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The Contributions of Molecular Vibrations and Higher Triplet Levels to the Intersystem Crossing Mechanism in Metal-Free Organic Emitters.

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Intense, simultaneous, room temperature phosphorescence (RTP) and thermally activated delayed fluorescence (TADF) is observed in a series of donor-acceptor-donor (D–A–D) molecules. This dual-luminescence is stronger in the “angular” isomers, compared to their “linear” regioisomers, which is consistent with an enhanced intersystem crossing (ISC) in the former. Herein, we demonstrate that the small energy gap between the triplet levels, T_1 – T_n , below the lowest singlet state, S_1 , in the “angular” regioisomers, enhances the coupling between S_1 and T_1 states and favors ISC and reverse ISC (rISC). This is consistent with a spin-vibronic mechanism. In the absence of this “triplet ladder”, due to the larger energy difference between T_1 and T_n in the “linear” regioisomers, the ISC and rISC are not efficient. Remarkably the enhancement on the ISC rate in the “angular” regioisomers is accompanied by an increase on the rate of internal conversion (IC). These results highlight the contributions of higher triplet excited states and molecular vibronic



coupling to harvest triplet states in organic compounds, and casts the TADF and RTP mechanisms into a common conceptual framework.

INTRODUCTION

Metal-free organic compounds with dual emission from singlet and triplet excited states, e.g. fluorescence and phosphorescence at room temperature (RTP) or prompt and delayed fluorescence due to thermally activated delayed fluorescence (TADF),¹ have attracted great attention in recent years due to their potential for application in photonic and optoelectronic devices, including sensing of organic solvents² and oxygen,^{3,4} optical thermometry,⁵ optical storage,⁶ mechano-luminochromic materials,⁷ counterfeit labeling,⁸ luminescent probes for biological imaging,⁹ and organic light emitting diodes (OLEDs).^{10,11}

Detailed investigations aimed at revealing the fundamental mechanisms of TADF and RTP are also topics of strong research focus, because they offer excellent opportunities to investigate the dynamics of excited triplet states in organic materials.¹² These studies can, therefore, provide valuable information about the role of molecular conformation and connectivity, and coupling between electronic states for promoting triplet harvesting. These aspects are still not completely understood, and are extremely relevant when the properties of the triplet excited state is of importance, such as in photon up-conversion due to triplet-triplet annihilation,¹³ triplet harvesting in OLEDs,^{14–16} and organic spintronics.¹⁷

One of the major problems affecting current state-of-the-art OLEDs is the large roll-off observed in most devices. This is primarily caused by the long lifetime of the emitting excitons used in TADF molecules and heavy metal complexes, and also makes current metal-free RTP emitters unsuitable for application in OLEDs.



Consequently, developing strategies to accelerate the triplet harvesting rate in both metal-free TADF and RTP emitters is a major objective, and requires detailed investigations of the intersystem crossing (ISC) and reverse intersystem crossing (rISC) mechanisms.

The observation of dual fluorescence-phosphorescence at room temperature in pure organic materials is a rare phenomenon.¹⁸ This is because good fluorescent emitters generally show very small yields of triplet formation. At the other extreme, strongly phosphorescent molecules, mostly containing heavy metals, have triplet yields close to 100%, and thus fluorescence is not observed.¹⁹ A delicate balance between the intersystem crossing rate, radiative decay rates and the non-radiative internal conversion (IC) pathways is therefore required to generate simultaneous emission from the singlet and triplet states,^{18,20} as shown schematically in Fig. 1a). However, in most purely organic compounds the rate of intersystem crossing, from S_1 to T_1 , k_{ISC} , and the rate for radiative decay from T_1 , k_{ph} , are very slow, and cannot compete with fluorescence. Therefore, to achieve RTP, the triplet formation yield has to be enhanced, and the non-radiative deactivation pathways from T_1 to S_0 need to be suppressed.

Concurrently, similar conditions are required for efficient TADF to be observed. However, the two mechanisms differ in that efficient reverse intersystem crossing, k_{rISC} , is required for TADF but not for RTP, see Fig. 1a). TADF and RTP are therefore often seen as two contrasting mechanisms. We show here that this is not the case. Instead, the TADF and RTP mechanisms share the same fundamental requirements at a molecular level, and TADF or RTP are different responses obtained in a common framework, which can be tuned by selecting the host properties. This reduces the emphasis upon the energy gap between the emitting



singlet and triplet states, and crucially introduces the complementarity between the two mechanisms that otherwise would not exist. This complementarity means that it is feasible that both the TADF and RTP mechanisms can be operative in the same molecule, as demonstrated from Cu(I) complexes,²¹ and recently in a pure organic emitter.²² However, the emergence of this dual luminescence is strongly dependent on the ability to fine tune the kinetic parameters, which can be achieved by the molecular structure of the emitter and the host in which it is dispersed. Interestingly, in dual emitters showing simultaneously both long-lived forms of luminescence, the ratiometric nature of the dual-emission would be preserved during the entire luminescence lifetime, which makes these materials very convenient in gated-acquisition luminescence methods for sensing/imaging. Here, the ratio between the RTP and TADF intensities is preserved even in the microsecond time range, and can be used as an internal calibration. This is in clear contrast with conventional RTP and TADF emitters where the prompt fluorescence decays in just a few nanoseconds, and the phosphorescence or delayed fluorescence lives for microseconds or milliseconds.

RESULTS AND DISCUSSION.

The development of pure organic RTP compounds has progressed significantly in recent years. These have adopted a number of different strategies, but have mainly targeted restricting the intramolecular vibrations of the emitters using intermolecular interactions, either in rigid hosts,^{10,23,24} crystalline form,^{8,25,26} or induced by aggregation.⁸ All of these methods restrict thermal vibrations and increase the phosphorescence lifetime and quantum yield. This suppresses the rate of T_1 non-radiative decay, which in organic materials is usually faster than the radiative rate. However, it is not clear whether or not the intersystem crossing rate is



also affected. Interestingly, RTP has been reported recently in a brominated fluorene derivative,²⁷ and deuterium substitution has been also used to diminish internal conversion due to molecular vibrations, and thus enhance RTP emission.^{20,24}

Despite this strong interest in RTP materials, dual fluorescence-phosphorescence at RT has only been reported in general when the emitter is dispersed in a few selected hosts, which are able to suppress vibrations, such as poly(vinylalcohol), poly(methylmethacrylate), β -estradiol, micelles, cyclodextrin or inorganic crystals.^{18,20} Remarkably, reports of RTP in conjugated hosts are extremely rare, which has limited their application in OLEDs. To the best of our knowledge only two publications report RTP in conjugated hosts.^{10,14}

A comprehensive investigation of molecular structure-property relationships is, therefore, still needed to guide synthetic strategies for the design of more efficient RTP emitters.^{28,29} There is no clear understanding how the host-guest interactions influence the intersystem crossing rate and the pathways for non-radiative decay of the excited states. For example, it is not clear why some molecules give intense RTP when dispersed in particular hosts but not in others, or why some molecular structures give intense RTP and structurally similar analogues give only fluorescence.^{20,29} This clearly shows that besides suppressing internal vibrations, rational molecular design of emitters and hosts can be used to tune the electronic energy levels and activate RTP. However, while manipulation of electronic properties to achieve RTP is very attractive, it is also extremely challenging.²⁹

In contrast, guidelines for designing efficient TADF molecules are beginning to emerge,^{1,30} and delayed fluorescence is now routinely observed in organic compounds that comprise covalently linked electron donor (D) and electron acceptor (A) units, in solution and also in several solid hosts, including those with extended



conjugation. In addition, the fine details determining the mechanism for efficient thermally activated reverse intersystem crossing in TADF emitters are beginning to be understood.^{16,31,32,33,34} This includes work illustrating that the energy alignment between the singlet state with charge transfer character and a local triplet state was proposed as a key requirement for efficient TADF.^{16,31} However, not all TADF materials may have a local triplet state in near resonance to the CT states, which may be particularly valid in blue TADF emitters, where the singlet emissive state has to be kept at high energies. Therefore, more general requirements are required to explain the observation of TADF. Moreover, in the design of most TADF emitters, the attempts to minimize the singlet-triplet energy gap often have a negative impact on the fluorescence yield. Strategies to design TADF molecules with “strong” fluorescence yield and shorter luminescence lifetimes are, therefore, required.³³ Moreover, as in OLEDs the triplet yield is fixed at 75%, there is still potential to improve the performance of OLED emitters by achieving a proper balance between the yield of reverse intersystem crossing and the fluorescence yield.³⁵ These are issues that need to be resolved and require detailed understanding of the mechanisms of triplet harvesting at the molecular level.

