

# Twisting strategy applied to *N,N*-diorganoquinacridones leads to organic chromophores exhibiting efficient solid-state fluorescence

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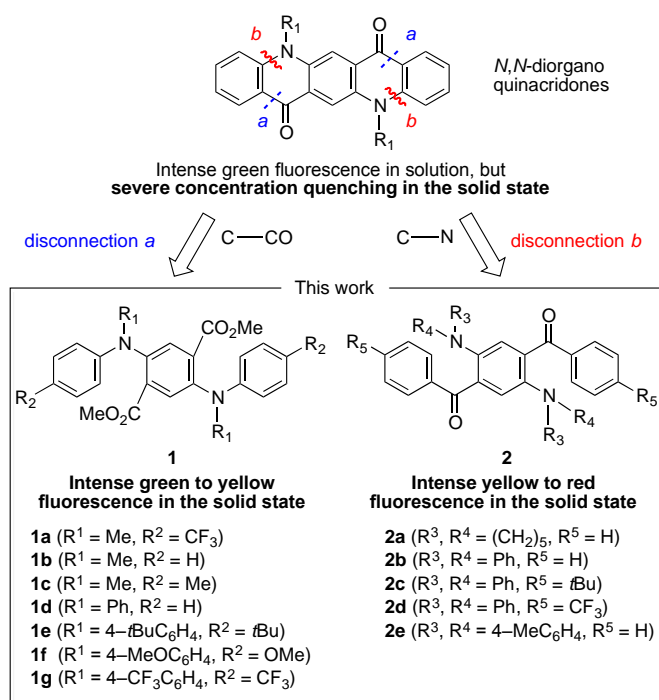
## ABSTRACT

A new molecular design of organic emitters exhibiting efficient solid-state fluorescence, which involves planarity breaking of *N,N*-diorganoquinacridones, is presented. The new design principle led to the development of dimethyl 2,5-diaminoterephthalates and 2,5-diamino-1,4-diaroylbenzenes, which emitted green to yellow and yellow to red light with high-to-excellent quantum yields, respectively. In addition, the photoluminescence properties of the diaroylbenzenes were dependent on the morphology and reversibly variable by thermal and solvent vapor stimuli.

Organic chromophores that exhibit fluorescence in the visible region with high efficiency in the solid state have attracted increasing attention in the field of functional materials. This is because the highly efficient solid-state emissions of organic fluorophores are essential for the advances needed to realize optoelectronic devices such as organic light-emitting diodes (OLEDs),<sup>1</sup> light-emitting field-effect transistors,<sup>2</sup> semiconductor lasers,<sup>3</sup> and fluorescent solid sensors.<sup>4</sup> However, most organic fluorophores are non-luminescent or only faintly emissive in the solid state due to concentration quenching induced by molecular aggregation, which is an inherent characteristic of organic solids. Hence, the molecular design and development of organic fluorophores exhibiting highly efficient solid-state emissions have met limited success,<sup>5</sup> but such fluorophores are of great importance with respect to both practical applications in optoelectronic devices and fundamental studies in solid-state molecular engineering. The reported molecular designs that render fluorophores efficiently luminescent in the solid state involves 1) the incorporation of a spiro framework; 2) the introduction of bulky substituents; 3) vicinal disubstitution of aryl/aryl or alkenyl/aryl groups on an aromatic ring or alkenyl moiety, and 4) the installation of electron-donating and -withdrawing groups at the molecular ends and/or middles.<sup>5a</sup> Recently, self-assembly-induced and exciplex-mediated emission enhancement of organic gels is also demonstrated.<sup>5b-5c</sup>

In this communication, we present a new guiding principle for the design of organic fluorophores that emit visible light in the solid state with high efficiency; this principle involves double disconnection of *N,N*-diorganoquinacridones, which show

intense green fluorescence in solution but undergo severe luminescence quenching in the solid state (Scheme 1). The



