciting molecular design strategy to achieve these requirements was introduced by Hatakeyama and co-workers and relies typically on p- and n-doped nanographenes.<sup>21,22</sup> Termed multiple resonance (or multiresonant) TADF (MR-TADF) emitters, these compounds possess small-to-moderate  $\Delta E_{ST}$  values, narrowband emission (with FWHM typically < 30 nm) and limited positive solvatochromism due to the short-range charge transfer (SRCT) nature of the emissive  $S_1$ .<sup>22</sup> Maximum external quantum efficiencies (EQE<sub>max</sub>) as high as 34% for deep blue,<sup>22</sup> 40% for sky-blue,<sup>23</sup> 35%<sup>24</sup> for green and 36% for red<sup>25</sup> have been achieved for OLEDs containing MR-TADF emitters comprised of boron-nitrogen (B/N) doped polycyclic aromatic frameworks. Recently, the library of MR-TADF emitters has been expanded to contain boron-oxygen (B/O)-based emitters<sup>26</sup> and carbonyl-nitrogen (C=O/N)-based emitters<sup>14–19</sup> as well as C=O/N/O,<sup>17,27,28</sup> C=O/N/S,<sup>17, 29</sup> and C=O/N/SO<sub>2</sub>-based<sup>29</sup> emitters.

While the recent conceptual progress in MR-TADF emitter design has delivered a burgeoning number of efficient B/N-doped compounds that emit across the entirety of the visible spectrum,<sup>30,31</sup> few tactics have been advanced to exploit the promising class of C=O/N-based MR-TADF emitters, which are represented by the emitter **DiKTa** (Fig. 1).<sup>14–19,32,33</sup> Recently,

Yasuda *et al.* reported a linearly extended C=O/N-based emitter, **QA-2** (Fig. 1) displaying TADF in 3 wt% PPCz doped film, with a  $\Delta E_{ST}$  of 0.19 eV.<sup>27</sup> Similar to the design strategy used for ***v*-DABNA**, where the central ring is functionalized with *meta*-disposed B- $\pi$ -B and N- $\pi$ -N groups, **QA-2** also contains *meta*-disposed C=O- $\pi$ -C=O and N- $\pi$ -N, which leads to an improvement of the TADF properties over the parent compound **DiKTa**, with a faster delayed lifetime (48  $\mu$ s in 3 wt% PPCz and 93.3  $\mu$ s in 5 wt% mCP, respectively). Despite the apparent increased conjugation length, a slightly blue-shifted  $\lambda_{PL}$  of 465 nm in toluene was observed for **QA-2**, compared to that of **DiKTa** ( $\Delta E_{ST}$  of 0.15 eV and  $\lambda_{PL}$  of 473 nm in toluene).<sup>15</sup> The OLED with **QA-2** showed an EQE<sub>max</sub> of 19.0%. Our group has recently reported a helically chiral isomer of **QA-2** that contains *meta*-disposed C=O- $\pi$ -C=O and N- $\pi$ -N skeleton. Enantiomers of **Hel-DiDiKTa** (Fig. 1)<sup>34</sup> display circularly polarized luminescence (CPL), with a  $|g_{PL}|$  of  $4 \times 10^{-4}$ , but likewise, its  $\lambda_{PL}$  (473 nm) is similar to that of **DiKTa**.

The exploitation of *para*-disposed functional groups to generate  $\pi$ -extended skeletons is a powerful strategy to tune the photophysical properties of MR-TADF emitters to the red.<sup>25,35</sup> Yasuda and co-workers have adopted this strategy where both the B- $\pi$ -B and N- $\pi$ -N-based groups are *para*-linked to construct red MR-TADF emitters such as **BBCz-R** (Fig. 1) that exhibits a  $\lambda_{PL}$  of 615 nm and a narrow FWHM of 21 nm (0.07 eV) in dilute toluene solution.<sup>31</sup> This strategy has also been used to generate B/O and B/S derivatives as shown in Fig. 1.<sup>36,37</sup> Remarkably, among the many MR-TADF emitters reported, comprehensive studies on compounds containing *para*-C=O- $\pi$ -C=O and N- $\pi$ -N frameworks have not as of yet been presented. In this context, expanding the chemical space explored in ketone-based MR-TADF emitters is desired.

Herein, we present two new isomeric  $\pi$ -extended C=O/N-based polycyclic aromatic compounds based on the annellation of two triangulene **DiKTa** units to form tetraketone structures. Differently from **QA-2** and **Hel-DiDiKTa**, the central ring in each of **Sym-DiDiKTa** and **Asym-DiDiKTa** is functionalized with *para*-C=O- $\pi$ -C=O and N- $\pi$ -N groups, these molecules represent

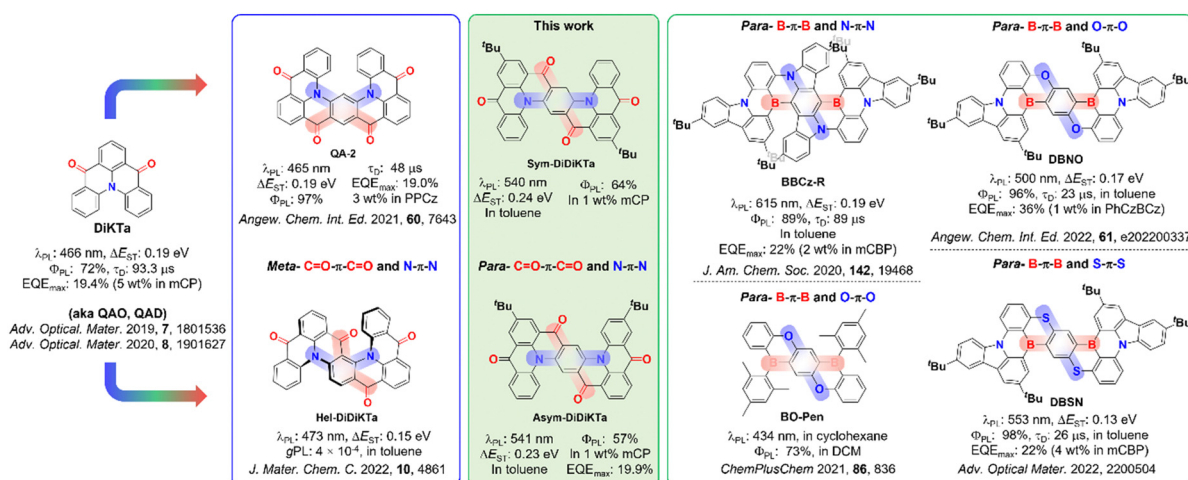


Fig. 1 Schematic representation of the molecular design discussed in this work.<sup>15,27,31,34,36,37,42</sup>



