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# Nonplanar structure accelerates reverse intersystem crossing of TADF emitters: nearly 40% EQE and relieved efficiency roll off†

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Exploring strategies to enhance reverse intersystem crossing (RISC) is of great significance to develop efficient thermally activated delayed fluorescent (TADF) molecules. In this study, we investigate the substantial impact of nonplanar structure on improving the rate of RISC ( $k_{\text{RISC}}$ ). Three emitters based on spiroacridine donors are developed to evaluate this hypothesis. All molecules exhibit high photoluminescent quantum yields (PLQYs) of 96–98% due to their rigid donor and acceptor. Leveraging the synergistic effects of heavy element effect and nonplanar geometry, S2-TRZ exhibits an accelerated  $k_{\text{RISC}}$  of  $24.2 \times 10^5 \text{ s}^{-1}$  compared to the  $11.1 \times 10^5 \text{ s}^{-1}$  of S1-TRZ, which solely incorporates heavy atoms. Additionally, O1-TRZ possesses a further lower  $k_{\text{RISC}}$  of  $9.42 \times 10^5 \text{ s}^{-1}$  because of the absence of these effects. Remarkably, owing to the high PLQYs and suitable TADF behaviors, devices based on these emitters exhibit state-of-the-art performance, including a maximum external quantum efficiency of up to 40.1% and maximum current efficiency of  $124.7 \text{ cd A}^{-1}$ . More importantly, devices utilizing S2-TRZ as an emitter achieve a relieved efficiency roll-off of only 7% under  $1000 \text{ cd m}^{-2}$ , in contrast to the 12% for O1-TRZ and 11% for S1-TRZ, respectively. These findings advance our fundamental understanding of TADF processes for high-performance electroluminescent devices.

## 1. Introduction

In the last few decades, thermally activated delayed fluorescent (TADF) emitters have emerged as highly promising materials for organic light emitting diodes (OLEDs) owing to their capability of boosting 75% triplet excitons to emit *via* reverse intersystem crossing (RISC) without the help of noble metals.<sup>1,2</sup> The advent of multi-resonance (MR) type emitters, characterized by narrow band emission, has positioned them as a burgeoning option aligning well with BT.2020 standards, a wide color gamut standard for the next generation of high resolution displays.<sup>3–7</sup> Nevertheless, MR emitters usually possess prolonged delayed fluorescence due to the large energy difference ( $\Delta E_{\text{ST}}$ ) between the first singlet ( $S_1$ ) and triplet excited state ( $T_1$ ), which impedes swift RISC processes. In response, sensitization techniques have been applied to augment the efficiency and durability of MR-emitter-based devices.<sup>8,9</sup> These sensitizers play a critical role in effectively recycling all excitons, thereby offering a new opportunity for leveraging conventional donor–acceptor (D–A)

type TADF compounds. Developing efficient D–A TADF emitters remains a paramount objective for realizing high performance OLEDs.

The primary task to craft an efficient TADF emitter lies in establishing a rapid RISC pathway, which necessitates a minimized  $\Delta E_{\text{ST}}$  and a large spin–orbit coupling (SOC), as delineated by Fermi's golden rule.<sup>10–12</sup> Various structural motifs, such as twisted D–A arrangements,<sup>13–19</sup> through-space charge transfer (TSCT),<sup>20–24</sup> or multi-resonance (MR) architectures,<sup>25–31</sup> have successfully achieved diminutive  $\Delta E_{\text{ST}}$  in pristine organic compounds. Indeed, the attainment of near-zero  $\Delta E_{\text{ST}}$  has been achieved in several works, which propelled the rate of RISC ( $k_{\text{RISC}}$ ) to a maximum of  $10^8 \text{ s}^{-1}$ .<sup>32</sup> However, conventional TADF emitters encounter an intrinsic limitation in adjusting  $k_{\text{RISC}}$  solely through  $\Delta E_{\text{ST}}$  manipulation, as its minimum value is constrained to zero. To further enhance  $k_{\text{RISC}}$ , attention must be directed to optimizing SOC. Hence, heavy elements such as sulfur (S),<sup>33–37</sup> selenium (Se),<sup>38–42</sup> or noble metals<sup>43–46</sup> have been incorporated, which elevate the  $k_{\text{RISC}}$  value and mitigate efficiency roll-off in devices. Notably, the introduction of heavy atoms will amplify the rate of intersystem crossing ( $k_{\text{ISC}}$ ) simultaneously, which will compete with the radiative decay process and potentially compromise photoluminescent quantum yield (PLQY). Therefore, despite the advantage of Se, its utilization in D–A type TADF compounds remains limited due to the inherent suppression of radiative decay caused by the

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diminished overlap between frontier molecular orbitals (FMOs). Exploring alternative strategies to enhance SOC represents a promising avenue for the design of efficient TADF emitters.

