

Exploring the Photocyclization Pathways of Styrylthiophenes in the Synthesis of Thiahelicenes: When the Theory and Experiment Meet

Bianca C. Baciú, José Antonio Vergés, and Albert Guijarro*



Cite This: *J. Org. Chem.* 2021, 86, 5668–5679



Read Online

ACCESS |



Metrics & More

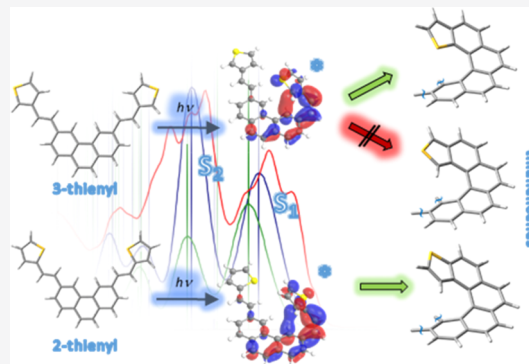


Article Recommendations



Supporting Information

ABSTRACT: The introduction of thiophene rings to the helical structure of carbohelicenes has electronic effects that may be used advantageously in organic electronics. The performance of these devices is highly dependent on the sulfur atom topology, so a precise knowledge of the synthetic routes that may afford isomeric structures is necessary. We have studied the photocyclization pathway of both 2- and 3-styrylthiophenes on their way to thiahelicenes by experiment and theory. To begin with, the synthesis of stereochemically well-defined 2- and 3-styrylthiophenes allowed us to register first, and simulate later, the UV–vis electronic spectra of these precursors. This information gave us access through time-dependent density functional theory calculations to the very nature of the excited states involved in the photocyclization step and from there to the regio- and stereochemical outcome of the reaction. For the widely known case of a 2-styrylthiophene derivative, the expected naphtho[2,1-*b*]thiophene type of ring fusion was predicted and experimentally observed by synthesis. On the contrary, 3-styrylthiophene derivatives have been seldom used in synthetic photocyclizations. Among the two possible structural outcomes, only the naphtho[1,2-*b*]thiophene type of ring fusion was found to be mechanistically sound, and this was actually the only compound observed by synthesis.



INTRODUCTION

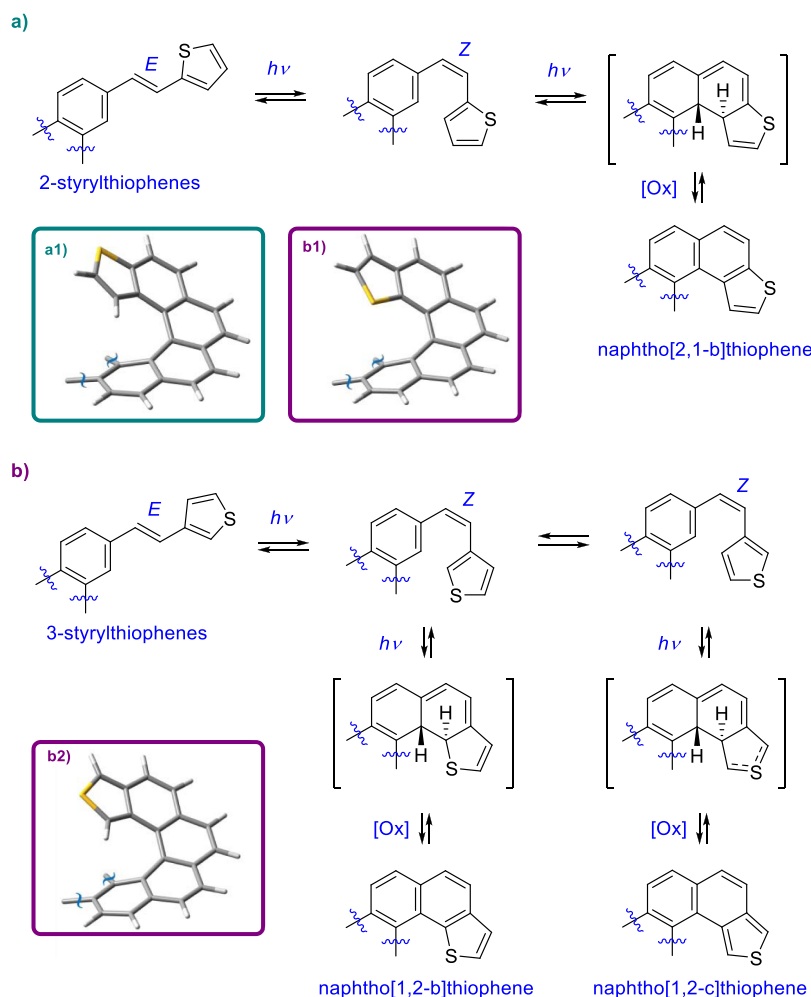
Helicenes display a robust, fully conjugated helical architecture that makes them prototypes of chiral carbon nanostructures, with increasing applications in organic electronics.¹ With the inclusion of thiophene rings into the helical structure, two noticeable effects occur. First, a stable doped state with improved conducting properties arises;^{2,3} second, an enhanced binding energy with metallic electrodes results.^{3–5} Both are the consequence of an enriched π -electron density induced by the sulfur atom. These two aspects are highly dependent on the sulfur atom position and bond topology generated therefrom within the thiahelicene,^{6,7} so synthetic routes must be explicit in that respect. In their seminal studies, Wynberg and coworkers used the photocyclization of 2-styrylthiophene derivatives as a straightforward way to access to thiahelicenes (Scheme 1a).⁸ Developed half a century ago, it remains a prevalent, still captivating method to achieve the type of ring fusion needed to construct helicenes starting from one of the simplest conceivable precursors, so a large number of thiophene-containing helicene structures have been obtained this way since then.⁹ The general accepted mechanism does not differ much from the classical Mallory reaction of stilbenes¹⁰ and involves a fast *Z*–*E* isomerization of the styrylthiophene precursor,¹¹ followed by a key photocyclization step, namely, photochemical electrocyclic reaction, to afford 9*a*,9*b*-dihydronaphthothiophenes as transient intermediates,¹² which in turn dehydrogenate in the reaction media by

means of an oxidizer to afford the aromatic naphthothiophene moiety (Scheme 1a). It became already apparent at that early stage that 3-styrylthiophenes were far less prolific precursors (Scheme 1b) to the point that there are barely no examples of this photocyclization strategy in the synthesis of thiahelicenes up to until very recently. Although highly under-represented, it is a valuable alternative for the construction of unusual thiophene-terminated helicenes. Focusing on the central features of this photocyclization and particularly on its backbone scaffold (Scheme 1b), it has been reported to afford not only naphtho[1,2-*b*]thiophenes¹³ but also its isomeric naphtho[1,2-*c*] counterpart,¹⁴ which may leave doubts about the actual natural outcome of this reaction. This is a critical yet obscure point that gives rise to different topologies in the sulfur atom positioning, and with it, to the entire electronic structure of the corresponding thiahelicene. This is an important matter for many reasons, but in particular, if the objective of the synthesis is to build conductive yet robust nanocontacts between a helicene and external metallic electrodes. It is

Received: January 19, 2021

Published: March 26, 2021



Scheme 1. Reaction Pathways for (a) 2- and (b) 3-Styrylthiophenes Based on the Widely Accepted Mechanism of Photocyclization^a

^aInset (a1) framed in green shows the 3D view of a thiophene-terminated helicenic structure originated from reactions in (a), while (b1) and (b2) framed in magenta would originate from reactions in (b).

known that quantum interference¹⁵ may lead to constructive or destructive interference, enhancing or suppressing conductance depending on the sulfur atom positioning. Within this scenario, we decided to study in depth this photocyclization in the framework of our studies pursuing different thiophene-terminated helicenes with special regard to the topology originated from the sulfur atom positioning in the key photochemical step. In line with those aims, this work is a combined experimental-theoretical study structured as follows: (a) first, we synthesized suitable 2- and 3-styrylthiophenes as stereochemically homogeneous precursors of thiahelicenes, and then, (b) we studied both experimentally and theoretically their absorption process (excitation) and subsequent key photocyclization step. Finally, (c) we isolated the corresponding thiahelicenes as reaction products and characterized them unequivocally to establish the proper reaction pathway. We decided to work with thiophene doubly terminated helicenes (or more specifically with dithiahelicenes **1** and **2**; Scheme 3) throughout all this mechanistic study for practical reasons, in line with our developing project on molecular solenoids. Their applications as single-molecule devices in organic electronics are currently under study.

