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Abstract
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Early Excited State Dynamics of 6-Styryl-Substituted Pyrylium Salts Exhibiting Dual Fluorescence

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A series of 6-styryl-2,4-diphenylpyrylium salts exhibiting dual fluorescence has been investigated by fluorescence up-conversion in conjunction with quantum chemical calculations. The short-wavelength emission is due to an excited state localized on the pyrylium fragment and the long-wavelength emission arises from a charge-transfer state delocalized over the whole molecule. The two fluorescing states do not exhibit a precursor−successor relationship. The rise time of the short-wavelength fluorescence is smaller than 200 fs, and that of the long-wavelength emission depends on the electron-donating property of the styryl group substituent. The rise is almost prompt with the weaker donors but is slower, wavelength and viscosity dependent with the strongest electron-donating group. A model involving a S2/S1 conical intersection is proposed to account for these observations.

Introduction

Pyrylium salts (PS) represent an important class of dyes that has found many applications such as laser and Q-switch dyes,1−7 organic luminophores,8 near-IR absorbing dyes,9 pH-sensors,10 and textile dyes.11 They have also been used in various synthetic procedures.12−15

On the other hand, several PS are characterized by unusual photophysical properties that have been discussed in many reports. In general, 2,4,6-trisubstituted PS with three identical substituents, e.g., 2,4,6-triaryl PS, as well as some blocked PS are considered as two-dimensional chromophoric systems with X and Y absorption bands.16−23 The photophysical and nonlinear properties of PS dimers that tend to be formed in low polarity solvents have been investigated in detail by Markovitsi and co-workers.20,23−25 Kossanyi and co-workers have reported the observation of dual emission from blocked PS and explained it in terms of two excited states with different geometries.26−28 The appearance of dual fluorescence has also been reported with other PS such as thiopyrylium salts and PS with biphenyl groups.19,29 Some authors have also assigned the anomalous fluorescence to a twisted intramolecular charge-transfer (TICT) state.16,26 In all these investigations with systems exhibiting dual emission, the time resolution of the measurements was not high enough to monitor the buildup dynamics of the anomalous fluorescence. Despite this, a precursor−successor relationship between the two fluorescent states was assumed. Finally, it should be noted that the presence of a nonemissive TICT state has also been invoked to account for the viscosity-dependent fluorescence quantum yield of a PS with an electron-donating group.21 This hypothesis is further supported by a recent investigation using ultrafast transient absorption and fluorescence up-conversion.30

In a previous paper,31 the unusual fluorescence properties of 6-styryl-2,4-disubstituted PS in solvents of different polarities have been reported. Their UV−vis absorption spectra are characterized by two well-separated bands. The high energy band with a maximum in the 340−400 nm spectral region is apparently strongly localized on the pyrylium fragment of the molecule and has been assigned to the S0−S2 transition. The low energy band, with a maximum in the 420−530 nm region, depending on the nature of the substituent in the 6-position of the pyrylium fragment, corresponds to the S0−S1 transition. Excitation of the S0−S2 transition gives rise to two distinct fluorescence bands, one that is located between the S0−S2 and the S0−S1 absorption bands and the other that is a mirror image of the S0−S1 absorption band. The analysis of the experimental data indicates that the low energy emission is the ordinary S1−S0 fluorescence associated with the whole conjugated system of the PS. On the other hand, the short-wavelength emission, located at an energy higher than that for the S0−S1 absorption band has been ascribed to a local fluorescence related to the pyrylium fragment of the molecule. Consequently, the high and low energy emission bands have been designated as L and D bands, for localized and delocalized, respectively.31

In nonprotic solvents, the decay of the L fluorescence was found to be monoexponential with a lifetime varying between 1.5 and 2 ns. On the other hand, the decay of the D fluorescence could not be reproduced with less than three exponential functions, with about 90% of the total amplitude associated with

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Figure 1. Structure of the PS investigated.

a 3.5-4 ns lifetime. However, no precursor-successor relationship between both emissions could be evidenced. In fact, no rising component of the D-fluorescence could be detected, the time resolution of the measurements being of the order of 300 ps.

