Two-photon-excited ultralong organic room temperature phosphorescence by dual-channel triplet harvesting†

Zhu Mao, † Zhan Yang, † Chao Xu, † Zongliang Xie, † Long Jiang, † Feng Long Gu, † Juan Zhao, † † Yi Zhang, † † † Matthew P. Aldred † and Zhenguo Chi † † †

Due to inefficient molecular design strategies, two-photon-excited ultralong organic room temperature phosphorescence (TPUOP) has not yet been reported in single-component materials. Herein, we present an innovative design method by dual-channel triplet harvesting to obtain the first bright TPUOP molecule with a lifetime of 0.84 s and a quantum efficiency of 16.6%. In compound o-Cz the donor and acceptor units are connected at the ortho position of benzophenone, showing intramolecular space charge transfer. Therefore, the two-photon absorption ability is improved due to the enhanced charge transfer character. Moreover, the small energy gap boosts dual-channel triplet harvesting via ultralong thermally activated delayed fluorescence and H-aggregation phosphorescence, which suppresses the long-lived triplet concentration quenching. Through two-photon absorption, a near-infrared laser (808 nm) is able to trigger the obvious ultralong emission under ambient conditions. This research work provides valuable guidance for designing near-infrared-excited ultralong organic room temperature phosphorescence materials.

Introduction

Ultralong organic room temperature phosphorescence (UOP) materials, in which the light generated can last from several seconds to hours after removal of the excitation source, have been attracting considerable research interest. These types of materials are important for certain practical applications such as in emergency lighting, anti-counterfeiting, sensors, and biological sciences. 1-5 Recently, strategies such as H-aggregation, crystallization, guest-host doping, intermolecular electronic coupling, and molecular self-assembly have all been adopted in order to promote the development of UOP materials. 6-13 However, the reported UOP materials are only triggered by a single-photon of ultraviolet or blue light, 14,15 restricting their actual utilization. Alternatively, two-photon absorption (TPA) utilizes the near-infrared (NIR) wavelengths that are more accessible for biological sciences due to higher spatial resolution, less phototoxicity and deeper tissue penetration. 16,17 Although progress has been made, systems exhibiting two-photon-induced ultralong emission are limited to inorganic and organic–inorganic hybrid materials, 18-21 and two-photon-excited ultralong organic room temperature phosphorescence (TPUOP) materials from purely organic single component materials are unfortunately not yet discovered. The design and synthesis of TPUOP materials still remains a challenge because of the lack of efficient molecular design strategies.

There are several aspects that need to be considered for the design of efficient TPUOP emitters. It is desirable for the material to possess the following: (i) a longer wavelength absorption so that excitation can be maintained in the NIR spectral region for efficient tissue penetration, (ii) a high phosphorescence efficiency (ΦT) in order to lower the excitation incident energy, and (iii) a long phosphorescence lifetime to avoid continuous excitation. For high ΦT one should suppress the non-radiative pathways and enlarge the charge delocalization for red-shifted absorption. Hence, recent research studies have focused on enhancing intermolecular interactions to suppress non-radiative pathways, and indeed have successfully achieved high ΦT and a long lifetime. 5-15 However, less attention has been paid to the intramolecular steric hindrance effects in reducing the excited-state vibrations and regulating the charge transfer characteristics of ultralong phosphorescent compounds.

† Electronic supplementary information (ESI) available. CCDC 1885292. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c9sc02282a

‡ Dr Z. Mao, Dr Z. Yang and Dr C. Xu contributed equally to this work.
In this contribution, we present the design and synthesis of a single-component organic molecule that exhibits TPUOP properties by dual-channel triplet harvesting. Based on p-Cz (Fig. 1a), which is a known UOP emitter, the more non-linear structural isomer o-Cz that incorporates its donor (D) and acceptor (A) units with an ortho-connectivity, exhibits a folded conformation with close spatial proximity of the D and A units (Fig. 1b). Consequently, intramolecular charge transfer (CT) can take place through space CT (SCT) and through bond CT (ICT) simultaneously, affording an enhanced CT character to increase the TPA ability. The SCT effect causes a reduction in the singlet (S1)-triplet (T1) energy gap (ΔE<sub>ST</sub>), which results in a highly efficient dual-channel triplet harvesting (Φ<sub>T</sub> = 16.6%) via intersystem crossing (ISC) for phosphorescence and reverse ISC (RISC) for ultraslong thermally activated delayed fluorescence (TADF). In addition, intramolecular D-A interactions confine the vibrations of the excited molecule that suppresses non-radiative pathways. Benefiting from the SCT feature, o-Cz exhibits a stronger CT character and effective triplet harvesting, and consequently we successfully obtained the first single-component TPUOP molecule. o-Cz exhibits obvious ultraslong emission for 4 s after removal of the NIR-light excitation source (Fig. 1d).