Herein, we explore factors influencing the rate of intersystem crossing and reverse intersystem crossing in pure organic emitters. We show that simple manipulations of the way electron donor (D) and electron acceptor (A) units are linked in D–A–D compounds have a profound influence on the activity of both RTP and TADF. When the emitters are dispersed in suitable hosts and depending on the polarity of the local surroundings, simultaneous RTP and TADF are observed. This illustrates the similarity of the two mechanisms and properties required to facilitate high intersystem crossing and reverse intersystem crossing rates in organic



molecules and brings TADF and RTP into a common conceptual framework. Crucially, these synergies can be used to create and guide the design of novel emitters.

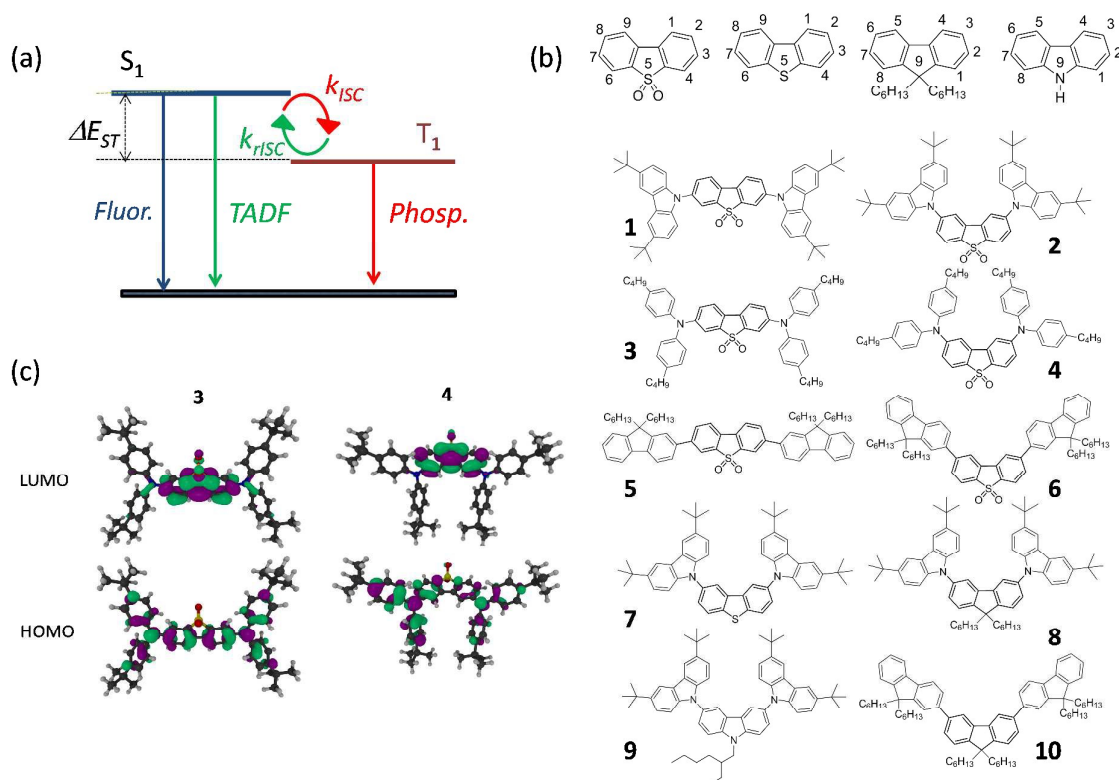


Figure 1 – (a) Energy level diagram describing the photophysical mechanisms that originate RTP and TADF emissions. (b) Molecular structures of the compounds used in this work, and numbering code for molecular isomers used in this work (top). (c) Frontier molecular orbitals HOMO and LUMO for compounds **3** and **4**.

Figure 1b shows the molecular structures of the compounds studied herein. Previously we have reported strong TADF emission in ethanol solution for compounds **2** and **4** at room temperature, and intense dual fluorescence-phosphorescence in compounds **2**, **4** and **6**, in ethanol solution at 100 K.³⁶ The dual fluorescence-phosphorescence in these compounds is so intense that it is observed in a normal fluorimeter without using gated luminescence methods. In contrast, their



regioisomers, **1**, **3** and **5**, exhibit no phosphorescence, even at low temperature and their delayed fluorescence in ethanol solution at 300 K is due to triplet-triplet annihilation,³⁶ (see Fig. S1 and S2). This markedly different isomer effect was not understood at the time, and an intermediate triplet level, between the singlet and lower triplet states, was introduced *ad-hoc* to justify the enhanced TADF in **2** and **4**. However, this reveals little information about the mechanism sustaining reverse intersystem crossing. In the present work, we unravel fundamental concepts on the way triplets are harvested in purely organic compounds, providing new understanding to the mechanisms of RTP and TADF in general.

The only difference in the molecular structure of these regioisomers is the linking position of the carbazole, diphenylamine and fluorene electron donor (D) units and the dibenzothiophene-*S,S*-dioxide electron acceptor unit (A). In **1**, **3** and **5**, the acceptor is substituted at positions C-3,7, whereas in **2**, **4** and **6**, the substituents are at positions C-2,8 of the A unit. For easy identification, these different isomers are hereafter referred to as “*linear*” and “*angular*”, respectively. Despite this seemingly small change, the effect on the dynamics of the excited states is very significant in the context of RTP and TADF luminescence. Compounds **7**, **8** and **9**, also with “*angular*” substitution, show intense dual fluorescence-phosphorescence at 100 K in ethanol solution, but in the “*angular*” compound **10** no phosphorescence is observed.³⁶ This illustrates that the presence of heteroatoms is crucial for the observation of dual fluorescence-phosphorescence, as previously identified by others.²⁸ However, this is not the only required structural feature for the observation of phosphorescence, as confirmed by the absence of strong phosphorescence in the “*linear*” compounds **1**, **3** and **5**. Intrinsic molecular-structure relations must therefore be tailored to create intense RTP and TADF emitters.²⁹



Absorption and emission in solution. The absorption and emission spectra of compounds **1** and **2**; **3** and **4**; **5** and **6**; and of **7**, **8**, **9** and **10** in hexane solution are shown in Fig. 2. The absorption of the “linear” compounds **1**, **3** and **5**, is clearly red-shifted compared to their “angular” counterparts **2**, **4** and **6** respectively. This is expected from the shorter conjugation length of the “angular” emitters, which pushes the singlet and triplet states to higher energies, see also Table 1. Compounds **7**, **8**, **9** and **10** are all with “angular” geometry, and based upon a carbazole donor unit, (except **10**), but differ in their acceptor units. All of these show blue shifted absorption, similar to **2**, **4** and **6**. They also have less excited state charge transfer character, as evidenced by their blue-shifted and well-resolved emission, see Fig. 2f. This is due to a lower electron affinity of fluorene, dibenzothiophene, and carbazole units compared to dibenzothiophene-*S,S*-dioxide unit.

