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An S-shaped double helicene showing both

fluorescence and circularly polarized

Matthew J. Fuchter * and Eli Zysman-Colman * * and

multi-resonance thermally activated delayed

We present the first example of a multi-resonant thermally activated delayed fluorescent (MR-TADF) extended helicene, Hel-DiDiKTa. This S-shaped double helicene exhibits sky-blue emission, a singlettriplet energy gap, ΔE_{ST} , of 0.15 eV and narrow emission at a peak maximum of 473 nm with a full-width at half-maximum of 44 nm in toluene. The MR-TADF character is confirmed by the small degree of positive solvatochromism and temperature-dependent increase in intensity of the delayed emission. The chiroptical properties of the separated enantiomers are similar to other large helicenes with comparable dissymmetry values, but with the added benefit of MR-TADF. (P)-Hel-DiDiKTa is stable towards enantiomerization, with a Gibbs free energy of activation for enantiomerization (ΔG_e^{\pm}) of 31 \pm 2 kcal mol^{-1} at 298 K, a value similar to other reported double helicenes. (P)-Hel-DiDiKTa is also thermally stable, with a 5% weight loss at 399 °C revealed by thermogravimetric analysis (TGA). Thus, this study further strengthens the burgeoning area of chiral TADF emitters for use in cutting-edge optoelectronic and photocatalytic molecules and materials.

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Introduction

Helicenes are a class of fused polycyclic aromatic frameworks that possess a helically chiral framework, 1,2 where overlapping rings render the enantiomers kinetically stable to racemization. Helicenes usually display strong circular dichroism (CD)³ and circularly polarized luminescence (CPL), 4-8 which has led them to be investigated in a number of different applications. ^{4,9} These include nonlinear optics, 10 chemical sensors, 11 asymmetric catalysis, 12,13 circularly polarised luminescent materials 14-16 and as ligands in Ir, Zn and Pt complexes,9 the latter of which have been employed in phosphorescent CP-OLEDs. 17,18 They have also been exploited as fluorescent emitters,5,19 chiral additives for induced chiral fluorescence polymers²⁰⁻²³ and as components within donor-acceptor (D-A) TADF emitters^{24,25} in CP-OLEDs. Beyond chiroptical properties the high solubility of helicenes in organic solvents (compared to planar acenes), coupled with their excellent thermal stability has led to a wider array of optoelectronic applications, including as hole-transport materials in photovoltaic devices.26-28

Thermally activated delayed fluorescence (TADF) materials have become increasingly attractive as sensors, 29 in lasers, 30

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photodetectors, 31 photocatalysis 32 and bioimaging 33 as well as both emitters and hosts in organic light-emitting diodes (OLEDs)³⁴ due to their ability to harvest 100% of electrically generated excitons to generate light. This is accomplished by conversion of triplet excitons into singlets via reverse intersystem crossing $(RISC)^{34-38}$ enabled by their small S_1 – T_1 energy gap, ΔE_{ST} . The dominant molecular design relies on a strongly twisted donor-acceptor architecture that minimises the overlap between the highest occupied molecular orbital (HOMO) and the lowest unoccupied molecular orbital (LUMO), resulting in a small ΔE_{ST} . This strategy also results in compounds with relatively large conformational flexibility and/or intramolecular/stretching vibrations, which in turn leads to broad emission characterized by a full width half maximum (FWHM) typically greater than 80 nm, resulting in poor color purity in resulting OLED devices.40 Hatakeyama and co-workers have advanced an alternative molecular design strategy to produce TADF materials with narrowband emission (FWHM < 40 nm), based on polycyclic aromatic hydrocarbons doped with electronaccepting B, and electron-donating N and O atoms. 41,42 These so-called multi-resonant TADF (MR-TADF) emitters rely on p- and n-doping of polycyclic aromatic hydrocarbons to limit the overlap between the HOMO and the LUMO electron-density and thus produce a small ΔE_{ST} , while their rigid molecular structure results in negligible conformational reorganization in the excited state, and narrowband emission. 40,42,43 There are now examples of blue, 44-48 green 48-52 and red 43,53,54 MR-TADF materials used as emitters in high-efficiency OLEDs. 55-57