**Results and discussion**

Compound o-Cz was synthesized by a CuI catalyzed Ullmann cross-coupling reaction of carbazole and 2BrBP (Fig. S1†). A suspension of 2BrBP (1 g, 3.83 mmol), carbazole (0.77 g, 4.60 mol), CuI (0.1 g) and 1,10-phenanthroline monohydrate (0.1 g) in DMF (10 mL) was stirred at reflux (150 °C) overnight under an argon atmosphere. The reaction mixture was cooled to room temperature and subsequently poured into water (100 mL), and the resulting precipitate was filtered and dried. The crude product was purified by flash column chromatography on silica gel (DCM : hexane = 1 : 2) to afford a yellowish solid o-Cz (0.72 g, yield 54.1%).

Time-dependent density functional theory (TD-DFT) was firstly performed to investigate the electronic distributions for the constitutional isomers p-Cz and o-Cz (extracted from the corresponding single crystal X-ray analysis). As depicted in Fig. S2, both the highest occupied molecular orbitals (HOMOs) and lowest unoccupied molecular orbitals (LUMOs) are separated on the carbazole (donor) and benzophenone (acceptor) units, respectively. Interestingly, the LUMO+3 of o-Cz is formed by the through-space orbital overlap of the donor and acceptor units and results in a smaller ΔE<sub>ST</sub> value of 0.21 eV, compared to p-Cz (0.40 eV). The reduced ΔE<sub>ST</sub> value suggests the prospects for rapid triplet population via ISC, which is favorable for generation of phosphorescence. Additionally, the reduced density gradient analysis reveals the significant intramolecular interactions in the folded o-Cz (Fig. 2b and S3, including the green region of D-A attractions and the brown region for steric hindrance). These interactions in o-Cz can effectively restrict the excited-state molecular vibrations and thus suppress non-radiative pathways.

In contrast, the interaction regions are negligible in the more linear p-Cz molecule (Fig. 2a). Moreover, the second hyperpolarizability values (γ), which are directly proportional to the TPA cross section, are 411.90 × 10<sup>−36</sup> esu and 6022.53 × 10<sup>−36</sup> esu for p-Cz and o-Cz, respectively, which indicates better TPA capacities of o-Cz.

Detailed analysis of the photophysical properties of o-Cz was carried out by UV-vis absorption, photoluminescence (PL), and temperature-dependent transient decay studies. The UV-vis absorption profiles of compounds o-Cz and p-Cz in dilute THF are similar (Fig. 3a). The main vibronic absorption bands ranging around 230–270 nm, 270–300 nm, and 300–350 nm are due to the localized π–π* transitions of the acceptor and donor moieties. The broad and weak absorptions around 350–400 nm are caused by the ICT of the two isomers. For o-Cz, the onset of absorption is red-shifted to 420 nm, which originates from the SCT with much lower oscillator strength.
the emission maximum of o-Cz in solvents with different polarities exhibits a significant red-shift as the solvent polarity increases, from 424 nm in non-polar hexane to 492 nm in polar DMSO. This result demonstrates the ICT character of o-Cz (Fig. S4†). The PL emission of o-Cz in THF solution exhibits a bathochromic shift with a narrower full width at half-maximum (FWHM) of 73 nm compared to that of the analogous p-Cz (90 nm), indicating a stronger CT character with less excited-state structural vibrations in o-Cz (Fig. 3b). In the solid state, the onset absorption of o-Cz is extended to the visible-light region around 460 nm, denoting further strengthening of the CT character in the aggregation state (Fig. 3c). Generally, the enhancement of the ICT effect is an effective strategy to increase the TPA ability.