Conformational twisting has long been recognized to trigger SOC in a conjugated  $\pi$ -electron system.<sup>47</sup> Nonplanar molecules usually possess a larger  $k_{\text{RISC}}$  than their planar counterparts. The spin-orbit Hamiltonian in central-field approximation could be written as<sup>48</sup>

$$\hat{H}_{\text{SO}} = \alpha_{\text{F}}^2 \sum_{\mu}^N \sum_i^n \frac{Z_{\mu}}{r_{i\mu}^3} \vec{L}_i \vec{S}_i \quad (1)$$

where  $\alpha_{\text{F}}$  represents the fine-structure constant.  $Z_{\mu}$  denotes the effective nuclear charge at nucleus  $\mu$ , and  $S$  and  $L$  are the spin and orbital angular momenta of electron  $i$ , respectively. Notably, in coplanar molecules, the SOC between  $\pi$ - $\pi^*$  orbitals yields relatively small values, whereas the 'out-of-plane' component  $L_z$  of the angular momentum operator  $L$  contributes to a non-negligible SOC. This phenomenon has been observed in polymer systems as well.<sup>49</sup> Recently, twisting geometry has been applied in boosting phosphorescence emission in pure organic molecules.<sup>50–53</sup> SOC could be effectively improved by a nonplanar structure. Additionally, You *et al.* reported that the nonplanar benzothiophene could accelerate  $k_{\text{RISC}}$  by reducing the  $\Delta E_{\text{ST}}$ .<sup>54</sup> Enlightened by these valuable experiences, the

exploration of such strategies will provide a new perspective in assembling practical TADF emitters.

Herein, to validate this hypothesis, we designed and synthesized a new emitter, named 10-(4-(4,6-diphenyl-1,3,5-triazin-2-yl)phenyl)-2,7-dimethyl-10*H*-spiro[acridine-9,12'-benzo-*b*]fluoreno[1,2-*d*]thiophene] (**S2-TRZ**), with a properly bent acridine (Fig. 1). The structurally fixed benzothiophene unit provides a substantial steric hindrance between the hydrogen atom and the acridine plane. Hence, a hooked acridine with a dihedral angle of 166° was acquired. In comparison, an isomer, 10-(4-(4,6-diphenyl-1,3,5-triazin-2-yl)phenyl)-2,7-dimethyl-10*H*-spiro[acridine-9,7'-benzo[*b*]fluoreno-[3,2-*d*]thiophene] (**S1-TRZ**), with the same benzothiophene substitution but a more planar acridine was assembled. Additionally, the oxygen (O) equipped analogs of **S1-TRZ**, 10-(4-(4,6-diphenyl-1,3,5-triazin-2-yl)phenyl)-2,7-dimethyl-10*H*-spiro[acridine-9,7'-fluoreno[2,3-*b*]benzofuran] (**O1-TRZ**), was also forged to unveil the effect of heavy atoms. All emitters exhibit high PLQY of 96–98%, relevant with the rigid D and A units. Benefiting from the synergy of the heavy element effect and nonplanar donor structure, **S2-TRZ** possesses the highest  $k_{\text{RISC}}$  of  $24.2 \times 10^5 \text{ s}^{-1}$ . Without the nonplanar structure, **S1-TRZ** with a solely heavy element effect holds a moderate  $k_{\text{RISC}}$  of  $11.1 \times 10^5 \text{ s}^{-1}$ . Furthermore, lacking the nonplanar structure of the sulfur-

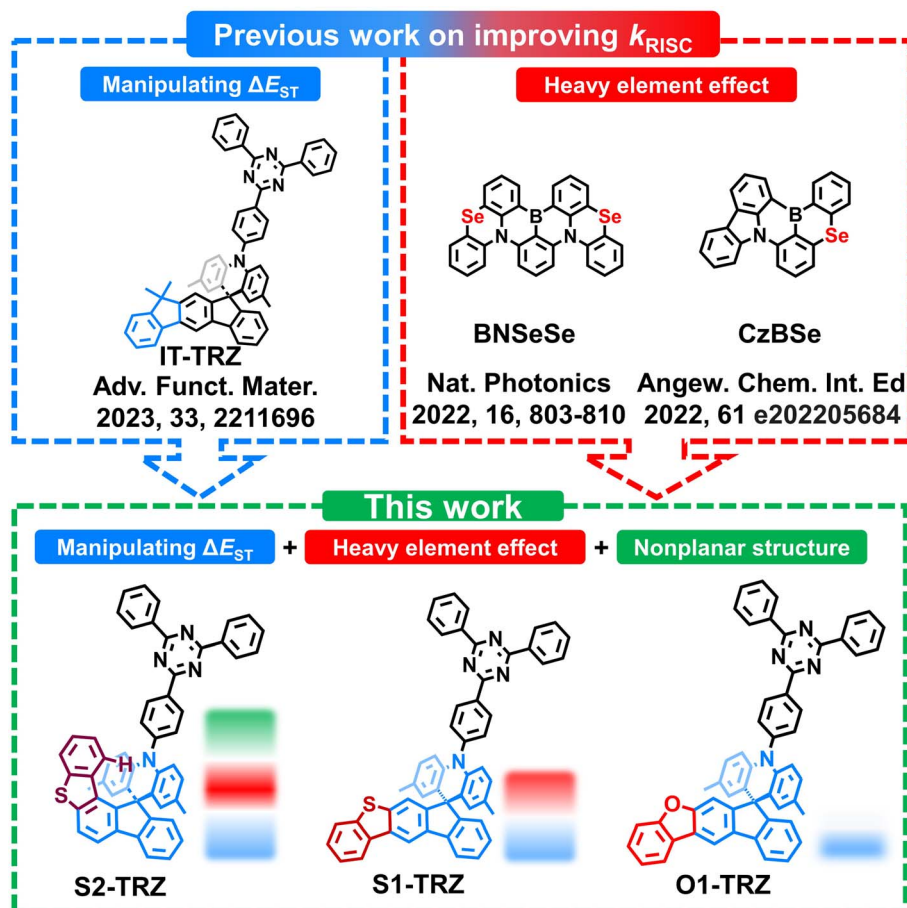


Fig. 1 Comparison of the reported work on the methods of enhancing  $k_{\text{RISC}}$ , molecular design and chemical structures of the related emitters.