We report here on an investigation of the early fluorescence dynamics of several 6-styryl-2,4-diphenyl PS (see Figure 1) using femtosecond fluorescence up-conversion to elucidate the conversion mechanism between the two emitting states. These PS differ by the electron-donating properties of the substituent on the styryl group. Therefore, the extent of charge-transfer (CT) character of the D transition can be expected to increase from compounds 2-4. For comparison, a PS dye with a methyl group instead of the styryl substituent (compound 1) and exhibiting only the L fluorescence was investigated as well. To get a deeper insight into the electronic character of the lowest electronic states of these dyes, quantum chemistry calculations on compounds 2 and 4 have also been performed. A possible model accounting for the “non-Kasha” fluorescence behavior of these dyes, which is also based on these calculations, will then be proposed.

Experimental Section

Apparatus. The fluorescence up-conversion setup has already been described in detail elsewhere. In brief, the frequency doubled output of a Kerr lens mode-locked Ti:Sapphire laser (Tsunami, Spectra-Physics) was used for excitation of the samples at 400 nm. This excitation wavelength was used for all measurements. The output pulses centered at 800 nm had a duration of about 100 fs and a repetition rate of 82 MHz. The pump intensity on the sample was around 10^14 photons cm^-2 pulse^-1. The polarization of the pump pulses was at a magic angle relative to that of the gate pulses. Experiments were carried out in a 0.4 mm thick rotating cell. The absorbance of the sample solutions was also measured by using a sample cell. Experiments were performed at 20 °C.

Samples. The synthesis of the pyrylium perchlorate salts 1-4 has been described in the literature. These compounds have been purified by repeated recrystallization and the amount of impurity has been estimated to be less than 1%. The solvents, dichloromethane (DCM), ethanol (EtOH) and butanol (BuOH) were of the highest commercially available purity and were used as received. All solutions were purged with argon before measurements. A substantial degradation of the sample solutions was observed after prolonged irradiation. Such a poor photo-stability of many PS dyes in solution is well-known, and therefore fresh samples were used for each measurement. To ensure that this degradation had no effect on the fluorescence up-conversion data, the time profiles obtained by scanning the time delay in both forward and backward directions were compared. Unless specified, all measurements have been performed at 20 ± 2 °C.

Computational Methods. The ground-state singlet gas-phase geometries of 2 and 4 were fully optimized at the density functional level of theory (DFT) using the BP86 and B3LYP functionals and a [3s2p1d] basis set. Electronic vertical excitation energies and oscillator strengths were then computed using time-dependent density functional theory (TDDFT) and configuration interaction over single excitations (CIS). For the TDDFT calculations, both the pure BP86 and hybrid B3LYP functionals were examined in conjunction with the [3s2p1d] basis set; for the CIS calculations, the INDO/S2 Hamiltonian was used. In addition, the gas-phase geometries of the first and second singlet excited states of 4 were optimized at the TDB3LYP[3s2p1d] level. Solvatochromic effects were computed at the VEM4.2 level of theory based on the CIS/INDO/S2 results. TDDFT calculations were carried out using Turbomole version 5.8.0 and ZINDO-MN version 1.2.

Results

Early Fluorescence Dynamics. The absorption and fluorescence spectra of the dyes 1-4 have been discussed in detail in ref 31, and the band maxima are listed in Table 1. As an example, the spectra of 4 are shown in Figure 2. Both the absorption and fluorescence spectra of the other dyes in the L band region are very similar. On the other hand, the D band shifts to shorter wavelength by going from 4 to 2. For the latter compound, the D band overlaps substantially with the L band, whereas it is absent with 1, where the styryl group is replaced by a methyl group.