RESULTS AND DISCUSSION

Synthesis of (*E,E*)-3,6-bis(2-(Thiophen-2-yl)vinyl)phenanthrene (7**) and (*E,E*)-3,6-bis(2-(Thiophen-3-yl)vinyl)phenanthrene (**8**) as Stereochemically Well-Defined Photochemical Precursors.** Stereochemically well-defined 2- and 3-styrylthiophene precursors were required to trace the following key photochemical step of the synthesis. In the simplest approach, the synthesis of an all-*trans* configuration of the double bonds was sought. It was carried out according to Scheme 2. First, the central symmetric fragment 3,6-dibromophenanthrene (**4**) was prepared from 4,4'-dibromostilbene (**3**) by means of a Mallory reaction.¹⁶ Stilbenic precursor **3** was easily prepared in one step from commercial 4-bromobenzyl bromide taking advantage of the dual electrophilic-nucleophilic role of this reagent in the presence of sodium *p*-toluenesulfonate under basic conditions (KOH).¹⁷ In situ elimination afforded *trans*-stilbene **3** in a good yield, which was photocyclized very efficiently using a 400 W high-pressure Hg lamp (see the Supporting Information) to **4**. Next, two arms containing the desired *E*-2- and *E*-3-thienylvinyl fragments, respectively, were attached to central fragment **4** via a Suzuki coupling, maintaining the stereochemical integrity of *E*-vinylborane reagents **5** and **6**

Scheme 2. Synthesis of All-*trans* (i.e., Configurationally Homogeneous) (*E,E*)-3,6-bis(2-(Thiophen-2-yl)vinyl)phenanthrene (7) and (*E,E*)-3,6-bis(3-(Thiophen-2-yl)vinyl)phenanthrene (8)

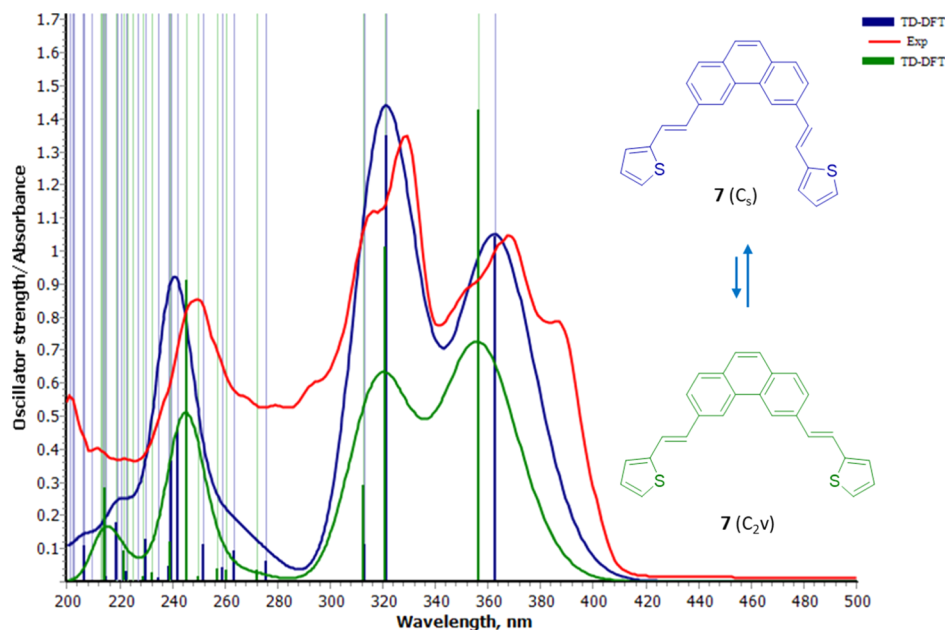
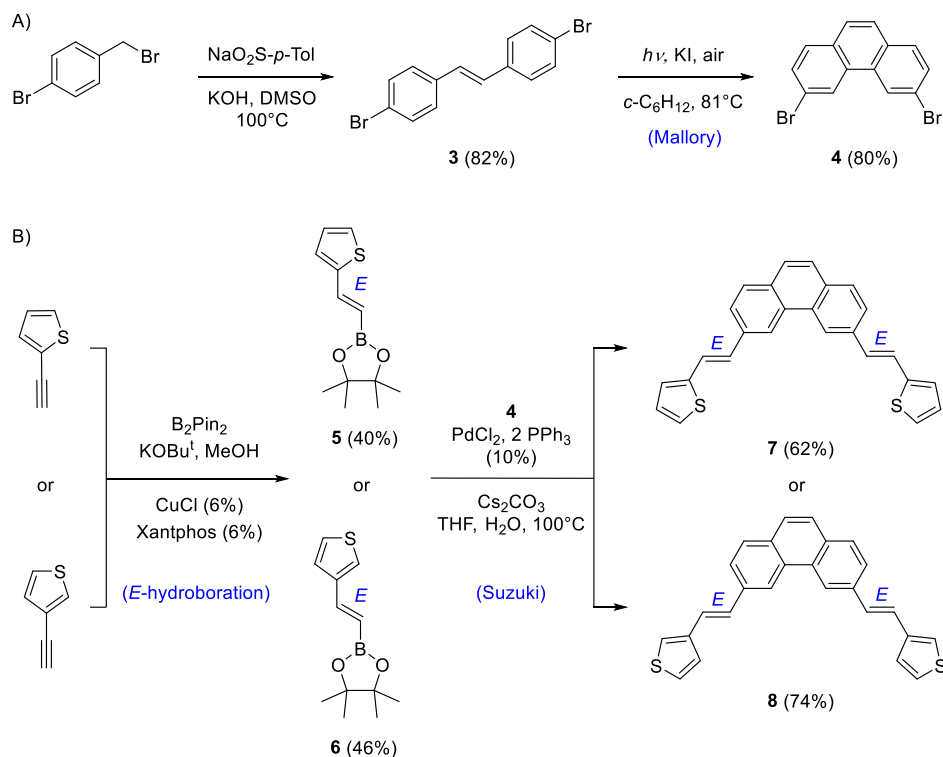


Figure 1. Experimental UV–vis spectrum of compound 7 (red line) in *n*-hexane. Calculated spectra of the most abundant conformer 7(C_s) (blue line) and second most abundant 7(C_{2v}) (green line, with its intensity halved, see the text) superimposed, as well as the calculated transitions (vertical lines) and oscillator strengths (thick vertical lines).

chosen for this purpose. These last reagents were conveniently prepared with a well-defined stereochemistry through a room-temperature CuCl-xantphos-catalyzed hydroboration of the commercially available acetylenic starting materials using bis(pinacolato)diboron (B_2Pin_2) and potassium *t*-butoxide in methanol.¹⁸

Understanding the Photochemical Reaction Pathway. Experimental UV–Vis Spectra of (*E,E*)-3,6-bis(2-

(Thiophen-2-yl)vinyl)phenanthrene (7) and (*E,E*)-3,6-bis(2-(Thiophen-3-yl)vinyl)phenanthrene (8). Styrylthiophene compounds 7 and 8 are the photochemical precursors of the targeted dithia[7]helicenes 1 and 2 in this study. The experimental UV–vis spectra of 7 and 8 in *n*-hexane are included in Figures 1 and 2 (red lines, see also the Supporting Information). In both cases, the absorption patterns share many similarities. For 2-thienylstyryl derivative 7, it consists of

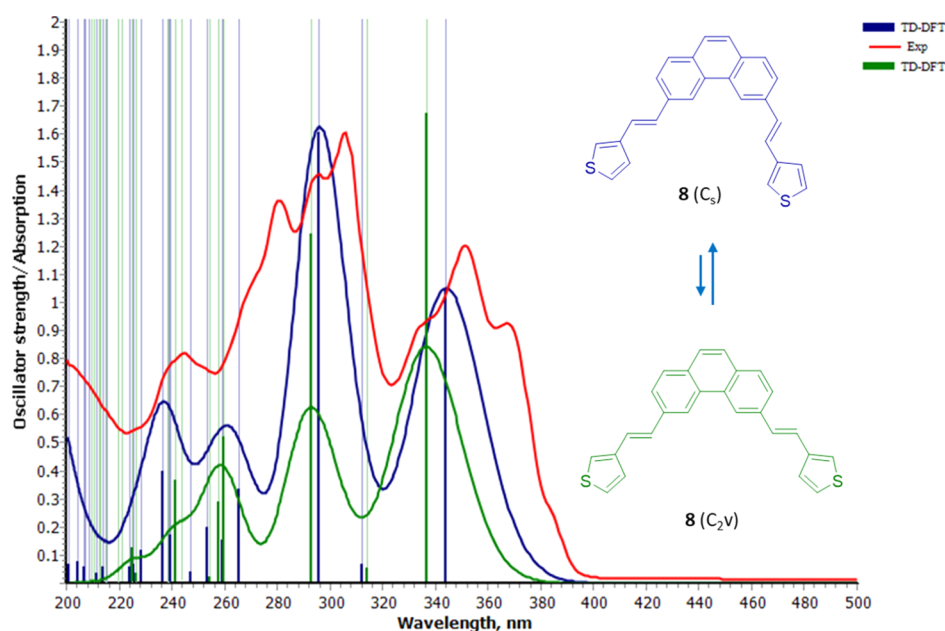


Figure 2. Experimental spectrum of compound **8** (red line) in *n*-hexane. Calculated spectra of the most abundant conformer **8**(C_s) (blue line) and second most abundant **8**(C_{2v}) (green line, with its intensity halved, see the text) superimposed, as well as the calculated transitions (vertical lines) and oscillator strengths (thick vertical lines).

Table 1. Calculated UV–Vis Spectrum of Major Conformations **7**(C_s) and **7**(C_{2v}). State Number, Wavelength (λ), Energy (E), Oscillator Strength (f), and Assignment of the Transitions

| state | λ (nm) | E (eV) | f | assignment |
|-----------------------|----------------|----------|------|-----------------------------|
| 7 (C_s) | | | | |
| S_1 | 362.6 | 3.419 | 1.04 | H \rightarrow L (78%) |
| S_2 | 321.6 | 3.855 | 1.35 | H \rightarrow L + 1 (46%) |
| S_3 | 313.0 | 3.961 | 0.11 | H \rightarrow L + 2 (46%) |
| S_7 | 252.0 | 4.920 | 0.11 | H \rightarrow L + 2 (46%) |
| S_8 | 242.0 | 5.123 | 0.46 | H \rightarrow L + 2 (44%) |
| S_9 | 239.7 | 5.173 | 0.36 | H \rightarrow L + 2 (27%) |
| S_{12} | 229.9 | 5.392 | 0.13 | H \rightarrow L + 2 (21%) |
| S_{18} | 219.0 | 5.662 | 0.18 | H \rightarrow L + 1 (28%) |
| S_{22} | 206.71 | 5.998 | 0.11 | H \rightarrow L + 1 (27%) |
| 7 (C_{2v}) | | | | |
| S_1 | 356.4 | 3.478 | 1.43 | H \rightarrow L (77%) |
| S_2 | 321.2 | 3.861 | 1.01 | H \rightarrow L + 1 (57%) |
| S_3 | 312.7 | 3.965 | 0.29 | H \rightarrow L + 2 (36%) |
| S_8 | 245.4 | 5.053 | 0.91 | H \rightarrow L + 2 (29%) |
| S_9 | 239.2 | 5.183 | 0.12 | H \rightarrow L + 2 (38%) |
| S_{18} | 214.3 | 5.786 | 0.28 | H \rightarrow L + 1 (22%) |

three main absorptions, the first centered at $\lambda_{\max 1} = 368.0$ nm, with a side peak at 385.8 nm (*i.e.*, 1253.7 cm^{-1} away from it) and a nearly symmetrical shoulder at higher energy evidencing a vibrational fine structure. There is another absorption at $\lambda_{\max 2} = 328.1$ nm, also with a shoulder at slightly higher energy, and additional absorptions at lower wavelengths, for example, $\lambda_{\max 3} = 250.0$ nm, less relevant for us since they are out of the reach of the reactor lamp.