**Scheme 1.** Molecular design of highly emissive organic solids on the basis of planarity breaking of *N,N*-diorganoquinacridones.

approach allows the development of dimethyl 2,5-bis(diorgano-amino)terephthalates **1** and 1,4-diaroyl-2,5-bis(diorgano-amino)-benzenes **2**, which exhibit efficient solid-state fluorescence in the green to yellow and yellow to red spectral regions, respectively. The high efficiency of the solid-state emissions is attributed to the non-planar molecular frameworks and the intramolecular charge-transfer (CT) character of the  $S_1$  states. Moreover, it is notable that **2** are fluorescent although benzophenone and its derivatives are generally non-fluorescent due to rapid intersystem crossing (ISC) from the  $S_1$  state to the  $T_1$  state.

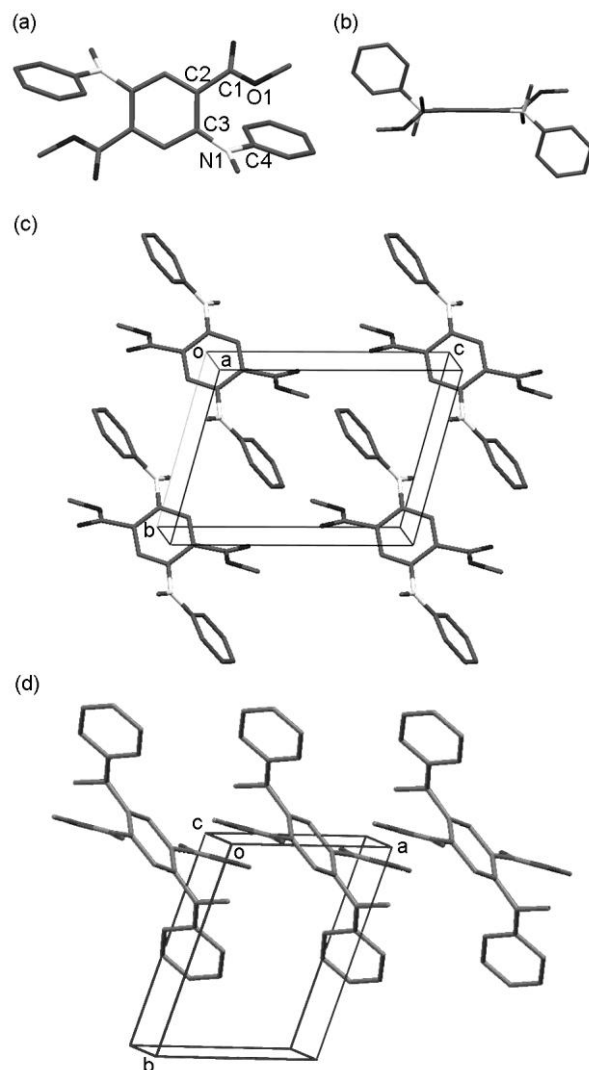
Organic chromophores that consist of planar and rigid  $\pi$ -conjugated systems generally exhibit luminescence with high efficiency *in solution*, where the chromophores are sufficiently isolated for them not to interact electronically with one another. The planarity of the molecular framework enhances  $\pi$ -conjugation, and the rigidity of the skeleton is beneficial in the restriction of both the stretching and twisting motions of each bond, which can result in radiationless deactivation of the excited states. However, planar polycyclic chromophores normally show a strong tendency to aggregate *in the solid state*, with a  $\pi$ - $\pi$  stacked geometry that causes severe luminescence quenching. Typical examples are *N,N*-diorganoquinacridones, which exhibit intense green-emissions in dilute solution, but are non- or weakly-emissive in the solid state and even in concentrated solution.<sup>6</sup> For example, the photoluminescence quantum yields of *N,N*-dibutyl-2,9-difluoroquinacridone were reportedly 0.97 in THF ( $1.0 \times 10^{-6}$  M) and 0.0019 in the microcrystals.<sup>6a</sup> They are therefore used as *dopants* in OLED devices.<sup>7</sup>