The areas of both L and D absorption bands of 4 in DCM are independent of temperature between -5 and +25 °C. The same behavior was observed with the D fluorescence band of the same dye. On the other hand, a 5% decrease of the area of its L emission band was found by going from -5 to +25 °C. The same small decrease was measured with 1. Although the L fluorescence maximum remains constant in the temperature range investigated, the D fluorescence band exhibits a hypsochromic shift of about 3 cm^-1 K^-1 with increasing temperature.

Figure 3 shows time profiles of the early fluorescence of 1 and 2 in DCM. The rise of all these profiles is essentially

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**Table 1: L and D Absorption and Emission Maxima (in nm, Excitation at 360 nm) of 1-4 in Various Solvents**

<table>
<thead>
<tr>
<th>Solvent</th>
<th>1</th>
<th>2</th>
<th>3</th>
<th>4</th>
</tr>
</thead>
<tbody>
<tr>
<td>DCM</td>
<td>L</td>
<td>L</td>
<td>D</td>
<td>L</td>
</tr>
<tr>
<td></td>
<td>λ_{max}</td>
<td>λ_{em}</td>
<td>λ_{max}</td>
<td>λ_{em}</td>
</tr>
<tr>
<td></td>
<td>383</td>
<td>383</td>
<td>470</td>
<td>383</td>
</tr>
<tr>
<td>EtOH</td>
<td>373</td>
<td>373</td>
<td>450</td>
<td>373</td>
</tr>
<tr>
<td>BuOH</td>
<td>373</td>
<td>373</td>
<td>555</td>
<td>440</td>
</tr>
<tr>
<td></td>
<td>373</td>
<td>373</td>
<td>430</td>
<td>373</td>
</tr>
</tbody>
</table>

---

**Figure 2.** Absorption (black) and fluorescence spectra (gray, excitation at 360 nm) of 4 in DCM.

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**Figure 3.** Time profiles of the early fluorescence behavior of 1-4 in DCM.
prompt; i.e., its time constant is shorter or equal to the fwhm of the instrument response function. After this rise, the L fluorescence at 430 nm of both 1 and 2 exhibits a weak decaying component with a time constant of the order of 1.5–2 ps (see Table 2). At the other wavelengths investigated, the fluorescence intensity remains constant within the time window of the measurement (10 ps).

In EtOH, the early dynamics of the L fluorescence of 1 and 2 is qualitatively the same as in DCM. On the other hand, the rise of the D fluorescence of 2 was found to be biphasic, with a prompt and a ~1.5 ps component. The amplitude of this slower component apparently increases with increasing wavelength. However, it should be noted that at 500 nm, both L and D emissions contribute to the signal. All the parameters obtained upon analysis of these time profiles are listed in Tables 2 and 3.

The early L fluorescence dynamics of 3 in both DCM and EtOH is essentially the same as that of 1 and 2. The rise of its D-fluorescence is also biphasic with a 1.5–2.3 ps component in both solvents. Because of the small amplitude of this slower component, the uncertainty on its time constant is relatively large.

As for the other dyes, the rise of the L fluorescence of 4 is essentially prompt. On the other hand, the early dynamics of the D fluorescence in both solvents differs substantially from that of 2 and 3. Indeed, it is not prompt and it exhibits a marked wavelength dependence. In DCM, the rise time increases from 180 to 360 fs by going from 600 to 650 nm. In EtOH, the rise of the D fluorescence is substantially slower and increases with wavelength as well, as shown in Figure 4A. To test whether the slower rise in EtOH is related to the higher viscosity of this solvent compared with DCM (η(DCM) = 0.44 cP, η(EtOH) = 1.2 cP at 20°C), the early fluorescence dynamics of 4 was also investigated in a more viscous solvent, namely BuOH (η(BuOH) = 3.3 cP at 20°C). As shown in Figure 4B, the rise of the fluorescence intensity is clearly slower than that in the other solvents and the wavelength dependence is still present. This rise could not be very well reproduced with a monoexponential function, especially above 650 nm and thus a biexponential function, especially above 650 nm and thus a biexponential function had to be used. The resulting best-fit parameters are listed in Table 3. The average rise times are also listed to enable comparison with the other solvents. Interestingly, a plot of the rise time measured as the longest wavelength as a function of solvent viscosity is almost perfectly linear with a slope of 0.3 ps/cP. Although the number of investigated solvents is small, this correlation is probably not coincidental.