the first ketone-based MR-TADF examples of their kind (Fig. 1).<sup>17–19,33</sup> **Sym-DiDiKTa** is composed of nine fully fused rings with symmetrically arranged *tert*-butyl substituents while the *tert*-butyl groups are disposed asymmetrically in **Asym-DiDiKTa**. Both emitters show moderate  $\Delta E_{ST}$  of around 0.23 eV, red-shifted emission at  $\sim 540$  nm compared to **DiKTa**, a small FWHM of 29 nm and delayed lifetime,  $\tau_d$ , of 4.6 ms and 3.0 ms for **Sym-DiDiKTa** and **Asym-DiDiKTa**, respectively.

Owing to their exciting properties, these compounds were employed as emitters in OLED devices, and to assess improvements in efficiency roll-off, they were also investigated as terminal emitters in hyperfluorescence (HF) OLEDs, which combine a narrowband emitter as terminal dopant with a co-deposited TADF emitter acting as an assistant dopant in a host matrix. Although MR-TADF OLEDs have achieved extremely high EQE<sub>max</sub>, an issue generally encountered is the severe efficiency roll-off that is mainly caused by quenching mechanisms such as singlet–triplet annihilation (STA) and triplet–triplet annihilation (TTA) due to the accumulation of triplet excitons as a result of their typically slow reverse intersystem-crossing rate ( $k_{RISC}$ ).<sup>17,38</sup> Therefore, a potential solution to circumvent this issue is to use MR-TADF compounds in combination with an efficient exciton harvesting assistant dopant in a HF device.<sup>39–41</sup> In these devices, exciton harvesting is managed by the donor–acceptor TADF assistant dopant and these excitons are transferred to the terminal MR-TADF emitter *via* a Förster

resonant energy transfer (FRET) mechanism. Emission from the terminal emitter then occurs, benefiting from the typically fast radiative decay and narrowband emission of the MR-TADF compound.<sup>39</sup> To ensure efficient FRET, there must be an effective overlap between the emission spectrum of the assistant dopant and the absorption spectrum of the emitter.

## Results and discussion

The two emitters were obtained *via* a four-step linear sequence (Fig. 2a). Compound **1** was obtained in good yield following a copper-catalysed Ullman coupling between the previously reported dimethyl 2,5-bis((4-*tert*-butyl)phenyl)amino)terephthalate<sup>43</sup> and methyl 2-iodobenzoate. Quantitative saponification yielded the key intermediate **2**, which then underwent a four-fold intramolecular Friedel–Crafts acylation of the *in situ*-prepared acyl chloride derivative in the presence of the Lewis acid  $AlCl_3$  to afford a mixture of two isomers, **Sym-DiDiKTa** and **Asym-DiDiKTa**, in 36% and 5% yield, respectively, after isolation by column chromatography. The compounds were further purified by gradient-temperature vacuum sublimation and their structure and purity were confirmed by a combination of NMR spectroscopy, high-resolution mass spectrometry, melting point determination and HPLC and elemental analysis.

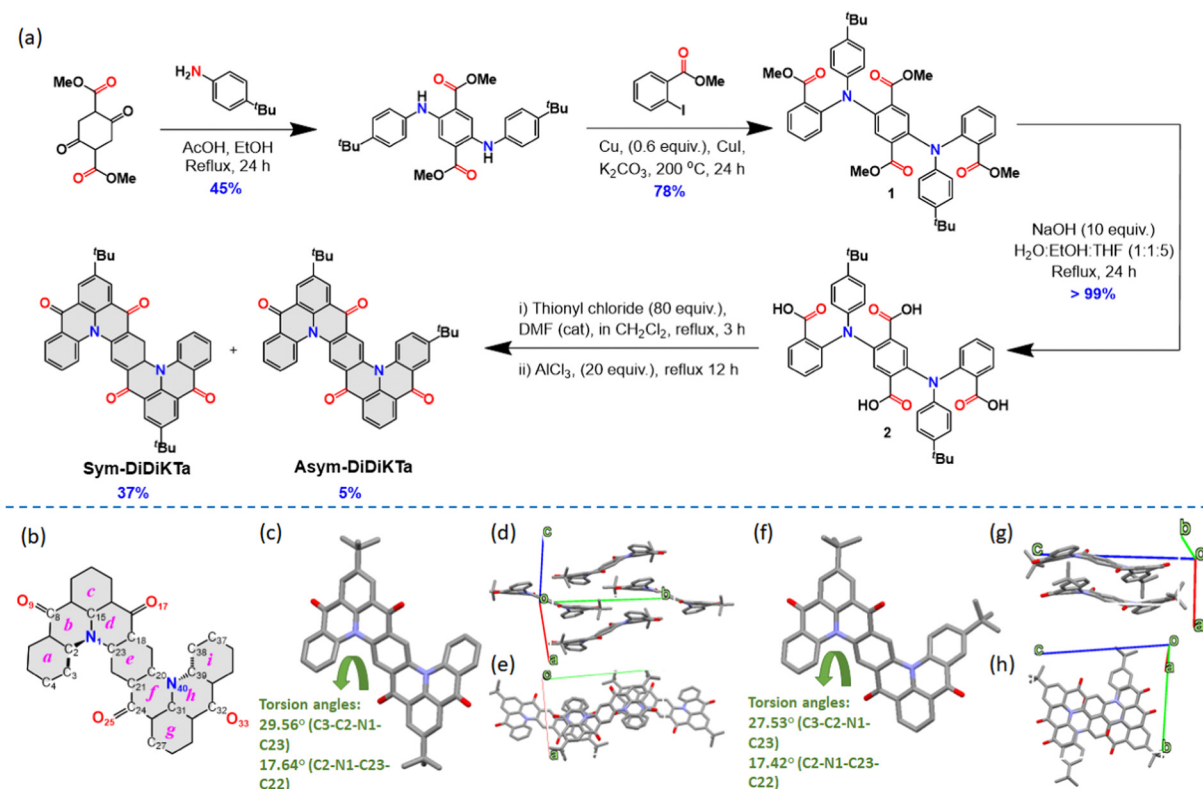


Fig. 2 (a) Synthesis of **Sym-DiDiKTa** and **Asym-DiDiKTa**. (b) Schematic of molecular structure of **Sym-DiDiKTa** and **Asym-DiDiKTa** showing atom labels (double bonds omitted for clarity). (c and f) View showing **Sym-DiDiKTa** and **Asym-DiDiKTa** in the crystal structure (diagonal view). (d and g) Side view of the packing diagram of **Sym-DiDiKTa** and **Asym-DiDiKTa**. (e and h) Top view of the packing diagram of **Sym-DiDiKTa** and **Asym-DiDiKTa**.