We have recently demonstrated that 1,4-bis(alkenyl)-2,5-dipiperidinobenzenes are highly emissive fluorophores in the solid state.<sup>8</sup> Through an investigation of their structural characteristics and photophysical properties, we concluded that breaking the high planarity of *N,N*-diorganoquinacridones by disconnection of the bonds linked to the outer benzene rings, while retaining the electronic structure consisting of a 1,4-diamino-2,5-dicarbonyl functionality on the central benzene core, would provide new chromophores exhibiting highly efficient visible-light fluorescence in the solid state if the two sets of *ortho*-linkages of the diorganoamino and carbonyl groups could induce a twisted conformation suitable for suppression of dense stacking in the ground state and a large Stokes shift leading to inhibition of Förster-type energy transfer from the excited state.

Based on this idea, there are two options for the disconnection (Scheme 1); one is cleavage of the C-CO bonds (indicated by dotted blue lines *a*), and the other involves fission of the C-N bonds (indicated by wavy red lines *b*). We initially designed dimethyl 2,5-bis(arylamino)terephthalates **1** as disconnection *a*, and prepared them in good to high yields from dimethyl 2,5-dibromoterephthalate and anilines by Pd-catalyzed amination or commercially available dimethyl 2,5-dioxocyclohexane-1,4-dicarboxylate via three steps (see the Supplementary data for details).<sup>9</sup>

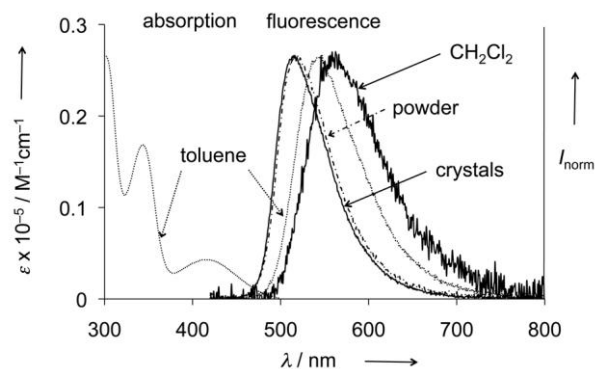
Single crystals of **1b** suitable for X-ray diffraction analysis were obtained by recrystallization from dichloromethane/hexane solution.<sup>9</sup> As shown in Figure 1, the phenyl and methyl groups of the amino moiety as well as the methoxy groups are positioned largely out of the central benzene plane ( $C2-C3-N1-C4 = 50.28^\circ$ ;  $C3-C2-C1-O1 = 40.45^\circ$ ) and the distances between the centroids of the central benzene rings in two adjacent molecules in the crystal lattice are over 6 Å; there is no intermolecular aromatic interaction like  $\pi$ - $\pi$  stacking to cause emission quenching. The molecular and crystal structures of **1d** also show similar distorted molecular conformations and the absence of  $\pi$ - $\pi$  stacking.<sup>9</sup>

Absorption and fluorescence spectra of **1d** in toluene are shown in Figure 2 as representative examples.<sup>9</sup> The absorption

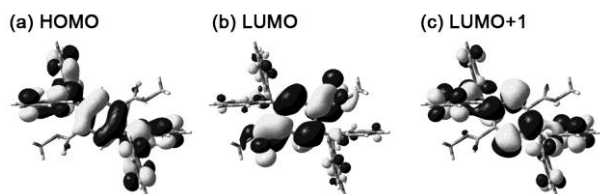


**Figure 1.** Molecular and crystal structures of **1b**: space group P-1 (triclinic),  $a = 6.0995(13)$ ,  $b = 8.5112(19)$ ,  $c = 11.134(2)$ ,  $\alpha = 95.713(4)^\circ$ ,  $\beta = 104.354(4)^\circ$ ,  $\gamma = 107.081(4)^\circ$ . Hydrogens are omitted for clarity. (a) top view, (b) side view, (c) packing view from *a* axis, and (d) packing view from *c* axis.