The study and exploitation of CPL has accelerated over the past decade due to a recognition that employing CPL-active compounds can lead to an increased efficiency of OLEDs and other optoelectronic devices. 17,58-62 Other technologies being investigated for the exploitation of CPL include quantum computing, 63,64 3D information displays, 23,65 spintronic devices, 66 transistors, 67 CPL lasers 68 and biological probes. 69,70 The CPL intensity of a chiral molecule can be quantified by the dissymmetry factor, g_{PL} , which is represented by eqn (1):.^{7,71,72}

$$g_{\rm PL} = \frac{I_{\rm L} - I_{\rm R}}{\frac{1}{2} (I_{\rm L} - I_{\rm R})} = \frac{4|\mu||m|\cos\theta}{|\mu|^2 + |m|^2} \tag{1}$$

Here, I_L and I_R refer to the intensity of left- and right-handed circularly polarised luminescence, respectively; μ is the electric transition dipole moment, m is the magnetic transition dipole moment and θ is the angle between the transition dipole moments. This expression reveals that the maximum attainable g_{PL} (which is ± 2) is obtained when the transition dipole moments, μ and m, have the same magnitude $(|\mu|/|m| = 1)$ and are parallel or antiparallel ($\cos \theta = \pm 1$) to each other. Unfortunately, chiral molecules that display high photoluminescence quantum yields $(\Phi_{\rm PL})$ typically show small $g_{\rm PL}$ and vice versa. ^{15,16} For instance, lanthanoid complexes typically show high g_{PL} (0.1–1)^{73–75} but in order to reach high brightness values⁷⁶ they need suitable antenna ligands to overcome the Laporte-forbidden nature of their transitions. In contrast, organic molecules can attain high absorption and emission intensities but typically show relatively

low $g_{\rm PL}$ ($\leq 10^{-2}$) due to their significantly larger electric transition dipole moments compared to their magnetic transition dipole moments, coupled with their relatively smaller molecular size. 7,8,77,78

Presently, there exists a small number of organic compounds displaying CP-TADF, with most of the reports emerging over the past few years. A number of these have been used in CP-TADF OLEDs. 18,60,62,79,80 showing maximum external quantum efficiencies (EQE_{max}) surpassing $30\%^{80}$ but with $g_{\rm EL}$ typically $\leq 10^{-3}$ (Fig. 1a). The overwhelming majority of compounds showing CP-TADF are based on a donor-acceptor design that incorporates a remote stereogenic unit (asymmetric centre or axis) to induce chirality, 18,60,61,81-83 and so do not show narrow emission spectra. Materials showing concomitantly narrow emission, TADF and CPL are of significant interest, and a class of compounds that can meet these requirements are chiral MR-TADF compounds. To date, there exists only a handful of reports of chiral MR-TADF compounds. Compounds OBN-2CN-BN and OBN-4CN-BN, 84 were based loosely on Hatakeyama's DABNA-1⁴¹ design and contain a peripheral chiral (R/S)octahydro-binaphthol group that is not directly implicated in the emissive short-range charge transfer excited state.84 A similar strategy was used for QAO-PhCz,85 by introducing a 9-phenyl-9Hcarbazole appended group to the MR-framework to lock the helical structure and obtain chirality. There are only two reports where the MR-TADF core structure shows intrinsic chirality and both are boron/nitrogen-based helicenes: compounds BN586 and OBN-Cz (aka 1a).87

The first CPL data reported for helicenoids belong to two bridged triphenylamine diketone derivatives, reported by Venkataraman et al., (renamed here as Hel-DiKTa-3 and Hel-**DiKTa-4**, Fig. 1b) with $g_{\rm PL}$: $-1.1 \times 10^{-3}/9 \times 10^{-4}$ at 453 nm for (M)-Hel-DiKTa-3 and (P)-Hel-DiKTa-3 and $-7 \times 10^{-4}/8 \times 10^{-4}$ at 478 nm for (M)-Hel-DiKTa-4 and (P)-Hel-DiKTa-4, respectively.14 Given our previous work on diketone-based MR-TADF emitters (namely DiKTa and DDiKTa, structures shown in Fig. S1, ESI†),44,49 we envisaged that developing a helical analogue of DiKTa would be an elegant strategy to obtain a CPL-active molecule without compromising MR-TADF. Our preliminary quantum calculations for Hel-DiKTa-2⁸⁸ (Fig. 1a, renamed here for clarity), however, revealed that its $\Delta E_{\rm ST}$ is too large (0.51 eV) for it to show MR-TADF due to triplet stabilization on the naphthalene group (vide infra).