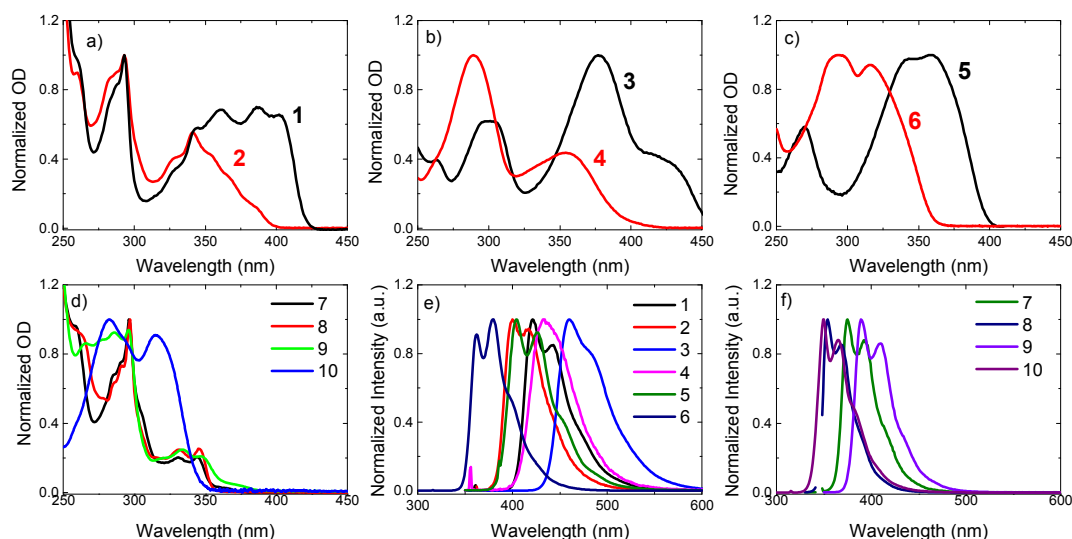


Figure 2 -Absorption spectra of (a) **1** and **2**, (b) **3** and **4**, and (c) **5** and **6**, and (d) **7**, **8**, **9** and **10**. Emission spectra of compounds (e) **1** to **6**, and (f) **7** to **10**. All spectra were collected in dilute hexane solution at room temperature.



Table 1 shows the photophysical data in solution for compounds **1** to **9**. The triplet energies of compounds **7** to **9** are high, above 2.8 eV, and in the case of **7** as high as 3.02 eV. For compounds **1** to **6**, the triplet state appears at lower energies, ranging from 2.20 eV in **3** to 2.70 eV in **2**. As a general trend, the “angular” substitution shifts the singlet and triplet states to higher energies relative to the “linear” counterparts, i.e. **2**, **4** and **6** have higher singlet and triplet energies than **1**, **3** and **5** respectively. However, as the change in the position of substitution affects the singlet and triplet energies almost equally, the singlet-triplet energy gaps of the corresponding isomers are very similar in non-polar hexane. Because the singlet and triplet states are affected in a similar manner, this indicates that both emitting states (S_1 and T_1) are of similar electronic character. In the present series, compounds **7** and **9** have the smallest $\Delta E_{S_1-T_1}$ gap, around 0.4 eV, and compounds **5**, **6** and **8** have the largest $\Delta E_{S_1-T_1}$ gap, above 0.8 eV.

Table 1 - singlet and triplet energies, S_1 and T_1 , fluorescence quantum yields, Φ_f , fluorescence lifetime, τ_f and triplet yields, Φ_T . All determined in hexane at room temperature, with exception of T_1 energies, which was determined at 80 K in ethanol solution.

	S_1 $\pm 0.02(\text{eV})$	T_1 $\pm 0.02(\text{eV})$	ΔE_{ST} $\pm 0.03(\text{eV})$	Φ_f ± 0.05	τ_f $\pm 0.05(\text{ns})$	Φ_T ± 0.1	$k_f \times 10^8$ (s^{-1})	$k_{ISC} \times 10^8$ (s^{-1})	$k_{IC} \times 10^8$ (s^{-1})
1	3.07	2.54	0.53	0.94	3.08 ^a	-	3.1	-	0.2
2	3.24	2.70	0.54	0.10	1.60 ^a	0.32	0.6	1.9	3.6
3	2.84	2.20	0.64	0.72	3.66 ^a	-	1.9	-	0.8
4	3.05	2.48	0.57	0.10	1.13 ^a	0.19	0.9	1.7	6.3
5	3.21	2.36	0.85	0.80	1.05 ^a	-	7.6	-	1.9
6	3.54	2.48	1.06	0.39	1.02 ^a	0.41	3.8	4.0	1.9
7	3.42	3.02	0.40	0.05	0.88	0.42	0.6	4.8	6.0
8	3.62	2.82	0.80	0.22	3.81	0.26	0.6	0.7	1.4
9	3.33	2.97	0.36	0.14	3.99	0.33	0.4	0.8	1.3

^(a)obtained from reference [36]

The fluorescence yields follow a clear pattern. The “angular” compounds that show intense phosphorescence at low temperature also show weak fluorescence. The fluorescence lifetimes are also significantly shorter in the “angular” substituted compounds, when compared with their “linear” counterparts, see compounds **1** to **6**



in Table 1. The yield of triplet formation is also significantly higher in the “angular” oligomers. While isomers **1**, **3** and **5** show no triplet population at room temperature in hexane solution, their “angular” analogues show large triplet yields. This occurs despite the singlet-triplet energy gap being very similar. Marked differences are also observed in the rate constants for radiative decay, k_f , intersystem crossing, k_{ISC} , and non-radiative internal conversion, k_{IC} . Remarkably, the radiative decay rate decreases significantly when going from “linear” to “angular” form. This drop in the radiative rates is accompanied by an increase on the intersystem crossing and internal conversion rates.

In summary, despite not significantly affecting the S_1 - T_1 energy gap, the “angular” substitution is very effective in decreasing the radiative rate and accelerating the rates of intersystem crossing and internal conversion. This explains the observation of strong phosphorescence in the “angular” isomers, as at low temperatures internal conversion is greatly suppressed.

The simultaneous increase in internal conversion and intersystem crossing is a general rule that points to a common intermediary state for both processes, i.e., a low lying CT state responsible for both increased internal conversion and intersystem crossing. However, closer inspection shows another important aspect to take into account when designing novel TADF/RTP emitters. Comparison of **2**, **4** and **6**, where the acceptor is conserved, shows that there is a significant increase in the rate of intersystem crossing for **6** with respect to **2** and **4**. This is accompanied by a decrease in the internal conversion rate to the ground state. While in **6** the donor and acceptor moieties are linked through a $C(sp^2)$ - $C(sp^2)$ bond, in **2** and **4**, a $N(sp^3)$ - $C(sp^2)$ bond performs the same function. This difference impacts both in the vibrational deactivation of the singlet state and the donor-acceptor coupling needed



to obtain the CT state. While the main degree of freedom in **6** is rotation around the C-C bond coupling the donor and acceptor groups, additional vibrational modes related to inversion around N (sp^3 - sp^2 transition) exist in **2** and **4**. The contribution of these additional vibrational modes can be further observed when comparing **2** and **4**. In **2**, the donor group is a carbazole unit, which is more rigid than the diphenylamine unit in **4**. Increasing the flexibility of the donor group (and on the donor-acceptor linking motif) yields higher k_{IC} and lower k_{ISC} .