containing analogs, **O1-TRZ** displays a lower  $k_{\text{RISC}}$  of  $9.42 \times 10^5 \text{ s}^{-1}$ . Notably, electroluminescent devices based on these emitters achieved a state-of-the-art external quantum efficiency (EQE) of 39.6% for **S2-TRZ**, 39.2% for **S1-TRZ** and even 40.1% for **O1-TRZ** thanks to the exceptional molecular properties. More importantly, **S2-TRZ** enables a significantly reduced efficiency roll-off of only 7% under  $1000 \text{ cd m}^{-2}$ , outperforming both **S1-TRZ** (11%) and **O1-TRZ** (12%).

## 2. Results and discussion

### 2.1. Molecular design, synthesis and thermal properties

To validate the impact of nonplanar structures on  $k_{\text{RISC}}$ , a delicate molecular design is imperative. Nonplanar motifs within D or A groups may affect their electron-donating or -withdrawing capabilities, potentially hindering charge transfer (CT) interactions. In some cases, such motifs could lead to unfavorable conformations, such as the quasi-axial (QA) geometry, which is generally recognized to result in less efficient TADF properties. Thus, there arises a need for a building block composed of a fixed geometry with controllable modulation. Enlightened by our previous works, spiro-acridines featuring a rigid crisscross geometry could be an excellent platform to establish a model compound. To further reveal the insights for manipulating the

TADF properties, the heavy element effect is also taken into consideration. Therefore, **S2-TRZ**, **S1-TRZ** and **O1-TRZ** are designed. All the compounds were synthesized *via* conventional Buchwald–Hartwig coupling between the bromo triazine acceptor and corresponding acridine donors (Scheme S1†). The donor parts were synthesized using a previously reported one-pot reaction method in good yields.<sup>55</sup> All the final compounds were identified by  $^1\text{H}$  NMR,  $^{13}\text{C}$  NMR, mass spectra and elemental analysis. Excellent thermal stability is demonstrated with decomposition temperatures of 390–445 °C as evidenced by thermal gravimetric analysis (TGA) measurements (Fig. S1†). No obvious glass transition temperatures ( $T_g$ s) are detected in the range of 50–400 °C.

### 2.2. Theoretical simulation

To examine the properties of ground and excited states, density functional theory (DFT) and time dependent DFT (TD-DFT) were performed at the PBE0/def2svp level. As shown in Fig. 2 and S2,† **S1-TRZ** and **O1-TRZ** hold a more planar acridine unit with a dihedral angle of 173° and 177°. Conversely, **S2-TRZ** exhibits a twisted acridine with a dihedral angle of 166°, which is attributed to the spatially close hydrogen atom forming a constrained interaction with the acridine plane. These interactions could be identified using reduced density gradient