Here we report an intrinsically helically chiral MR-TADF molecule that is CPL-active and is based on an extended helicene structure, Hel-DiDiKTa (Fig. 1b); an S-shaped double [4]helicene. Despite a large current interest in multiple helicenes, chiroptical properties of these molecules remain understudied.⁸⁹ We report in-depth structural, photophysical and chiroptical data for Hel-DiDiKTa and believe our study paves the way for further chiral MR-TADF molecules, based on extended or multiple helicene frameworks.

The synthesis of Hel-DiDiKTa is shown in Scheme 1a. A Cu-catalysed Ullman coupling between m-phenylenediamine and methyl 2-iodobenzoate afforded intermediate 1 in moderate yield (43%). Saponification proceeded quantitatively to yield 2,

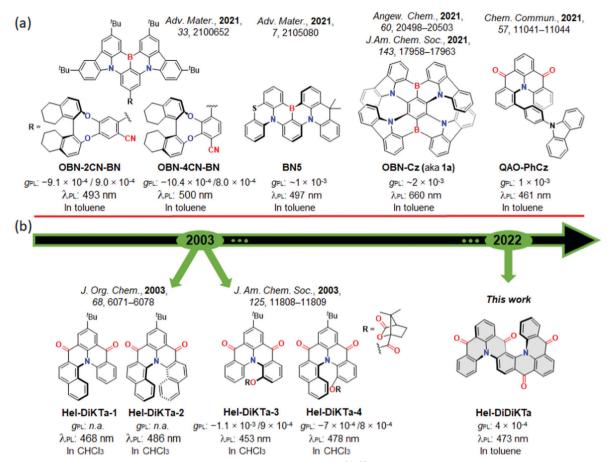
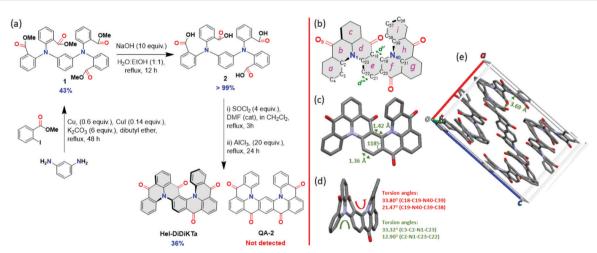


Fig. 1 (a) Structure of prior examples of MR-TADF materials able to display CPL. 84-87 (b) Structure of prior examples of triangulene helicenes and Hel-DiDiKTa. 14,88

which was then subjected to a four-fold intramolecular Friedel-Crafts acylation of the in situ-prepared acyl chloride derivative in the presence of the Lewis acid AlCl₃ to afford the title compound,

(rac)-Hel-DiDiKTa, in 36% yield as a racemate; the C-shaped compound QA-2, which was recently reported by Yasuda et al. 45 using a different synthetic route (and crystallised as a non-chiral



Scheme 1 (a) Synthesis of Hel-DiDiKTa. (b) Schematic of molecular structure of Hel-DiDiKTa showing atom labels (double bonds omitted for clarity). (c) View showing Hel-DiDiKTa in the crystal structure (front view) (internal bond angles: 119.42(10)° (C18-C19-C20), 118.73(10)° (C19-C20-C21), 122.54(10)° (C20-C21-C22), 119.41(10)° (C21-C22-C23), 119.12(9)° (C22-C23-C18) and 119.39(9)° (C23-C18-C19)) and (d) side view showing the helical pitch of Hel-DiDiKTa (torsion angles: 33.32° (C3-C2-N1-C23), 12.90° (C2-N1-C23-C22), 33.80° (C18-C19-N40-C39) and 21.47° (C19-N40-C39-C38)) (H atoms omitted for clarity). (e) Packing diagram of Hel-DiDiKTa viewed down the crystallographic b-axis.

meso isomer), was not detected. We performed Density Functional Theory (DFT) and coupled cluster calculations to probe which of the two products, (rac)-Hel-DiDiKTa or QA-2, is more stable in the ground state. QA-2 was calculated to be more stable than **Hel-DiDiKTa** by between 18 and 29 kJ mol⁻¹, irrespective of the methodology applied (Fig. S19 and Table S7, ESI†), thus (rac)-Hel-DiDiKTa represents the kinetic product. The identity and purity of Hel-DiDiKTa was established by a combination of NMR spectroscopy, HRMS, melting point determination, HPLC measurements and elemental analysis. Additionally, single crystals of sufficient quality were grown by slow evaporation of a CH₂Cl₂: EtOAc solution (7:3 ratio), enabling determination of the single crystal structure.