Single crystals were isolated following the gradient-temperature vacuum sublimation. Crystallographic data of **Sym-DiDiKta** and **Asym-DiDiKta** revealed that both molecules display very similar structures with each **DiKta** unit showing a helical-like structure. The torsion angles generated by this helical section are similar (torsion angles C3–C2–N1–C23): 29.6(5)° for **Sym-DiDiKta** and 27.9(3) and 28.1(3)° for **Asym-DiDiKta** (Fig. 2c and f). Both compounds pack following a one-dimensional slipped  $\pi$ - $\pi$  stacking motif with interplanar chains along the *c*-axis for **Sym-DiDiKta** [centroid...centroid distance 3.5876(18) Å] and along the *a*-axis for **Asym-DiDiKta** [centroid...centroid distances 3.4697(13)–3.7685(13) Å]. The disposition of the *tert*-butyl groups in the compounds influences their packing. In the case of **Sym-DiDiKta**, adjacent molecules are partially superimposed, forming  $\pi$ -stacked chain along the *c*-axis with the molecules stacked in a slipped manner (Fig. 2d and e). In contrast, for **Asym-DiDiKta**, adjacent molecules display a greater degree of overlap and more  $\pi$ -stacking (Fig. 2g), resulting in symmetric pairs of adjacent molecules with the *tert*-butyl groups pointing in the opposite direction (Fig. 2h). Additional  $\pi$ -stacking interactions link these to form  $\pi$ -stacked chains along the *a*-axis. The distance between the central ring (*e*) between two nearest molecules is 9.1422(9) Å for **Sym-DiDiKta**, whereas for **Asym-DiDiKta** it is only 3.4699(17) Å. We also obtained single crystals of **Sym-DiDiKta** from slow evaporation of a CH<sub>2</sub>Cl<sub>2</sub>:MeOH solution (8:2 ratio). Under these conditions, the crystal structure is comprised  $\pi$ -stacked columns of molecules along the crystallographic *a*-axis (Fig. S13a and b, ESI†), with greater superimposition than in the previous case (distance between the central rings (*e*) of 3.8500(2) Å). This is achieved with the aid of MeOH molecules forming both strong and weak hydrogen bonds between and within the columns.

We have previously demonstrated that DFT methods are not appropriate for accurately predicting the excited state properties of MR-TADF materials.<sup>44</sup> Explicit inclusion of second-order electronic correlation effects using wavefunction-based methods such as spin component scaling second-order approximate Coupled-Cluster (SCS-CC2) calculations with the cc-pVDZ basis set addresses this problem, resulting in accurate  $\Delta E_{ST}$  prediction for MR-TADF materials.<sup>45</sup> The two isomers

possess identical  $S_1$  and  $T_1$  energies of 2.93 eV and 2.69 eV, respectively, and thus a  $\Delta E_{ST}$  of 0.24 eV (Fig. 3), which is similar to previously reported ketone-containing MR-TADF materials.<sup>46</sup> Compared to the parent compound **DiKta** ( $\Delta E_{ST}$  = 0.27 eV and  $S_1$  = 3.45 eV), there is a modest decrease in  $\Delta E_{ST}$  and significant stabilization of  $S_1$ . The smaller computed  $\Delta E_{ST}$  in **Sym-DiDiKta** to **Asym-DiDiKta** relative to **DiKta** is due to the increase in the  $\pi$ -conjugation of these materials compared to **DiKta** resulting in a lowering of the exchange energy, as has been documented in other extended MR-TADF systems, such as **v-DABNA** and **OAB-ABP-1**.<sup>26,46</sup> The predicted stabilization of  $S_1$  results from the nitrogen atoms being positioned *para* to each other, which positively reinforce their electron-donating character, as has been previously reported.<sup>18,41</sup> The oscillator strength associated with the  $S_0$ – $S_1$  transition is significant at 0.26 and 0.27 for **Sym-DiDiKta** and **Asym-DiDiKta**, respectively. The difference density plots of  $S_1$  and  $T_1$  of each emitter highlight the alternating pattern of increasing and decreasing electronic density on neighbouring atoms that produce the emissive short-range charge transfer (SRCT) excited state, which is characteristic of MR-TADF materials.<sup>46</sup>

Cyclic voltammetry (CV) and differential pulse voltammetry (DPV) recorded in degassed dichloromethane (Fig. 4a and b) document a reversible reduction wave at  $E_{red}$  = –0.96 and –0.94 V vs. SCE for **Sym-DiDiKta** and **Asym-DiDiKta**, respectively. The oxidation wave of **Sym-DiDiKta** is more reversible than that of **Asym-DiDiKta** (both at 1.53 V vs. SCE), revealing the impact of the regiochemistry that the *tert*-butyl groups play in terms of electrochemical stability. The corresponding HOMO and LUMO levels are nearly identical, at –5.87 eV for both compounds and –3.38 eV for **Sym-DiDiKta** and –3.40 eV for **Asym-DiDiKta**, respectively, which align well with the gas-phase DFT calculations at the PBE0/6-31G(d,p) level of theory (–5.94 eV and –2.69 eV, respectively, Table S11 and Fig. S20, ESI†). The experimentally determined HOMO levels of **Sym-DiDiKta** and **Asym-DiDiKta** are similar to that of **DiKta** (–5.93 eV), while the LUMO levels are significantly stabilized (LUMO of **DiKta** = –3.11 eV).<sup>15</sup> The electrochemical data are summarized in Table S9 (ESI†).