maxima appeared at 343 nm ( $\epsilon = 16800$ ) and 415 nm ( $\epsilon = 4300$ ), and green fluorescence was observed with an emission maximum at 547 nm and a quantum yield of 0.40. Thus, there was no overlap of the absorption and fluorescence spectra, suggesting that photo-excitation induced the large structural change in **1d**. When a solvent was changed from toluene to dichloromethane, the fluorescence spectra red-shifted ( $\lambda_{em} = 564$  nm,  $\Phi_f = 0.11$ ) as shown in Figure 2.<sup>9</sup>



**Figure 2.** Absorption and fluorescence spectra of **1d**.



**Figure 3.** HOMO (a), LUMO (b), and LUMO+1 (c) diagrams of **1d**.

The HOMO, LUMO, and LUMO+1 of **1d** obtained by density functional theory (DFT) calculations at the B3LYP/6-31G(d) level are depicted in Figure 3.<sup>10</sup> The HOMO is developed over the Ph<sub>2</sub>N–benzene–NPh<sub>2</sub> framework, while the MeO<sub>2</sub>C–benzene–CO<sub>2</sub>Me unit contributes to the LUMO. Time-dependent DFT calculations reveal that the lowest energy transition is governed mainly by the HOMO–LUMO transition.<sup>11</sup> Thus, the absorption band at 415 nm can be ascribed to an intramolecular CT transition, which is consistent with the large Stokes shift and the solvatochromism.

The photoluminescence properties of **1** in the crystalline and powder states are summarized in Table 1, together with those in toluene (the spectra are shown in Supplementary data). Upon photo-irradiation, solid **1** exhibited intense solid-state fluorescence ranging from green to yellow, with good to high quantum yields ( $\Phi_f = 0.32$ – $0.76$ ). The fluorescence spectra of solid **1d** are shown in Figure 2.<sup>9</sup> Some photographs of the irradiated samples are shown in Figure 4. There is no big difference between the emission maxima in the crystalline and powder states for each compound. In all cases,  $\Phi_f$  values in solution were much lower than those in the solid state; **1** were found to exhibit aggregation-induced emission (AIE).<sup>12</sup> The serious non-radiative decay in solution is presumably attributable to the intramolecular free motion of the amino and methoxycarbonyl moieties. Diethyl bis(4-trifluoromethylphenylamino)terephthalate and diethyl bis[3,5-bis(trifluoromethyl)phenylamino]terephthalate are reported to display unique thermal stimuli-induced switching of solid-state fluorescence.<sup>13</sup> Since the quantum yields of the ArNH-substituted terephthalates in the solid state ranged from 0.066 to 0.410, it is apparent that the presence of the second carbonaceous substituents on the nitrogens of **1** is essential for attaining high solid-state quantum yields in molecular design based on 2,5-diaminoterephthalate frameworks.<sup>14</sup>

**Table 1.** Photoluminescence data of **1**.

Entry	<b>1</b>	$\lambda_{em}$ [nm] <sup>a</sup> ( $\Phi_f$ ) <sup>b</sup>		
		microcrystals <sup>c</sup>	powder <sup>d</sup>	solution <sup>e</sup>
1	<b>1a</b>	523 (0.47)	527 (0.32)	514 (0.15)
2	<b>1b</b>	539 (0.64)	544 (0.60)	544 (0.03)
3	<b>1c</b>	553 (0.47)	549 (0.43)	– <sup>f</sup>
4	<b>1d</b>	519 (0.44)	516 (0.71)	547 (0.40)
5	<b>1e</b>	572 (0.67)	565 (0.76)	556 (0.18)
6	<b>1f</b>	571 (0.60)	568 (0.63)	– <sup>f</sup>
7	<b>1g</b>	504 (0.76)	508 (0.72)	499 (0.54)

<sup>a</sup> Emission maxima upon photo-irradiation ( $\lambda_{ex} = 400$  nm).