Crystallographic data of Hel-DiDiKTa obtained by single crystal X-ray diffraction revealed that the compound crystalizes as a racemate. The helical pitch through each of the two [4]helicenes varies between 12.90(15) and 33.80(15)° (torsion angles C3-C2-N1-C23, C2-N1-C23-C22, C18-C19-N40-C39 and C19-N40-C39-C38) and is similar to that found in DiKTa (27.2°) (Scheme 1d). Typically, in helicenes, the benzene ring at the centre of the helical structure is distorted, with two very different C-C bond lengths; a longer C-C bond facing the inner helix and opposite it a much shorter one facing the outer helix (distances d' and d'' in Scheme 1b). This also leads to internal bond angles that are considerably different from that of an ideal planar benzene ring. This is not usually the case for S-shaped double helicenes, 90-93 and as such, in Hel-DiDiKTa, the

central benzene e-ring (Scheme 1b) has internal bond angles between 118.73(10) and 122.54(10)° and C-C bond distances between 1.3641(17) and 1.4207(15) Å (Scheme 1c). There is only one $\pi - \pi$ stacking interaction between adjacent molecules (packing diagram displayed in Scheme 1e), the c-ring interacting with the g-ring [centroid...centroid distance 3.6875(6) Å], resulting in π -stacked chains running along the *b*-axis.

Cyclic voltammetry (CV) and differential pulse voltammetry (DPV) recorded in degassed dichloromethane (Fig. 2a) revealed an irreversible oxidation wave at $E_{\rm ox}$ = 1.81 V νs . SCE and a reversible reduction wave at $E_{\rm red}$ = -1.16 V vs. SCE. The corresponding HOMO and LUMO energy levels are -6.15 and -3.18 eV, respectively. The LUMO is only slightly stabilized compared to that of the parent compounds DiKTa (-3.11 eV)while the HOMO is considerably more stabilized than that of **DiKTa** (-5.93 eV). The electrochemical data are summarized in Table S6 (ESI†).

The UV-vis absorption spectrum of (rac)-Hel-DiDiKTa in toluene (Fig. 2b and Fig. S8a, ESI†) is significantly different than that of DiKTa. There is a high-energy, high-intensity band at 360 nm ($\varepsilon = 10\,917~{\rm M}^{-1}~{\rm cm}^{-1}$) assigned by Spin-Component Scaling second-order approximate Coupled-Cluster (SCS-CC2) calculations as a π - π * transition centered on one half of the emitter (Fig. S23, ESI†). There is a higher-intensity, low-energy band at 425 nm (ε = 21 140 M⁻¹ cm⁻¹) assigned as a short-range charge-transfer (SRCT) transition, which, according to the calculations, is associated with a transition to the S2 state at

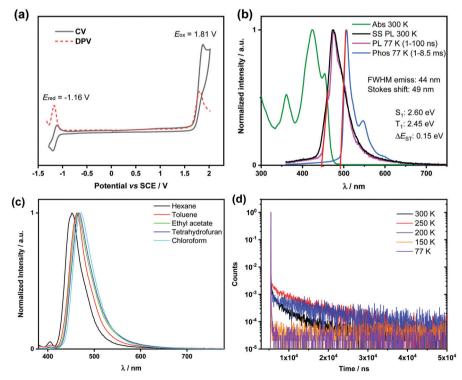


Fig. 2 Optoelectronic characterization of (rac)-Hel-DiDiKTa: (a) cyclic and differential pulse voltammograms in degassed CH₂Cl₂ with 0.1 m [ⁿBu₄N]PF₆ as the supporting electrolyte and Fc/Fc⁺ as the internal reference (Fc/Fc⁺ = 0.46 V vs. SCE).⁹⁴ (b) Absorption (green line) and steady-state (SS) PL spectra obtained in toluene at 300 K (black line) and 77 K (pink line), and phosphorescence (Phos.) spectrum in toluene glass at 77 K (blue line) (delay: 1 ms; gate time: 8.5 ms, $\lambda_{\text{exc}} = 343$ nm). (c) Solvatochromic PL study ($\lambda_{\text{exc}} = 350$ nm). (d) Temperature-dependent lifetime for 1 wt% (rac)-Hel-DiDiKTa in mCP.