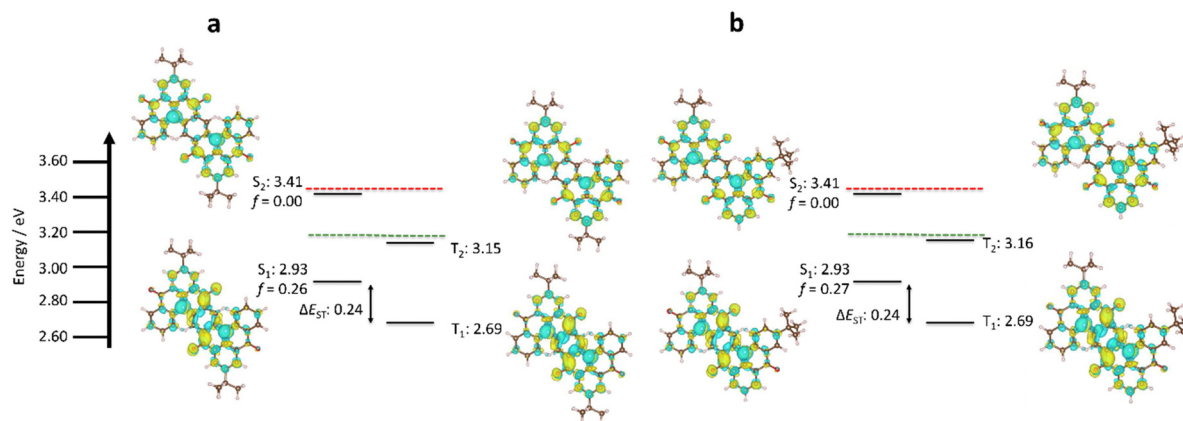


Fig. 3 Difference density picture and excited state energies of **Sym-DiDiKta** (a) and **Asym-DiDiKta** (b) calculated at SCS-CC2/cc-pVDZ, where red and green lines indicate  $S_1$  and  $T_1$  energies respectively of **DiKta**, (isovalue = 0.001).





The monomolecular photophysical properties of **Sym-DiDiKta** and **Asym-DiDiKta** were first studied in dilute toluene solution (Fig. 4c, d and Table 1). The UV-vis absorption spectra of both emitters in toluene are expectedly nearly identical, yet show significant differences compared to that of **DiKta** in terms of the SRCT absorption band's wavelength (around 435 nm for **DiKta**<sup>14,15</sup> but around 515 nm for **Sym-DiDiKta** and **Asym-DiDiKta**). The molar absorptivity,  $\epsilon$ , of the SRCT band for **Sym-DiDiKta** and **Asym-DiDiKta** is 26,325 M<sup>-1</sup> cm<sup>-1</sup> and 23,458 M<sup>-1</sup> cm<sup>-1</sup>, respectively, assigned to the transition to  $S_1$  according to the SCS-CC2 calculations (Fig. 3). The  $\epsilon$  is somewhat larger than reported for **DiKta** ( $\epsilon$  of **DiKta** = 21 000 M<sup>-1</sup> cm<sup>-1</sup>),<sup>15</sup> which is consistent with the trends in the calculated oscillator strength where  $f$  = 0.20 for **DiKta**, compared to 0.26 and 0.27 for **Sym-DiDiKta** and **Asym-DiDiKta**, respectively. The absorption spectra of both **Sym-DiDiKta** and **Asym-DiDiKta** also show a high-energy shoulder at 485 nm ( $\epsilon$  = 11 680 and 10 238 M<sup>-1</sup> cm<sup>-1</sup> for **Sym-DiDiKta** and **Asym-DiDiKta**, respectively), which likely originates from a vibronic band and not a transition to a higher-lying singlet state given that computed energies of the  $S_2$  state in each of these two compounds is *ca.* 0.48 eV higher in energy and the  $S_0$ - $S_2$  transition possesses negligible oscillator strength.<sup>16</sup> There is also a high-

energy, high-intensity band at 373 nm ( $\epsilon$  = 31 900 M<sup>-1</sup> cm<sup>-1</sup> and 28 527 M<sup>-1</sup> cm<sup>-1</sup> for **Sym-DiDiKta** and **Asym-DiDiKta**, respectively, Fig. 4c and d), which is assigned to transitions to  $S_4$  and  $S_5$ , for **Sym-DiDiKta** and **Asym-DiDiKta**, respectively (Table S12, ESI<sup>†</sup>) based on the comparison with the SCS-CC2 simulated absorption spectra (Fig. S21 and Table S12, ESI<sup>†</sup>). The difference density plots of these states (Fig. S21, ESI<sup>†</sup>) show that there is only minimal density situated on the central phenyl ring and more density situated on the carbonyl groups compared with the difference density pattern of the  $S_1$  state (*vide supra*).

Both compounds display green emission in dilute toluene with emission maxima,  $\lambda_{PL}$ , of 540 nm and 541 nm for **Sym-DiDiKta** and **Asym-DiDiKta**, respectively, which are *ca.* 90 nm red-shifted from that of **DiKta** ( $\lambda_{PL}$  = 453 nm). This bathochromic shift is corroborated by the SCS-CC2 calculations.<sup>19,20,31,48</sup> Both compounds display narrow PL spectra at room temperature (FWHM = 29 nm), and small Stokes shifts of *ca.* 25 nm, which confirms the small degree of geometrical reorganization in the excited state owing to their conformationally rigid structure (Fig. 2c and d). The small degree of positive solvatochromism (Fig. S15 and Tables S2, S3, ESI<sup>†</sup>) reflects the SRCT character of the emissive excited state.<sup>15</sup> The  $S_1$  excited state

