Spiroconjugated Tetraaminospirenes as Donors in Color-Tunable Charge-Transfer Emitters with Donor-Acceptor Structure

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Dedicated to Professor Rolf Gleiter on the occasion of his 85th birthday

Abstract: Charge-transfer emitters are attractive due to their color tunability and potentially high photoluminescence quantum yields (PLQYs). We herein present tetraaminospirenes as donor moieties, which, in combination with a variety of acceptors, furnished 12 charge-transfer emitters with a range of emission colors and PLQYs of up to 99%. The spatial separation of their frontier molecular orbitals was obtained through careful structural design, and two DA structures were confirmed by X-ray crystallography. A range of photophysical measurements supported by DFT calculations shed light on the optoelectronic properties of this new family of spiro-NN-donor-acceptor dyes.

Introduction

Small molecules with donor-acceptor (DA) character are attractive as active materials for optoelectronic applications. They are used in organic light-emitting diodes (OLEDs), dye-sensitized solar cells, as fluorescence-based sensors, UV absorbers and nonlinear optical materials, to name a few.[1–9] In DA-compounds, photoexcitation can lead to an intramolecular charge transfer (CT) of an electron from the donor to the acceptor part of the molecule.[10] Introducing spiro units can be beneficial in organic DA-type emitters. Due to their rigid three-dimensional structure π–π-stacking and excimer formation can be effectively suppressed, since they often form non-crystalline, amorphous structures in the solid-state.[11,12] In OLED emitters, such interactions can lead to significant redshifts in photoluminescence and electroluminescence, making them less practicable for commercial use.[13] Another interesting feature in spiro-compounds is the possibility of spiroconjugation,[14,15] defined as the homoconjugative through-space interaction between two π-systems arranged perpendicular to each other and connected by a common tetrahedral atom.[16–19] Its extent depends on the orbital overlap between the two π-systems and hence on the substitution pattern around the spiro center.[20] Several examples of spiro-conjugated CT emitters have been reported to date, where donor and acceptor are separated by a spiro center.[14,21–29] We recently reported on spiro-donor-acceptor dyes, with 1,3-indandione and derivatives thereof as acceptors and aromatic methylated diamines, amino thiolis, aromatic thiolis, or aromatic alcohols as donor units.[10,30] Although intramolecular CT was observed in the absorbance spectra, these dyes did not show fluorescence due to fast vibrational energy deactivation of the excited states. We herein report the first use of heterospiropyrans 1 and 2 as donors for charge-transfer dyes with DA-character (Figure 1). Heterosporane 2 was first reported in 1968 by Quast and Schmitt,[32] and naphthyl-based 1 in 1986 by Gleiter and Uschmann.[33] Through photo-electron spectroscopic investigations, Gleiter et al. attested strong spiro-interactions in 2 due to significant orbital overlap between the two halves of the molecule.[13,34] We combined these donors 1 and 2 with five different acceptors and used three modes of connection between donor and acceptor (Figure 2).
compounds were confirmed by X-ray crystallography. The resulting 12 in total spiro-NN-DA compounds showed strong photoluminescence both in solution and in thin films. We investigated in particular the influence of the connection mode between donor and acceptor on the photophysical properties and found the meta-connection to result in the highest photoluminescence quantum yields of up to 99%.

Results and Discussion

Syntheses

The synthesis of the mono-halogenated derivatives of heterospiranes 1 and 2, namely 8a and 8b, 9 and 12, are shown in Scheme 1. The heterospirane frameworks are typically synthesized through condensation of a 2-chlorobenzimidazolium salt and methylaed o-phenylenediamine or 1,8-naphthyldiamine, thereby constructing the spiro center.[32,33]