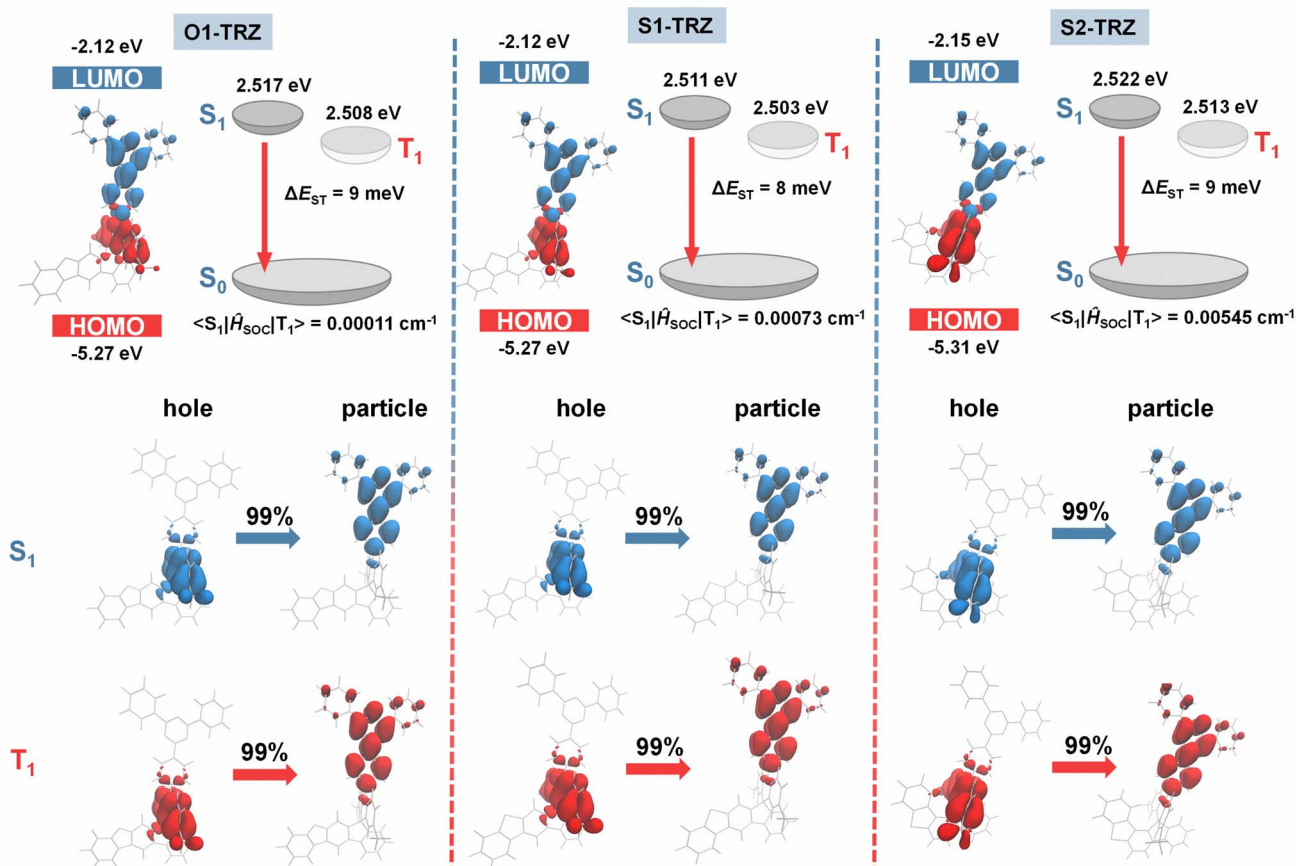


Fig. 2 Theoretical simulated distribution and energy levels of HOMO/LUMO, and NTO distribution of the particle/hole of  $S_1$  and  $T_1$  of **O1-TRZ**, **S1-TRZ** and **S2-TRZ** at the PBE0/def2svp level. The figures are extracted using Multiwfn.<sup>56</sup>



(RDG) analysis (Fig. S3†). Strong van der Waals interactions (green isosurface) and large hindrance (brown isosurface) are clearly presented. Despite the enforced bending of acridine in **S2-TRZ**, the rigid crisscross structure prevents excessive deformation, favoring a quasi-equatorial (QE) geometry. In contrast, both **S1-TRZ** and **O1-TRZ** exhibit predictable QE patterns. In return, all emitters show large twisting between D and A with torsion angles of 85° for **S2-TRZ**, 88° for **S1-TRZ** and 87° for **O1-TRZ**, respectively. Hence, well separated highest occupied molecular orbitals (HOMOs) and lowest unoccupied molecular orbitals (LUMOs) are established in all three molecules. LUMOs predominantly reside on triazine with similar energy levels of -2.15 eV for **S2-TRZ** and -2.12 eV for both **S1-TRZ** and **O1-TRZ**, while HOMOs mostly spread on acridine units with comparable values of -5.27 eV for both **S1-TRZ** and **O1-TRZ**, and -5.31 eV for **S2-TRZ**. All  $S_1$  and  $T_1$  of the three compounds exhibit CT features as revealed by the distribution of their natural transition orbitals (NTOs). Taking **S2-TRZ** for example, 'hole' NTOs localized on acridine with a small portion on the spiro-junction, while 'particle' NTOs extend over the acceptor unit. The good separation of 'hole' and 'particle' NTOs suggests small  $\Delta E_{ST}$ , leading to predicted values of 9 meV for **S2-TRZ**, 8 meV for **S1-TRZ** and 9 meV for **O1-TRZ**, respectively, which may facilitate a good TADF feature. To distinguish the effect of the hooked donor, the SOC constant was calculated to be 0.00545  $\text{cm}^{-1}$  for **S2-TRZ**, 0.00073  $\text{cm}^{-1}$  for **S1-TRZ** and 0.00011  $\text{cm}^{-1}$  for **O1-TRZ**, respectively (Table 1). **S1-TRZ** exhibits a slightly higher SOC than **O1-TRZ**, hinting at a weak heavy element effect. In sharp comparison, **S2-TRZ** manifests a much larger SOC value than that of **S1-TRZ**, highlighting the pronounced impact of nonplanar structure in enhancing SOC. Theoretical simulation suggests that these emitters could exhibit efficient TADF properties, and more encouragingly, the nonplanar donor could provide an effective enhancement on SOC.

### 2.3. Photophysical properties

Photophysical properties were assessed in dilute toluene solution ( $10^{-5}$  M) and doped films (10 wt% in dibenzo[*b,d*]furan-2,8-diylbis(diphenylphosphine oxide) (PPF)). As shown in Fig. 3, the intense absorption below 320 nm is assigned to the  $\pi$ - $\pi^*$  transition. The peaks between 320 and 370 nm are attributed to the  $n$ - $\pi^*$  transition of the donor. Different from **O1-TRZ** and **S1-TRZ** that exhibit structural and sharp peaks, **S2-TRZ** shows a broadened absorption in this region. To gain a deep insight into this phenomenon, we have recorded the absorption of the donor unit. As shown in Fig. S5†, the  $n$ - $\pi^*$  transition of these emitters fits well with the absorption of their donors, which confirms that these differences are related to the donor. It is reported that the molecular absorption becomes structureless

