N-rich Electron Acceptor: Triplet Harvesting in Multichromophoric Pyridoquinoxaline and Pyridopyrazine-based Organic Emitters

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Abstract: Control of nonradiative deactivation of triplet states and tuning the singlet-triplet energy gap (ΔE_ST) are the major challenges to develop materials exhibiting thermally activated delayed fluorescence (TADF) and room temperature phosphorescence (RTP). Herein, we propose a new class of multichromophoric tridonor-acceptor (D_A−A) compounds with rigid and flexible π-spacer having N-rich pyridoquinoxaline (PQ) and pyridopyrazine (PZ) acceptor core, respectively. The molecule with carbazole (Cz) donors at meta to quinoxaline (OX) nitrogen of rigid PQ core exhibits TADF. Whereas, the variation of the linkage position of Cz to PQ as well as twisted and flexible PQ core show predominantly RTP due to relatively higher singlet-triplet energy gap (ΔE_ST). Increasing the donor strength with phenoxazine (PO) in PQ system leads to simultaneous TADF and RTP. Further, we demonstrate the promising scope of all-organic triplet harvesting materials in solid-state security encryption.

Smart organic donor-acceptor (D−A) based multichromophores have gained profound interest in recent years owing to the efficient harvesting of the triplet excited states.[1] The triplet state could be emissive by thermally activated delayed fluorescence (TADF) or the radiative decay at ambient conditions, commonly known as room temperature phosphorescence (RTP).[2] Such emitters are making their way in a wide variety of photonic applications like organic light-emitting diodes (OLEDs),[3] α) sensors,[4] security devices,[5] time-resolved luminescence imaging,[6] etc. The key challenges are to control the dynamic collisional quenching of triplet excited state and lowering the energy gap between the excited singlet (S1) and triplet (T1) states (ΔE_ST).[6a, 7] The lower ΔE_ST value promotes the photon upconversion from excited triplet states to the emissive singlet states through reverse intersystem crossing (RISC) leading to TADF.[10] The excited state intramolecular charge transfer (ICT) in donor-acceptor (D−A) molecules with spatially-separated the highest occupied molecular orbitals (HOMOs) and the lowest unoccupied molecular orbitals (LUMOs) is conducive to low ΔE_ST values. The CT interactions also favour a strong spin-orbit coupling (SOC) through the mixing of isostructural singlet and triplet electronic excited states.[8] Consequently, a high rate of intersystem crossing (ISC) along with stable triplet excited states leads to persistent or ultralong RTP in certain D−A compounds.[6a, 7]

The CT interactions depend not only on the molecular geometry but also vary with the relative orientation of D and A units.[9, 9] ICT significantly differs among positional isomers with different substitution positions of donors on the acceptor core.[10] Chen et al. demonstrated the switching between TADF and RTP in phenothiazine-benzophenone based D−A compounds invoking the concept of twisted intramolecular charge transfer (TICT) and planarized intramolecular charge transfer (PICT).[11] The TICT-based probe was shown to exhibit TADF. RTP was observed in a more rigid conformer exhibiting PICT. Bryce and coworkers observed RTP in phenothiazine–dibenzo[b,d]thiophene-S,S-dioxide based D−A−D pair by introducing steric hindrance on the donor units.[2b] The flexibility and the rigidity of the molecular backbone in the D−A−A pair significantly influence the charge transfer that may tune the TADF and RTP. Xie et al. reported delayed emission with high quantum efficiency in the red region (λem = 600 nm) with rigid planar dibenzo[a,c]phenazine as the acceptor core and 9,9-dimethylacridan as donor units.[11] Interestingly, the emission wavelength was tuned from green to red by increasing the number of donor units from D−A, D−A−A, to D−A architecture by controlling the ICT effect. Amidst several recent reports, a molecular design strategy unraveling the abstruse role of the central acceptor core along with conformational analysis in multichromophoric organic emitters for tunable and efficient TADF and RTP is still intriguing albeit challenging.