To obtain mono-halogenated derivatives of these heterospires as functional building blocks, which can be employed in Pd-catalyzed cross-coupling reactions, we developed synthetic routes to the halogenated chlorobenzimidazolium salts 3a and 3b (Scheme 1A). Their syntheses commenced from 4-bromo-1-fluoro-2-nitrobenzene (4a) and 1-chloro-2-fluoro-3-nitrobenzene (4b) by nucleophilic aromatic substitution of the fluoride with methylamine, followed by reduction of the nitro to an amino group with tin chloride. This furnished 5a and 5b in excellent yields of 90% and 98% over two steps, respectively. The following condensation with 1,1-carbonyldimazole afforded dihydrobenzimidazolones 6a and 6b. Reaction with phosphoryl chloride led to imidoyl chlorides 7a and 7b, which were methylated with trimethyloxonium tetrafluoroborate, providing the halogenated chlorobenzimidazolium salts 3a and 3b. With 77%, the overall yield for 3a over five steps from 4a is high, while the lower overall 19% yield for 3b from 4b was mostly due to the poor yield in the condensation reaction of 5b with 1,1-carbonyldimazole. The molecular structures of intermediate 4-bromo-N-methyl-2-nitroaniline, 7a, as well as 3a in the solid state were characterized by X-ray crystallography (see structures in Scheme 1A).

For the synthesis of “meta”-brominated heterospiranes 8a and 9, the benzimidazolium salt 3a was reacted with naphthyl-based diamin 10 and phenylene diamine 11, respectively, using triethylamine as base in acetonitrile (Scheme 1B). For N\textsubscript{2},N\textsubscript{2}dimethyl phenylenediamine (10), we developed an improved four-step synthetic route from 1,8-diaminonaphthalene, shown in the Supporting Information, and 11 was synthesized according to the literature.[35] Due to the air and thermal sensitivity of diamines 10 and 11 this reaction had to be performed with degassed solvents at room temperature. Furthermore, the correct ratio of the solvents acetonitrile/triethylamine was critical to the yields of 8a and 9. Suzuki-Miyaura-coupling of 8a with (2-chlorophenyl)boronic acid allowed introducing the 1,2-phenylene spacer unit in 12. “Ortho”-chlorinated 8b was obtained in a similar manner by condensation of benzimidazolium salt 3b with N\textsubscript{2},N\textsubscript{2}dimeth-

The type of connection between the donor and acceptor can significantly influence the optoelectronic properties of the CT-dyes. For naphthyl-based donor 1, all three connection modes were employed, in which the acceptors are either connected “ortho” to the spiro unit (1-o-acceptor), “meta” (1-m-acceptor), or via a 1,2-phenylene spacer (1-ms-acceptor). With donor 2, only this third connection mode via the phenylene spacer was employed in DA compounds 2-ms-acceptor. As acceptors, we chose dibenzo[b]thiophenedioxide (DBTO), diphenyl sulfone (DPS), thianthrenetetroxide (TTO), benzonitrile (BN), and diphenyloxadiazole (DPOD) (Figure 2).

Since mono-halogenated derivatives of heterospiranes 1 and 2 were unknown prior to this study, we developed synthetic strategies to these functional building blocks, which we employed in Suzuki-Miyaura cross-coupling reactions to furnish the resulting DA compounds. The structures of two DA...
ylnaphthalene-1,8-diamine (10) in a good yield of 68%. The molecular structure of 8a in the solid-state was confirmed by X-ray diffraction analysis (see Scheme 1B).

To obtain the borylated acceptors A-Bpin with A = DBTO, DPS and TTO required for the cross-coupling reactions with the heterospiranes, we performed Miyaura-borylations of the respective bromides. This is shown in Scheme 2A for DBTO-Br and TTO-Br (for their synthesis, see Supporting Information) or has been reported in the literature for DPS-Bpin.

Benzonitrile-based acceptor 13 with attached 1,2-phenylene spacer was afforded through Suzuki-Miyaura-coupling of 1,2-dibromobenzene with benzonitrile-boronic ester BN-Bpin followed by Miyaura-borylation of the remaining aryl bromide (Scheme 2B). For DPOD-based acceptor 14, the reaction of N'-benzoyl-4-bromobenzohydrazide (15) with POCl₃ afforded the corresponding oxadiazole, which was subjected to Miyaura borylation conditions to furnish DPOD-Bpin. Suzuki-Miyaura-coupling of 1,2-dibromobenzene followed by Miyaura-borylation of the remaining aryl bromide furnished DPOD-based 14 with an incorporated 1,2-phenylene spacer in good overall yield.

With the halogenated heterospiranes and borylated acceptors in hand, we next synthesized 12 DA-compounds through Suzuki-Miyaura-cross-coupling reactions, as shown in Scheme 3.