Quantum Chemistry Calculations. To understand in more detail the electronic character of the various excited singlet states of these compounds, TDDFT calculations were carried out for gas-phase molecular geometries optimized using the BP86 and B3LYP functionals. In addition, CIS calculations at the INDO/S2 level of theory were also performed. Both levels of theory agree that the frontier orbitals of the styrylpyrylium system,
**Figure 5.** Frontier molecular orbitals (HOMO−1, HOMO, LUMO, and LUMO+1) for 4 computed at the B3LYP/[3s2p1d] level of theory. Orbital contours are drawn at the 0.05 a.u. level.

**TABLE 4: Parameters Obtained from the Fit of a Biexponential Function to the Time Profiles of the Early Fluorescence of 4 in BuOH upon 400 nm Excitation**

<table>
<thead>
<tr>
<th>λ (nm)</th>
<th>band</th>
<th>$\lambda$</th>
<th>$\tau_1$ (ps)</th>
<th>$\tau_2$ (ps)</th>
<th>$\tau^o$ (ps)</th>
</tr>
</thead>
<tbody>
<tr>
<td>450</td>
<td>L</td>
<td>−1</td>
<td>0.19</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>630</td>
<td>D</td>
<td>−0.66</td>
<td>0.39</td>
<td>−0.34</td>
<td>1.0</td>
</tr>
<tr>
<td>650</td>
<td>D</td>
<td>−0.60</td>
<td>0.42</td>
<td>−0.40</td>
<td>1.43</td>
</tr>
<tr>
<td>670</td>
<td>D</td>
<td>−0.60</td>
<td>0.46</td>
<td>−0.40</td>
<td>1.9</td>
</tr>
<tr>
<td>690</td>
<td>D</td>
<td>−0.55</td>
<td>0.47</td>
<td>−0.45</td>
<td>2.1</td>
</tr>
</tbody>
</table>

* Only the components with a time constant shorter than 10 ps are listed.

**TABLE 5: Excitation Energies, $E$ (eV), Wavelengths, $\lambda$ (nm), and Oscillator Strengths, $f$ (a.u.), for Vertical Excitations in 2 and 4 at Various Levels of Theory**

<table>
<thead>
<tr>
<th>transition</th>
<th>CIS/INDO/S2</th>
<th>TDBP86</th>
<th>TDB3LYP</th>
</tr>
</thead>
<tbody>
<tr>
<td>$S_0 \rightarrow S_2$</td>
<td>3.02 410 [401]$^a$</td>
<td>1.11 2.35 527 0.42 2.70 460 0.68</td>
<td>3.02 410 [401]$^a$</td>
</tr>
<tr>
<td>$S_0 \rightarrow S_3$</td>
<td>3.80 326 [316]</td>
<td>0.98 2.49 497 0.02 3.14 395 0.01</td>
<td>3.80 326 [316]</td>
</tr>
<tr>
<td>$S_0 \rightarrow S_4$</td>
<td>4.15 299 [291]</td>
<td>0.02 2.65 468 0.01 3.21 386 0.26</td>
<td>4.15 299 [291]</td>
</tr>
<tr>
<td>$S_0 \rightarrow S_1$</td>
<td>2.75 451 [432]</td>
<td>1.33 2.06 601 0.38 2.45 506 0.73</td>
<td>2.75 451 [432]</td>
</tr>
<tr>
<td>$S_0 \rightarrow S_3$</td>
<td>3.77 329 [316]</td>
<td>0.60 2.66 466 0.37 3.15 393 0.31</td>
<td>3.77 329 [316]</td>
</tr>
<tr>
<td>$S_0 \rightarrow S_4$</td>
<td>4.00 310 [301]</td>
<td>0.37 2.74 453 0.01 3.32 373 0.56</td>
<td>4.00 310 [301]</td>
</tr>
</tbody>
</table>

* Excitation wavelengths in brackets in this column refer to VEM4.2 predictions in alcohol solution.