and wide as the rigidity and planarity decrease.<sup>57</sup> Therefore, the broadened  $n$ - $\pi^*$  transition of **S2-TRZ** could probably be attributed to the decrease of the planarity and rigidity of the acridine part, which confirms the bending of the acridine as revealed by theoretical analysis. The weak peaks at 410 nm originate from the CT absorption, which suggests the tight correlation between the D and A units. Structureless blueish green emissions around 504–513 nm are observed in doped films. The energy levels of  $S_1$  ( $E_{S_1}$ )/ $T_1$  ( $E_{T_1}$ ) are calculated to be 2.72/2.67 eV for **O1-TRZ**, 2.69/2.63 eV for **S1-TRZ** and 2.73/2.67 eV for **S2-TRZ**, resulting in small  $\Delta E_{ST}$  values of 0.05, 0.06 and 0.06 eV, respectively. Moreover, FI of these emitters was also recorded at room temperature. As shown in Fig. S4,† **O1-TRZ** and **S1-TRZ** show slightly redshifted spectra and smaller  $E_{S_1}$  of 2.66 eV for **O1-TRZ** and 2.67 eV for **S1-TRZ**, respectively. Meanwhile, **S2-TRZ** exhibits the same  $E_{S_1}$  at room temperature and 77 K, indicating **S2-TRZ** possesses less relaxation of  $S_1$  in films. Proper TADF is corroborated *via* transient PL measurement, revealing distinct prompt (PF) and DF components in the decay profiles (Fig. 3b). Fitting well with two-exponential decay, lifetimes of PF ( $\tau_p$ )/DF ( $\tau_d$ ) are deduced to be 28 ns/2.1  $\mu$ s, for **O1-TRZ**, 24 ns/1.7  $\mu$ s for **S1-TRZ** and 25 ns/2.3  $\mu$ s for **S2-TRZ**, respectively. And the ratios of PF/DF ( $R_p/R_d$ ) are 48%/52% for **O1-TRZ**, 49%/51% for **S1-TRZ** and 15%/82% for **S2-TRZ**. The higher  $R_p/R_d$  may be attributed to the higher SOC value of **S2-TRZ**, which endows it with fast ISC and RISC processes. ISC competes with radiative decay and leads to more singlet excitons converting to triplet ones and enlarges the component of DF. Additionally, all emitters show high PLQY ranging from 96 to 98%. Rate constants are further calculated using the equations provided in the ESI† and summarized in Table 2. Compared to **O1-TRZ**, **S1-TRZ** exhibits a shortened  $\tau_p$  and  $\tau_d$ , demonstrating higher  $k_r$  of  $1.95 \times 10^7 \text{ s}^{-1}$  and  $k_{RISC}$  of  $11.1 \times 10^5 \text{ s}^{-1}$ , respectively. This enhancement could be related to the introduction of S. Since S is introduced far from the D–A center, the heavy element effect could be weak as predicted by theoretical simulation. In contrast, **S2-TRZ** displays a DF-dominated TADF feature with  $R_p/R_d$  of 18%/82%, leading to significantly improved  $k_{RISC}$  of  $24.2 \times 10^5 \text{ s}^{-1}$ . This substantial difference is attributed to the formation of the nonplanar donor, underscoring the significance of structural factors in TADF emitters. The HOMO energy levels of these compounds were determined using cyclic voltammetry (CV) and calculated to be 5.17 eV for **S2-TRZ**, 5.18 eV for **S1-TRZ** and 5.20 eV for **O1-TRZ**, respectively, suggesting that the nonplanar donor show minimal impact on electron donating ability. Thus, the enhanced  $k_{RISC}$  is mainly contributed by the bended structure. Photophysical analysis confirms the formation of a nonplanar donor and the accelerated RISC channels as suggested by theoretical simulation.

Table 1 Theoretical results and thermal properties of **S2-TRZ**, **S1-TRZ** and **O1-TRZ**

Compound	HOMO (eV)	LUMO (eV)	$S_1$ (eV)	$T_1$ (eV)	$\Delta E_{ST}$ (meV)	SOC ( $\text{cm}^{-1}$ )	$T_d$ ( $^\circ\text{C}$ )
<b>S2-TRZ</b>	-5.31	-2.15	2.522	2.513	9	0.00545	445
<b>S1-TRZ</b>	-5.27	-2.12	2.511	2.503	8	0.00073	404
<b>O1-TRZ</b>	-5.27	-2.12	2.517	2.508	9	0.00011	390