Structural properties

The layering of dichloromethane solutions with n-pentane provided single crystals suitable for X-ray diffraction analysis of 1-ms-DPS and 1-o-TTO. The molecular structures are shown in Figure 3 and demonstrate a twist between donor and acceptor unit. 1-ms-DPS possesses dihedral angles of 57.3° between the heterospirane donor and the 1,2-phenylene spacer and of 53.8° between the latter and the diphenylsulfone acceptor. In 1-o-TTO, the torsional angle between donor and acceptor amounts to 46.1°. Interesting are also the distances between donor and acceptor units for the intramolecular CT process. In 1-o-TTO the distance between the S-atom of the acceptor and the spirocenter of the donor amounts to 8.3 Å, while this distance decreases to 6.8 Å for 1-ms-DPS.

To incorporate the 1,2-phenylene spacer in the meta-position of the heterospirane donors, we employed two different strategies. For acceptors BN and DPOD, the spacer was first attached to the acceptor unit BN or DPOD (see Scheme 2) and then cross-coupled with bromo-heterospirane 8a to furnish 1-ms-BN and 1-ms-DPOD. The same protocol was followed using benzo-based heterospirane 9 for the benzonitrile acceptor and afforded DA-compound 2-ms-BN. In the second strategy, we used the 1,2-phenylene-extended heterospirane 12 in Suzuki-Miyaura-couplings with acceptors A-Bpin, resulting in DA-compounds 1-ms-DBTO, 1-ms-DPS, and 1-ms-TTO.
To obtain structural information on all spiro-NN-DA compounds, all 12 were optimized in the gas phase using the PBEh-3c composite scheme.\textsuperscript{[38]} As for 1-\textit{ms}-DPS and 1-\textit{o}-TTO the other compounds also showed a twist between donor and acceptor (or to the \textit{ortho}-phenylene bridges). The torsional angles between donor and acceptor $\theta_{D-A}$ for the \textit{m}- and \textit{o}-connection mode or between donor/acceptor and the spacer ($\theta_{D-S}$ and $\theta_{A-S}$) for the \textit{ms}-connection are listed in Table 1. The largest torsion angles with values of ca. 50°–60° were found for the DA compounds with a \textit{meta}-spacer in between donor and acceptor and for those with an \textit{ortho}-connection to the donor. The direct \textit{meta}-connection resulted in smaller values of 37°–39° due to decreased steric repulsion. Large torsion angles between donor and acceptor help in localizing the electron densities of the HOMO and LUMO states on donor and acceptor moieties, respectively, but can also diminish the oscillator strengths of the charge-transfer absorptions.

**Optoelectronic and photophysical properties**

DFT calculations provided information on the optoelectronic properties of the 12 spiro-NN-DA compounds. Selected calculated optoelectronic properties are listed in Table 2. The calculated HOMO energies are fairly similar for all compounds (between $-5.13$ eV and $-4.25$ eV), since the electron densities of the HOMOs are located on the spiro-NN-donor moieties (see Figure 4 for the HOMO and LUMO plots of three selected DA compounds, HOMO and LUMO plots of all other compounds can be found in the Supporting Information). The calculated LUMO energies depend on the type of acceptor chosen, and...
the LUMO electron densities are located on the acceptors in each case. The calculated HOMO-LUMO-energy differences ($E_{gap}$) are related to the (emission) colors of the compounds and increase for the same donor-acceptor pair from the meta-spacer over the ortho to the meta-connection. Of interest are further

The photoluminescence quantum yields (PLQY) in toluene solution were high for most spiro-NN-DA compounds (Table 3). In all cases, performing the measurement in degassed and oxygen-free toluene led to a significant increase in PLQY ($\text{PLQY}_{\text{film}}$) with values of 90 % or higher for seven of the 12 compounds. This indicates that the excited states participating in the photochemistry of the compounds are deactivated by oxygen. Usually, these are triplet excitons quenched by triplet $\text{O}_2$, but it could also be singlet excitons quenched by singlet $\text{O}_2$. The meta-connected DA compounds 1-m-DBTO, 1-m-DPS,

$$\Delta E_{ST} = E_{ST} - E_{S} - E_{T}$$

the splitting $\Delta E_{ST}$ between the lowest energy triplet ($T_1$) and singlet state ($S_0$) as well as the oscillator strength $f$. A large value $f$ is beneficial to ensure efficient radiative decay from $S_0$ to the ground state, while a small energy gap $\Delta E_{ST} < 0.1 \text{ eV}$ can enable thermal up-conversion from the triplet to the singlet excited state by reverse intersystem-crossing (RISC).