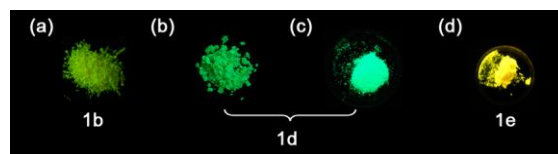
<sup>b</sup> Absolute quantum yield determined with a calibrated integrating sphere system.

<sup>c</sup> Prepared by recrystallization from dichloromethane/hexane.

<sup>d</sup> Prepared by evaporation (600 hPa, 40 °C) of CH<sub>2</sub>Cl<sub>2</sub> solution.

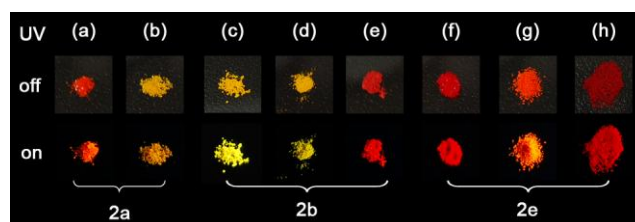
<sup>e</sup> 1 × 10<sup>−5</sup> M in toluene.

<sup>f</sup> No fluorescence was observed.



**Figure 4.** Photoluminescence images of **1**: (a) crystals of **1b**, (b) crystals of **1d**, (c) powder of **1d**, and (d) powder of **1e**.

Diaminodiaroylbenzenes **2**, designed as disconnection *b* (Scheme 1), were synthesized as colored solids from 2,5-dibromoterephthaloyl dichloride via three steps (Figure 5, (a)–(h), the upper photographs).<sup>9</sup> During purification and isolation, we observed that the colors of the isolated solids **2b** and **2e** were very sensitive to the preparative history of each solid sample. Recrystallization of **2b** from THF/MeOH solution and slow evaporation (220 hPa, 40 °C) of a THF solution gave yellow microcrystals and powder (Figures 5c and 5d), respectively; quick evaporation (60 hPa, 40 °C) afforded a red powder (Figure 5e). DSC analysis of **2b** in red powder showed an exothermic phase-transition peak at 127 °C.<sup>9</sup> When the red powder was heated at 130 °C, it changed color to yellow. Recrystallization of **2e** from THF/MeOH solution provided red microcrystals (Figure 5f), and quick evaporation of a THF solution left a mixture of orange and red powders. Interestingly, all of the red powder in the mixture turned orange (Figure 5g) upon heating the mixture at 100 °C.<sup>9</sup> In turn, the resulting orange powder gradually and completely turned red (Figure 5h) when the powder was exposed to dichloromethane vapor for 24 h.<sup>15</sup> The color-changing process was found to be reversible. These observations indicate that the photophysical properties of **2b** and **2e** in the solid state are very sensitive to subtle differences in morphology.



**Figure 5.** Optical and fluorescence images of **2**: (a) crystals of **2a**, (b) powder of **2a**, (c) crystals of **2b**, (d) yellow powder of **2b**, (e) red powder of **2b**, (f) crystals of **2e**, (g) orange powder of **2e**, and (h) red powder of **2e** under natural light (the upper photograph of each entry) and a UV lamp (the lower photograph of each entry).