For 3-thienylstyryl derivative **8**, the spectrum also consists of three main absorptions, the first two displaying a clear vibrational fine structure again. The first band centered at $\lambda_{\max 1} = 350.9$ nm has a side peak at 366.9 nm (*i.e.*, 1242.7 cm^{-1} away from it) and also a nearly symmetrical shoulder at higher energy. A second main absorption occurs at $\lambda_{\max 2} = 306.0$ nm, with two consecutive well-resolved vibrational peaks at 295.5

and 280.7 nm, and an additional smaller band at $\lambda_{\max 3} = 244.4$ nm that is nonrelevant for our purposes, complete the experimental spectrum.

With this information in hands, we calibrated our quantum chemistry methods and began the theoretical simulation of the electronic spectrum.

Conformational Equilibria of **7 and **8**.** First, we studied the conformational equilibria of **7** and **8**. There are four single bonds in each molecule, giving rise to a quite significant amount of $2^4 = 16$ conformations by rotation, which finally get reduced by symmetry arguments to 10 nonequivalent molecular conformations (only flat, fully delocalized structures were taken into account at this stage). A complete account of all the conformers, their energies, and vibrational analysis can be found in the [Supporting Information](#). From them, a

Table 2. Calculated UV–Vis Spectrum of Major Conformations **8**(C_s) and **8**(C_{2v}). State Number, Wavelength (λ), Energy (E), Oscillator Strength (f), and Assignment of the Transitions

| state | λ (nm) | E (eV) | f | assignment | | |
|-----------------------------|----------------|----------|------|---------------------|---------------------|---------------------|
| 8 (C _s) | | | | | | |
| S ₁ | 344.1 | 3.603 | 1.05 | H → L (82%) | H – 1 → L + 1 (10%) | |
| S ₃ | 295.8 | 4.192 | 1.61 | H → L + 1 (50%) | H – 1 → L (38%) | |
| S ₄ | 265.3 | 4.673 | 0.34 | H – 1 → L + 2 (30%) | H – 2 → L + 2 (16%) | H – 3 → L (10%) |
| S ₅ | 259.0 | 4.787 | 0.15 | H → L + 2 (32%) | H – 2 → L (16%) | H – 1 → L (15%) |
| S ₆ | 253.3 | 4.896 | 0.20 | H – 2 → L + 1 (28%) | H – 2 → L + 2 (13%) | |
| S ₈ | 239.1 | 5.185 | 0.17 | H → L + 8 (21%) | H – 1 → L + 6 (19%) | |
| S ₁₀ | 236.5 | 5.242 | 0.40 | H – 2 → L + 2 (36%) | H – 1 → L + 1 (27%) | H – 3 → L (12%) |
| S ₁₁ | 228.4 | 5.429 | 0.12 | H – 2 → L + 1 (28%) | H – 1 → L + 2 (18%) | H – 2 → L + 2 (15%) |
| S ₃₀ | 196.6 | 6.307 | 0.38 | H – 5 → L + 8 (25%) | H – 4 → L + 6 (19%) | H – 4 → L (11%) |
| 8 (C _{2v}) | | | | | | |
| S ₁ | 336.6 | 3.684 | 1.67 | H → L (82%) | H – 1 → L + 1 (11%) | |
| S ₃ | 292.7 | 4.236 | 1.25 | H – 1 → L (52%) | H → L + 1 (27%) | H → L + 2 (11%) |
| S ₄ | 259.4 | 4.780 | 0.52 | H – 1 → L + 2 (47%) | H – 2 → L + 2 (18%) | H – 3 → L (11%) |
| S ₅ | 257.6 | 4.813 | 0.29 | H – 2 → L + 1 (39%) | H → L + 12 (16%) | H – 1 → L + 1 (14%) |
| S ₈ | 241.2 | 5.142 | 0.37 | H – 1 → L + 1 (38%) | H – 2 → L + 1 (28%) | H – 3 → L (13%) |
| S ₁₂ | 225.0 | 5.511 | 0.13 | H – 4 → L (27%) | H – 4 → L + 6 (13%) | H – 5 → L + 1 (11%) |

Maxwell–Boltzmann distribution reveals two prevailing conformations very close in energy (<0.4 kcal/mol, see the [Supporting Information](#)) that stand out, accounting for more than 60% of the overall conformational population. They are represented in [Figures 1](#) and [2](#) (inset).

The calculated spectral shapes of **7**(C_s) and **8**(C_s) fit reasonably well with the experimental spectra, reproducing $\lambda_{\text{max}1}$, $\lambda_{\text{max}2}$, and somewhat less the more complex band at $\lambda_{\text{max}3}$, as well as the intensities of their absorbances. Their populations are doubled with respect to the minor C_{2v} conformers since they have degenerated structures and benefit from the statistical effect of a smaller symmetry number (C_s compared to the C_{2v} point group).¹⁹ For the minor conformers **7**(C_{2v}) and **8**(C_{2v}), the calculated transitions are of similar energy to the C_s ones, while their intensities are mismatched (green vs blue oscillator strengths, vertical thick lines, and overall calculated spectral shapes). Thus, by comparison with the experimental spectrum, it is evident that the statistical effect due to the symmetry overcomes the rather negligible difference in energy calculated between both conformers. We halved the simulated spectrum for C_{2v} conformers in [Figures 1](#) and [2](#) to stress this argument. We have therefore identified the transitions that give rise to the relevant absorption bands at $\lambda_{\text{max}1}$ and $\lambda_{\text{max}2}$, always keeping in mind that the experimental absorptions have a more complex pattern originated from vibronic coupling; this simulation is out of the reach of this work.

Calculated Electronic Spectra of the Major Conformations of **7 and **8**.** The assignment of the electronic spectra for the most representative conformations **7**(C_s) and **8**(C_s) and minor counterparts **7**(C_{2v}) and **8**(C_{2v}) calculated by time-dependent density functional theory (TDDFT) at the wB97XD/6-311++G(2d,2p) level of theory in *n*-hexane as the polarizable continuum model (PCM) is collected in [Tables 1](#) and [2](#). We confine our discussion to the transitions with oscillator strengths $f > 0.10$ and contributions with more than 10% weight.

Let us focus now on the key transitions of the simulated spectra. For **7**, the vertical transition to the S₁ state from the ground state S₀ for any of its major conformations is a $\pi \rightarrow -\pi^*$ type of excitation, mainly described by the highest

occupied molecular orbital (HOMO) → lowest unoccupied molecular orbital (LUMO) transition combined with a HOMO – 1 → LUMO + 1 in a much smaller weight along with other even minor unreported contributions ([Table 1](#)). In [Figure 3a](#), we can see this transition for the major conformer **7**(C_v) at $\lambda_1 = 362.6$ nm as a natural transition orbital (NTO),²⁰ representing the best hole/particle pair representation of the transition density matrix for that excited state. The MO diagram in the middle shows the electronic configurations that span the actual excited-state wavefunction. The composition of this peak is very similar for both conformers C_s and C_{2v} (and indeed for the rest of the less representative conformers studied) and corresponds to the experimental $\lambda_{\text{max}1} = 368.0$ nm. As can be seen in [Figure 3](#), the binding pair of carbons are the C(4) phenanthrene position and the C(3) of thiophene (green arrows), with a typical *supra-antara* stereochemistry as well as a weakened stilbenic double bond (antibonding interaction). This will enable its dihedral rotation and meeting of the reacting pair of carbons as the reactions further progress in the S₁ surface. The main characteristics of these NTOs are conserved for both the *E*- and *Z*-isomers, the excited state of which will eventually drive to the expected naphtho[2,1-*b*]thiophene type of ring fusion as we will see later.

The following calculated vertical transition is S₂, located at $\lambda_2 = 321.6$ nm for the major conformer **7**(C_v) and nearly the same for the minor, and corresponds to the experimental $\lambda_{\text{max}2} = 328.1$ nm. A description of the MOs involved in this vertical transition to the S₂ state for **7** can be found in the [Supporting Information](#). This state also provides an analogous reaction pathway with a regio- and stereochemical outcome akin to the above explained for S₁. Both S₁ and S₂ states can explain the photochemical reaction outcome, although the lowest excited state is usually the relevant one in terms of photochemical production based on half-life arguments (Kasha's rule,²¹ extended to chemical deactivation).