As previously stated, the change in the substitution from “*linear*” to “*angular*” has the strongest influence on the yield of triplet formation, giving stronger triplet yields in **2**, **4** and **6** and activating intense phosphorescence that is not observed in the linear isomers **1**, **3** and **5**. Strong dual fluorescence-phosphorescence is also observed in ethanol solution at low temperature in compounds **7**, **8** and **9**, see Fig. S1, showing that the “*angular*” substitution is equally effective at giving dual fluorescence-phosphorescence, even if sulfone functionality is not present. However it is noted that when the carbazole donors in **8** are replaced by fluorene units, i.e. no heteroatoms exist in the trifluorene structure (**10**), the phosphorescence is switched off, and only fluorescence is observed. The presence of heteroatoms in the “*angular*” molecular structures is, therefore, of fundamental importance for the observation of strong phosphorescence in these materials, but not a sole requirement.

Compounds **2** and **4** in ethanol solution have a singlet state of charge transfer character (CT) at much lower energies than the emissive S_1 state in hexane and other non-polar environments. In this case, the $\Delta E_{S_1-T_1}$ energy gap becomes very small in polar environments, around 0.1 eV and 0.21 eV in ethanol, for **2** and **4**, respectively. As the reverse intersystem crossing rate, k_{rISC} , and the intersystem crossing rate, k_{ISC} , are proportional to each other,³⁵ the “*angular*” isomers have also



stronger reverse intersystem crossing rates than their “linear” counterparts. This isomer effect and the smaller $\Delta E_{S_1-T_1}$ energy gap created under the influence of the stronger polarity in ethanol, explains why these compounds show strong TADF in ethanol solutions, while compounds **1** and **3**, are not TADF active.³⁶ Compound **2** was used previously in OLEDs giving EQEs of 14%.³⁷

The correlation between the ISC and IC rates that is observed is also important when comparing compounds **2**, **7**, **8** and **9**. These compounds all have the same carbazole donor, but different acceptors, where the presence of C, N, SO₂ and S promotes increasing ISC. Figure 3 shows that the rates of ISC and IC in these compounds show a clear positive correlation, i.e. increasing the ISC rate is linked with increasing the IC rate. This points to a similar mechanism and would suggest that the vibrational modes responsible for IC also work to enhance ISC, by spin-vibronic coupling and mixing low lying triplet states and S₁. This is consistent with recent findings where considering T_n, as well as S₁, T₁ and S₀ are essential to understand the exciton decay dynamics in TADF emitters.^{12,16}

It is also important to highlight that this effect is not correlated with the charge transfer character of the singlet and triplet states. Figure 3, shows a graph of the IC rate against the ISC rate for compounds **2**, **7**, **8** and **9**. The charge transfer character of the singlet excited state, as evaluated by the fluorescence solvatochromism, is represented by the size of each dot. Compound **2** with the strongest CT character in the series shows intermediate values for both ISC and IC rates. Therefore, the variation in the ISC and IC rates is not due to increasing the CT character of the excited state.



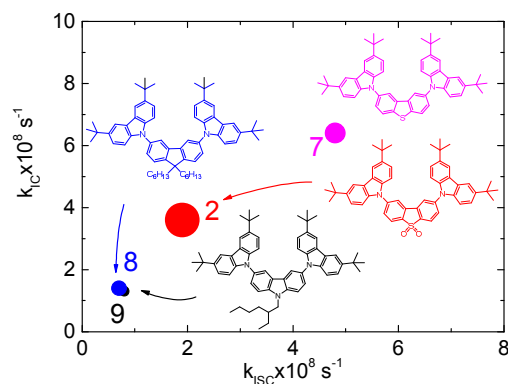


Figure 3 –Dependence of the IC rate with the ISC rate in the series of compounds **2**, **7**, **8** and **9**. All with the same donor (carbazole) but different acceptor. The size of the dot represents the CT character of the singlet state as evaluated by the solvatochromism in the fluorescence spectra.

The data in Table 1 gives significant information regarding the way the change in molecular connectivity influences the intersystem crossing and internal conversion rates. However, we have still not explained what causes the intersystem crossing rate to increase in the “angular” compounds. This crucial point is addressed in the next sections.

Room Temperature Phosphorescence in zeonex films. The steady state emission spectra of compound **2** dispersed in zeonex film is shown at 300 K in Fig. 4a), and in Fig. 4b) represented as a function of temperature.



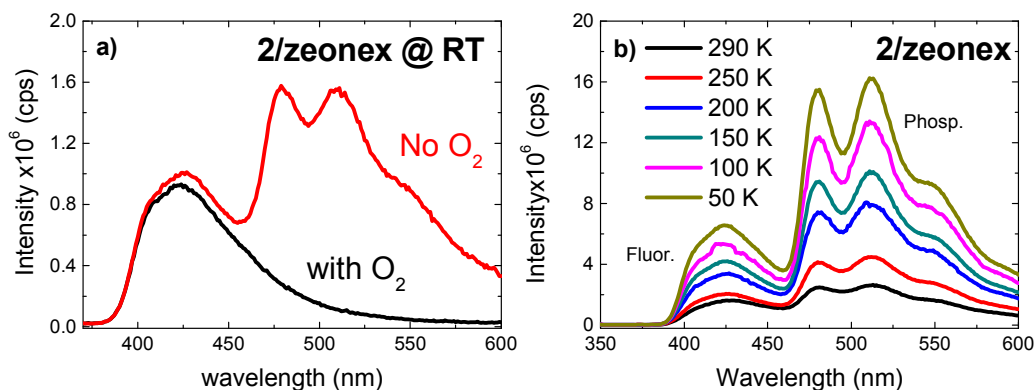


Figure 4 –Steady-state emission spectra of **2** in zeonex at (a) room temperature, and (b) as a function of temperature.

In degassed conditions, compound **2** dispersed in zeonex gives dual emission, i.e. strong fluorescence and phosphorescence at room temperature. The room temperature phosphorescence (RTP) emission is so strong that it is easily measured in a normal fluorimeter, without the need of using time-gated acquisition methods and is higher in intensity than the fluorescence. Strong RTP is also observed for compounds **4**, **6**, **7** and **8**, see Fig. 5, but no RTP is observed in compounds **1**, **3**, **5**, and **9**. However, the fluorescence in **9** is strongly quenched by oxygen, see Fig. S3. This indicates that there is a significant delayed fluorescence (DF) contribution to the fluorescence in **9**, which involves triplet states. Compound **9** has the smallest $\Delta E_{S_1-T_1}$ energy gap in the series, and triplet harvesting through reverse intersystem crossing may be active. Note that the excited states of **9** have no CT character.

A clear trend emerges from Fig. 5. Intense RTP is observed only for the "angular" compounds and is much stronger in **2**, **4** and **7** than in **6** and **8**. In the latter compound the low RTP emission is explained by a weaker ISC rate, which could be potentially due to the lack of heteroatoms in the fluorene acceptor. However, this is not the case in **6**. For this compound, the ISC rate is strong, and the IC rate is



weaker than in **2** and **4**. Therefore, compound **6** should be a strong RTP emitter, but it is not. When compared with the other “*angular*” compounds, **6** is different due to a higher radiative rate in the S_1 , k_f , and significantly larger $\Delta E_{S_1-T_1}$ gap. Moreover, compound **6** shows very long phosphorescence lifetime in ethanol at 100 K.³⁶ This indicates that the radiative rate of **6** in the T_1 state is significantly slower than in compounds **2** and **4**, which contributes to the decrease in the phosphorescence yield.