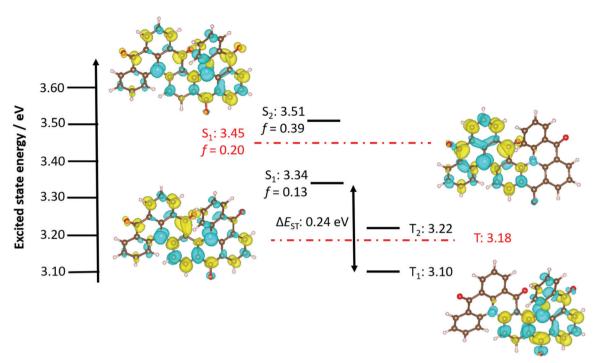


Fig. 3 Difference density plots computed at SCS-CC2/cc-pVDZ level considering only one-electronic transition contributions for (rac)-Hel-DiDiKTa. The red dashed lines show the energy levels for **DiKTa**.

3.51 eV. The band at 453 nm is associated with a transition to the S_1 state, and has a lower ε of 14 292 M^{-1} cm⁻¹ than the band at 425 nm; a trend in line with the lower calculated oscillator strength, f, of 0.13 for S_1 compared to 0.39 for S_2 . This assignment is further corroborated by the energy difference between these states, calculated to be 0.17 eV (Fig. 3) and measured to be 0.18 eV (Fig. 2b). S₁ is also SRCT in nature, with S₁ centered more around the middle of the molecule and S2 associated with electronic density across the entire molecule, hence the oscillator strength changes (Fig. 3). This low-energy S₁ band is red-shifted compared to that of **DiKTa** (433 nm), with lower molar absorptivity (ε of **DiKTa** $\sim 21\,000\,\mathrm{M}^{-1}\,\mathrm{cm}^{-1})^{44}$ in line with the change in energy of the S₁ state and the lower oscillator strength, which are 3.45 eV and 0.20 for DiKTa, and 3.34 eV and 0.13 for Hel-DiDiKTa (Fig. 3).

The room temperature steady-state (SS) photoluminescence (PL) spectrum of (rac)-Hel-DiDiKTa, recorded in toluene, peaks at 473 nm, which is slightly red-shifted compared to that of **DiKTa** (λ_{PL} = 453 nm). The PL spectrum is narrow (FWHM = 44 nm) and shows a small Stokes shift of 49 nm, both of which reflect the conformationally rigid structure and the small degree of geometrical relaxation in the excited state (Fig. 2b). This FWHM value is considerably smaller than many of the helicene compounds reported in the literature (often 50 -80 nm). 1,16,19,81,91-93,95 The PL spectrum at 77 K is nearly identical to that at room temperature. Furthermore, a very small degree of positive solvatochromism is observed (Fig. 2c and Table S1, ESI†), behavior that is a hallmark of MR-TADF compounds.44,96 Unlike para-disposed nitrogen and ketone groups that leads to a significant bathochromic shift in the emission compared to their meta-functionalized analogue, as evidenced in the case of the emitters DMQA and QA-1 (c.f.,

Fig. S1, ESI†) recently reported by Yasuda et al., 45 the emission energy of (rac)-Hel-DiDiKTa changes very little compared to that of **DiKTa**. This is despite of the increased conjugation length of Hel-DiDiKTa and is due to the meta-disposition of the amine and ketone groups. The S1 and T1, determined from the peaks of the prompt fluorescence and phosphorescence spectra in toluene at 77 K, respectively, are 2.60 and 2.45 eV, leading to a moderate $\Delta E_{\rm ST}$ of 0.15 eV. These values are in reasonable accord with the predicted by SCS-CC2 calculations (ΔE_{ST} = 0.24 eV) and with those of other reported ketone-containing MR-TADF compounds, including DiKTa (0.20 eV). 44,45,49 The $\Phi_{\rm PL}$ in toluene is very low at 1% and is due to significant non-radiative decay that is not uncommon in helicene compounds. No delayed emission was detected by timeresolved PL measurements in toluene (Fig. S13, ESI†). Recently, Wu et al. highlighted that many MR-TADF emitters, including N/ketone-based emitters, may not show TADF behavior in solvents while TADF can be observed in a suitable host due to exciplex-like host-emitter interactions. 97 The SS PL spectrum is only slightly affected by the presence of air (Fig. S9, ESI†). The $\Phi_{\rm PL}$ improved to 4.1% as a 1 wt% doped film in 1,3-bis (N-carbazolyl)benzene (mCP) and to 6.2% in 1 wt% doped film in PMMA. We note that the $\Phi_{\rm PL}$ remained essentially constant at ~5% regardless of the doping concentration in mCP (Table S2, ESI†). These results are similar to the Φ_{PL} values reported for unsubstituted helicenes. 16