We now move to the more interesting case **8**. For **8**, again the vertical transition S₀ → S₁ for any of its major conformations is a $\pi \rightarrow -\pi^*$ type of excitation, mainly described by a HOMO → LUMO transition combined with a HOMO – 1 → LUMO + 1 in a much smaller weight ([Table 2](#)). In [Figure 3b](#), we can see this transition for the major

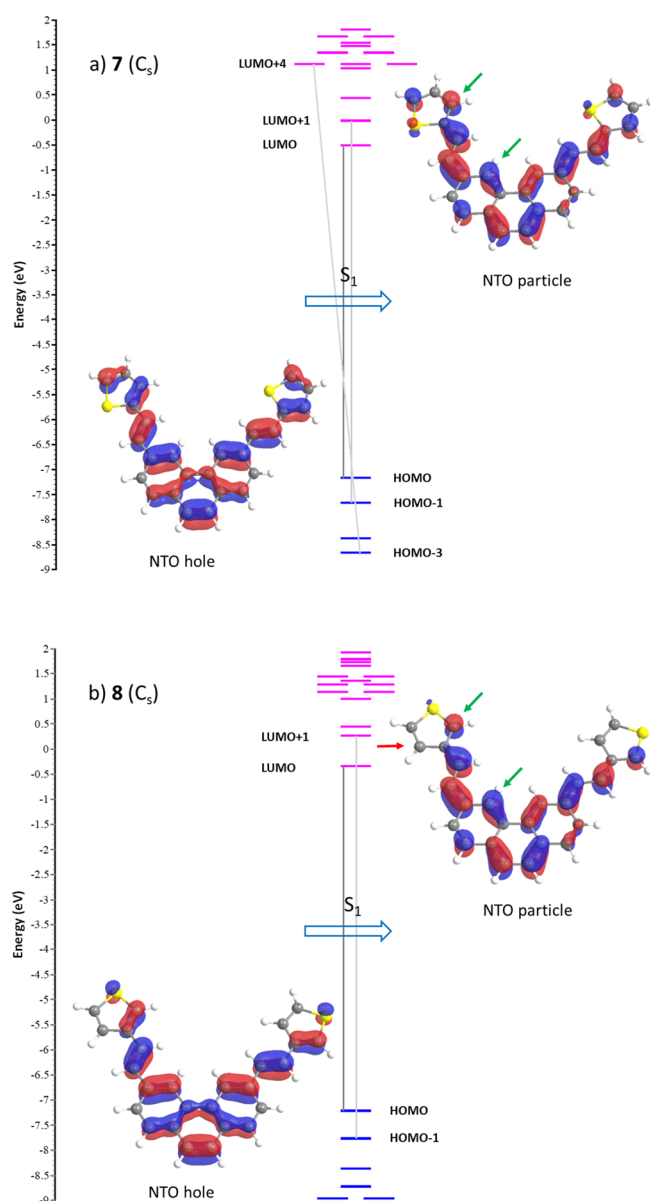


Figure 3. Vertical transition to the first excited state S_1 for the major conformations of (a) **7**(C_s) and (b) **8**(C_s), represented as the hole and particle NTO pair [in this case, HOMO \rightarrow LUMO weighing 78 and 82% of the transition for (a,b), respectively], and the diagram of MOs involved in the excited states. Green arrows signal the reacting pair of carbons.

conformer **8**(C_v) at $\lambda_1 = 344.1$ nm as an NTO, along with the MO diagram of the transition. Just as in the former case, the composition of this peak is very similar for both conformers C_s and C_{2v} and the rest of the less representative conformers and corresponds to the experimental $\lambda_{\text{max}1} = 350.9$ nm. It can be deduced from Figure 3b that the binding pair of carbons are the C(4) phenanthrene position and the C(2) of thiophene (green arrows), while the potentially competing C(4) is kinetically unreactive (red arrow) through this excitation pathway. Once evolved to the Z -conformation, its excited state will eventually lead to the expected naphtho[1,2-*b*]thiophene type of ring fusion but not to the naphtho[1,2-*c*] one (see also structures in Scheme 1), as we will see later.

The following calculated vertical transition of relevance is S_3 , located at $\lambda_2 = 295.8$ nm for the major conformer **8**(C_v) and

very close for the minor one (C_{2v}), and corresponds to the experimental $\lambda_{\text{max}2} = 306.0$ nm. Similar arguments as those explained before hold in here. A description of the MOs involved in this vertical transition to the S_3 state for **8** can be found in the Supporting Information. Again, this state also provides an analogous reaction pathway with a regio- and stereochemical outcome akin to the above explained for S_1 .

Calculated Excited States of 7 and 8 and the Photochemical Reaction Pathway. The above-described vertical transitions may evolve within the excited-state hypersurface to minimal energy conformations and eventually to a suitable excited Z -configuration for the cyclization step. We have tracked this evolution in the excited state S_1 for **7** and **8**, starting from the most symmetric conformations (C_{2v}) for simplicity. Calculations were done by density functional theory (DFT) and TDDFT at the wB97XD/6-311++G(2d,2p) level of theory in *n*-hexane as the PCM. Results are collected in Figure 4 and the Supporting Information. Upon vertical excitation, **8** (C_{2v}) (**I** in Figure 4) in the first place evolves to a planar excited state **I*** (relative minimum) that further relaxes to **IIa*** (absolute minimum in S_1), while **IIb*** (also a relative minimum) is some 5.1 kcal/mol above the former. Both are the two relevant conformers of the excited state S_1 with a twisted but near to Z -configuration of the stilbenic double bond. The study of the changes in the electron density upon excitation in **IIa*** and **IIb*** by NTO analysis is revealing. **IIa*** has an important binding interaction between the carbons C(4) of phenanthrene and C(2) of thiophene, which imparts extra stability to this excited state and entails an incipient sigma bond formation (Figure 5a, green arrow). After internal conversion and vibrational relaxation to the ground state, it may lead to the dihydro-intermediate **III** or alternatively the Z -isomer **IIa** (Figure 4, green reaction pathway). A typical *supra-antara* stereochemistry and a weakened stilbenic double bond (C=C=C-C dihedral angles of 30 and 31° for **IIa*** and **IIb***, compared to nearly 0° for the ground-state Z -configurations) are in agreement with the expected reaction course. Conversely, for **IIb***, the lack of an efficient overlap destabilizes this conformation and prevents the evolution toward sigma-bonded products (Figure 4, red reaction pathway and Figure 5b, red arrow). Binding through the C(4) of thiophene would require a dihydro-intermediate having an expanded valence shell for sulfur (represented in Scheme 1b by a dotted bond), a rare pattern in organosulfur chemistry that has defied every attempt we made trying to model it, as it always evolves toward the stable Z -double bond conformation **IIb**.

Starting from **7**, an allowed reaction pathway analogous in every way to that described above is found. The corresponding energy diagram for **7** can be found in the Supporting Information. Overall, these mechanistic scenarios corroborate the anticipated reaction pathways outlined in Figure 3 at the vertical transition stage.

Final Photochemical Synthesis of Dithia[7]helicenes Starting from 7 and 8. *Bis*-stilbenic compounds **7** and **8** were subjected to irradiation using a 400 W high-pressure Hg lamp to undergo a sequential double photocyclization under typical Mallory–Katz conditions²² (Scheme 3 and the Supporting Information). The reaction afforded the expected 3,14-dithia[7]helicene-1 (or *exo*-dithia[7]helicene-1) for **7**. A distinctive fingerprint useful in the structural elucidation of this type of thiahelicenes by ¹H NMR is the two doublets originated by the coupling of the vicinal hydrogens of the

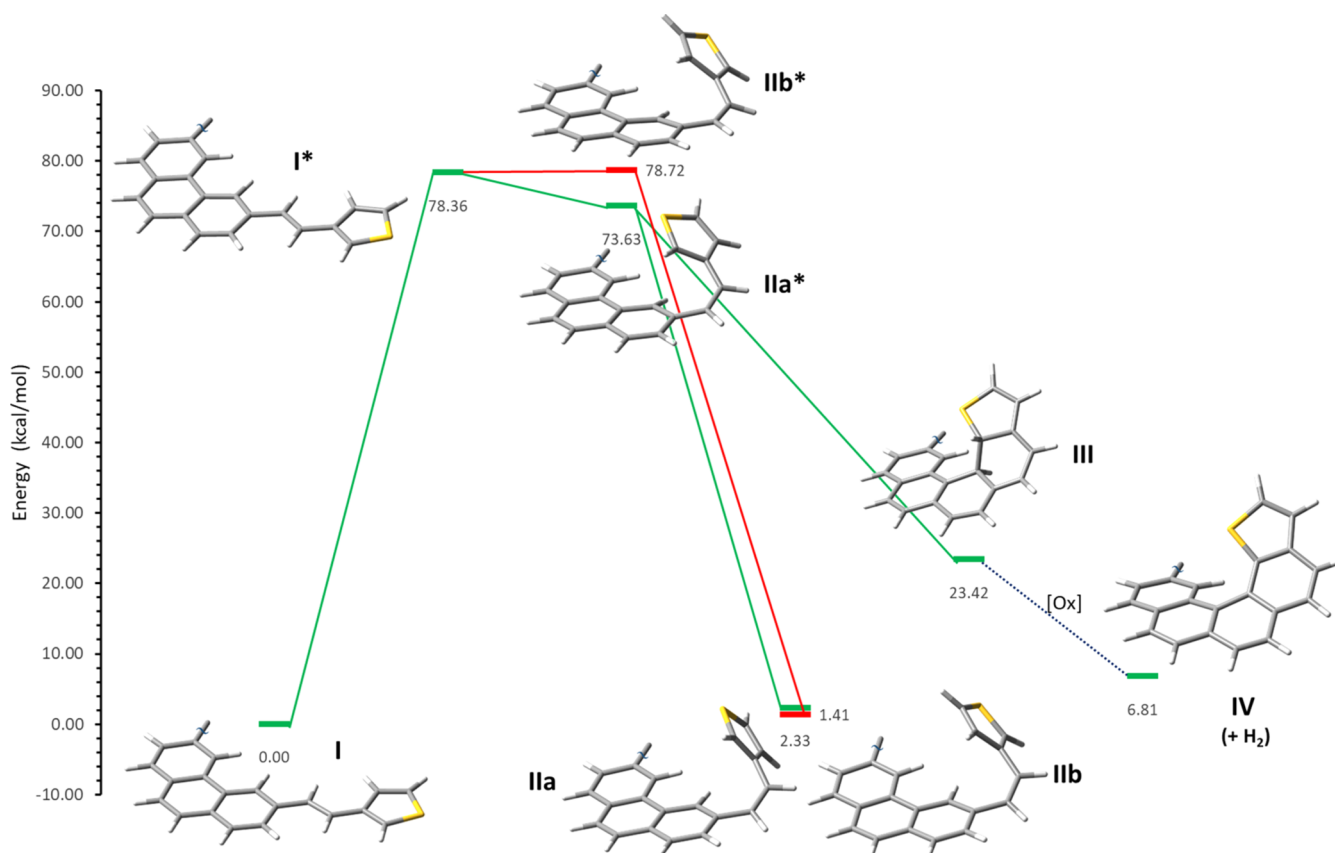
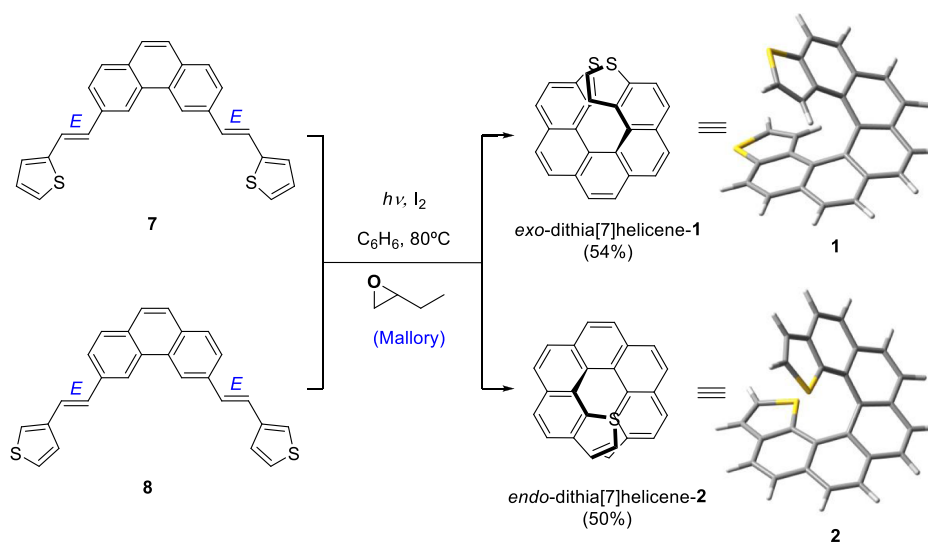