For the spiro-NN-DA compounds with donor 1 investigated herein, these values are proportionate to each other. With the meta-spacer in between donor and acceptor, the smallest $\Delta E_{ST}$ values between 8–118 meV are predicted with at the same time low oscillator strengths $f = 0.0005$–0.0050. The ortho-connection leads to intermediate values of $\Delta E_{ST} = 20$–280 meV and $f = 0.0045$–0.0105, while for the direct meta-connection large singlet-triplet splittings between 200 and 490 meV were calculated with high oscillator strengths of 0.0997–0.2918.

As can be seen from the solution fluorescence spectra in Figure 5, the emission colors of the spiro-NN-DA compounds ranged from blue to red with emission maxima between 445 and 643 nm (see also Table 3). The highest energy blue emissions were found for 1-o-DPS and 1-m-DPS with the diphenylsulfone acceptor. The benzotriazole, diphenyloxazolone, and dibenzothiophenedioxide acceptors led to green emissions with the exception of 1-m-DBTO (yellow). The thianthrenetetrmine acceptor caused a bathochromic shift to yellow (1-m-DBTO), orange (1-o-DBTO), or red emissions (1-m-TTO). As DFT calculations had predicted, for the same acceptor, the band gap of the DA compounds decreased from the meta- over the ortho-connection to the meta-spacer.

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| Compound | $\theta_{D}$ | $\theta_{A}$ | $\theta_{DA}$ | $\theta_{meta}$ |
|----------|-------------|-------------|-------------|--------------|
| 1-m-DBTO | 54.3°       | 51.3°       | 1-o-DBTO    | 60.9°        |
| 1-m-DPS  | 54.1°       | 51.7°       | 1-o-DPS     | 61.7°        |
| 1-m-DPS  | 56.3°       | 49.6°       | 1-o-TTO     | 58.4°        |
| 1-m-DBTO | 55.7°       | 56.0°       | 1-m-DBTO    | 38.9°        |
| 1-m-TTO  | 54.2°       | 57.3°       | 1-m-DPS     | 37.8°        |
| 2-m-BN   | 53.1°       | 51.3°       | 1-m-TTO     | 37.3°        |

| Compound | $E_{HOMO-calc}$ | $E_{LUMO-calc}$ | $E_{gap}$ | $\Delta E_{ST}$ | $f$ |
|----------|----------------|----------------|-----------|-----------------|-----|
| 1-m-DBTO | 2.26           | 2.82           | 0.58      | 0.0005          | 0.0005 |
| 1-m-DPS  | 1.60           | 3.41           | 1.81      | 0.0050          | 0.0050 |
| 1-m-TTO  | 1.83           | 3.16           | 2.38      | 0.0045          | 0.0045 |
| 2-m-BN   | 1.69           | 3.24           | 2.45      | 0.0060          | 0.0060 |
| 1-o-DBTO | 1.95           | 3.02           | 2.56      | 0.0105          | 0.0105 |
| 1-o-DPS  | 2.38           | 2.63           | 0.25      | 0.0005          | 0.0005 |
| 1-o-TTO  | 2.25           | 2.72           | 0.47      | 0.0005          | 0.0005 |
| 1-m-DBTO | 2.82           | 3.16           | 0.34      | 0.0005          | 0.0005 |
| 1-m-DPS  | 2.82           | 3.16           | 0.34      | 0.0005          | 0.0005 |
| 1-m-TTO  | 2.82           | 3.16           | 0.34      | 0.0005          | 0.0005 |
| 2-m-BN   | 2.82           | 3.16           | 0.34      | 0.0005          | 0.0005 |

$\Delta E_{ST} = E_{ST} - E_{S} - E_{T}$

$\Delta E_{ST}$ values between 8–118 meV are predicted with the same time low oscillator strengths $f = 0.0005$–0.0050. The ortho-connection leads to intermediate values of $\Delta E_{ST} = 20$–280 meV and $f = 0.0045$–0.0105, while for the direct meta-connection large singlet-triplet splittings between 200 and 490 meV were calculated with high oscillator strengths of 0.0997–0.2918.
and 1-\textit{m}-TTO featured the highest PLQYs, likely related to the higher oscillator strengths of the first singlet excitations and probably also the emissions.