In mCP (1 wt% of emitter) the SS spectrum peaks at 477 nm (Fig. S10a, ESI†), the FWHM value is 50 nm, and, in contrast to what is seen in toluene solution (Fig. S9, ESI†), both the PL intensity and lifetime of (rac)-Hel-DiDiKTa decrease moderately in the presence of O₂ (Fig. S10a and b, ESI†).

Temperature-dependent time-resolved PL measurements revealed an increase in the contribution of the delayed component of the emission decay with increasing temperature (Fig. 2d and Table S4, ESI†), which corroborates the TADF character of **Hel-DiDiKTa.** The S_1 , T_1 and ΔE_{ST} values, determined from the peaks of the prompt fluorescence and phosphorescence spectra at 77 K, in mCP are 2.59, 2.44 and 0.15 eV, respectively, while in PMMA, these are essentially the same at 2.58, 2.45 and 0.13 eV. respectively. The temperature-dependent steady-state PL emission of (rac)-Hel-DiDiKTa was measured in 1% mCP, and the data are shown in Fig. S12 (ESI†). Analysis of the evolution of the SS PL of (rac)-Hel-DiDiKTa with decreasing temperature reveals a competition between phosphorescence and delayed fluorescence, with the phosphorescence intensity increasing considerably at lower temperatures (150 K and 77 K, Fig. S12, ESI†), which indicates that non-radiative decay is suppressed at lower temperatures and the phosphorescence becomes more competitive (Table S4, ESI†). We also observed from the temperature dependent transient decay profile of (rac)-Hel-DiDiKTa (1 wt% in mCP, Fig. 2d) that the delayed fluorescence intensity increases upon going from 300 K to 250 K, indicating decreased non-radiative decay due to suppression of molecular vibrations at 250 K. The delayed fluorescence intensity then decreases considerably upon reaching 77 K, which we ascribe to the quenching of the TADF (Table 1). The photophysical data is summarized in Table 1.

Density Functional Theory (DFT) calculations were first performed to optimize the geometry of the ground state, taking the crystal structure geometry as the starting point. Compared to DiKTa (-6.20 eV), the calculated HOMO energy is slightly stabilized to -6.26 eV for **Hel-DiDiKTa** (Table S8, ESI†). A larger stabilization of the LUMO energy exists at -2.53 eV (for **DiKTa**, $E_{\text{LUMO}} = -2.23 \text{ eV}$). The relative HOMO/LUMO levels corroborate the experimental values of -5.93 eV/-3.11 eV for DiKTa and -6.15 eV/-3.18 eV for **Hel-DiDiKTa**. Previously, we have highlighted that DFT is not appropriate as it does not accurately predict the excited state energies of MR-TADF emitters. 96 We thus used Spin-Component Scaling second-order approximate Coupled-Cluster (SCS-CC2) calculations, which we have shown previously to provide an excellent set of predictions of $\Delta E_{\rm ST}$ and S₁ energies. 44,49,98,99 SCS-CC2 calculations predict the S₁ state to be 3.34 eV and the T_1 state to be 3.10 eV, resulting in a ΔE_{ST} of 0.24 eV, which is slightly smaller than that calculated for DiKTa $(\Delta E_{\rm ST} = 0.27 \text{ eV})$. The difference density plots are shown in Fig. 3

and reveal how the electronic density evolves from the ground state to the excited states. The pattern of the difference density plots for S₁ and S₂ is indicative of SRCT states involving the entirety of the helicene structure, while the plots for T₁ and T₂ likewise show the alternating increase and decrease in electron density pattern, but this is localized on different fragments of the helicene corresponding to the DiKTa core structure. The differences in orbital type between the singlet and triplet excited states ensure that RISC can occur directly between these states; 100 the presence of an intermediate triplet excited state should also facilitate RISC. 101-103 The SRCT character of the S₁ and T_1 states explains the moderate ΔE_{ST} . We also performed SCS-CC2 calculations on the previously reported compound Hel-?A3B2 tlsb -0.045w?>DiKTa-2. This compound possesses a very large $\Delta E_{\rm ST}$ of 0.51 eV (Table S8, ESI†) due to triplet stabilisation by the naphthalene group 104 and so is unlikely to show TADF (Fig. S21, ESI†).