Herein, we have examined the role of N-rich acceptor units along with π-spacers in terms of flexibility/ rigidity and twisted/planar geometry on emission properties of D−A−A molecules at different timescale. The central core of pyridoquinoxaline (PQ) or pyridopyrazine (PZ) serves as facile acceptor units owing to the multiple nitrogen centers, which induces strong SOC, and thus enhance the ISC (Figure 1).[2c] Furthermore, the adjacent phenyl spacers also offer a distinctive synthetic advantage in regulating the molecular flexibility/ rigidity of the central core. The molecule with carbazole (Cz) donors at meta to quinoxaline (OX) nitrogen of rigid PQ core showed enhanced TADF due to low ΔE_ST. However, the higher ΔE_ST due to variation of the linkage position of Cz (para to OX nitrogen) to PQ and PZ core resulted in predominant RTP. The simultaneous TADF and RTP were achieved employing PQ as donors in PZ system. Tuning TADF and RTP through a rational design strategy of new D−A architecture, as demonstrated in the present communication, would contribute to develop advanced optoelectronic materials for security encryption and identification.

Figure 1. Design strategy of triplet harvesting materials: representative electron donor (D) and acceptor (A) based charge transfer compounds with D−A, D−A−A and D−A architecture. (b) Schematic illustration of our approach representing planar dibenzo[b,d]phenazine (T-shaped: PGT) and Y-shaped: PQV) and twisted diphenyldipyridopyrazine (FPZV) acceptor units, and (c) the molecular structures of tridonor-acceptor (D−A−D) compounds with carbazole (Cz) and phenoxazine (PO) as donor units: PQTc2, PQVCz, PZVc2 and PZVPO; D−A linkage, π-spacer and donor strength are varied.
Quinoxaline is a well-known electron acceptor unit due to the presence of two electron-negative sp² hybridized nitrogen atoms. Phenazine also possesses similar electron-deficient characteristics as quinoxaline. The presence of another nitrogen-containing heterocycle to the quinoxaline unit leads to the PQ motif, with better electron accepting capability than quinoxaline or phenazine. Similarly, PZ is more electron deficient than pyrazine. The suitable disposition of π-spacers (phenyl groups) leads to the development of dibenzopyridoquinoxaline and diphenylpyridopyrazine. The phenyl π-spacers do not alter the electron-withdrawing capabilities of PQ or PZ, rather allow synthetic flexibility and a range of viable options to introduce the donor substituents. Henceforth, dibenzopyridoquinoxaline and diphenylpyridopyrazine are abbreviated as PQ and PZ, respectively. The significant difference between PQ and PZ core lies in the molecular flexibility/ rigidity induced by the linkage pattern of phenyl spacers (Figure 1b). Donor substitutions at 2,7 and 3,6- positions of PQ lead to T and V-shaped molecular geometry. We introduce multiple electron donor units covalently linked with central PQ and PZ core for the development of a new series of multichromophoric D₃-A molecules. Considering the efficient electron-donating properties, high thermal stability and low-cost, Cz and PO were chosen as the donor units. Palladium-catalyzed Buchwald-Hartwig cross-coupling reactions (C-N coupling) of tribromo-PQT, PQV, and PZV with Cz and PO resulted in PQTCz, PQVCz, PZVCz, and PZVPO (Figure 1c). All the compounds were characterized by NMR spectroscopic and mass spectrometric analysis.