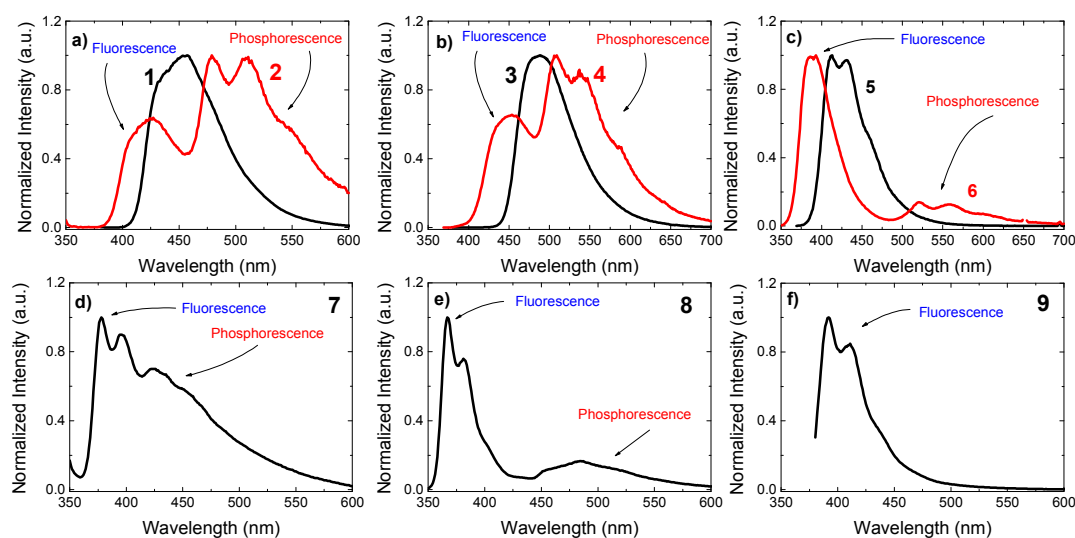


Figure 5—Steady-state emission spectra for compounds **1** to **9** dispersed in zeonex at RT, in vacuum.

Remarkably, compounds that are substituted at the C-3,7 positions of the dibenzothiophene-*S,S*-dioxide unit, i.e. “*linear*”, show no RTP at all. This is evidenced in Fig. S8, where compounds **1** and **3** show no RTP emission even when dispersed in poly(4-bromostyrene), a host that is known to enhance triplet formation due to the external heavy-atom effect.³⁸ Therefore, the “*angular*” substitution and the presence of the low atomic weight heteroatoms, sulfur or nitrogen, are clearly fundamental for strong RTP observation in pure organic materials.



The strong quenching effect of oxygen on the red-shifted emission band in compounds **2**, **4**, **6**, **7** and **8**, shown in Fig. S3, confirms that the extra emission band is due to phosphorescence at room temperature. Also in zeonex, compounds **2** and **4**, show significantly shorter fluorescence decays than **1** and **3**, see Fig. S4 in SI. This is entirely consistent with the stronger triplet formation yield, see Table 1.

We turn now to discuss compounds **1–4**, where the regioisomerization effect is strongest, to understand the reasons why these compounds have such marked differences in regard of their RTP emission. A summary of the photophysical data in zeonex film, concerning the observation of RTP, is given in Table 2. With decreasing temperature the intensities of both the fluorescence and phosphorescence bands increase, see Fig. 4b). This clearly indicates that the quenching effect due to vibrations is being suppressed. Moreover, while the phosphorescence in the presence of oxygen is, as expected, completely quenched, see Fig. S3, there is practically no difference between the intensities of the fluorescence band of spectra collected in the presence of oxygen and in degassed conditions (with the exception of compound **9**, as discussed). This shows that triplet harvesting due to TADF is not operative in these compounds when dispersed in zeonex.

Table 2—Fluorescence and phosphorescence quantum yields, Φ_f and Φ_{ph} respectively, fluorescence and phosphorescence lifetimes, τ_f and τ_p respectively; rates of radiative decay for the S_1 and T_1 states, k_f and k_{ph} respectively; rates of internal conversion, k_{IC} , and intersystem crossing, k_{ISC} , in the S_1 ; and rate of intersystem crossing from T_1 to S_0 , $k_{ISC}^{T_1}$. All determined in zeonex films at room temperature.

	Φ_f ± 0.05	Φ_{ph} ± 0.05	Φ_{ph}/Φ_f ± 0.07	τ_f ± 0.03 (ns)	τ_p ± 0.5 (ms)	$k_f \times 10^8$ (s^{-1})	$k_{ISC} \times 10^8$ (s^{-1}) ^b	$k_{IC} \times 10^8$ (s^{-1}) ^b	k_{ph} (s^{-1}) ^b	$k_{ISC}^{T_1}$ (s^{-1}) ^b
1	0.93	-	-	4.75	-	1.9	-	0.1		
2	0.09	0.26	2.9	1.68	34.3 ^a	0.5	1.9	3.5	23.7	5.5
3	0.56	-	-	4.71	-	1.2	-	0.9		
4	0.06	0.14	2.3	0.98	29.8	0.6	1.9	7.6	24.7	8.9

^aThe phosphorescence decay in **2** is bi-exponential, the value given is the average lifetime.

^bDetermined using the triplet yield in solution.



Table 2 shows a consistent trend with the data discussed in Table 1. The intersystem crossing and internal conversion rates are higher in the “*angular*” isomers when compared with their “*linear*” counterparts, even in solid zeonex film.

Calculations. The Kohn-Sham HOMO-LUMO orbitals for compounds **1–4** are shown in Fig. S6. These dominate the lowest transition responsible for the S_1 state. All four compounds show charge transfer (CT) character, which can be quantified using the absolute orbital overlap $O = \langle |\psi_{\text{LUMO}}| | |\psi_{\text{HOMO}}| \rangle$, for which smaller overlaps implies stronger CT character.³⁹ Therefore the CT character at the ground state geometry is larger for compound **2** ($O=0.41$) than **1** ($O=0.49$), and for compound **4** ($O=0.58$) than **3** ($O=0.67$). To determine the excited state and emission properties, the molecules were optimized in the lowest triplet state using Unrestricted Kohn-Sham (UKS). Importantly, the oscillator strength for the S_1 state is over an order of magnitude larger for compound **1** ($E_{S_1}=2.64$ eV, $f=0.9053$) than **2** ($E_{S_1}=2.75$ eV, $f=0.0276$). This corresponds to a radiative lifetime for **1** of ~ 3 ns, in good agreement with the experimentally determined fluorescence lifetime, shown in Table 1. In contrast, the smaller oscillator strength of **2** corresponds to an excited state lifetime of ~ 50 ns. As the fluorescence lifetime reported for compound **2** is < 2 ns (Table 1), this illustrates that the S_1 state in **2** is susceptible to other processes, namely ISC, consistent with the observation of phosphorescence in **2**. In addition, the energetics agree well with the maxima of the emission spectra shown in Figure 5. The same trend is observed for structures **3** ($E_{S_1}=2.77$ eV, $f=1.0109$) and **4** ($E_{S_1}=2.79$ eV, $f=0.020$). Again the oscillator strength of the S_1 states is much smaller in **4** than in **3**, allowing ISC to out-compete radiative decay.