Figure 4. Diagram of relative energy showing the photochemical pathways that **8** (C_{2v}) (represented here as **I**) may follow upon $S_0 \rightarrow S_1$ excitation. Calculated absolute and relative minima located in the ground and excited states (*) are also represented. An allowed route that goes through **IIa*** leading both to $E \rightleftharpoons Z$ isomerization ($I \rightleftharpoons IIa$) and to the cyclized product ($I \rightleftharpoons III$) is shown in green. Conformer **IIb*** leading only to $E \rightleftharpoons Z$ isomerization ($I \rightleftharpoons IIb$) is shown in red. No stationary cyclized intermediate was found that could lead to the helicenic final product.

Scheme 3. Photochemical Synthesis of Dithia[7]helicenes 1 and 2 via a Double Photocyclization of bis-Stilbenic Precursors 7 and 8



thiophene ring. They are easily spotted in the spectrum for two reasons. Even though the chemical shifts of thiophene hydrogens fall within the chemical shift of a typical aromatic hydrogen (e.g., benzene), within the framework of a helicenic structure, they are shifted upfield as a consequence of the anisotropic effect of the rest of the underlying aromatic structure. Also, the coupling constant (J) of these vicinal

hydrogens is substantially smaller and easily identified from the rest of the aromatic signals. The result is a pair of doublets in the 6–7 ppm region with J around 5–6 Hz. In the case of **1**, there are two doublets at 6.60 (d, $J = 5.6$ Hz, 2H) and 6.24 (dd, $J = 5.6, 0.5$ Hz, 2H), clearly identifying a naphtho[2,1-*b*]thiophene type of ring fusion (Figure 6a). For **8**, the only compound that could be isolated has two doublets at 7.06 (d, J

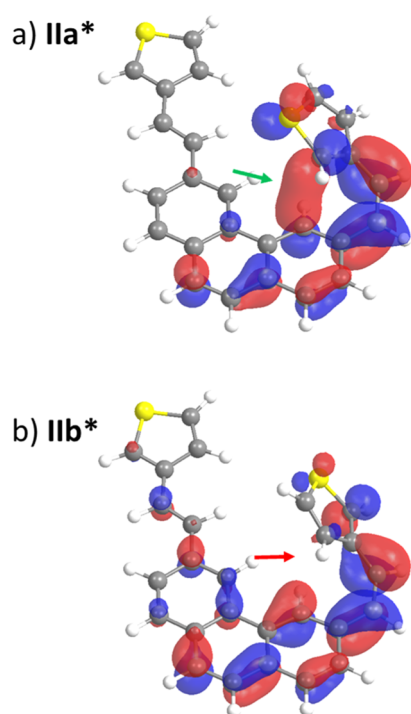


Figure 5. Relevant minima **IIa*** and **IIb*** of the first excited state S_1 of **8** and changes in the electron density represented as the particle NTO. A developing sigma-bonding interaction in **IIa*** (green arrow) and the lack of the same in **IIb*** (red arrow) is shown.

= 5.4 Hz, 2H) and 6.80 (d, $J = 5.4$ Hz, 2H), consistent with a naphtho[1,2-*b*]thiophene type of ring fusion as in **2** (Figure 6b), but not with a naphtho[1,2-*c*]thiophene one, which would display a much smaller J (estimated $J = 2.7$ – 2.8 Hz from gauge-invariant atomic orbital DFT calculations, scaled using experimental J_s from Figure 6). The isolated compound was hence identified as 1,16-dithia[7]helicene-2 (or *endo*-dithia[7]helicene-2). Other than the dithia[7]helicene or starting material, only intractable, unidentifiable products by NMR were obtained in different trials whether the reactions were taken to completion or not (*i.e.*, to the disappearance of the starting material). An unequivocal characterization of **1** and **2** by means of X-ray crystallography was recently reported by our group.⁵

CONCLUSIONS

In summary, the photocyclization mechanism of 2- and 3-styrylthiophenes to afford naphtho[2,1-*b*]thiophene and naphtho[1,2-*b*]thiophene type of ring fusions, respectively, has been studied combining the conclusive information resulting from synthesis/structural elucidation with DFT and TDDFT methods to unveil mechanisms. In the first place, we carried out the syntheses of (*E,E*)-**7** and (*E,E*)-**8** and obtained their experimental UV–vis spectra. Then, we simulated these electronic spectra and analyzed the nature of their vertical electronic transitions by TDDFT. This gave us access to the excited states involved in the reaction. These excited states were geometrically optimized and analyzed again to gain information on the photochemical intermediates of the reaction. The regiochemical and stereochemical outcome of the photocyclization step is revealed at this point. Finally, back to the synthesis, the actual photocyclization of **7** and **8** afforded the theoretically anticipated dithiahelicenes **1** and **2**, the

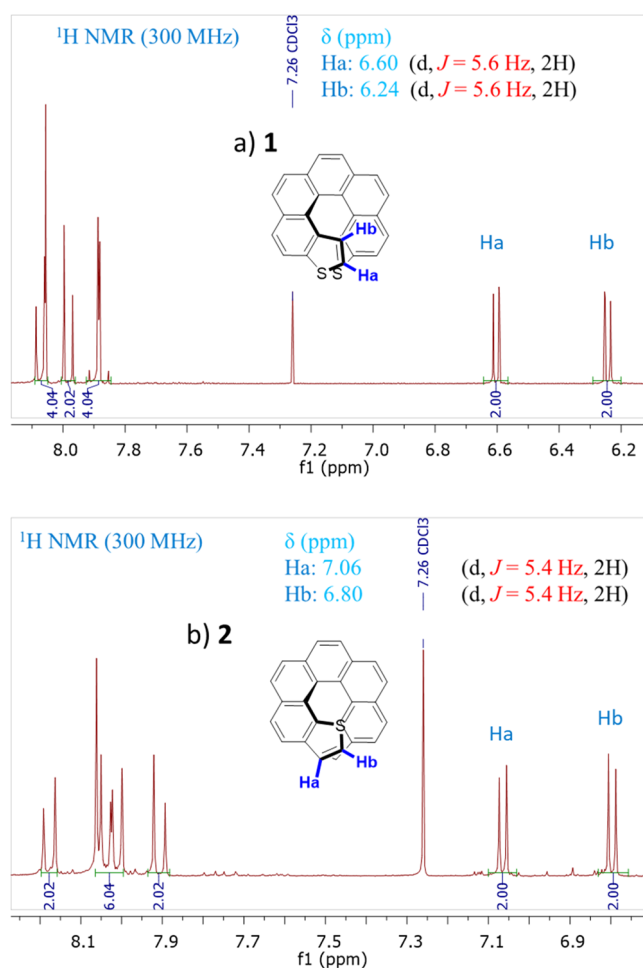


Figure 6. Terminal thiophene hydrogens show a distinctive pattern by ^1H NMR when forming parts of helicenes, consisting of an upfield chemical shift and a characteristic coupling constant of these vicinal hydrogens, which makes them easily recognizable. This pattern is shown in (a) for **1** and in (b) for **2**, corroborating their naphtho[2,1-*b*]thiophene and naphtho[1,2-*b*]thiophene type of ring fusion, respectively.

structure of which could be unambiguously assigned by ^1H NMR spectroscopy. Full structural characterization of the resulting dithiahelicenes leaves little doubts about the findings here reported, which show the potential of excited-state quantum mechanical methods in structural and mechanistic elucidation tasks.

EXPERIMENTAL SECTION

General Methods. Unless otherwise stated, commercially available starting materials and solvents for chromatography and recrystallization were used without further purification. We dried and distilled over Na/K alloy tetrahydrofuran (THF), benzene, and cyclohexane immediately before using them in synthesis or photochemistry. Commercially unavailable reagents were synthesized by different means; they are explained below each one separately.