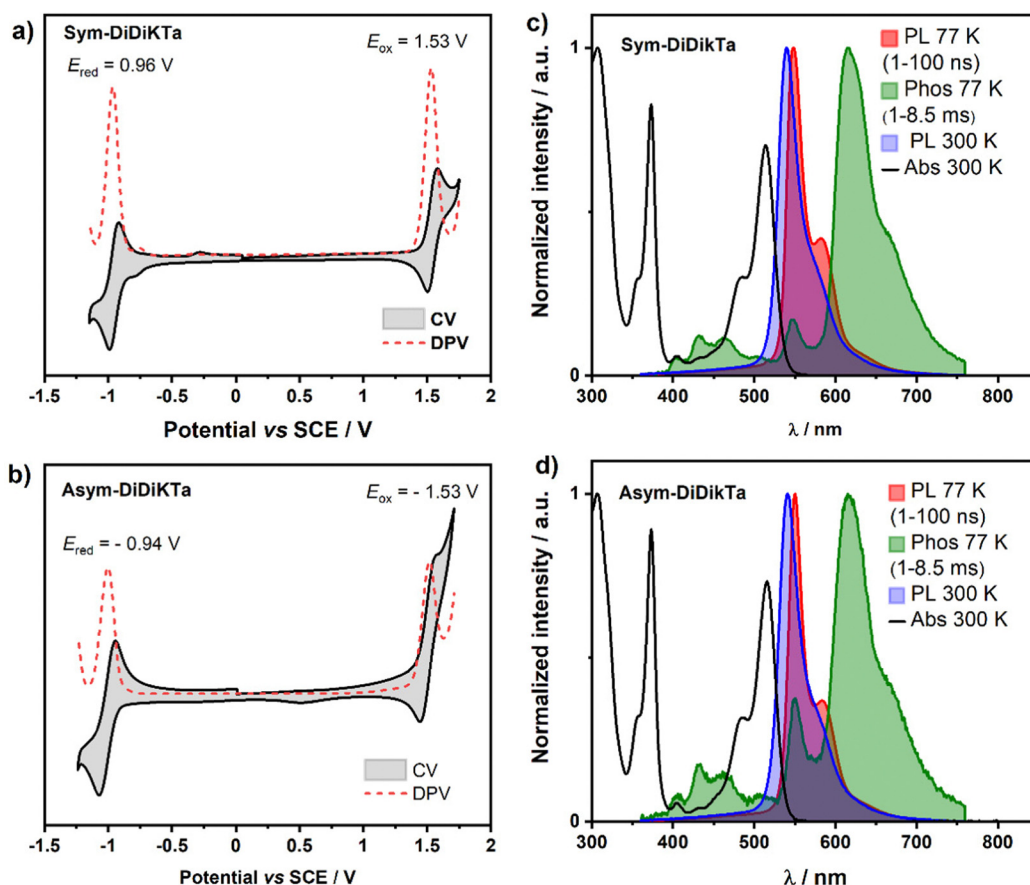


Fig. 4 Optoelectronic characterization of **Sym-DiDiKta** and **Asym-DiDiKta**: (a and b) cyclic and differential pulse voltammograms of **Sym-DiDiKta** and **Asym-DiDiKta**, respectively, in degassed CH<sub>2</sub>Cl<sub>2</sub> with 0.1 M [<sup>n</sup>Bu<sub>4</sub>N]PF<sub>6</sub> as the supporting electrolyte and Fc/Fc<sup>+</sup> as the internal reference (Fc/Fc<sup>+</sup> = 0.46 V vs SCE).<sup>47</sup> (c and d) Absorption (black line), steady-state PL spectra obtained in toluene at 300 K (blue line) and 77 K (red line; delay: 1 ns; gate time: 100 ns,  $\lambda_{exc}$  = 343 nm), and phosphorescence (Phos.; delay: 1 ms; gate time: 8.5 ms,  $\lambda_{exc}$  = 343 nm) spectra in toluene glass at 77 K (green olive line) of **Sym-DiDiKta** and **Asym-DiDiKta**, respectively.



Table 1 Optoelectronic properties of **Sym-DiDiKTa** and **Asym-DiDiKTa**

	In toluene							In film			
	$\lambda_{\text{abs}}^a/\text{nm}$	$\epsilon^a/\text{M}^{-1}\text{cm}^{-1}$	$\lambda_{\text{PL}}^b/\text{nm}$	$\Phi_{\text{PL}}^c$ in $\text{N}_2$ (air) $^c/\%$	FWHM $^d$ /nm (eV)	$S_1^e$ /eV	$T_1^e$ /eV	$\Delta E_{\text{ST}}^f$ /eV	$\lambda_{\text{PL}}/\text{nm}$	$\Phi_{\text{PL}}/\%$	FWHM $^d$ /nm (eV)
<b>Sym-DiDiKTa</b>	373/485/515	31900/11680/26325	540	69 (65)	29 (0.12)	2.26	2.02	0.24	542 $^g$	64	35 (0.14) $^g$
<b>Asym-DiDiKTa</b>	373/485/516	28527/10238/23458	541	53 (50)	29 (0.12)	2.25	2.02	0.23	547 $^g$	57	35 (0.14) $^g$

$^a$  UV-vis absorption of CT transition.  $^b$  Prompt emission in toluene degassing with  $\text{N}_2$ .  $^c$  Photoluminescence quantum yield in toluene relative to quinine sulfate in 1N  $\text{H}_2\text{SO}_4$  ( $\Phi_{\text{PL}} = 54.6\%$ ).  $^d$  Full-width at half-maximum.  $^e$  Obtained using an integrating sphere under  $\text{N}_2$ .  $^f$  Energy gap between  $S_1$  and  $T_1$  calculated from the difference of the peaks of the fluorescence and phosphorescence spectra in toluene glass at 77 K.  $^g$  1 wt% **Sym-DiDiKTa** and **Asym-DiDiKTa** doped in mCP.

energy levels were determined to be 2.26 and 2.25 eV, respectively, for **Sym-DiDiKTa** and **Asym-DiDiKTa** and the  $T_1$  excited state energy levels are identical at 2.02 eV, each obtained from the  $\lambda_{\text{PL}}$  of the respective prompt fluorescence and phosphorescence spectra in toluene glass at 77 K. We note that the nature of  $T_1$  and  $S_1$  are identical based on the difference density plots (Fig. 3). The corresponding  $\Delta E_{\text{ST}}$  values are 0.24 eV and 0.23 eV, respectively, for **Sym-DiDiKTa** and **Asym-DiDiKTa**. These values match with those predicted by SCS-CC2 calculations ( $\Delta E_{\text{ST}} = 0.24$  eV for both compounds). The PL quantum yield,  $\Phi_{\text{PL}}$ , in toluene is 70% under  $\text{N}_2$ , which decreases to 64% in air for **Sym-DiDiKTa** and 53% under  $\text{N}_2$  and 50% in air for **Asym-DiDiKTa**. The  $\Phi_{\text{PL}}$  values were next measured for vacuum-deposited 1 wt% doped thin films in 1,3-bis(*N*-carbazolyl)benzene (mCP). The  $\Phi_{\text{PL}}$  values were determined to be 64% and 57% for **Sym-DiDiKTa** and **Asym-DiDiKTa**, respectively (Table S5, ESI $^\ddagger$ ). The  $\Phi_{\text{PL}}$  values closely resemble those obtained in dilute toluene, which implies that non-

radiative decay due to vibrations is not significant in these compounds.