**Figure 6.** Optimized geometries for the $S_0$ and $S_1$ states of 4 at the TDB3LYP/[3s2p1d] level of theory.

but the predicted energetics are substantially to the red and to the blue of experiment, respectively. These results are quite consistent with the known tendencies of (i) TDDFT to underestimate the energy separation between ground states and CT excited states (particularly for pure density functionals such as BP86) and (ii) CIS/INDO/S2 to overestimate the transition energies associated with CT excited states and delocalized π excitations.48 However, the CIS/INDO/S2 results are useful insofar as they permit an analysis of solvatochromic effects in alcohol at the VEM4.2 level. In general, the effects are predicted to be relatively small: a 10–20 nm blue shift is predicted for the first three excitations. If these solvatochromic shifts are added to the gas-phase B3LYP predictions, the agreement with experiment remains good.

According to these calculations, both $S_0$–$S_2$ and $S_0$–$S_3$ transitions contribute to the broad absorption band between 300 and 400 nm. This explains why the L fluorescence band is not really its mirror image. Indeed, the L fluorescence band is much narrower (fwhm ~3000 vs ~6350 cm$^{-1}$), and the pronounced shoulder around 345 nm has no equivalent in the fluorescence spectrum.

When the geometries of the first two excited states of 4 are optimized, the lower energy state twists about the styryl moiety and minimizes to a classic TICT structure (Figure 6). Vertical emission from this state is predicted to occur at approximately 1200 nm at the TDB3LYP level, which is substantially to the red of experiment. However, this large red shift is consistent with the known tendency mentioned above of TDDFT to underestimate the energy of CT excited states. Because, with the twisted geometry, CT is effectively complete, it is not surprising that this inaccuracy arises even for the hybrid B3LYP functional. The $S_2$ state, on the other hand, remains planar after relaxation. The major difference between $S_0$ and $S_2$ equilibrium geometries is the twist angle of the phenyl group in the 4 position, which changes from 8° in $S_0$ to 1° in $S_2$. Vertical emission from this state is predicted to occur at about 430 nm, which is in good agreement with experiment.

**Discussion**

Molecules exhibiting dual fluorescence are relatively scarce, in agreement with Kasha’s rule. Most observations of dual emission are due to molecules experiencing a chemical process, such as electron or proton transfer, or a substantial conformational change on the excited-state surface. It should be noted that in these cases, the excited state responsible for the low energy fluorescence cannot be directly populated upon optical excitation. These cases cannot really be considered as exceptions of Kasha’s rule as the fluorescence always occurs from the lowest singlet excited state. The situation is different for the PS dyes investigated here, because their absorption spectra exhibit both L and D bands and therefore the low energy fluorescence can be generated upon direct excitation in the D
band. Therefore, L and D emissions can be considered as $S_2 \rightarrow S_0$ and $S_1 \rightarrow S_0$ emissions, respectively. Zn-tetraphenylporphine (ZnTPP) is most probably the best known molecule, which exhibits fluorescence from both $S_2$ and $S_1$ states. Time-resolved measurements have shown that these emissions have a precursor–successor relationship, indicating the population of the $S_1$ state upon internal conversion from $S_2$.

The striking feature of the styrylpyrylium dyes 2–4 compared to ZnTPP is the absence of precursor–successor relationship between the $S_2$ (L) and $S_1$ (D) emission profiles. Indeed, the rise time of the D fluorescence of 2–4 is much faster than the decay time of the L fluorescence.