To further interrogate the mechanism that is causing enhanced ISC in **2** and **4**, we turn to the energy diagram in Fig. 6. Within each pair, the “*angular*” substituted



compounds, **2** and **4**, show in the calculations consistently smaller $\Delta E_{S_1-T_1}$ gaps than their “*linear*” counterparts. However, this does not “*per se*” explain the observation of strong RTP emission (or TADF emission). When comparing across the four structures, compound **4**, a good RTP emitter in zeonex (and a good TADF emitter in ethanol solution at room temperature) has $\Delta E_{S_1-T_1}$ of 0.66 eV, whereas compound **1**, which do not show neither RTP or TADF in any medium, has similar $\Delta E_{S_1-T_1}$ of 0.64 eV. Therefore, clearly the $\Delta E_{S_1-T_1}$ gap is not the dominant effect, and probably more pertinent is the gap T_1-T_n , between the manifold of triplet states closest to the S_1 state, from where a clear pattern emerges between the “*linear*” and “*angular*” compounds.

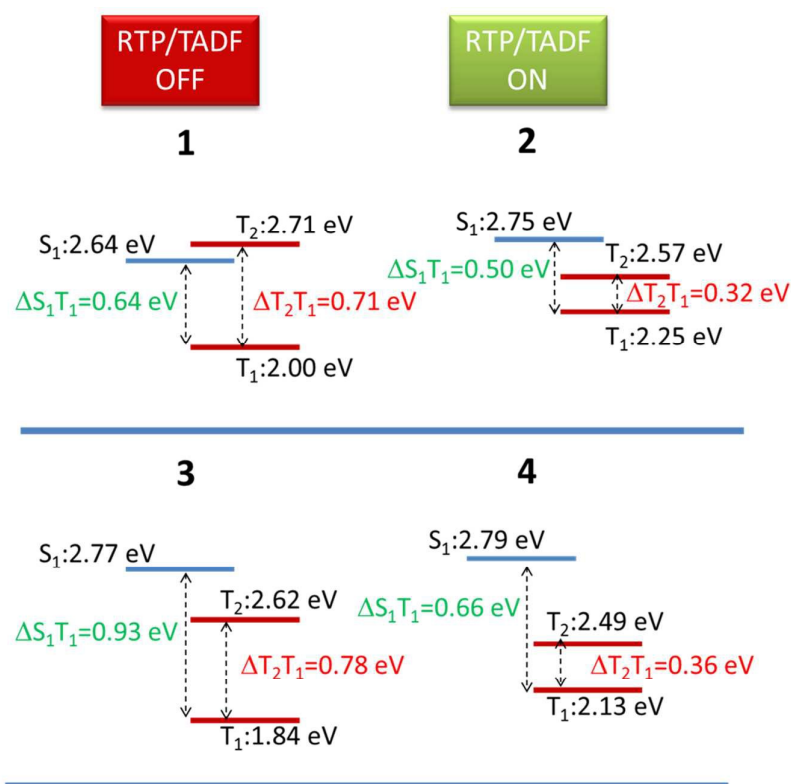


Figure 6-Energy level diagram of the singlet and triplet states in compounds **1–4**, obtained from TDDFT calculations.



For both **1** and **3**, which do not exhibit RTP or TADF, the gap between the T_1 and T_2 states is large (>0.7 eV), whereas for **2** and **4**, both good RTP/TADF emitters the gap between T_1 and T_2 is small (~ 0.3 eV). This provides the potential for efficient internal conversion between the triplet states and a decay route of the initially populated triplet to the lowest, T_1 , triplet state to emit, according to Kasha's rule. In this case, recent work has also demonstrated that the spin-vibronic mechanism, involving both spin orbit coupling and nonadiabatic coupling^{31,32} significantly increases the rate of ISC. The close proximity of all of the triplet states make this mechanism much more likely in **2** and **4**, and in combination with the slower radiative rates of the S_1 state in **2** and **4**, these data sets provide strong support for the observation and mechanism for RTP. This mechanism also explains why compounds **2** and **4** are good TADF emitters in ethanol. In this case, the strong solvent polarity shifts the S_1 to lower energies, without affecting significantly the energy of the T_1 state, which is not of CT character. This makes the $\Delta E_{S_1-T_1}$ gap smaller and facilitates rISC. To test this hypothesis compounds **2** and **4** were dispersed in hosts of stronger polarity than zeonex aiming to shift the S_1 to lower energies. With the correct tuning of the energy levels dual luminescence, RTP and TADF, is observed, as shown below. This again casts the TADF and RTP mechanisms into a common conceptual framework, not previously described.

Dual luminescence, RTP and TADF, in organic hosts. Here, compounds **7** and **9**, and two other large band gap hosts TSPO1,⁴⁰ and poly(4-bromostyrene),³⁸ are used as hosts in blends with compounds **2** and **4**; see Fig. S7 for the molecular structures of TSPO1,⁴⁰ and poly(4-bromostyrene). All of these host materials have a triplet energy higher than **2** and **4**, and are able to act as donors to promote energy transfer



to compounds **2** and **4**. TSP01 in particular has been used as a host in blue OLEDs.⁴⁰

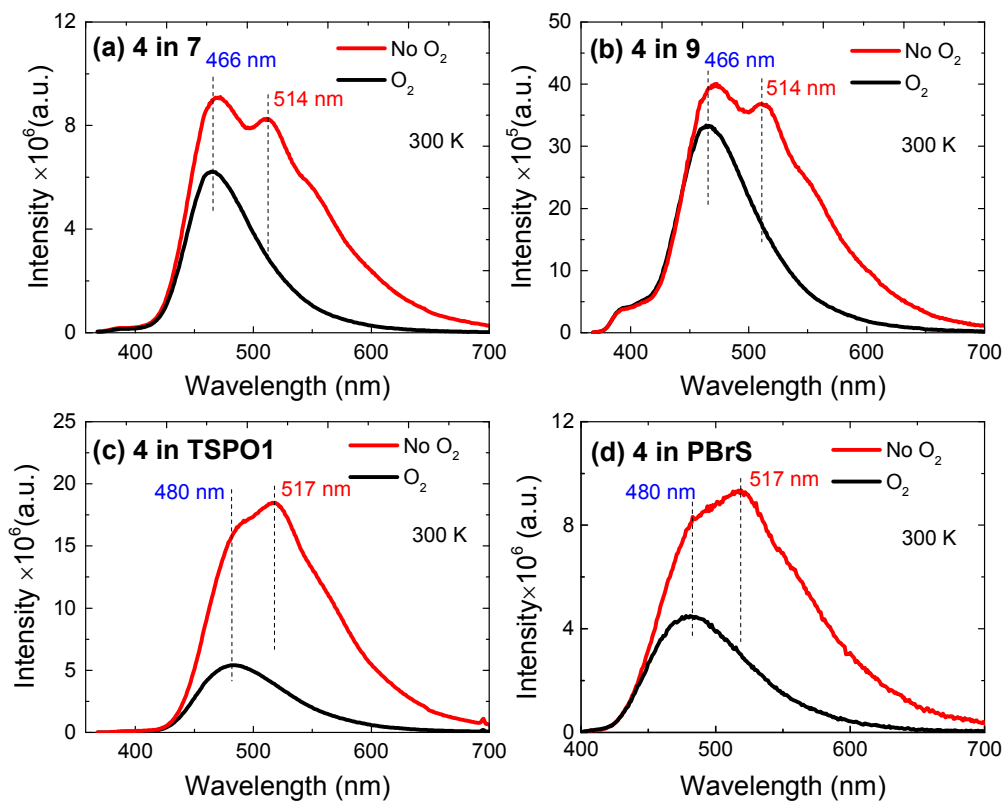


Figure 7-Steady-state emission spectra of compound **4** dispersed in different hosts (a) **7**, (b) **9**, (c) TSP01 and (d) poly(4-bromostyrene), identified as PBrS, in vacuum (red) and in air equilibrated (black) conditions.