We next investigated the solid-state photophysical properties of spin-coated 1 wt% **Sym-DiDiKTa** and **Asym-DiDiKTa** doped thin films in mCP. The low doping concentration was selected to mitigate the potential for undesired aggregation in the films. As shown in Fig. 5a and c, at 300 K, **Sym-DiDiKTa** and **Asym-DiDiKTa** show a similar emission profile with  $\lambda_{\text{PL}}$  at 542 and 547 nm, respectively, values that are close to the prompt emission maximum in toluene. A small FWHM of 35 nm (0.13 eV) was calculated for both compounds in the films, which is slightly broader than the FWHM of 29 nm (0.12 eV) determined in toluene, indicating the influence of the matrix on the conformational stabilization of the excited states. Another aspect to consider are concentration-dependent aggregation effects since the concentration of emitters at 1 wt% doping in the film is higher than that of the  $10^{-6}$  M solution.

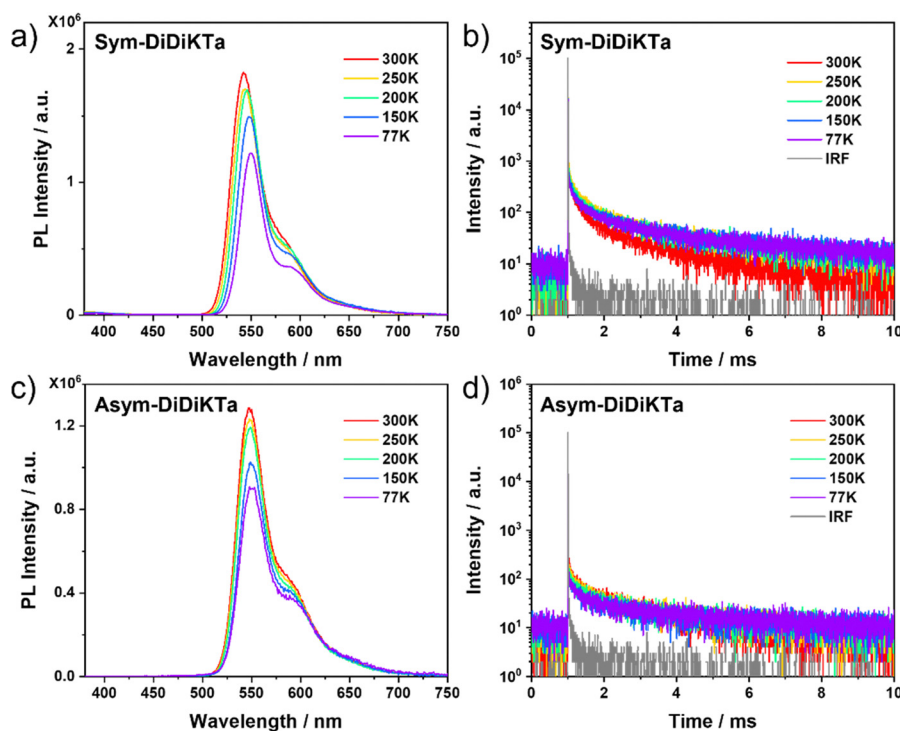


Fig. 5 (a and c) Temperature-dependent PL study of 1 wt% of **Sym-DiDiKTa** and **Asym-DiDiKTa**, respectively, in mCP,  $\lambda_{\text{exc}} = 350$  nm. (b and d) Temperature-dependent time-resolved PL decays for 1 wt% **Sym-DiDiKTa** and **Asym-DiDiKTa**, respectively, in mCP matrix ( $\lambda_{\text{exc}} = 378$  nm).



The average  $\tau_d$  values are 4.6 ms and 3.0 ms for **Sym-DiDiKTa** and **Asym-DiDiKTa**, respectively (Fig. 5 and Table S7, ESI†). We then compared the oxygen dependence of the PL spectrum at room temperature. The PL spectrum in air shows a small decrease in intensity compared to the spectrum under vacuum (Fig. S17a and b, ESI†), indicating a weak involvement of triplet excited states. The temperature-dependent PL spectra (Fig. 5a and c) and temperature-dependent time-resolved PL decays (Fig. 5b and d) of both emitters document the characteristic decrease in intensity and in the contribution of the delayed emission, respectively, with decreasing temperature. This could be due to the MR-TADF behavior only being apparent in a suitable host matrix due to exciplex-like host-emitter interactions.<sup>29</sup>

## OLED characterization

We fabricated vacuum-deposited OLEDs using **Sym-DiDiKTa** and **Asym-DiDiKTa** as the emitters. The following device configuration was used: indium tin oxide (ITO)-coated glass (100 nm)/TAPC (35 nm)/TCTA (10 nm)/mCP: 3 wt% of **Sym-DiDiKTa** or **Asym-DiDiKTa** (20 nm)/TmPyPB (30 nm)/LiQ (2 nm)/Al (100 nm). 1,1-Bis[(di-4-tolylamino)phenyl]cyclohexane (TAPC) is the hole-transporting layer, 4,4',4'-tris(carbazol-9-yl)triphenylamine (TCTA) is used for electron-blocking and host layers, 1,3,5-tris(3-pyridyl-3-phenyl)benzene (TmPyPB) is the electron-transporting layer, and lithium 8-hydroxyquinolinolate (LiQ) and aluminum are the electron injection and cathode layers, respectively. Fig. 6 and Table 2 summarize all the device characteristics. TADF-only devices based on **Sym-DiDiKTa** (Device I) and **Asym-DiDiKTa** (Device II) emitted green narrow-band electroluminescence,  $\lambda_{EL}$ , at 543 and 544 nm, respectively, values that are consistent to their corresponding  $\lambda_{PL}$ . The FWHMs of both Devices I and II are 36 nm, which resulted in CIE coordinates of (0.362, 0.623) and (0.376, 0.613), respectively. The  $EQE_{max}$  of Device I is 9.8%, which is slightly lower than the 10.5% measured for Device II. We hypothesized that the moderate  $\Phi_{PL}$  in the mCP-doped films limits the  $EQE_{max}$  values in Devices I and II. The moderate  $\Phi_{PL}$  may be due to the long-lived delayed fluorescence that permits a greater probability that non-

radiative decay processes such as triplet-polaron quenching and triplet-triplet annihilation will contribute to the decay of the excitons. Both devices suffered from severe efficiency roll-off issues, which in part are due to the long delayed fluorescence lifetimes of **Sym-DiDiKTa** and **Asym-DiDiKTa**. Moreover, the linear decrease in the EQE – current density curve indicates possible exciton-polaron annihilation processes, which can be ascribed to trap formation by the emitters.