With this morphology-sensitive behavior of **2** in mind, we measured the photoluminescence of the crystals and powders. The results are summarized in Table 2, along with the results obtained from measurements in toluene. All of the prepared solid samples emitted brilliantly visible light ranging from yellow to deep red (Figure 5, (a)–(h), the lower photographs). Comparison of the solid-state emissions with those in solution confirms that **2** are also AIE-active chromophores. There was no overlap of the absorption and emission spectra of **2**; in other words, large Stokes shifts were observed, as illustrated by the example of **2b** in Figure 6. Changing a solvent from toluene to dichloromethane induced a red-shift of the fluorescence spectra as observed with **1** (see the Supplementary data for details).<sup>9</sup> X-ray single crystal analysis of **2a** revealed that the piperidyl and benzoyl groups were largely out of the central benzene plane.<sup>9</sup> DFT calculations indicate that the photoluminescence is also governed by intramolecular CT transitions (Figure 7).<sup>10</sup> The emissions appeared in a longer wavelength region than those of **1**; this can be ascribed to the enhanced intramolecular CT character of the transitions as a result of the stronger electron-

**Table 2.** Photoluminescence data of **2**.<sup>a</sup>

Entry	<b>2</b>	color and shape	$\lambda_{em}$ [nm] <sup>b</sup>	$\Phi_f$ <sup>c</sup>
1	<b>2a</b>	orange crystals <sup>d</sup>	608	0.38
2		orange powder <sup>e</sup>	588	0.14
3		(in toluene) <sup>f</sup>	– <sup>g</sup>	– <sup>g</sup>
4	<b>2b</b>	yellow crystals <sup>d</sup>	562	0.61
5		yellow powder <sup>h</sup>	565	0.35
6		red powder <sup>e</sup>	637	0.22
7		(in toluene) <sup>f</sup>	602	0.20
8	<b>2c</b>	orange crystals <sup>d</sup>	614	0.60
9		orange powder <sup>e</sup>	610	0.44
10		(in toluene) <sup>f</sup>	603	0.23
11	<b>2d</b>	red crystals <sup>d</sup>	640	0.33
12		red powder <sup>e</sup>	640	0.18
13		(in toluene) <sup>f</sup>	633	0.14
14	<b>2e</b>	red crystals <sup>d</sup>	650	0.47
15		orange powder <sup>i</sup>	610	0.30
16		red powder <sup>l</sup>	657	0.20
17		(in toluene) <sup>f</sup>	623	0.16

<sup>a</sup> Irradiation was effected at 400 nm for **2a** and 450 nm for **2b–2e**.

<sup>b</sup> Emission maxima.

<sup>c</sup> Absolute quantum yield determined with a calibrated integrating sphere system.

<sup>d</sup> Prepared by recrystallization from THF/ MeOH solution.

<sup>e</sup> Prepared by evaporation (60 hPa, 40 °C) of THF solution.

<sup>f</sup>  $2 \times 10^{-5}$  M in toluene.

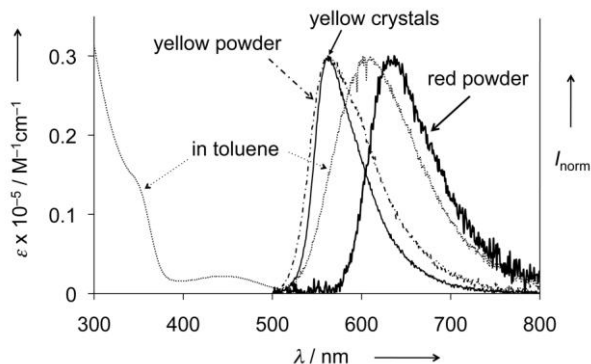
<sup>g</sup> No fluorescence was observed.

<sup>h</sup> Prepared by slow evaporation (220 hPa, 40 °C) of THF solution.

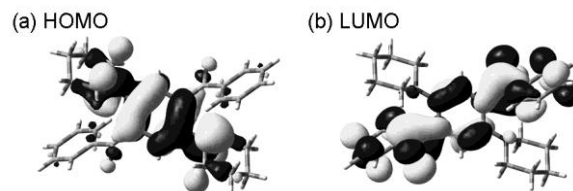
<sup>i</sup> Prepared by evaporation (60 hPa, 40 °C) of THF solution followed by heating at 100 °C.