In Figure 7, the steady-state emission spectra of compound **4** in blends with the hosts mentioned above are compared in vacuum and air. The emission spectra are obtained at 300 K in a normal fluorimeter. The presence of dual-luminescence is clearly observed. The phosphorescence band in **4** peaks at around 515 nm in the four hosts, 7 nm below the phosphorescence maximum observed in zeonex. The fluorescence band peaks at 466 nm for blends of **4** with **7** and **9** and at 480 nm in



TSPO1 and PBrS. This represents a shift of 16 and 30 nm to the red in comparison with the fluorescence maximum in zeonex at 450 nm.

Interestingly, when compared with the emission in zeonex, where no variation on the intensity of the fluorescence band is observed when oxygen is removed, in blends of **4** with **7**, **9**, TSPO1 and in PBrS, a marked increase on the fluorescence intensity occurs upon removing oxygen. This is observed together with the phosphorescence emission, and indicates that triplet states are being harvested in two different ways: one due to reverse intersystem crossing, giving delayed fluorescence (DF), and the other by directly decaying radiatively to the ground state, giving rise to room temperature phosphorescence (RTP). For **4** dispersed in TSPO1 the overall luminescence decreases more than four-fold upon introducing oxygen.

The S_1 and T_1 energies of **4** in zeonex, determined from the onset of the fluorescence and phosphorescence bands, are 3.05 eV and 2.63 eV, respectively. For blends of **4** in **7** and **9**, the S_1 shifts to 2.92 eV and the triplet state shows no shift. In blends of **4** with TSPO1 and PBrS, both the S_1 and T_1 energies shift to 2.82 eV and 2.53 eV, respectively. Therefore, compared with the situation in zeonex, the S_1 energy of **4** has shifted by 0.13 eV in **7** and **9**, and by 0.23 eV in TSPO1 and PBrS. However, the triplet energy exhibits no shift for blends of **4** with **7** and **9**, and shifts by only 0.1 eV in TSPO1 and PBrS. Therefore, the $\Delta E_{S_1-T_1}$ gap decreases from 0.42 eV in zeonex to approx. 0.29 eV in all the other hosts. This explains the turn-on of the delayed fluorescence, and the observation of dual luminescence, TADF and RTP, as shown in Figure 8. However, it is stressed that given the manifold of triplet states below S_1 in the case of **4**, we cannot determine at present from which triplet the S_1 is being thermally activated, only that the significant quenching of



fluorescence on addition of oxygen reveals an emission mechanism which passed through the triplet states.

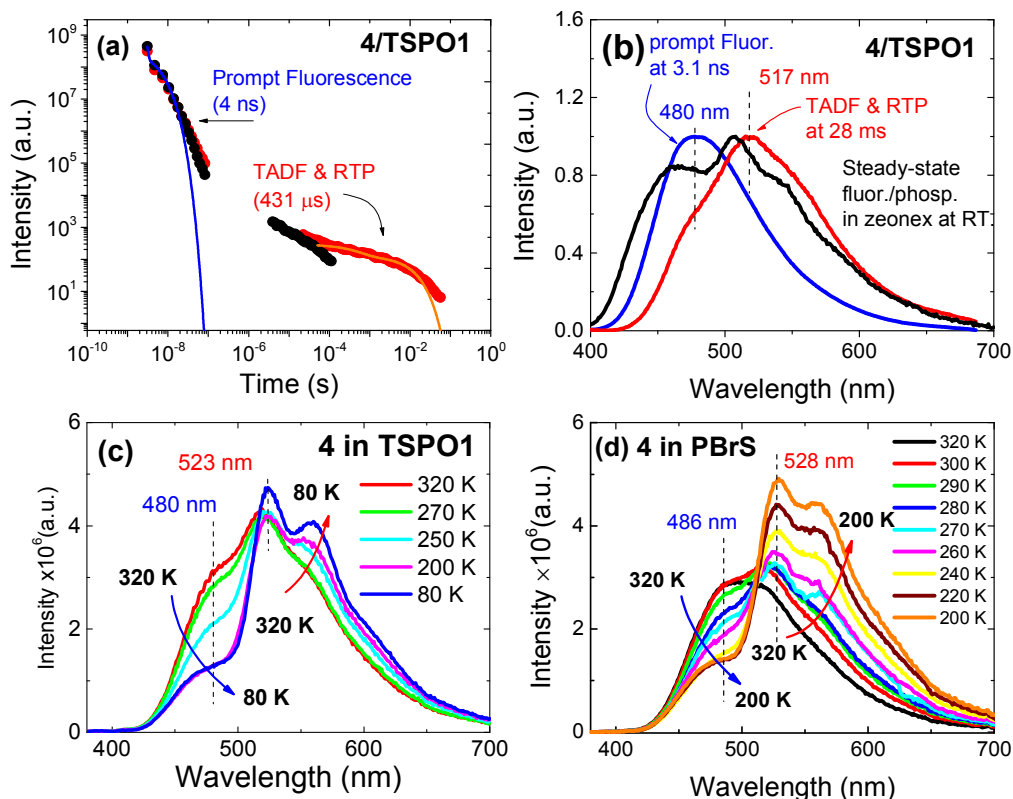


Figure 8-(a) Time resolved luminescence decay of **4** in TSP01, in vacuum (red) and in the presence of oxygen (black), showing the decay of the prompt fluorescence and long-lived TADF and RTP. (b) Time-resolved spectra of **4** in TSP01, showing the spectra of the prompt fluorescence (blue), and long-lived TADF and RTP (red). Both spectra are compared with the steady-state emission of **4** in zeonex. (c,d) Steady-state emission spectra of **4** in TSP01 (c) and **4** in PBrS as a function of temperature (d). The contributions of TADF and RTP are clearly observed in both situations, confirming the presence of dual long-lived luminescence.

The time-resolved luminescence of compound **4** in TSP01 is shown in Fig. 8a. The luminescence decays, obtained in vacuum and in presence of oxygen, show a fast decay component due to prompt fluorescence. This component shows no



variation in the presence of oxygen. A long-lived component due to TADF and RTP is also observed. This is completely quenched by oxygen, confirming that it originated from triplet states. In Fig. 8b, the time-resolved emission spectra obtained in the prompt fluorescence and TADF/RTP regions are compared with the steady-state emission obtained in zeonex. The spectrum at early times is dominated by the fluorescence band, as expected, whereas the delayed component is dominated by TADF and RTP, appearing with a similar emission spectrum to the prompt fluorescence and phosphorescence.

Figs. 8c and 8d show the steady-state emission spectra of **4** dispersed in TSPO1 and PBrS, respectively. In both cases dual emission is observed, with a fluorescence band peaking at 480 nm, which is due to TADF, as confirmed by its temperature dependence, decreasing in intensity at low temperatures, and well resolved phosphorescence peaking at 523 nm in TSPO1, and at 528 nm in PBrS, which, in contrast, increases in intensity with decreasing temperature. This convincingly confirms the dual nature of the luminescence of compound **4** in these hosts. Similar observations were made for blends of compound **2** and are shown in Figs. S9 to S11.