To improve the device performance, we implemented a hyperfluorescence (HF) strategy in Device III with the following device structure: indium tin oxide (ITO)-coated glass (100 nm)/HAT-CN (10 nm)/Tris-PCz (30 nm)/mCBP (5 nm)/mCBP: 20 wt% 4CzIPN: 3 wt% **Asym-DiDiKTa** (30 nm)/T2T (10 nm)/BPy-TP2 (40 nm)/LiQ (2 nm)/Al (100 nm), where 1,4,5,8,9,11-hexaazatriphenyl-enehexacarbonitrile (HAT-CN) is the hole-injection layer, 9-phenyl-3,6-bis(9-phenyl-9H-carbazol-3-yl)-9H-carbazole (Tris-PCz) is the hole-transporting layer, mCBP is used in the exciton-blocking and host layers, 1,2,3,5-tetrakis(carbazol-9-yl)-4,6-dicyanobenzene (4CzIPN) is a green TADF assistant dopant, 2,4,6-tris(biphenyl-3-yl)-1,3,5-triazine (T2T) is the hole-blocking layer and 2,7-di(2,2'-bipyridin-5-yl)triphenylene (BPy-TP2) is the electron-transporting layer, and lithium 8-hydroxyquinolinolate (LiQ) and Al are the electron injection and cathode layers, respectively. Gratifyingly, the device performance improved and an  $EQE_{max}$  of 19.9% was achieved along with an enhanced maximum brightness of 53625 cd m<sup>-2</sup>. The HF device not only resulted in a better EQE, but also reduced the efficiency roll-off when compared to Device II. The  $\lambda_{EL}$  is red-shifted to 548 nm and there is a slightly larger FWHM of 56 nm. The broader FWHM may originate from an incomplete energy transfer from 4CzIPN to **Asym-DiDiKTa**.

## LEC devices

The light-emitting electrochemical cell (LEC) can be comprised of solely air-stable materials and feature a very simple and robust device structure in the form of a single-layer active material sandwiched between two electrodes. This renders the LEC technology highly fit for scalable and cost-efficient printing and coating fabrication under ambient air.<sup>49,50</sup> The characteristic feature

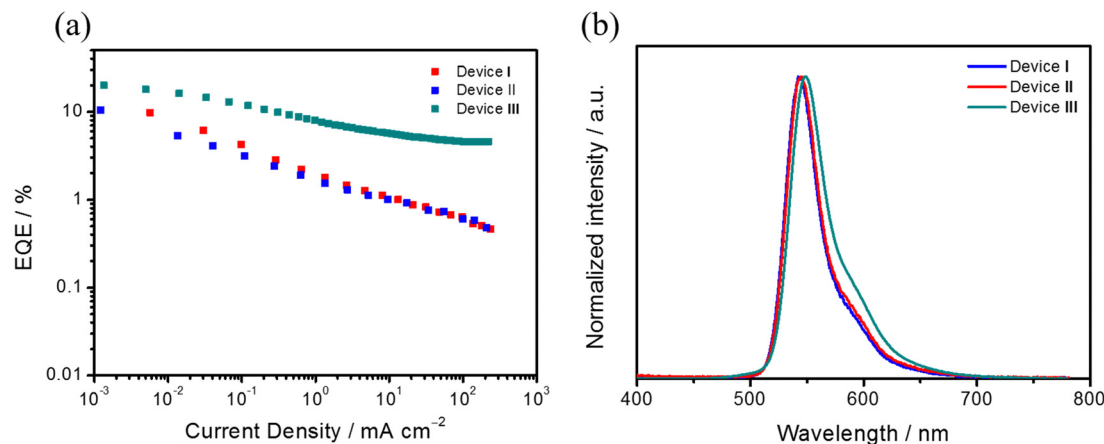


Fig. 6 (a) EQE versus current density curves of Devices I–III. (b) Electroluminescence spectra of Devices I–III at 500 cd m<sup>-2</sup>.



Table 2 Summary of device characteristics

Device	Dopant	$V_{\text{on}}/V^a$	EQE/% <sup>b</sup>	$\lambda_{\text{EL}}/\text{nm}^c$	Max. brightness ( $\text{cd m}^{-2}$ ) <sup>d</sup>	FWHM/nm <sup>c</sup>	CIE (x, y) <sup>c</sup>
I	3 wt% <b>Sym-DiDiKTa</b>	3.5	9.8/1.8/0.8	543	4310	36	(0.359, 0.624)
II	3 wt% <b>Asym-DiDiKTa</b>	4	10.5/1.3/0.8	544	3989	36	(0.369, 0.616)
III	3 wt% <b>Asym-DiDiKTa</b> : 20 wt% <b>4CzIPN</b>	3.2	19.9/9.9/6.4	548	53625	56	(0.397, 0.592)

<sup>a</sup> Voltage at 1  $\text{cd m}^{-2}$ . <sup>b</sup> Value at maximum, 100  $\text{cd m}^{-2}$  and 1000  $\text{cd m}^{-2}$ . <sup>c</sup> Value at 500  $\text{cd m}^{-2}$ . <sup>d</sup> Maximum brightness.

of LEC devices is the combination of mobile ions with the emissive organic semiconductor within the emissive layer.<sup>51,52</sup> These mobile ions redistribute when a voltage is applied, and enable p-type electrochemical doping of the organic semiconductor at the anode and n-type doping at the cathode. With time, these doping regions grow in size and make contact under the formation of a p-n junction. The fact that **Sym-DiDiKTa** displays highly reversible electrochemical oxidation and reduction behaviour in the cyclic voltammetry experiments (Fig. 4a) suggests that it could be fit for the task of the emissive organic semiconductor in a LEC device.