The D₂-A compounds absorb predominantly in the UV-region along with a prominent broad absorption band in the visible region. The lower-energy absorption bands are attributed to the extended π-ionization between the donor and the acceptor units. Whereas, the higher energy absorption bands are ascribed to π-π transitions. Strong fluorescence with broad emission bands were observed for PQTCz, PQVCz, and PZVCz in toluene (5 μM) at room temperature (RT, Figure 2a-c). A relatively weak emission was noticeable for PZVPO in toluene at RT (Figure 2d, strong D-A pair, dark TICT, *vide infra*). On increasing the solvent polarity from toluene to dichloromethane, a gradual bathochromic shift in the emission maxima was observed for all compounds. The emission maxima were found to be independent of the excitation wavelengths. The spectral broadening of the emission bands with increasing solvent polarity and positive solvatochromic behaviour are the two characteristic features indicative of strong ICT in the D₂-A compounds. Notably, the Stokes shift in PQTCz (m-substitution) is higher than PQVCz and PZVCz (p-substitution) in all solvents, which is in agreement with the more facile ICT in the former. The largest Stokes shift (150 nm, 5245 cm⁻¹) exhibited by PZVPO among the D₂-A compounds suggests the efficient CT interactions due to stronger donor characteristics of PO over Cz. The Lippert-Mataga plots further substantiate the ICT behaviour.

The fluorescence decays of all the compounds were measured in the solvents with varying polarity. A biexponential fluorescence decay for all the compounds at RT is possibly due to the emission from the locally excited (LE) and the charge transfer state. To get the decay associated emission spectra for a particular

![Figure 2](image-url)

**Figure 2** Time-resolved emission spectra (TRES) from nanosecond to millisecond timescale: normalized emission spectra of D₂-A compounds in toluene at room temperature (a-d), in toluene at 77 K (e-h) and in polymethyl methacrylate (PMMA) films at room temperature (RT: 22-25°C) (i-j), respectively for (a, e, i) PQTCz (Inset: spectrum at 1.5 ns depicting LE emission). (b, f, j) PQVCz, (c, g, k) PZVCz and (d, h, l) PZVPO. Spectra were recorded using λex = 465 nm [steady state (SS), 450 W Xe lamp], λex = 468 nm [pico second (ps) diode laser] and 461 nm [millisecond (ms) spectraLED] for PQTCz, PQVCz and PZVPO, respectively and λex = 420 nm (SS, 450 W Xe lamp), λex = 410 nm (ps diode laser) and 417 nm (ms spectraLED) for PZVCz. The solid bars indicate the similarity of emission peak position of D₂-A compounds at different timescale in solution at RT, 77K, and in polymer film.
timescale, the emission decays at 350 to 750 nm with the interval of 5 nm were measured. The reconstructed time-resolved emission spectra (TRES) of PQTCz at 1.5 ns and 18.9 ns show two distinct spectral profiles (Figure 2a). The emission spectrum obtained at 1.5 ns is similar to that of Cz emission in toluene (Figure 2a). It refers to the LE emission. The emission at 18.9 ns resembles the steady-state (SS) emission and is attributed to the charge transfer (CT) fluorescence (Figure 2a). Similarly, the TRES of PQVCz, PZVCz, and PZVPO in toluene at ambient conditions are shown in Figure 2b-d.

The phosphorescence spectra and associated decays of D2-A compounds were recorded at an ultralow temperature in liquid nitrogen (77 K) using 450 W Xe lamp [steady-state (ss)], picosecond (ps) diode laser and millisecond (ms) spectral LED as the excitation source. The ss emission spectra of D2-A compounds at 77 K showed emission bands with multiple humps (Figure 2e-h) and were significantly different compared to those obtained at RT (Figure 2a-d). The biexponential emission decays at 77 K at fast timescale were similar to that observed at RT due to the fluorescence from LE and CT states (Figure 2a-d). However, the decay measurements at slower timescale at 77 K revealed ultralong phosphorescence ranging from few milliseconds to 0.5 seconds (Figure 2e-h).