Conclusions

In summary, we have shown that the ISC and rISC mechanisms are enhanced in pure organic molecules by using a simple strategy based on changing the molecular connectivity of carbazole, diphenylamine and fluorene donors to a dibenzothiophene-S,S-dioxide acceptor, from “*linear*” to “*angular*”. The latter substitution of the A unit, breaks effective molecular conjugation and shifts the singlet and triplet states to higher energy. However, more importantly, the “*angular*” substitution gives rise to a small energy gap



between triplet levels below S_1 and to a slower radiative rate, making the S_1 state susceptible to other decay processes such as ISC and IC. In contrast, in the “*linear*” compounds the radiative decay rate of the S_1 state is an order of magnitude faster and in addition, the energy separation between the consecutive triplet states is much larger. The small energy difference between triplet states below S_1 , present in the “*angular*” isomers has profound implications to the ISC and rISC mechanisms. Nonadiabatic vibronic coupling between the T_1 and the upper triplet levels lying below S_1 , creates a fast T_1 - T_n equilibrium, and strong mixing between these states. This enhances coupling between S_1 and T_1 , as described recently.³² In the absence of the strong mixing between the triplet states lying below S_1 , due to the larger energy difference between T_1 and T_2 , for compounds **1** and **3**, the ISC is not efficient, and in combination with the increased S_1 radiative decay rate produces a lower triplet yield and consequently RTP and TADF are not observed, even when **1** and **3** are dispersed in poly(4-bromostyrene), a host where the external heavy-atom effect is used to promote intersystem crossing. Finally, using the host to fine-tune the energy difference between S_1 and T_1 , dual long-lived luminescence, TADF and RTP in **2** and **4** are activated simultaneously in different hosts. This work, therefore, gives guidelines that can be used to design novel molecules showing alternative routes to harvest triplet states and enhance their potential for application in photonic and optoelectronic devices.

EXPERIMENTAL METHODS

Optical Spectroscopy. Absorption and emission spectra were collected using a UV-3600 double beam spectrophotometer (Shimadzu), and a Fluorolog fluorescence spectrometer (Jobin Yvon). Time resolved fluorescence decays were collected using



the picosecond time correlated single photon counting technique (impulse response function, IRF: 21 ps). The vertical polarization excitation source was the second or third harmonic from a picosecond Ti:sapphire laser (Coherent). Emission was collected using a polarizer at magic angle and detected by a double subtractive monochromator, (Acton Research Corporation), coupled to a micro-channel plate photo multiplier tube (Hamamatsu R3809U-50). Signal acquisition was performed using a TCSPC module (Becker & Hickl SPC-630). Temperature dependent measurements were acquired using a model liquid nitrogen cryostat (Janis Research). Phosphorescence, time resolved spectra and decays were recorded using nanosecond gated luminescence and lifetime measurements (from 400 ps to 1 s) using a high energy pulsed Nd:YAG laser emitting at 355 nm (EKSPLA), or a pulsed N₂ laser emitting at 337 nm. Emission was focused onto a spectrograph and detected on a sensitive gated iCCD camera (Stanford Computer Optics) having sub-nanosecond resolution. Solution measurements used concentrations in the 10⁻⁵–10⁻⁴ M range, and samples were degassed using 5 freeze/thaw cycles, when required. Films for optical characterization were prepared in zeonex and other hosts by drop-cast or spin coated in a quartz substrate with emitter/host ratio of (1:20 w/w).

Flash Photolysis. Triplet-formation quantum yields and lifetimes were measured with a flash photolysis setup composed of a LKS.60 ns laser photolysis spectrometer from Applied Photophysics, with a Brilliant Q-Switch Nd:YAG laser from Quantel, using the third harmonics ($\lambda_{\text{ex}} = 355$ nm, laser pulse half-width equal to 6 ns). First-order kinetics was observed for the decay of the lowest triplet state (T-T annihilation was prevented by the low excitation energy and/or low analyte concentration), see Fig. S12 in SI. The triplet lifetimes were measured at absorbance ca. 0.1 in degassed *n*-hexane. The transient spectra were obtained by monitoring the optical



density change at intervals of 10 nm over the 280-600 nm range and averaging at least 5 decays at each wavelength. The Φ_T values were measured by comparing the change of absorbance of β -carotene triplet ($\lambda = 510$ nm) sensitized by energy transfer from a substrate, ΔOD , and from 2-naphthone, $\Delta OD(N)$, used as standard ($\Phi_T(N) = 0.46$)⁴¹ under the same experimental conditions (absorbance at 355 nm and concentration of β -carotene)].⁴² The efficiency of energy transfer from the substrate, P_{ET} , and from 2-naphthone, $P_{ET}(N)$, was also taken into account by measuring the triplet lifetime of the donor in the absence (τ_T^0) and in the presence (τ_T') of β -carotene:

$$P_{ET} = \frac{\tau_T^0 - \tau_T'}{\tau_T^0} \quad (1)$$

Thus, Φ_T values are given by equation (2)

$$\phi_T = \phi_T(N) \frac{\Delta OD}{\Delta OD(N)} \frac{P_{ET}(N)}{P_{ET}} \quad (2)$$

For compounds with T-T absorption at 510 nm, ΔOD in the absence of β -carotene was subtracted in the final value. All measurements were carried out at 20 ± 2 °C; the solutions were deaerated by using the freeze-pump-thaw method through a minimum of 5 cycles.

Computational Details. The geometries of compounds **1-4** were optimized using (U)DFT(PBE0)⁴³⁻⁴⁵ and a 6-31g* basis set as implemented within the Gaussian09 quantum chemistry package without any symmetry constraints.⁴⁶ These static S_0 ground state optimized geometries were confirmed as true minima based on no imaginary frequencies found in frequency calculations. Subsequently, excited state energies and oscillator strengths were calculated using TDDFT(PBE0) within the Tamm-Damcoff approximation.⁴⁷ Spin orbit coupling matrix elements were computed



on the optimized 'gas-phase' ground state geometries using linear response time-dependent density functional theory (LR-TDDFT) using the Tamm–Damcoff approximation and the PBE0 exchange and correlation functional as implemented within the Qchem code.⁴⁸ The lowest six singlet and six triplet states were calculated in every TDDFT calculation. The emission properties were mimicked by optimizing the gas-phase molecules **1-4** in their lowest T_1 states using UKS DFT and the computed emission data are listed in the supporting information. These T_1 optimized geometries were also confirmed as true minima by frequency calculations.

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Author contributions

R.H. performed the majority of the optical measurements, with P.L.S., M.K.E., and E.C. performed photoinduced absorption studies and luminescence spectra in zeonex. J.A. and J.C.L. determined the fluorescence yields in zeonex and the triplet formation yields. P. D. performed cyclovoltammetry experiments. M.N.B.S. investigated the temperature dependence of the luminescence of compound **2** in zeonex. J.W. synthesized molecules **1-4** under the guidance of M.R.B.. M.A.F. performed preliminary calculations and revised the manuscript. The full physical model was devised by J.C.L., T.J.P. and F.B.D., and it was theoretically tested within a quantum chemistry framework by T.N. and T.J.P.. M.R.B. initially designed the regioisomeric emitter structures; F.B.D. conceived the original idea and the optical experiments for investigation, and analysed experimental data. F.B.D. also wrote the manuscript, which was revised by T.J. P., J. W., M.R.B., and J.C.L..

Data Availability



Data supporting this publication is openly available under an 'Open Data Commons Open Database License'. Additional metadata are available at: <http://dx.doi.org/10.17634/153015-3>. Please contact Newcastle Research Data Service at rdm@ncl.ac.uk for access instructions.'

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