We fabricated and characterized LEC devices with the following configuration: indium tin oxide (140 nm)/poly(3,4-ethylenedioxythiophene) polystyrene sulfonate (40 nm)/26DCzPPy (44 wt%):POT2T (44 wt%):**Sym-DiDiKTa** (4 wt%):THABF<sub>4</sub> (8 wt%) (100 nm)/Al (100 nm), where THABF<sub>4</sub> (tetrahexylammonium tetrafluoroborate) is the ionic liquid electrolyte that contributes the mobile ions, and 6-bis(3-carbazol-9-yl)pyridine (26DCzPPy):(1,3,5-triazine-2,4,6-triyl)tris(benzene-3,1-diyl)tris(diphenylphosphine oxide) (POT2T) is a blend-host matrix, which was introduced to enable for the formation of a uniform solution-processed thin film and to suppress losses by exciton-polaron<sup>53</sup> and exciton-exciton quenching.

Fig. 7a presents the steady-state EL spectrum recorded during the driving with a constant current density of 77  $\text{mA cm}^{-2}$ , with the  $\lambda_{\text{EL}}$  at 551 nm and the FWHM being 60 nm. The broadening of the EL spectrum of the LEC device in comparison to the PL spectra in Fig. 5a indicates the formation of exciplexes with the blend host. This conclusion is supported by the fact that the FWHM of the EL spectrum is essentially independent of the guest concentration

for a guest concentration range of 1 to 8 wt% (Fig. S22, ESI†). Fig. 7b details the voltage and luminance transients recorded during the early stages of the constant-current driving. The observed initial decrease of the voltage is a characteristic indicator of conductivity-enhancing electrochemical doping, whereas the increase in luminance is in line with the gradual formation of a p-n junction, where electrons and holes can recombine efficiently into excitons. Importantly, these observations imply that the **Sym-DiDiKTa** emitter can be *in situ* electrochemically p- and n-type doped during LEC operation. This conclusion is further supported by the fact that the LEC device delivered a significant luminance of 300  $\text{cd m}^{-2}$  despite being equipped with an air-stable Al cathode in direct contact with the active emissive material. Finally, the luminance turn-on time is a direct indicator of the ion mobility in the active material,<sup>54,55</sup> and the comparatively fast turn-on time of less than 2 s to 100  $\text{cd m}^{-2}$  demonstrates that the ion mobility in the active material is high.

## Conclusion

In summary, new amine/carbonyl-based MR-TADF materials were developed by an approach that expands the  $\pi$ -conjugated backbone to show green-yellow emission. The regio-functionalization of the *tert*-butyl groups was shown to significantly impact the single-crystal packing structure. With a  $\Delta E_{\text{ST}}$  of around 0.23 eV in toluene, the narrowband emitters, FWHM, of 29 nm, display TADF activity with a  $\tau_d$  of 4.6 ms and 3.0 ms for **Sym-DiDiKTa** and **Asym-DiDiKTa**, respectively. Application

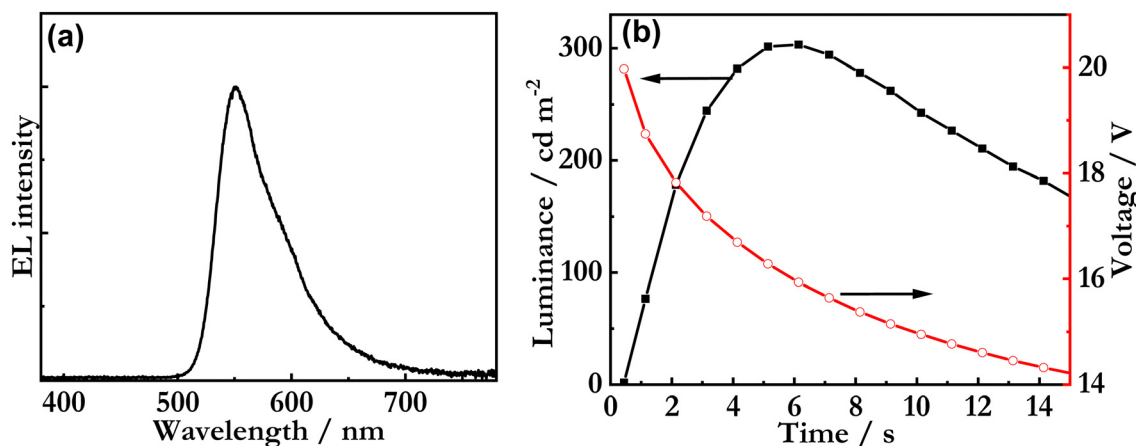


Fig. 7 (a) The steady-state EL spectrum. (b) The temporal evolution of the luminance (left y-axis, solid black squares) and the drive voltage (right y-axis, open red circles) for the ITO/PEDOT:PSS/26DCzPPy:POT2T:**Sym-DiDiKTa**:THABF<sub>4</sub>/Al LEC device. The LEC devices were driven by a constant current density of 77  $\text{mA cm}^{-2}$ .





in OLEDs provided devices with electroluminescence maxima at around 544 nm, and an EQE<sub>max</sub> of *circa* 10% which low value is attributed to the long delayed lifetimes of this class. Upon further studies in hyperfluorescence devices, an EQE of 19.9% was finally obtained, showing the promise of this class of molecules towards this strategy. We also show the first examples of a LEC device incorporating an MR-TADF emitter. With a  $\lambda_{\text{EL}}$  at 551 nm and a larger FWHM of 60 nm due to formation of exciplexes with the host, the device based on **Sym-DiDiKta** delivered a significant luminance of 300 cd m<sup>-2</sup> and high ion mobility with a fast turn-on time of less than 2 s to 100 cd m<sup>-2</sup>.

## Conflicts of interest

The authors declare that they have no conflict of interest.

## Acknowledgements

The St Andrews team would like to thank EPSRC (EP/P010482/1) and the Leverhulme Trust (RPG-2016-047) for financial support. Computational resources have been provided by the Consortium des Équipements de Calcul Intensif (CÉCI), funded by the Fonds de la Recherche Scientifiques de Belgique (F. R. S.-FNRS) under Grant no. 2.5020.11. Y. O. acknowledges funding by the Fonds de la Recherche Scientifique-FNRS under Grant no. F.4534.21 (MIS-IMAGINE). The authors also acknowledge Ms N. Nakamura and Ms K. Kusuhara for their technical assistance with this research. This work was supported financially by the JSPS Core-to-Core Program (grant number: JPJSCCA20180005), Kyulux Inc, the Swedish Research Council, the Swedish Energy Agency, and Bertil & Britt Svenssons stiftelse för belysningsteknik.

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