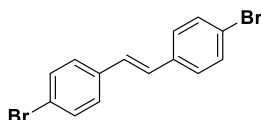
Gas chromatography analyses (GLC) were carried out with a Hewlett Packard HP-5890 instrument equipped with a flame ionization detector and a 30 m HP-5 capillary column (0.32 mm diameter and 0.25 μm film thickness) using nitrogen as carrier gas (12 psi). Column chromatography was performed with Merck silica gel 60 (0.040–0.063 μm , 240–400 mesh). Thin-layer chromatography (TLC) was performed on precoated silica gel plates (Merck 60, F254, 0.25 mm). TLC detection was done by UV₂₅₄ light, and R_f

values are given under these conditions. NMR spectra were recorded on a Bruker Avance 300 and a Bruker Avance 400 (300 and 400 MHz for ^1H NMR and 75 and 100 MHz for ^{13}C NMR, respectively, broadband proton decoupling was applied during the acquisition of $^{13}\text{C}\{^1\text{H}\}$ spectra) using CDCl_3 as a solvent and tetramethylsilane (TMS) as an internal standard. Chemical shifts (δ) are given in ppm versus TMS. Infrared (IR) analysis was performed with a JASCO FT/IR 4100 spectrophotometer equipped with an attenuated total reflection component. Low-resolution mass spectrometry was performed using the electron impact (EI) mode at 70 eV in an AGILENT 5973N mass spectrometer coupled with an AGILENT 6890N gas chromatographer. High-resolution mass spectrometry (HRMS) analyses were carried out in an AGILENT 7200 using the EI mode at 70 eV by quadrupole time-of-flight (Q-TOF). Melting points were performed with a Reichert Thermovar polarizing light microscope and a melting point apparatus and have been corrected. A double-beam UV-vis spectrophotometer (Shimadzu UV-1603) was used for recording the electronic spectra.

Photochemistry. A 400 W high-pressure mercury lamp (Osram HQL MBF-U) was modified by cutting away the outer glass envelope from the screw base (preserving the inner quartz arc tube containing Hg) and was mounted in a porcelain lamp holder provided with an aluminum reflector. The lamp was connected to a corresponding power unit, and the light beam was focused to a number of 100 mL Schlenk's single-wall borosilicate tubes (0.6 mm wall thickness) placed some 10 cm away from the source, provided with magnetic stirring and a vertical condenser refrigerated with a recirculating chiller (Huber MPC-K6) using a 30% ethylene glycol–water mixture as a coolant. The chemical hood was lined with aluminum foil to avoid unwanted exposure to UV radiation. We used cyclohexane or benzene under reflux as solvents.

Synthesis and Characterization of Compounds. Experimental Procedures for the Synthesis of Intermediates and Final Products

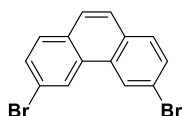
*(E)-1,2-bis(4-Bromophenyl)ethene (3).*²³ This compound was prepared by the procedure described in the literature.²³ In a 100



mL round flask, 4-bromobenzyl bromide (5.00 g, 20 mmol, 2 equiv), sodium *p*-toluenesulfonate (1.78 g, 10 mmol, 1 equiv), and KOH (1.68 g, 30 mmol, 3 equiv) were added, and then 25 mL of dimethyl sulfoxide (DMSO) was added. The mixture was heated in an oil bath under reflux (100 °C) for 24 h. After cooling to room temperature, the solvent (DMSO) was removed by vacuum distillation, and a brown solid was obtained. The product was purified by column chromatography on silica gel (hexane) to afford a white solid (2.772 g, 82% yield). A different way to purify the product was by recrystallization from $\text{EtOH}/\text{CHCl}_3$ (3:1), obtaining **3** as white crystals.

White solid; mp 195.3 °C (corrected); R_f = 0.53 (hexane); ^1H NMR (CDCl_3 , 300 MHz): δ = 7.51–7.46 (m, 4H), 7.39–7.34 (m, 4H), and 7.02 (s, 2H). $^{13}\text{C}\{^1\text{H}\}$ NMR (CDCl_3 , 75 MHz): δ = 136.1, 132.0, 128.3, 128.2, and 121.8. MS (EI) m/z : 339.90 (M^+ + 4, 31.7), 337.90 (M^+ + 2, 61.2), 335.90 (M^+ , 32.5), 281.05 (22.2), 258.00 (6.8), 207.05 (53.4), 180.00 (2.7), 179.10 (17.9), 178.10 (100), 176.10 (24.7), 152.15 (10.3), 126.05 (3.3), 89.05 (23.0), and 88.10 (21.0). IR (neat) ν_{max} : 3020, 2924, 1581, 1481, 1404, 1323, 1215, 1068, 999, 968, 822, and 710 cm^{-1} .

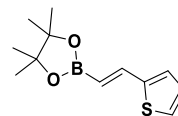
*3,6-Dibromophenanthrene (4).*²⁴ This compound was prepared using the main photochemical setup described above. A solution/



suspension of *(E)*-1,2-bis(4-bromophenyl)ethene (33.6 mg, 0.1 mmol; 1 equiv) and KI (16.6 mg, 0.1 mmol; 1 equiv) in cyclohexane (100 mL) was prepared in a Schlenk tube provided with a vertical condenser open to the air connected to a chiller. The recirculating chiller was turned on, and the mixture was irradiated with a 400 W high-pressure Hg lamp for 3–4 h under reflux. The progress of the reaction was monitored by GLC. After the reaction was completed, the crude mixture was washed with NaHSO_3 , dried under magnesium sulfate, and filtered, and the solvent was evaporated under reduced pressure (15 Torr). The product was purified by column chromatography of silica gel (hexane) to afford a white solid (26.8 mg, 80% purified yield).

White solid; mp 191.6 °C (corrected); R_f = 0.53 (hexane); ^1H NMR (CDCl_3 , 300 MHz): δ = 8.72 (d, J = 1.7 Hz, 2H), 7.76 (d, J = 8.3 Hz, 2H), and 7.73–7.69 (m, 4H). $^{13}\text{C}\{^1\text{H}\}$ NMR (CDCl_3 , 75 MHz): δ = 131.0, 130.9, 130.6, 130.2, 126.9, 125.7, and 121.4. MS (EI) m/z : 337.90 (M^+ + 4, 50.0), 335.90 (M^+ + 2, 100), 333.90 (M^+ , 50.2), 176.10 (8.1), 150.10 (10), and 88.05 (29). IR (neat) ν_{max} : 2923, 2851, 1901, 1586, 1494, 1429, 1407, 1381, 1153, 1106, 1069, 1017, 856, 846, 835, and 769 cm^{-1} .

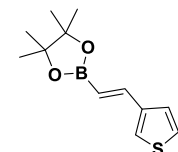
*(E)-4,4,5,5-Tetramethyl-2-(2-(thiophen-2-yl)vinyl)-1,3,2-dioxaborolane (5).*²⁵ This compound was prepared by adapting a



procedure from the literature.¹⁸ In an oven-dried Schlenk tube CuCl (2.96 mg; 0.03 mmol; 0.06 equiv), NaOt-Bu (5.76 mg; 0.06 mmol; 0.12 equiv) and xantphos ligand (17.35 mg; 0.03 mmol; 0.06 equiv) were added, and the reaction mixture was then subjected to three cycles of vacuum/argon. Then, 1 mL of dry THF was injected, and the solution was stirred for 30 min at room temperature. Next, bis(pinacolato)diboron (253.94 mg; 1 mmol; 2 equiv) in 0.5 mL of dry THF was added. The solution was stirred for 10 min at room temperature, and 2-ethynylthiophene (54.08 mg; 0.5 mmol; 1 equiv) was added, followed by MeOH (42 μL , 1 mmol). The reaction mixture was stirred at room temperature overnight (no starting material was detected by TLC). Then, it was filtered through a pad of celite, and the residue was purified by preparative TLC (silica gel, hexane–EtOAc 9:1) obtaining a pale yellow oil (47.2 mg, 40% yield).

Pale yellow oil; R_f = 0.51 (hexane–EtOAc 9:1); ^1H NMR (CDCl_3 , 300 MHz): δ = 7.47 (d, J = 18.1 Hz, 1H), 7.24 (d, J = 5.1 Hz, 1H), 7.08 (d, J = 3.5 Hz, 1H), 6.99 (dd, J = 5.0, 3.6 Hz, 1H), 5.91 (d, J = 18.1 Hz, 1H), and 1.30 (s, 12H). $^{13}\text{C}\{^1\text{H}\}$ NMR (CDCl_3 , 75 MHz): δ = 144.1, 141.9, 127.8, 127.8, 126.4, 83.5, and 24.9. MS (EI) m/z : 238.1 (M^+ + 2, 5.6), 237.1 (M^+ + 1, 13.4), 236.1 (M^+ , 88.8), 235 (M^+ – 1, 23.2), 221.1 (20.5), 193.05 (7.3), 178.05 (10.7), 163.00 (26.8), 151.10 (41.1), 136.00 (100), 111.00 (27.9), 85.10 (8.9), and 57.10 (5.2). IR (neat) ν_{max} : 2974, 2911, 2168, 1616, 1520, 1458, 1423, 1373, 1327, 1234, 1146, 976, 910, 849, and 733 cm^{-1} .

*(E)-4,4,5,5-Tetramethyl-2-(2-(thiophen-3-yl)vinyl)-1,3,2-dioxaborolane (6).*²⁵ This compound was prepared following the previous

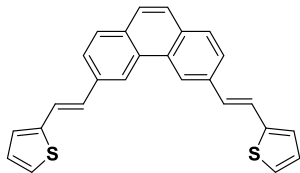


procedure, replacing the terminal alkyne by 3-ethynylthiophene as a pale yellow oil (54.3 mg, 46% yield).