The phosphorescence emission maxima of PQTCz were observed at 585 and 590 nm with decay time 400 and 195 ms, respectively (Figure 2e). PQVCz emits with peaks at 565 and 590 nm with decay time 465 and 221 ms, respectively (Figure 2f). The phosphorescence emission peak maxima of PQVCz were noticeable at 535 and 545 nm with decay time 381 and 225 ms, respectively (Figure 2g). The phosphorescence decay time of PZVPO is 60 and 350 ms with a peak centered at 585 nm and 595 nm, respectively (Figure 2g), which are very close to the fluorescence bands observed in toluene at RT (585 and 610 nm, Figure 2d). The multieponential phosphorescence decays suggest more than one triplet charge-transfer state (ICT) due to the presence of multiple donor units. The lower ΔEST values (<0.5 eV) of D2-A compounds based on experimental data indicate the possible thermally activated RISC. The long-lived emission decay at RT with the oxygen-free condition of D2-A compounds in toluene revealed the prominent delayed emission characteristics of PQTCz. However, no longer component emission was observed for PQVCz, PZVCz, and PZVPO under the same condition.

Further, the D2-A compounds were doped in polymethyl methacrylate (PMMA) matrix to ascertain the triplet harvesting characteristics in the rigid media. All the compounds emit (ss) in the distinct regions of the visible spectrum (Figure 2i-l), ranging from cyan (PZVCz) to green (PQTCz and PQVCz) to red (PZVPO). The decays at the ns timescale monitored at the emission maxima indicate multiple emitting species at RT. It could be due to the emission from the singlet (LE, ICT), triplet (T1) states, and aggregates.23, 5a, 7c, 18 PQTCz film at RT exhibits emission peaks at 565 and 580 nm, with the decay time 139 and 42 μs, respectively (Figure 2i). The emission maximum at 565 nm obtained at μs timescale matches with TRES at RT in toluene (18.9 ns), indicating TADF in polymer film (Figure 2a and 2l). Whereas, it shows phosphorescence peak at 590 nm at 77 K in toluene (Figure 2e). The TRES measured using sensitive gated intensified charge-coupled device (ICCD) camera further ascertain the TADF of PQTCz film (Figure 3a). Unlike the toluene solution at RT, PQVCz film displays distinct longer component emission at 525 and 550 nm at decay time 414 and 135 μs, respectively (Figure 2j). It also shows phosphorescence (77 K) with peak maxima at 565 and 590 nm (Figure 2l). D-A compounds with flexible acceptor cores are likely to exhibit delayed fluorescence.19 We anticipated the same for D2-A compounds with flexible PZ core. The peaks at 510 and 545 nm with a respective decay time of 85 and 11 μs of PQVCz film (Figure 2k) resemble with phosphorescence peaks at 77 K (Figure 2g). Thus, PQVCz in PMMA film exhibits RTP. Similarly, PZVPO film shows long-lived decays with peaks at 570 and 590 nm having decay time 18.9 and 3.8 μs, respectively (Figure 2l). The origin of long-lived decay in PZVPO film at ambient conditions is presumably due to mixed RTP and delayed fluorescence (vide infra). Further, the laser power-dependent emission spectra using ICCD camera and decay were recorded to elucidate the long-lived components of D2-A compounds in a rigid polymer matrix (Figure 3a, b). A linear correlation of emission intensity with laser power for all the compounds signifies the unimolecular decay processes (Figure 3c), ruling out the triplet-triplet annihilation, as expected in TADF or RTP.

Figure 3 (a) Time-resolved emission spectra (TRES) with varying laser power (pulse width = 5 - 7 ns) of PMMA films of PQTCz at ambient conditions. Spectra were recorded using intensified charge-coupled device (ICCD) camera (λem = 355 nm). (b) The emission decay kinetics (λem = 355 nm, λex = 570 nm) of PQTCz-PMMA film with varying laser power at ambient conditions. (c) The linear fitting of the emission intensity of PMMA films of D2-A compounds as a function of excitation power (λem = 355 nm); the linear dependence of emission characteristic with laser power suggests unimolecular decay process.