<sup>j</sup> Prepared by evaporation (60 hPa, 40 °C) of THF solution followed by heating at 100 °C, and then exposure to a vapor of dichloromethane for 24 h.

withdrawing effects of aroyl moieties compared with those of methoxycarbonyl groups. Since red-emissive organic fluorophores in the neat solid state generally have a strong tendency to cause aggregation, which results in severe concentration quenching, it is a formidable challenge to attain quantum yields exceeding 0.3 with regard to red emissions from neat organic solids.<sup>16</sup> It is therefore noteworthy that **2d** and **2e** in crystals exhibits red emission at 640 nm and 650 nm with excellent quantum yields of 0.33 and 0.47, respectively.



**Figure 6.** Absorption spectra of **2b** in toluene and fluorescence spectra of **2b** in toluene, as crystals, and as a powder.



**Figure 7.** HOMO (a) and LUMO (b) diagrams of **2a**.

It should be noted that the luminescence from **2** originates in radiative decay of the  $S_1$  state, i.e., *fluorescence*; this is confirmed by lifetime measurements;  $\tau(\mathbf{2a}, \text{crystal}) = 15.2$  ns and 5.9 ns;  $\tau(\mathbf{2a}, \text{powder}) = 8.1$  ns;  $\tau(\mathbf{2b}, \text{crystal}) = 22.5$  ns;  $\tau(\mathbf{2b}, \text{powder}) = 10.3$  ns and 4.7 ns.<sup>17</sup> Generally, diaryl ketones in which the orbital configuration of the  $S_1$  state is assigned to  $n-\pi^*$  transitions of the ketone carbonyl group are almost non-fluorescent ( $\Phi_f = \sim 0.01$  to 0.0001) because ISC from the  $S_1$  state to the  $T_1$  state is much faster than the relatively slow radiative decay.<sup>18</sup> For example, benzophenone exhibits no fluorescence in solution at room temperature and shows phosphorescence at low temperatures in a frozen solvent due to highly efficient ISC ( $\Phi_{ISC} = 1.0$ ).<sup>19</sup> Room-temperature phosphorescence from crystals of benzophenone and its derivatives has also been demonstrated very recently.<sup>20</sup> Compared with these features of phosphorescence emissions from diaryl ketones, the observed efficient fluorescence from **2**, which possesses two diaryl ketone moieties, clearly implies that the intramolecular CT transitions induced by the amino groups substituted at the *ortho*-positions mainly govern the electronic structure of the  $S_1$  state in favor of the  $n-\pi^*$  transitions of the ketone carbonyls. This is consistent with nearly no distribution of the HOMO on the carbonyl oxygen and significant contribution of the amino groups to the HOMO (Figure 7).

In summary, we have demonstrated that dimethyl 2,5-bis(diorganoamino)terephthalates and 1,4-diaroyl-2,5-bis(diorganoamino)benzenes in the solid state exhibit intense green to yellow and yellow to red fluorescence, respectively, with good to high quantum yields.<sup>21</sup> In particular, it is remarkable that an excellent quantum yield of solid-state red fluorescence is attained with 1,4-dibenzoyl-2,5-bis(4-methylphenyl)benzene ( $\Phi_f = 0.47$  with an emission maximum of 650 nm). The molecular design of these new solid fluorophores originates in the idea of planarity breaking of *N,N*-diorganoquinacridones, and the success of the design is now prompting us to apply this molecular design concept to other types of planar dopant emitters, such as coumarins, in order to extend the validity of the design principle. Furthermore, we have found that 1,4-diaroyl-2,5-bis(diorganoamino)benzenes constitute a new class of polymorphic solids that exhibit morphology-sensitive luminescence. These findings make 2,5-bis(diorganoamino)-1,4-diaroylbenzenes potentially attractive as organic emitting materials, not only for OLEDs, but also for sensing and switching devices that utilize solid-state luminescence as an output.<sup>22</sup> Further studies on these polymorphic structures and their relationship with photophysical properties are in progress.

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