The crystalline powder of o-Cz (Fig. S5†) exhibits an obvious UOP feature under ambient conditions after turning off the UV and NIR light excitation (Fig. 1d). Both the steady-state PL and delayed spectra of the crystalline o-Cz show two emission bands located around 472 nm (structureless with a CT character) and 522–700 nm (fine vibrations with a locally excited character) at room temperature (Fig. 3d). However, the two emission bands exhibit lifetimes of 0.79 s (472 nm) and 0.84 s (542 nm) (Fig. 3e), respectively, with totally different electronic configurations (CT and locally excited (LE) characters). We speculate that the different-emissive origins are responsible for the dual bands in o-Cz. In a further experiment, the excitation–emission matrix spectra (Fig. S6 and S7a†) show that the two emission bands can be excited from 260 nm to 450 nm, with the strongest excitation at 428 nm. The long-lived emissions of o-Cz can be triggered by both ultraviolet and visible light. The normalized steady-state PL spectra (Fig. S7b†) with different excitation wavelengths are not overlapped around the LE zone (522–680 nm). Additionally, the delayed emission spectra (normalized at a delay time of 8 ms) show distinct intensity around the CT zone (472 nm) with UV (365 nm) and visible light (405 nm) excitation (Fig. S8†). These results indicate that the dual-channel of CT and LE long-lived emission originate from different emissive species in the crystalline o-Cz. The emission intensities of the dual peaks in o-Cz show only a slight decrease in an oxygen atmosphere compared to that in a vacuum (Fig. S9†), which
could be due to the crystalline morphology with relatively larger crystal dimensions than usual. However, the emission intensities of the dual peaks do show a slight decrease in an oxygen atmosphere compared to vacuum, indicating the involvement of triplet states in the overall luminescence of the molecule.

To gain detailed information regarding the photophysical properties of the dual bands with ultralong emission, the time-resolved PL of crystalline o-Cz was recorded at elevated temperature. In general, when the temperature increases the phosphorescence intensity should decrease dramatically due to thermal quenching. Abnormally, the delayed emission intensity of o-Cz enhances as the temperature increases, especially for the CT bands, exhibiting linear plots of ln I (emission intensity) versus temperature (Fig. 3f and h). Within this temperature range, no thermally induced phase transitions occur (Fig. S10†). The decay curves (collected at 472 nm) of o-Cz displays prompt and delayed components (Fig. 3g and S11†) and the proportions of the delayed component increase significantly with shortened lifetime as the temperature increases from 297 K to 343 K. In addition, the intensity-dependent PL spectra (Fig. S12 and S13†) show a slope of 0.98 at low excitation density, which excludes consistent with the TADF phenomenon, the triplet annihilation process. These results are consistent with the TADF phenomenon but show a much longer lifetime. With regard to TADF, the appropriate energy level distributions are essential for efficient RISC from the lowest triplet state to the singlet state. The ΔE_ST values of o-Cz obtained in dilute THF solution, as well as the crystalline state (Fig. S14 and S15†), are 0.28 eV and 0.17 eV, respectively, which are close to the theoretical calculations (0.20 eV and 0.21 eV, respectively). The obtained values are sufficient for thermal activation. Therefore, thermally-activated RISC from T1 to S1 is expected to harvest triplet excitons and result in the enhancement of the structureless CT band around 472 nm with longer lifetime. The long-lived LE emission of o-Cz located around 522–700 nm shows red-shifted emission compared to the monomeric phosphorescence in dilute THF solution (Fig. S14b†), suggesting intermolecular electronic coupling within the crystalline lattice. Accordingly, single crystal X-ray diffraction analysis was performed to investigate the detailed molecular packing of o-Cz in the unit cell. o-Cz shows an ordered and condensed arrangement (Fig. S17 and S18†). The visual analyses of non-covalent interactions were carried out through an independent gradient model, which reveal the abundant interaction regions (green color) around adjacent molecules, as shown in Fig. S19†. Interestingly, the carbazole moieties stack tightly in a herringbone motif through the van der Waals interaction modes of C–H···π (2.894 Å) (Fig. S19†), which are driven by the electrostatic interaction between the electron-deficient hydrogen atoms and the π-electron density. The herringbone packing modes prevent further molecular close-packing upon excitation and lead to predominantly H-aggregate dimers. It has been reported that carbazole carbazole dimers in the solid-state can cause unwanted color variations in emission that could have negative implications for the device performance of TADF-OLEDs. The ortho-connectivity molecular design that is present in compound o-Cz should prevent these unwanted modes of dimerization and thereby be more desirable for TADF-based OLED devices with improved color purity. The H-dimers generate the intermolecular electronic coupling that can be verified by the TD-DFT result (Fig. S20†), affording the ultralong phosphorescence at 542 nm (T1) with a LE-character. The two emission bands with the CT and LE character demonstrate the existence of dual-channel triplet harvesting pathways (TADF and H-aggregation) in o-Cz, which minimize the triplet concentration quenching and result in a high ΦT of 16.6%. The dual-channel triplet harvesting shows a distinct advantage in the reduction of long-lived triplet quenching since the exciton concentration is very high with two-photon-excitation upon NIR laser irradiation.