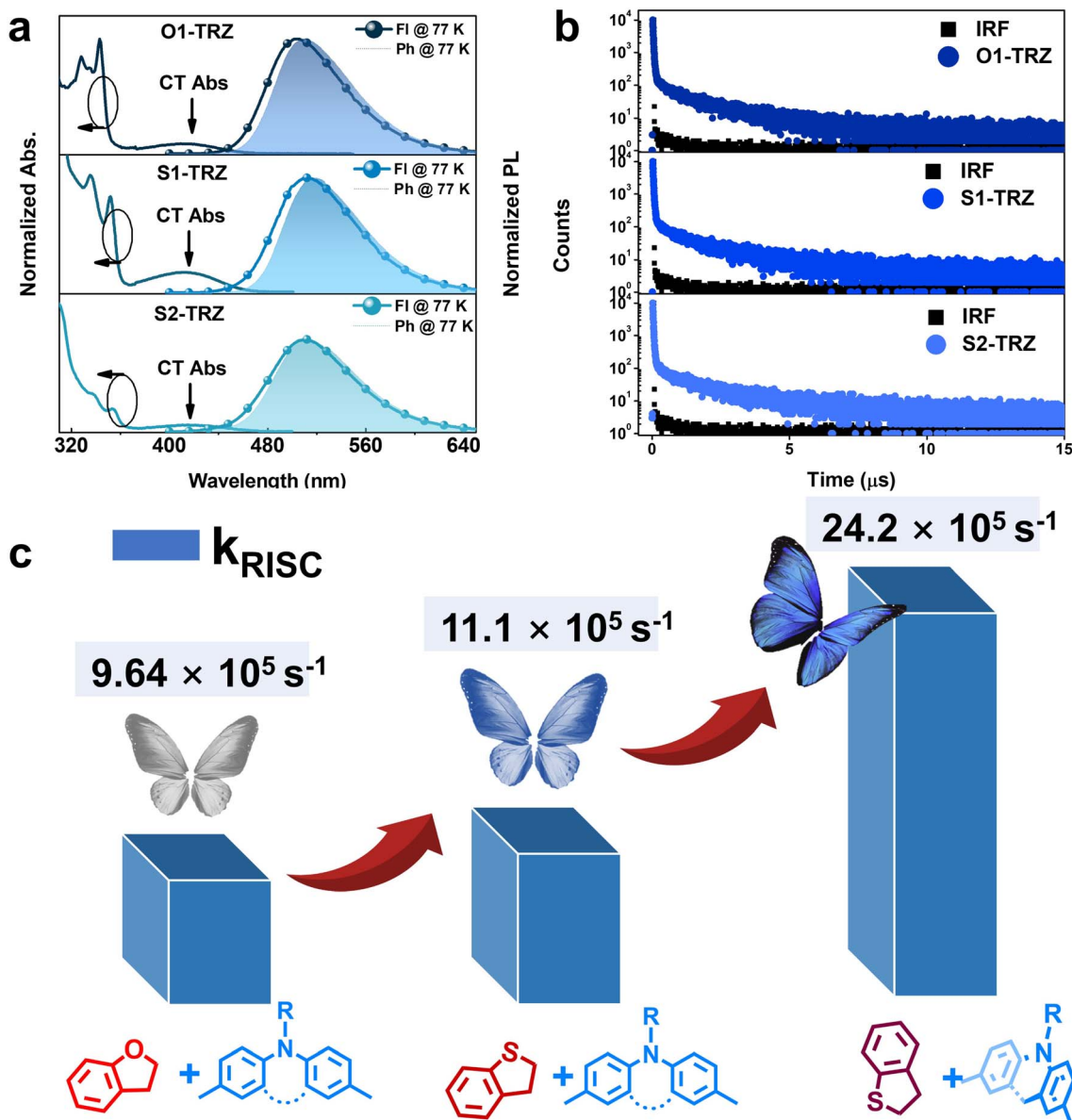


Fig. 3 (a) Absorption in toluene ( $10^{-5}$  M), fluorescence (FI) and phosphorescence (Ph) at 77 K in doped films (10 wt% in PPF) of O1-TRZ, S1-TRZ and S2-TRZ. (b) Transient PL decay in doped films (10 wt% in PPF) of O1-TRZ, S1-TRZ and S2-TRZ. (c) Comparison of  $k_{\text{RISC}}$  and structure characteristics of O1-TRZ, S1-TRZ and S2-TRZ.

Table 2 TADF properties of O1-TRZ, S1-TRZ and S2-TRZ in doped films (10 wt% in PPF)

Emitter	$\tau_p$ (ns)	$\tau_d$ ( $\mu\text{s}$ )	$\Phi_{\text{PL}}$ (%)	$\Phi_p$ (%)	$\Phi_d$ (%)	$k_r$ ( $10^7 \text{ s}^{-1}$ )	$k_{\text{ISC}}$ ( $10^7 \text{ s}^{-1}$ )	$k_{\text{RISC}}$ ( $10^5 \text{ s}^{-1}$ )	$k_{\text{nr}}$ ( $10^5 \text{ s}^{-1}$ )
O1-TRZ	28	2.1	98	47	51	1.69	1.87	9.64	3.45
S1-TRZ	24	1.7	96	47	49	1.95	2.09	11.1	8.03
S2-TRZ	25	2.3	97	17	80	0.70	3.27	24.2	2.15

#### 2.4. Electroluminescent properties

To evaluate the impact of such variation on electroluminescent (EL) properties, devices based on following structures of ITO/HATCN (5 nm)/TAPC (30 nm)/TCTA (15 nm)/mCBP (10 nm)/PPF:emitters (10–20 wt%, 25 nm)/PPF (10 nm)/ANT-BIZ (30

nm)/Liq (2 nm)/Al were fabricated. The corresponding EL profiles are listed in Fig. 4, S6<sup>†</sup> and Table 3. Initially, these emitters were examined at a doping concentration of 10 wt% to reduce the concentration-induced quenching. Single EL emission around 504–513 nm suggests the good confinement of