(rac)-Hel-DiDiKTa was separated into its P- and Menantiomers using peak recycling chiral HPLC (cHPLC; Fig. S14, ESI†), demonstrating high enantiomeric excess (>99% and >97%, respectively; Fig. S15, ESI†). The circular dichroism (CD) spectra for the separated enantiomers were recorded in toluene (Fig. 4a) and showed mirror-image CD signals (Fig. S17, ESI† right), which exhibited multiple bisignate cotton effects throughout the ground state absorption spectral range (300-480 nm). Time-dependent DFT calculation (PBE0/6-31g(d,p)) was employed to predict the CD spectra and assign the absolute configuration (P and M) of Hel-DiDiKTa (Fig. S24, ESI†). The TD-DFT predicted CD spectra shown in Fig. S24 (ESI†) align with the CD spectra obtained experimentally. A maximum $|g_{abs}|$ of 2.8×10^{-3} was observed at 320 nm (Fig. S17, ESI†). These results are of a similar order of magnitude to other chiral small molecule TADF systems.2,14 Circularly polarized photoluminescence (CPPL) spectra (Fig. S17, ESI†) display mirror image monosignate bands peaking at around 465 nm, confirming the transference of chirality to the excited state, with a maximum $|g_{\rm PL}|$ of 4 \times 10⁻⁴ for both enantiomers (Fig. S18, ESI†). These values are also similar to other helicenes and chiral small molecule TADF systems reported in the literature, 76 and align with those reported for helicenoids with two bridged triphenylamine diketone derivatives $(10^{-4}-10^{-3})$, see Fig. 1). As expected for helicenes⁷ where the ground and emitting excited state geometries are generally similar, the sign and magnitude of $g_{\rm PL}$ align with those of $g_{\rm abs}$ calculated at around 455 nm.

Table 1 Optoelectronic properties of (rac)-Hel-DiDiKTa

In toluene								In film					
$\lambda_{\rm abs}{}^a$ /nm	$\varepsilon^a/\mathrm{M}^{-1}~\mathrm{cm}^{-1}$		$\Phi_{\rm PL}$ in N ₂ (air) ^c /%			T_1^e/eV	$\Delta E_{\rm ST}^{f}/{\rm eV}$		$\Phi_{\rm PL}$ in N ₂ (air) /%		S ₁ /eV	T ₁ /eV	$\Delta E_{ m ST}$ /eV
360/425/453	10 917/21 140/14 292	473	1.34 (1.27)	44 (0.25)	2.60	2.45	0.15	478^{g} 480^{h}	$4.1 (4.0)^g$ $6.2 (5.4)^h$	$52 (0.29)^g$ $58 (0.33)^h$	2.59^{g} 2.58^{h}	2.44^{g} 2.45^{h}	$0.15^g \ 0.13^h$

^a UV-vis absorption band of interest. ^b PL in toluene degassing with N₂. ^c Photoluminescence quantum yield in toluene relative to quinine sulfate in 1 N $_2$ SO₄ ($\Phi_{PL} = 54.6\%$). d Full width at half maximum. Obtained using an integrating sphere under N₂. f Energy gap between S₁ and T₁ calculated from the difference of the peaks of the fluorescence and phosphorescence spectra in toluene glass at 77 K. g 1 wt% (rac)-Hel-DiDiKTa doped in mCP. h 1 wt% (rac)-Hel-DiDiKTa doped in PMMA.