Furthermore, the natural transition orbital (NTO) simulations and spin-orbital coupling (SOC) calculations of the excited states on both o-Cz (Fig. S21†) and p-Cz (Fig. S22†) were carried out to evaluate the (R)ISC channels, which play pivotal roles in TADF and phosphorescence materials. As illustrated in Fig. 4b, the two excited triplet states (T1 and T2) are located below the singlet state (S1) in the folded molecule o-Cz with narrow gaps of 0.04 eV (S1 to T1) and 0.21 eV (S1 to T2). In addition, the S1 state exhibits pure π–π* configuration, while the T1 and T2 consist of

![Fig. 4](https://example.com/fig4.png)

**Fig. 4** Theoretically-calculated energy levels and spin–orbit coupling constants between S0 and lower-lying Tn with the corresponding crystal geometries of (a) p-Cz and (b) o-Cz. Solid black arrows represent major channels with ξ over 1 cm⁻¹, and dashed black arrows represent minor channels with ξ less than 1 cm⁻¹.
The remaining triplet excitons are "trapped" by the H-aggregation \( (T_1') \) in the crystal lattice that results in the long-lived LE-state emission. The triplet excitons are harvested in a dual-channel (TADF and "H-trap") to reduce the concentration quenching of the long-lived excitons. Due to the intra- and intermolecular interactions in the crystal lattice, \( \alpha\text{-Cz} \) exhibits ultralong emission due to the suppression of non-radiative pathways. In contrast, the more linear molecule \( \beta\text{-Cz} \) exhibits a much lower phosphorescence efficiency (\( \Phi_T = 0.3\% \)) and vibronic fluorescence which indicates a decreased CT effect in the crystalline state (Fig. S27†), resulting in the faint emission after the NIR laser irradiation (Fig. S28†).

Conclusions

In conclusion, we designed and synthesized a spatially folded D–A type TPUOP molecule \( \alpha\text{-Cz} \), which exhibits SCT features and can be excited by UV, visible, and NIR light. The enhanced CT character increases the TPA ability and the reduced \( \Delta E_{CT} \) facilitates both ISC and RISC processes to increase the triplet harvesting in a dual-channel. Meanwhile, the non-radiative pathways are suppressed due to the synergy of intra- and intermolecular interactions. These features of \( \alpha\text{-Cz} \) contribute to an ultralong lifetime of 0.84 s and a high \( \Phi_T \) of 16.6\%, which is among the best reported for heavy-atom-free UOP materials. Most importantly, we demonstrate that the \( \alpha\text{-Cz} \) is an appropriate TPUOP molecule due to the enhanced CT character and efficient triplet harvesting. To the best of our knowledge, \( \alpha\text{-Cz} \) is the first example of a single component TPUOP molecule. This molecular architecture may provide a new platform for designing more TPUOP molecules to broaden their applications.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was financially supported by the National Natural Science Foundation of China (NSFC: 51803242, 51733010, 61605253 and 21672267), the Science and Technology Planning Project of Guangdong (2015B090913003), the China Postdoctoral Science Foundation (2017M620395) and the Fundamental Research Funds for the Central Universities. We appreciate Dr Shuang Li for help with the measurement of two-photon induced emission spectra and Dr Bingjia Xu for the further understanding of the results.
Notes and references