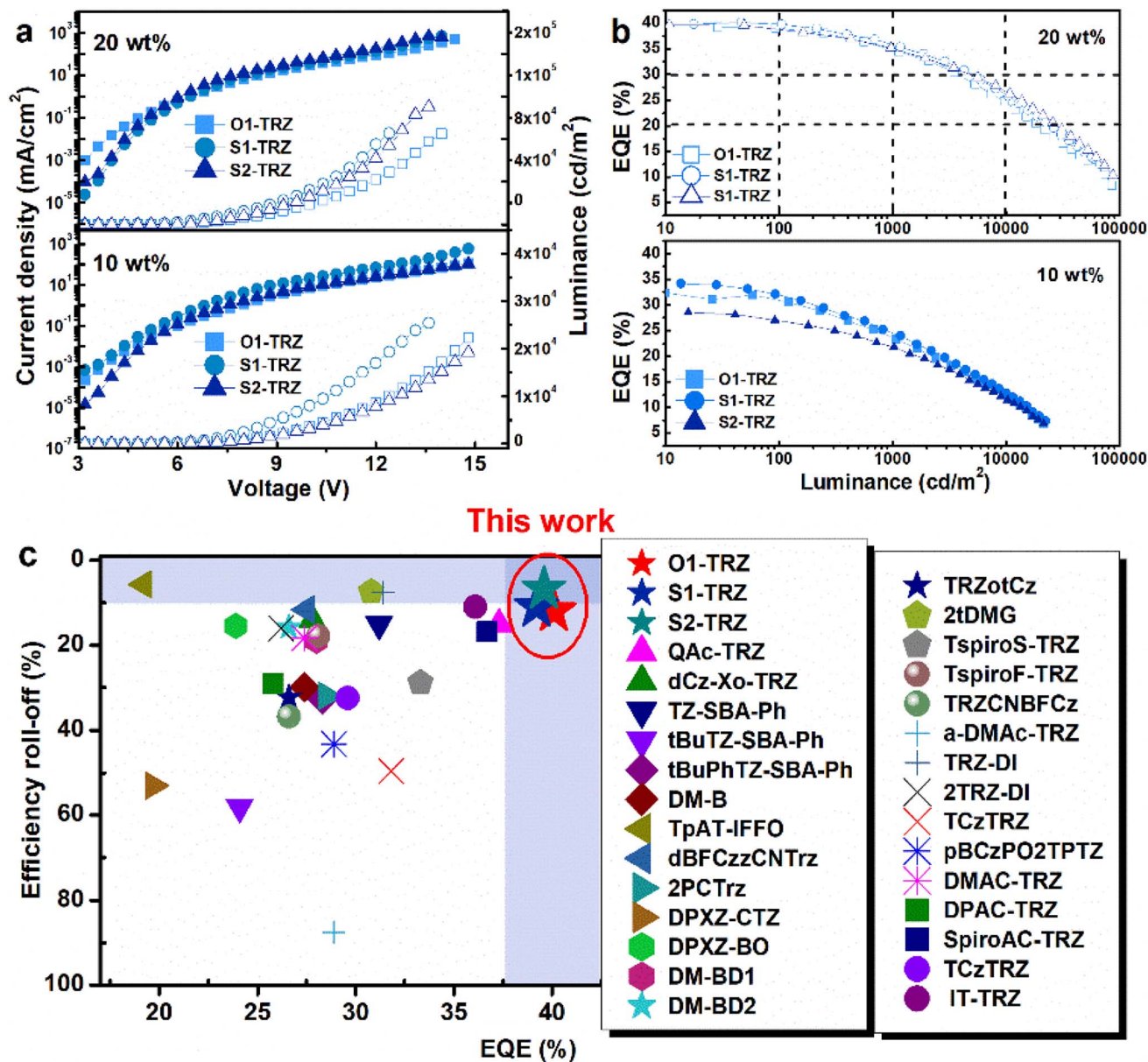


Fig. 4 (a) Current density–voltage–luminance ( $J$ – $V$ – $L$ ) profiles; (b) EQE vs. luminance characteristics. Inset of devices based on O1-TRZ, S1-TRZ and S2-TRZ at 10 and 20 wt% doping concentrations. (c) Comparison of EQE and efficiency roll-off under  $1000 \text{ cd m}^{-2}$  between reported emitters.

Table 3 EL data of O1-TRZ, S1-TRZ and S2-TRZ based devices at different doping concentrations

Emitting layer	EL (nm)	$V_{\text{on}}$ (V)	$L_{\text{max}}$ ( $\text{cd m}^{-2}$ )	$\text{CE}^a$ ( $\text{cd A}^{-1}$ )	$\text{EQE}^a$ (%)	Roll-off <sup>b</sup> (%)	$\text{CIE}^c$ (x, y)
O1-TRZ (10 wt%):PPF	506	3.2	22 324	98.6/91.4/67.0	34.1/31.9/24.5	6/28	0.27, 0.50
S1-TRZ (10 wt%):PPF	506	3.2	25 376	95.2/88.3/66.4	33.3/31.2/23.6	6/29	0.27, 0.50
S2-TRZ (10 wt%):PPF	512	3.2	21 452	85.3/80.5/64.0	29.4/27.0/21.8	8/26	0.30, 0.53
O1-TRZ (20 wt%):PPF	515	3.2	85 230	90.4/78.2/57.8	40.1/39.8/35.3	0.7/12	0.29, 0.53
S1-TRZ (20 wt%):PPF	516	3.2	85 476	122.7/120.9/106.2	39.2/39.0/34.8	0.5/11	0.30, 0.54
S2-TRZ (20 wt%):PPF	519	3.2	110 267	124.7/124.0/115.3	39.6/39.5/36.9	0.2/7	0.31, 0.55