TD-DFT calculations, coupled with crystal structure investigations, shed light on the intriguing photophysics of D2-A compounds (Figure 4a-c). The computed energy levels of the lowest singlet (E_S = 2.12 eV) and triplet (E_T = 1.99 eV) excited states of PQTCz are very close to the values obtained from the experimental data (Figure 4a). The lower ΔEST (0.13 eV) due to strong CT facilitated by the decoupled HOMO-LUMO levels enables the photon upconversion through RISC (Figure 4a). Thus, a prominent TADF was observed in the PQTCz film. Additionally, the similar nature of transition configurations and isosurfaces of S1 and other triplet excited states (T1 and T2) increase the probability of SOC, leading to high triplet yield.20 Whereas, a higher ΔEST (0.29 eV) and a greater number of energy transition channels (S1 to T1, T2, and T3) were observed for PQVCz leading to phosphorescence at ambient conditions (Figure 4a). On the other hand, PQVCz exhibits higher ΔEST (0.26 eV) and multiple ISC channels between S1 and T1 (Figure 4a), and phosphorescence was observed from PQVCz film. The molecular structure of PZVPO obtained from crystal structure analysis revealed a highly twisted geometry (Figure 4b). The dihedral angles between the central PZ core and the phenyl linker groups are 144.77° and 118.79°, while the dihedral angle between the π-spacers and phenoxazine donor unit is close to 80°. The donor (PO) units are connected directly to the acceptor core exhibiting a dihedral angle of 127.35°. Furthermore, a pair of CH···π (2.86 Å) and CH···HC (2.31 Å) intermolecular interactions between 11-
spacer aromatic phenyl groups generate a dimer like twisted structure along a axis (Figure 4c). These strong intermolecular interactions in the crystals of PZVPO could reduce energy loss through nonradiative relaxation channels. Additionally, a very low $\Delta E_{ST}$ value (0.02 eV) owing to the strong CT interactions coupled with long-lived triplet state contributed to the delayed fluorescence and radiative triplet emission in the rigid matrix at ambient conditions (Figure 4a).

Tunable fluorescence and RTP with the afterglow feature of D$_2$A compounds in polymer films were demonstrated through the encryption of FML (Figure 4d, written using PQVCz: F, PZVCz: M, and PZVPO: L). Greenish-yellow, cyan, and red fluorescence to greenish-orange, yellowish-green, and very weak red RTP, were observed, respectively, from PQVCz, PZVCz, PZVPO embedded PMMA films (Figure 4d). Further, we developed a facile information encryption pattern employing PZVCz thin film under ambient conditions (Figure 4e). The color-changing events were recorded using an image capturing Canon EOS camera. The film upon UV irradiation ($\lambda_{ex} = 365$ nm) showed intense cyan fluorescence. A distinctly contrasting greenish-yellow phosphorescence was observed by naked eye after turning off the UV light source (Figure 4e). Thus, the encrypted information is visible multiple times through switching on/off the UV light.

In summary, we developed a new class of multichromophoric tridonor-acceptor compounds with rigid and flexible π-spacers based on pyridoquinazoline and pyridopyrazine acceptor core, respectively. A strategy for tuning the TADF and RTP emission by controlling the charge transfer interactions through the variation of molecular geometry, as well as the relative orientation of donor and acceptor units, was demonstrated. The lower $\Delta E_{ST}$ value upon the $m$-substitution of donors on rigid pyridoquinazoline core facilitates RISC leading to delayed fluorescence. Whereas, the relatively higher $\Delta E_{ST}$ values and close packing induced by the $p$-substitution of donors on both pyridoquinazoline and flexible pyridopyrazine acceptor core, led to predominantly RTP. The potential afterglow feature of D$_2$A compounds was shown for security data encryption and identification. The present study elucidating the subtle role of D-A linkages coupled with the rigidity and flexibility of the new N-rich central acceptor core lays a promising foundation for further development of all-organic materials for triplet harvesting.

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