We next measured PLQY in thin films with 10 wt% spiro-NN-DA in \textit{m}-CBP (3,3:\textprime-di(9H-carbazol-9-yl)-1,1:\textprime-biphenyl) matrix. Interestingly, the quantum yields were lower than in solution for most compounds. The spiro-NN-DA compounds with the shortest wavelength absorptions featured the highest PLQYs in the thin-films (54–66\%), namely 1-\textit{ms}-BN and 2-\textit{ms}-BN, 1-\textit{ms}-DPOD, and all three compounds with the diphenylsulfone acceptor 1-\textit{ms}-DPS, 1-\textit{o}-DPS, and 1-\textit{m}-DPS. This may be related to an energy alignment with the \textit{m}-CPB matrix for these molecules, which has to efficiently transfer its excited state energy to the emitter molecule. The fact that the PLQY of the spiro-NN-DA emitters decreased in the matrix may indicate that the CT state is reached through an intermolecular electron-transfer process rather than an intramolecular one.

We next measured the transient photoluminescent decay for all 12 spiro-NN-DA emitters to investigate whether an indication for efficient RISC was found, which would result in a delayed component. In all cases, only first-order decay was observed, corresponding to a prompt fluorescence component with lifetimes $\tau = 2–14$ ns. To rationalize this observation, we experimentally determined the $S_1-T_1$ energy splittings $\Delta E_{ST}$ from the onsets of the solution fluorescence (at rt) and phosphorescence spectra (at 77 K, see Supporting Information) (Table 3). With values $\Delta E_{ST}$ between 280 and 770 meV, these energy differences are significantly higher than the calculated ones and too large to be efficiently overcome by thermally activated RISC. Such a deviation between calculated and experimental values has been reported in other instances.\[41\]

Hence, the fact that the PLQY increased upon deoxygenation of the solutions was probably due to singlet exciton quenching by singlet O$_2$, which was thereby inhibited. Still, their high PLQYs in solution and tunable emission colors make the spiro-NN-DA compounds herein attractive as fluorescent emitters.

**Conclusions**

In summary, we developed efficient synthetic routes to two monofunctionalized tetraaminospirenes as spiroconjugated donor units. Palladium-catalyzed cross-coupling to a variety of acceptor moieties in three different modes of connection between donor and acceptor afforded 12 spiro-NN-DA compounds. Their structures were investigated by X-ray crystallography and/or DFT calculations, confirming spatial separation of the donor and acceptor part and as a consequence of the HOMO and LUMO. Photophysical measurements provided photoluminescence quantum yields of up to 99\%, depending on the connection mode between the donor and acceptor unit, and a range of emission colors. Time-resolved emission spectroscopy showed that these compounds featured a prompt fluorescence with lifetimes up to 14 ns. The spiroconjugated tetraamine donors presented herein enable access to structurally and optoelectronically interesting DA-type emitters.
Experimental Section

Experimental Details can be found in the Supporting Information. Deposition Numbers 1949447 (for 1-ms-DPS), 1949585 (for 1-o-TTO), 1912248 (for 3a), 1962741 (for 7a), 2041007 (for 14a), 2046064 (for 51, structure see Supporting Information), and 2043983 (for 53, structure see Supporting Information) contains the supplementary crystallographic data for this paper. These data are provided free of charge by the joint Cambridge Crystallographic Data Centre and Fachinformationszentrum Karlsruhe Access Structures service.

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Conflict of Interest

The authors declare no conflict of interest.

Data Availability Statement

The data that support the findings of this study are available from the corresponding author upon reasonable request.

Keywords: amines · charge transfer · donor-acceptor systems · fluorescence · spiro compounds

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