Before proposing a model that could account for this observation, we will discuss the origin of the slower rising or decaying components, $\tau_2$, observed in both L and D emission in DCM and EtOH. According to Tables 2 and 3, $\tau_2$ amounts to about 2 ps independently of the PS dye, the solvent and the emission wavelength. Its amplitude, $A_2$, is wavelength dependent: when different from zero, it is positive on the blue side of the L fluorescence band, indicating a decay and negative at longer wavelength (see 4 in DCM). Moreover, its magnitude tends to increase with wavelength (see 2 in EtOH). These two observations suggest that this component is not related to population dynamics. If it were the case, $A_2$ should always be positive in the L band and negative in the D band, as expected for a transfer of population from the L to the D states. Moreover, the observation of this component with compound 1 clearly indicates a different origin. The wavelength dependence of $A_2$ rather suggests a temporal evolution of the shape of the fluorescence spectrum. Considering the value of $\tau_2$, two processes can be envisaged: solvent relaxation and vibrational cooling. According to dynamic Stokes shift measurements performed with coumarin 153, the solvation dynamics of DCM is biphasic with an ultrafast inertial component (100 fs and 50% relative amplitude) and a slower diffusive component of 1 ps. For EtOH, three time constants have been reported: 390 fs (25%), 5 ps (20%), and 30 ps. These values do not agree very well with those of $\tau_2$ found here. Therefore, dynamic Stokes shift seems not to be the main origin of this component.

Vibrational cooling is known to lead to a narrowing of the fluorescence spectrum. This process appears as a decaying component on the edges of the fluorescence band and as a rise in the central region. Vibrational cooling takes place on different time scales, going from less than a picosecond to a few tens of picoseconds, depending apparently on the amount of vibrational excess energy. The small time constant measured here is in agreement with a small excess excitation energy. The solvent dependence of this process is still not totally understood, although the slower cooling component has been reported to depend on the thermal diffusivity of the environment. A joint contribution of both solvation and vibrational cooling dynamics to this component cannot be excluded.

The absence of a D band with 1 and the dependence of the D band maximum on the electron-donating properties of the styryl substituent indicate a CT character of this transition. This is fully supported by the calculations, which can very well reproduce the change of the absorption maximum of the D band by going from 2 to 4. TDB3LYP and CIS/INDO/S2 both agree that the $S_0 \rightarrow S_1$ transition for 2 is blue shifted by about 40 nm relative to that for 4, whereas the other two excitations are less sensitive to this substitution (as expected, because the $S_0 \rightarrow S_2$ and $S_0 \rightarrow S_3$ transitions are substantially more localized on the pyrylium fragment that is common to both 2 and 4).

Moreover, triphenylpyrylium salts are well-known as electron-transfer photosensitizers, and are characterized by a relatively low reduction potential. Kossanyi and co-workers have reported the observation of CT absorption and fluorescence with donor–acceptor complexes composed of aromatic hydrocarbons as electron donors and of 1 as electron acceptor. The shape of these CT bands is very similar to those of the D bands measured here with 2–4. Moreover, local excitation of 1 was found to be followed by both local and CT emissions as observed here for 2–4. Therefore, the D band can be unambiguously assigned to a CT transition. As the relative intensities of L and D bands of 2–4 are independent of the dye concentration, the formation of intermolecular donor–acceptor complexes can be safely excluded. Moreover, the absence of a temperature effect on the absorption spectrum of 4 also rules out the existence of an equilibrium between several conformations in the ground state.