Pale yellow oil; R_f = 0.52 (hexane–EtOAc 9:1); ^1H NMR (CDCl_3 , 300 MHz): δ = 7.38 (d, J = 18.4 Hz, 1H), 7.32–7.28 (m, 2H), 7.28–7.24 (m, 1H), 5.95 (d, J = 18.3 Hz, 1H), and 1.30 (s, 12H). $^{13}\text{C}\{^1\text{H}\}$ NMR (CDCl_3 , 75 MHz): δ = 143.3, 141.3, 126.2, 125.1, 125.0, 83.4, and 24.9. MS (EI) m/z : 238.10 (M^+ + 2, 4.1), 237.10 (M^+ + 1, 10.4), 236.10 (M^+ , 70.2), 235.10 (M^+ – 1, 15.6), 221.10 (17.5), 192.10 (7.2), 178.10 (26.5), 163.05 (54.8), 150.10 (26.2), 136.00 (100),

110.05 (12.1), 85.05 (9.4), and 57.10 (5.5). IR (neat) ν_{\max} : 2927, 2858, 2167, 2025, 1724, 1624, 1516, 1458, 1331, 1261, 1146, 991, 849, 771, and 690 cm^{-1} .

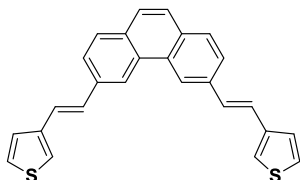
3,6-bis((E)-2-(Thiophen-2-yl)vinyl)phenanthrene (7). This compound was prepared by adapting to our substrates a Suzuki coupling



described in the literature.²⁶ In an oven-dried pressure tube, PdCl₂ (17.73 mg; 0.1 mmol; 0.20 equiv), PPh₃ (52.45 mg; 0.2 mmol; 0.40 equiv), Cs₂CO₃ (977.46 mg; 3 mmol; 6 equiv), and 3,6-bibromophenanthrene (168.3 mg; 0.5 mmol; 1 equiv) were added. The tube was sealed with a septum, and after being subjected to three cycles of vacuum/argon, 4,4,5,5-tetramethyl-2-(2-(thiophen-2-yl)-vinyl)-1,3,2-dioxaborolane (360.3 mg; 1.5 mmol; 3 equiv) dissolved in 3.6 mL of THF was added with a syringe, followed by 0.4 mL of H₂O. The threaded tube was closed and heated in an oil bath at 85 °C for 20–24 h. The reaction was monitored by TLC. After the reaction was completed, it was extracted using 10 mL of H₂O and 3 × 10 mL of CH₂Cl₂. A rather insoluble solid in suspension was observed in the organic phase. The product, yellow solid, was separated by filtration. On the other hand, the combined extracts were solvent-evaporated and purified by column chromatography with silica gel (hexane–EtOAc 9:1) to obtain an additional fraction of the same yellow solid which was incorporated to the formerly filtered solid. The product was obtained as a yellow solid (122.3 mg, 62% yield).

Yellow solid; mp 203.8 °C (corrected); R_f = 0.55 (hexane–EtOAc 9:1); ¹H NMR (CDCl₃, 400 MHz): δ = 8.66 (d, J = 0.65 Hz, 2H), 7.84 (d, J = 8.3 Hz, 2H), 7.77 (dd, J = 8.4, 1.4 Hz, 2H), 7.67 (s, 2H), 7.46 (d, J = 16.0 Hz, 2H), 7.26 (d, J = 3.5 Hz, 2H), 7.23 (d, J = 16.0 Hz, 2H) 7.17 (d, J = 3.5 Hz, 2H), and 7.06 (dd, J = 5.1, 3.5 Hz, 2H).²⁷ MS (EI, DIP) m/z : 396.2 (M^+ + 2, 24.7), 395.2 (M^+ + 1, 58.6), 394.2 (M^+ , 100), 359.1 (7.1), 308.1 (10.3), 276.1 (8.7), 197.1 (18.8), 179.3 (5.8), and 154.2 (3.4). IR (neat) ν_{\max} : 3097, 1786, 1608, 1508, 1423, 1342, 1192, 949, 887, 845, and 698 cm^{-1} . HRMS (EI/Q-TOF) m/z : [M]⁺ calcd for C₂₆H₁₈S₂, 394.0850; found, 394.0853.

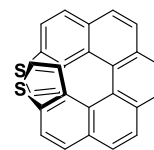
3,6-bis-((E)-2-(Thiophen-3-yl)vinyl)phenanthrene (8). This compound was prepared following the previous procedure but using



4,4,5,5-tetramethyl-2-(2-(thiophen-3-yl)vinyl)-1,3,2-dioxaborolane as the starting reagent. The product was obtained as a yellow solid (146.0 mg, 74% yield).

Yellow solid in a 74% yield; mp 193.4 °C (corrected); R_f = 0.60 (hexane–EtOAc 9:1); ¹H NMR (CDCl₃, 400 MHz): δ = 8.66 (s, 2H), 7.84–7.75 (m, 4H), 7.65 (s, 2H), 7.45 (d, J = 4.2 Hz, 2H), 7.40–7.29 (m, 6H), and 7.23 (d, J = 16.0 Hz, 2H).²⁷ MS (EI, DIP) m/z : 396.1 (M^+ + 2, 13.0), 395.1 (M^+ + 1, 30.3), 394.1 (M^+ , 100), 360.1 (9.8), 309.1 (14.1), 284.1 (9.4), and 197.0 (9.1). IR (neat) ν_{\max} : 3089, 1608, 1408, 1203, 1087, 957, 845, 771, and 690 cm^{-1} . HRMS (EI/Q-TOF) m/z : [M]⁺ calcd for C₂₆H₁₈S₂, 394.0850; found, 394.0848.

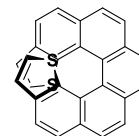
3,14-Dithia[7]helicene (1).^{5,14} The reaction was run in parallel in two oven-dried Schlenk tubes, loaded with a suspension of 3,6-bis-((E)-2-(thiophen-2-yl)vinyl)phenanthrene (19.73 mg; 0.05 mmol; 1 equiv), I₂ (38.07 mg; 0.15 mmol; 3 equiv), and 1,2-epoxybutane (360.50 mg; 5 mmol; 100 equiv) in 100 mL of benzene each. The tubes were provided with vertical condensers connected to a chiller, and Ar was bubbled into the solution during the reaction. The



recirculation chiller was turned on, and the mixture was irradiated with a 400 W high-pressure Hg lamp for 3 h under reflux. The progress of the reaction was monitored by TLC. After the reaction was completed, it was washed with aqueous NaHSO₃, dried over magnesium sulfate, and filtered, and the solvent was evaporated under a reduced pressure (15 Torr). The residue was purified by column chromatography on silica gel (hexane–CH₂Cl₂ 8:2) to obtain a yellow solid. A 54% yield was calculated by ¹H NMR using 1,3,5-trimethoxybenzene (99%) as the internal standard. Recrystallization from EtOAc afforded slightly brown, near colorless crystals (6.8 mg, 35% yield).

Light brown, near colorless crystals; R_f = 0.48 (hexane/AcOEt 9:1); ¹H NMR (CDCl₃, 300 MHz): δ = 8.07 (d, J = 8.9 Hz, 2H), 8.06 (s, 2H), 7.98 (d, J = 8.4 Hz, 2H), 7.93–7.84 (m, 4H), 6.60 (d, J = 5.6 Hz, 2H), and 6.24 (d, J = 5.6, 2H). ¹³C{¹H} NMR (CDCl₃, 75 MHz): δ = 138.6, 135.3, 132.2, 129.9, 127.9, 127.9, 127.1, 125.4, 125.1, 124.6, 124.2, 123.0, and 121.4.

1,16-Dithia[7]helicene (2).⁵ In this case, a suspension of 3,6-bis-((E)-2-(thiophen-3-yl)vinyl)phenanthrene (19.72 mg; 0.05 mmol;



1 equiv), I₂ (38.07 mg; 0.15 mmol, 3 equiv), and 1,2-epoxybutane (360.50 mg; 5 mmol; 100 equiv) in benzene was irradiated following the same overall methodology described above for the *exo* isomer. A 50% yield was calculated by ¹H NMR using 1,3,5-trimethoxybenzene (99%) as the internal standard. Recrystallization from EtOAc afforded yellow crystals (5.8 mg, 30% yield).

Yellow crystals; R_f = 0.35 (hexane/AcOEt 9:1); ¹H NMR (CDCl₃, 300 MHz): δ = 8.18 (d, J = 8.4 Hz, 2H), 8.07–7.99 (m, 6H), 7.91 (d, J = 8.4 Hz, 2H), 7.06 (d, J = 5.4 Hz, 2H), and 6.80 (d, J = 5.4 Hz, 2H). ¹³C{¹H} NMR (CDCl₃, 75 MHz): δ = 138.2, 136.5, 132.8, 130.6, 128.8, 128.3, 126.8, 125.6, 125.2, 124.1, 123.1, 122.8, and 122.6.

Calculation Details. For the UV–vis spectral simulation, we performed a series of DFT geometry optimizations to understand the conformational equilibria of 7 and 8, followed by TDDFT on the major conformers in order to find the most appropriate functional for this job. Our experimental UV–vis spectra were used as a benchmark in all cases for spectral fitting. In general, all the methods tested provided comparable results and a reasonable fit, indicating that this is not an intrinsically difficult spectral simulation. The double hybrid functional with a dispersion correction term wB97X-D,²⁸ in *n*-hexane as the PCM, was finally chosen for both DFT and TDDFT calculations. People's split-valence quasi triple- ζ , in the valence shell basis set and addition of both polarization and diffuse functions, 6-311++G(d,p) was used in the ground state for geometry optimization and vibrational analysis, while a more expanded basis set, 6-311++G(2df,2pd), was used for UV–vis spectral calculations.²⁹ These calculations were performed using Gaussian09 suite of programs.³⁰ For the excited states, optimization and vibrational analysis were carried out with Gaussian16,³¹ which provides the tools to perform analytical vibrational analysis to TDDFT-optimized excited states.

■ ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.joc.1c00147>.