<sup>a</sup> Maximum values at 100/1000  $\text{cd m}^{-2}$ . <sup>b</sup> Efficiency roll-off under 100/1000  $\text{cd m}^{-2}$ . <sup>c</sup> CIE coordinates under 6 V.

excitons. All devices exhibit turn-on voltages of 3.2 V and high luminance exceeding  $21\,000 \text{ cd m}^{-2}$ . Maximum EQEs ( $\text{EQE}_{\text{max}}$ ) of over 29.4% are achieved with the highest value of 34.1% in

the O1-TRZ based device. Compared with O1-TRZ, the lower  $\text{EQE}_{\text{max}}$  of the S1-TRZ based device may be attributed to its lower PLQY. And the low  $\text{EQE}_{\text{max}}$  in the S2-TRZ based device



could result from its low  $k_r$ . Further probing the efficiency roll-off offers similar values of 6–8% under  $100 \text{ cd m}^{-2}$  and 26–29% under  $1000 \text{ cd m}^{-2}$ , which are relatively large. The electron-withdrawing nature of PPF as the host matrix may have led to carrier imbalances in the emissive layer. Therefore, devices with concentration optimization were evaluated at 20 wt%. Remarkably, all devices achieve improved performance with a significantly enhanced luminance of 85 000 to  $110\,000 \text{ cd m}^{-2}$  and increased  $\text{EQE}_{\text{max}}$  values of 39.6% for **S2-TRZ**, 39.2% for **S1-TRZ** and 40.1% for **O1-TRZ**. These results demonstrate their sufficient exciton utilization at a doping concentration of 20 wt%. Besides high PLQY and efficient TADF features, the excellent EL performances also benefit from high molecular horizontal dipole ratios, which will enhance the out-coupling efficiency. The p-polarized PL spectra are shown in Fig. S7,† suggesting that these emitters exhibit horizontal dipole ratios in the range of 80–97%. Such high values will greatly enhance the out-coupling efficiency and boost the EQE to nearly 40%. Notably, efficiency roll-off is substantially reduced with negligible roll-offs under  $100 \text{ cd m}^{-2}$ . At a higher luminance level, **O1-TRZ** and **S1-TRZ** achieve similar roll-off of 12% and 11% under  $1000 \text{ cd m}^{-2}$ , respectively, while **S2-TRZ** shows a small value of only 7%. Compared to **O1-TRZ**, **S1-TRZ** with slightly larger  $k_{\text{RISC}}$  produces a comparable roll-off. **S2-TRZ** features a higher  $k_{\text{RISC}}$  than **S1-TRZ**, experiencing a much gentler roll-off. The synergistic effect of the heavy element effect and nonplanar structure contributes to a significant enhancement of the RISC channel, facilitating the rapid consumption of excitons.

### 3. Conclusion

In conclusion, we have disclosed the great impact of nonplanar structure on accelerating the RISC channel in TADF emitters. By combining cross-coupled spiroacridine and structurally fixed compulsory hindrance, **S2-TRZ** with a properly hooked acridine is successfully established. Analogs of **S1-TRZ** composed of a coplanar acridine and **O1-TRZ** with an O atom are designed to distinguish the effect of the nonplanar geometry and heavy element effect, respectively. All compounds exhibit high PLQYs of 96–98% and small  $\Delta E_{\text{ST}}$ s of 8–9 meV, indicating excellent TADF behaviors. **S2-TRZ** with both heavy atoms and a nonplanar structure shows the highest  $k_{\text{RISC}}$  of  $24.2 \times 10^5 \text{ s}^{-1}$ , whereas **S1-TRZ** possesses a lower  $k_{\text{RISC}}$  of  $11.1 \times 10^5 \text{ s}^{-1}$  due to the absence of a nonplanar acridine. Lacking these benefits, **O1-TRZ** displays the lowest  $k_{\text{RISC}}$  of  $9.64 \times 10^5 \text{ s}^{-1}$ . Encouragingly, all emitters exhibit amazing EL performance with  $\text{EQE}_{\text{max}}$  of 39.6% for **S2-TRZ**, 39.2% for **S1-TRZ** and even 40.1% for **O1-TRZ**, respectively, alongside ultra-high luminance of up to  $100\,000 \text{ cd m}^{-2}$ . More importantly, **S2-TRZ** with a large value of  $k_{\text{RISC}}$  exhibits the lowest roll-off of 7% under  $1000 \text{ cd m}^{-2}$ , indicating the importance of the nonplanar structure in enhancing RISC for excellent EL performance. Overall, the effectiveness of nonplanar geometry in accelerating the RISC channel has been theoretically and experimentally validated. This strategy represents a valuable alternative to manipulating SOC besides the heavy atom effect. This work provides guidance for designing practical TADF emitters.

### Data availability

All data supporting the findings of this study are presented in the article and ESI.† Additional data are available from the corresponding author upon reasonable request.

### Author contributions

H. Liu designed the study. C. L. Yang supervised the project. Y. Liu and G. H. Chen performed the experiments. Y. Meng, H. Peng and J. S. Miao collected and analyzed the data. All the authors discussed the results and co-wrote the manuscript.

### Conflicts of interest

There are no conflicts to declare.

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