The fact that both L and D emissions exhibit an almost prompt rise and a distinct decay in the nanosecond time scale could be explained with an arrangement of potential-energy surfaces as shown in Figure 7. For the sake of simplicity, harmonic potentials have been assumed, but of course, their actual shape may differ substantially depending on the nature of the coordinate $Q$. Excitation in the L band prepares the excited state population away from equilibrium. Upon relaxation, a fraction of this population remains on the L potential, whereas the other crosses to the D potential. After thermalization, both populations are irreversibly “trapped” at the bottom of their potential and thus the two relaxed states are decoupled. In a multidimensional picture, the intersection region shown in Figure 7 corresponds to a conical intersection. This picture is partially supported by TDDFT geometry optimization of the first two excited states of 4. However, according to these calculations, the D state remains below the L state even after relaxation of the latter’s geometry, and D emission at this geometry is predicted to occur at 510 nm. This result contradicts the above conical intersection hypothesis. However, the CT nature of $S_1$ causes its potential-energy surface to be predicted to be anomalously low at the TDDFT level. It is thus entirely possible that with more accurate energetics, a conical intersection between the $S_1(D)$ and $S_2(L)$ states might occur sufficiently near the ground-state geometry such that relaxation along the formally $S_2$ surface would result.
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in a minimum that, at its own geometry, corresponds to a formally S1 state from which L fluorescence can occur. Unfortunately, the levels of theory that would be required to offer more quantitative support than TDDFT for this phenomenon are not yet computationally practical for polyaromatic systems of this size.

As the transition from the L to D state can be considered as an intramolecular CT process, the coordinate Q should involve both solvent and intramolecular modes as it is generally the case for electron-transfer processes.68 The TDDFT calculations point out the twisting motion of the styryl group as an important coordinate. The occurrence of such a large amplitude motion can explain the viscosity dependence of the D fluorescence rise time measured with 4. The wavelength-dependent rise of the D fluorescence found with this dye could be explained by a conical intersection located far from the D equilibrium state. Therefore, long-wavelength D emission appears only after relaxation of the D state population. On the other hand, the location of the conical intersection close to the equilibrium geometry of the L state is supported by the prompt rise of the L emission. It is, however, not absolutely clear whether the substantial structural change predicted for the CT state can be completed within a picosecond or less. The actual twist angle of the styryl group might be not as pronounced as that calculated. Here again it appears that a more definitive answer calls for higher level of theory. This is, however, a difficult problem even for smaller molecules such as the well-known 4-(dimethylamino)benzonitrile.69

Solvent modes could also contribute to the viscosity dependence of the D fluorescence rise measured with 4. The average solvation times of DCM, EtOH and BuOH amount to 0.56, 16 and 63 ps, respectively, and are thus much larger than the fluorescence rise time. However, solvation dynamics is well-known to be multiphasic and even viscous solvents have an apparent contribution to a nonradiative deactivation pathway with a small temperature dependence.

Both $\Delta E_{LD}$ and $\Delta Q_{LD}$ are the largest with 4, and therefore a slower and wavelength-dependent rise time can be observed. On the other hand, the close to prompt D fluorescence rise of PS 2 and 3 can be explained by a smaller $\Delta Q_{LD}$ and hence by a conical intersection located closer to the equilibrium position of the D state.

The absence of a temperature dependence of the D fluorescence band measured with 4 shows that the D state is not thermally accessible from the L state and reciprocally, in agreement with this model. The small temperature effect observed with the L fluorescence of both 4 and 1 indicates that it is not related to the D state. The slightly larger L fluorescence quantum yield at lower temperature is most probably connected to a nonradiative deactivation pathway with a small temperature dependence.

Concluding Remarks

We have reported the first investigation of the anomalous fluorescence of PS dyes using ultrafast spectroscopy. The data show clearly the absence of a precursor—successor relationship between the two emitting states. These styrylpyrylium dyes thus exhibit exceptional photophysical properties. In this case, the violation of Kasha’s rule is due to the fact that both thermally equilibrated states are totally uncoupled. The presence of a conical intersection seems to be the most plausible explanation for the quasi-prompt rise of the fluorescence from both states. This hypothesis is partially supported by TDDFT calculations. However, higher level calculations, which are presently not computationally practical, would be required to obtain a deeper insight into the unusual photophysics of these dyes. Moreover, it would be interesting to perform similar measurements with other PS, which also exhibit dual fluorescence, to find out whether the unusual behavior observed here is a general feature of PS.

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