^1H (1–8) and $^{13}\text{C}\{^1\text{H}\}$ NMR (1–6) spectra, melting point corrections applied and UV–vis spectra (7,8) of these compounds, calculation details on the simulation of the UV–vis spectra, conformational analysis, NTOs of the relevant UV–vis bands, energies of the optimized excited states, and NTO analysis (PDF)

AUTHOR INFORMATION

Corresponding Author

Albert Guijarro – *Departamento de Química Orgánica and Instituto Universitario de Síntesis Orgánica, Campus de San Vicente del Raspeig, Universidad de Alicante, 03080 Alicante, Spain*; orcid.org/0000-0001-9196-8436; Email: aguijarro@ua.es

Authors

Bianca C. Baciú – *Departamento de Química Orgánica and Instituto Universitario de Síntesis Orgánica, Campus de San Vicente del Raspeig, Universidad de Alicante, 03080 Alicante, Spain*

José Antonio Vergés – *Departamento de Teoría y Simulación de Materiales, Instituto de Ciencia de Materiales de Madrid (CSIC), 28049 Madrid, Spain*; orcid.org/0000-0002-2937-4802

Complete contact information is available at: <https://pubs.acs.org/10.1021/acs.joc.1c00147>

Notes

The authors declare no competing financial interest.

ACKNOWLEDGMENTS

Financial support by the Spanish Ministerio de Economía y Competitividad (MAT2016-78625-C2-2-P), the Ministerio de Ciencia, Innovación y Universidades (PID2019-109539GB-C4 and PGC2018-096955-B-C44), the Generalitat Valenciana (PROMETEO/2017/139), and finally, the University of Alicante (VIGROB-285) is gratefully acknowledged. The computational resources provided by the Department of Applied Physics of the University of Alicante are greatly appreciated.

REFERENCES

- Chen, C.-F.; Shen, Y. *Helicene Chemistry. From Synthesis to Applications*; Springer-Verlag: Berlin, Heidelberg, Germany, 2017, pp. 231–245. ISBN 978-3-662-53168-6.
- Roncali, J. Synthetic Principles for Bandgap Control in Linear π -Conjugated Systems. *Chem. Rev.* **1997**, *97*, 173–206.
- Tagami, K.; Tsukada, M.; Wada, Y.; Iwasaki, T.; Nishide, H. Electronic transport of benzothiophene-based chiral molecular solenoids studied by theoretical simulations. *J. Chem. Phys.* **2003**, *119*, 7491–7497.
- Boscoboinik, J. A.; Kohlmeyer, R. R.; Chen, J.; Tysoe, W. T. Efficient transport of gold atoms with a scanning tunneling microscopy tip and a linker molecule. *Langmuir* **2011**, *27*, 9337–9344.
- Baciú, B. C.; de Ara, T.; Sabater, C.; Untiedt, C.; Guijarro, A. Helical nanostructures for organic electronics: the role of topological sulfur in ad-hoc synthesized dithia[7]helicenes studied in the solid and on a gold surface. *Nanoscale Adv.* **2020**, *2*, 1921–1926.
- (a) Wex, B.; Kaafarani, B. R.; Schroeder, R.; Majewski, L. A.; Burckel, P.; Grell, M.; Neckers, D. C. New organic semiconductors and their device performance as a function of thiophene orientation. *J. Mater. Chem.* **2006**, *16*, 1121–1124. (b) Hu, P.; Ye, J.; Jiang, H. Which isomer is better for charge transport: anti- or syn-? *J. Mater. Chem. C* **2019**, *7*, 5858–5873.
- Zhang, Y.; Ye, G.; Soni, S.; Qiu, X.; Krijger, T. L.; Jonkman, H. T.; Carloti, M.; Sauter, E.; Zharnikov, M.; Chiechi, R. C. Controlling destructive quantum interference in tunneling junctions comprising self-assembled monolayers via bond topology and functional groups. *Chem. Sci.* **2018**, *9*, 4414–4423.
- Wynberg, H.; Groen, M. B.; Schadenberg, H. Synthesis and resolution of some heterohelicenes. *J. Org. Chem.* **1971**, *36*, 2797–2809.
- Licandro, E.; Cauteruccio, S.; Dova, D. Thiahelicenes: from basic knowledge to applications. *Adv. Heterocycl. Chem.* **2016**, *118*, 1–46.
- Mallory, F. B.; Mallory, C. W. Photocyclization of stilbenes and related molecules. *Organic Reactions*; John Wiley & Sons, Inc., 1984; Vol. 30, Chapter 1, p 3.
- Bartocci, G.; Galiazzo, G.; Ginocchetti, G.; Mazzucato, U.; Spalletti, A. Effect of thienyl groups on the photoisomerization and rotamerism of symmetric and asymmetric diaryl-ethenes and diaryl-butadienes. *Photochem. Photobiol. Sci.* **2004**, *3*, 870–877.
- Shi, Y.-G.; Møllerup, S. K.; Yuan, K.; Hu, G.-F.; Sauriol, F.; Peng, T.; Wang, N.; Chen, P.; Wang, S. Stabilising fleeting intermediates of stilbene photocyclization with amino-borane functionalisation: the rare isolation of persistent dihydrophenanthrenes and their [1,5] H-shift isomers. *Chem. Sci.* **2018**, *9*, 3844–3855.
- (a) Tedjamulia, M. L.; Stuart, J. G.; Tominaga, Y.; Castle, R. N.; Lee, M. L. The synthesis of naphtho[1,2-b]thiophene and all of the eight isomers of monomethylnaphtho[1,2-b]thiophene. *J. Heterocycl. Chem.* **1984**, *21*, 1215–1219. (b) Song, K.; Wu, L.-Z.; Yang, C.-H.; Tung, C.-H. Photocyclization and photooxidation of 3-styrylthiophene. *Tetrahedron Lett.* **2000**, *41*, 1951–1954.
- Moussa, S.; Aloui, F.; Ben Hassine, B. Synthesis and optoelectronic properties of some new thiahelicenes. *Synth. Commun.* **2011**, *41*, 1006–1016.
- Lambert, C. J. Basic Concepts of Quantum Interference and Electron Transport in Single-Molecule Electronics. *Chem. Soc. Rev.* **2015**, *44*, 875–888.
- Matsushima, T.; Kobayashi, S.; Watanabe, S. Air-driven potassium iodide-mediated oxidative photocyclization of stilbene derivatives. *J. Org. Chem.* **2016**, *81*, 7799–7806.
- Zhao, F.; Luo, J.; Tan, Q.; Liao, Y.; Peng, S.; Deng, G.-J. Sodium sulfinate-mediated trans-stilbene formation from benzylic halides. *Adv. Synth. Catal.* **2012**, *354*, 1914–1918.
- Lee, J.-E.; Kwon, J.; Yun, J. Copper-catalyzed addition of diboron reagents to α,β -acetylenic esters: efficient synthesis of β -Boryl- α,β -Ethylenic Esters. *Chem. Commun.* **2008**, 733–734.
- Symmetry number, in IUPAC. *Compendium of Chemical Terminology*, 2nd ed.; (the “Gold Book”). Compiled by A. D. McNaught and A. Wilkinson; Blackwell Scientific Publications: Oxford, 1997. Online version (2019) created by S. J. Chalk. ISBN 0-9678550-9-8.
- Martin, R. L. Natural transition orbitals. *J. Chem. Phys.* **2003**, *118*, 4775–4777.
- Kasha rule, in IUPAC. *Compendium of Chemical Terminology*, 2nd ed.; (the “Gold Book”). Compiled by A. D. McNaught and A. Wilkinson; Blackwell Scientific Publications: Oxford, 1997. Online version (2019) created by S. J. Chalk. ISBN 0-9678550-9-8.
- Jørgensen, K. B. Photochemical oxidative cyclisation of stilbenes and stilbenoids—The Mallory reaction. *Molecules* **2010**, *15*, 4334–4358.
- Zhao, F.; Luo, J.; Tan, Q.; Liao, Y.; Peng, S.; Deng, G.-J. Sodium Sulfinate-Mediated Trans-Stilbene Formation from Benzylic Halides. *Adv. Synth. Catal.* **2012**, *354*, 1914–1918.
- Talele, H. R.; Chaudhary, A. R.; Patel, P. R.; Bedekar, A. V. Expedient Synthesis of Helicenes using an Improved Protocol of Photocyclodehydrogenation of Stilbenes. *Arkivoc* **2011**, *2011*, 15–37.
- Zeng, X.; Gong, C.; Guo, H.; Xu, H.; Zhang, J.; Xie, J. Efficient heterogeneous hydroboration of alkynes: enhancing the catalytic

activity by Cu(0) incorporated CuFe₂O₄ nanoparticles. *New J. Chem.* **2018**, *42*, 17346–17350.

(26) Molander, G. A.; Brown, A. R. Suzuki–Miyaura Cross-Coupling Reactions of Potassium Vinyltrifluoroborate with Aryl and Heteroaryl Electrophiles. *J. Org. Chem.* **2006**, *71*, 9681–9686.

(27) Limited solubility of large flat aromatic compounds like **7** and **8** prevents recording their ¹³C NMR spectra.

(28) Chai, J.-D.; Head-Gordon, M. Long-range corrected hybrid density functionals with damped atom–atom dispersion corrections. *Phys. Chem. Chem. Phys.* **2008**, *10*, 6615–6620.

(29) Krishnan, R.; Binkley, J. S.; Seeger, R.; Pople, J. A. Self-consistent molecular orbital methods. XX. A basis set for correlated wave functions. *J. Chem. Phys.* **1980**, *72*, 650–654.

(30) Frisch, M. J.; Trucks, G. W.; Schlegel, H. B.; Scuseria, G. E.; Robb, M. A.; Cheeseman, J. R.; Scalmani, G.; Barone, V.; Mennucci, B.; Petersson, G. A.; et al. *Gaussian 09*, Revision C.01; Gaussian, Inc.: Wallingford, CT, 2009.

(31) Frisch, M. J.; Trucks, G. W.; Schlegel, H. B.; Scuseria, G. E.; Robb, M. A.; Cheeseman, J. R.; Scalmani, G.; Barone, V.; Petersson, G. A.; Nakatsuji, H.; et al. *Gaussian 16*, Revision A.03; Gaussian, Inc.: Wallingford, CT, 2016.