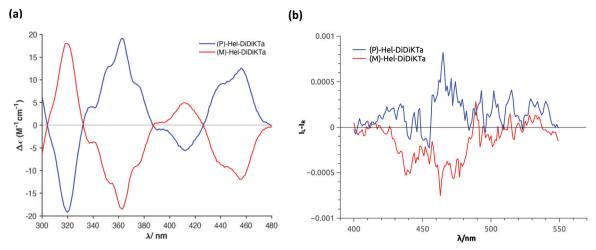


Fig. 4 (a) Circular dichroism and (b) CPL spectra for (P)-Hel-DiDiKTa (blue) and (M)-Hel-DiDiKTa (red) in toluene

Upon optimization of the molecular structure at the lowest-energy singlet excited state (S₁) at the PBE0/6-31g(d,p) level of theory, we calculated a $g_{\rm PL}$ value of 6.94 \times 10⁻⁴ (Table S9, ESI†), which aligns closely with the measured g_{PL} . The main limiting factor for the magnitude of g_{PL} , based on our calculations, is the nearly perpendicular alignment of the $|\mu|$ and |m| vectors $[\cos \theta] =$ 0.116918 ($\theta = 83.28573$)]. Although the magnitude of |m| is low at $0.348671679 \times 10^{-20}$ esu cm⁻¹, compared to $|\mu|$ (234.811511 \times 10⁻²⁰ erg G⁻¹), it is within the range typically observed for helicenes. 3,15,16,95 This result suggests that our molecular design is promising to generate strong CPL MR-TADF emitters if the alignment between $|\mu|$ and |m| can be solved.^{3,15,16,95} While these molecules do not have suitable $\Phi_{\rm PL}$ to warrant exploration in OLEDs, they provide valuable insight into CP-TADF material design, particularly integrating MR-TADF into helicenes.

After the resolution of both enantiomers, the enantiomerisation process was studied. The CD signal decay from (P)-Hel-**DiDiKTa** in *p*-xylene at different temperatures (T = 75 °C, 80 °C, 85 °C and 90 °C) was monitored over time (see Fig. S25-S28, ESI†). An inversion barrier characterised by $\Delta H_{\rm e}^{\ddagger}$ = 39 \pm 1 (kcal mol^{-1}) and $\Delta S_e^{\ddagger} = 26 \pm 1$ (cal K⁻¹ mol^{-1}) was found (see Table S10, ESI†). This results in a Gibbs free energy of activation for the enantiomerisation process ($\Delta G_{\rm e}^{\ddagger}$) of 31±2 (kcal mol⁻¹) at 298 K. (P)-Hel-DiDiKTa possesses a similar inversion barrier (298 K) to other double helicenes found in the literature (Table S10, ESI†). 105,106 (P)-Hel-DiDiKTa also appears to have a similar enantiomerisation barrier to other bridged phenylamines previously reported (Table S10, ESI†). 107,108 Moreover, when compared to [5] and [6]-helicene (f and g in Table S10, ESI†), (P)-Hel-DiDiKTa possesses a similar barrier as [6]helicene. 109 The half-life obtained for the enantiomerisation is ca. 310 years at room temperature.

Not only is (P)-Hel-DiDiKTa stable to enantiomerization but the compound is also stable towards thermal decomposition. Thermogravimetric analysis (TGA) showed that no decomposition was observed in the temperature range of the enantiomerization study and the decomposition temperature was found to be 399 °C, corresponding to 5% weight loss (Fig. S30, ESI†).

In summary, we report a novel S-shaped triphenylamine diketone double [4]helicene that exhibits MR-TADF. Unlike the previously reported triphenylamine diketone helicene (Hel-DiKTa-2), Hel-DiDiKTa was predicted by SCS-CC2 calculations to show a $\Delta E_{\rm ST}$ appropriate for TADF, which was confirmed experimentally (0.15 eV in 1 wt% doped mCP film). As expected from a MR-TADF emitter, the emission peak at 473 nm is narrow, with a FWHM of 44 nm in toluene, a value considerably smaller than often observed for helicenoids. In 1 wt% mCP the emission peak at 477 nm shows a FWHM value of 50 nm. The TADF behavior of Hel-DiDiKTa was confirmed by temperaturedependent time-resolved photoluminescence measurements, and the chiroptical behavior of the enantiomers was determined, with a maximum $|g_{abs}|$ of 2.8×10^{-3} and a $|g_{PL}|$ of 4 \times 10⁻⁴. While the low $\Phi_{\rm PL}$ precluded the use of this compound as an emitter in OLEDs, we believe that this report will serve as a foundation to develop intrinsically chiral, CPL-active MR-TADF molecules for OLEDs. Further work is undergoing in our lab to improve the optoelectronic and chiroptical properties of chiral MR-TADF compounds. Finally, the enantiomerization studies and the thermogravimetric analysis demonstrate the high inherent stability of (P)-Hel-DiDiKTa.

Conflicts of interest

The authors declare that they have no conflict of interest.

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